

National Inventory Report Sweden 2013

Greenhouse Gas Emission Inventories 1990–2011
– Submitted under the United Nations Framework
Convention on Climate Change and the Kyoto Protocol



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Emissions and removals within the Land Use, Land Use Change
and Forestry sector are significant within the Swedish greenhouse gas inventory.

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Preface

According to Articles 4 and 12 of the United Nations Framework Convention on Climate Change (UNFCCC), parties are required to, on an annual basis, submit national inventories of anthropogenic emissions by sources and removals by sinks of all greenhouse gases not controlled by the Montreal Protocol. This report is also a submission under the Kyoto Protocol.

This is Sweden's National Inventory Report (NIR) for the year 2013 and it is written in line with the guidelines of the Annotated outline of the National Inventory Report including reporting elements under the Kyoto Protocol. It contains national greenhouse gas emission inventories for the period 1990 to 2011, and descriptions of methods used to produce the estimates. The methods used to calculate the emissions and removals are in accordance with the Revised IPCC 1996 Guidelines for National Greenhouse Gas Inventories and the IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories. The report is prepared in accordance with the Reporting Guidelines, agreed by the UNFCCC at the eighth session of the Conference of the Parties (COP) in New Delhi 2002 and subsequent decisions.

This inventory is coordinated, on behalf of the Swedish Ministry of Environment, by the Swedish Environmental Protection Agency.

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Contents

PREFACE	3
AUTHORS	4
SAMMANFATTNING	13
(Summary in Swedish)	13
S.1 Bakgrund	13
S.2 Sammanfattning av nationella utsläpp och upptag samt trender, inklusive KP-LULUCF	13
S.2.1 Växthusgaser	13
S.2.2 KP-LULUCF	14
S.3 Översikt över utsläppsberäkningar och trender sektorsvis, inklusive KP-LULUCF	16
S.3.1 Växthusgaser	16
S.3.2 KP-LULUCF	18
S.4 Översikt av utsläppsberäkningar och trender för indirekta växthusgaser och SO ₂	19
EXECUTIVE SUMMARY	20
ES.1 Background information	20
ES.2 Summary of National Emissions and Removal Related Trends, including KP-LULUCF	20
ES.2.1 GHG inventory	20
ES.2.2 KP-LULUCF activities	21
ES.3 Overview of Source and Sink Category Emission Estimates and Trends, including KP-LULUCF	23
ES.3.1 GHG inventory	23
ES.3.2 KP-LULUCF activities	25
ES.4 Overview of Emission Estimates and Trends of Indirect GHGs and SO ₂	25
SUMMARY IN ARABIC	27
1 INTRODUCTION	33
1.1 Background Information	33
1.1.1 Climate change	33
1.1.2 Greenhouse gas inventories	35
1.1.3 Supplementary information required under Article 7, paragraph 1, of the Kyoto Protocol	36
1.1.4 Sweden's commitment under the Kyoto Protocol and the EU Burden Sharing Decision	36
1.1.5 National emission targets	37

1.2	Institutional arrangements	38
1.2.1	Legal arrangements	38
1.2.2	Institutional arrangements	39
1.3	Inventory planning, preparation and management	42
1.3.1	Quality system	42
1.3.2	Training, awareness and skills	45
1.3.3	Inventory planning (PLAN)	45
1.3.4	Inventory preparation (DO)	46
1.3.5	QA/QC procedures and extensive review of GHG inventory and KP-LULUCF inventory (CHECK)	48
1.4	Brief general description of methodologies and data sources used	51
1.4.1	GHG inventory	51
1.4.2	KP-LULUCF inventory	53
1.5	Brief description of key categories, including for KP-LULUCF key categories	53
1.5.1	GHG inventory (including and excluding LULUCF)	53
1.5.2	KP-LULUCF inventory	55
1.6	Information on QA/QC	55
1.6.1	QA/QC Procedures	55
1.6.2	Verification activities	55
1.6.3	Treatment of confidentiality issues	55
1.7	General uncertainty evaluation	55
1.7.1	GHG inventory	55
1.7.2	KP-LULUCF activities	58
1.7.3	General assessment of completeness	59
2	TRENDS IN GREENHOUSE GAS EMISSIONS	61
2.1	Description and interpretation of emission trends for aggregated greenhouse gas emissions	61
2.1.1	Overview of emissions trends per sector	62
2.2	Description and interpretation of emission trends by gas	62
2.2.1	Carbon dioxide	63
2.2.2	Methane	64
2.2.3	Nitrous oxide	64
2.2.4	Fluorinated greenhouse gases	65
2.3	Description and interpretation of emissions by category	65
2.3.1	Energy	66
2.3.2	Industrial processes	78
2.3.3	Solvents and other products use	81
2.3.4	Agriculture	82
2.3.5	Land Use, Land Use Change and Forestry – LULUCF	84
2.3.6	Waste	86
2.3.7	International bunkers	87
2.4	Description and interpretation of emission trends for indirect greenhouse gases and SO ₂	88
2.4.1	NMVOC	88

2.4.2	NO _x	89
2.4.3	CO	90
2.4.4	SO ₂	91
2.4.5	Description and interpretation of emission trends for KP-LULUCF inventory in aggregate and by activity, and by gas	92
3	ENERGY (CRF SECTOR 1)	93
3.1	Overview of sector	93
3.2	Fuel combustion (CRF 1.A)	95
3.2.1	Comparison of the sectoral approach with the reference approach	96
3.2.2	International bunker fuels	97
3.2.3	Feedstocks and non-energy use of fuels	99
3.2.4	CO ₂ capture from flue gases and subsequent CO ₂ storage	99
3.2.5	Country-specific issues	99
3.2.6	Public electricity and heat production (CRF 1.A.1.a)	100
3.2.7	Petroleum refining (CRF 1.A.1.b)	104
3.2.8	Manufacture of solid fuels and other energy industries (CRF 1.A.1.c)	106
3.2.9	Iron and steel (CRF 1.A.2.a)	108
3.2.10	Non-Ferrous Metals (CRF 1.A.2.b)	111
3.2.11	Chemicals (CRF 1.A.2.c)	112
3.2.12	Pulp, Paper and Print (CRF 1.A.2.d)	115
3.2.13	Food Processing, Beverages and Tobacco (CRF 1.A.2.e)	117
3.2.14	Other Industries (CRF 1.A.2.f)	118
3.2.15	Civil Aviation (CRF 1.A.3.a)	122
3.2.16	Road transport (CRF 1.A.3.b)	125
3.2.17	Railways (CRF 1.A.3.c)	129
3.2.18	Navigation (CRF 1.A.3.d)	130
3.2.19	Other transportation (CRF 1.A.3.e)	134
3.2.20	Commercial/institutional (CRF 1.A.4.a)	136
3.2.21	Residential (CRF 1.A.4.b)	139
3.2.22	Agriculture/forestry/fisheries (CRF 1.A.4.c)	141
3.2.23	Other stationary (CRF 1.A.5.a)	143
3.2.24	Other mobile (CRF 1.A.5.b)	143
3.3	Fugitive emissions from solid fuels and oil and natural gas (CRF 1.B)	145
3.3.1	Fugitive emissions from solid fuels (CRF 1.B.1)	145
3.3.2	Oil and natural gas (CRF 1.B.2)	146
4	INDUSTRIAL PROCESSES (CRF SECTOR 2)	156
4.1	Overview of sector	156
4.2	Mineral products (CRF 2.A)	158
4.2.1	Cement production (CRF 2.A.1)	158
4.2.2	Lime production (CRF 2.A.2)	162
4.2.3	Limestone and dolomite use (CRF 2.A.3)	167

4.2.4	Soda ash use (CRF 2.A.4)	171
4.2.5	Asphalt roofing (CRF 2.A.5)	172
4.2.6	Road paving with asphalt (CRF 2.A.6)	174
4.2.7	Other (CRF 2.A.7)	175
4.3	Chemical industry (CRF 2.B)	179
4.3.1	Ammonia production (CRF 2.B.1)	179
4.3.2	Nitric acid production (CRF 2.B.2)	180
4.3.3	Carbide production (CRF 2.B.4)	182
4.3.4	Other (CRF 2.B.5)	186
4.4	Metal production (CRF 2.C)	188
4.4.1	Iron and steel production (CRF 2.C.1)	188
4.4.2	Ferroalloy production (CRF 2.C.2)	197
4.4.3	Aluminium production (CRF 2.C.3)	200
4.4.4	SF ₆ used in aluminium and magnesium foundries (CRF 2.C.4)	204
4.4.5	Other metal production (CRF 2.C.5)	204
4.5	Other production (CRF 2.D)	206
4.5.1	Pulp and paper (CRF 2.D.1)	206
4.5.2	Food and drink (CRF 2.D.2)	207
4.6	Production of Halocarbons and SF ₆ (CRF 2.E)	210
4.7	Consumption of Halocarbons and SF ₆ (CRF 2.F)	210
4.7.1	Refrigeration and air conditioning equipment (2.F.1)	211
4.7.2	Foam blowing (2.F.2)	214
4.7.3	Fire extinguishers (2.F.3)	216
4.7.4	Aerosols/metered dose inhalers (2.F.4)	217
4.7.5	Solvents (2.F.5)	219
4.7.6	Other applications using ODS substitutes (2.F.6)	219
4.7.7	Semiconductor manufacture (2.F.7)	219
4.7.8	Electrical equipment (2.F.8)	220
4.7.9	Other (2.F.9)	222
4.8	Consumption of Halocarbons and SF ₆ Potential Emissions (CRF 2.F.P)	224
4.8.1	Potential emissions	224
4.9	Other, CRF 2G	224
5	SOLVENT AND OTHER PRODUCT USE (CRF SECTOR 3)	225
5.1	Overview of sector	225
5.2	Paint application (CRF 3.A)	227
5.2.1	Source category description	227
5.2.2	Methodological issues	228
5.2.3	Uncertainties and time-series consistency	228
5.2.4	Source-specific QA/QC and verification	228
5.2.5	Source-specific recalculations	228
5.2.6	Source-specific planned improvements	228

5.3	Degreasing and Dry cleaning (CRF 3.B)	229
5.3.1	Source category description	229
5.3.2	Methodological issues	229
5.3.3	Uncertainties and time-series consistency	229
5.3.4	Source-specific QA/QC and verification	229
5.3.5	Source-specific recalculations	230
5.3.6	Source-specific planned improvements	230
5.4	Chemical products, Manufacture and Processing (CRF 3.C)	230
5.4.1	Source category description	230
5.4.2	Methodological issues	230
5.4.3	Uncertainties and time-series consistency	231
5.4.4	Source-specific QA/QC and verification	231
5.4.5	Source-specific recalculations	231
5.4.6	Source-specific planned improvements	231
5.5	Other (CRF 3.D)	231
5.5.1	Source category description	231
5.5.2	Methodological issues	232
5.5.3	Uncertainties and time-series consistency	232
5.5.4	Source-specific QA/QC and verification	233
5.5.5	Source-specific recalculations	233
5.5.6	Source-specific planned improvements	233
6	AGRICULTURE (CRF SECTOR 4)	234
6.1	Overview of sector	234
6.2	Enteric Fermentation (CRF 4.A)	237
6.2.1	Source category description	237
6.2.2	Methodological issues	237
6.2.3	Uncertainties and time-series consistency	241
6.2.4	Source-specific QA/QC and verification	241
6.2.5	Source-specific recalculations	241
6.2.6	Source-specific planned improvements	241
6.3	Manure Management (CRF 4.B)	243
6.3.1	Source category description	243
6.3.2	Methodological issues	243
6.3.3	Uncertainties and time-series consistency	249
6.3.4	Source-specific QA/QC and verification	249
6.3.5	Source-specific recalculations	249
6.3.6	Source-specific planned improvements	249
6.4	Agricultural Soils (CRF 4.D)	249
6.4.1	Direct Soil Emissions (CRF 4.D.1)	250
6.4.2	Pasture, Range and Paddock Manure (CRF 4.D.2)	265
6.4.3	Indirect Emissions (CRF 4.D.3)	266
6.4.4	Other (CRF 4.D.4)	269

7	LAND USE, LAND-USE CHANGE AND FORESTRY (CRF SECTOR 5)	271
7.1	Overview of LULUCF	271
7.2	Description of categories 5A, 5B, 5C, 5D, 5E and 5F	278
7.2.1	Characteristics of categories	278
7.2.2	Information on approaches used for representing land areas and on land-use databases used for the inventory preparation	278
7.2.3	Land-use definitions and the classification systems used and their correspondence to the LULUCF categories	285
7.2.4	Definition of carbon Pools, CRF 5A, 5B, 5C, 5D, 5E and 5F	288
7.2.5	Emissions of N ₂ O, CO ₂ and CH ₄ , CRF 5(I), 5(II), 5(III), 5(IV) and 5(V)	289
7.3	Methodological issues	291
7.3.1	CRF-tables 5A, 5B, 5C, 5D, 5E and 5F	291
7.3.2	CRF 5(I), 5(II), 5(III), 5(IV) and 5(V)	294
7.4	Uncertainties and time series consistency	295
7.4.1	Uncertainties	295
7.4.2	Living biomass, CRF 5A, 5B, 5C, 5D, 5E and 5F	296
7.4.3	Dead organic matter, CRF 5A, 5B, 5C, 5D, 5E and 5F	297
7.4.4	Soil organic carbon, CRF 5A, 5B, 5C, 5D, 5E and 5F	298
7.4.5	Other CO ₂ emissions, CRF 5(IV) and 5(V)	298
7.4.6	N ₂ O and CH ₄ emissions, CRF 5(I), 5(III) and 5(V)	299
7.4.7	Completeness	299
7.4.8	Time series consistency and verification	299
7.5	QA/QC	300
7.5.1	Quality assurance	300
7.5.2	Quality control	300
7.6	Source-specific Recalculations	301
7.7	Coming improvements	303
7.7.1	Informal reporting of HWP	303
8	WASTE (CRF SECTOR 6)	305
8.1	Overview of sector	305
8.2	Solid waste disposal on land (CRF 6.A)	307
8.2.1	Managed waste disposal on land (CRF 6.A.1)	309
8.3	Waste water handling (CRF 6.B)	326
8.3.1	Industrial, domestic and commercial wastewater (CRF 6.B.1 and CRF 6.B.2)	326
8.4	Waste incineration (CRF 6.C)	335
8.4.1	Source category description	335
8.4.2	Methodological issues	335
8.4.3	Uncertainties and time-series consistency	337
8.4.4	Source-specific QA/QC and verification	337
8.4.5	Source-specific recalculations	338
8.4.6	Source-specific planned improvements	338

9	OTHER	339
10	RECALCULATIONS AND IMPROVEMENTS	340
10.1	Explanations and justifications for recalculations	340
10.2	Implications for emission levels	340
10.2.1	Energy, CRF 1	340
10.2.2	Industrial processes, CRF 2	341
10.2.3	Solvents and other products use, CRF 3	342
10.2.4	Agriculture, CRF 4	342
10.2.5	LULUCF, CRF 5	342
10.2.6	Waste, CRF 6	343
10.3	Implications for emission trends	345
10.4	Recalculations and other changes made in response to the UNFCCC review process	346
10.5	Major changes in methodological descriptions	366
11	KP-LULUCF	369
11.1	General information	369
11.1.1	Definitions of forest and any other criteria	372
11.1.2	Elected activities under Article 3, paragraph 4, of the Kyoto Protocol	373
11.1.3	Description of how the definitions of each activity under Article 3.3 and each elected activity under Article 3.4 have been implemented and applied consistently over time	373
11.1.4	Descriptions of precedence conditions and/or hierarchy among Article 3.4 activities, and how they have been consistently applied in determining how land was classified.	375
11.2	Land-related information	375
11.2.1	Spatial assessment unit used for determining the area of the units of land under Article 3.3	375
11.2.2	Methodology used to develop the land use matrix	375
11.2.3	Maps and/or database to identify the geographical locations, and the system of identification codes for the geographical locations	376
11.3	Activity-specific information	377
11.3.1	Methods for carbon stock change and GHG emission and removal estimates	377
11.4	Article 3.3	383
11.4.1	Information that demonstrates that activities under Article 3.3 began on or after 1 January 1990 and before 31 December 2012 and are direct human-induced	383
11.4.2	Information on how harvesting or forest disturbance that is followed by the re-establishment of forest is distinguished from deforestation	383

11.4.3	Information on the size and geographical location of forest areas that have lost forest cover but which are not yet classified as deforested	384
11.5	Article 3.4	384
11.5.1	Information that demonstrates that activities under Article 3.4 have occurred since 1 January 1990 and are human induced	384
11.5.2	Information relating to Cropland Management, Grazing Land Management, and Revegetation, if elected, for the base year	384
11.5.3	Information relating to Forest Management	385
11.6	Other information	385
11.6.1	Key category analysis for Article 3.3 activities and any elected activities under Article 3.4	385
11.7	Information relating to Article 6	385
12	INFORMATION ON ACCOUNTING OF KYOTO UNITS	386
12.1	Background information	386
12.2	Summary of information reported in the SEF tables	388
12.3	Discrepancies and notifications	388
12.4	Publicly accessible information	389
12.5	Calculation of the commitment period reserve (CPR)	390
12.5.1	Assigned Amount	390
12.5.2	Commitment Period Reserve (CPR)	390
12.6	KP-LULUCF accounting	391
13	INFORMATION ON CHANGES IN NATIONAL SYSTEM	392
14	INFORMATION ON CHANGES IN NATIONAL REGISTRY	393
15	INFORMATION ON MINIMIZATION OF ADVERSE IMPACTS IN ACCORDANCE WITH ARTICLE 3, PARAGRAPH 14	397
16	OTHER INFORMATION	401
16.1	References	401
16.2	Units and Abbreviations	422

Sammanfattning

(Summary in Swedish)

S.1 Bakgrund

Växthusgaser har alltid funnits i atmosfären, men på grund av mänsklig aktivitet har koncentrationen av många av dem ökat, vilket intensifierar växthuseffekten. 1988 bildades Intergovernmental Panel on Climate Change (IPCC) och två år senare konstaterade de att antropogen klimatpåverkan utgjorde ett globalt hot och efterfrågade en internationell överenskommelse för att hantera problemet. FN:s generalförsamling inledde förhandlingar om en ramkonvention kring klimatförändringar (UNFCCC), vilken trädde i kraft 1994. Det långsiktiga målet är att stabilisera halterna av växthusgaser i atmosfären på en nivå som förhindrar skadliga antropogena klimatförändringar från att äga rum. Det viktigaste tillägget till konventionen förhandlades fram i Kyoto, Japan, 1997. Kyotoprotokollet innebär bindande åtaganden gällande utsläppsmängder för Annex I-parterna, vilket innebär att dessa länders utsläpp av växthusgaser under åren 2008-2012 i medeltal ska vara minst 5 % lägre än under basåret 1990.

Enligt Artikel 4 och 12 i UNFCCC måste Annex I-parterna årligen rapportera sina utsläpp från källor och upptag i sänkor för alla växthusgaser som inte omfattas/kontrolleras av Montrealprotokollet. Rapporteringen ska innefatta utsläppssiffror i ett speciellt format (CRF) och en nationell inventeringsrapport (NIR).

Denna rapport utgör Sveriges NIR 2013. Rapporten omfattar utsläpp till luft av de direkta växthusgaserna CO₂, CH₄, N₂O, HFC, PFC, SF₆ och de indirekta växthusgaserna NO_x, CO, NMVOC och SO₂. Rapporten innehåller information om Sveriges inventering av växthusgaser för alla år från 1990 till 2011, inklusive beskrivningar av metoder, datakällor, osäkerheter, den kvalitets-säkring och kvalitetsstyrning (QA/QC) som görs och en trendanalys.

De data, såsom emissioner, aktivitetsdata och emissionsfaktorer som UNFCCC efterfrågar återfinns i CRF-tabeller tillsammans med denna rapport.

S.2 Sammanfattning av nationella utsläpp och upptag samt trender, inklusive KP-LULUCF

S.2.1 Växthusgaser

Totala utsläpp av växthusgaser i Sverige exklusive LULUCF, uttryckt i kol-dioxidekvivalenter, var ca 61,4 miljoner ton år 2011 (Tabell S 1), vilket är en minskning med ca 4 miljoner ton jämfört med 2010. Osäkerheten i skattingen är ca 4,5 %. Utsläppen har minskat med ca 16 %, eller ca 11 miljoner ton, mellan 1990 och 2011. Osäkerheten i trenden är $\pm 2,0$ %, dvs. med 95 % sannolikhet är den faktiska minskningen mellan 14 % - 18 %.

Nettopptaget för sektorn Markanvändning, Förändrad markanvändning och Skogsbruk (LULUCF) har beräknats till ca 35 miljoner ton koldioxidekvivalenter 2011 (Tabell S 1). Den årliga förändringen i sänkan är mycket liten i förhållande till poolernas storlekar (levande biomassa, dött organiskt material och markkol). Beräkningarna är på grund av detta behäftade med stora osäkerheter. Osäkerheterna tillsammans med löpande metodutveckling kan ge stora förändringar i rapporterade värden mellan submissionerna.

Utsläppen av koldioxid var ca 48,7 miljoner ton år 2011 vilket är drygt 14 % lägre jämfört med 1990 (Tabell S 1). Energisektorn, inklusive transporter, står för ca 88 % av de totala koldioxidutsläppen och är därmed den största källan till koldioxidutsläpp i Sverige. Koldioxid står för ca 79 % av de totala utsläppen av växthusgaser.

Metanutsläpp (CH_4) kommer framför allt från jordbruk och avfallsdeponier och var knappt 5 miljoner ton 2011 räknat som koldioxidekvivalenter (Tabell S 1). Sedan 1990 har utsläppen av metan minskat med ca 28 %, vilket främst beror på åtgärder inom avfallssektorn och jordbrukssektorn.

2011 var de totala utsläppen av lustgas (N_2O) ca 6,7 miljoner ton räknat som koldioxidekvivalenter (Tabell S 1), vilket är en minskning med ca 20 % jämfört med 1990. Utsläpp av lustgas kommer huvudsakligen från jordbrukssektorn, men också från energiproduktion, industriprocesser och hantering av avloppsvatten. Jordbrukssektorn står för den största delen av minskningen.

Totala utsläppen av fluorerade gaser (PFCs, HFCs och SF_6) 2011 var ca 1 miljon ton uttryckt i koldioxidekvivalenter (Tabell S 1). Detta innebär en ökning av utsläppen med 116 % jämfört med 1990. Ökningen beror främst på att ozonförstörande ämnen ersatts av HFCs.

S.2.2 KP-LULUCF

Sverige rapporterar artikel 3.3 och 3.4 under Kyoto protokollet. För dessa aktiviteter har Sverige valt att bokföra upptag/utsläpp för hela åtagandeperioden (och ej på årsbasis).

Aktiviteter under artikel 3.3, nybeskogning /återbeskogning (AR), är relativt ovanliga i Sverige. Sedan 2008 har arealen nybeskogad/återbeskogad mark ökat med ca 5 000 ha per år. Upptaget för nybeskogning /återbeskogning (AR) har beräknats till 1,66 miljoner ton CO_2 ekvivalenter 2011. Andelen mark som avskogats (D), har ökat något från 2008. Det totala utsläppet 2011 har beräknats till 2,6 miljoner ton CO_2 ekvivalenter. Under artikel 3.4 har Sverige valt bokföring av skogsbruk. För denna aktivitet är upptaget ca 37,6 miljoner ton CO_2 ekvivalenter 2011. Rapporteringen under Kyotoprotokollet artikel 3.3 och 3.4 harmonierar med UNFCCC rapporteringen för skogsmark och mark som konverterats till skogsmark (arean).

Tabell S.1. Utsläpp av växthusgaser ämnesvis (Gg CO₂ ekvivalenter).

UTSLÄPP AV VÄXTHUSGASER	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
CO ₂ inkl. netto CO ₂ från LULUCF	19 688	27 232	18 529	19 332	20 247	23 938	26 273	26 044	18 750	20 609	17 044	13 394	21 464	13 377
CO ₂ exkl. netto CO ₂ från LULUCF	56 954	58 872	54 145	55 078	55 993	56 608	55 768	53 231	53 194	51 972	50 006	46 405	52 302	48 726
CH ₄ inkl. CH ₄ från LULUCF	6 940	6 829	6 255	6 214	6 033	5 888	5 910	5 780	5 711	5 474	5 283	5 172	5 077	4 987
CH ₄ exkl. CH ₄ från LULUCF	6 938	6 827	6 252	6 211	6 028	5 882	5 904	5 775	5 699	5 471	5 270	5 170	5 076	4 985
N ₂ O inkl. N ₂ O från LULUCF	8 449	8 132	7 675	7 493	7 385	7 347	7 314	7 165	7 191	6 930	7 078	6 895	7 170	6 797
N ₂ O exkl. N ₂ O från LULUCF	8 370	8 070	7 602	7 420	7 314	7 270	7 232	7 073	7 096	6 825	6 956	6 779	7 033	6 682
HFCs	4	132	568	615	666	710	769	789	818	838	867	869	845	813
PFCs	377	343	241	236	261	258	254	257	245	248	225	35	158	183
SF ₆	107	127	94	111	104	69	81	142	111	151	84	81	73	60
Total (inkl. LULUCF)	35 566	42 795	33 360	34 001	34 696	38 210	40 601	40 178	32 827	34 250	30 581	26 446	34 787	26 217
Total (exkl. LULUCF)	72 750	74 371	68 902	69 670	70 366	70 797	70 009	67 268	67 164	65 506	63 407	59 338	65 487	61 449

Tabell S.2. Utsläpp av växthusgaser sektorsvis (Gg CO₂ ekvivalenter).

GREENHOUSE GAS SOURCE AND SINK CATEGORIES CO ₂ -eq (Gg)	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
1. Energy	53 670	55 464	50 584	51 460	52 242	53 179	51 960	49 604	49 608	48 249	46 401	44 508	48 804	45 015
2. Industrial Processes	6 330	6 644	6 812	6 810	6 951	6 679	7 071	6 976	6 969	6 922	6 804	4 986	6 810	6 661
3. Solvent and Other Product Use	332	309	278	269	276	292	311	303	299	281	288	270	289	289
4. Agriculture	8 997	8 722	8 313	8 260	8 171	8 060	8 094	7 954	7 932	7 856	7 915	7 683	7 786	7 772
5. Land Use, Land-Use Change and Forestry	-37 184	-31 576	-35 541	-35 669	-35 670	-32 587	-29 408	-27 091	-34 337	-31 255	-32 826	-32 891	-30 701	-35 232
6. Waste	3 421	3 233	2 916	2 871	2 726	2 586	2 572	2 431	2 356	2 197	2 000	1 892	1 798	1 713
7. Other	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO
Total (including LULUCF)(5)	35 566	42 795	33 360	34 001	34 696	38 210	40 601	40 178	32 827	34 250	30 581	26 446	34 787	26 217

S.3 Översikt över utsläppsberäkningar och trender sektorsvis, inklusive KP-LULUCF

S.3.1 Växthusgaser

De metoder som använts för att beräkna utsläpp och upptag överensstämmer med 'Revised IPCC 1996 Guidelines for National Greenhouse Gas Inventories' och 'IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories'. Inventeringen och rapporten är framtagen i enlighet med 'Reporting Guidelines', som beslutades av UNFCCC under den åttonde sammankomsten av Conference of the Parties (COP) i New Delhi 2002 och efterföljande beslut.

De sektorer som innefattas av inventeringen och de källor som används för aktivitetsdata och/eller utsläppsdata presenteras i Tabell S 3. Utsläppen är hämtade direkt från dessa datakällor eller beräknade baserat på aktivitetsdata.

Tabell S.3. CRF sektorer och datakällor som används i inventeringen.

CRF	Sektor	Primär källa till aktivitetsdata/utsläppsdata
1	Energi	
	-Stationär förbränning	Statistiska undersökningar av energiförbrukning
	-Transport	Transportmyndigheter
2	Industriprocesser	Miljörapporter
		Statistiska undersökningar av energiförbrukning
		Direktkontakt med företag
		EU:s utsläppshandelssystem
3	Lösningsmedel och annan Produktanvändning	Nationella data från Produktregistret på Kemikalieinspektionen
		Nationella experter
		Miljörapporter
4	Jordbruk	Officiella statistiska rapporter
		Organisationer och Forskare
5	Förändrad Markanvändning och Skogsbruk	Sveriges lantbruksuniversitet
		Skogsstyrelsen
6	Avfall	Avfall Sverige (fd RVF)
		Skogsindustrierna
		SCB
		Naturvårdsverket
		Miljörapporter

Utsläppen från energisektorn varierar på grund av temperatur- och nederbördsförhållanden samt det ekonomiska konjunkturläget men trenden för perioden 1990-2011 är minskande utsläpp. Utsläppen av växthusgaser från energisektorn inklusive transporter var ca 45 miljoner ton i koldioxidekvivalenter 2011 (Tabell S 2), vilket är ca 73 % av de totala utsläppen. Utsläpp av växthusgaser från vägtrafiken ökade med ca 6 % från 1990 till 2011. Utsläppen av växthusgaser har under 2011 minskat jämfört med 2010.

Det är framförallt utsläppen från el och värmeproduktion och transporter som minskat mellan 2010 och 2011.

Utsläppen från produktion av el och fjärrvärme år 2011 har minskat med ca 22 % (till ca 8,3 miljoner ton) jämfört med 2010 (ca 10,6 miljoner ton). Några bakomliggande orsaker är minskat behov av uppvärmning på grund av varmare väder, god tillgång på vattenkraft och en något ökad kärnkraftsproduktion.

Utsläppen av växthusgaser från inrikes transporter minskade med ca 2 % (till ca 20 miljoner ton) under 2011 jämfört med 2010. Ökad andel förnybara bränslen och bränseleffektivare fordon bidrar till de minskande utsläppen.

För industriprocesser är koldioxid den dominerande växthusgasen med 82 %, sedan kommer fluorerade växthusgaser med ca 16 % dikväveoxid med ca 2 % och metan med 0,2 %. Utsläppen kommer framför allt från produktion av järn och stål samt mineralindustrin. De totala utsläppen från industriprocesser var omkring 6,7 miljoner ton koldioxidekvivalenter år 2011 (Tabell S 2), vilket motsvarar ca 11 % av Sveriges totala utsläpp. Utsläppen från denna sektor minskade med ca 0,15 miljon ton koldioxidekvivalenter eller ca 2 % mellan 2010 och 2011. Denna minskning är inte relaterad till någon nedgång i produktionsvolymen, utan beror på ombyggnad av två anläggningar som producerar salpetersyra, vilket ledde till en betydande minskad av N₂O (ca 87 % jämfört med 2010). Sedan 1990 har utsläppen från industriprocesser varierat, vilket framför allt beror på att produktionsvolymerna varierar den ekonomiska konjunkturen. 2011 var utsläppen 5 % högre än 1990.

Användning av Lösningsmedel och andra produkter ger huvudsakligen upphov till utsläpp av flyktiga organiska ämnen (MNVOCs), lustgas (N₂O) och en del koldioxid. 2011 var utsläppen av koldioxid och lustgas drygt 0,3 miljoner ton uttryckt i koldioxidekvivalenter (Tabell S 2), vilket utgör knappt 0,5 % av de totala växthusgasutsläppen. Jämfört med 1990 har utsläppen i denna sektor minskat med ca 13 %. Ca 9 % av koldioxidutsläppen kommer från användningen av färg, även om dessa utsläpp har minskat p.g.a. en övergång till vattenbaserade färger.

Jordbruk är den största källan till utsläpp av lustgas och metan. 2011 var de totala utsläppen från jordbrukssektorn ca 7,8 miljoner ton uttryckt i koldioxid-ekvivalenter (Tabell S 2) varav ca 63 % utgjordes av N₂O och ca 37 % av CH₄. Det är en minskning med ca 14 % jämfört med 1990. Utsläpp av metan kommer framför allt från boskapens matsmältningsprocesser och gödselhantering. De viktigaste anledningar till de minskade utsläppen är en minskad boskaphållning och en minskad användning av mineralgödsel-medel i jordbruket. Lustgas kommer framför allt från omvandling av kväve i jorden, vilken påverkas av användning av gödsel och handelsgödsel och odling av kvävefixerande växter. Mer än hälften (57 %) av sektorns utsläpp kommer från jordbruksmark.

Nettoupptaget för LULUCF under 2011 uppskattas till omkring 35 CO₂ miljon ton. Nettoupptaget ökade något mellan 2010 till 2011 med ca 4,5 CO₂ miljon ton (Tabell S 2). Den långsiktiga trenden med minskande nettoupptag i levande biomassa beror främst på ökad avverkning, medan ökningen i bruttoupptag härrör från trädens tillväxt. Om tillväxten är mindre än avverkningen minskar sänkan.

Avfallsdeponier är den näst största källan till utsläpp av metan. Av avfallssektorns utsläpp under 2011 dominerar metanutsläppen från avfallsdeponier med ca 87 % medan lustgas från avloppsvatten står för 9 % och koldioxidutsläppen från förbränning av farligt avfall för ca 3 %. 2011 var de totala utsläppen från avfallssektorn drygt 1,7 miljoner ton (Tabell S 2) uttryckt i koldioxidekvivalenter, vilket motsvarar 2,8 % av de totala växthusgasutsläppen. Utsläppen från sektorn har minskat med 50 % jämfört med 1990. Utvinning av deponigas, deponiförbud och deponiskatter är huvudorsakerna till utsläppsminskningen.

De totala växthusgasutsläppen i Sverige uppgick 2011 till ca 61,4 miljoner ton koldioxidekvivalenter varav utsläpp icke inkluderade i utsläppshandel stod för 41,6 miljoner ton. Detta innebär att de svenska utsläppen år 2011 var 16 % lägre än för basåret.

S.3.2 KP-LULUCF

I och med att aktiviteterna under artikel 3.3 (beskogning och avskogning) är relativt ovanliga är det svårt att uttala sig om skillnader mellan år. Netto-upptag respektive emissioner ligger på ungefär samma nivåer som i tidigare submission.

När det gäller aktiviteten skogsbruk under artikel 3.4 så har nettoinlagringen efter att den minskat lite de sista åren åter uppnått ungefär samma nivå som 1990, cirka 28 miljoner ton koldioxidekvivalenter.

S.4 Översikt av utsläppsberäkningar och trender för indirekta växthusgaser och SO₂

Utsläppen av kväveoxider (NO_x) var drygt 146 kton 2011 (Tabell S 4), vilket är en minskning med ca 46 % jämfört med 1990. Vägtrafikens utsläpp av NO_x har minskat med 57 % mellan 1990 och 2011. Utsläppen har minskat med ca 7 % mellan 2010 och 2011. De största källorna till utsläpp av kväveoxider är vägtrafik, arbetsmaskiner, sjöfart och el- och värmeproduktion. I tätorter är vägtrafiken den största källan till kväveoxidutsläpp, men införandet av katalysatorer i bilar och den påföljande successivt mer skärpta avgasstandarderna har bidragit till en generell minskning av kväveoxidnivåer i tätbebyggda områden. Den ökande användningen av fjärrvärme och NO_x-avgiften i början på 1990-talet har också resulterat i stora minskningar av kväveoxidutsläpp från energisektorn.

Tabell S.4. Utsläpp av indirekta växthusgaser och SO₂ (kton).

Gas	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
NO _x	271	247	211	201	193	191	184	180	176	170	163	152	154	146
CO	1 280	1 127	826	785	739	725	678	669	629	617	607	606	587	571
NM VOC	359	278	224	213	205	206	202	199	195	193	189	184	183	177
SO ₂	105	69	42	41	41	41	37	36	36	32	30	29	32	30

Utsläppen av kolmonoxid (CO) har minskat från 1 280 kton 1990 till 571 kton 2011 (Tabell S 4), en reduktion på 55 %. Omkring 96 % av utsläppen kommer från energisektorn varav ca 39 % kommer från transport.

Utsläpp av flyktiga organiska ämnen (NMVOC) var ca 177 kton 2011 (Tabell S 4), vilket är en minskning med 49 % jämfört med 1990. De huvudsakliga källorna till NMVOC är vägtrafik, vedeldning inom bostadssektorn och produkter innehållande lösningsmedel. Frivilliga miljöstandarder för nya installationer av vedeldningspannor och minskade utsläpp från produkter med lösningsmedel har bidragit till utsläppsminskningen.

Utsläppen av svaveldioxid (SO₂) har minskat från 105 kton 1990 till knappt 30 kton 2011 (Tabell S 4), en reduktion på ca 72 %. Minskningen beror framför allt på övergång till lågsvavelhaltiga bränslen, för såväl vägtrafik som uppvärmning. Svavelskatt, som infördes 1991, spelar en stor roll för utvecklingen. Svaveldioxidutsläpp härrör främst från energiproduktion, transporter och industriprocesser.

Executive summary

ES.1 Background information

Greenhouse gases have always been present in the atmosphere, but now concentrations of several of them are rising as a result of human activity, which intensifies the greenhouse effect. An Intergovernmental Panel on Climate Change (IPCC) was established in 1988 and two years later they came up with the conclusion that anthropogenic climate change is a global threat and asked for an international agreement to deal with the problem. The United Nations started negotiations to create a framework convention on climate change (UNFCCC), which came into force in 1994. The long-term goal is to stabilize the amount of greenhouse gases in the atmosphere at a level where harmful anthropogenic climate changes is prevented. The most important addition to the convention was negotiated in 1997 in Kyoto, Japan. The Kyoto protocol involves binding obligations for the Annex I parties to decrease their emissions of greenhouse gases (GHG) with at least 5% during 2008-2012 compared to the base year 1990. According to Articles 4 and 12 of the United Nations Framework Convention on Climate Change (UNFCCC), Annex I parties are required to annually submit national inventories of anthropogenic emissions by sources and removals by sinks of all greenhouse gases not controlled by the Montreal Protocol. The submission of inventories should including emissions in the Common Reporting Format (CRF) and a National Inventory Report (NIR).

This report constitutes Sweden's NIR 2013 for anthropogenic emissions of direct greenhouse gases CO₂, CH₄, N₂O, HFC, PFC, SF₆ and indirect greenhouse gases NO_x, CO, NMVOC and SO₂. The report contains information on Sweden's inventories of greenhouse gases for all years from 1990 to 2011, including descriptions of methods, data sources, uncertainties, the quality assurance and quality control (QA/QC) activities carried out and a trend analysis.

Electronic data on emissions, activity data and emission factors in the Common Reporting Format (CRF) requested by the UNFCCC are provided together with this report.

ES.2 Summary of National Emissions and Removal Related Trends, including KP-LULUCF

ES.2.1 GHG inventory

Total greenhouse gas emissions in Sweden excluding LULUCF, expressed in carbon dioxide equivalents, were about 61.4 million tonnes for 2011 with an uncertainty of 4.5% (Table ES 1). The emission has decreased by about 4 million tonnes compared to 2010. Emissions have decreased by about 16% or approximately 11 million tonnes between 1990 and 2011. The uncertainty

of the trend of national total greenhouse gas emissions excluding LULUCF was $\pm 2.0\%$. The uncertainty in the trend is a percentage point range relative to the inventory trend and should be interpreted as $\pm 2.0\%$ to the estimated percentage difference between total GHG emissions 1990 and 2011, i.e. there is a 95% probability that the decrease in GHG emissions in Sweden 1990 to 2011 is between 14% and 18%.

In 2011, the net uptake of the land use, land-use change and forestry (LULUCF) has been estimated to 35 million tonnes carbon dioxide equivalent (Table S1). The annual change of the sink is very small compared to the size of pools (living biomass, dead organic matter and land-based carbon). For this reason, the estimations are associated with significant uncertainties. The uncertainties and change of methods may result in substantial changes on annual values and between different submissions.

Emissions of CO₂ were around 48.7 million tonnes in 2011, which is about 14% lower than in 1990 (Table ES 1). About 88% of total carbon dioxide emissions come from the energy sector, including transport, which is the largest source of carbon dioxide in Sweden. Carbon dioxide's share of the total GHG emissions is approximately 79%.

Emissions of methane (CH₄) arise mainly from agriculture and landfill sites, and in 2011 were approximately 5 million tonnes CO₂-equivalents (Table ES 1). Since 1990, emissions have decreased about 28%, primarily due to measures implemented in the waste sector and agriculture.

In 2011, total emissions of nitrous oxide (N₂O) were around 6.7 million tonnes, CO₂-equivalent (Table ES 1), a reduction of 20% compared to 1990. Emissions arise mainly from agriculture, but also from energy production, wastewater handling and industrial processes. The agricultural sector accounts for the bulk of the N₂O decline.

Total emissions of fluorinated gases (PFCs, HFCs and SF₆) in 2011 were approximately one million tonnes expressed in carbon dioxide equivalents (Table ES 1). This corresponds to an increase of about 116% compared to 1990. The increase is due to the replacement of the ozone-depleting substances by HFCs.

ES.2.2 KP-LULUCF activities

Sweden reports Articles 3.3 and 3.4 of the Kyoto Protocol. For these activities, Sweden has chosen to report the uptake/emission for the entire commitment period but not on an annual basis.

The activities under Article 3.3, afforestation / reforestation (AR), are relatively rare in Sweden. Since 2008 the area afforested / reforested land increased by about 5,000 ha per year. The uptake for all pools has been estimated to 1.66 million tonnes CO₂ equivalents in 2011. The proportion of land that is deforested (D), increased slightly compared to 2008. The total emission in 2011 is estimated to 2.6 million tonnes CO₂ equivalents. Under Article 3.4, Sweden has chosen accounting of forestry. For this activity, the uptake is approximately 37.6 million tons of CO₂ equivalents in 2011. Reporting under the Kyoto Protocol Article 3.3 and 3.4 is in line with the UNFCCC reporting for forest and land converted to forest land (area).

Table ES.1. Greenhouse gas emissions by gas (Gg CO₂ equivalents).

GREENHOUSE GAS EMISSIONS CO ₂ -eq (Gg)	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
CO ₂ emissions including net CO ₂ from LULUCF	19,688	27,232	18,529	19,332	20,247	23,938	26,273	26,044	18,750	20,609	17,044	13,394	21,464	13,377
CO ₂ emissions excluding net CO ₂ from LULUCF	56,954	58,872	54,145	55,078	55,993	56,608	55,768	53,231	53,194	51,972	50,006	46,405	52,302	48,726
CH ₄ emissions including CH ₄ from LULUCF	6,940	6,829	6,255	6,214	6,033	5,888	5,910	5,780	5,711	5,474	5,283	5,172	5,077	4,987
CH ₄ emissions excluding CH ₄ from LULUCF	6,938	6,827	6,252	6,211	6,028	5,882	5,904	5,775	5,699	5,471	5,270	5,170	5,076	4,985
N ₂ O emissions including N ₂ O from LULUCF	8,449	8,132	7,675	7,493	7,385	7,347	7,314	7,165	7,191	6,930	7,078	6,895	7,170	6,797
N ₂ O emissions excluding N ₂ O from LULUCF	8,370	8,070	7,602	7,420	7,314	7,270	7,232	7,073	7,096	6,825	6,956	6,779	7,033	6,682
HFCs	4	132	568	615	666	710	769	789	818	838	867	869	845	813
PFCs	377	343	241	236	261	258	254	257	245	248	225	35	158	183
SF6	107	127	94	111	104	69	81	142	111	151	84	81	73	60
Total (including LULUCF)	35,566	42,795	33,360	34,001	34,696	38,210	40,601	40,178	32,827	34,250	30,581	26,446	34,787	26,217
Total (excluding LULUCF)	72,750	74,371	68,902	69,670	70,366	70,797	70,009	67,268	67,164	65,506	63,407	59,338	65,487	61,449

Table ES.2. Greenhouse gas emissions by sector (Gg CO₂ equivalents).

GREENHOUSE GAS SOURCE AND SINK CATEGORIES CO ₂ -eq (Gg)	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
1. Energy	53,670	55,464	50,584	51,460	52,242	53,179	51,960	49,604	49,608	48,249	46,401	44,508	48,804	45,015
2. Industrial Processes	6,330	6,644	6,812	6,810	6,951	6,679	7,071	6,976	6,969	6,922	6,804	4,986	6,810	6,661
3. Solvent and Other Product Use	332	309	278	269	276	292	311	303	299	281	288	270	289	289
4. Agriculture	8,997	8,722	8,313	8,260	8,171	8,060	8,094	7,954	7,932	7,856	7,915	7,683	7,786	7,772
5. Land Use, Land-Use Change and Forestry	-37,184	-31,576	-35,541	-35,669	-35,670	-32,587	-29,408	-27,091	-34,337	-31,255	-32,826	-32,891	-30,701	-35,232
6. Waste	3,421	3,233	2,916	2,871	2,726	2,586	2,572	2,431	2,356	2,197	2,000	1,892	1,798	1,713
7. Other	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO
Total (including LULUCF)(5)	35,566	42,795	33,360	34,001	34,696	38,210	40,601	40,178	32,827	34,250	30,581	26,446	34,787	26,217

ES.3 Overview of Source and Sink Category Emission Estimates and Trends, including KP-LULUCF

ES.3.1 GHG inventory

The methods used to calculate the emissions and removals are in accordance with the Revised IPCC 1996 Guidelines for National Greenhouse Gas Inventories and the IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories. The report is prepared in accordance with the Reporting Guidelines, agreed by the UNFCCC at the eighth session of the Conference of the Parties (COP) in New Delhi 2002 and subsequent decisions.

The sectors included in the inventory and the main sources used for activity data and/or emission data are presented in Table ES 3. The emissions are collected directly from these data sources, or calculated based on activity data.

Table ES.3. CRF sectors and data sources used in the inventory.

CRF	Sector	Main source for activity/emission data
1	Energy	
	-Stationary combustion	Statistical survey on energy consumption
	-Transport	Transport authorities
2	Industrial processes	Environmental reports
		Statistical survey on energy consumption
		Direct contact with companies
		CO ₂ Data from the European trading scheme (ETS)
3	Solvent and Other Product Use	National data from the Products register at the Swedish Chemicals Agency
		National experts
		Environmental reports
4	Agriculture	Official statistical reports
		Organisations and researchers
5	Land Use Change and Forestry	Swedish University of Agricultural Sciences
		Swedish Forest Agency
6	Waste	Swedish Association of Waste Management
		The Swedish Forest Industries Federation
		Statistics Sweden
		Swedish Environmental Protection Agency
		Environmental reports

Emissions from the energy sector vary due to temperature and precipitation conditions and the state of the economy, but the trend for the period 1990-2011 is generally decreasing emissions.

Greenhouse gas emissions from the energy sector were about 45 million tonnes, expressed as carbon dioxide equivalents in 2011 (Table ES 2), which is equivalent to 73% of the total emissions. From 1990 to 2011, there was an increase of about 6% for greenhouse gas emissions from road traffic. In 2011

greenhouse gases emissions have decreased compared to 2010. The reason is mainly decreased emissions from electricity and heat production and transports.

Carbon dioxide emissions from production of electricity and district heating in 2011 have decreased by about 22% (about 8.3 million tonnes) compared to 2010 (about 10.6 million tonnes carbon dioxide equivalents). Some of the underlying causes are decreased need of heating due to warmer weather, high hydropower production and a slight increase in nuclear production.

For industrial processes, the most dominant greenhouse gas is carbon dioxide, which makes about 82% of the emissions for this sector, followed by fluorinated greenhouse gases (16%), nitrous oxide (about 2%) and methane (0.2%).

Emissions from industrial processes in Sweden are primarily derived from production of iron and steel. Total emissions in 2011 were approximately 6.7 million tonnes CO₂ equivalents (Table ES 2), which makes approximately 11% of the national total emissions. The overall emissions from this sector decreased by 0.15 million tonnes of carbon dioxide equivalents or about 2% between 2010 and 2011. This decrease is not related to any drop in production volume, but to upgrading of two factories with nitric acid production which led to a significant decreased of N₂O-emission (about -87% compared to 2010). Since 1990, emissions in this sector have varied primarily because of variation in production volumes / economic cycles. In 2011, emissions were 5% higher than in 1990.

The use of Solvents and Other products mainly gives rise to emissions of volatile organic substances (VOCs), nitrous oxides (N₂O) and some carbon dioxide. In 2011, emissions of carbon dioxide and nitrous oxide were almost 0.3 million tonnes CO₂ equivalents (Table ES 2), which corresponds to about 0.5% of total greenhouse gas emissions. Compared to 1990, emissions have decreased with about 13%. Almost 9% of carbon dioxide emissions arise from paint application, even though these emissions have decreased due to a transition to water-based paints.

Agriculture is the largest source of methane and nitrous oxide emissions. In 2011, total greenhouse gas emissions were 7.8 million tonnes CO₂ equivalents (Table ES 2) of which 63% consisted of nitrous oxide and 37% of methane. Emissions decreased by about 14% compared to 1990. Methane emissions arise primarily from the digestive processes of cattle and from manure management. The most important reasons for the reduced emissions are reduced livestock keeping and reduced application of N-fertilisers in agriculture. Nitrous oxide emissions originate mainly from transformation of nitrogen that takes place in the ground, which is influenced by the use of manure and commercial fertiliser and the cultivation of nitrogen-fixing crops. More than a half (57%) of the sector's emissions comes from agricultural land.

The net removal for LULUCF in 2011 is estimated to around 35 million tonnes of CO₂. Net uptake increased slightly between 2010 and 2011 with about 4.5 million tons of CO₂ (Table S2). The long-term trend of decreasing

net absorption in living biomass is mainly due to an increase in logging, while the increase in gross uptake derived from tree growth. If growth is less than felling, a decrease in net removal is obtained.

Emissions from the waste sector in 2011 is dominated by methane emissions from landfills. Methane makes up to 87%, while nitrous oxide emissions from waste water account for 9% and carbon dioxide emissions from the incineration of hazardous waste is about 3%. In 2011, total emissions from the waste sector were about 1.7 million tonnes CO₂ equivalents (Table ES 2), or 2.8% of national total GHG emissions. Compared to 1990, emissions in 2011 have decreased by about 50%. The collection of landfill gas, a ban on landfill deposit for organic material and the introduction of a landfill tax have played a key role for the decrease in emissions.

The total emissions of greenhouse gases in 2011 were about 61.5 million tonnes expressed as carbon dioxide equivalents and the emissions that come from sectors outside the trading system were 43.6 million tonnes. This means that emissions in Sweden in 2011 were 16% below the base year emissions.

ES.3.2 KP-LULUCF activities

As the activities under Article 3.3 (afforestation and deforestation) are relatively uncommon in Sweden, it is difficult to draw any conclusions on the differences between years. The net uptakes and emissions are approximately at the same levels as in previous years.

Net removals related to Forest management under Article 3.4, decreased during the previous years. This is mainly an effect of reduced felling.

ES.4 Overview of Emission Estimates and Trends of Indirect GHGs and SO₂

Emissions of nitrogen oxides (NO_x) were about 146 ktonnes in 2011 (Table ES 4), a reduction of 46% compared to 1990. Road traffic emissions of NO_x have decreased by 57% between 1990 and 2011 and by about 7% between 2010 and 2011. The largest sources of emissions of nitrogen oxides are road traffic, mobile machinery, maritime transport and electricity and heating production. In urban areas, road traffic is the most significant contributor to emissions of nitrogen oxides, but the introduction of catalytic converters in the late 1980's and the subsequent successively more stringent emission standards have contributed to a general reduction of nitrogen oxide levels in urban areas. The increased use of district heating and the "NO_x charge" of the early 1990s have also resulted in a great reduction of emissions of nitrogen oxides from the energy sector.

Table ES.4. Emissions of indirect greenhouse gases and SO₂ (ktonnes).

Gas	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
NO _x	271	247	211	201	193	191	184	180	176	170	163	152	154	146
CO	1,280	1,127	826	785	739	725	678	669	629	617	607	606	587	571
NMVOC	359	278	224	213	205	206	202	199	195	193	189	184	183	177
SO ₂	105	69	42	41	41	41	37	36	36	32	30	29	32	30

Emissions of carbon monoxide (CO) have decreased from 1280 ktonnes in 1990 to 571 ktonnes in 2011 (Table ES 4), a reduction of 55%. About 96% of emissions come from energy sector of which 39% come from transport.

Emissions of volatile organic compounds (NMVOC) were 177 ktonnes in 2011 (Table ES 4), a decrease by one-half compared to 1990. The main contributors to NMVOC emissions are road traffic, wood combustion in the residential sector and products containing solvents. Voluntary environmental standards for new installations of wood-burning boilers and reduced emissions from products containing solvent have contributed to the decrease in emissions.

Emissions of sulphur dioxide (SO₂) have decreased from 105 ktonnes in 1990 to almost 30 ktonnes in 2011 (Table ES 4), a reduction of about 72%. Sulphur dioxide emissions derive from the energy, transport and industrial sectors. The reduction is mainly due to a transfer from fuels with high sulphur levels to low-sulphur fuels, both for road traffic and heating. A tax on sulphur, introduced in 1991, has been important in this transition.

Summary in Arabic

ملخص انبعاثات غازات الدفيئة في السويد لعام 2011

مقدمة

إن التغيرات المناخية التي نشهدها في عصرنا الحالي يعود سببها إلى زيادة تراكيز غازات الدفيئة (CO_2 , CH_4 , N_2O , HFC, PFC, SF_6) في الغلاف الجوي. لقد كانت هذه الغازات موجودة دائماً وبشكل طبيعي في الغلاف الجوي ولكن بتركيز قليلة. أظهرت البيانات والتحليل الكيميائية للغلاف الجوي في العقود الماضية إزدياداً مضطرباً لتراكيز هذه الغازات مقارنةً عما كانت عليه في العقود الأخيرة. وبالرغم من وجود هذه الغازات دائماً وبشكل طبيعي في الغلاف الجوي، فإن تراكيز بعضاً منها أخذت في الارتفاع نتيجة لإزدياد النشاط البشري متمثلاً في حرق المشتقات البترولية للحصول على الطاقة ومن خلال النشاطات الزراعية، والذي بدوره يفاقم من حدة ظاهرة الاحتباس الحراري.

ونتيجةً لذلك، وفي عام 1988 أنشئ الفريق الحكومي الدولي المعني بتغير المناخ (IPCC) لمتابعة هذه الظاهرة. وبعد ذلك بعامين خرج الفريق بنتيجة مفادها أن التغيرات المناخية هي من صنع الإنسان وهي تمثل تهديداً عالمياً للبشرية. لقد كانت هناك الحاجة إلى اتفاق دولي للتعامل مع هذه المشكلة. لذا بدأت الأمم المتحدة مفاوضات لإنشاء الاتفاقية الإطارية بشأن تغير المناخ (UNFCCC)، والتي دخلت حيز التنفيذ عام 1994. كان الهدف من هذه الاتفاقية هو تحقيق استقرار، وعلى المدى الطويل، لتراكيز غازات الدفيئة في الغلاف الجوي وعند مستوى يمكن فيها تجنب التغيرات المناخية الضارة والتي تنشأ من صنع الإنسان. في عام 1997 وفي كيوتو، اليابان، تم التفاوض لإضافة أكثر البنود أهمية بالنسبة للاتفاقية وهو بروتوكول كيوتو، والذي يشمل على التزامات ملزمة للبلدان المدرجة في المرفق الأول من الاتفاقية لخفض انبعاثات هذه الدول من الغازات المسببة للاحتباس الحراري (غازات الدفيئة) إلى ما لا يقل عن 5% خلال 2008-2012 مقارنة مع سنة الأساس 1990.

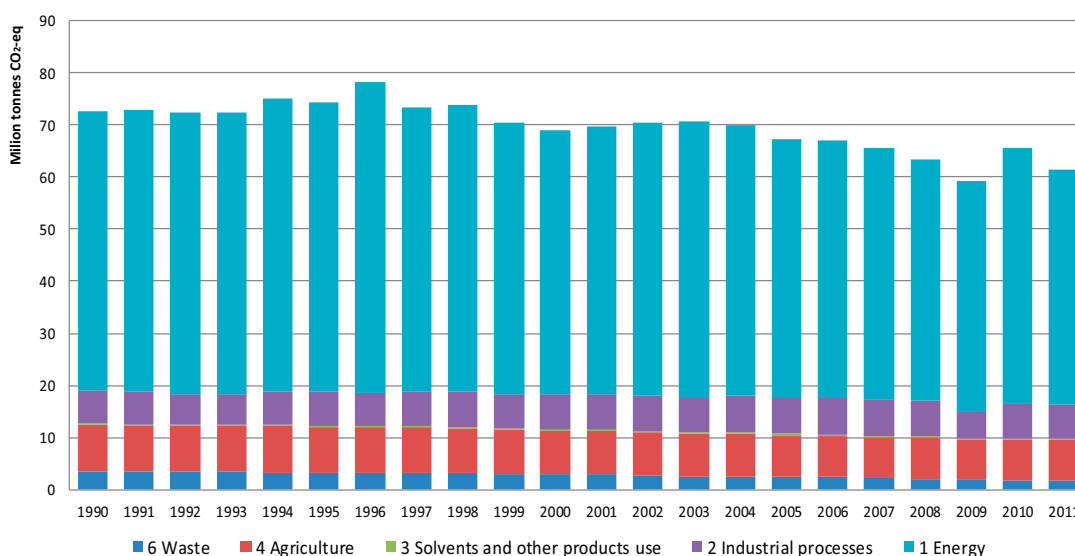
وفقاً للمادتين 4 و 12 من اتفاقية الأمم المتحدة الإطارية بشأن تغير المناخ (UNFCCC)، يطلب من الأطراف أن تقدم سنوياً قوائم الجرد الوطنية للانبعاثات البشرية المصدر بحسب مصادرها وعمليات إزاحتها بواسطة المصارف لجميع غازات الدفيئة غير الخاضعة لبروتوكول مونتريال. وينبغي أيضاً تقديم قوائم الجرد بما في ذلك الانبعاثات في نموذج الإبلاغ الموحد (CRF) وتقرير الجرد الوطني (NIR) National Inventory Report.

يشكل هذا التقرير (NIR submission 2013) جرداً للانبعاثات غازات الدفيئة المباشرة (CO_2 , CH_4 , N_2O , HFC, PFC, SF_6) في الغلاف الجوي وغير المباشرة (NMVOC، NO_x ، SO_2 ، CO) الناتجة من النشاطات البشرية في السويد لعام 2011 بالإضافة إلى معلومات عن قوائم الجرد لغازات الاحتباس الحراري لجميع السنوات من 1990 إلى 2011، بما في ذلك وصفا للطرق التحليل و معلومات أخرى متعلقة.

نظرة عامة عن تقديرات انبعاثات غازات الدفيئة المباشرة واتجاهاتها

قوائم جرد غازات الدفيئة

بلغ مجموع انبعاثات غازات الدفيئة في السويد لعام 2011 ما يقارب 61.5 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون، منخفضاً بنحو 4 ملايين طن مقارنةً بعام 2010. وتراجعت الانبعاثات بنسبة حوالي 16% أو ما يقرب من 11 مليون طن بين عامي 1990 و 2011. في عام 2011، تم تقدير صافي امتصاص غاز ثاني أكسيد الكربون بواسطة الغابات والأراضي الحرجية بحوالي 35 مليون طن مكافئ ثاني أكسيد الكربون (أنظر إلى الرسم البياني والجدول 1 و 2).



الرسم البياني مقادير الانبعاثات الصادرة من القطاعات المختلفة ما عدا قطاع تغيير إستخدام الأراضي والحرجة (LULUCF).

GREENHOUSE GAS EMISSIONS CO ₂ -eq (Gg)	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
CO ₂ emissions including net CO ₂ from LULUCF	19,688	27,232	18,529	19,332	20,247	23,938	26,273	26,044	18,750	20,609	17,044	13,394	21,464	13,377
CO ₂ emissions excluding net CO ₂ from LULUCF	56,954	58,872	54,145	55,078	55,993	56,608	55,768	53,231	53,194	51,972	50,006	46,405	52,302	48,726
CH ₄ emissions including CH ₄ from LULUCF	6,940	6,829	6,255	6,214	6,033	5,888	5,910	5,780	5,711	5,474	5,283	5,172	5,077	4,987
CH ₄ emissions excluding CH ₄ from LULUCF	6,938	6,827	6,252	6,211	6,028	5,882	5,904	5,775	5,699	5,471	5,270	5,170	5,076	4,985
N ₂ O emissions including N ₂ O from LULUCF	8,449	8,132	7,675	7,493	7,385	7,347	7,314	7,165	7,191	6,930	7,078	6,895	7,170	6,797
N ₂ O emissions excluding N ₂ O from LULUCF	8,370	8,070	7,602	7,420	7,314	7,270	7,232	7,073	7,096	6,825	6,956	6,779	7,033	6,682
HFCs	4	132	568	615	666	710	769	789	818	838	867	869	845	813
PFCs	377	343	241	236	261	258	254	257	245	248	225	35	158	183
SF ₆	107	127	94	111	104	69	81	142	111	151	84	81	73	60
Total (including LULUCF)	35,566	42,795	33,360	34,001	34,696	38,210	40,601	40,178	32,827	34,250	30,581	26,446	34,787	26,217
Total (excluding LULUCF)	72,750	74,371	68,902	69,670	70,366	70,797	70,009	67,268	67,164	65,506	63,407	59,338	65,487	61,449

1- أَلْجَدُول يَبِين مَقَادِير الْاَنْبِعَاثَات وَالْاِزَالَة لِغَازَات الدَّفِيئَة الْمَخْتَلِفَة فِي السُّوَيْد.

GREENHOUSE GAS SOURCE AND SINK CATEGORIES CO ₂ -eq (Gg)	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
1. Energy	53,670	55,464	50,584	51,460	52,242	53,179	51,960	49,604	49,608	48,249	46,401	44,508	48,804	45,015
2. Industrial Processes	6,330	6,644	6,812	6,810	6,951	6,679	7,071	6,976	6,969	6,922	6,804	4,986	6,810	6,661
3. Solvent and Other Product Use	332	309	278	269	276	292	311	303	299	281	288	270	289	289
4. Agriculture	8,997	8,722	8,313	8,260	8,171	8,060	8,094	7,954	7,932	7,856	7,915	7,683	7,786	7,772
5. Land Use, Land-Use Change and Forestry	-37,184	-31,576	-35,541	-35,669	-35,670	-32,587	-29,408	-27,091	-34,337	-31,255	-32,826	-32,891	-30,701	-35,232
6. Waste	3,421	3,233	2,916	2,871	2,726	2,586	2,572	2,431	2,356	2,197	2,000	1,892	1,798	1,713
7. Other	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO
Total (including LULUCF)(5)	35,566	42,795	33,360	34,001	34,696	38,210	40,601	40,178	32,827	34,250	30,581	26,446	34,787	26,217

2- أجدول يبين مقادير الانبعاثات والازالة الصادرة من القطاعات المختلفة في السويد

غاز ثاني أكسيد الكربون (CO₂)

بلغت انبعاثات CO₂ حوالي 48.7 مليون طن في عام 2011، منخفضاً بما يقارب 14٪ مقارنةً بعام 1990. ويستحوذ قطاع الطاقة، بما في ذلك قطاع النقل، على حوالي 88٪ من إجمالي الانبعاثات، لذا يعتبر هذا القطاع المصدر الأكبر لغاز ثاني أكسيد الكربون في السويد. ويمثل هذا الغاز حوالي 79 بالمئة من الانبعاثات الإجمالية لغازات الدفيئة.

غاز الميثان (CH₄)

ينبعث غاز الميثان (CH₄) بشكل أساسي من النشاطات الزراعية ومواقع مكبات النفايات حيث يتكون غاز الميثان خلال عمليات الهضم للحيوانات المجترة وعمليات التخمر لروث هذه الحيوانات في المزارع. أما في قطاع النفايات فيتكون الميثان نتيجة تخمر النفايات العضوية. كان مجمل الانبعاث في عام 2011 بحوالي 5 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون، منخفضاً حوالي 28٪ مقارنة مع عام 1990، ويرجع هذا الانخفاض في المقام الأول إلى التدابير المنفذة في قطاع النفايات والزراعة والتي تحد من تكون الميثان من هذين القطاعين.

أكسيد النيتروز (N₂O)

في عام 2011، بلغت انبعاثات أكسيد النيتروز (N₂O) حوالي 6.7 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون، منخفضاً بنسبة 20٪ مقارنة مع عام 1990. ينشأ هذا الغاز أساساً من قطاع الزراعة وخصوصاً من تخمر روث حيوانات المزارع ومن خلال انبعاث هذا الغاز من السماد العضوي وغير العضوي المستعمل لتحسين مستوى المحاصيل الزراعية. كما يتكون هذا الغاز خلال عمليات إحتراق الوقود المتعلقة في إنتاج الطاقة ومن خلال معالجة مياه الصرف الصحي والعمليات الصناعية. ويمثل القطاع الزراعي الجزء الأكبر من الانخفاض وذلك للتدابير المنفذة في القطاع للحد من انبعاث هذا الغاز.

غازات الكربون المشبعة بالفلور

بلغ مجموع الانبعاثات من الغازات المفلورة (مركبات الكربون المشبعة بالفلور، ومركبات الكربون SF₆) في عام 2011 ما يقرب من مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون. ويتوافق هذا مع زيادة قدرها حوالي 116٪ مقارنة بعام 1990. وتعزى هذه الزيادة إلى الاستعاضة عن المواد المستنفدة للأوزون من مركبات الكربون المشبعة بالكلور والفلور. القطاع الوحيد المسؤول عن هذه الانبعاثات هو قطاع الصناعة حيث تدخل مركبات الكربون المشبعة بالفلور في العديد من الصناعات مثل الاسفنج والمطاط الصناعي.

نظرة عامة عن تقديرات الانبعاثات والإزالة لغازات الدفيئة من القطاعات المختلفة

قطاع الطاقة

تتأثر كميات الانبعاثات الناجمة من قطاع الطاقة بمعدل درجات الحرارة السنوية (شتاء بارد أو معتدل نسبياً) و معدل هطول الأمطار (وهذا يؤثر على معدل إنتاج الطاقة الكهرومائية) وحالة الاقتصاد (معدل الانتاج الصناعي وتأثره بالإنتعاش أو الركود). عموماً فإن اتجاه كميات الانبعاثات من قطاع الطاقة للفترة 1990-2011 يشير إلى انخفاض مستمر. في عام 2011 انخفضت انبعاثات الغازات المسببة للاحتباس الحراري مقارنة بعام 2010 وذلك لتناقص الانبعاثات الناجمة عن إنتاج الطاقة والنقل. لقد بلغ مجموع انبعاثات غازات الدفيئة من قطاع الطاقة بما في ذلك النقل لعام 2011 حوالي 45 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون. وهذا ما يعادل 73٪ من إجمالي الانبعاثات. وفي الفترة ما بين عام 1990 إلى 2011 إزدادت كميات الانبعاثات الصادرة من قطاع المواصلات (حركة المرور على الطرق) بحوالي 6٪.

انخفضت انبعاثات ثاني أكسيد الكربون من إنتاج الكهرباء وطاقة التدفئة في عام 2011 بنحو 22٪ مقارنةً مع عام 2010 ويرجع هذا إلى الاستخدام العالي نسبياً للطاقة الكهرومائية والطاقة النووية (نظراً لعدم إنتاج أية انبعاثات من هذه المصادر) وبدوره قلة استخدام الوقود الأحفوري.

قطاع الصناعة

يعتبر ثاني أكسيد الكربون أبرز غازات الدفيئة الذي ينبعث من قطاع الصناعة (حوالي 82٪) تليه الغازات المفلورة (بنسبة 16٪)، وأكسيد النيتروز وغاز والميثان بنسبٍ ضئيلة.

معظم هذه الانبعاثات من هذا القطاع تصدر أساساً من العمليات الصناعية المتعلقة من إنتاج الحديد والصلب في السويد. قدرت الانبعاثات الإجمالية من هذا القطاع في عام 2011 ما يقرب من 6.7 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون، مما يجعل هذا القطاع يمثل ما يقرب من 11٪ من اجمالي الانبعاثات الوطنية. لقد انخفضت الانبعاثات الكلية من هذا القطاع بين عامي 2010 و 2011 بحوالي 0.15 مليون طن مكافئ ثاني أكسيد الكربون أو حوالي 2٪. لم يكن هذا الانخفاض نتيجة لانخفاض في حجم الإنتاج، وإنما كان سببه تطوير اثنتين من الوحدات المسؤولة عن إنتاج حمض النيتريك في إحدى المنشآت مما أدى إلى انخفاض كبير (حوالي 87٪ مقارنة مع 2010) في انبعاثات غاز أكسيد النيتروز. منذ عام 1990 وكميات الانبعاثات في تذبذب وذلك نتيجة لتذبذب حجم الإنتاج والذي يتأثر بحالة الاقتصاد العالمي. بقي القول أن انبعاثات غازات الدفيئة من هذا القطاع عام 2011، كانت أعلى بمستوى 5٪ مقارنةً بعام 1990.

قطاع المذيبات واستخدام والمنتجات

ينشأ من هذا القطاع انبعاثات المركبات العضوية المتطايرة (NMVOC) وأكسيد النيتروز وبعض ثاني أكسيد الكربون. في عام 2011 بلغ تقريباً مجموع الانبعاثات 0.3 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون. وهذه الكمية تعادل حوالي 0.5٪ من إجمالي انبعاثات غازات الدفيئة في السويد. وبالمقارنة مع عام 1990، انخفضت الانبعاثات بحوالي 13٪. والسبب الرئيسي يعود إلى استخدام الدهانات التي تتركز على الماء في تركيبها بدلاً عن التي تتركز على الزيت.

قطاع الزراعة

قطاع الزراعة هو أكبر مصدر لانبعاثات غاز الميثان وغاز أكسيد النيتروز. في عام 2011، بلغ مجموع انبعاثات هذا القطاع حوالي 7.8 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون. ويشكل أكسيد النيتروز وغاز الميثان 63٪ و 37٪ على التوالي لهذا القطاع. انخفضت الانبعاثات لعام 2011 بنسبة حوالي 14٪ مقارنة بعام 1990 وذلك لانخفاض عدد مزارع تربية المواشي في السويد. وينشأ غاز الميثان أساساً من خلال عمليات الهضم وكذلك خلال عمليات التخمر لروث الماشية. أما انبعاثات أكسيد النيتروز فتنشأ أساساً من تبيخ هذا الغاز نتيجة استخدام الأسمدة العضوية والتجارية وزراعة المحاصيل المثبتة للنيتروجين. ويأتي أكثر من نصف (57٪) هذه الانبعاثات من الأراضي الزراعية.

قطاع تغيير استخدام الأراضي والحراجة (LULUCF)

في عام 2011 قدر صافي إزالة ثاني أكسيد الكربون من قطاع تغيير استخدام الأراضي والحراجة (LULUCF) بحوالي 35 مليون طن. وقد زاد صافي الإزالة قليلاً بحوالي 4.5 مليون عام 2011 مقارنةً مع 2010.

قطاع النفايات

يهيمن غاز الميثان على الغازات الأخرى التي تنشأ من قطاع النفايات حيث بلغت انبعاثات الميثان في عام 2011 بنحو 87٪ من إجمالي القطاع. في حين بلغت انبعاثات أكسيد النيتروز من المياه العادمة حوالي 9٪ وانبعاثات ثاني أكسيد الكربون الناتجة عن حرق النفايات نحو 3٪. وفي عام 2011 كانت مجموع الانبعاثات من هذا قطاع ما يقرب من 1.7 مليون طن مكافئ محسوبة بما يعادلها من ثاني أكسيد الكربون. وتشكل هذه الكمية حوالي 2.8٪ من إجمالي انبعاثات غازات الدفيئة الوطنية. بالمقارنة مع عام 1990 فقد انخفضت الانبعاثات بنحو 50٪ وذلك نتيجة لجمع غاز الميثان من مكبات النفايات حيث يتكون هذا الغاز خلال عمليات التخمر للنفايات العضوية. وقد لعبت مجموعة من القوانين من الحد من انبعاثات الميثان من النفايات كقرص حظراً على التخلص مباشرةً من النفايات العضوية في المكبات وإدخال ضريبة على طمر النفايات دوراً رئيسياً لانخفاض في الانبعاثات

نظرة عامة عن تقديرات انبعاثات غازات الدفيئة الغير المباشرة واتجاهاتها

أكاسيد النيتروجين (NOx)

قدرت كميات انبعاثات أكاسيد النيتروجين بحوالي 146 كيلو طن في عام 2011 (أنظر إلى الجدول 3).

بتناقص 46٪ مقارنة بعام 1990. كما انخفضت انبعاثات أكاسيد النيتروجين من حركة المرور على الطرق بنسبة 57٪ بين عامي 1990 و 2011 وبنسبة حوالي 7٪ بين عامي 2010 و 2011. تعتبر حركة المرور على الطرق في المدن والمركبات الكبيرة المتنقلة والنقل البحري والإنتاج الكهربائي والتدفئة من أكبر مصادر انبعاثات أكاسيد النيتروجين. ونظراً لأن حركة المرور على الطرق هي المساهم الأكبر لهذه الانبعاثات فقد أدخلت في أواخر عام 1980 تعديلات على المركبات المتنقلة وما رافقها من معايير أكثر صرامة للحد من هذه الانبعاثات كلها أسهمت في خفض عام لمستويات أكسيد النيتروجين. وقد أدت زيادة استخدام التدفئة

المركزية المعتمدة على الوقود في أوائل تسعينات القرن الماضي إلى سن قانون يعرف "بضريبة انبعاثات أكاسيد النيتروجين" أدى هذا أيضا إلى انخفاض كبير لانبعاثات أكاسيد النيتروجين من قطاع الطاقة.

أول أكسيد الكربون (CO)

انخفضت انبعاثات أول أكسيد الكربون من 1280 كيلو طن في 1990 إلى 571 كيلو طن في عام 2011، بخفض قدره 55%. يتكون غاز أول أكسيد الكربون من خلال عمليات حرق الوقود في قطاع الطاقة. ينتج هذا القطاع حوالي 96% من الانبعاثات حيث يأتي 39% تقريبا من وسائل النقل والتي تنضوي تحت قطاع الطاقة.

المركبات العضوية المتطايرة (NMVOC)

قدرت كميات انبعاثات المركبات العضوية المتطايرة عام 2011 بحوالي 177 كيلو طن، بانخفاض مقداره النصف مقارنة بعام 1990. تعتبر وسائل النقل على الطرق المساهم الرئيسي في انبعاثات المركبات العضوية المتطايرة. كما أن احتراق الحطب للتدفئة في القطاع السكني وكذلك استخدام المذيبات في المنتجات الصناعية ينشأ عنهما انبعاث المركبات العضوية المتطايرة بنسب متفاوتة. وقد أسهمت المعايير البيئية الغير إلزامية المتبعة في تقنيات موافد الحطب الجديدة من خفض هذه الانبعاثات بشكل محسوس.

ثاني أكسيد الكبريت (SO₂)

انخفضت انبعاثات ثاني أكسيد الكبريت من 105 كيلو طن في عام 1990 إلى ما يقرب من 30 كيلو طن في عام 2011 أي بنسبة 72% تقريبا. ينبعث ثاني أكسيد الكبريت قطاعات الطاقة والنقل والصناعة. ويعود هذا الانخفاض أساسا إلى استخدام لأنواع الوقود المنخفضة الكبريت سواء وقود المركبات أو وقود التدفئة. كما أن فرض ضريبة على الوقود الذي يحتوي على نسب عالية من الكبريت عام 1991 ساهم في الحد من هذه الانبعاثات.

Gas	1990	1995	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
NOx	271	247	211	201	193	191	184	180	176	170	163	152	154	146
CO	1,280	1,127	826	785	739	725	678	669	629	617	607	606	587	571
NMVOC	359	278	224	213	205	206	202	199	195	193	189	184	183	177
SO ₂	105	69	42	41	41	41	37	36	36	32	30	29	32	30

3- أجدول يبين مقادير انبعاثات غازات الدفيئة الغير المباشرة الصادرة من القطاعات المختلفة في السويد.

PART 1: ANNUAL INVENTORY SUBMISSION 2013

1 Introduction

According to Articles 4 and 12 of the United Nations Framework Convention on Climate Change (UNFCCC), Annex I Parties are required to annually submit national inventories of anthropogenic emissions by sources and removals by sinks of all greenhouse gases not controlled by the Montreal Protocol. The inventory submitted to the UNFCCC Parties, through the secretariat, should include emissions in the Common Reporting Format (CRF) and a National Inventory Report (NIR).

This report constitutes Sweden's NIR for submission 2013. The report contains information on Sweden's inventories for all years from 1990 to 2011 including descriptions of methods, data sources, uncertainties, quality assurance and quality control (QA/QC) activities carried out, and a trend analysis. In order to ensure the transparency, consistency, comparability, completeness and accuracy of the inventory, the report contains information on inventories for all years from the base year to the year of the current annual inventory submission.

This section presents background information on climate change and greenhouse gas (GHG) inventories. It also contains a description of institutional arrangements for the inventory preparation, brief descriptions of the process of inventory preparation, methodologies and data sources used and the key sources in the Swedish inventory. Finally there is information about the progress of quality assurance/quality control (QA/QC) work, the general uncertainties in the inventory and on the completeness of inventoried emissions.

1.1 Background Information

1.1.1 Climate change

In consequence of scientific indications that human activities influence the climate and an increasing public awareness about local and global environmental issues during the middle of the 1980s, climate change was brought up on the political agenda. The Intergovernmental Panel on Climate Change (IPCC) was established in 1988 and two years later they concluded that anthropogenic climate change was a global threat and asked for an international agreement to deal with the problem.

The United Nations started negotiations to create a framework convention on climate change (UNFCCC), which came into force in 1994. A decade later UNFCCC had 188 member states (including EU as a part). The long-term goal is to stabilize the amount of greenhouse gases in the atmosphere

at a level where harmful anthropogenic climate changes are prevented. After the UNFCCC came into force, the framework convention has developed and every year a Conference of the Parties (COP) is held. The most important addition to the convention was negotiated in 1997 in Kyoto, Japan. The Kyoto protocol involves binding obligations for the Annex I countries (including all EU member states and other industrialized countries). Together the emissions of greenhouse gases in these countries should be at least 5 % lower during 2008-2012 compared to the base year 1990 (for fluorinated greenhouse gases it is allowed to use 1995 as a base year).

In the spring 2002 Sweden, together with the other EU member states, ratified the Kyoto protocol and the 16th of February 2005 it came into force. EU and its member states uses a paragraph in the Kyoto protocol which gives them the right to, instead of national emission objective, have a joint EU objectives of a decrease in emissions with 8 %. Within EU the 8 is shared among the member states in accordance with the burden sharing agreement¹. For Sweden the agreement involves an allowed increase in emissions of 4 %. Sweden has chosen to go beyond the EU target. Reduced Climate Impact is one of the 16 Swedish Environmental Quality Objectives and is supported by long and short-term emissions targets. In 2009, Sweden adopted a vision that in 2050 the country will have no net emissions of greenhouse gases in the atmosphere. A comprehensive reduction strategy is set out to 2020.

2008–2012: Swedish greenhouse gas emissions will decrease by four per cent in comparison with 1990.

2020: Greenhouse gas emissions in Sweden (this applies to activities outside the emissions trading scheme) will decrease by 40 per cent in comparison with 1990.

2050: The vision is for Sweden in 2050 not to have any net emissions of greenhouse gases into the atmosphere

The objective also involve that Sweden should encourage the global work to aim at the objective to stabilize the concentration of greenhouse gases in the atmosphere at a level that would prevent dangerous anthropogenic interference with the climate system.

Some of the gases in the earth's atmosphere have an ability to absorb infrared radiation (heat). They do not prevent sunlight reaching the earth's surface and warming it, but they do trap some of the infrared outgoing radiation. Without the natural greenhouse effect of the atmosphere, the surface of our planet would be almost 35°C colder than it is now.

Greenhouse gases (i.e. gases which contribute to the greenhouse effect) have always been present in the atmosphere, but now concentrations of several of them are rising as a result of human activity. This is intensifying

¹ 2002/358/EC

the greenhouse effect. The IPCC sums up the cause of the climate change we have witnessed over the last 50 years by stating that it is impossible to explain other than as the result of anthropogenic emissions of greenhouse gases.

Apart from carbon dioxide, other greenhouse gases are being emitted in larger quantities now than in pre-industrial times. These gases include nitrous oxide and methane. Ground-level ozone also contributes to the greenhouse effect. The amount of ozone forming in the lower atmosphere has increased as a result of emissions of nitrogen oxides, hydrocarbons and carbon monoxide.

Entirely new, man-made greenhouse gases that are entering the atmosphere cause further intensification of the greenhouse effect. These include, in particular, a number of substances containing fluorine, among them HFCs (compounds of hydrogen, fluorine and carbon). HFCs are used instead of the ozone layer depleting CFCs (freons) in refrigerators and other applications, and their use is on the increase.

Compared with carbon dioxide, all other greenhouse gases occur at very low concentrations. Per molecule, however, these substances are much more effective as greenhouse gases than carbon dioxide, which means that they still make a considerable contribution to the greenhouse effect. Furthermore, some of the fluorine compounds have such a long atmospheric lifetime that they will contribute to the greenhouse effect for ten thousands of years to come.

The threat of climate change is considered to be one of the most serious environmental problems faced by humankind.

1.1.2 Greenhouse gas inventories

The inventory covers anthropogenic emissions of direct greenhouse gases CO₂, CH₄, N₂O, HFC, PFC, SF₆ and indirect greenhouse gases NO_x, CO, NMVOC and SO₂. Indirect means that they do not contribute directly to the greenhouse effect, but that their presence in the atmosphere may influence climate in different ways. Ozone (O₃) is also a greenhouse gas but, since it is formed by the chemical reactions of nitrogen oxides, hydrocarbons and/or carbon monoxide, a separate report is not necessary.

The obligations of the Kyoto protocol have led to an increased need for international supervision of the emissions reported by the parties. The Kyoto protocol therefore contains rules for how emissions should be estimated, reported and reviewed. Emissions of the direct greenhouse gases CO₂, N₂O, CH₄, HFCs, PFCs and SF₆ are calculated as CO₂ equivalents and added to produce a total. Together with the direct greenhouse gases, also the emissions of NO_x, CO, NMVOC and SO₂ are reported to UNFCCC. These gases are not included in the obligations of the Kyoto protocol. When a method used to estimate emissions is improved, a need to recalculate the whole time series may arise in order to maintain consistency. This means that data presented can be changed in the next submission.

1.1.3 Supplementary information required under Article 7, paragraph 1, of the Kyoto Protocol

Sweden provides supplementary information under Article 7 of the Kyoto Protocol for the Land Use, Land-Use Change and Forestry sector. The inventory for Kyoto protocol activities encompass emissions/ removals originating from the activities Afforestation and Reforestation (AR), Deforestation (D) and Forest management (FM) under article 3.3 and 3.4 under the Kyoto protocol, respectively. FM covers a major part of the Swedish land area whereas ARD are quite uncommon in Sweden.

To a large extent the KP-reporting of FM and AR harmonize with the UNFCCC-reporting of Forest land and land converted to Forest land. Small discrepancies occur regarding the accumulation of reported land areas as described in section 11.

In addition to the reporting of carbon pool changes, direct N₂O emissions from N fertilization and emissions from forest fires are reported under FM. Forest fires – both natural and wildfires – are uncommon and, this far, has not been registered on AR-land.

N₂O emissions from disturbance associated with land use conversion from Forest land to Cropland are reported under D.

1.1.4 Sweden's commitment under the Kyoto Protocol and the EU Burden Sharing Decision

According to Sweden's commitment under the Kyoto Protocol and the EU burden sharing decision, Sweden's greenhouse gas emissions must not exceed 104% of the emissions in the base year. The base year is 1990 for all emissions, except for fluorinated greenhouse gases, which is 1995. The base year's emissions were 72.2 million tonnes carbon dioxide equivalents, when the assigned amount was established. The assigned amount for Sweden is calculated to 75 million tonnes per year as an average (Assigned Amount Units (AAU)). In addition, Sweden can also credit itself with a net removal of carbon dioxide of a maximum 2.13 million tonnes per year (Removal Units (RMU)), according to article 3.3 and 3.4 in the Kyoto Protocol. This means that Sweden's emissions of greenhouse gases will be allowed to comprise a maximum of 77.13 million tonnes per year on average for 2008-2012. Of these emission allowances, an average of approximately 22.3 million tonnes per year has been allocated within the EU emissions trading scheme.

Total greenhouse gas emissions in 2011 were 61.4 million tonnes of carbon dioxide equivalents in Sweden, see Table 1-1. The allocation for the trading sector was 22.7 million tonnes in 2011 and the emissions of the trading sector amounted to 19.8 million tonnes in 2011. The emissions from sectors outside the trading scheme were 41.6 million tonnes carbon dioxide equivalents. This means that the total emissions in Sweden in 2011 were around 17% below the assigned amount, including the net effect of allocation to the EU ETS and the carbon sink. This implies that Sweden will meet its commitment. The national surplus of Assigned Amount Units and Removal Units was around 13 million

tonnes in 2008, 2009 and 2011 and 11 million tonnes in 2010. Note that these figures are uncertain and only preliminary since the final calculations on target fulfilment will be made in 2014.

Table 1-1 Preliminary table on Sweden's greenhouse gas emissions in relation to the Kyoto Protocol and EU ETS

Mt CO₂-eq. (subm 2013)	2008	2009	2010	2011
Non EU ETS Emissions	43.3	41.8	42.8	41.6
EU ETS Verified emissions	20.1	17.5	22.7	19.8
Allowances EU ETS	20.8	22.1	23.5	22.7
Total emissions	63.4	59.3	65.5	61.4
Total emissions incl. net effect of EU ETS	64.1	63.9	66.3	64.3
Assigned amount incl LULUCF	77.13	77.13	77.13	77.13
Emissions incl. effect of EU ETS and LULUCF relative to Assigned Amount	17%	17%	14%	17%

1.1.5 National emission targets

The Swedish target according to the 2009 climate policy resolution of the Swedish Parliament is for greenhouse gas emissions by non-trading sectors to decrease by 40% or around 20 million tonnes between 1990 and 2020 (of which one third can be reduced by emission reductions in other countries). This decrease can be interpreted as a decrease of 33% between 2005 and 2020. Emissions from the non-trading sectors totalled 46.2 million tonnes in 2005 calculated with the same scope of the trading scheme as in 2008-2012 (including aviation), see Table 1-2. Non EU ETS emissions totalled 41.6 million tonnes in 2011, which is a decrease of 10% compared with 2005.

Table 1-2 Reduction of Sweden's emissions between 2005 and 2011

Mt CO₂-eq.	2005 (scope 2008-12)	2008	2009	2010	2011	2005-2011 (%)
Non EU ETS	46.2	43.3	41.8	42.8	41.6	-10
EU ETS	21.1	20.1	17.5	22.7	19.8	-6
Total emissions	67.3	63.4	59.3	65.5	61.4	-9

Furthermore, for the period 2008-2012, Sweden has set a national target of reducing greenhouse gas emissions with at least 4% compared with the 1990 level. This target will be achieved, since already the mean level for 2008-2011 is 14% below the 1990 level.

Sweden's commitment for the sectors outside the EU Emissions Trading Scheme according to the EU's Climate and Energy Package is that emissions have to decrease by 17% between 2005 and 2020 (trading period 2008-2012).

1.2 Institutional arrangements

Under Article 5 of the Kyoto Protocol each party in Annex 1 has to introduce a national system for estimating anthropogenic emissions by sources and removals by sinks of all greenhouse gases not controlled by the Montreal Protocol by 1 January 2007. The national system is to be designed in compliance with UNFCCC decision 20/CP.7. Under the terms of Decision No. 280/2004/EC of the European Parliament and of the Council, the national system has to be in place by the end of 2005. The national system has to ensure the function of all the institutional, legal and procedural arrangements required to calculate emissions and removals of greenhouse gases.

The Swedish national system came into force on 1 January 2006 and its aim is to ensure that climate reporting to the secretariat of the Convention (UNFCCC) and the European Commission complies with specified requirements. This means, among other things,

- estimating and reporting anthropogenic GHG emissions and removals in accordance with the Kyoto Protocol,
- assisting Sweden in meeting its commitments under the Kyoto Protocol,
- facilitating the review of submitted information,
- ensuring and improving the quality of the Swedish inventory and
- guaranteeing that submitted data is officially approved.

The national system ensures annual preparation and reporting of the national inventory and of supplementary information in a timely manner and that the inventory fulfills all quality criteria, i.e. is transparent, accurate, consistent, comparable and complete.

The KP-reporting of LULUCF uses the same institutional arrangements, national system and corresponding QA/QC procedures as for the UNFCCC reporting.

1.2.1 Legal arrangements

Ordinance (2005:626) Concerning Climate Reporting provides the basis for the Swedish national system and describes the roles and responsibilities of the government agencies in the context of climate reporting. Through this, sufficient capacity for timely performance is ensured.

There are legislations in Sweden which is not primarily intended to apply to climate reporting but indirectly supports the work by providing a basis for the estimation of greenhouse gas emissions and removals. Under Chapter 26 Section 19 of the Environmental Code (1998:808), there is an obligation for annual environmental reports to be submitted for certain environmentally hazardous activities so that government agencies can undertake supervision.

The General Statistics Act (SFS 2001: 99) and the associated Ordinance (2001:100) Concerning Official Statistics impose an obligation on companies and other organizations to submit annual data. The data then serve as a basis for estimating greenhouse gas emissions and removals in several sectors.

According to Directive 2003/87/EC and national Act (2004:1199) on emission trading, emission data for plants included in the emission trading system should be reported annually. These data are used as a supplementary source within this greenhouse gas inventory.

1.2.2 Institutional arrangements

Sections 6-19 of the Ordinance (2005:626) Concerning Climate Reporting describe the tasks of the government agencies in the context of the yearly inventory and reporting activity. The illustration in Figure 1-1 and Table 1-3 and the associated text below describe in broad terms which organizations are involved in the work of compiling documentation for the yearly inventory report and for other reporting to the European Commission and the UNFCCC. Depending on the role of the government agencies in climate-reporting activity, this responsibility may range for example from supplying data and producing emission factors/calorific values to carrying out calculations to estimate emissions or conducting a national peer review (red). In addition to what is described in the Ordinance, the Swedish Environmental Protection Agency (Swedish EPA) engages the SMED consortium as consultants with expert skills to conduct the inventory and reporting in the area of climate change.

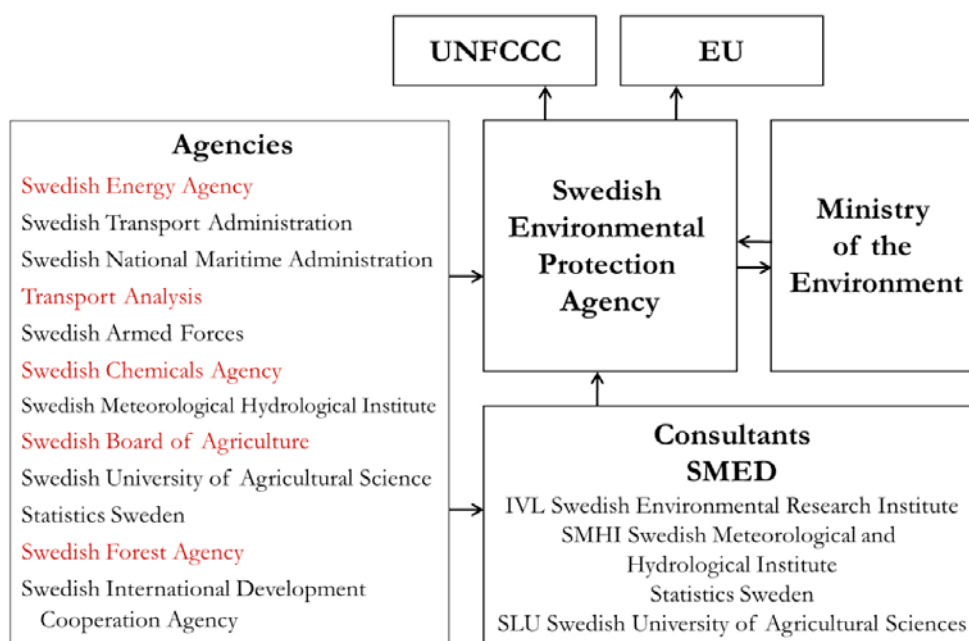


Figure 1-1 The Swedish national system.

1.2.2.1 SINGLE NATIONAL ENTITY

The Swedish Ministry of Environment is the single national entity and has overall responsibility for the inventory.

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UNFCCC focal point

Ms. Nilla Thomson: nilla.thomson@regeringskansliet.se

Responsible for reporting to EU

Ms. Stina Gustafsson: stina.gustafsson@regeringskansliet.se

1.2.2.2 SWEDISH EPA RESPONSIBILITIES

The Swedish EPA is responsible for co-ordinating the activities for producing the inventory, maintaining the reporting system and also for the final quality control and quality assurance of the inventory.

The Swedish EPA sends the inventory to Ministry of the Environment and – on behalf of the Ministry of Environment – submits the inventory to the EU and to the UNFCCC. Finally, the Swedish EPA is responsible for national publication of the greenhouse gas inventory.

The National inventory compiler at the Swedish EPA is Ms. Maria Lidén.

1.2.2.3 AGENCIES RESPONSIBILITIES

Agencies responsibilities according to Ordinance (2005:626) Concerning Climate Reporting is described in Table 1-3 below.

1.2.2.4 THE SMED CONSORTIUM

The Swedish EPA engages consultants with expert skills to conduct the inventory and reporting in the area of climate change. During the spring of 2005, the Swedish EPA completed a negotiated procurement of services under the terms of the Public Procurement Act. After procurement had been completed, a framework contract was signed with the consortium Swedish Environmental Emissions Data (SMED)², consisting of the Swedish Meteorological and Hydrological Institute (SMHI), Statistics Sweden (SCB), the Swedish University of Agricultural Sciences (SLU) and the Swedish Environmental Research Institute (IVL). The contract between the Swedish EPA and SMED runs for nine years and thus covers the whole first commitment period under the Kyoto Protocol.

² <http://www.smed.se/>

SMED receives data and documentation from responsible authorities as described above and produces most of the data and documentation in the Swedish inventory. The regular inventory work is organized as a project involving all SMED organizations. The project is run by a project management team with one person from each organization. The Swedish Meteorological and Hydrological Institute is main responsible for production of gridded emission data. Statistics Sweden is main responsible for the energy sector, the agriculture sector and parts of the waste sector, but is also involved in industrial processes since these are closely connected to the energy sector. The Swedish University of Agricultural Sciences is responsible for the LULUCF sector. The Swedish Environmental Research Institute is main responsible for the industrial process sector, the solvents and other products use sector and also parts of the waste sector and energy sector.

On behalf of the Swedish EPA, SMED also conducts development projects necessary for improving the inventory.

Table 1-3. Agencies responsibilities according to Ordinance (2005:626) Concerning Climate Reporting

Sector	Data and documentation provided by	Peer review conducted by	Other responsibilities
Energy	Swedish Energy Agency the Swedish Transport Administration, the Swedish Transport Agency, the National Maritime Administration the Swedish Armed Forces	Swedish Energy Agency (energy sector excluding transports) Transport Analysis (transports)	The Swedish Energy Agency also assists the Swedish EPA in the work of developing documentation concerning flexible mechanisms and emissions projections as well as extracts from and information about changes to the national register.
Industrial Processes	Swedish Chemicals Agency (fluorinated greenhouse gases)	The Swedish EPA (CO ₂ , CH ₄ and N ₂ O) Swedish Chemicals Agency (fluorinated greenhouse gases)	
Solvents and Other Product Use	Swedish Chemicals Agency	Swedish Chemicals Agency	
Agriculture	Swedish Board of Agriculture Statistics Sweden	Swedish Board of Agriculture	
Land Use, Land-Use Change And Forestry Sector	Swedish University of Agricultural Sciences, Statistics Sweden, the Swedish Forest Agency, the Swedish Meteorological and Hydrological Institute (SMHI) and the Swedish Board of Agriculture	Swedish Forest Agency Swedish Board of Agriculture (agriculture related parts)	
Waste		Swedish EPA	
Reporting relating to efforts in developing countries			The Swedish International Development Cooperation Agency (Sida) is responsible for presenting documentation to the Swedish EPA.

1.3 Inventory planning, preparation and management

The present Swedish greenhouse gas inventory and KP-LULUCF inventory was compiled according to the recommendations for inventories set out in the UNFCCC reporting guidelines according to Decision 18/CP.8, the Common Reporting Format (CRF), Decision 13/CP.9, the new CRF for the Land Use Change and Forestry Sector, the IPCC 1996 Guidelines for National Greenhouse Gas Inventories, which specify the reporting obligations according to Articles 4 and 12 of the UNFCCC (IPCC Guidelines, 1996) as well as the IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories (IPCC GPG, 2000) and the IPCC Good Practice Guidance for Land Use, Land-Use Change and Forestry (IPCC GPG-LULUCF, 2003).

It could be noted that the greenhouse gas inventory is integrated with the inventory of air pollutants for reporting to the UNECE (CLRTAP). This assures effective use of resources and consistency between the reporting to the UNFCCC and to the CLRTAP.

1.3.1 Quality system

In order to fulfill the obligations of reporting to the UNFCCC and the EU, the Swedish EPA has set up a quality system as part of the national system. The structure of the quality system follows the PDCA cycle (Plan, Do, Check, Act) illustrated in figure 1-2 below. This is an adopted model for how systematic quality and environmental management activity is to be undertaken according to international standards to ensure that quality is maintained and developed.



Figure 1-2 Structure of the quality system.

The quality system includes several procedures such as training of staff, inventory planning and preparation, QA/QC procedures, publication, data storage, and follow-up and improvements. All QA/QC procedures are documented in a QA/QC plan³. The QA/QC plan also includes a scheduled time frame describing the different stages of the inventory from its initial development to final reporting. The quality system ensures that the inventory is systematically planned, prepared and followed up in accordance with specified quality requirements so that the inventory is continuously developed and improved.

The responsibilities of the Swedish EPA and the other government agencies for the quality system are described in Ordinance (2005:626) Concerning Climate Reporting. Under Section 3, the Swedish EPA and other government agencies which take part in the climate-reporting work have to ensure that the methodologies applied in the reporting and inventories of emissions and removals attain the quality required for it to be possible for Swedish climate reporting to be done in the correct manner and with correct information.

The government agencies have to have internal routines to plan, prepare, check and act/follow up the quality work and consult one another with the aim of developing and maintaining a coordinated quality system.

The responsibility of SMED to maintain and develop an internal quality system is described in the framework contract between the Swedish EPA and the consultants. The SMED quality system is described in a detailed manual including several appendices.⁴ It is updated annually and lists all quality control steps that must be undertaken during inventory work (Tier 1 and where appropriate Tier 2). It also includes descriptions of roles and responsibilities, of databases and models, work manuals for each CRF category and documented procedures for uncertainty and key source analyses, as well as procedures for handling and responding to UNFCCC's review of the Swedish inventory. It also handles follow-up and improvement by procedures of non-conformity reporting and collection of improvement needs from all stages of the annual inventory cycle. This results in a planning document, which is used as a basis for planning and selecting further actions to improve the inventory.

The illustration in Figure 1-3 below shows a process description of the annual Swedish inventory.

³ Swedish EPA, National Greenhouse Gas and Air Pollutants Inventory System in Sweden

⁴ Manual for SMED:s Quality System in the Swedish Air Emission Inventories, available at www.smed.se

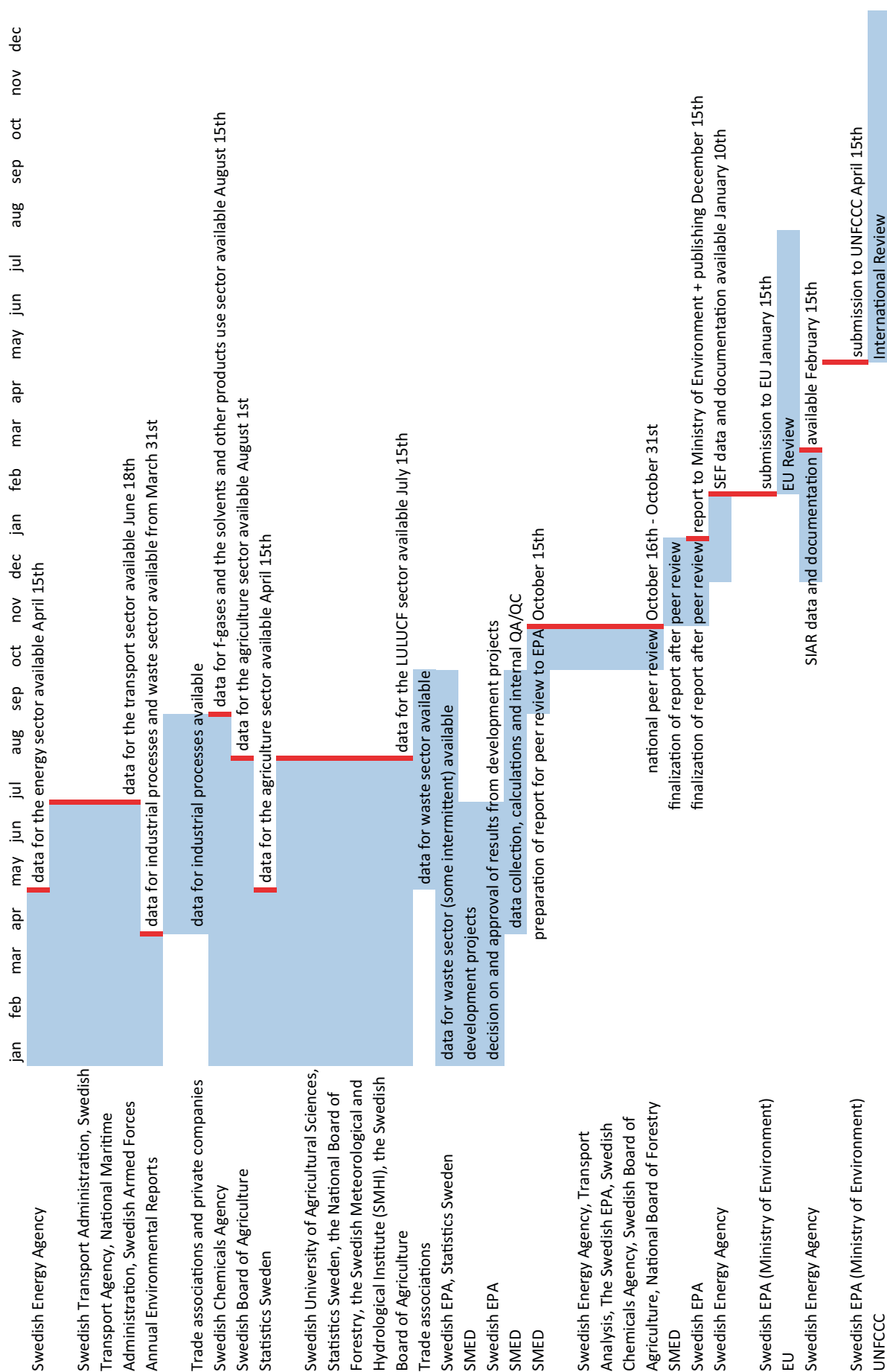


Figure 1-3 Overview of inventory planning, preparation and management.

1.3.2 Training, awareness and skills

To meet quality criteria in the IPCC and UNFCCC guidelines, experts from different government agencies are participating in the inventory according to SFS 2005:626. By involving these agencies, it is ensured that the best expertise available in the country is involved.

Skills on the part of SMED are ensured in accordance with the requirements laid down in the framework contract between the Swedish EPA and the consultants. The levels of consultant's skills are continuously reviewed.

1.3.3 Inventory planning (PLAN)

Planning of the inventory for submission in year x starts in the fall of year x-2 when the Swedish EPA gets the preliminary budget for year x-1. General priorities for the coming year are set by the Swedish EPA based on

- recommendations from international review not yet implemented in the inventory
- recommendations from national peer review not yet implemented in the inventory
- key category analysis (focus on major sources/sinks)
- uncertainty analysis (focus on sources/sinks that contributes significantly to the uncertainty of the inventory)
- ideas from SMED and the Swedish EPA on how to improve quality and effectiveness of the inventory
- new international and national requirements, decisions and guidelines

Priorities are distributed to SMED approximately in October.

Based on the priorities and on detailed information in the list on suggestions on improvements (see section 1.3.5.5 below), SMED compiles a list of suggested development projects for the coming years. The list of suggested development projects is discussed between SMED and the Swedish EPA. During the winter the Swedish EPA decides on what projects should be performed.

In January-June (approximately) SMED is working with development projects. Reports on the results and recommendations for implementation in the inventory are delivered to the Swedish EPA who then decides how these new methods/activity data/emission factors should be implemented in the inventory. In order to be able to implement results in the current inventory with sufficient QA/QC, the Swedish EPA has to decide on implementation in June.

From time to time, there is a need to change data provided by responsible authorities as discussed above. The Swedish EPA each year contacts responsible authorities and discusses needs for updates.

1.3.4 Inventory preparation (DO)

SMED gather data and information from various government agencies, organizations and companies over the period from April to August with the aim of being able to carry out emission calculations. The calculations are performed in models, statistics programs and calculation programs in April to September. Over the period from September to October, the material is put together in a reporting format. A short description of data collection and processing for each sector is provided below. See sections 3-8 for a detailed description. Preparation of the inventory is documented in detailed work documentation, which serves as instructions for inventory compilers to ensure quality and consistency, and also serves as information in the national peer review process.

1.3.4.1 ENERGY- STATIONARY COMBUSTION

Energy industries: Data from quarterly fuel statistics, a total survey conducted by Statistics Sweden at plant level and by fuel type. For some petroleum refining plants, data from the European Union Emission Trading Scheme (ETS) is used.

Manufacturing industries: Data mainly from the quarterly fuel statistics, a sample survey conducted by Statistics Sweden. In some cases data from the industrial energy statistics or ETS is used as a complement. All data is at plant level and by fuel type.

Other sectors: Data from official statistical reports prepared by Statistics Sweden at national level and by fuel type.

Activity data is multiplied by thermal values, mainly from Statistics Sweden, and emission factors provided by the Swedish Energy Agency and the Swedish EPA.

1.3.4.2 ENERGY- MOBILE COMBUSTION

Data on fuel consumption at national level and by fuel type is collected from Statistics Sweden and used in combination with emissions data and fuel data from the National Road Administration, the National Rail Administration, the Civil Aviation Administration and the Swedish Military. Activity data is multiplied by thermal values, mainly provided by Statistics Sweden, and emission factors provided by the responsible authorities.

1.3.4.3 ENERGY – FUGITIVE EMISSIONS

For flaring in refineries and chemical industries, activity data and CO₂ emissions from ETS are used for 2005 and later. In earlier years, data was collected through personal contacts with the facilities. Activity data and CO₂ emissions from hydrogen production in oil refineries are taken from ETS and reported under CRF 1B2a1 in line with 2006 IPCC Guidelines. For non-CO₂ emissions, regular emission factors for stationary combustion are used.

Activity data for transfer losses of gas works gas are taken from the annual energy balances. Emission factors for stationary combustion are used.

Fugitive emissions from refineries and from storage of petroleum products at storage depots are mainly compiled from the facilities' environmental reports. Estimates of fugitive emissions from gasoline stations are calculated from fuel data provided by the National Road Administration.

1.3.4.4 INDUSTRIAL PROCESSES

The reported data for industrial processes is mainly based on information from environmental reports. According to Swedish environmental legislation, operators performing environmentally hazardous activities that require a permit by law are required to compile and send an annual environmental report to their supervisory authority. The County Administrative Boards audit the data from the operators' environmental reports.

The data in the environmental reports refer to emissions derived from plant specific measurements or estimates such as mass balances. The use of default emission factors is limited.

In some cases, when there are a large number of smaller companies within a specific sector, and all the environmental reports are not available, a combination of information available from environmental reports and production statistics at national level is used to estimate national emissions. Emission factors used are usually derived nationally based on available information from some facilities in a specific sector, and applied to the national level. The use of default emission factors is limited.

For most CO₂ emissions from industrial processes, activity data on e.g. the produced amount of clinker, limestone, etc. is collected directly from the operators. In some cases data on CO₂ emissions from the European trading scheme is used for 2005 and later years. Activity data on fuels used in CO₂ emitting processes are collected from the same surveys as those used for energy emissions for manufacturing industries, as described above.

Emissions of fluorinated greenhouse gases are estimated based on national import and export statistics from the Swedish Chemicals Agency.

1.3.4.5 SOLVENT AND OTHER PRODUCT USE

Data used for estimating emissions from solvent and other product use are based on emission factors and national activity data obtained from the Products Register kept by the Swedish Chemicals Agency.

1.3.4.6 AGRICULTURE

Data on animal numbers, crop areas, yields, sales of manure, manure management and stable periods are taken from official statistical reports published by the Swedish Board of Agriculture and Statistics Sweden. Some complementary information is collected from organisations and researchers, such as the Swedish Dairy Association, Swedish Poultry Meat Association, SLU and the Swedish Institute of Agricultural and Environmental Engineering.

1.3.4.7 LAND USE, LAND USE CHANGE AND FORESTRY

Estimates presented in the LULUCF sector are mainly based on data from the SLU. The SLU is responsible for the National Forest Inventory, which focuses on living biomass, and for the Swedish Forest Soil Inventory, that focuses on dry organic matter and on soil organic carbon. The two inventories are integrated and use the same infra-structure for the field sample.

1.3.4.8 WASTE

Statistics on deposited waste quantities, methane recovery and nitrogen emissions from wastewater handling, are provided by the Swedish Association of Waste Management (Avfall Sverige, former RVF), Statistics Sweden, the Swedish Forest Industries Federation and the Swedish EPA. If new data on organic content in household waste or other relevant research is published, such reports are also considered. Profu, an independent research and consultant company in the areas of energy, environment and waste management, provides estimates of deposited organic fractions of industrial waste.

Emissions reported for waste incineration are compiled from the facilities' annual environmental reports.

1.3.5 QA/QC procedures and extensive review of GHG inventory and KP-LULUCF inventory (CHECK)

1.3.5.1 QUALITY CONTROL

Quality control is the check that is made during the inventory on different types of data, emission factors and calculations that have been made. The quality control takes place according to general requirements (Tier 1) which apply to all types of data used as support material for the reporting, and specific requirements for quality control (Tier 2) which are applied to certain types of data and/or emission sources. In this inventory, general Tier 1 QC measures, according to Table 8.1 in IPCC Good Practice Guidance (2000), have been carried out as follows:

- Transcription errors in data input
- Calculations are made correctly
- Units and conversion factors are correct
- Integrity of database files
- Consistency in data between source categories
- Correct movement of inventory data between processing steps
- Recalculations, checked and documented
- Completeness check
- Comparison of last submission's estimates to previous estimates
- Documentation of changes that may influence uncertainty estimates

In addition, source specific Tier 2 QC procedures are carried out for several categories (Table 1-4).

All QC measures performed are documented by SMED in QC checklists for each CRF code or group of codes. After completion of the initial compilation of the inventory, a QC-team within SMED reviews all QC checklists. In addition, the project management team performs checks of submission data using the functionality of the CRF Reporter (i.e. checks of completeness, time-series consistency and recalculation explanations).

Table 1-4. Source specific Tier 2 QC procedures carried out in the inventory.

CRF		Action
1.A, 1.B and parts of 2	Energy amounts and emissions of CO ₂	Analysis of differences between the sectoral and reference approach. In order to check activity data and EF, several quality control projects have been carried out over time comparing the inventory data with information from environmental reports and EU ETS data.
1.B	Fugitive emissions and flaring of CO ₂ , CH ₄ and N ₂ O	Measured emissions from flaring are checked to assure that the quality is sufficiently high. Trends for activity data and emissions are compared and analysed.
2.A.1	Cement production, process emissions of CO ₂	Emissions are calculated both using the bottom-up and the top-down method, the results have been compared and differences explained. It is also stated that emission factors and activity data used are in accordance with internationally accepted methods.
2.A.2	Lime production, process emissions of CO ₂	Emissions are calculated using both the bottom-up and the top-down method, the results have been compared and differences explained.
2.B.2	N ₂ O-emissions from Nitric Acid production	Bottom-up production data could not be compared to official data since official data were not available in the statistical database. Only one company produces nitric acid. Calculation methods, abatement technique and production capacity is based on information achieved directly from the company.
2.C.1	Iron and steel production	Activity data are checked with fuel combustion data in order to avoid double counting of emissions or omissions. Activity data is also compared to trade statistics. IEF are compared to IPCC default values.
2.C.3	PFC emissions from aluminium production	Documented process information obtained directly from the company enable plant-specific data checks.
2.F	Consumption of halocarbons and SF ₆	Differences between country specific emission factors and default emission factors from IPCC Guidelines are documented.

When the reporting tables and the NIR are completed by SMED, a quality coordinator performs the final quality control before delivery of the inventory to the Swedish EPA.

1.3.5.2 QUALITY ASSURANCE

Key categories should be subject to external peer review according to the Tier 2 of the Good Practice Guidance. The Swedish QA/QC system includes national peer reviews by sectoral authorities. The peer review is defined in the Ordinance (2005:626) Concerning Climate Reporting and is, for all sectors, conducted by a person who has not taken part in the inventory preparation. The Swedish EPA is responsible for coordinating the annual peer review. This means, among other things, ensuring that the peer reviewers have received the necessary training.

The peer review includes methodology and emissions factors used, as well as comparisons of activity and emission data with other national statistics. The reviewers also identify areas for improvement, which consolidates the basis for improvements in coming submissions. Results from the national peer review are documented in review reports. Recommendations from the review reports are collected to the list of suggested improvements described in section 1.3.5.5.

The UNFCCC secretariat administers an international peer review of Swedish reporting after submission. Recommendations from the review reports are collected to the list of suggested improvements described in section 1.3.5.5. See also section 10.

The submission will also be reviewed by the EU. Recommendations from this review will be handled in the same way as recommendations from the UNFCCC review and the national peer review.

1.3.5.3 FINALIZATION, PUBLICATION AND SUBMISSION OF THE INVENTORY

The Swedish Environmental Protection Agency delivers the greenhouse gas inventory and the KP LULUCF inventory to the Ministry of Environment 20 working days before the reporting date, which is in mid-December. At the same time, the inventory is published nationally⁵.

The Swedish EPA, on behalf of the Ministry of Environment, submits the inventory to the European Commission on January 15th and to the UNFCCC on April 15th. Reported data in the submission of year t relates to emissions year $t-2$, in other words emissions which took place during 2011 are reported in early 2013.

1.3.5.4 DATA STORAGE

A system for handling emission data, entitled Technical Production System (TPS)⁶, has been developed and was implemented for the first time in submission 2007. It supports data input from text files and Microsoft Excel sheets, and provides different types of quality gateways. For instance the system makes it possible for multiple users such as the SMED consortium and the national independent reviewers to view data, plot time series and make comparisons between different years and submissions. For all CRF categories and sub-categories, time series from 1990 onwards of emission data, activity data, and implied emission factors where relevant can be presented. The system also allows for different types of data output, e.g. to the CRF Reporter or to MS Excel. Finally, TPS is used for data archiving of each submission. For access to the TPS, login with password is requested.

The CRF-tables were generated using the export function in CRF Reporter.

⁵ www.naturvardsverket.se

⁶ <https://tps.naturvardsverket.se/>

In addition to TPS, documentation, data and all calculations for each submission are stored at each organizations servers and, for collective use and archiving, at two projects at Projectplace⁷. One project is for documents shared between Swedish EPA, other involved agencies and SMED and the other project is primarily for SMEDs use however the Swedish EPA also has access to the project. At Projectplace, all documents are stored in versions, in other words when documents are changed a new version is automatically created. This function ensures that important information is not lost and facilitates backtracking of changes. Login with password is requested for access to projects at Projectplace.

1.3.5.5 FOLLOW-UP AND IMPROVEMENT (ACT)

Each year, all comments received from national and international reviews that are not already addressed and also ideas from SMED and the Swedish EPA are compiled into a list for suggestions on improvements. From this list, development projects are formed each year as describes in section 1.3.3. All suggestions not implemented one year is kept on the list for next year.

Each year, the Swedish EPA follows up on delivered data from responsible agencies to ensure correct and appropriate data for next submission.

Development of TPS such as additional functions etc is organized in a similar way as for the inventory: Ideas are compiled into a list, and from this list issues to be implemented are prioritized.

1.4 Brief general description of methodologies and data sources used

1.4.1 GHG inventory

Emission estimates are mainly based on activity data from national or official Swedish statistics, e.g. energy statistics, European Union Emission Trading Scheme (EU ETS)⁸, environmental reports⁹, agricultural and forestry statistics, as well as data on production (e.g. cement) and consumption (e.g. fluorinated gases: F-gases) obtained directly from the major producers and consumers, respectively.

Emission factors and thermal values used are either developed nationally or are internationally recommended default factors.

The methodologies used for Sweden's greenhouse gas emissions inventory are in accordance with the Revised 1996 IPCC Guidelines for National Greenhouse Gas Inventories (IPCC Guidelines)¹⁰ and, in general, in line with IPCC's Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories (Good Practice Guidance)¹¹ and IPCC's Good

⁷ www.projectplace.com

⁸ See Annex 8.1

⁹ See Annex 8.3

¹⁰ The IPCC Guidelines can be found at: <http://www.ipcc-nggip.iges.or.jp/public/gl/invs1.htm>

¹¹ The Good Practice Guidance can be found at: <http://www.ipcc-nggip.iges.or.jp/public/gp/english/>

Practice Guidance for Land Use, Land-Use Change and Forestry (GPG-LULUCF)¹². Some parts of the methodologies are taken directly from the IPCC Guidelines, the Good Practice Guidance and the EMEP/EEA air pollutant emission inventory guidebook (formerly called the EMEP CORINAIR emission inventory guidebook).¹³ Information from the 2006 IPCC Guidelines for National Greenhouse Gas Inventories (2006 IPCC Guidelines)¹⁴ is used in some parts of the inventory.

In Table 1.5, all Tier methods used, which differ from Tier methods recommended in IPCC Guidelines or Good Practice Guidance, are presented. There is also a brief explanation of why the recommended methods have not been used. Note that for sectors where no specific recommendations are made in the IPCC Guidelines or Good Practice Guidance, these sectors are not included in Table 1.5. For an overview of the methods used in all sectors, see Summary 3 in the CRF tables and in each sector section, where a more detailed explanation on data sources and methodologies is given.

Table 1-5. Methods used that differ from recommended methods in the IPCC Guidelines or Good Practice Guidance for all sectors.

Sector	Used method Tier	IPCC Guidelines method Tier	Explanation
Energy:Emissions of CH ₄ and N ₂ O from navigation	1	2	Reliable data required for Tier 2 is currently not available (various engine types etc).
Industrial processes: Emissions of PFC from aluminium production	2	3	No measurements are performed, so Tier 3 cannot be applied. The method used is Tier 2.
Industrial processes: Emissions of SF ₆ from electrical insulation	2a	3	There is not enough information available to perform Tier 3.
Industrial processes: Semiconductor manufacture	1	2	There is not enough information available to perform Tier 2.
Waste:N ₂ O from waste water	National	1	Sweden uses national statistics on nitrogen emissions. Nitrogen emissions are only model calculated for the rural population.

SMED has carried out the calculations. In a few cases, estimates are based on expert judgements.

The combined effect of various greenhouse gases has been calculated using global warming potential factors (GWP). These are developed by the IPCC and are used as a means of comparing the relative significance of various gases in terms of their greenhouse effect, expressed as carbon dioxide equivalents.

Emission factors and thermal values for the energy sector are provided in Annex 2.

¹² The GPG-LULUCF can be found at: <http://www.ipcc-nggip.iges.or.jp/public/gpglulucf/gpglulucf.html>

¹³ The EMEP/Corinair Guidebook can be found at: <http://tfeip-secretariat.org/unece.htm>

¹⁴ The 2006 IPCC Guidelines can be found at: <http://www.ipcc-nggip.iges.or.jp/public/2006gl/index.html>

1.4.2 KP-LULUCF inventory

The same base methodology, emission factors and data sources is used for the reporting of LULUCF under the KP as for the reporting under UNFCCC.

Data from the Swedish National Forest Inventory (NFI) have been used for developing the land use matrix and is consistent with the data used for developing the land use matrix under the UNFCCC-reporting. The main difference is that activities are reported under the KP while land use categories are reported under the UNFCCC.

The carbon pool changes associated to the activities reported under the Kyoto protocol is estimated in exactly the same way as under the UNFCCC reporting, using the stock change method and area based sampling for most of the carbon pools. However, the living biomass is reported separately for above-ground and below-ground biomass, respectively, and the Dead organic matter is reported separately for Litter and Dead wood.

1.5 Brief description of key categories, including for KP-LULUCF key categories

1.5.1 GHG inventory (including and excluding LULUCF)

According to Good Practice Guidance, key categories in a national inventory should be identified in order to prioritize the efforts in improving the quality of the inventory estimates. Key categories are defined as sources and/or sinks that have “a significant influence on a country’s total inventory of direct greenhouse gases in terms of the absolute level, the trend, or the uncertainty in emissions and removals”. The identification of level and trend key categories is done in two tiers, tier 1 assessment and tier 2 assessment. The two tier assessments differ in the sense that the tier 2 assessment includes information of uncertainties. According to Good Practice Guidance, the results from the tier 2 assessment should be utilized if the tier 2 results differ from the tier 1 results. Hence, in the Swedish inventory, the results from the tier 2 assessment is the basis for applying and describing higher tier methodologies and when prioritizing efforts in improving the quality of the inventory estimates.

The resulting tier 1 and tier 2 key categories are presented in CRF table 7 and Table 1-6. In Annex 1 the methodology is discussed in detail and corresponding background tables, according to tables 7.A1 - 7.A3 of the Good Practice Guidance are presented.

As can be seen in Table 1-6 most tier 2 key categories are part of the tier 1 assessment. A few additional categories are however identified as key due to their high uncertainty, e.g. N₂O from CRF 1.A.4.b (Residential) and CH₄ from CRF 1.B.2 (Oil and Natural Gas). Note that it is common that the tier 2 assessment render fewer key categories than the tier 1 assessment.

Table 1-6. Tier 1 and tier 2 key categories 2011 in terms of level and trend

IPCC Source Category	GHG	Including LULUCF		Excluding LULUCF	
		Tier 1	Tier 2	Tier 1	Tier 2
1.AA.1.A (Public Electricity and Heat Production)	CH ₄			T	T
1.AA.1.A (Public Electricity and Heat Production)	CO ₂	LT	LT	LT	LT
1.AA.1.A (Public Electricity and Heat Production)	N ₂ O	LT	T	LT	LT
1.AA.1.B (Petroleum Refining)	CO ₂	LT	LT	LT	LT
1.AA.1.C (Manufacture of Solid Fuels and Other Energy Industries)	CO ₂	LT		LT	
1.AA.2.A (Iron and Steel)	CO ₂	LT		LT	
1.AA.2.C (Chemicals)	CO ₂	LT		LT	LT
1.AA.2.D (Pulp, Paper and Print)	CO ₂	LT		LT	T
1.AA.2.E (Food Processing, Beverages and Tobacco)	CO ₂	LT		LT	
1.AA.2.F (Other Manufacturing Industries and Construction)	CO ₂	LT	L	LT	L
1.AA.2.F (Other Manufacturing Industries and Construction)	N ₂ O			L	L
1.AA.3.A (Civil Aviation)	CO ₂	L		L	
1.AA.3.B (Road Transportation)	CH ₄			T	T
1.AA.3.B (Road Transportation)	CO ₂	LT	LT	LT	LT
1.AA.3.C (Railways)	CO ₂			LT	
1.AA.3.D (Navigation)	CO ₂	T		LT	
1.AA.3.E (Other Transportation)	CO ₂			LT	
1.AA.4.A (Commercial/Institutional)	CO ₂	LT	T	LT	LT
1.AA.4.B (Residential)	CH ₄		LT	L	LT
1.AA.4.B (Residential)	CO ₂	LT	LT	LT	LT
1.AA.4.B (Residential)	N ₂ O				L
1.AA.4.C (Agriculture/Forestry/Fisheries)	CH ₄				T
1.AA.4.C (Agriculture/Forestry/Fisheries)	CO ₂	LT		LT	L
1.AA.5.B (Military Use)	CO ₂	T		T	T
1.B.2 (Oil and Natural Gas)	CH ₄		LT		LT
1.B.2 (Oil and Natural Gas)	CO ₂	LT	T	LT	LT
2.A.1 (Cement production)	CO ₂	LT		LT	L
2.A.2 (Lime Production)	CO ₂	LT		LT	
2.B.2 (Nitric Acid Production)	N ₂ O	T		T	T
2.C.1 (Iron and Steel Production)	CO ₂	LT	LT	LT	LT
2.C.2 (Ferroalloys Production)	CO ₂			T	
2.C.3 (Aluminium production)	PFC			T	T
2.F.1 (Refrigeration and Air Conditioning Equipment)	HFC	LT	LT	LT	LT
4.A (Enteric Fermentation)	CH ₄	LT	LT	LT	L
4.B (Manure Management)	CH ₄	T		LT	
4.B (Manure Management)	N ₂ O	L	L	LT	LT
4.D.1 (Direct Soil Emissions)	N ₂ O	LT	LT	LT	LT
4.D.2 (Pasture, Range and Paddock Manure)	N ₂ O	L	LT	LT	LT
4.D.3 (Indirect Emissions)	N ₂ O	L	L	LT	LT
4.D.4 (Agricultural Soils. Other)	N ₂ O	LT	LT	LT	LT
5.A (Forest Land)	CO ₂	LT	LT		
5.B (Cropland)	CO ₂	LT	LT		
5.B (Cropland)	N ₂ O		T		
5.C (Grassland)	CO ₂	T	T		
5.E (Settlements)	CO ₂	LT	LT		
6.A (Solid Waste Disposal on Land)	CH ₄	LT	LT	LT	LT
6.B (Wastewater Handling)	CH ₄		LT	L	LT

L Level. T Trend.

1.5.2 KP-LULUCF inventory

The key category assessment for KP-LULUCF is found in section 11.6.1.

Carbon dioxide emissions for land use categories Forest land, Cropland, Grassland and Settlements are considered key categories under the UNFCCC. Emissions under categories 5I, 5IV and 5V are never identified as key categories for any gas under the UNFCCC. Under the UNFCCC, 5III N₂O emissions from land use conversions to Cropland was identified as a key-category for trend including uncertainty.

Activities Forest management, Afforestation/ Reforestation and Deforestation were considered key-categories (CO₂). Part of 5III N₂O emissions from land use conversions to Cropland corresponds to D from Forest to Cropland. However, this category was not identified as key-category because most of the emissions under the UNFCCC refer to land use conversions from Grasslands to Cropland and due to a “conservative” high assumed uncertainty (100%). Every key category is estimated using higher tiers.

1.6 Information on QA/QC

See section 1.3.

1.6.1 QA/QC Procedures

See section 1.3.5.

1.6.2 Verification activities

See section 1.3.5.

1.6.3 Treatment of confidentiality issues

In the inventory, several data sources are confidential at micro level (e.g. plant level). This is for example the case for statistical surveys of fuel consumption used in Energy (CRF 1) and data from the Products Register at the Swedish Chemicals Agency used in Solvent and other product use (CRF 3). Results published in the inventory are aggregated, and because of this no confidentiality issues remains in the CRF or in the NIR.

1.7 General uncertainty evaluation

1.7.1 GHG inventory

An uncertainty analysis has been performed according to the Tier 1 method, described in detail in Annex 7 and Good Practice Guidance section 6.3.2. The analysis has been performed both including LULUCF and excluding LULUCF. According to the IPCC Guidelines, uncertainty estimates are an essential part of an emission inventory. They should be derived for each variable used in the inventory (measured emissions, activity data and emission factors) and

aggregated into uncertainty estimates in total national emissions and emission changes over time (trends). The 2006 IPCC Guidelines identify that: “An uncertainty analysis should be seen, first and foremost, as a means to help prioritise national efforts to reduce the uncertainty of inventories in the future, and guide decisions on methodological choice”.

During 2005, a SMED study was performed, aiming at improving the transparency and quality in the present uncertainty estimates in the Swedish National Greenhouse Gas Inventory by making the underlying documentation and structures for uncertainty estimates more consistent and traceable. This will facilitate easier replication and updating of results as well as enable internal and external reviews of assigned uncertainties. To simplify the methodology, there have not been any adjustments for correlation between gases, even though many of them have the same activity data and therefore are correlated. The study is described in Annex 7 and in detail in a SMED report.¹⁵

The Good Practice Guidance Tier 1 method is based on emission estimates and uncertainty coefficients for activity data and emission factors. The analysis was done for the sectors Energy, Industrial Processes, Solvent and Other Product Use, Agriculture, LULUCF and Waste. Uncertainty coefficients have in many cases been assigned based on expert judgement or on default uncertainty estimates provided in the Good Practice Guidance, since not enough background data was available to make actual statistical uncertainty calculations. Hence, care should be taken when interpreting and assessing the uncertainty results.

Uncertainty estimates have been performed for the base year 1990 and 2011 for direct greenhouse gases, e.g. CO₂, CH₄, N₂O and F-gases and are presented as 95% confidence intervals.

When reporting the results in the NIR, uncertainties are presented on the same aggregation level as the key categories. The purpose is to facilitate combined use of the two analyses, since both aims at showing what parts of the inventory are especially important and/or weak. This is very important information when planning future inventories and, above all, using and evaluating the inventory results.

The 2005 study did not include improvement of single uncertainties, for instance by contacting external experts for better information on uncertainties on different sources. Further work considering uncertainties will focus on such improvements. During each development project, uncertainties in estimated activity data and emission factors are overhauled and revised when needed.

In conjunction with the tier 1 uncertainty calculations of emission levels for 1990 and 2011, uncertainty introduced to the trend 1990-2011 is calculated following the IPCC tier 1 method.

¹⁵ Gustafsson, 2005

1.7.1.1 RESULTS

The results of the uncertainty calculations according to the tier 1 uncertainty approach are presented in Annex 7. The overall uncertainty for 2011 GHG emissions (in CO₂ equivalents) in Sweden is calculated to be $\pm 4.5\%$, excluding LULUCF (Figure 1-4). The overall uncertainty largely stems from uncertainty in the agricultural sector (CRF 4). The national estimated uncertainty neither include corrections for the correlation that may exist between gases (i.e. based on the same activity data), nor include corrections for non-reported sources. Therefore, the actual uncertainty of the estimated emissions per compound and of the aggregated greenhouse gas emissions will be somewhat different.

As can be seen in Figure 1-4, when including LULUCF in national total emissions the uncertainty is higher ($\pm 32\%$), and this is an effect of large and relatively uncertain carbon sinks.

Table 1-7 shows the ten sources with the largest uncertainty contributions in the Swedish inventory for 2011, excluding LULUCF.

The uncertainty of the trend of national total greenhouse gas emissions excluding LULUCF was $\pm 2.0\%$. The uncertainty in the trend is a percentage point range relative to the inventory trend and should be interpreted as $\pm 2.0\%$ to the estimated percentage difference between total GHG emissions 1990 and 2011, i.e. there is a 95% probability that the decrease in GHG emissions in Sweden 1990 to 2011 is between 13.5 % and 17.5 %.

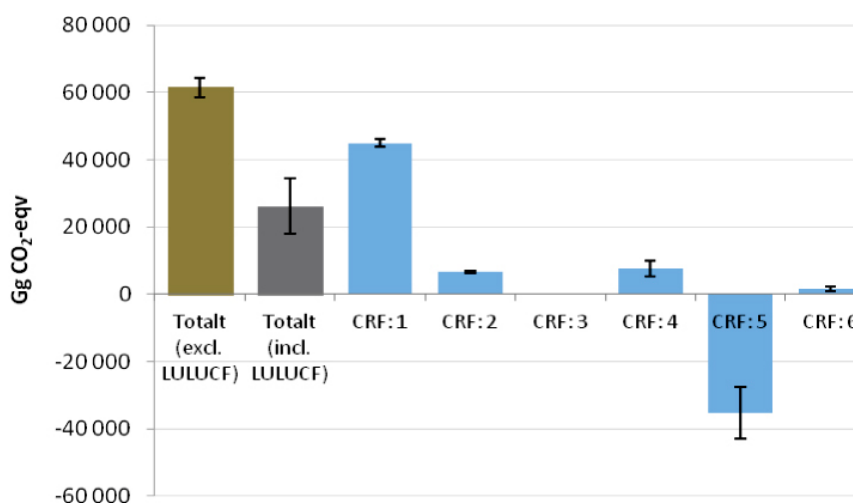


Figure 1-4 Uncertainty estimates, as 95% confidence intervals, in national total emissions (excluding and including LULUCF) and by sector.

Table 1-7. The ten sources with the largest uncertainty contributions in the Swedish inventory for 2011, excluding LULUCF.

IPCC Source Category	GHG	Year 2011 emissions or removals (Gg CO ₂ -eqv)	Activity data uncertainty (%)	Emission factor uncertainty (%)	Combined uncertainty (%)	Relative contribution to variance in year 2011 (%)
4.D.1 (Direct Soil Emissions)	N ₂ O	2,493	15	66	67	37%
4.D.4 (Agricultural Soils. Other)	N ₂ O	686	35	150	154	15%
4.D.3 (Indirect Emissions)	N ₂ O	828	28	122	125	14%
4.D.2 (Pasture, Range and Paddock Manure)	N ₂ O	440	35	150	154	6%
6.A (Solid Waste Disposal on Land)	CH ₄	1,193	25	50	56	6%
1.B.2 (Oil and Natural Gas)	CH ₄	109	0	583	583	5%
1.AA.3.B (Road Transportation)	CO ₂	18,412	2	2	3	5%
1.AA.1.A (Public Electricity and Heat Production)	CO ₂	7,756	1	7	7	4%
4.A (Enteric Fermentation)	CH ₄	2,578	2	11	12	1%
1.AA.2.F (Other Manufacturing Industries and Construction)	CO ₂	4,546	5	3	6	1%

1.7.2 KP-LULUCF activities

Estimates of carbon stock changes are based on the same underlying data as the reporting under the UNFCCC. These estimates originate mainly from a sampling design with the intention to keep systematic errors as low as possible. The systematic error is reduced by using representative functions, by direct measurements in field and laboratory measurements. We assume that the major source of uncertainty arises from random variation due to sampling. The sampling error is estimated using statistical theory for living biomass and partly for other carbon pools (all Tier 3). A consistent methodology for estimating carbon pools has been used from 1990 and onwards. Therefore, we expect the uncertainty to be the same for all years where all sample units are used to estimate the annual change. The uncertainties for other categories are based on expert judgment.

1.7.3 General assessment of completeness

In the following section the completeness of the GHG inventory and the KP-LULUCF inventory is described.

1.7.3.1 GHG INVENTORY

The inventory covers emissions and sinks in Sweden. All greenhouse gases are covered. The general completeness for each sector is discussed below. Detailed information is presented in Annex 5.

1.7.3.2 ENERGY

Estimated emissions are considered to be complete for most sources. Emissions of CH₄ and N₂O from liquid bio fuels used in military transportation are however not estimated. There might also still be some lack in completeness as regards in-house generated fuels in the chemical industry and in smaller companies.

1.7.3.3 INDUSTRIAL PROCESSES

For most sources, and particularly for the most important ones, the estimates are in accordance with the requirements concerning completeness as laid out in the Good Practice Guidance. However, some exceptions do exist. These are primarily in sub-sectors with a large number of smaller facilities with minor emissions.

Data is complete for all greenhouse gases, possibly with the exception of CH₄ for a few sources, e.g. within the chemical industry.

1.7.3.4 SOLVENT AND OTHER PRODUCT USE

The estimated emissions from solvent and product use are considered to be complete, since a new method was developed during 2005 in order to obtain all activity data concerning the sector from the Products register at the Swedish Chemicals Agency.

The estimated emissions of N₂O are also considered to be complete, since national data from the Products register is used in the inventory.

1.7.3.5 AGRICULTURE

All relevant agricultural emissions and sources are reported in the inventory. Reindeer, which are normally not considered as a part of the agricultural sector, are included in the inventory. There are, however, some marginal animal groups, which are not included, such as fur-bearing animals (minks, foxes and chinchillas). These groups are very small and there is no methodology developed for estimating their GHG emissions.

All sales of fertilizers are included in the inventory, also quantities used in other sectors. N-fixing crops used in temporary grass fields, and sludge used as fertilizer is also included. This means that all anthropogenic inputs to agricultural soils are covered.

1.7.3.6 LAND USE, LAND USE CHANGE AND FORESTRY

All land areas are inventoried in the field except high mountains, military impediments and urban land. We believe that their relative importance for the Swedish GHG inventory is small.

The inventory of the LULUCF-sector is complete in the sense that all carbon pools and other sources, defined based on the IPCC GPG for LULUCF, are reported for land use categories that are considered managed.

The reporting of woody biomass stocks refers to above and below ground parts of trees taller than 1.3 m. Other vegetation such as shrubs and herbs are not reported. Emissions/removals from below ground biomass of dead stump systems are from this submission included in the dead organic matter pool.

1.7.3.7 WASTE

The effects of possible leakage of methane and nitrous oxide from the wastewater treatment processes have not been estimated. All other data are complete.

1.7.3.8 KP-LULUCF

Sweden has elected the activity Forest management (FM) under Article 3.4 of the Kyoto Protocol (KP). All carbon pools as well as associated mandatory activities (such as fertilization of forest land, biomass burning and conversion to cropland) are reported for activities under article 3.3 and under FM.

2 Trends in greenhouse gas emissions

2.1 Description and interpretation of emission trends for aggregated greenhouse gas emissions

Total greenhouse gas emissions in Sweden, calculated as carbon dioxide equivalents, totalled 61.4 million tonnes (excl. LULUCF) in 2011, see Figure 2-1. Aggregated greenhouse gas emissions varied over the period but in all cases were less than the 1990 level during the period 1999-2011. Emissions decreased by 11 million tonnes or 16 % between 1990 and 2011. The uncertainty range in the trend is ± 2.0 % relative to the inventory trend. The decreasing emissions in the energy sector, in agriculture, and in the waste sector contribute to the decreasing trend in total emissions from 1999.

Emissions in year 2009 and 2010 were quite extreme. During the autumn of 2008, an economic downturn began which was deepened during 2009 and has affected a number of sectors. The recession has led to that many industrial sectors have reduced production considerably, with diminished emissions in 2009 as a consequence. In 2010 most of the industries have started to recover from the economic downturn. The cold winters in 2010, both in the beginning and in the end of the year, increased the need for heating. In combination with lower nuclear production than normal this led to increased use of fossil fuels in the electricity and heat production. In 2011 the emissions decreased compared to year 2010 due to a warmer winter and that the economic development in the industry somewhat stabilized.

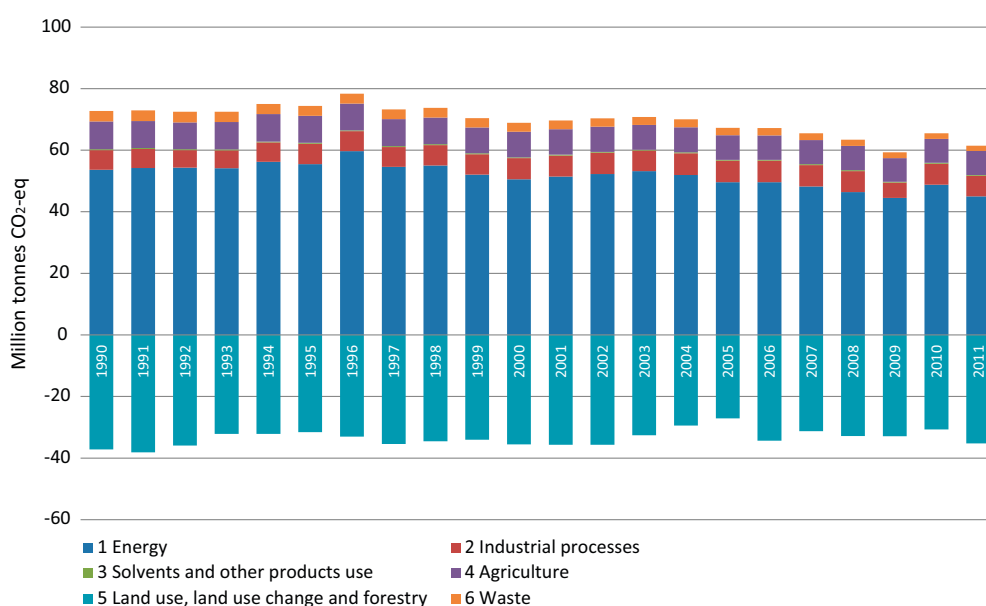


Figure 2-1 Total emissions and removals of greenhouse gases by sectors.

The Land Use, Land-Use Change and Forestry sector (LULUCF) resulted in annual net removals in Sweden during the period 1990-2011. The size of the sink varied over the period.

Yearly growth in GDP averaged around 4 % over the period 1990 to 2011. The growth in GDP has been positive for all years except between 1991 and 1992, and between 2008 and 2009. The overall growth between 1990 to 2011 was around 40 %. Despite this economic growth, the emissions, reported to the UNFCCC, have been reduced during the period. The emissions decreased with 16 % during the same period of time.

2.1.1 Overview of emissions trends per sector

Emissions of greenhouse gases have developed differently in different sectors over the period from 1990 to 2011, see Figure 2-2. The sectors which contributed mostly to the overall decrease is the sector other sectors (CRF 1.A.4), manufacturing industries and construction (CRF 1.A.2), waste (CRF 6) and agriculture (CRF 4).

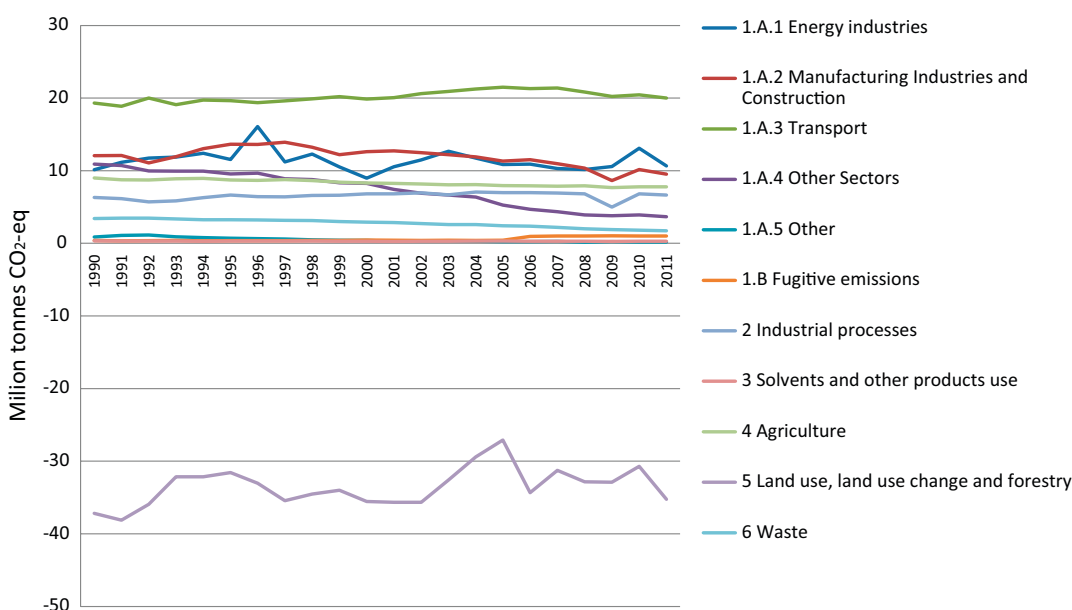


Figure 2-2 Total emissions of all greenhouse gases calculated as CO₂-equivalents by sectors.

2.2 Description and interpretation of emission trends by gas

In 2011, emissions (excl. LULUCF) of carbon dioxide totalled 48.7 million tonnes, which is equivalent to almost 80 % of aggregated greenhouse gas emissions counted as carbon dioxide equivalents, see Figure 2-3. Emissions of methane were 5.0 million tonnes of carbon dioxide equivalents and account for just over 8 % of emissions, while emissions of nitrous oxide totalled 6.7 million tonnes, equivalent to almost 11 %. Emissions of fluorinated green-

house gases amounted to almost 2 % or 1.1 million tonnes of carbon dioxide equivalents of the aggregated greenhouse gas emissions. The shares of the different greenhouses gases were roughly the same over the period 1990 to 2011.

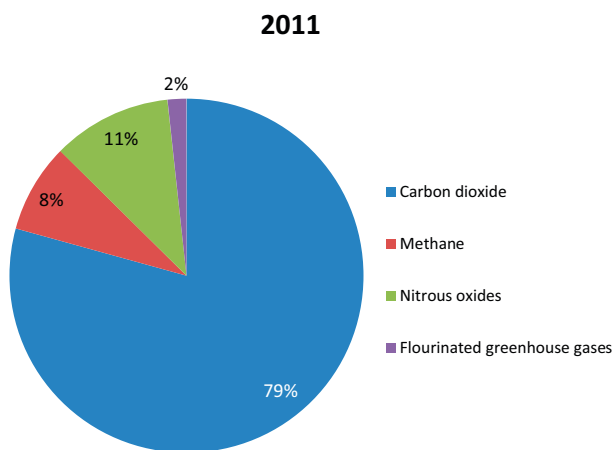


Figure 2-3 Share of greenhouse gases in emissions in year 2011, in carbon dioxide equivalents.

2.2.1 Carbon dioxide

In 2011, the carbon dioxide (CO₂) emissions in Sweden totalled 48.7 million tonnes, excluding LULUCF, see Figure 2-4. 88 % of the carbon dioxide emissions originated from the energy sector (CRF 1), 11 % originated from industrial processes (CRF 2), and the remaining less than 1% originates from solvent and other product use (CRF 3) and waste (CRF 6). Emissions were 14 % lower in 2011 than in 1990, the energy sector stands for the largest reduction.

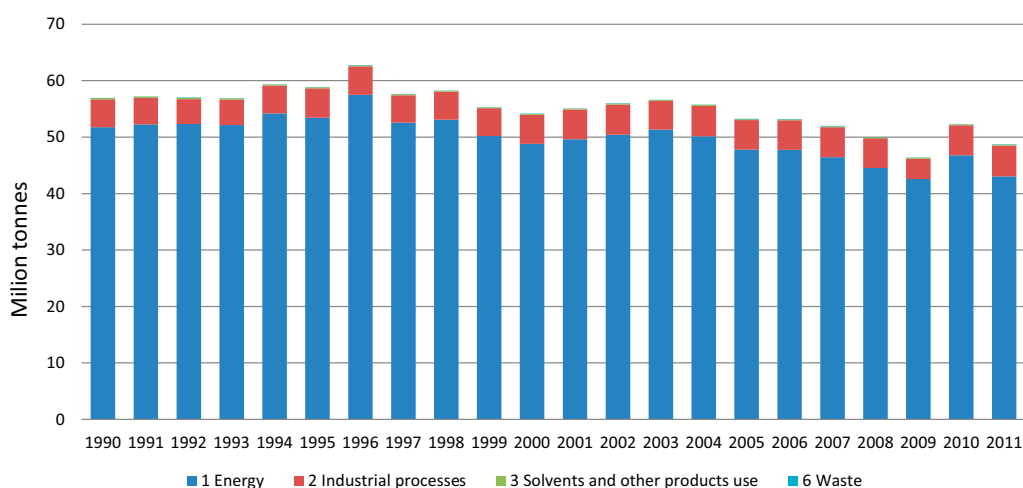


Figure 2-4 Total emissions of CO₂ from different sectors in million tonnes.

2.2.2 Methane

Almost 60 % of the emissions of methane (CH_4) originate from agriculture (CRF 4). Around 30 % is emitted in the waste sector (CRF 6) and 12 % is emitted in the energy sector (CRF 1). The total emissions of methane, excluding emissions from LULUCF, totalled 237 ktonnes in 2011, which is equivalent to 5.0 million tonnes calculated as carbon dioxide equivalents or 8 % of total greenhouse gas emissions (Figure 2-5). Emissions have decreased by 28 % since 1990, largely due to measures taken in the waste sector.

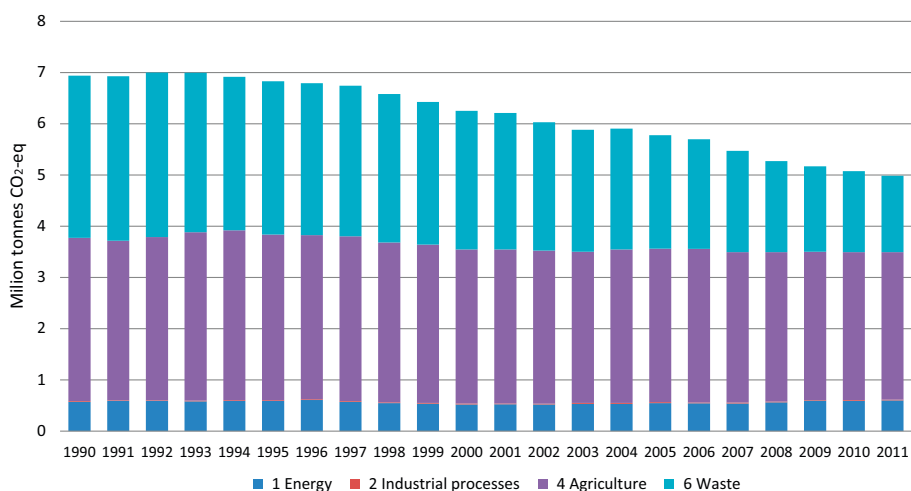


Figure 2-5 Total emissions of CH_4 from different sectors, calculated as CO_2 -equivalents.

2.2.3 Nitrous oxide

In 2011, emissions of nitrous oxide (N_2O) totalled 22 ktonnes or almost 7 million tonnes of carbon dioxide equivalents (excl. LULUCF), see Figure 2-6. All sectors produce nitrous oxide emissions, but the emissions mainly originate from the agriculture sector, which accounted for 73 % of nitrous oxide emissions. Compared with 1990, emissions have decreased by 20 %, and it is primarily emissions from the agriculture sector that account for the decrease.

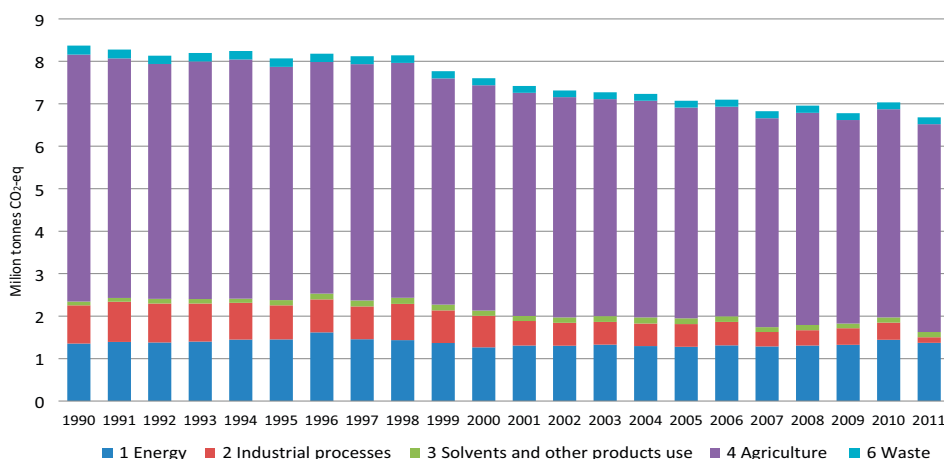


Figure 2-6 Total emissions of N_2O from different sectors calculated as CO_2 -equivalents.

2.2.4 Fluorinated greenhouse gases

Emissions of fluorinated greenhouse gases (F-gases), HFC, PFC, SF₆, are reported in the industrial processes sector. Total emissions of fluorinated greenhouse gases in 2011 amounted to 1.1 million tonnes calculated as carbon dioxide equivalents and account for almost 2 % of the total greenhouse gas emissions, see Figure 2-7. The emissions of f-gases increased with 116 % between 1990 and 2011.

The increasing emissions of f-gases are mainly due to increased emissions of HFCs. Emissions of HFCs increased in particular, from 4 ktonnes of carbon dioxide equivalents in 1990 to 813 ktonnes in 2011. PFCs emissions, on the other hand, have decreased. In 1990 emissions of PFCs amounted to 377 ktonnes of carbon dioxide equivalents, and in 2011 they had decreased to around 183 ktonnes. Emissions of SF₆ varied between 1990 and 2010. In 1990 they totalled 107 ktonnes and in 2011 they amounted to 60 ktonnes of carbon dioxide equivalents.

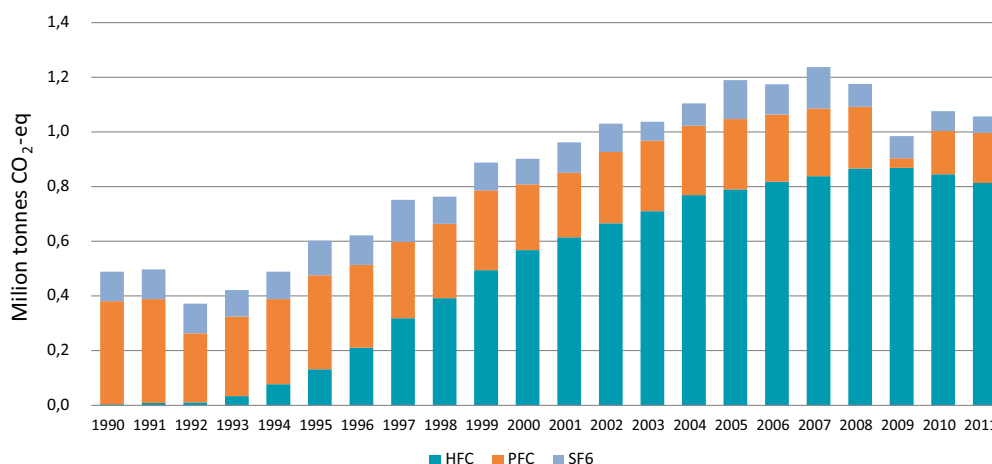


Figure 2-7 Total emissions of SF₆, PFC and HFC, calculated as CO₂-equivalents.

2.3 Description and interpretation of emissions by category

In 2011 emissions from the Energy sector (CRF 1) made up 72 % of the total emissions, see Figure 2-8. The Energy sector includes Transport (CRF 1.A.3) (33 % of total emissions), Energy industries (CRF 1.A.1) (17%) and combustion in Manufacturing Industries (CRF 1.A.2) (15%). Other sectors (CRF 1.A.4) with (6 %) and Fugitive emissions (CRF 1.B) (2%) are also part of the Energy sector. Agriculture (CRF 4) accounted for 13 % of total emissions and Industrial processes (CRF 2) for 11 % of total emissions. Waste (CRF 6) contributes with 3% of the emissions and Solvents and other product use (CRF 3) counts for less than 0.5 %.

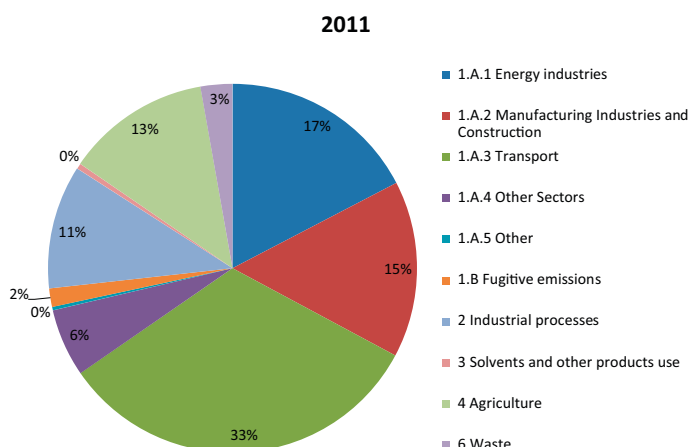


Figure 2-8 Greenhouse gas emissions by sector (2011).

2.3.1 Energy

Emissions from the energy sector include emissions from the production of electricity and district heating, refineries, manufacture of solid fuels, manufacturing industries, transports, other sectors (including commercial/institutional, residential, agriculture, forestry and fisheries), other (military transports), and fugitive emissions.

Emissions from the transport sector (1A3) dominates followed by energy industries (1A) and combustion from manufacturing industries and construction (1A2), which account for almost equal shares of total emissions within the energy sector in 2011 (Figure 2-9). Important sources are production of electricity and heating within energy industries sector (1A) and heating in the residential and commercial/institutional sector included in “other sectors” (1A4).

In the energy sector there has been a reduction in total emissions over the period 1990-2011, see Figure 2-10. Emissions have decreased from 53.7 million tonnes carbon dioxide equivalents in 1990 to 45.0 in 2011. This is a decrease with 16% compared to 1990, and the decrease is principally due to decreased use of oil for heating in residential and commercial/institutional, included in “Other Sectors” (1A4), and district heating being increasingly based on biomass fuels. Between 2011 and 2010 there was a decrease in emissions with 8 %.

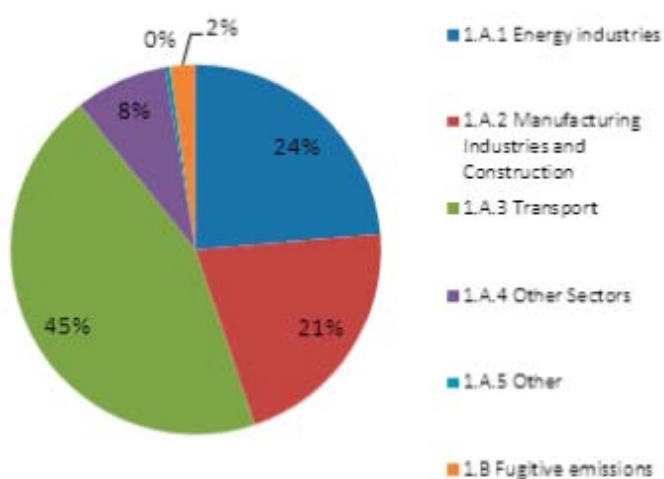


Figure 2-9 Share of emissions (2011) for the Energy sector, by subsector.

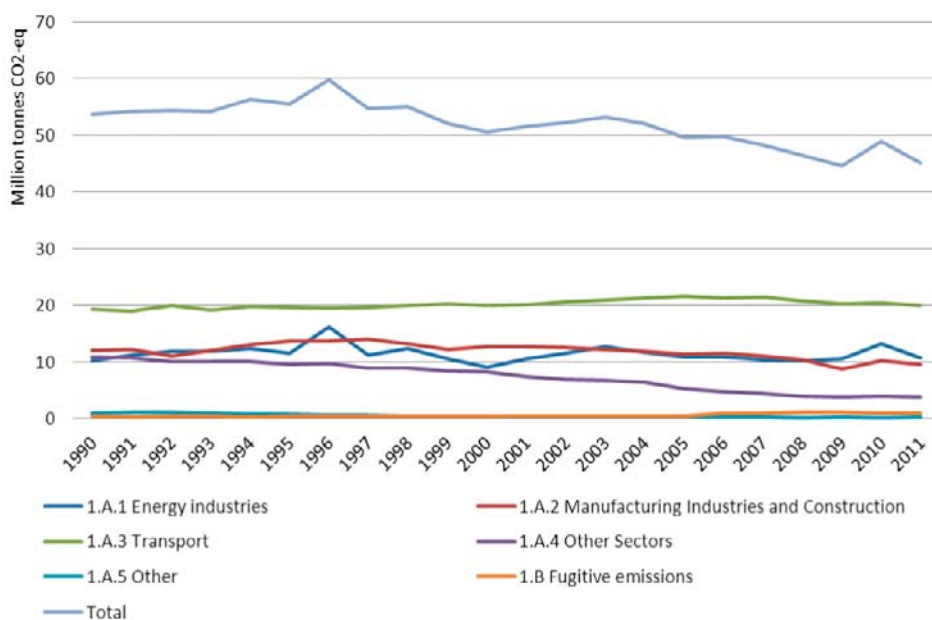


Figure 2-10 Emissions from the Energy sector, total and by subsector.

2.3.1.1 ENERGY INDUSTRIES (CRF 1A)

Total emissions for energy industries (1A) is 10.7 million tonnes carbon dioxide equivalents in 2011 (Figure 2-1). Energy industries are dominated by the electricity and heat production (1A1a) which stand for the larger part of the emissions (8.3 million tonnes). Emissions from Refineries (1A1b) contributes with 2.0 million tonnes and Manufacture of solid fuels (1A1c) 0.4 million tonnes in 2011.

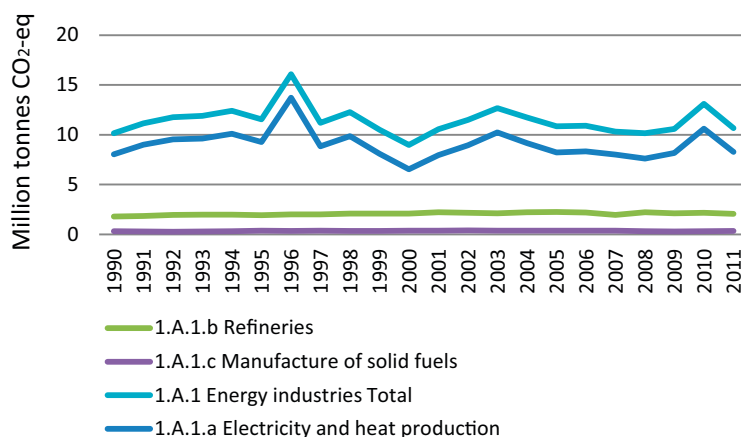


Figure 2-11 Emissions from Energy industries, total and by subsector.

2.3.1.1.1 ELECTRICITY AND HEAT PRODUCTION (CRF1A1A)

Emissions from production of electricity and heat production totalled around 8.3 million tonnes of carbon dioxide equivalents in 2011, which is a decrease by 22 % compared to 2010. The 2011 emissions are around the same level as the emissions in 2009, and slightly higher than the emission level in 1990 (3 %). The emissions from electricity and heat production vary over time and no apparent trend can be seen between 1990 and 2011.

Temperature and precipitation conditions, which vary between years, have an impact on hydropower production and heating needs in individual years and thus lead to a variation in emissions between years. This is illustrated by the high emissions in 1996, which was a cold and dry year, and by the low emissions in 2000, which was a warm year with heavy precipitation and thus good availability of hydropower.

In years of low hydropower production the emissions are affected by which kind of electricity production the shortage is offset by. Therefore the deficient production of hydropower in 2003 was offset by imports of electricity while shortage of hydropower in 1996, which was another year of low hydropower production, was partially offset by increased oil condensing production. The increased possibility to import electricity from Nordic countries might therefore lead to decreased emissions in certain years.

Emissions are also affected by the increase in iron and steel production which has taken place since 1990, as residual gases from this industry are used to produce electricity and district heating.

High prices on fossil fuels and introduction of the electricity certificates system, which improves the profitability of renewable electricity production, have had a reducing effect on the emissions¹⁶.

During the period 1990-2011, the supply of district heating increased with around 50 %¹⁷. On the other hand, emissions have not increased significantly as the expansion has principally taken place through increased use of biomass fuels at the same time as the use of oil and coal has decreased. Energy and carbon dioxide taxes have, among other things, contributed to this trend¹⁸.

The total production of electricity in 2011 was around the same level as in 2010, but the electricity demand was lower in 2011¹⁹. Main reasons for reduced use of electricity were decreased heating demand due to warm weather and a certain industrial slowdown²⁰. In Sweden there was a net export of electricity in 2011 (7 TWh), while in 2010 there was a net import (5 TWh). In the Nordic countries there was still a net importing (5 TWh) in 2011, however lower than in 2010 (19 TWh)²¹. The production in Swedish nuclear reactors during 2011 was higher than in 2009 and 2010, but still lower than the expected average production,²² Inflow to reservoirs during 2011 was higher than normal; even though reservoirs were lower than average in the beginning of the year. An early spring flood and a large amount of precipitation were the reasons behind the high inflow²³. Wind-power production continued to increase sharply. The increase between 2011 and 2010 has been 74 % and the total increase from 6 GWh in 1990 to 6,078 GWh in 2011.²⁴ Combustion of natural gas for production of electricity and district heating increased significantly between 2008 and 2010. In 2011 however, usage of natural gas decreased back to around the same level as in 2009²⁵, probably because of lower electricity and heating demand. The production of district heating decreased in 2011 compared to 2010 (with 6 TWh) and the use of oil and natural gas decreased the most²⁶.

2011 was estimated as 13 % warmer than a “normal” year²⁷, whereas 2010 was estimated as 14 % colder than a “normal” year. The decrease due to warmer weather has been analyzed with a normal-correction calculation method, which concludes that emissions in 2011 were lower than they would

¹⁶ Swedish Energy Agency, 2012b

¹⁷ Swedish Energy Agency, 2012b

¹⁸ Ministry of the Environment Sweden, 2009.

¹⁹ Swedish Energy Agency, 2012b

²⁰ Swedenergy, 2012

²¹ Swedenergy, 2012

²² Swedish Energy Agency, 2012b

²³ Swedenergy, 2012

²⁴ Swedish Energy Agency, 2012b

²⁵ Swedish Energy Agency, 2012b and Swedish Energy Agency, 2012c

²⁶ Swedish Energy Agency, 2012b

²⁷ Swedish Energy Agency, 2012b

have been a “normal” year (Figure 2-12). Emissions from electricity and heat production (1A1a) and residential and commercial/institutional (1A4a-b) are included. Temperature and precipitation are some of the included parameters and for more information about the method used, see Annex 8:2.

All together these factors lead to lower emissions from both electricity and heating in 2011 compared to the emissions in 2010. The warm weather, favourable conditions from hydropower production and increased production in nuclear power plants are some of the important factors behind this development.

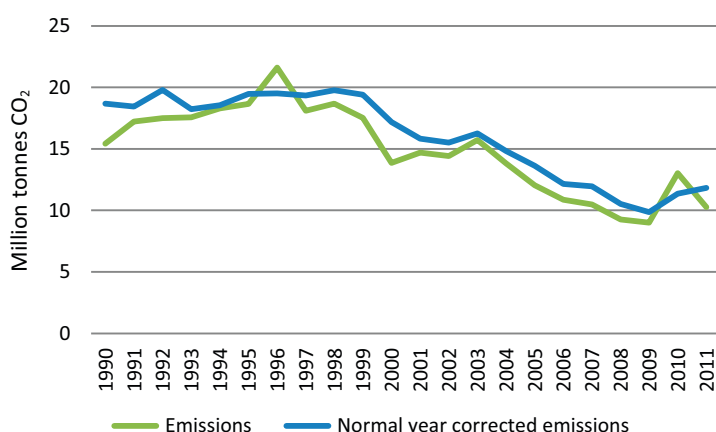


Figure 2-12. Actual and normal-year corrected fossil CO₂-emissions for heating of buildings and electricity production in Sweden for the years 1990-2011. Included sectors are production of electricity and heat (1A1a) and residential and commercial/institutional (1A4a-b). For the year 2011 preliminary statistics on fossil fuel consumption is used

2.3.1.1.2 Refineries (CRF 1A1b)

Production of refined products increased in Sweden during the period, which increased emissions with 14 %, between 1990 and 2011. The emissions were 5 % lower in 2011 than in 2010. Generally the emissions have been around 2 million tonnes carbon dioxide equivalents since 1990 (Figure 2-11).

2.3.1.2 MANUFACTURING INDUSTRIES AND CONSTRUCTION (CRF 1A2)

Emissions from combustion in the industrial sector were around 9.5 million tonnes carbon dioxide equivalents in 2011. Emissions in 2011 were 21 % lower than in 1990, but they have varied upwards and downwards over the years, principally due to economic fluctuations and replacement of oil with electricity or biofuels partly depending on the difference in relative prices between electricity and oil. A small number of energy-intensive industries account for a large proportion of carbon dioxide emissions in the sector. The iron and steel industry (1A2a), the pulp and paper industry (1A2d) and

chemical industry (1A2c) account for almost equal shares in the emissions – 16 %, 13 % and 13% of emissions in this sector (1A2) in 2011. Other industries (1A2F) stand for 52 % of the emissions in 2011. This sector is heterogeneous and includes for instance mining industry and non-metallic mineral production. Off road vehicles and working machinery from all industries are allocated to Other industries (1A2f).

There is a downward trend in total emissions from combustion in manufacturing industries between 2002 and 2011, and one reason is reduced emissions from the pulp and paper industry, caused by fuel substitution from fossil fuels to biofuels, see Figure 2-13. Also other manufacturing industries have made a transition to electricity or biofuels, but the conversion in the pulp and paper sector have the largest impact on emissions. Stationary combustion from “other industries” also has a decreasing trend. The decreases were larger in 2008 and 2009 during the economic recession, but in 2010 production volumes, energy demand and emissions increased again. In 2011, emissions and energy demand decreased²⁸ and there was a slight decrease in production in some industries compared to 2010.

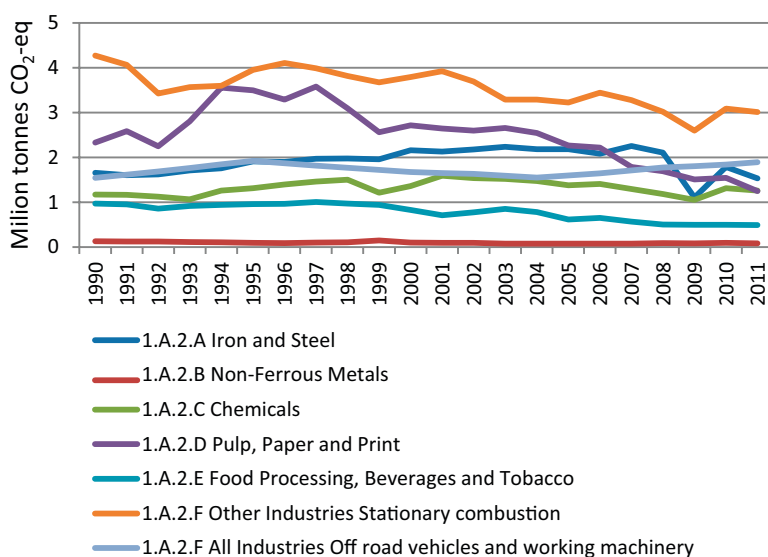


Figure 2-13 Emissions from combustion in manufacturing industries by subsectors.

²⁸ Swedish Energy Agency, 2012b

Viewed over a longer period from 1970 on, industry has reduced its use of oil and increased its use of electricity. Between 1990 and 2010 the use of oil decreased with 36 %²⁹. In 2011 the usage of oil for energy purposes within the industry decreased to the same level as in 2009. Biofuels and electricity is now the main energy sources within the industry and stood in 2010 for respectively 37 % and 36 % of the final energy use.³⁰ Use of biofuels is most frequent in the pulp and paper sector.

2.3.1.3 FUGITIVE EMISSIONS FROM FUELS (CRF1B)

Emissions from the fugitive emissions sector come for example from refineries. Emissions were 1.0 million tonnes of carbon dioxide in 2011 (Figure 2-14). The increase of fugitive emissions from oil (1B2A) since 2006, is due to that two new hydrogen production plants were taken into operation.

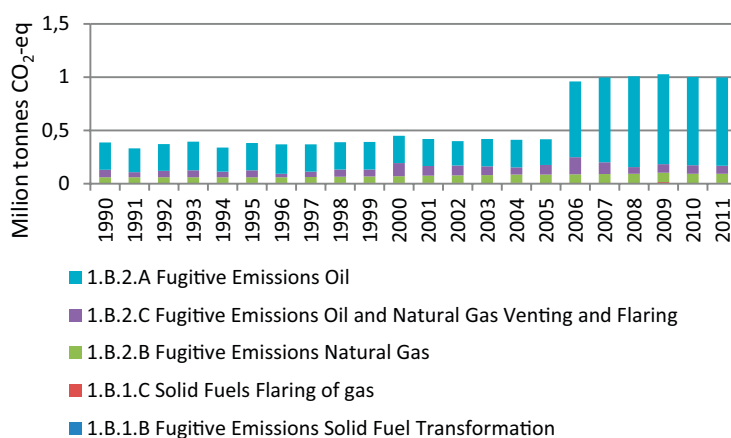


Figure 2-14 Emissions of all greenhouse gases from Fugitive emissions, total and by subsector.

²⁹ Swedish Energy Agency, 2012b and Swedish Energy Agency, 2012c

³⁰ Swedish Energy Agency, 2012b

2.3.1.4 TRANSPORT (CRF 1.A.3)

Greenhouse gas emissions from domestic transport totalled 20.0 million tonnes of carbon dioxide equivalents in 2011 (Figure 2-15), which is 3.6 % higher than the 1990 level. After a peak in emissions around 2005, emissions have decreased slightly. Greenhouse gas emissions 2011 from road transportation were 18.6 million tonnes, from domestic aviation 0.5 million tonnes, from domestic navigation 0.5 million tonnes, from railways 0.1 million tonnes and from other machinery 0.3 million tonnes.

While emissions of carbon dioxide account for the by far largest share of greenhouse gas emissions from the transport sector, methane and nitrous oxide contribute with a very small share. Methane emissions totalled 0.05 million tonnes of carbon dioxide equivalents in 2011 and have fallen by 73 % since 1990 as a result of better exhaust emission control. Nitrous oxide emissions totalled 0.16 million tonnes of carbon dioxide equivalents in 2011. Emissions of nitrous oxide increased from 1990 to 2000 in connection with the switch to cars fitted with catalytic converters. Emissions decreased during the early 2000s with better exhaust treatment technology, and have remained at a fairly steady level during the latest years.

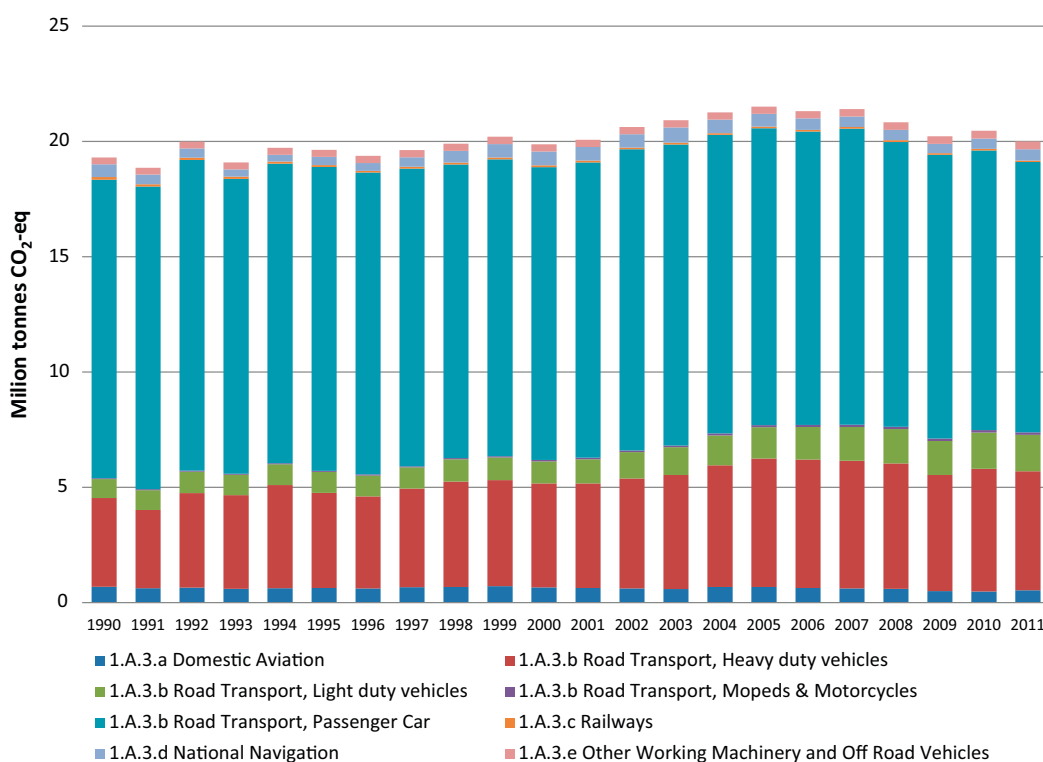


Figure 2-15 Emissions of CO₂ eq. from the transport sector, total and per subsector.

2.3.1.4.1 *Road transport*

Road transportation accounts for the greatest share of the transport sector's greenhouse gas emissions and totalled 18.6 million tonnes of carbon dioxide equivalents in 2011, which is 5.2 % higher than 1990 emissions. Emissions increased from 1990 to 2005 before stagnating, and has since then been decreasing somewhat as a result of an increased share of renewable fuels and reduced fuel consumption in combination with the economic downturn in 2008-2009.

The increase up to the mid-2000s followed the increase in traffic and transport mileage. It was principally transport mileage with heavy duty vehicles and to some extent with light duty vehicles that increased. There was insignificant improvement in energy efficiency in vehicles during this period. The proportion of diesel-powered light duty vehicles has increased continuously since 1990, and the same change in the market began in the late 1990s for cars, steadily strengthening to the present day. The switch from petrol-powered to diesel-powered cars is leading to greater energy efficiency, which since the mid-2000s has been reinforced by a general improvement in fuel efficiency for new cars which has counteracted the increase in fuel consumption to which the growth in transport has led. The aggregate effect has been sharply increased consumption of diesel fuel and reduced volumes of petrol. The economic downturn in 2008 - 2009 also led to reduced emissions from diesel-powered heavy duty vehicles in 2009 and to somewhat lower overall use of diesel fuel in the road transport sector.

Several factors have been significant in limiting emissions from road transport. Among other factors, increased fuel taxes, tax exemption for transport biofuels, carbon dioxide-based vehicle tax, tax relief for green cars and green car rebates, together with rising market price for petrol and diesel, have contributed to more fuel-efficient cars, an increased number of fuel-flexible cars and consequently a switch to renewable fuels. The use of renewable fuels has been principally boosted by the fact that since 2004 they have been exempt from carbon dioxide tax and energy tax, along with a law from 2006 requiring every major petrol station to provide a renewable fuel. Large-scale blending of ethanol into petrol began in 2003, with the result that almost all petrol sold in Sweden now contains 5% ethanol. Blending of FAME into diesel has also increased, accounting for the proportionally largest increase among the biofuels in 2011. (SPBI 2012). The use of biofuels for fuel-flexible cars also increased from 2004 to 2011, particularly ethanol (E85), although in 2009 the use of E85 decreased somewhat as a result of high ethanol prices and lower petrol prices.

2.3.1.4.2 *Domestic aviation*

In 2011, emissions from domestic aviation totalled 0.5 million tonnes of carbon dioxide equivalents (Figure 2-16), which is 22 % lower than the level for 1990. However, the emissions varied during the period. Domestic aviation has decreased since the mid-2000s because the share of train and to some

extent car journeys has increased. The fact that more people are choosing train or car rather than flying for domestic travel is considered to be due in part to a decline in the availability of short-haul aviation and to new security requirements and routines having reduced the advantages of flying in terms of speed and flexibility. In 2011, however, there was a 10 % increase in the emissions. This increase was likely related to weather – the cold and snowy winter, in combination with the problems of the railway sector in handling the situation, caused railway passengers as well as car drivers to use aviation.

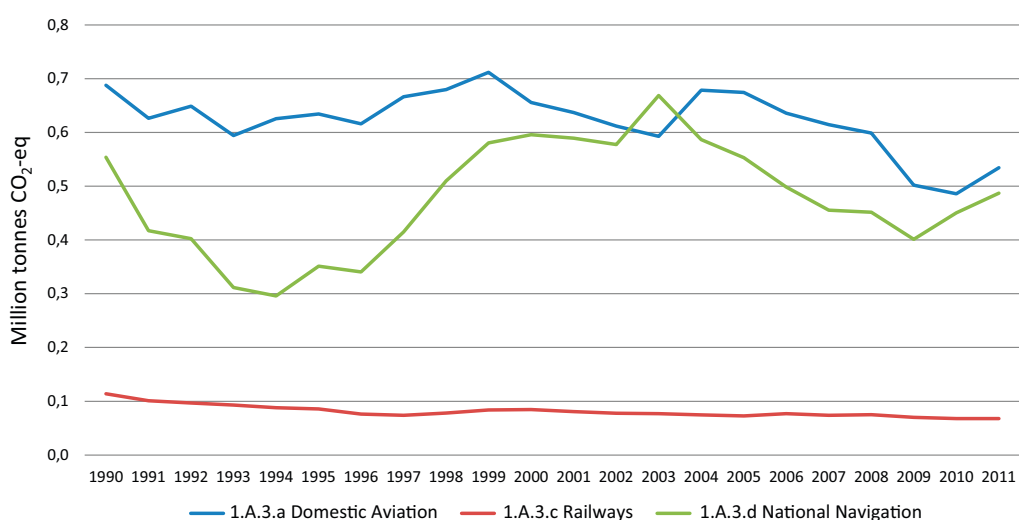


Figure 2-16 Emission of CO₂ eq. from aviation, navigation and railways.

2.3.1.4.3 Navigation

Emissions for domestic navigation are estimated at 0.6 million tonnes of carbon dioxide equivalents in 2011 (Figure 2-16), which is around 12% lower than in 1990. Emissions have been increasing after a decrease between 2003 and 2009.

2.3.1.4.4 Railways

Railway greenhouse gas emissions from diesel-powered trains account for a marginal share of transport sector emissions and have decreased by around 40 % since 1990 (Figure 2-16).

2.3.1.5 OTHER SECTORS (CRF1A4)

Other Sectors include combustion in the residential, commercial/institutional and agriculture, forestry and fisheries sector. Mobile emissions from working machinery and off road vehicles are also included in the sectors residential and agriculture, forestry and fisheries.

The emissions in Other Sectors were approximately 3.7 million tonnes of carbon dioxide equivalents in 2011, which is a decrease with 7% compared to 2010. In comparison to 1990 the decrease is 67 %. The reduction is due to

a strong decrease in emissions from the residential and commercial/institutional (1A4A and 1A4B) sectors between 1990 and 2011 (Figure 2-17), depending on a large decrease in total use of fossil fuels.

It is mainly energy input for heating and warm water within the residential and commercial/institutional sectors that is decreasing. There are several reasons for this development: the shift from oil to district heating and electricity heating, increased usage of heat pumps and pellet fired boilers and measures to increase energy efficiency. The amount of heat pumps has increased considerably and almost half of all one and two-dwelling buildings had some kind of heat pump in 2011³¹. Another contributing factor to the favourable development has been the generally warm weather since 1990.

The emissions in 2011 decreased slightly from 2010 partly due to warmer weather and decreased heating demand. More information about the weather in 2011 and normal corrected emissions can be found in 2.3.1.1.1 and in Figure 2-12.

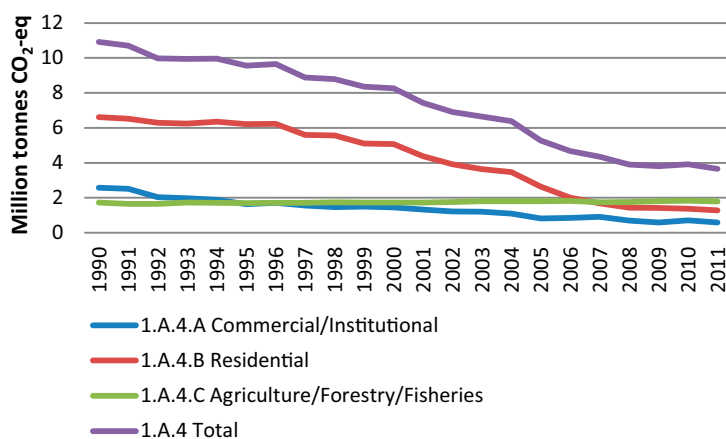


Figure 2-17 Emissions from Other sectors, total and by subsector.

In 2011 the share of biofuels and district heating has increased some and there is a continued decrease in usage of oil in one and two-dwelling buildings. Less than 1 % of one and two-dwelling buildings have oil as their sole source of heating in 2011, which is a decrease from 2010. This development also contributes to the decrease in emissions during 2011. The large decrease in emissions from the residential sector has resulted in a substantial increase in the share of emissions from biofuels (Figure 2-18). There is also a continued decrease in energy consumption for heating per unit of floor space area in 2011 for these houses.³²

³¹ Swedish Energy Agency, 2012b

³² Swedish Energy Agency, 2012.

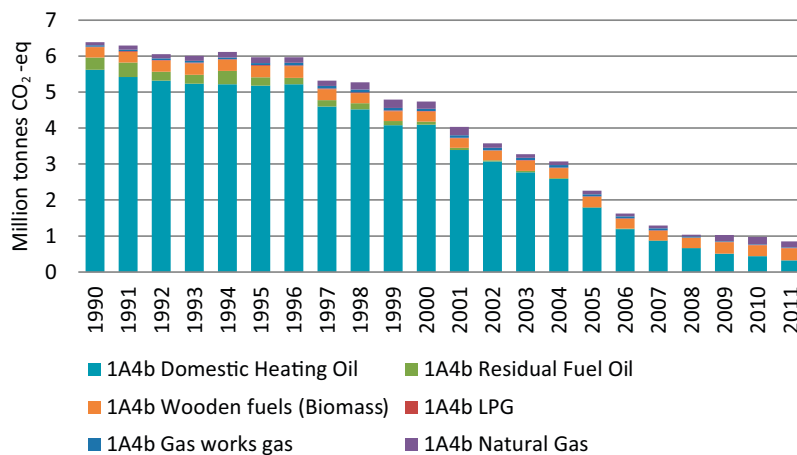


Figure 2-18 Emissions from Residential sector (1A4b) by fuel type, stationary combustion.

Working machinery and off road vehicles are also included in the Residential sector (1A4b). These emissions are slightly increasing, but are very low, around 0.4 million tonnes carbon dioxide equivalents in 2011.

In agriculture, forestry and fisheries (1A4c) emissions from stationary combustion are decreasing, while mobile emissions from working machinery and off road vehicles in agriculture and forestry increase slightly (Figure 2-19). In fisheries there is instead a decrease in mobile emissions.

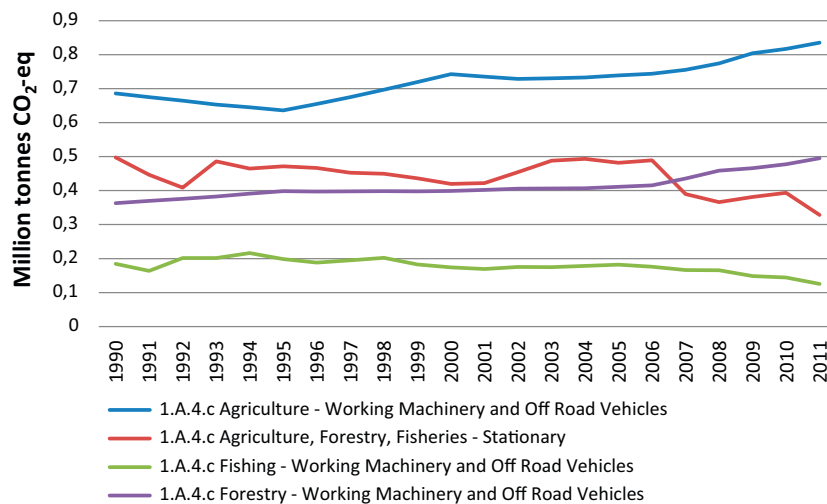


Figure 2-19 Emissions from agriculture, forestry and fisheries, mobile and stationary combustion.

2.3.1.6 EMISSIONS OF METHANE AND NITROUS OXIDE FROM THE ENERGY SECTOR EXCLUDING TRANSPORTS

Only a small proportion of emissions from the energy sector are emissions of methane and nitrous oxide. Almost 5 % of emissions from the energy sector are emissions of nitrous oxide, and approximately 2 % are emissions of methane.

Methane emissions from the energy sector excl. transport have increased by 42 % between 1990 and 2011. The increased emissions are principally due to increased use of biofuels in the residential and commercial/institutional service sector and in the production of electricity and district heating. Almost two-thirds of emissions originate from the residential and commercial/institutional service sector, and including energy use in agriculture. Also electricity and district heating had increased methane emissions principally due to increase in total emissions.

Nitrous oxide emissions decreased slightly between 2010 and 2011, however it is still an increase with about 7 % compared to 1990.

2.3.2 Industrial processes

Greenhouse gas emissions within industrial processes (CRF 2) stem from the materials in the processes. The combustion of fuels, for production or heating purposes, is reported in the energy sector (CRF 1). To cover all industry related emissions one must take into consideration emissions from CRF 1 and CRF 2.

Greenhouse gas emissions from the industrial processes sector are 0.3 million tonnes CO₂ equivalents higher in 2011 compared to 1990. The increase is approximately 5% from 6.3 million tonnes of CO₂ equivalents in to 6.7 million tonnes CO₂ equivalents. The emissions from industrial processes make up 11% of the aggregated total emissions in 2011. The trend is mainly affected by increased emissions of HFCs and CO₂, but also in decreased emissions of N₂O and PFCs.

The main sources for emissions in the industrial processes are the production of iron and steel and the cement and lime industries, see Figure 2-20. Other sources are the usage of coke in blast furnaces, of dolomite and limestone in production in the mineral industry and of coal in the reduction of copper. There are also emissions of fluorinated greenhouse gases in this sector.

Carbon dioxide emissions is the major contributor in the sector with 82 %, followed by fluorinated greenhouse gases at approximately 16 %, nitrous oxide at approximately 1.9 % and methane at 0.2 %, counted as CO₂ equivalents.

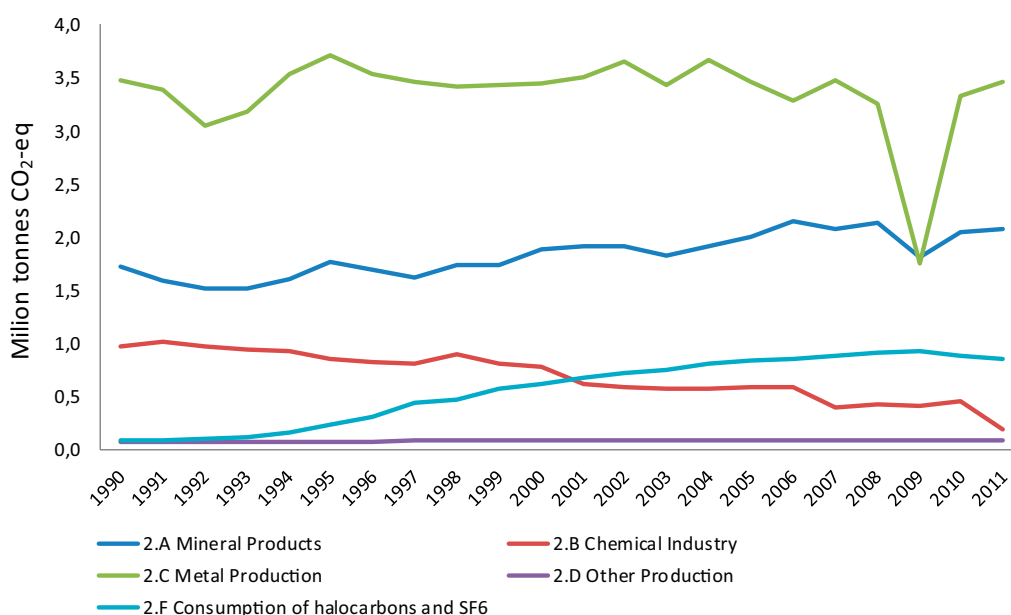


Figure 2-20 Emissions from the Industrial processes sector, per subsector.

Total emissions in this sector have varied somewhat since 1990, mainly due to variation in production volumes and economic fluctuations. Except for 2009, which was an extreme year due to the economic downturn, there has been a slow decrease in total emissions from the sector since 2004. The development, however, differ for different industries. Emissions from the mineral industry have increased, for example, while those from the chemical industry have decreased over the same period. The emissions from the consumption of halocarbons and SF₆ has been increasing since 1992 and starting to stagnate around 2008.

The main subsector is metal production (CRF 2C). Emissions in metal production have been at about the same level during the period from 1990 to 2011, except for 2009. Total emissions from the iron and steel industry have increased by 28 % during the entire period spanning 1990-2008, and emissions have increased most in recent years. The iron and steel industry was severely affected by the economic downturn in 2008-2009, and emissions in 2009 were 38 % lower than in 2008. In 2010 the industry recovered and the emissions were 4 % lower than in 2008. A recalculation of the emissions in metal production will be performed during 2013 which might have an effect on emission trends in the sector during the years from 2005 and onwards.

The subsector mineral products (CRF 2.A) is the second largest subsector and shows an increasing trend between 1990 and 2008. This was principally due to improving economic conditions in the building sector, both in Sweden and in other countries to which cement is exported. Emissions decreased in 2009 as a result of a decline in production due to the economic downturn. However, the emissions increased again in 2010 and 2011 but not to the same level as in 2008.

Emissions from the chemical industry (CRF 2.B) decreased by 80% during the period 1990-2011. The decrease since 2007 is mainly due to a new treatment technology that has been installed and has resulted in reduced emissions of nitrous oxide in the nitric acid production.

2.3.2.1 FLUORINATED GREENHOUSE GASES (HFC, PFC, SF₆)

Fluorinated greenhouse gases, HFC, PFC and SF₆, are used in a number of applications. Most emissions of fluorinated greenhouse gases in Sweden today are due to leakage from refrigeration and air-conditioning systems and air conditioning in vehicles. Other sources are foam manufacturing, medical inhalers, aluminium production and magnesium foundries. Total fluorinated greenhouse gas emissions in 2011 amounted to 1.1 million tonnes calculated as carbon dioxide equivalents and account for almost 2 % of total emissions, see Figure 2-21.

Emissions of fluorinated gases are increasing from 1992 to 2007. After 2007 there is a decrease in emissions, with 2009 showing low emissions due to the economic downturn. The increase in the total emissions of fluorinated greenhouse gases is due to the increase in HFC emissions. One explanation for the increase is that HFCs in many cases have replaced the ozone-depleting substances CFCs and HCFCs as refrigerants. The ozone-depleting substances decreased sharply between 1990 and 2009 and have now been more or less completely phased out. Another explanation for the increase in HFCs is that the number of refrigeration and air-conditioning systems, air conditioning in vehicles and heat pumps has increased, particularly in the recent years.

PFC emissions mainly originate from the aluminum industry, and these emissions have decreased in recent years as a result of investments in new technology since 2007. The new technology was used exclusively in 2009, and in combination with lower production as a result of the weakness in the economy, this led to a sharp decrease in emissions.

SF₆ emissions have remained at approximately the same level but have varied somewhat but shown a decreasing trend from 2007. One explanation to the decrease is an EU Regulation (EG 842/2006) introduced in 2006. Under this Regulation, certain fluorinated greenhouse gases were banned at various times between 2006 and 2009. Other emissions of fluorinated greenhouse gases have also started to decrease as a result of the EU Regulation.

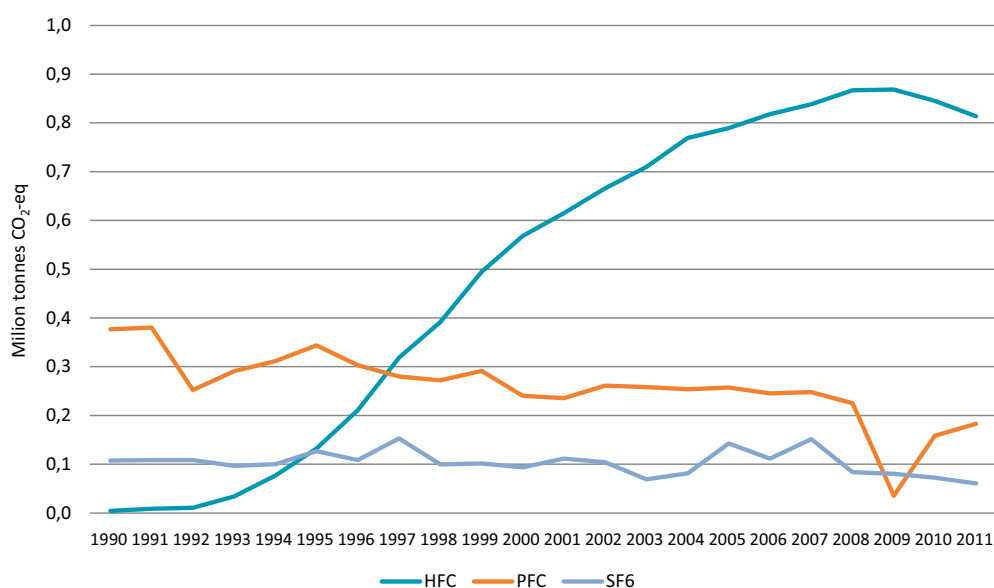


Figure 2-21 Emissions of fluorinated greenhouse gases per gas.

2.3.3 Solvents and other products use

The use of solvents leads to emissions of volatile organic compounds. The carbon content of these emissions is assumed, according to the reporting guidelines, to be oxidised to carbon dioxide. The use of other products such as spray cans and gas springs, leads to emissions of nitrous oxide. Emissions of carbon dioxide and nitrous oxide calculated as carbon dioxide equivalents in 2011 totalled 0.3 million tonnes, which is 0.5 % of the total greenhouse gas emissions, see Figure 2-22. In comparison with 1990, total emissions from this sector have decreased by 13 %.

As shown in Figure 2-22 the emission trends in the subsectors differs between sectors. Subsector Other (3.D) is increasing over the period while the emissions in paint application (3.A) is decreasing. The decrease in subsector paint application is due to a shift to water-based paints.

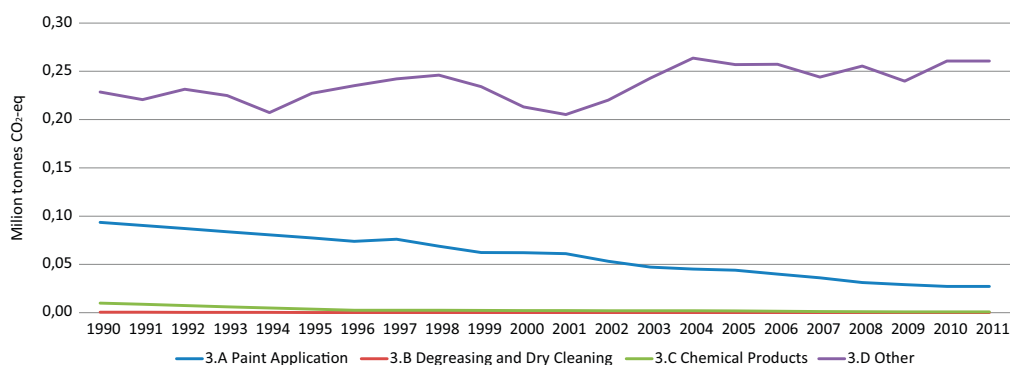


Figure 2-22 Emissions from the use of solvents and other products, per subsector.

2.3.4 Agriculture

Agricultural activities contribute directly to emissions of greenhouse gases through a variety of processes in which methane (CH_4) and nitrous oxide (N_2O) are the primary greenhouse gases emitted. In addition, agriculture is the largest source of methane and nitrous oxide emissions. The main sources of emissions of these two gases in Sweden are domestic livestock activities with enteric fermentation in domestic livestock, livestock manure management and agricultural soil management activities, such as fertilizer application. Cattle produce the major part of methane emission from enteric fermentation, while other types of livestock are relatively of low significance in Sweden. Activities related to agricultural soil management are the major source of N_2O emissions. Emission from burning of agricultural residues does not exist as this activity is legally prohibited. In addition rice is not cultivated and savannah does not exist in Sweden.

The Agriculture sector contributes with about 13 % (about 7.8 million tonnes $\text{CO}_2\text{-eq}$) of the overall greenhouse gases in 2011, of which around 63 % was made up of nitrous oxide and almost 37 % of methane.

Aggregated emissions from various agricultural activities decreased by 6 % (0.54 million ton $\text{CO}_2\text{-eq}$) over the period 2000-2011, and they have fallen by 13 % (1.2 million ton $\text{CO}_2\text{-eq}$) since 1990, see Figure 2-23. The main reasons for the decreasing trend are: firstly the number of cattle has decreased, resulting in lower methane release, and secondly lower application of nitrogen fertiliser to agricultural land has resulted in decreased release of nitrous oxide.

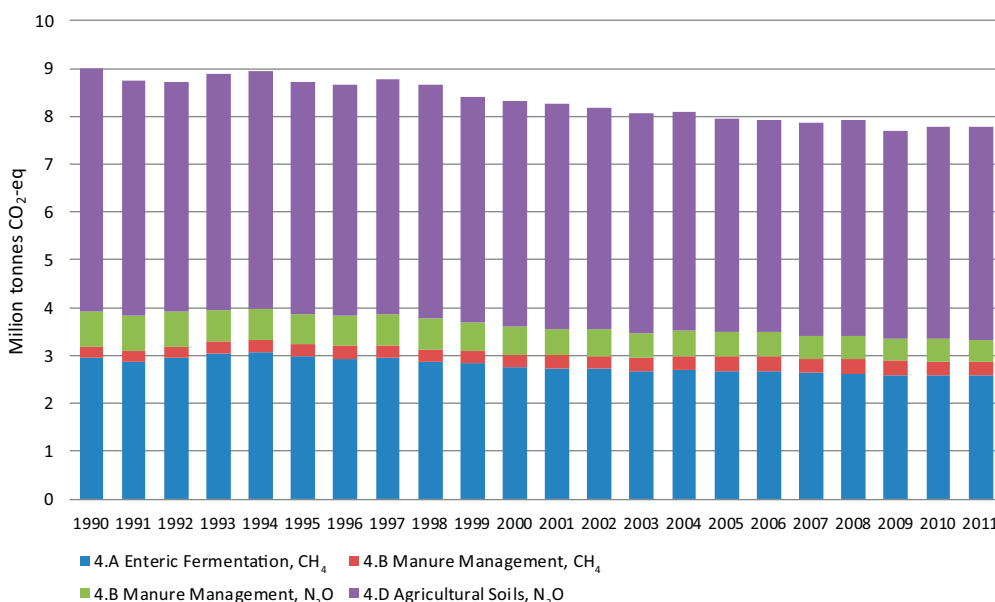


Figure 2-23 Emissions from agriculture, total and per gas and subsector.

2.3.4.1 ENTERIC FERMENTATION AND MANURE MANAGEMENT (CRF4.A AND 4.B)

Methane emissions come principally from the digestion and manure of cattle, while other types of livestock are of relatively low significance. In 2011, about one-third (33 % or 2.6 million ton CO₂-eq.) of the emissions in agriculture are related to digestion of livestock, i.e. enteric fermentation, see Figure 2-23. Methane emission due to enteric fermentation has decreased by about 12 % over the period 1990-2011. The most important reason for the decreased emissions is reduced activities within livestock farming. For example, the population of cattle has decreased by 12 % from 1.718 million head to 1.537 million head over the period 1990-2011.

Emission of methane and nitrous oxide from manure of cattle, swine, horses, goats, sheep, poultry and reindeer are included in the Swedish inventory. Emissions related to manure management constitute less than 10% (or 0.75 million ton CO₂-eq.) of the total emission of the sector. Emissions have decreased by about 23% since 1990. This is due to the fact that the quantity of farmyard manure is declining principally as a consequence of the decreasing number of dairy cows. The expansion of slurry management for pigs and dairy cows has also reduced emissions.

A large reduction in dairy cattle population (about 8 %) took place in 1990 and 1991, when a large number of farms abandoned milk production. Some of these farms changed to extensive meat production with the aid of government conversion grants, and the number of beef cattle therefore, increased by about 15% during the first half of the 1990s. Following Sweden's accession to the EU in 1995, the EU's Common Agricultural Policy (CAP) stabilised livestock numbers for livestock that have the right for subsidies, for example, cattle. The long-term trend is nevertheless for a successive decrease in both cattle and swine, while the number of horses, sheep and chickens for slaughter has increased. Overall, this signifies a decrease in methane emissions from both livestock and their manure. It is worth to mention that methane emissions per dairy cow in 2011 have increased by about 10 % compared to 1990, due to increased milk yield, greater quantity of manure and a higher proportion of slurry management. Milk yield per cow in 2011 has increased by one-third since 1990. The average milk yield per cow is about 9000 litres per year.

2.3.4.2 AGRICULTURAL SOILS (CRF 4.D)

Where large applications of fertilizer are combined with soil conditions favorable to denitrification, large amounts of nitrous oxide can be produced and emitted to the atmosphere. The emission of nitrous oxide originates as a result of microbial nitrification-denitrification processes in agricultural soil. Similarly, the often poorly controlled use of animal waste as fertilizer can lead to substantial emissions of nitrous oxide from agricultural soils.

A major source of nitrous oxide emission from agricultural soils comes principally from the supply and conversion of nitrogen from synthetic fertilizer use, animal waste and sewage sludge use as fertilizer. The need for greater crop

yields is the main driver to a large increase in the use of fertilizers. Cultivation of peat soils, also results in significant release of nitrous oxide, as does conversion of the nitrogen that leaches to lakes and watercourses.

In 2011, N₂O emissions from agricultural soils were responsible for about 57 % (or 4.4 million ton CO₂-eq.) of the total emissions of the agriculture sector (Figure 2-23). The emissions show a significant long-term trend as they are highly sensitive to the amount of nitrogen applied to soils. The emissions have decreased by about 12 % since 1990 due to decreased application of mineral fertilizer and farmyard manure.

2.3.5 Land Use, Land Use Change and Forestry – LULUCF

The Sector Land Use, Land Use Change and Forestry (LULUCF) constituted an annual net removal in Sweden during the whole period 1990-2011, see Figure 2-24. During the period the net removals have varied between 27 and 38 million tonnes of carbon dioxide equivalents but the long-term trend points to a small decrease in net removals from the sector. The decrease in net removal is mainly due to an increased harvest rate and over the last years the decrease was also enhanced due to the effect of two severe storms in the beginning of 2005 and in 2007, respectively. The storm in 2005 brought down a large quantity of timber. According to the Swedish National Board of Forestry, gross fellings ranged between 64 Mm³sk and 96 Mm³sk over the period 1990 to 2011, with the exception of 2005 when felling, including wood felled by storms, was estimated at 122 Mm³sk.

However, the uncertainty is greater in data for 2011 than 2008 since the number of sample plots gradually decreases, and extrapolations are needed for a major part of the sample plots.

The total size, variation and trend of the net removals in the LULUCF-sector are mainly affected by the carbon stock change in the Forest land. The carbon stock change in living biomass in the forest counts up for the major part of these net removals. The net removal in living biomass in Forest land varied between approximately 22 and 39 million tonnes of carbon dioxide during the period 1990-2011 with the highest figures in the beginning of the period and with a small decrease during the last years. Cropland is responsible for emissions of carbon dioxide when organogenic soils are cultivated. Emissions varied during the period 1990 to 2011 between 1 and 2.5 million tonnes of carbon dioxide. The subsectors grassland, wetlands and settlements account for very small areas (and associated carbon stock changes) compared to forest land, which leads to higher uncertainty in data. The carbon stock change in grassland and wetlands is small. Emissions from settlements were in the range of 1-2.9 million tonnes of carbon dioxide during the period 1990-2011.

The net removal in the LULUCF-sector is calculated as the total carbon stock change in the three carbon pools of living biomass, dead organic matter (dead wood and litter including the humus layer of soil) and soil organic carbon for different land use categories. The three carbon pools living biomass, dead

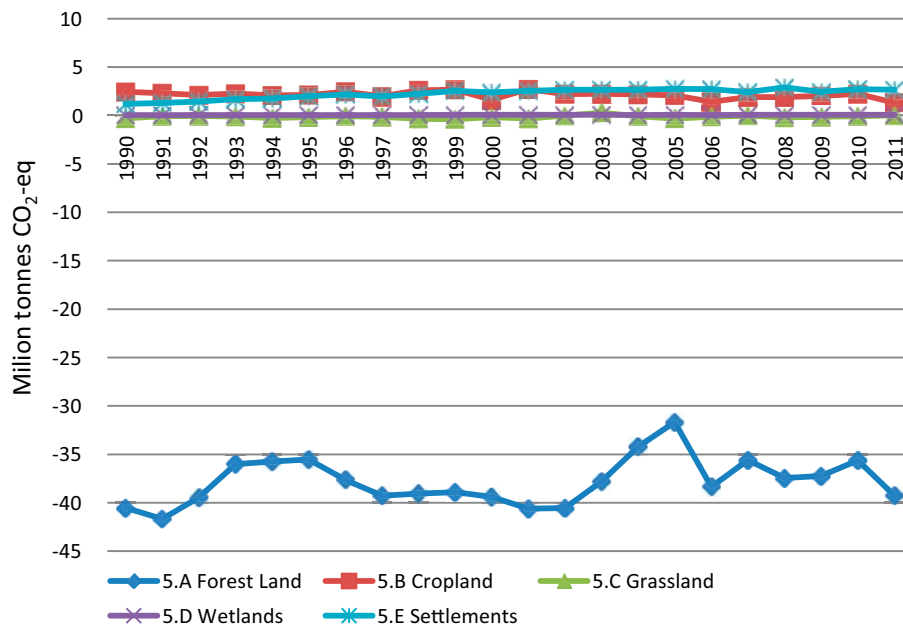


Figure 2-24 Emissions (+) and removals (-) of greenhouse gases from the LULUCF sector.

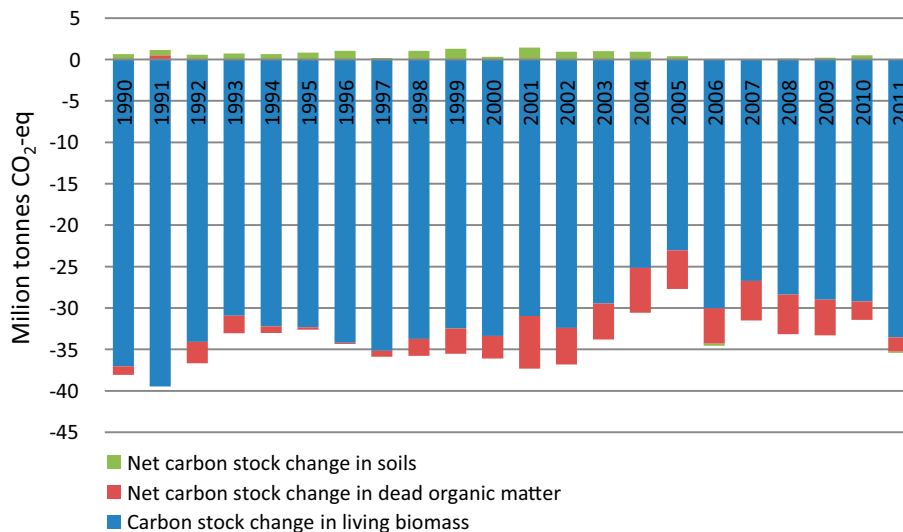


Figure 2-25 Emissions (+) and removals (-) of carbon dioxide from different carbon pools.

organic matter and soil organic carbon contributes as an aggregate to a net removal, see Figure 2-25. In addition, emissions of N_2O from fertilization and disturbance associated with conversion to cropland, CO_2 emissions from lime application and N_2O , CH_4 and CO_2 from biomass burning are calculated and contributes to a very small source with small emissions.

2.3.6 Waste

Total emissions from the waste sector (CRF 6) in 2011 amounted to around 1.7 million tonnes of carbon dioxide equivalents or almost 3 % of total greenhouse gas emissions. In comparison with 1990, emissions were about 50 % lower in 2011. Emissions from the waste sector come mostly from landfills (about 70 %) are dominated by methane emissions while emissions from waste-water account for 26% and the rest of the emissions come from incineration of hazardous waste. Emissions by subsectors are shown in Figure 2-26.

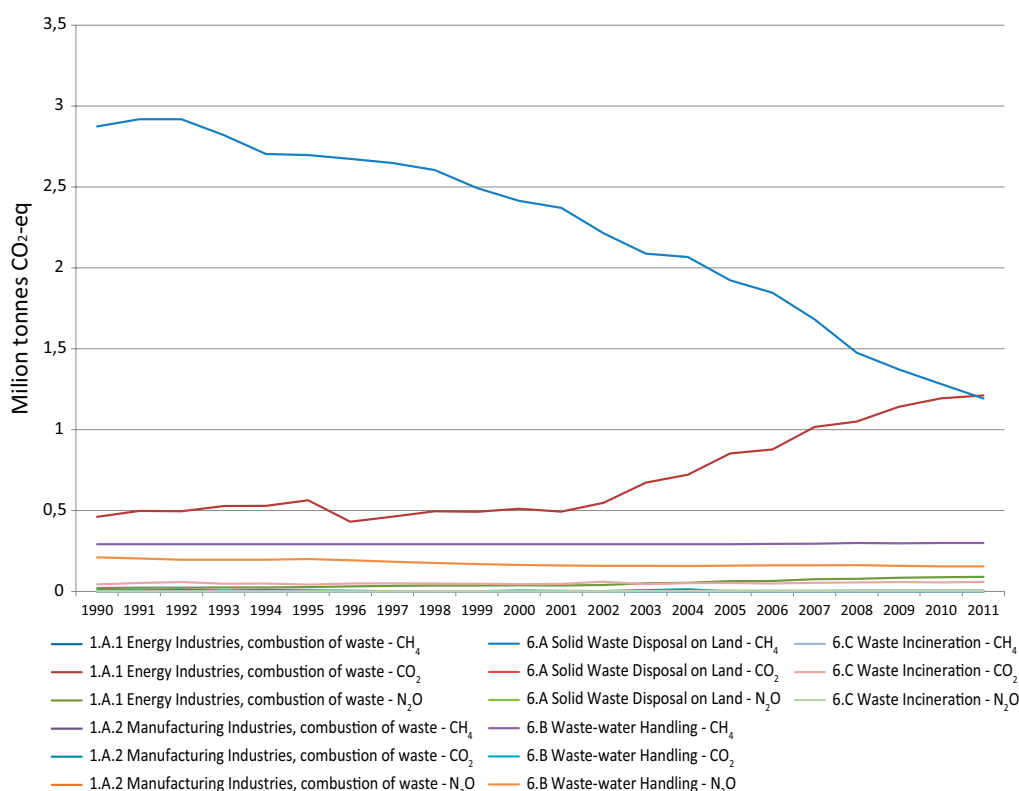


Figure 2-26 Emissions from the waste sector, total and per subsector.

Landfills are the largest source for emissions of methane gas, after livestock farming, as methane is formed when organic waste is placed in landfills. Methane emissions have declined steadily since the early 1990s, partly because the amount of organic material in landfills has declined and partly due to the increased collection and management of methane gas from landfills.

Several policy instruments have been significant in this trend. During the 1990s there was, for instance, the introduction of producer responsibility for a number of different groups of articles, for example packaging, waste paper, office paper and tyres. It is also believed that the demand for municipal waste planning that was introduced in 1991 has contributed to the expansion of methane collection and to the reduction in the amount of degradable material deposited in landfills.

A tax on landfill waste was introduced in 2000, and bans on the landfill disposal of combustible waste (in 2002) and organic material (in 2005) have subsequently been introduced. The bans have contributed to a substantial change in waste management in Sweden. The landfilling of other types of waste has also fallen sharply.

Emissions from wastewater handling were less than 0.5 million tonnes of carbon dioxide equivalents in 2011 and accounted for 0.7 % of national total emissions. Emissions have fallen by about 10 % since 1990 due to improved sewage treatment.

Emissions from incineration of waste were around 0.065 million tonnes CO₂ equivalents in 2011. Emissions have increased somewhat in recent years in comparison with the level of emissions from 1990 to 2002. The increase in emissions is due to an increased quantity of waste being incinerated as capacity has increased.

2.3.7 International bunkers

Emissions of greenhouse gases from international bunkering of fuels amounted to around 8.3 million tonnes of carbon dioxide equivalents in 2011, see Figure 2-27. This includes refuelling in Sweden by international navigation and international aviation and emissions from this fuel are not included in the reporting of the total emissions from Sweden which is calculated in the Kyoto Protocol commitments. International bunkering of fuel is substantially greater than fuel use for domestic navigation and aviation.

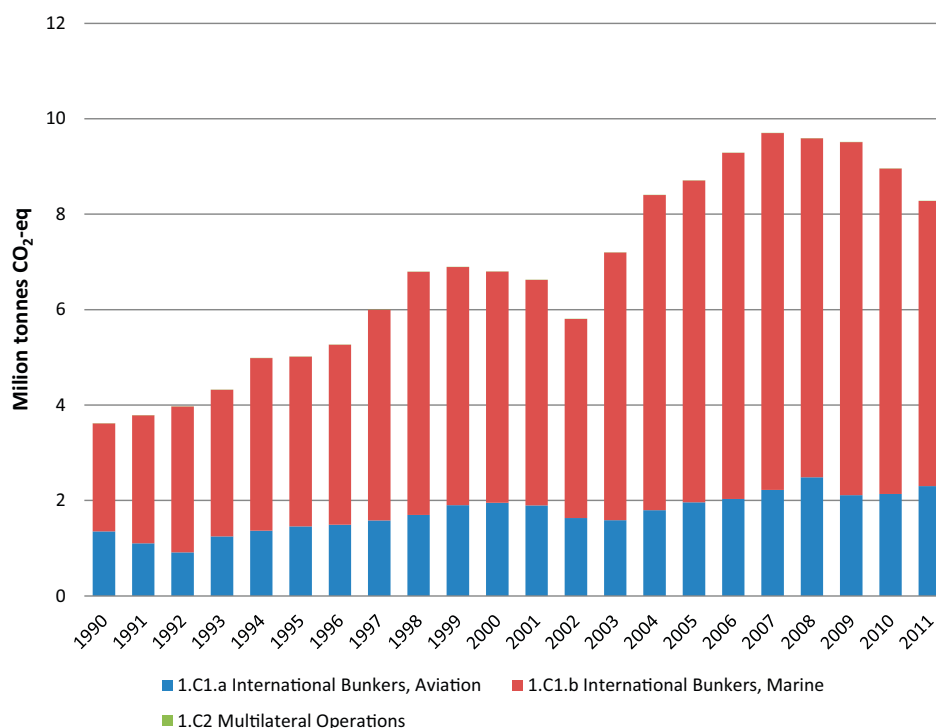


Figure 2-27 Emissions from international bunkers, total and per subsector.

Emissions from international navigation totalled 6.0 million tonnes of carbon dioxide equivalents in 2011. This is a decrease of 12% compared with 2010 but an increase of 164% since 1990. Part of the explanation for the increase is that international freight transport activity has increased due to an increased quantity of freight and globalisation of trade and the systems of production having led to freight being transported over longer distances. Another explanation could be that Swedish refineries are producing low-sulphur residual fuel oil, which fulfils strict environmental requirements, and this has led to more shipping lines having chosen to refuel in Sweden. The fluctuations in bunkered volumes between different years also depends on the price of fuel in Sweden in comparison with other ports in other countries.

Greenhouse gas emission from international aviation bunkers was 2.3 million tonnes of carbon dioxide equivalents in 2011. This is an increase by 8 % compared with 2010 and an increase of 70 % since 1990. Emissions from international bunkering of aviation have varied over time. The long-term trend is increasing although there have been declines in the early 1990s as well as at the beginning of the 2000s and in 2009.

2.4 Description and interpretation of emission trends for indirect greenhouse gases and SO₂

2.4.1 NMVOC

In 2011, the emissions of non-methane volatile organic compounds (NMVOCs) totalled around 177 ktonnes. The energy sector (mainly road traffic, wood combustion in the residential sector) and solvent and other product use sector are the dominant sources of emissions contributing with 48 % and 44 %, respectively Figure 2-28. In addition, some industrial activities (such as pulp and paper, and food and drink) are also significant for NMVOC emissions.

The emissions have declined sharply by almost one-half since 1990. The decline is clearly visible in the energy sector and solvent and other product use sector which accounts for about 63 % and 2 %, respectively, compared to 1990.

Emission from road traffic shows the greatest reduction in emission (about 82 % since 1990) due to the introduction of new exhaust emission requirements. In addition, the voluntary environmental standards for new installations of wood-fired boilers and reduced emissions from products containing solvents have also contributed to lower emissions during the last two decades. Fugitive emissions from oil refineries have also declined by more than one-half since 1990.

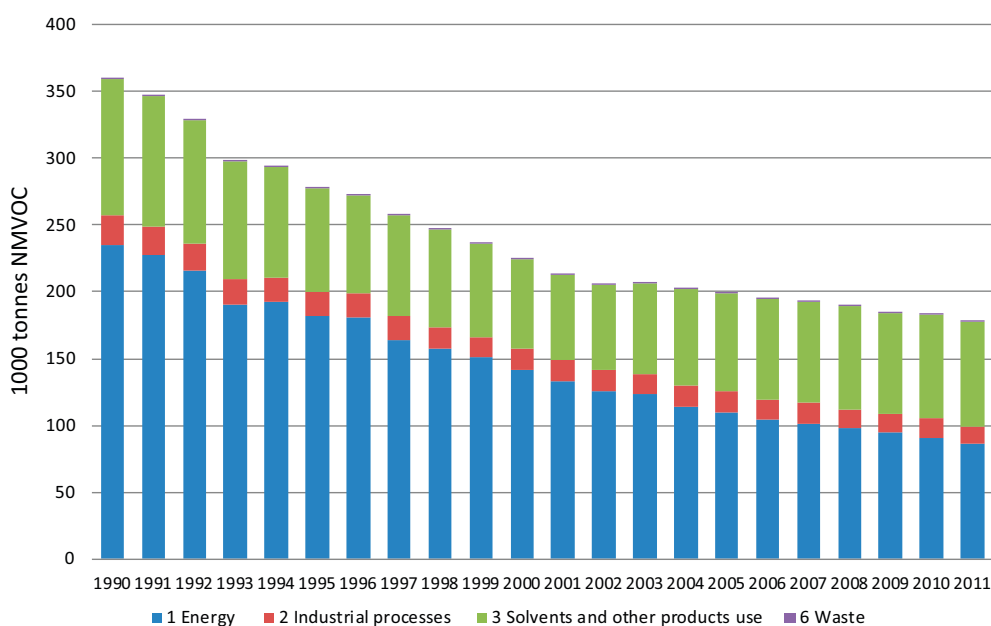


Figure 2-28 Total emissions of NMVOC and emissions from different sectors.

2.4.2 NO_x

Emissions of nitrogen oxides (NO_x) amounted to about 146 ktonnes in 2011, a decrease by 46 % in comparison to 1990, see Figure 2-29. Nitrogen oxides are formed in all combustion in the energy and transport sectors, and the largest emission sources are road traffic, machinery, navigation and production of electricity and heating. Emissions of nitrogen oxides from the energy sector were about 131 ktonnes which accounts for about 90 % of the total emissions. However, emissions from transport (within the energy sector) were about 72 ktonnes or about 50 % of total emissions. NO_x emissions from the energy sector have declined by about 48 % since 1990 whereas emissions from transport have declined by about 54 %.

Traffic is a large source of emissions of nitrogen oxides, and the emissions come largely from road traffic which accounts for 61 ktonnes in 2011. The introduction of catalytic converters in cars and the subsequent successive tightening of exhaust emission requirements have contributed to a general decrease in concentrations of nitrogen oxides in urban areas. Road-traffic emissions of NO_x decreased by 57 % between 1990 and 2011 and decreased by 7 % between 2010 and 2011.

Approximately 10 % of NO_x emissions in the energy sector in 2011 came from electricity and district heating production. Annual NO_x emissions from this source show significant long-term trend fluctuations between 1990 and 2011, although overall emissions were 1 % lower in 2011 than in 1990. The trend of NO_x emissions from this source is related to weather patterns in Sweden, such as (temperature and precipitation) and consequently the need for heating, which affects the need for combustion-based production of electricity.

The next largest source of emissions is the industrial processes sector which accounts for about 10 % of NO_x emission mainly from pulp and paper industry.

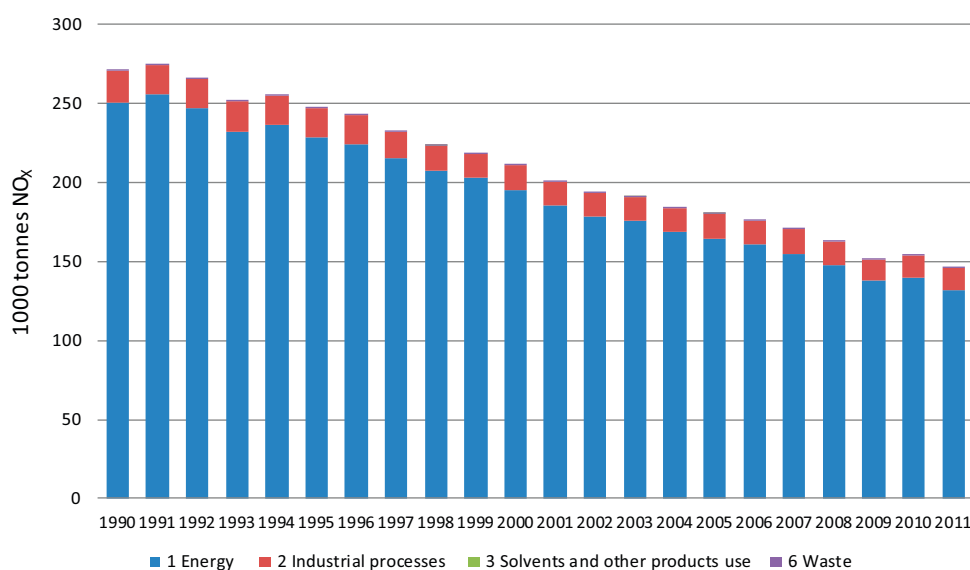


Figure 2-29 Total emissions of NOx and emissions from the different sectors.

2.4.3 CO

Emissions of carbon monoxide have developed in the same way as NOx emissions. Emissions have decreased by about one-half, from 1,280 ktonnes in 1990 to 571 ktonnes in 2011, see Figure 2-30.

Around 96 % of emissions come from the energy sector in which transport and household energy are the largest sources and contribute by about 40 % and 36 %, respectively. Carbon monoxide emissions from the energy sector decreased from around 1,258 ktonnes in 1990 to around 547 ktonnes in 2011.

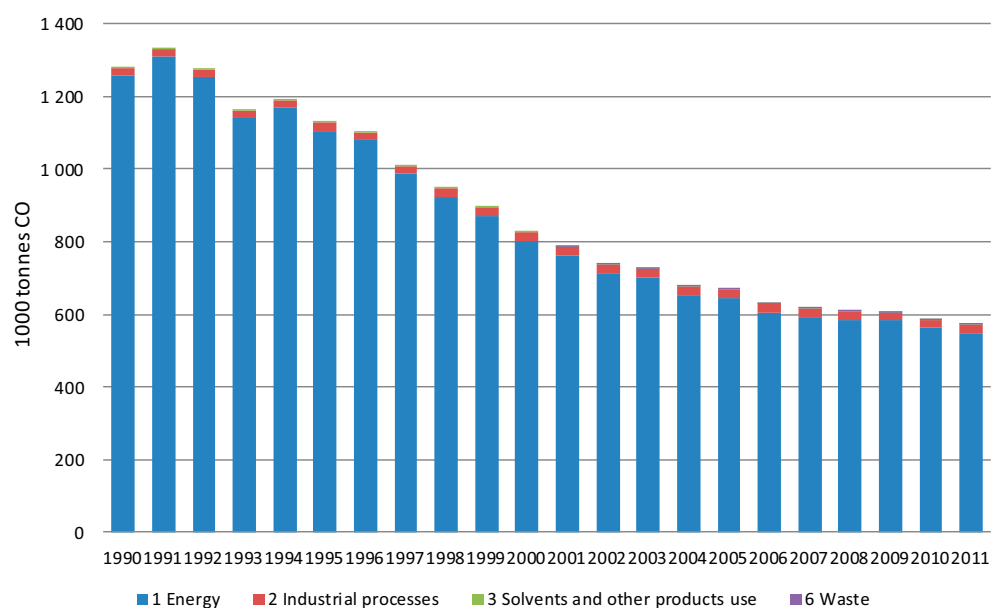


Figure 2-30 Emissions of CO from different sectors.

2.4.4 SO₂

Sulphur dioxide (SO₂) emissions come from energy, transports industrial processes and combustion of hazardous waste. Emissions are declining since 1990, see Figure 2-31. In 2011, emissions totalled about 30 ktonnes, a decrease by about 72 % compared to 1990. The continued decrease is due to a shift from fuels with high-sulphur levels to low-sulphur fuels, for vehicles, industry and production of electricity and district heating.

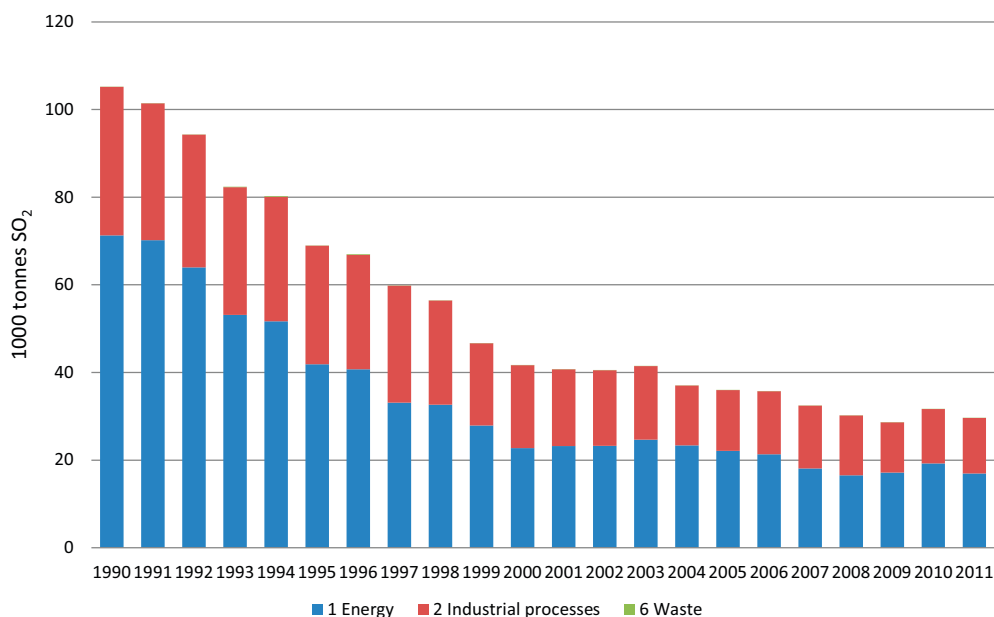


Figure 2-31 Total emissions of SO₂ and emissions from different sectors.

Energy sector emissions of sulphur dioxide (excl. transport) continued to decrease during the 1990s and in 2011 emissions totalled 17 ktonnes, a decrease of 76 % compared with 1990. The continued decrease is due to a shift from fuels with higher sulphur levels to low-sulphur fuels, both for industry and for production of electricity and district heating. The sulphur tax introduced in 1991 has been significant in this shift. Other factors which contribute to reduced emissions include the consideration of industries under the Environmental Code.

Road traffic emissions of SO₂ have fallen by 99 % since 1990 as a result of lower sulphur levels in motor fuels, and totalled 0.1 ktonnes in 2011. Sulphur emissions from domestic navigation have decreased by 73 % since 1990 and are now 1.4 ktonnes due to a switch to oils with lower sulphur content.

2.4.5 Description and interpretation of emission trends for KP-LULUCF inventory in aggregate and by activity, and by gas

Since the KP-LULUCF has been reported for only three years there is not so much of a trend to describe. However, the reporting under Forest management is strongly linked to the reporting of Forest land remaining forest land under the UNFCCC-reporting so to get a picture of the long term trend it is recommended to read section 2.3.5 which describes the trend in the LULUCF-sector.

3 Energy (CRF sector 1)

3.1 Overview of sector

The energy sector includes emissions from fuel combustion (CRF 1.A) and fugitive emissions from fuel production and handling (CRF 1.B). Energy consumption per capita is high in Sweden compared to other OECD countries. This is because of the availability of natural resources such as forests and hydropower, which led to the early and rapid expansion of energy-intensive industries. Sweden's geographical location, with low mean annual temperatures also explains the high demand for energy for heating. The energy sector, including transport, has long accounted for the major part of Swedish greenhouse gas emissions, and emissions of carbon dioxide dominate overwhelmingly in this sector. However, carbon dioxide emissions per capita are relatively low in Sweden compared with other industrialized nations. This is due to a relatively high use of hydropower and nuclear power and low use of fossil fuels, as well as the use of energy and carbon dioxide taxation for limiting the emissions of carbon dioxide.³³

It can be seen in Figure 3-1 that in the energy sector, emissions of CO₂ contribute about 96 % of total greenhouse gas emissions (in CO₂ equivalents) 2011. Emissions of total greenhouse gases from the energy sector have decreased by 16.1 % from 53,670 Gg CO₂ equivalents in 1990 to 45,015 Gg CO₂ equivalents in 2011, mainly due to reduced fossil fuel consumption in the residential sector (CRF 1.A.4) (Figure 3-2).

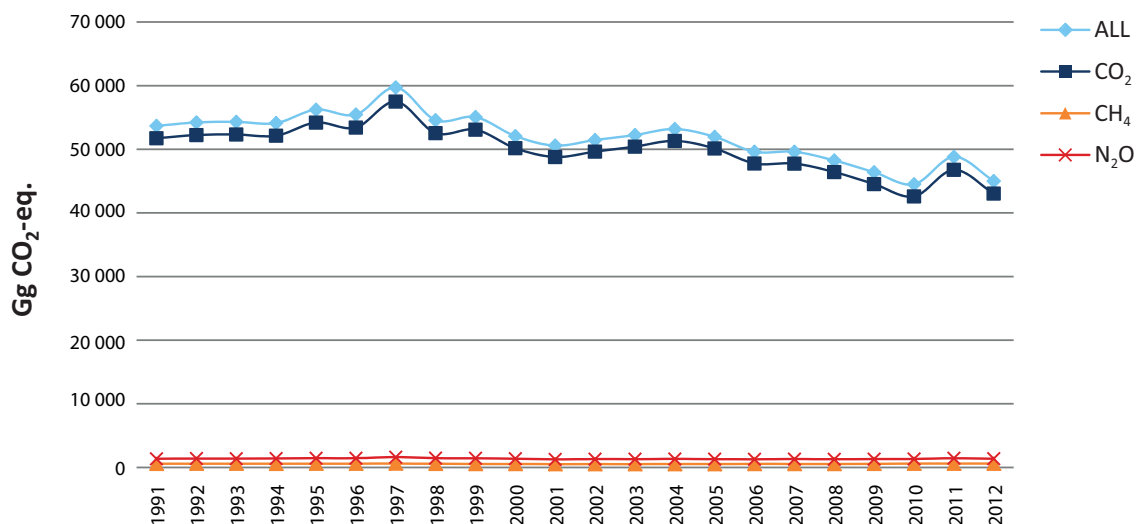


Figure 3-1. Total emissions of all greenhouse gases calculated as CO₂ equivalents from CRF 1 Energy.

³³ Ministry of the Environment, 2001

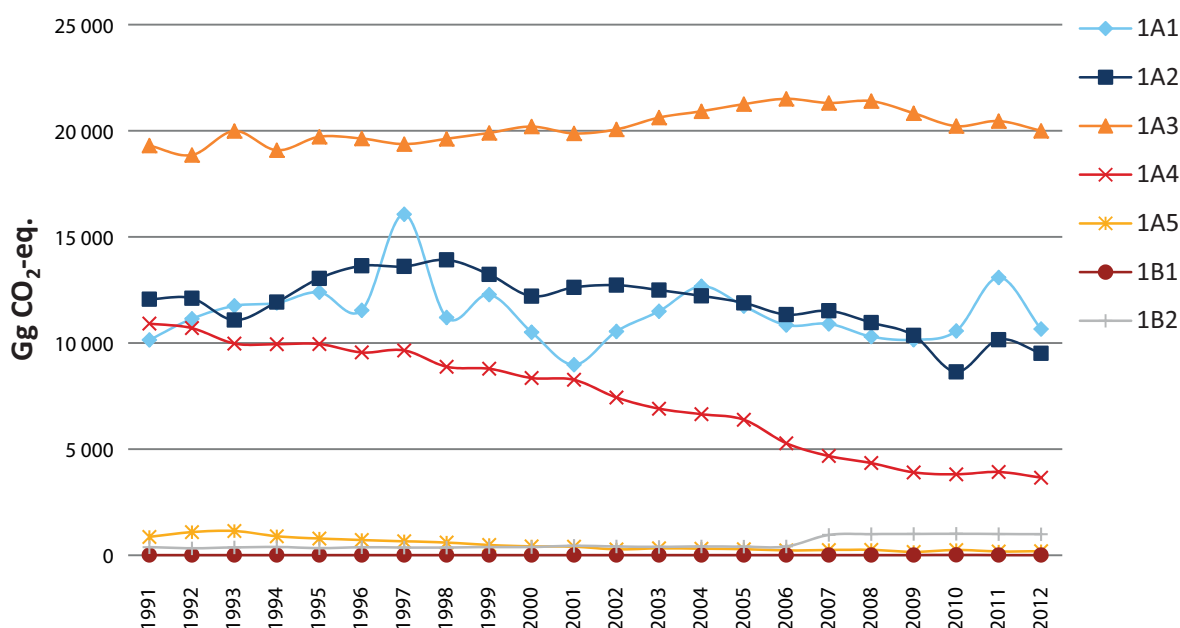


Figure 3-2. Total emissions of all greenhouse gases calculated as CO₂ equivalents from the different sub-sectors within the Energy sector.

1A1 Energy industries. 1A2 Manufacturing industries and construction. 1A3 Transport. 1A4 Other sectors. 1A5 Other. 1B1 Solid fuels (fugitive). 1B2 Oil and natural gas (fugitive).

As shown in Figure 3-2, the transport sector (CRF 1.A.3) accounts for the largest, and increasing, part of the GHG emissions from the energy sector. Emissions from public electricity and heat production (CRF 1.A.1) varies mainly because of temperature variations between years. As mentioned earlier, the emissions from residential heating (CRF 1.A.4) are decreasing due to reduced consumption of fossil fuels. In manufacturing industries and construction (CRF 1.A.2), the three largest industries in terms of fuel consumption are the pulp and paper industry, the chemical industry and the iron and steel industry. Despite rising industrial production, oil consumption has fallen sharply since 1970. This has been possible due to increased use of electricity and improved energy efficiency.

The large increase in emissions from CRF 1.A.1 in 2010 is mostly due to the cold winters that year and low production of nuclear energy, which meant that the demand of electricity and heat had to be met by combustion based energy. In 2011, conditions were less extreme and emissions especially from electricity and heat production decreased considerably. The dip in emissions from manufacturing industries and construction in 2009 reflects the economic conditions resulting in lower demand of e.g. iron and steel. The recent increase in fugitive emissions from oil and natural gas (CRF 1.B.2) is caused by hydrogen production facilities put into operation at two of the oil refineries in 2005 and 2006 respectively.

Table 3-1 shows the impact of recalculations reported in submission 2013 for GHG emissions by sector and sub-sectors for 1990, 1995, 2000 and 2005-2010. The recalculations are mainly due to:

- Revised activity data for residual fuel oil in 1.AA.3.D for 2009 and 2010 and corresponding emissions recalculated.
- Emissions of CH₄ and CO₂ from venting of natural gas in the transmission network was added to the already reported fugitive emissions in CRF 1.B.2.B.3. This is in line with IPCC GPG. Default emission factors from GPG were used to estimate venting emissions.
- Revised activity data for the Other sectors (1.A.4 and parts of 1.A.2) for 2009-2010
- Combustion of coke for production of heat used in carbide manufacturing has been included in CRF 1.AA.2.C in submission 2013

More detailed descriptions of the recalculations are found under sector specific sections below.

Table 3-1 Impact of recalculations of GHG emissions submission 2013 in the energy sector (1990, 1995, 2000, 2005-2010).

Impact of recalculations submission 2012 (Gg CO ₂ eq.)									
CRF	1A1	1A2	1A3	1A4	1A5	1B1	1B2	Total CRF 1	% CRF 1
1990	NA	22	-3	2	35	NA	8	63	0.1%
1995	NA	18	-1	2	-6	NA	8	22	0.0%
2000	NA	14	13	-7	-4	NA	11	27	0.1%
2005	NA	17	15	-5	-2	NA	14	39	0.1%
2006	NA	5	17	-8	-2	NA	15	28	0.1%
2007	0	9	61	-4	-2	NA	16	80	0.2%
2008	0	7	-29	13	1	NA	16	9	0.0%
2009	-2	2	-133	-16	1	NA	16	-132	-0.3%
2010	0	33	-285	-317	-1	NA	16	-554	-1.1%

0: value less than 0.5. NA: no recalculation is performed.

3.2 Fuel combustion (CRF 1.A)

Emissions from fuel combustion, CRF 1.A, are allocated to a number of sub-sectors. In CRF 1.A.1, emissions from energy industries, e.g. public electricity and heat production plants, combustion activities within oil refineries, and combustion related to solid fuel production, i.e. coke ovens, are reported. CRF 1.A.2 includes combustion-related emissions in manufacturing industries and construction. Emissions from working machinery within the construction sector are allocated to CRF 1.A.2, but apart from that, CRF 1.A.2 includes only stationary combustion. Emissions from transports (aviation, road traffic, railways and navigation) are reported in CRF 1.A.3. Only domestic activities are included in CRF 1.A.3; emissions from international aviation and navigation are allocated to the category 1C1, international bunkers, and not included in

the national total. In CRF 1.A.4, emissions from households, service, agriculture, forestry and fisheries are reported. CRF 1.A.5 is intended for “other” combustion, which in the Swedish inventory includes emissions from military operations. Flaring, e.g. combustion where the energy is lost and not used, is reported in CRF 1B.

Emissions from fuel combustion in Sweden are, if not specifically otherwise stated, determined as the product of fuel consumption, thermal value and emission factors (EF) as shown in the formula:

$$\text{Emissions}_{\text{fuels}} (\text{unit}) = \sum \text{Fuel consumption} (\text{unit}) * \text{thermal value}_{\text{fuels}} * \text{EF}_{\text{fuels}}$$

Different Tiers are used for different sub-sectors as discussed in sections below.

Please note that some fuel types are used in industrial processes rather than for energy purposes. This is the case for black liquor in the paper- and pulp industry and for coal and coke in the metal industry. Emissions from these fuels are thus accounted for under CRF 2 and methods used are described in section 4.

3.2.1 Comparison of the sectoral approach with the reference approach

A detailed discussion on the reference approach and its differences compared to the sectoral approach is provided in Annex 4. Figure 3-3 shows the differences in fuel consumption and CO₂ emissions between the Reference and Sectoral Approach for the whole time series 1990-2011.

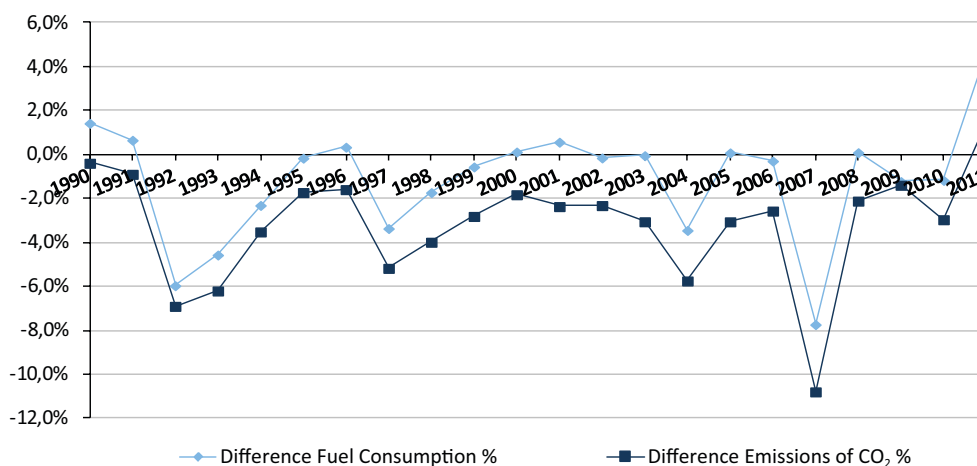


Figure 3-3 Differences between Reference Approach and Sectoral Approach (Reference minus Sectoral).

It is obvious that fuel consumption and CO₂ emissions from the Sectoral Approach exceed the Reference Approach for most years. For a number of years the difference is larger than ± 2 %. Data on fuel group level indicates that this problem is primarily related to liquid fuels. Statistical differences in energy balances contribute to a large share of differences especially in the early 1990's. Large differences for 1997, 1998, 2001 and 2007 are mainly attributed to liquid fuels and most notably crude oil and refinery feedstocks. For solid fuels, the difference in energy and emissions, respectively, is not consistent between years. On total level, this has the effect that differences expressed as energy consumption are less negative than differences expressed as CO₂ emissions, as shown in Figure 3-3. This is related to solid fuels used in the iron and steel industry. Large amounts of energy are lost in coke ovens and blast furnaces when coking coal is transformed to coke and coke oven gas, and coke is transformed to blast furnace gas. This means that the reported amounts of coke oven gas, blast furnace gas and steel converter gas reported in the sectoral approach contains all the carbon but only parts of the energy reported as coal in the reference approach, which gives a large difference between the reference and sectoral approach for consumption of solid fuels. However, the time series for both liquid and solid fuels still show very large variations after this correction. For gaseous fuels, the differences are low for most years, except for the period 2004-2008 when the amounts of natural gas used as feedstock cannot be reported due to confidentiality reasons. Further explanations are given in Annex 4.

3.2.2 International bunker fuels

This sector includes emissions from refuelling in Sweden used for international navigation and international aviation. All gases are covered.

According to IPCC guidelines, international bunkers are not included in national totals. To evaluate Swedish emissions, international bunkers are of course important, especially as international bunkering of fuel is substantially greater than the fuel use for domestic navigation and aviation. Emissions have increased significantly since 1990 due to among other things increased travelling and increased transportation of goods. See also section 2.

3.2.2.1 INTERNATIONAL BUNKERS, CRF 1C1

Emissions from international bunkers for aviation and navigation are not included in the national total, but instead reported separately as a memo item in CRF 1.C.1, in accordance with the 1996 revised IPCC Guidelines.

The ERT (Expert Review Team) has noticed that the data reported to the IEA are generally higher than what is reported to the UNFCCC. A study in 2010 showed that the differences between the IEA (International Energy

Agency) and the UNFCCC reporting can to some extent be explained by revision policies of the different reporting obligations. Since the UNFCCC has a high demand on consistency of time series, efforts are made to ensure high quality of times series.³⁴

International bunkers from aviation are fuels purchased in Sweden and used for flights to destinations abroad. This includes the whole flight cycle, i.e. both LTO (Landing and Take-Off) and Cruise, see also Annex 2.

Activity data from the Swedish Transport Agency (STA), former Swedish Civil Aviation Authority (SCAA), starts from 1994. With regards to bunkering the data has been estimated using 1994 for CO₂ emissions. The share of LTO is estimated using an average for the years 1995 to 2001.

International bunkers from navigation are defined as fuels bought in Sweden, by Swedish or foreign-registered ships, and used for transport to non-Swedish destinations. The division on international and domestic fuels is based on information from the monthly survey on supply and delivery of petroleum products.

In 2011, the fuel consumption by national and international navigation has been studied in a project and the results are presented in the report “Emissions from navigation and fishing including international bunkers”³⁵. Fuel data in the Monthly fuel, gas and inventory statistics has been analyzed. The fuel data is collected from oil companies and other providers of petroleum products and coal. The survey also collects stock data from companies with a large consumption of oil in the manufacturing industries and energy industries. The population consists of approximately 70 companies, and all of them are included in the survey.

In 1993, 24 companies reported fuel amounts for domestic navigation and 15 companies reported amounts for international maritime bunkers. During the period 1993-2010 many companies have been closed or taken over by larger companies. In 2010, very few companies remained – only eight companies reported fuel amounts for Domestic navigation or International maritime bunkers. With such a small population, it was possible to study each company carefully. Suppliers of significant quantities of bunker oil (~100,000 m³ – 2,000,000 m³) have been examined more closely.

The response from the companies was very good and produced reliable information. Data on domestic and international bunker fuel in the Monthly fuel, gas and inventory statistics have been found to be of high quality. As a consequence of that VAT is applied on national fuel consumption, but not on international bunkers, all respondents to the survey are able to separate these fuel amounts with high accuracy. Fuels used for domestic and international navigation have been separated correctly and in line with IPCC Guidelines.

³⁴ Hedlund & Lidén 2010

³⁵ Eklund et al. 2011. Emissions from navigation and fishing including international bunkers

3.2.2.2 MULTILATERAL OPERATIONS, CRF 1.C.2

Emission from multilateral operations are not included in the national total but instead reported separately as a memo item in CRF 1.C.2, in accordance with the 1996 revised IPCC Guidelines. These emissions are calculated based on information from the military on the amount of fuel purchased in Sweden but used abroad by Swedish forces participating in international operations.

3.2.3 Feedstocks and non-energy use of fuels

Activity data on feedstocks and non-energy use of fuels is collected from the quarterly fuel statistics. As also noted in Annex 2 section 1.1.1, in the survey form for the quarterly fuel statistics, respondents are among many other things asked to specify whether fuels are used as raw materials or for energy purposes. This facilitates the use of data for CRF table 1.A.d, non-energy use of fuels. As mentioned in section 3.2.1, data on natural gas used as feedstock cannot be reported for the years 2004-2008 due to confidentiality reasons (this activity started in 2004, and for the years 2009 and later, the company using natural gas as feedstock has given permission to publish this data. It is not possible to get a “retroactive” permission to publish data reported in the survey before 2009).

Data on carbon from coke, bound in produced ferroalloys is collected directly from the only ferroalloy producer and is added to the remaining data on carbon from coke. Estimates of carbon stored are derived by multiplying given energy amount with emission factors for CO₂ (as given in Annex 2, section 1.2) multiplied by 12/44 (the weight of one atom of carbon is by definition 12/44 the weight of one molecule of CO₂).

CO₂ emissions derived from non-energy use of fuels and reported under CRF 1.B and CRF 2 (e.g. flaring of gases and iron and steel process emissions) are added under CRF 1.A.d and linked to the CRF 1.A.b as carbon stored (see Annex 4). The parameter “fraction of carbon stored” has been set to 1.00 for all fuels. This is done because otherwise the emissions corresponding to CRF 2 and 1.B in the sectoral approach would not be accounted for in CRF 1.A.d but in 1.A.b, which would cause systematic differences in the comparison 1.A.c.

3.2.4 CO₂ capture from flue gases and subsequent CO₂ storage

So far, storage of CO₂ does not occur in Sweden³⁶. There are, however, several research projects going on where CO₂ is captured from flue gases at a pilot scale.³⁷

3.2.5 Country-specific issues

No country-specific issues are reported in this submission.

³⁶ Geological Survey of Sweden, 2010

³⁷ E-on 2010-11-04, Fortum 2010-11-04

3.2.6 Public electricity and heat production (CRF 1.A.1.a)

3.2.6.1 SOURCE CATEGORY DESCRIPTION

Swedish production of electricity is characterized by large proportions of hydropower and nuclear energy. Only a small share of electricity production is based on fuels used in conventional power plants. Public electricity and heat use vary between years, due to variations in ambient temperatures for instance. In addition, production of electricity based on fuels depends to a large extent on the actual weather conditions. Years with dry weather and cold winters have a significant effect on the use of fuel in electricity production since less electricity can be produced by means of hydropower and more electricity is needed for heating. The largest emissions from electricity production were thus in 1996, due to very dry and cold weather. The winters 2009/2010 and 2010/2011 were unusually cold, which lead to an increase in fuel consumption particularly in 2010. Liquid fuels and natural gas account for most of the increase, although the increase in natural gas use can to a large extent be explained by the fact that new gas fuelled facilities have been taken into operation. The use of solid fuels also increased substantially between 2009 and 2010, but in this case the explanation is the recovery from the dip in production in the iron and steel industry in 2009, which thus affected the amounts of energy gases sold to the public electricity and heat production plants.

In Sweden, electricity and district heating are used to a large extent to heat homes and commercial premises. Increased use of district heating since 1990 to heat homes and commercial/industrial premises has led to increased energy efficiency and thus lower emissions. Emissions of methane and nitrous oxide have increased from electricity and heat production because of the increased burning of biomass fuels.

Electricity is an important energy source in the manufacturing industry, where the most important industries are the pulp and paper and the steel industry.

Production of district heating is currently to a large extent based on biomass and waste. There has been a change from fossil fuels towards biomass since 1990. In 1990, 15 % of fuels used were biomass and 15 % was waste. In 2011, 55 % of all fuels used for district heating were biomass while waste accounted for 22 %. ³⁸ These proportions have been quite similar during the last six years. Since 1990, there has been a large increase in the use of district heating from 89 PJ (1990) to 173 PJ (2011)³⁹ but, due to the more frequent use of biomass, greenhouse gas emissions from district heating were only slightly larger than in 1990.

³⁸ All numbers are according to data used in the Greenhouse gas inventory this submission.

³⁹ Statistics Sweden EN11SM 1202 (Electricity supply, district heating and supply of natural and gasworks gas 2011. Preliminary data).

The number and distribution of Swedish power stations in 2010 are presented in Table 3-2⁴⁰. Changes since 1990 in number of plants and their installed effect have been minor in the electricity sector, but the number of plants that only produce district heating has increased.

Table 3-2. Number and distribution of Swedish energy stations, 2010.

Type of plant	Number of plants	Gross Production GWh	Gross Production TJ
Total power stations	2,721	148,255	533,718
Power generation not based on fuels	2,543	70,780	254,808
Windpower	1,663	3,502	12,607
Hydropower	880	67,278	254,808
Power generation based on fuels	178	77,474	278,906
Nuclear power	3	57,728	207,821
Conv. thermal power	175	19,746	71,086
- Manufacturing industries, ISIC 10-37	39		
- Energy plants, ISIC 40	108		
- Others	28		

A summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-3.

Table 3-3. Summary of source category description, CRF 1.A.1.a.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1A1a	CO ₂	X	X		T2	CS	Yes
	CH ₄		X		T2	CS	Yes
	N ₂ O	X	X		T2	CS	Yes

CS Country Specific. T2 Tier 2.

3.2.6.2 METHODOLOGICAL ISSUES

Plant specific activity data and country- and sector-specific emission factors are used, which is considered to be Tier 2 methodology.

The activity data source for emissions in CRF 1.A.1.a is the quarterly fuel statistics, further described in Annex 2. Emission factors, also further described in Annex 2, are generally country specific, but in a few cases plant specific emission factors are used. For energy gases purchased from the iron and steel works and combusted by public electricity and heat production

⁴⁰ Data for 2011 currently not available (i.e., these facts are not included in the preliminary data). Statistics Sweden EN11SM 1201 (Electricity supply, district heating and supply of natural and gasworks gas 2010).

plants, CO₂ emission estimates provided by the iron and steel works are used, which results in aggregate year specific implied emission factors for blast furnace gas, coke oven gas and steel converter gas.

The most important fuels in recent years are wooden fuels followed by domestic waste. Greenhouse gas emission factors for these fuels are national and were developed in a study made in 2004⁴¹.

Emissions from energy plants integrated with the iron and steel industry are allocated to CRF 1.A.2.a. This is discussed in chapter 3.2.9 and in detail in chapter 4.4.1.

3.2.6.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The quarterly fuel statistics is a total survey for ISIC (International Standard Industrial Classification of All Economic Activities) 40 and the response rate is almost 100 %. This provides the inventory with data of very good quality, accurate, complete, consistent and with very low uncertainties.

The trend in fuel consumption in this sector varies depending on the production of hydroelectric power and weather variations between years. The greatest changes in fuel consumption are for biomass fuels, where the consumption has increased significantly due to for instance increased district heating. It can also be noted that the use of natural gas in this sector was much higher in 2009-2011 than earlier years. The reason is that the number of gas-fuelled facilities has increased.

The variations in IEFs (implied emission factors) between years are normally small. The IEFs for solid fuels, however, are considerably more variable than for other fuel types due to the variable supply of energy gases from the iron and steel industry. As blast furnace gas has a much higher CO₂ EF than other solid fuels, the share of blast furnace gas has a very large influence on the aggregate CO₂ IEF for solid fuels. As the production in the iron and steel industry was much lower in 2009 than in other recent years, the share of blast furnace gas in CRF 1.A.1.a dropped, which explains the drop in CO₂ IEF for solid fuels in 1.A.1.a in 2009. The IEFs for the group other fuels also vary between years because the emission factors for municipal solid waste are different from the emission factors for other fuels in this group. In recent years, municipal waste accounts for 90-92% of the consumption of "other fuels". The remaining 8-10% are in most cases specified as "recycled fuel", but before 2007 there is no such information. As the composition of "recycled fuel" is unknown, there are no specific emission factors for this fuel, so the regular emission factors for "other non-specified fuels" are used. The CO₂ emission factor for this fuel is considerably higher than the emission factor for municipal waste, as the fossil fraction is assumed to be higher in unspecified fuels. The emission factors are discussed in Annex 2.

⁴¹ Boström et al, 2004

Emissions of NO_x and SO₂ and in relation to fuel consumption are also slightly variable between years due to variations in fuel mix. In the latest years, especially the SO₂ emissions in relation to fuel consumption have decreased due to a shift from residual fuel oils towards natural gas.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. Wooden fuels are the most common fuels in this sector, but as CO₂ from biomass is not included in the sectoral total of GHG emissions, CO₂ from combustion of peat, blast furnace gas and “other fuels” accounts for the largest contributions to the aggregate uncertainty of GHG emissions in CRF 1A1a. The activity data uncertainties are relatively low, 2 % for peat and blast furnace gas and 10 % for “other fuels”. The CO₂ emission factor uncertainties are 20 % and 100 %, respectively, and thus account for the greater part of the aggregate uncertainties. Activity data uncertainties are assigned by expert judgements made by persons at the energy statistics department of Statistics Sweden. Emission factor uncertainties have been assigned by national experts on emissions from stationary combustion.

3.2.6.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All quality procedures according to the Swedish QA/QC plan (including the Manual for SMED's Quality System in the Air Emission Inventories) have been implemented during the work with this submission.

All Tier 1 general inventory level QC procedures and all QC procedures listed in GPG section 8.1.7.4 applicable to this sector are used. The activity data has, of course, been subject to QA/QC procedures prior to the publishing of quarterly fuel statistics. In addition, the consumption of every type of fuel in the last year is checked and compared with previous years. If large variations are discovered for certain fuels, the consumption of these fuels is studied on facility level and if necessary, the staff responsible for the quarterly fuel survey is contacted for an explanation. IEFs are calculated per fuel, substance and CRF-code and checked against the emission factors to make sure that no calculation errors have occurred when emissions were computed.

The time series for all revised data have been studied carefully in search for outliers and to make sure that levels are reasonable. Data has, when possible, been compared with information from companies' legal environmental reports and/or other independent sources. Remarks in recent review reports from the UNFCCC have been carefully read and taken into account whenever time limits allow. The results are verified by calculating CO₂ emissions with the reference approach, and comparing results with the sectoral approach (see Annex 4).

During 2011, there was a study⁴² comparing the currently used quarterly fuel statistics with two other data sources, and the conclusion was that the quarterly fuel statistics is of very good quality, and also the only data source that is ready in time for use for the last emission year.

3.2.6.5 SOURCE-SPECIFIC RECALCULATIONS

No major recalculation was made in this sector in submission 2013, see Table 3-4. During the QC-procedure it was discovered that the year specific emission factors for CO₂ for natural gas 2007-2010 were not updated in submission 2012 and this minor error was corrected in submission 2013.

Table 3-4. Recalculation of CO₂ in CRF 1A1a, gaseous fuels, submission 2013 compared to submission 2012.

Year	2007	2008	2009	2010
Gg CO ₂	-0.3	-0.2	-2.4	-0.3

3.2.6.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.7 Petroleum refining (CRF 1.A.1.b)

3.2.7.1 SOURCE CATEGORY DESCRIPTION

Refineries process crude oil into a variety of hydrocarbon products such as gasoline and kerosene. During the refining process, dissolved gases are separated, some of which may be leaked or vented during processing and consequently reported under CRF 1.B.2. There are five refineries in Sweden. Three of these refineries produce fuel products such as gasoline, diesel and heating oils. The other two refineries mainly produce bitumen products and naphthenic special oils. One facility has a catalytic cracker; two facilities have hydrogen production plants and four of the facilities have sulphur recovery plants. The fuel consumption in this sector consists mainly of refinery gas, which is a by-product in the refining process. The use has increased since the 1990's due to higher demand of refined products. For the last four years, fuel consumption has been quite constant.

A summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-5.

⁴² Eklund & Kanlén 2011

Table 3-5. Summary of source category description, CRF 1A1b.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1A1b	CO ₂	X	X		T2	CS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. T2 Tier 2.

3.2.7.2 METHODOLOGICAL ISSUES

Activity data for the five refineries was collected directly from each company for 1990-1999, since the industrial energy statistics and quarterly fuel statistics did not account for all fuels produced within refineries during these years. The corresponding energy content of all fuels was also collected and individual thermal values were calculated for each operator and fuel. For 2000-2004, e.g. before the EU Emission Trading System (ETS) was established, energy statistics was used as the data quality was improved compared to the 1990's and is considered to be sufficient for these years. As a result of a specific SMED study during 2006⁴³, data from ETS are used for four refinery plants for 2005 and later years. For the fifth plant data from environmental reports were used. In 2008 and later years, the quality of ETS data is considered to be very high for all five of the refineries, and thus this is the primary data source for the GHG inventory. However, one of the refineries reports refinery gas and natural gas aggregated in the ETS data, and for this facility, data from the environmental reports are used to allocate the proper amount of this fuel to gaseous fuels. Environmental reports are used for verification for all five refineries. For refinery gas, plant specific CO₂ emission factors reported to the ETS are used for 2008 and later, since they are considered to be more accurate than the older standard emission factor.

During the national peer review remarks have been made that gaseous fuels are reported as "NO" for 2003 and questioned if this is the correct notation key. Investigations of activity data files used in earlier submissions show that in 2001 to 2003, sweet gas (a by-product from the cryogen plant) was probably miscoded as natural gas in submission 2005. Data for 2003 has been revised in later submissions, i.e. sweet gas has been re-coded as refinery gas. Environmental reports show that natural gas has been used in CRF 1.A.1.b in 2004 and later, but not in 2003, and hence "NO" is considered to be the correct notation key for 2003. The environmental reports for 2001-2002 are no longer available, and hence there is not enough information to recode the natural gas reported in 2001 and 2002, even though it might be miscoded refinery gas.

⁴³ Backman & Gustafsson, 2006

In addition to the five refineries, there are a few other plants with the Swedish Standard Industrial Classification 232, petroleum refining, which should be reported under refineries according to IPCC guidelines. For these plants, activity data from the quarterly fuel statistics are used together with national emission factors. It should be noted that the five refineries account for more than 99 % of the fuel consumption and emissions reported in CRF 1.A.1.b.

3.2.7.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The use of so many different sources for this sector could of course lead to consistency problems. Data used in the inventory is however analysed and no (significant) signs of inconsistency have been found. E.g. the slight dip in fuel consumption in 2007 is visible in all available data sources and is thus real and not caused by the shifting of data sources.

The implied emission factor for CO₂ for refinery gas is slightly lower for 2008-2011 when plant specific emission factors are used. However, as the national emission factor used for earlier years is based on information from the refineries, the decreasing IEF is considered to reflect actual technology improvements.

CO₂ from refinery gas is by far the largest source of uncertainty due to the fact that refinery gas accounts for about 90% of the energy from fuel combustion in this sector. The emission factor uncertainty is 5% and the activity data uncertainty is 10 %. The assigned uncertainties are based on information directly from the facilities. They are updated regularly but not annually.

3.2.7.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for CRF 1.A.1.b as for 1.A.1.a described above. For each of the five refineries, ETS data for the latest year are verified against the refineries' legal environmental reports.

3.2.7.5 SOURCE-SPECIFIC RECALCULATIONS

No recalculations other than the correction of CO₂ emission factors for natural gas mentioned in section 3.2.6.5 have been carried out. The difference compared to submission 2012 is less than 0.1 Gg CO₂ per year.

3.2.7.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.8 **Manufacture of solid fuels and other energy industries (CRF 1.A.1.c)**

3.2.8.1 SOURCE CATEGORY DESCRIPTION

Most emissions in this sector arise from two plants belonging to one company, producing coke to be used in blast furnaces for production of iron. The plants are integrated into the iron and steel production industry. Other fuel combustion in manufacturing of solid fuels and all fossil fuel combustion in manufac-

turing of nuclear fuels are also included in CRF 1A1c. The trend is related to the amounts of iron and steel produced, and hence there was a dip in 2009. Apart from this, emissions in recent years are quite similar to the emissions in the 1990's.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-6.

Table 3-6. Summary of source category description, CRF 1A1c.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1A1c	CO ₂				T2	CS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. T2 Tier 2.

3.2.8.2 METHODOLOGICAL ISSUES

The methodology for estimating emissions from the iron and steel industry was thoroughly revised in submission 2010. Activity data is taken from environmental reports and CO₂ emissions are estimated from carbon balances calculated by the facilities' own experts. Emissions of N₂O, CH₄, NMVOC and CO are estimated with the general T2 methodology with national emission factors. Estimates of emissions of SO₂ and NO_x are available from environmental reports on an aggregate level, and these emissions are distributed over the different CRF codes (1.A.1.c, 1.A.2.a, 1.B.1.c and 2.C.1, SO₂ also 2.B.5 and 1.B.1.b) according to the activity data distribution. The new methodology is described in detail in the methodological issues section about CRF 2.C.1.2.

Activity data for facilities manufacturing nuclear fuels is collected from industrial energy statistics for 1990 - 1996 and 2000 - 2002 and from quarterly fuel statistics for 1997 - 1999 and from 2003 onwards. For more details on the surveys see Annex 2.

3.2.8.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The time series is considered to be very consistent as all data on emissions from the coke producing plants has been collected directly from the facilities. The inter-annual variations in IEFs for solid fuels are caused by variations in the relative amounts of blast furnace gas and coke oven gas, respectively, between years. The composition of each gas is also quite variable, and this is another explanation to the fluctuating IEF's. Solid fuel consumption decreased considerably in 2009 due to lower production of coke caused by lower demand of primary iron and steel. In 2010, the demand increased and thus the fuel consumption increased to about the same level as before 2009. Consumption of liquid fuels has decreased since 2006 and the consumption of biomass is small and fairly constant. (Liquid fuels and biomass are, of course, not used in the coke ovens but in the other facilities that are allocated to 1.A.1.c due to their ISIC classification).

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. CO₂ from blast furnace gas and coke oven gas are the dominating sources of uncertainty.

3.2.8.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

The estimation of emissions from coke production is based on carbon balance calculations and the methodology is thoroughly described in chapter 4.

3.2.8.5 SOURCE-SPECIFIC RECALCULATIONS

A minor allocation error in submission 2012 was corrected which resulted in 0.003 Gg higher emissions of NO_x in 2010 compared to submission 2012.

3.2.8.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.9 Iron and steel (CRF 1.A.2.a)

3.2.9.1 SOURCE CATEGORY DESCRIPTION

The iron and steel industry is, together with the pulp and paper industry and the chemical industry, one of the most energy intensive industrial branches in Sweden. There are three primary steel works that base their production on iron ore pellets procuring either steel or iron powder. There are also 10 secondary steel plants producing steel based on scrap iron. In 2009, fuel consumption in the iron and steel industry fell sharply as a consequence of decreased production due to the global recession. In 2010, production and fuel consumption recovered to more “normal” levels. The Swedish iron and steel works produced 4.8 million tonnes of steel in total in 2010, which is 73% more than in 2009. In 2011, the production was 4.9 million tonnes.⁴⁴ The trend of the fuel combustion is increasing slightly since 1990 due to higher production of iron and steel products.

A summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3.7.

Table 3-7. Summary of source category description, CRF 1.A.2.a.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.2.a	CO ₂				T2,T3	CS, PS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. PS Plant Specific. T2 Tier 2. T3 Tier 3.

⁴⁴ The Swedish Steel Producers' Association, 2012-09-13

3.2.9.2 METHODOLOGICAL ISSUES IRON AND STEEL, CRF 1.A.2.A

During 2009, a new methodology was applied for the two largest primary iron and steel works. This is described briefly in section 3.2.8.2 above and in detail in section 3.2.9.2.1 below.

For companies with less than 10 employees the Tier 1 method is used, since current data does not allow the Tier 2 methods to be used. Emissions from companies with less than 10 employees are allocated to CRF 1.A.2.f because the model estimate of fuel consumption within small companies is only produced on an aggregate level and not separated by ISIC code.

Activity data for all facilities apart from the two largest ones mentioned above is, if not otherwise stated, collected from industrial energy statistics for 1990-1996 and 2000-2002, and from quarterly fuel statistics for 1997-1999 and 2003 onwards, further described in Annex 2.

Emissions reported from primary steel works and other iron and steel works are reported in both CRF 1A2a and in CRF 2.C.1 since some emission arises from fuel combustion and some from reducing agents in the process. The text in this section is hence closely connected to the text in the section CRF 2.C.1.1 (secondary steel) and CRF 2.C.1.2 (primary pig iron and steel).

3.2.9.2.1 *Primary iron and steel works*

In Sweden, there are two plants for integrated primary iron and steel production, i.e. basing their production on iron ore pellets. The integrated iron and steel production consists of material flows between coke oven, blast furnace and steelworks, and in one plant, rolling mill (see Table 3-8). Emissions from fuel combustion (oils, LPG (Liquefied Petroleum Gas) and recovered energy gases, i.e. coke oven gas and blast furnace gas) used in the rolling mills and for in-house power and heat production is allocated to this sub-sector in accordance with the IPCC Guidelines. The allocation approach used in submission 2013 has been used since submission 2010. From one of the facilities, large amounts of recovered energy gases are sold to a public heat and power plant, and the emissions from combustion of these gases are hence reported in CRF 1.A.1.a.

Table 3-8. Allocation of fuel consumption and CO₂ emissions in 2011 from iron ore based iron and steel industry on different CRF codes.

CRF	Plant station	Fuel consumption 2011 (TJ)	CO ₂ emissions 2011 (Gg)
1A1c	Coke Oven	4,489	345
1A2a	Combustion in Rolling Mills + Power and Heat Production	3,603	726
1B1c	Flare in Coke Oven (COG)	128	6
2C1.2	Blast Furnace + Steelworks (including Flaring of BFG and LD-gas)	NA	2,436
Total			3,512

3.2.9.2.2 *Secondary iron and steel works*

Except for the primary iron ore based iron and steel works, this sector includes emissions from for instance electric arc furnaces plants, iron ore pellet plants and iron powder plants.

3.2.9.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

For the two largest facilities, the time series is very consistent as all data is based on information from these facilities, and they have checked that the calculated emission and energy data is accurate. In 2012 it was concluded that the emissions of CO₂ from coke used in blast furnaces at one of the facilities is overestimated for the years 2005 and later, see section 4.4.1.3. However, the overestimation seems to affect only CRF 2.C.1.2 and not CRF 1.A.1c or 1.A.2.a. This issue will be investigated before next submission. For CRF 1.A.2.a in total, the time series is also considered to be consistent, despite the fact that the quarterly fuel survey is used for most years and the yearly industrial energy survey for some years. The quarterly fuel survey data is weighted to cover the same population as the yearly industrial energy survey. A discussion on the reasons for changing data sources can be found in Annex 2.

The CO₂ implied emission factors for solid fuels in CRF 1.A.2.a are higher than for solid fuels in other industries, since a large proportion of the fuel used is blast furnace gas which has a high carbon content compared to other solid fuels. This also implies that the IEF varies between years, and it is considerably lower in 2009 than recent years because of the drop in blast furnace gas consumption. This explains the fact aggregate CO₂ IEF for CRF 1.A.2.a is considerably lower in 2009 than in earlier years. See also section 0.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. CO₂ from blast furnace gas, coke oven gas and LPG are the largest sources of uncertainty in this sector. For these fuels the activity data uncertainty is 5 %. The CO₂ emission factor uncertainty is 20% for blast furnace gas and coke oven gas, and 5% for LPG.

3.2.9.4 SOURCE SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for CRF 1.A.2.a as for 1.A.1.a described above. In addition to this, fuel consumption for the year t-2 is verified against the annual industrial energy survey on an aggregate level to check that the weight factors for the year t-1 are reasonable. For the two largest facilities, all data is collected directly from the company.

3.2.9.5 SOURCE SPECIFIC RECALCULATIONS

Minor errors have been corrected; the revision of emission factors for natural gas described in section 3.2.6.5. and the reallocation of NO_x emissions 2010 mentioned in section 3.2.8.5

3.2.9.6 SOURCE SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.10 Non-Ferrous Metals (CRF 1.A.2.b)

3.2.10.1 SOURCE CATEGORY DESCRIPTION

This source category covers combustion-related emissions from seven aluminium producers (ISIC 27420), six copper producers (ISIC 27440) and five facilities producing various other metals. More detailed descriptions are given in section 4.

Fuel consumption shows a decreasing trend for the period 1990-2002, but from 2003 onwards, the inter-annual variations in fuel consumption for energy production are relatively small and the copper- and aluminium producers account for about 40 % each. The most common fuel is LPG followed by natural gas. Smaller amounts of heating oils are also used.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-9.

Table 3-9. Summary of source category description, CRF 1.A.2.b.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.2.b	CO ₂				T2	CS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. T2 Tier 2.

3.2.10.2 METHODOLOGICAL ISSUES

Activity data is taken from industrial energy statistics for 1990-1996 and 2000-2002, and from quarterly fuel statistics for 1997-1999 and 2003 and later. For more details on these surveys see Annex 2.

As for all subcategories to CRF 1.A.2, for companies with less than 10 employees the Tier 1 method is used, since current data does not allow the Tier 2 methods to be used. Emissions from companies with less than 10 employees are allocated to CRF 1.A.2.f.

3.2.10.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As for CRF 1.A.2.a, time series consistency despite the changes in activity data source is discussed in Annex 2. In 1999 there is a large jump in the time series due to increased consumed amounts of natural gas. This has been identified as a possible reporting error for one facility, but original raw data from 1999 is no longer available and hence revision is not possible.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. In 1990, the largest contribution to the aggregate uncertainty arises from CO₂ from “other solid fossil fuels” due to the fact that the emission factor uncertainty for this quite unspecified fuel is as high as 100%. In later years, this fuel is not used in CRF 1.A.2.b, and CO₂ from LPG accounts for most of the uncertainty. The uncertainty is 5%, both in activity data and in the CO₂ emission factor for this fuel. Activity data uncertainties are assigned by expert judgements by staff at the energy statistics department of Statistics Sweden. Emission factor uncertainties have been assigned by national experts on emissions from stationary combustion.

3.2.10.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

The same QA/QC procedures are used for CRF 1.A.2.b as for 1.A.2.a described above. In addition to this, a detailed quality study of the non-ferrous metal industry was performed in 2010.⁴⁵ This study showed no reasons for revisions in CRF 1.A.2.b.

3.2.10.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

3.2.10.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.11 Chemicals (CRF 1.A.2.c)

3.2.11.1 SOURCE CATEGORY DESCRIPTION

The chemical industry produces a number of different products such as chemicals, plastics, solvents, petrochemical products etc. In total, around 50 plants are included, of which ten uses more than 90 % of the energy according to the activity data used for emission calculations for this sector. The fuel consumption trend is increasing since 1990, especially for liquid fuels, mainly due to increased use within the basic plastic industry. Throughout the time series, liquid fuels account for about 80% of the energy and gaseous fuels for 10-15%.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-10.

⁴⁵ Danielsson & Nyström, 2010

Table 3-10. Summary of source category description, CRF 1.A.2.c.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.2.c	CO ₂	X	X		T2	CS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. T2 Tier 2.

3.2.11.2 METHODOLOGICAL ISSUES

Activity data is, with exceptions mentioned below, collected from industrial energy statistics for 1990-1996 and 2000-2002, and from quarterly fuel statistics for 1997-1999 and 2003 and later. For more details on these surveys see Annex 2.

Generally, plants classified as ISIC Division 24 according to ISIC Rev.3⁴⁶ in the energy statistics are included in this sector, as recommended in IPCC 1996 Revised Guidelines.

For one of the largest facilities, including two plants, ETS data is the activity data source for 2008 and later. Before 2008, this facility was not fully covered by energy statistics or ETS data, so environmental reports and several energy surveys were used in order to get complete data for this important facility.

One calcium carbide manufacturing facility uses coke both as a fuel and as a reductant in the production process. In submission 2013, it was revealed that the reporting of this coke consumption is not properly allocated in the energy statistics, and several years the total amounts reported were obviously too low. For this reason, activity data from environmental reports and in later years ETS are used for this coke consumption in submission 2013.

As in other subcategories of CRF 1A2, for companies with less than 10 employees the Tier 1 method is used, since current data does not allow the Tier 2 methods to be used. Emissions from companies with less than 10 employees are allocated to CRF 1.A.2.f.

According to environmental reports, the “other petroleum fuels” used in this sector is a process by-product consisting mainly of methane. The fuel is produced at one facility and used by several chemical industries in the same municipality. ERT has remarked that this fuel is probably partly originating from natural gas, which is also indicated by the environmental reports. It has, however, not been possible to determine how much of the gas mixture that should be allocated to gaseous fuels, so in submission 2013, all consumption of this fuel is allocated to liquid fuels. This is the same allocation that has been used in previous submissions. Both natural gas and petroleum products are used as feedstock, and hence the by products as well as the actual desired products are partly of liquid origin and partly of gaseous origin. The major part of the raw material is, however, of liquid origin. This assumption is sup-

⁴⁶ United Nations Statistics Division, 2010

ported by the comparison between the reference and sectoral approach for gaseous fuels, which indicates that the amount of suspectedly misallocated fuel is very small.

In 2011, a consistent time series of the CO₂ emission factor for this fuel was developed in cooperation with the facility that produces the fuel and hence it is plant specific. The emission factor used in submission 2011, namely 55 kg CO₂/GJ, was verified by the company for the period 1990-2000. In 1999 to 2001, the process that produces the gas was gradually modified by technological improvements, resulting in an altered composition of the fuel. The proportion of hydrogen increased, which gave a higher calorific value and lower CO₂ emissions. The company also provided a time series of CO₂ emissions covering the period 2001-2010, which was used to calculate the year specific emission factors. These new emission factors were implemented in submission 2012, and thus the inconsistent time series used in submission 2011, where the “old” emission factor was used 1990-2007 and the considerably lower emission factors reported to ETS were used for 2008-2009, has now been corrected. For non-CO₂ emissions, emission factors for natural gas are used as no specific emission factors are available and both fuels consist mainly of methane.

3.2.11.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As for CRF 1A2a and 1A2b, time series consistency despite the changes in activity data source is discussed in Annex 2.

As mentioned above, fuel consumption in 2011 was higher than in 1990. However, since 2003 there is no distinct trend and the fuel consumption in 2009 was about 10 % lower than in 2008, which reflects the decreasing demand for this sector in 2009. In 2010-2011, fuel consumption increased to roughly the same level as in 2006-2007.

As noted by the ERT, the implied emission factors for “other fuels” are variable, especially in the early years. This is explained by the fact that municipal waste has occasionally been combusted within the chemical industry, and most years also “other non-specified fuels”. As these fuels have very different emission factors for CO₂, the relative amounts of these two fuels cause inter-annual variations in IEFs. The outlier value of 28.4 kg/GJ in 1992 is explained by the fact that a small amount of municipal waste was combusted that year, but no “other non-specified fuels”. It should be noted that the group “other fuels” accounts for a relatively low share of the emissions compared to other fuel groups; typically around 5% of the emissions of fossil CO₂ within CRF 1.A.2.c.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. CO₂ from methane-based gas mixtures accounts for most of the uncertainty. The uncertainty in activity data is 5 % and the emission factor uncertainty is assumed to be 10 % based on the variation in plant specific values. Activity data uncertainties are assigned by expert judgements by staff at the energy statistics department of Statistics Sweden.

3.2.11.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for CRF 1A2c as for 1.A.2.a–b described above. For the largest plants in terms of emissions and fuel consumption, both environmental reports and ETS data are used for verification of the estimates based on energy statistics.

In the development project in 2010⁴⁷ mentioned above, the activity data time series 1990–2008 for all fuel types and all facilities within the chemical industry were thoroughly reviewed. Reported emissions and activity data in CRF 1 and 2 were analysed on plant level and verified against environmental reports and when necessary, the plants were contacted for explanations or complementary data.

3.2.11.5 SOURCE-SPECIFIC RECALCULATIONS

As mentioned above, activity data for coke combustion has been revised for the whole time series. The differences in emissions compared to submission 2012 expressed as CO₂-equivalents are shown in Table 3-11.

Table 3-11 Difference in GHG-emissions submission 2013-2012, CO₂ equivalents, CRF 1.A.2c

Year	1990	1995	2000	2005	2006	2007	2008	2009	2010
Gg CO ₂	22.1	18.2	14.3	16.9	4.8	9.2	7.9	2.3	13.8
Percent of 1A2c	1.9%	1.4%	1.1%	1.2%	0.3%	0.7%	0.7%	0.2%	1.1%

Apart from this, only minor revisions of the CO₂ emission factors for natural gas 2007–2010 have been made as described in section 3.2.6.5.

3.2.11.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.12 Pulp, Paper and Print (CRF 1.A.2.d)

3.2.12.1 SOURCE CATEGORY DESCRIPTION

In 2011 there were 39 paper mill plants, 140 sawmills (production capacity >10,000 m³/year) and 41 pulp industry plants in Sweden. In total, they were producing 11.3 million tonnes of paper, 16.8 million m³ of sawn timber and 11.9 million tonnes of pulp.⁴⁸ Since 1990, production has had an increasing trend, but not in the latest few years. There is no apparent trend in total fuel consumption since 1990, but in recent years, the share of energy from biomass fuels has increased, from 68 % of fuel consumption in 2007 to 78 % in 2011.

⁴⁷ Gustafsson, Nyström & Gerner, 2010

⁴⁸ The Swedish Forest Industries Federation, 2012-09-13 http://skogsindustrierna.org/branschen/statistik_7/statistik_om_skogsindustri

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-12.

Table 3-12. Summary of source category description, CRF 1.A.2.d

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.2.d	CO ₂		X		T2	CS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. T2 Tier 2.

3.2.12.2 METHODOLOGICAL ISSUES

Emissions from processes in the Pulp, paper and print industry are reported under CRF 2D1 according to IPCC Guidelines. See chapter 4.5.

Activity data is collected from industrial energy statistics for 1990-1996 and 2000-2002, and from quarterly fuel statistics for 1997-1999 and 2003 and later. For more details on these surveys see Annex 2.

As for CRF 1.A.2 in general, for companies with less than 10 employees the Tier 1 method is used, since current data does not allow the Tier 2 methods to be used. Emissions from companies with less than 10 employees are allocated to CRF 1A2f.

Emissions from combustion of sulphur lyes are presently not reported in CRF 1A2d as this activity has been considered as an industrial process, despite the fact that the process heat is used for heat and electricity production. The emissions of CH₄, N₂O and indirect greenhouse gases from the processes in which the sulphur lyes are consumed, are reported in CRF 2.

3.2.12.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As for CRF 1A2 in general, time series consistency despite the changes in activity data source is discussed in Annex 2. The fluctuating IEFs for liquid fuels reflect variations in fuel mix. In the 1990s, petroleum coke was used in some facilities, and in the latest years, combustion of heavy heating oils has decreased a bit. Fuels classified as “other fuels” are scarcely occurring in this CRF category, and as in 1A2c, the large variations in IEFs are caused by occasional use of municipal waste.

In recent years, the relative amount of biomass has increased and the relative amounts of liquid fuels, especially residential fuel oil, have decreased. One effect of the increasing share of biomass is that emissions of fossil CO₂ per TJ of total fuel consumption is decreasing.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. N₂O from wooden fuels and CO₂ from residual fuel oil are the greatest contributors to the aggregate uncertainty in this sector. The activity data uncertainty is 2 % for all years for both of these fuels. The N₂O emission factor uncertainty for wood

is 40 % and the CO₂ emission factor for residual fuel oil is 1 %. Activity data uncertainties are assigned by expert judgements made by persons in the energy statistics department at Statistics Sweden. Emission factor uncertainties have been assigned by national experts on emissions from stationary combustion.

3.2.12.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for CRF 1A2d as for 1A1a and 1A2a–c described above.

3.2.12.5 SOURCE-SPECIFIC RECALCULATIONS

Only minor corrections of emission factors for CO₂ from natural gas 2007-2010 have been made as described in section 3.2.6.5.

3.2.12.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.13 Food Processing, Beverages and Tobacco (CRF 1.A.2.e)

3.2.13.1 SOURCE CATEGORY DESCRIPTION

The food and drink industry is the fourth largest branch of industry measured as production value and number of employees. There are about 3,200 companies, of which only around 650 have more than 10 employees.⁴⁹ The largest number of companies and employees are found in the bakery industry, but the most energy intensive branch is the sugar industry which accounts for about 25 % of the fuel consumption in 1A2e. Dairies, breweries, producers of refined vegetable fats and potato products are other industries with significant fuel consumption (around 7-12 % each of the fuel consumption in 1.A.2.e). The fuel consumption varies between years. A slight decrease can be observed since 1990. In later years, gaseous fuels account for 45-49 % and liquid fuels account for about 38-40 % of the total fuel consumption.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-13.

Table 3-13. Summary of source category description, CRF 1.A.2.e

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.2.e	CO ₂				T2	CS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. T2 Tier 2.

⁴⁹ The Swedish Food Federation 2009-09-16

3.2.13.2 METHODOLOGICAL ISSUES

Activity data is collected from industrial energy statistics for 1990-1996 and 2000-2002, and from quarterly fuel statistics for 1997-1999 and 2003 and later. For more details on these surveys see Annex 2. For companies with less than 10 employees the Tier 1 method is used, since current data does not allow the Tier 2 methods to be used. Emissions from companies with less than 10 employees are allocated to CRF 1.A.2.f.

3.2.13.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As for CRF 1.A.2 in general, time series consistency despite the changes in activity data source is discussed in Annex 2. The IEFs are slightly variable between years due to variations in fuel mix. The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. In the early 1990's, CO₂ from residual fuel oil was the largest source of uncertainty, followed by CO₂ from natural gas. In recent years, CO₂ from natural gas accounts for most of the uncertainty. For both fuels, the activity data uncertainty is 5 %. CO₂ emission factor uncertainty is 1 % and 5% for residual fuel oil and natural gas, respectively. Activity data uncertainties are assigned by expert judgements made by persons in the energy statistics department in Statistics Sweden. Emission factor uncertainties have been assigned by national experts on emissions from stationary combustion.

3.2.13.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Generally, the same QA/QC procedures are applied for 1.A.2.e as for other 1.A.2 categories described above.

3.2.13.5 SOURCE-SPECIFIC RECALCULATIONS

Only minor corrections of data have been made, such as the correction of the CO₂ emission factors for natural gas 2007-2010 described in earlier sections.

3.2.13.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.14 Other Industries (CRF 1.A.2.f)

3.2.14.1 SOURCE CATEGORY DESCRIPTION

This source category is by nature quite heterogeneous. Both stationary and mobile emission sources are included. The stationary sources included are combustion within ISIC 10-37 except from the branches separately reported in 1.A.2.a-1.A.2.e, and stationary combustion within all companies with less than 10 employees regardless of branch, and stationary combustion within the construction sector. The mobile emission sources included in this sector are off-road vehicles and working machinery used in the construction and manufacturing industry.

In terms of stationary fuel combustion and emissions, three branches of industry are dominating; non-metallic mineral production (ISIC 26), manufacturing of wood products (ISIC 20), and mining industry (ISIC 13). In ISIC 20, however, biomass fuels are dominating and hence the emissions of fossil CO₂ from this branch of industry are low. The construction industry also accounts for a significant share of fuel consumption and emissions. The fuel consumption varies between years, but for stationary combustion within 1.A.2.f in total, it has decreased slightly since 1990. Liquid and biomass fuels account for most of the decrease. For mobile combustion, i.e. off road vehicles and working machinery, fuel consumption in 2010 was about 22 % higher than in 1990 but the trend has been slightly variable.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-14.

Table 3-14. Summary of source category description, CRF 1.A.2.f

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.2.f	CO ₂	X			T1,T2	CS	Yes
	CH ₄				T1,T2	CS	Yes
	N ₂ O	X			T1,T2	CS	Yes

CS Country Specific. T1 Tier 1. T2 Tier 2.

3.2.14.2 METHODOLOGICAL ISSUES

For emissions from stationary combustion, the Tier 2 method is used with the following exception: For the construction industry and for companies with less than 10 employees the Tier 1 method is used, since current data does not allow the Tier 2 method to be used. Stationary fuel combustion in the construction sector is shown below (Table 3-15).

Table 3-15 Stationary fuel combustion in the construction sector (part of 1A2f), TJ.

Year	LPG	Domestic heating oil	Residual fuel oil	Natural gas
1990	46	5,051	420	39
1996	46	4,692	382	79
2000	46	4,621	382	40
2005	46	1,254	265	501
2006	46	1,329	272	507
2007	46	1,391	269	507
2008	92	1,440	263	498
2009	92	1,469	247	466
2010	92	1,724	278	525
2011	85	1,269	342	431

Emissions from stationary combustion in mining and quarrying and in the manufacturing of various products such as textiles, wearing apparel, leather, wood and wood products, rubber and plastics products, other non-metallic mineral products, fabricated metal products and manufacturing of different types of machinery, are calculated with activity data from the industrial energy statistics for 1990-1996 and 2000-2002, and from the quarterly fuel statistics for 1997-1999 and 2003 and later. For more details on these surveys see Annex 2.

Emissions from all companies with less than 10 employees are estimated and reported under CRF 1.A.2.f. Activity data are collected from Statistics Sweden⁵⁰. Emissions are minor and with current data not possible to separate on different industry sectors.

Emissions from stationary combustion in the construction industry are calculated with activity data from Statistics Sweden.⁵¹ The methodology used for this sub-category is the same as for stationary combustion in the Other sector, see section 3.2.20.4.1. Activity data is basically from the annual energy balances, except for the latest emission year. However, the data in Table 3-15 differ slightly from the official energy balances due to use of slightly different calorific values especially for earlier years.

Since 2002, for one glassworks plant, it is no longer possible to separate combustion emissions of SO₂ from process emissions. The reason is that the facility has restructured its environmental report, and only reports emissions of SO₂ on an aggregate level. The median value for the share of process related SO₂ emissions to the total SO₂ emissions is 2 % for the years 1990 - 2001. Emission data reported in the plants environmental report are considered to be more accurate than emissions calculated from fuel combustion with standard emission factors. Thus for practical reasons, SO₂ and NO_x emission data available from environmental reports are reported in CRF 2A7. All other energy related emissions for this facility are reported in CRF 1A2f. For 2008 and later, activity data for the three plants within the cement production industry is taken from the EU ETS system.

Emissions from mobile combustion refer to off-road vehicles and other machinery including various mobile vehicles and machines as for example tractors, dumpers, cranes, excavators, generators, wheel loaders, sorting works, pump unit and any other mobile machine in industry that run on petroleum fuels. The model used to estimate emissions from off-road vehicles and other machineries is considered to correspond to Tier 2. The model is further explained in Annex 2.

⁵⁰ Statistics Sweden, EN20SM 1990-2011. See also Annex 2.

⁵¹ Statistics Sweden, EN20SM 1990-2011. See also Annex 2.

3.2.14.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

As for CRF 1A2 in general, time series consistency despite the changes in activity data source is discussed in Annex 2. As for other categories in CRF 1.A.2, the IEFs are vary slightly between years due to variations in fuel mix. In earlier submissions, the EC (European Commission) has asked for clarification of the drop in wood consumption in 2000 compared to earlier years. This issue has not been prioritized, but since the annual wood consumption 2001-2009 is considerably lower than in the 1990s, there is no reason to believe that the activity data for 2000 is incorrect.

CO₂ from diesel and heating oils are the largest sources of uncertainty in GHG emissions within CRF 1.A.2.f. The activity data uncertainty for all heating oils within this sector is as high as 20 % on an aggregate level, due to the fact that emissions from the construction sector and small industries are estimated with the Tier 1 method. The activity data uncertainty for diesel combusted in off-road vehicles and working machinery is 5 % and for gasoline 3 %.

3.2.14.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Generally, the same QA/QC procedures are applied for 1A2f as for other 1.A.2 categories described above. In some earlier submissions, extensive QA/QC and verification efforts have been made for the other sectors including the construction industry. This is described in section 3.2.20.4.1 below.

3.2.14.5 SOURCE-SPECIFIC RECALCULATIONS

Activity data for the construction sector and enterprises with less than 10 employees in all industrial branches have been revised for 2009-2010 following revisions of the annual energy balances. Compared to submission 2012, stationary fuel consumption in 1.A.2.f is 0.03 % higher in 2009 and 0.42 % higher in 2010 (i.e. very minor revisions). The difference in emissions is 0.6 and 19.2 Gg CO₂ equivalents respectively.

The methodology for estimating emissions from off-road vehicles and working machinery was revised in submission 2012. The revision did not imply an updated methodology, but aimed to simplify the use of the model and at the same time update some emission factors, activity data and the allocation of emissions to different sectors. Allocation of emissions from off-road vehicles and working machinery is based on a report by Flodström (et al)⁵². This is the most recent inventory including an allocation of working machinery to sectors in Sweden.

⁵² Flodström et al 2004. Uppdatering av utsläpp till luft från arbetsfordon och arbetsredskap för Sveriges internationella rapportering.

3.2.14.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.15 Civil Aviation (CRF 1.A.3.a)

3.2.15.1 SOURCE CATEGORY DESCRIPTION

Presently data are provided for a total of 40 airports with regular and/or chartered air traffic. The national government administers 13 of these airports, while the remaining 27 are private and/or administered by local government.⁵³ The traffic routed through governmental airports account for about 90 % of the total fuel consumption within the civil aviation sector.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-16.

Table 3-16. Summary of source category description, CRF 1.A.3.a.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.3.a	CO ₂				T1	CS	Yes
	CH ₄				T2, M	D, M	Yes
	N ₂ O				T2, M	D, M	Yes

CS Country Specific. T1 Tier 1. T2 Tier 2. M Model. D Default.

3.2.15.2 METHODOLOGICAL ISSUES

Sweden uses Tier 1 to estimate emissions of CO₂ and SO₂, Tier 2 to estimate emissions of CH₄, NMVOC and N₂O and Tier 3a for all other gases.

Emissions from aviation in agricultural and forestry sectors are currently reported together with domestic aviation. Emissions from military use of aviation fuels are reported under Other – mobile sources (CRF 1.A.5.b).

The emissions from aviation reported to the UNFCCC are estimated using both data on supply and delivery of petroleum products from Statistics Sweden (see Annex 2) together with fuel- and emission data reported by the Swedish Transport Agency (STrA).

The Swedish Transport Agency (STrA) includes the traffic from a number of non-governmental airports in their estimates from 2005 and from all Swedish airports from 2006. Since 2010 there is no separate reporting on emissions from governmental respectively private airports, instead a total is reported. STrA publishes information on aviation emissions for 2011, from all Swedish airports with regular and chartered flights, in their annual environmental report.

⁵³ Swedish Transport Agency.

The fuel consumption and emissions published by STrA are calculated by the Swedish Defence Research Agency (FOI) by using an estimation model. STrA provide FOI statistics on the number of flights between cities, type of aircraft, amount of fuel needed for different flights, emissions per fuel on specific flights based on data on aircraft performance during different phases of the flight and the distance between destinations. This data is used in the model to calculate emissions and amounts of burnt fuel for whole flights as well as for aircraft movements below 3,000 ft at the airports, the so called LTO cycle.

Due to the fact that the Swedish airports generally are smaller than international airports in other countries, taxi times are much shorter for domestic flights and climb-out and take-off times are often shorter as well. Hence traffic from Swedish airports needs less fuel and give rise to lower emissions compared to the International Civil Aviation Organization (ICAO) standards that the IPCC guidelines follow.⁵⁴ For international flights, ICAO standard taxi time has been used for the part of the LTO cycle occurring on international airports.⁵⁵

The results from the emission calculations are aggregated into four groups: domestic landing and take-off (LTO), domestic cruise, international LTO and international cruise. This is in line with the IPCC guidelines and data of good quality exists from 1995 and onwards.

The methodology for calculating national emissions is the same for all years with a few exceptions for earlier years. Emissions of CO₂ are based on fuel delivery statistics and thermal values from 2006 IPCC Guidelines and emission factors from the Swedish EPA. Quotas for distributing the CO₂ emissions to domestic and international LTO and cruise are based on estimations of energy consumption from the Swedish Transport Agency.

All non-CO₂ emissions for 1990-1994 were calculated by SMED in cooperation with the STrA due to the lack of activity data. Emissions of CO for 1990-1994 were estimated by using the ratio between CO and CO₂ in 1995 (4.85 % of CO₂ emissions). Emissions of NO_x are calculated in a similar way. The mean value of the ratio between NO_x and CO₂ emissions in 1995-2004 is used for 1990-1994 (4.03 % of CO₂ emissions). Emissions of HC for 1990-1994 are calculated by extrapolation.

From 1995 and onwards, emissions of SO₂, NO_x, CO and HC are estimated by the Swedish Defense Research Agency (FOI) on behalf of STrA and adjusted to match the delivered amount of aviation fuels. Emissions of NMVOC and CH₄ are estimated based on the emissions of HC and emission factors from the IPCC guidelines.

N₂O emissions from LTO are estimated using information from STrA on the number of LTO cycles together with emission factors from IPCC. N₂O emissions from cruise are based on delivered amounts of fuel for cruise activities estimated by FOI together with emission factors according to the IPCC guidelines. Emissions of N₂O are adjusted to be in line with fuel delivery statistics.

⁵⁴ Gustafsson, 2005.

⁵⁵ Näs, 2005.

In 2006, the STrA responded to the governmental call to reduce response burden on statistical compilations. As a result, private aviation as well as educational training flights are no longer covered in the STrA reports on fuel consumption and emissions from aviation as from 2007. However, as the estimated emissions from aviation are adjusted to match the delivered amount of aviation fuels, the emissions from private aviation as well as from educational training flights will consequently be included.

The emissions of NMVOC have decreased noticeably in the last years as a result of a specific type of airplane (MD-80/82), which is a major contributor to these emissions, is phased out.

3.2.15.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

In order to maintain consistency with the time-series the estimation procedures have been developed as described above. However, due to the fact that some of the estimations are not based on activity data but on other factors as LTO cycles, a certain degree of uncertainty exists.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

Time series are checked for consistency and recalculations are verified.

3.2.15.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All Tier 1 general inventory level QC procedures and all QC procedures listed in GPG section 8.1.7.4 applicable to this sector are used. The activity data has, of course, been subject to QA/QC procedures prior to the publishing. In addition, the consumption of every type of fuel in the last year is checked and compared with previous years. If large variations are discovered for certain fuels, responsible staff is contacted for an explanation. IEFs are calculated per fuel, substance and CRF-code and checked against the emission factors to make sure that no calculation errors have occurred when emissions were computed.

3.2.15.5 SOURCE-SPECIFIC RECALCULATIONS

The ratio of aviation gasoline for national aviation was underestimated in submission 2012 and the ratio for aviation kerosene was as a result overestimated. This has been adjusted in submission 2013. The actual change in emissions is minor for both aviation gasoline and aviation kerosene.

The database used by FOI, in their calculations of fuel consumption and emissions from aviation, has been updated for the years 2008-2011. The most prominent difference is a better coherence between the engines used in the model calculations and the engines employed by aircrafts in reality. The estimation of distances in the database is also updated.

The improvements of the database have foremost resulted in decreased emissions of NO_x but increased emissions of CO. FOI judged that the changes to the database have a negligible impact on the results before 2008.

3.2.15.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.16 Road transport (CRF 1.A.3.b)

3.2.16.1 SOURCE CATEGORY DESCRIPTION

Road transport is the single largest source category contributing to the total national greenhouse gas emissions (excluding LULUCF) in Sweden, all years. It includes several vehicle categories: Passenger cars, Light duty vehicles, Heavy duty vehicles and Mopeds & Motorcycles. Gasoline has previously been the most common fuel used for road transports, but as from 2011 the amount of diesel used is on the same level as gasoline and the emissions of CO₂ from diesel has for the first time ever surpassed the emissions of CO₂ from gasoline.

Ethanol used for road transport has increased by nearly fifty per cent since 2006. The main part of the ethanol used for road transport is used as a blending component for gasoline, but the part of the ethanol used as a pure biofuel increases steadily.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-17.

Table 3-17. Summary of source category description, CRF 1.A.3.b.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.3.b	CO ₂	X	X		T2	CS	Yes
	CH ₄		X		T2, M	CS, M	Yes
	N ₂ O				T2, M	CS, M	Yes

CS Country Specific. T2 Tier 2. M Model.

3.2.16.2 METHODOLOGICAL ISSUES

Emissions of CO₂ and SO₂ from road traffic are based on statistics on supply and delivery of petroleum products (see Annex 2) and national emission factors in accordance with the IPCC Guidelines Tier 2. Emissions of CO₂ from combustion of gasoline and diesel are based on thermal values from SPBI⁵⁶ and country-specific emission factors from Statistics Sweden and the Swedish EPA as shown in Annex 2.

Prior to submission 2007, emissions of SO₂ from diesel and gasoline were based on the maximum allowed sulphur content of different environmental classes. Data on maximum allowed sulphur content was provided by SPBI. As from submission 2007, emissions of SO₂ are based on the actual sulphur content for the different environmental classes of petrol and diesel fuel. The data

⁵⁶ Swedish petroleum institute. www.spbi.se

on actual sulphur content, provided by the Swedish Transport Administration (STA), is based on estimates made by VTI⁵⁷ for the years 1990-2001, and on fuel analysis from SPBI from 2001 and onwards.

The amount of gasoline and the emissions of CO₂ and SO₂ from the use of gasoline by "Bifuel E85/petrol" vehicles are reported under "Passenger cars" and the fuel type "gasoline". The rest of the emissions generated by "Bifuel E85/petrol" vehicles are estimated under respectively vehicle category.

CO₂ and SO₂ from natural gas and biofuels are estimated using statistics on deliveries for natural gas, biogas, ethanol and FAME (Fatty Acid Methyl Ester). The thermal value and emission factor for CO₂ from ethanol was changed as from submission 2012 for all years⁵⁸. This resulted in a decrease of the CO₂ emissions by 1.48 % every year. The thermal value and emission factor for FAME, as well as for natural gas, have been provided by the SEPA. Emissions of CO₂ from biogas, ethanol (including ethanol admixture) and FAME are reported as biomass and not included in the national totals.

Emissions of all other substances are estimated by the Swedish Transport Administration (STA) by using a road traffic emission model HBEFA 3.1, except for particles from motorbikes which are estimated by EMEP/EEA⁵⁹ methodology. As from submission 2012 the road traffic emission model HBEFA 3.1 is used by STA to estimate emission from road traffic instead of ARTEMIS⁶⁰.

HBEFA 3.1 is updated yearly with new information regarding emission factors, vehicle fleet, composition of the fuel and the current traffic work. In submission 2013 there has been an extended review of traffic work and the driving distances were updated for all vehicle types all the way back to 1999. One consequence has been a substantial reallocation of fuels between different vehicle categories. But even though there has been a reallocation of fuel, it has not affected the total amount of emissions in any greater degree.

The HBEFA model has also been updated with information regarding the use of air conditioning in passenger cars for all years, which was not available prior to submission 2013. The use of air conditioning usually increases the fuel consumption and consequently the emissions. Data regarding the number of passenger cars equipped with particle filters has also been updated.

The updated and revised data in HBEFA has resulted in both increased and decreased emissions; however emissions of each gas do not show any significant difference as compared to submission 2012.

Data from HBEFA is separated by fuel type and four vehicle types: Passenger cars, Light commercial vehicles, Heavy-duty vehicles (including bus) and Mopeds & Motorcycles. Estimated fuel consumption per fuel and vehicle type is used to proportionally allocate national fuel statistics over those categories.

⁵⁷ Swedish Road- and Transport Research Institute. www.vti.se

⁵⁸ Fridell et al. 2010. "Uppdatering av klimatrelaterade emissionsfaktorer".

⁵⁹ European Monitoring and Evaluation Programme / European Environmental Agency

⁶⁰ Assessment and Reliability of Transport Emission Models and Inventory Systems

The fuel consumption and CO₂ emissions estimated by the Swedish Transport Administration (STA) differ slightly from those reported to the UNFCCC. The STA aims to describe what is emitted on Swedish roads, regardless of where the fuel was bought or the nationality of the vehicles. According to IPCC Guidelines, the inventory should only account for emissions from fuel purchased in Sweden. An overview of the two different objectives is presented in Table 3-18.

Table 3-18. Emissions from road transport reported by the STA and in the CRF.

Fuel bought in	Traffic on Swedish roads	Traffic in Sweden, not on roads	Traffic to/from other country	Traffic in other countries
Sweden	CRF 1.A.3.b STA	CRF 1.A.3.b	CRF 1.A.3.b * STA to the Swedish border	CRF 1.A.3.b *
Other country	STA	Not reported	STA to the Swedish border	Not reported

* Since the IPCC Guidelines do not consider international bunkers for road transportation, all emissions from road traffic and fuel bought in Sweden are considered to be domestic and thus reported under CRF 1A3b.

Emission of N₂O and CH₄ from low blend ethanol-gasoline admixtures are included in the emissions from gasoline for all vehicles in the HBEFA model. HBEFA does however not calculate the emissions of N₂O and CH₄ from ethanol in E85 cars or buses running on bio alcohol.

As from submission 2011 emissions of CH₄ and N₂O from ethanol used by E85 passenger cars and HDV (buses) are estimated by SMED. An implied emission factor derived from gasoline consumption and the related emissions of N₂O and CH₄ from passenger cars in HBEFA is used together with the amount of ethanol used in E85 in cars and buses.

Emissions of N₂O and CH₄ from natural gas and biogas are also missing in HBEFA. As from submission 2012 emissions of CH₄ and N₂O from natural gas and biogas are also estimated by SMED. Activity data from national statistics on delivered amounts of natural gas and biogas is used together with national emission factors⁶¹.

Military transport emissions are reported under CRF 1A5b to be in accordance with the IPCC Guidelines, except for Military road transport which is included in road traffic emissions estimated by HBEFA 3.1.

Emissions in HBEFA for each vehicle type are reduced by proportional amount equal to the weight of the fuel consumption reported by the Swedish Armed Forces relative to the fuel consumption from national statistics allocated to civil road transport.

⁶¹ Paulrud, Fridell Stripplé. Uppdatering av klimatrelaterade emissionsfaktorer. IVL report 2nd edition 2009

3.2.16.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Activity data for natural gas and ethanol is available from 1990, while reliable activity data for biogas exists from 1996 and for FAME from 1999.

One important basic parameter for the HBEFA model is vehicle-km, which is calculated through another model. This second model is based on the mileage driven by the vehicle noted at time of MOT (annual testing of the vehicle).

A passenger car that goes through MOT in the beginning of 2011 has driven the most part during 2010. If the development of traffic is without interruption this issue is not a problem for the calculations. However if a sudden event occurs, such as a drop in the economy, it will not be shown as clearly in the development of vehicle mileage as in statistics on fuel consumption.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

Time series are checked for consistency and recalculations are verified.

3.2.16.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All Tier 1 general inventory level QC procedures and all QC procedures listed in GPG section 8.1.7.4 applicable to this sector are used. The activity data has, of course, been subject to QA/QC procedures prior to the publishing of quarterly fuel statistics. In addition to this, the consumption of every type of fuel in the last year is checked and compared with previous years. If large variations are discovered for certain fuels, responsible staff is contacted for an explanation. IEFs are calculated per fuel, substance and CRF-code and checked against the emission factors to make sure that no calculation errors have occurred when emissions were computed.

3.2.16.5 SOURCE-SPECIFIC RECALCULATIONS

The updated road transport emission model HBEFA 3.1 has resulted in revised emissions except for CO₂ and SO₂, since these are based on statistics of fuel deliveries in Sweden. The updated and revised data in HBEFA has resulted in both increased and decreased emissions, however emissions of each gas do not show any significant difference as compared to submission 2012.

The emissions of N₂O and CH₄ from ethanol were overestimated in submission 2012. The emissions of N₂O and CH₄ from low blended ethanol in gasoline vehicles were already estimated in HBEFA. As the emissions of N₂O and CH₄ from both low blended ethanol and ethanol in E85 vehicles and buses also were calculated separately by SMED, emissions from low blended ethanol were double counted in submission 2012. This has been corrected in submission 2013.

In submission 2012 the emissions of CH₄ and N₂O from low blended gasoline in E85 vehicles were not included. This is corrected in submission 2013. These two adjustments cancel each other out to a certain extent, but the emissions of CH₄ and N₂O are slightly increased in submission 2013.

The consumption of ethanol for road traffic by the military has been added as from 2007.

As from submission 2013 the CH₄ and N₂O emissions from HBEFA are no longer adjusted to be in line with fuel delivery statistics, as these emissions are based on other factors such as mileage, type of vehicle, speed limits etc. These factors are considered in the road traffic model HBEFA.

Correction of previous minor erroneous data also occurs.

3.2.16.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.17 Railways (CRF 1.A.3.c)

3.2.17.1 SOURCE CATEGORY DESCRIPTION

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-19.

Table 3-19. Summary of source category description, CRF 1.A.3.c.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.3.c	CO ₂				T2	CS	Yes
	CH ₄				T2	CR, CS	Yes
	N ₂ O				T1	CR, CS	Yes

CS Country Specific. CR CORINAIR. T2 Tier 2.

The majority of all railway traffic in Sweden runs on electricity. Only a small part runs on other fuels i.e. diesel fuel. According to IPCC's guidelines emissions related to the use of electricity for railway should not be included in this sector. Production of electricity is accounted for in CRF 1A1A, regardless of where it's consumed.

3.2.17.2 METHODOLOGICAL ISSUES

Both Tier 1 and Tier 2 methods are used. Information on emissions from railways is provided by the Swedish Transport Administration (STA). STA estimates the amount of diesel consumed as well as the emissions of CO₂, SO₂, NO_x, NMVOC, CH₄, CO, HC and N₂O.

Emission estimates are based on diesel consumption together with emission factors from three different sources. Emission factors used for calculating CO₂ emissions are supplied by the Swedish Petroleum and Biofuel Institute⁶², whereas emission factors used for HC, NO_x and CO estimates are provided by the Swedish Environmental Research Institute (IVL). Remaining emissions are calculated based on default emission factors from EMEP/CORINAIR.

⁶² www.spbi.se August 2005

3.2.17.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Overall, the emissions for CRF 1.A.3.C is consistent over time and associated with low uncertainties. The estimate of diesel consumption is based on fees paid by the rail operators and is considered to be of very high quality.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

Time series are checked for consistency and recalculations are verified.

3.2.17.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All quality procedures according to the Swedish QA/QC plan (including the Manual for SMED's Quality System in the Air Emission Inventories) have been implemented during the work with this submission.

All Tier 1 general inventory level QC procedures and all QC procedures listed in GPG section 8.1.7.4 applicable to this sector are used. The activity data has, of course, been subject to QA/QC procedures prior to the publishing of quarterly fuel statistics. In addition to this, the consumption of every type of fuel in the last year is checked and compared with previous years. If large variations are discovered for certain fuels, the staff responsible is contacted for an explanation. IEFs are calculated per fuel, substance and CRF-code and checked against the emission factors to make sure that no calculation errors have occurred when emissions were computed.

3.2.17.5 SOURCE-SPECIFIC RECALCULATIONS

Correction of previous minor erroneous data also occurs.

3.2.17.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.18 Navigation (CRF 1.A.3.d)

3.2.18.1 SOURCE CATEGORY DESCRIPTION

The sector covers domestic navigation and leisure boats. Emissions from fuels that are purchased in Sweden but used abroad are reported separately as international bunker emissions.

CO₂ emission from navigation does not show any particular trend, but fluctuates over time.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-20.

Table 3-20. Summary of source category description, CRF 1.A.3.d.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.3.d	CO ₂				T1	CS	Yes
	CH ₄				T1	CR, CS	Yes
	N ₂ O				T1	CR, CS	Yes

CS Country Specific. CR CORINAIR. T1 Tier 1.

3.2.18.2 METHODOLOGICAL ISSUES

Emissions from national navigation are estimated using Tier 1.

Emissions from domestic navigation are calculated based on the amount of fuels that are purchased and consumed in Sweden.⁶³ Emissions from fuels that are purchased in Sweden but used abroad are reported separately as international bunker emissions. The allocation of emissions from navigation is summarized in Table 3-21.

Table 3-21. Reporting of emissions from navigation, according to the Good Practice Guidance.

Fuel bought in	Traffic between Swedish harbours	Traffic between Swedish and international harbours	Traffic between two international harbours
Sweden	Domestic, 1A3d	International bunkers, 1C	International bunkers, 1C
Other country	Not included	Not included	Not included

Emissions from gas/diesel oil and residual fuel oils, for 1990-2002, are calculated using emission factors from a SMED study from 2004⁶⁴. Emissions for 2003 and 2004 have been estimated using the emissions factors for 2002 while emissions for 2005 and later years have been calculated using emissions factors provided by the Swedish Maritime Administration (SMA). The emission factors provided by the SMA are based on the mentioned study by SMED.

Emissions of CO₂ and SO₂ from leisure boats are based on gasoline consumption, together with the same thermal values and emission factors as for civil road traffic. Emissions of NO_x, NMVOC, CH₄, CO and N₂O are all based on gasoline consumption together with emission factors from CORINAIR for gasoline.

The gasoline consumption from leisure boats in Sweden is based on a survey carried out by SMED in 2005⁶⁵. The results from the survey indicate no evidence of any trend in gasoline consumption and the result of the survey (gasoline consumption by leisure boats is estimated to 32,500 m³/year)⁶⁶ is being used as a volume estimate for the whole time period.

⁶³ Statistics Sweden EN31SM

⁶⁴ Cooper and Gustafsson, 2004.

⁶⁵ Gustafsson, 2005.

⁶⁶ Statistics Sweden, 2005a.

Emissions of NO_x, NMVOC, CH₄, CO and N₂O from leisure boats also depend on the ratio between 2-stroke and 4-stroke engines. The estimated ratios between the two are based on a study by Statistics Sweden⁶⁷ from 2005. The study indicates that there is a larger share of 4-stroke engines in 2004 than in 1990. Based on the assumption that the move towards a larger number of 4-stroke engines has been gradual between since 1990, the ratio for each year between 1990 and 2004 has been estimated by interpolation. From 2005 and onwards, the ratio between 2- and 4-stroke engines is assumed to be the same as for 2004.

The Swedish Maritime Administration also report emissions from domestic navigation. These can however not be compared with emissions from the Swedish national inventory since the former include emissions from the whole Baltic Sea region.

3.2.18.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Fuel consumption based on energy statistics from Statistics Sweden's shows fluctuations for which it has been difficult to find natural explanations. See Figure 3-4.

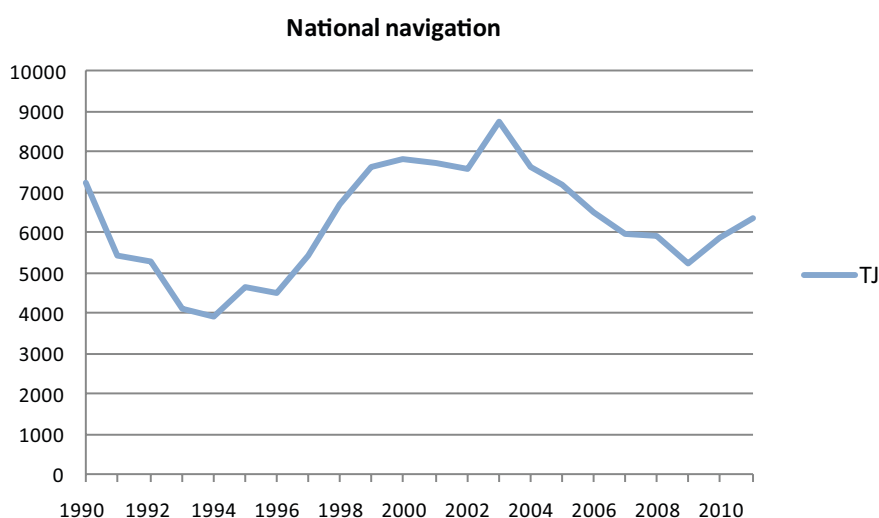


Figure 3-4. Fuel consumption by national navigation

⁶⁷ Statistics Sweden, 2005a.

As a result the fuel consumption by national and international navigation has been reviewed in a SMED study⁶⁸. Fuel data in the Monthly fuel, gas and inventory statistics, which is used as activity data for estimating emissions for national navigation and international maritime bunkers, has been analyzed. The fuel data is collected from oil companies and other sellers who have stocks of petroleum products and coal. The survey also collects stock data from companies with a large consumption of oil in the manufacturing industries and energy industries. The population consists of approximately 70 companies, and all of them are included in the survey.

In 1993, 24 companies reported fuel amounts for domestic navigation and 15 companies reported amounts for international maritime bunkers. During the period 1993-2010 many companies have been closed or taken over by larger companies. In 2010, very few companies remained – only eight companies reported fuel amounts for Domestic navigation or International maritime bunkers. With such a small population, it was possible to study each company carefully. Suppliers of significant quantities of bunker oil (~100,000 m³ – 2,000,000 m³) have been examined more closely. The response from the companies was very good and produced reliable information.

As a consequence of that VAT is applied on national fuel consumption, but not on international bunkers, all respondents to the survey are able to separate these fuel amounts with high accuracy. Fuels used for domestic and international navigation have been separated correctly and in line with IPCC Guidelines.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

3.2.18.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All Tier 1 general inventory level QC procedures and all QC procedures listed in GPG section 8.1.7.4 applicable to this sector are used. The activity data has, of course, been subject to QA/QC procedures prior to the publishing of statistics on fuel purchased. In addition to this, the consumption of every type of fuel in the last year is checked and compared with previous years. If large variations are discovered for certain fuels, responsible staff is contacted for an explanation. IEFs are calculated per fuel, substance and CRF-code and checked against the emission factors to make sure that no calculation errors have occurred when emissions were computed.

An attempt was made to verify the emissions for domestic shipping by comparison with an alternative, independent bottom-up calculation. The bottom-up calculation includes all ship movements in the waters around Sweden. Ship positioning data is gathered using the AIS (Automatic Identification System), which is a complement to radar that provides positions and some static information for almost all ships found in the Baltic and the North Sea. The calcula-

⁶⁸ Eklund et al. 2011. Emissions from navigation and fishing including international bunkers

tions distinguish domestic shipping from international shipping by tracking each ship from its origin to its destination harbour. A route is classified as domestic if origin and destination is within the same country. Where the ship refuels is not possible to distinguish using this method, which causes a slight difference to the reporting guidelines. However, for the purpose of verification this difference is considered to be of little importance. Emission factors are assigned individually for each ship depending on its technical properties. The power output, fuel consumption and emissions are estimated with 5 minute resolution for all ships carrying an AIS transponder. For the years 2009-2011, about 40,000 unique transponder id's are registered by AIS.

The results from the bottom-up calculation show higher emissions than reported emissions from domestic navigation. This is probably related to fishing vessels (reported under CRF 1A4c) and military ships (1A5b). Further studies should also include fishing and military ships to get the whole picture. The data needs to be further analyzed.

3.2.18.5 SOURCE-SPECIFIC RECALCULATIONS

The reported data on residual fuel oil from domestic navigation for 2009 and 2010 are uncertain, due to a merge between refineries during this period. Data is very high and suspected to be partly double reported and it is unfortunately not possible to reproduce the accurate values. As a consequence the decision was made to interpolate the data for 2009 and 2010 based on data 2008 and 2011 and the emissions for these years have been affected. As from 2011 the data on residual fuel oil is considered to be reliable.

Correction of previous minor erroneous data also occurs.

3.2.18.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.19 Other transportation (CRF 1.A.3.e)

3.2.19.1 SOURCE CATEGORY DESCRIPTION

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-22.

Table 3-22. Summary of source category description, CRF 1.A.3.e.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.3.e	CO ₂				T2	CS	Yes
	CH ₄				T3	M	Yes
	N ₂ O				T3	M	Yes

CS Country Specific. T3 Tier 3. M Model.

Emissions reported under CRF 1.A.3.e refer to emissions from off-road vehicles and working machinery including various mobile vehicles and machinery as for example refrigerating plants, wheel loaders, lawn movers, excavators, trimmers, snow blowers and other mobile machine that run on liquid fuels.

3.2.19.2 METHODOLOGICAL ISSUES

The model used to estimate emissions from off-road vehicles and other machineries is considered to correspond to Tier 3 for all emissions, except for CO₂ and SO₂ which are estimated according to Tier 2. The model is further explained in Annex 2.

The model was implemented for the first time in submission 2009 and was reviewed in 2010. In 2012 the model was revised aiming to simplify the use of the model and at the same time update emission factors, activity data and the allocation of emissions to different sectors. Allocation of emissions from off-road vehicles and working machinery is based on a report by Flodström (et al)⁶⁹. This is the most recent inventory including an allocation of working machinery to sectors in Sweden.

Emissions from off-road vehicles and other machinery are also reported under CRF 1.A.2.f, 1.A.4.b and 1.A.4.c, in line with IPCC Guidelines, see Table 3-23.

Table 3-23. Distribution of emissions from off-road vehicles and other machinery

Category	CRF	Definition IPCC Guidelines
Industry	1.A.2.f	Mobile machineries in industry that run on petroleum fuels, as for example tractors, dumpers, cranes, excavators, generators, wheel loaders, sorting works, pumpunits etc.
Residential	1.A.4.b	All emissions from mobile fuel combustion in households, as for example tractors, lawn movers, snow mobiles, forklifts, trimmers, chainsaws and forklifts
Agriculture, Forestry	1.A.4.c	Emissions from mobile fuel combustion in agriculture and forestry, as for example loader-excavator, tractor, harvester, clearing saw etc. Highway agricultural transportation is excluded.
Other	1.A.3.e ii	Combustion emissions from all remaining transport activities including ground activities in airports and harbours, and off-road activities not otherwise reported under 1.A.4.c or 1.A.2.f.

3.2.19.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The calculations are based on a model that takes into consideration emission regulations according to EU legislation in g kWh⁻¹, differences between regulation and value measured at certification, transient use (i.e. difference between static test cycle and real use of the machine), emission deterioration with age and differences between certification fuel and Swedish diesel of type “MK1”. The model does not consider market fluctuations.

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

⁶⁹ Flodström et al 2004. Uppdatering av utsläpp till luft från arbetsfordon och arbetsredskap för Sveriges internationella rapportering.

3.2.19.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

The model is rather new (implemented the first time in submission 2009). During 2010 the model underwent a second verification. Time series are checked for consistency and recalculations are verified.

3.2.19.5 SOURCE-SPECIFIC RECALCULATIONS

No source specific recalculations were made in submission 2013.

3.2.19.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.20 Commercial/institutional (CRF 1.A.4.a)

3.2.20.1 SOURCE CATEGORY DESCRIPTION

This category includes stationary combustion for heating of premises used for commercial and institutional activities. Emissions from mobile combustion used in these activities are included in residential (CRF 1.A.4.b), as it is currently not possible to separate mobile combustion in these two categories from one another.

The heated area was 135 million m² in 2010⁷⁰. Around 30 % of this area consists of schools. Since 1990, the total consumption of fuels for heating of premises has decreased significantly due to the increased use of district heating. In the early 1990s, the total annual fuel consumption in this sector was around 35,000 TJ, around year 2000 it had decreased to about 20,000 TJ, and in 2011 it was around 11,000 TJ. Liquid fuels account for most of the decrease. Over 70 % of the area was heated solely with district heating in 2010. The corresponding share in 1990 was about 55 %.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-24.

Table 3-24. Summary of source category description, CRF 1.A.4.a.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.4.a	CO ₂	X	X		T1	CS	Yes
	CH ₄				T1	CS	Yes
	N ₂ O				T1	CS	Yes

CS Country Specific. T1 Tier 1.

⁷⁰ Swedish Energy Agency, ES 2011:8. Data for 2011 currently not available.

3.2.20.2 METHODOLOGICAL ISSUES

For stationary combustion within CRF 1.A.4.a, all activity data is on national level by fuel type and estimated emissions are therefore considered to correspond to Tier 1. The data source for activity data is the annual energy balance, which for this sector is mainly based on premises statistics that is further described in section 3.2.20.4.1 and in Annex 2.

3.2.20.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. CO₂ from domestic heating oil is the largest uncertainty source. The activity data and emission factor uncertainties are 20 % and 1 % respectively. The large activity data uncertainty is due to the use of Tier 1 methodology with data from the annual energy balances.

3.2.20.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In submission 2005 and earlier, there were large uncertainties in estimation models and allocation methods for fuel in the other sectors and CRF 1.A.2.f, construction. In 2005, a study was performed by SMED, aiming at identifying and analyzing the methods and models applied for each sub-sector and determine whether they were in line with the IPCC guideline recommendations.⁷¹ In addition, each fuel was traced back to its original source in order to determine whether it had been correctly allocated on stationary and mobile combustion.

The results from the study show good agreement with IPCC guideline recommendations. All fuels but biomass had little or no changes in methodologies, and where changes occurred, no significant inconsistencies in fuel consumption time series were detected. However, for biomass, several significant inconsistencies were identified leading to recalculations of activity data and emissions in CRF 1.A.4.a and 1.A.4.b⁷². Due to these recalculations there are obvious inconsistencies between the national energy balances and the national emission inventory data for years before 2005.

Furthermore, all fuels proved to be correctly allocated on stationary and mobile combustion. All diesel oil and gasoline reported under Other sectors in the energy balances is allocated to mobile combustion, while all the other fuels are related to stationary combustion.

3.2.20.4.1 Activity data for stationary combustion in other sectors

For stationary combustion within the Other sectors, the only complete and readily available activity data source is the energy balance. Table 3-25 below shows the consumption of different fuels in 2010 in the different sub-categories

⁷¹ Gustafsson, et al. 2005.

⁷² Paulrud et al. 2005.

within the other sector, and those parts of CRF 1.A.2.f where the energy balance is also used, e.g. construction and small industrial facilities. This data is from the annual energy balance, which is not yet completed for 2011.

Table 3-25. Excerpt from energy balance sheet. Fuel consumption in 2010, TJ

Subsector	Biomass	LPG	Domestic heating oil	Residual fuel oils	Natural gas	Gas works gas
Agriculture, fisheries, 1A4c	6,200	100	3,100	200	500	-
Forestry, 1A4c	-	0	600	300	0	-
Small industrial plants, 1A2f	-	300	1,100	600	200	-
Construction, 1A2f	-	100	1,700	300	500	-
Service, 1A4a	1,700	3,700	3,100	0	3,800	200
Residential, 1A4b	47,100	200	5,900	0	3,600	300

In 2008 all available methods to estimate emissions from stationary combustion in other sectors were overhauled in a SMED study⁷³. The main problem is still that the timeline for the GHG inventory is too short for using final data for other sectors and construction for the latest year. All available alternatives have specific problems including higher uncertainties etc. discussed in the study. The method that was considered to give the best data was using annual energy balances for all years when available, and for the latest year make a model estimate of fuel combustion that adjusts the amounts from the year before with the trend in the preliminary quarterly fuel statistics, as exemplified for 2007 in the equation below:

Estimate 2007 = Annual statistics 2006 * preliminary quarterly fuel statistics 2007 / quarterly fuel statistics 2006

Since emissions for the most recent years are based on this model estimate, uncertainties are a bit higher for this year. Emissions for the most recent years will be revised in next submission when annual statistics are available.

Since 2002, and in particular since 2004, the consumption of biomass fuels has increased in this sector. This is partly explained by the general shift from liquid to biomass fuels in recent years. However, a data check performed in 2009 showed that the data for biomass use in the commercial/institutional sector in the energy balances might not be complete. Further investigations were planned to submission 2011, but this issue was not prioritised since no suitable alternative or complementary data sources were found.

In submission 2010 it was noted that the consumption of biomass, liquid fuels and gaseous fuels within this sector was higher in 2007 than in 2006 and 2008. In submission 2011, the activity data for 2007 and 2008 were revised due to revisions in the energy balances (as described above). The fuel consumption in 2007 is still relatively high. The input data to the energy balances for this sector has not been available for analysis. However, the activity data

⁷³ Lidén and Gerner, 2008

uncertainty is high in this sector and the time series 1990-2010 shows that inter-annual variations in total fuel consumption can be large. Thus the fuel consumption in 2007 is considered to be high, maybe as a result of the large uncertainty, but not erroneous as no calculation errors have been found.

3.2.20.5 SOURCE-SPECIFIC RECALCULATIONS

As indicated in the previous section, activity data has been revised for all fuels for years 2009 and 2010. The greenhouse gas emissions in submission 2013 was 4 Gg CO₂ equivalents higher in 2009 and 32 Gg lower in 2010 compared to submission 2012.

3.2.20.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.21 Residential (CRF 1.A.4.b)

3.2.21.1 SOURCE CATEGORY DESCRIPTION

In this category both stationary and mobile combustion occur. Stationary combustion of fuels within residential decreased by around 50% between 1990 and 2008, mainly due to a continuous increase in district heating use. Most of this change occurred before 2006; however, the use of heating oils is still decreasing while combustion of wood, wood chips and pellets has increased in recent years. In 2009-2010, fuel consumption increased due to the cold winters these years, especially in 2010. Despite this, the consumption of heating oil continued to decrease while consumption of wooden fuels and natural gas increased quite considerably.

Mobile combustion in commercial and institutional services is included in the residential sector, as it is currently not possible to separate mobile combustion in these two sectors from one another. Emissions from mobile combustion refer to emissions from off-road vehicles and other machinery including various mobile vehicles and machines as for example tractors, lawn movers, snow mobiles, forklifts, trimmers, chainsaws and forklifts.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-26.

Table 3-26. Summary of source category description, CRF 1.A.4.b.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.4.b	CO ₂	X	X		T1,T2	CS	Yes
	CH ₄	X	X		T1,T3	CS, M	Yes
	N ₂ O	X			T1,T3	CS, M	Yes

CS Country Specific. M Model. T1 Tier 1. T3 Tier 3.

3.2.21.2 METHODOLOGICAL ISSUES

For stationary combustion, all activity data is on national level by fuel type and estimated emissions are therefore considered to correspond to Tier 1.

For stationary combustion, the main data source is the annual energy balances. One- and two-dwellings statistics, Holiday cottages statistics and Multi-dwellings statistics are used as a complementary data source to get more details on biomass combustion. Biomass fuel consumption for heating residences are surveyed on the three most common combustion technologies: boiler, stoves and open fire places. Since 1998 biomass activity data is separated on wood logs, pellets/briquettes and wood chips/saw dust. Historical biomass data has been estimated by inter- and extrapolation.

Estimation models and allocation methods for fuel in the Other sectors as discussed in section 3.2.20 and use of preliminary data for stationary combustion in other sectors as discussed in section 3.2.20 also applies to CRF 1.A.4.b.

The model used to estimate emissions from off-road vehicles and other machineries is considered to correspond to Tier 3 for all emissions, except for CO₂ and SO₂ which are estimated according to Tier 2. The model is further explained in Annex 2.

3.2.21.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. CO₂ from domestic heating oil is the largest uncertainty source. The activity data and emission factor uncertainties are 20% and 1% respectively. The large activity data uncertainty is due to the use of Tier 1 methodology with indata from the annual energy balances.

The time series for 1.A.4.b is considered to be consistent as there haven't been any major changes in methodology or indata to the energy balances that affect this category. The estimates for the last year, however, are somewhat inconsistent due to the issues described in section 3.2.20. The CO₂ IEF for liquid fuels shows a decreasing trend because the share of residual fuel oil is decreasing. The CH₄ IEF for biomass is slightly fluctuating between years due to variations in type of biomass and technology as described in section 3.2.21.2.

3.2.21.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

See section 3.2.20.4

3.2.21.5 SOURCE-SPECIFIC RECALCULATIONS

Following revisions of the energy balances, the activity data for stationary combustion within 1.A.4.b was revised for all fuels 2009-2010. Compared to submission 2012, greenhouse gas emissions are 5 Gg higher in 2009 and 229 Gg lower in 2010. No source specific recalculations were made for off-road vehicles and working machinery in submission 2013.

3.2.21.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.2.22 Agriculture/forestry/fisheries (CRF 1.A.4.c)

3.2.22.1 SOURCE CATEGORY DESCRIPTION

This category includes emissions from stationary combustion for heating purposes and mobile combustion in off-road vehicles and working machinery within agriculture and forestry, and fishing vessels. The structure of the agricultural sector in Sweden is described in chapter 6.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-27.

Table 3-27. Summary of source category description, CRF 1.A.4.c.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.4.c	CO ₂	X			T1,T2	CS	Yes
	CH ₄		X		T1,T3	CS, M	Yes
	N ₂ O				T1,T3	CS, M	Yes

CS Country Specific. M Model. T1 Tier 1. T3 Tier 3.

3.2.22.2 METHODOLOGICAL ISSUES

For stationary combustion, all activity data is on national level by fuel type and estimated emissions are therefore considered to correspond to Tier 1.

For stationary combustion, activity data is taken from the annual energy balances, which for this sector are based on models and results from a survey from 1985 and repeated in 2007 (see Other statistics from Statistics Sweden in Annex 2).

Estimation models and allocation methods for fuel in the Other sectors as discussed in section 3.2.20, and use of preliminary data for stationary combustion in other sectors as discussed in section 3.2.20.4.1 also applies to CRF 1.A.4.c.

The model used to estimate emissions from off-road vehicles and other machineries is considered to correspond to Tier 3 for all emissions, except for CO₂ and SO₂ which are estimated according to Tier 2. The model is further explained in Annex 2.

Emissions from Fisheries, CRF 1.A.4.c, were first reported in submission 2006. The estimated fuel consumption is based on a survey on energy consumption within the fishing industry by Statistics Sweden⁷⁴ together with data on the Swedish fishing fleets' total installed effect in kW from the Swedish

⁷⁴ Statistics Sweden, 2006 ENFT0601.

Agency for Marine and Water Management (SwAM). The estimate on fuel consumption provided by Statistics Sweden refer to 2005, and for the previous and following years the fuel consumption is estimated by adjusting the 2005 value according to the development in total installed effect.

The emissions factors used to estimate emissions from Fisheries are based on a SMED study from 2005⁷⁵, producing emission factors for CO₂, SO₂, NO_x, NMVOC, CH₄ and N₂O for 1990-2004. From 2005 estimates are based on the same consumption estimate and emission factors as for 2004. However, from 2007 and onwards the emission factors for SO₂ from fisheries are assumed to be the same as for domestic navigation, which are updated every year.

Emissions from fisheries are derived under the assumption that the fishing fleet operates using medium speed diesel engines running on marine distillate fuel. The emission abatement technologies used by the fleet (e.g. Selective Catalytic Reduction (SCR) for NO_x reduction) is assumed to be negligible.

3.2.22.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

The sharp increase in use of biomass in 2003 is due to a revision in submission 2009, where improved data was used for 2003 and later years. There is no information available to improve data from 2002 and earlier years. Emissions in 1990 are considered to be of a sufficient quality as they are based on the 1985 survey mentioned above, which was reasonably recent in 1990. The time series for liquid, solid and gaseous fuels are considered to be consistent. Since almost ten years, solid fuels are not used in this sector.

3.2.22.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

See section 3.2.20.4

3.2.22.5 SOURCE-SPECIFIC RECALCULATIONS

Following revisions of the energy balances, the activity data for stationary combustion within 1A4c was revised for all fuels 2009-2010. Compared to submission 2012, greenhouse gas emissions in submission 2013 are 7 Gg higher in 2009 and 103 Gg lower in 2010.

No source specific recalculations were made for off-road vehicles and working machinery in submission 2013.

3.2.22.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

⁷⁵ Cooper et al., 2005a.

3.2.23 Other stationary (CRF 1.A.5.a)

3.2.23.1 SOURCE CATEGORY DESCRIPTION

No emissions are reported in this sector.

3.2.23.2 METHODOLOGICAL ISSUES

-

3.2.23.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

-

3.2.23.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

-

3.2.23.5 SOURCE-SPECIFIC RECALCULATIONS

-

3.2.23.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

-

3.2.24 Other mobile (CRF 1.A.5.b)

3.2.24.1 SOURCE CATEGORY DESCRIPTION

CRF 1A5b includes emissions from military transports. Emissions from military transports have decreased over the years 1990-2008 due to a decrease in activity. However in 2009 the Swedish military increased its consumption jet kerosene compared to 2008, but in 2010 it went back down again (from 77,894 m³ to 48,692 m³).

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-28.

Table 3-28. Summary of source category description, CRF 1.A.5.b.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.A.5.b	CO ₂		X		T1	CS	Yes
	CH ₄				T1	CS	No, see Annex 5
	N ₂ O				T1	CS	No, see Annex 5

CS Country Specific. T1 Tier 1.

3.2.24.2 METHODOLOGICAL ISSUES

Emissions from military transport are based on data on fuel consumption including all military activities and are considered to correspond to Tier 1. Fuel consumption from some more administrative military activities, such as the Swedish Defence Material Administration (FMV), the Swedish

Fortification Department (FORTV), the Swedish Defence Research Agency (FOI) and the National Defence Radio Institute (FRA), are not included in the calculations.

A special estimation for the use of FAME was conducted by the military for the years 1999-2001. None has been done for the other years.

CH₄ and N₂O emissions from the military are both based on a top-down approach, using fuel consumption (for aviation and navigation) and a bottom-up approach, using data from the HBEFA model (road transport). Hence, estimates are considered to be both Tier 1 and Tier 2. Emissions from military aviation are based on an average of LTO and cruise emission factors. Emissions from military navigation are estimated using emission factors from civil navigation. Emissions from the use of diesel oil by military stationed abroad is reported under Multilateral operations, CRF 1.C.2.

Military road transport is included in the road traffic emissions estimated by the HBEFA model. To subtract and separate emissions from military transport from emissions from civil road transport, emissions according to the HBEFA model for each vehicle type are reduced by an amount equal to the weight of the fuel consumption reported by the Swedish Armed Forces relative to the fuel consumption from national statistics allocated to civil road transport, according to:

$$\text{Equation 0-1: } A = B - \Sigma((C-D)/C * E_i)$$

A = Military transport emissions

B = Total HBEFA emissions

C = Total fuel consumption National Statistics

D = Military fuel consumption Swedish Armed Forces

E_i = HBEFA emissions per vehicle type

3.2.24.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7.

3.2.24.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source specific procedures have been made.

3.2.24.5 SOURCE-SPECIFIC RECALCULATIONS

The consumption of ethanol for road traffic by the military has been added as from 2007 as well as gasoline for military navigation for the years 1990-2001 in submission 2013.

The thermal value and the emission factor for CO₂ for aviation gasoline was used instead of the correct values for jet gasoline for the consumption of jet gasoline by the military for the years 1990-1993. This was adjusted in submission 2013 and resulted in increased emission of CO₂ ranging between 33-43 Gg.

Correction of previous minor erroneous data also occurs.

3.2.24.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.3 Fugitive emissions from solid fuels and oil and natural gas (CRF 1.B)

During all stages from extraction of fossil fuels to final use, escape or release of gaseous fuels, volatile components or absorbed gases may occur. These fugitive emissions are intentional or unintentional escapes and releases of gases from extraction point to final oxidation. In particular, they may arise from the production, processing, transmission, storage and use of fuels, and include emissions from combustion only where it does not support a productive activity (e.g. flaring).

Fugitive emissions in Sweden stem from flaring of fuels in the various categories (iron and steel industry, the chemical industry, refineries and the pulp and paper industry), hydrogen production, transport of crude oil, transmission losses of gas works gas, storage and handling of oil in refineries, depots and gasoline distribution, and natural gas transmission and distribution.

3.3.1 Fugitive emissions from solid fuels (CRF 1.B.1)

3.3.1.1 SOURCE CATEGORY DESCRIPTION

There are no coalmines in Sweden and hence no fugitive emissions from coalmines occur.

Emissions of SO₂ from quenching and extinction at coke ovens are reported in CRF 1.B.1.b. Flaring of coke oven gas from the coke oven is reported in CRF 1.B.1.c since submission 2004. Since submission 2010, flaring of blast furnace gas in the blast furnace and steel converter gas in the steel converter are reported under CRF 2.C.1. CRF 1.B.1 is not really designed to include flaring, but since CRF 1B2 only refers to liquid and gaseous fuels, it is not possible to report flaring of coke oven gas in CRF Table 1.B.2. The amounts of flared gas vary considerably between years, and in 2009 it was unusually high, resulting in increasing emissions in CRF 1.B.1. According to environmental reports⁷⁶, coke oven gas is flared when the production is temporarily stopped because of urgent needs of reparation of equipment or other maintenance measures.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-29.

⁷⁶ SSAB, 2008 and 2009

Table 3-29. Summary of source category description, CRF 1.B.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.B.1	CO ₂				T3	PS	Yes
	CH ₄				T2	CS	Yes
	N ₂ O				T2	CS	Yes

CS Country Specific. PS Plant Specific. T2 Tier 2. T3 Tier 3

3.3.1.2 METHODOLOGICAL ISSUES

The estimation of emissions from flaring of coke oven gas is included in the carbon balance calculations and other plant specific calculations made in cooperation with the two facilities, see section 4.4. Data concerning SO₂ emissions from quenching and extinction at coke ovens are obtained directly from the operators of the two facilities.

3.3.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. The extent of flaring is by nature very variable between years, and the uncertainties in activity data and emission factors are high compared to other activities.

3.3.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

See section 0

3.3.1.5 SOURCE-SPECIFIC RECALCULATIONS

No recalculations have been performed in CRF 1.B.1.b or 1.B.1.c in submission 2013.

3.3.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

3.3.2 Oil and natural gas (CRF 1.B.2)

3.3.2.1 SOURCE CATEGORY DESCRIPTION

In the Swedish inventory, fugitive emissions from a number of different activities related to production and handling of liquid fuels and natural gas are reported in this sector. These activities include hydrogen production at oil refineries (1.B.2.a.1), crude oil transport (1.B.2.a.3), activities in refineries such as catalytic, desulphurisation and storage and handling of oil (1.B.2.a.4), gasoline handling and distribution (1.B.2.a.5), fugitive emissions from transmission and distribution of gas works gas (1.B.2.a.5), fugitive emissions from transmission and distribution of natural gas (1.B.2.b.2 and 1.B.2.b.3), and flaring of liquid fuels (1.B.2.c.2).

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 3-30.

Table 3-30. Summary of source category description, CRF 1.B.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
1.B.2	CO ₂	X	X		T1, T2, T3	D, CS, PS	Yes
	CH ₄	X	X		CS, T1, T2	D, CS, PS	Yes
	N ₂ O				T1, T2	CS	Yes

CS Country Specific. PS Plant Specific. T1 Tier 1. T2 Tier 2. T3 Tier 3. D Default

3.3.2.2 METHODOLOGICAL ISSUES

3.3.2.2.1 *Hydrogen production plants at refineries, CRF 1.B.2.A.1*

According to 2006 IPCC Guidelines, emissions from hydrogen production plants should be reported in CRF 1.B.2.A.1. Since 2005, one such facility is in operation in Sweden, and another one was taken into operation in 2006. Emissions from these facilities are reported in CRF 1.B.2.A.1 in accordance with the 2006 IPCC Guidelines.

Both CO₂ and non-CO₂ emissions are estimated using the Tier 2 method. Activity data as consumed amount of fuels (butane gas and naphtha, respectively for the two plants) and CO₂ emissions are taken from the company's report to the EU ETS system. Non-CO₂ emissions are calculated with this plant specific activity data. National emission factors are used for butane, whereas national emission factors for "other petroleum fuels" are used for naphtha due to lack of specific emission factors for this fuel.

3.3.2.2.2 *Transport, CRF 1B2A3*

Crude oil is transported to and from Sweden by tankers. In response to recommendations from the UNFCCC expert review teams in submission 2010, Sweden estimates emissions of CH₄ from transport of crude oil from submission 2010 and onwards. National statistics available from Statistics Sweden on imported and exported amounts of crude oil is used as activity data. The activity data is corresponding to the data in the Reference Approach. Since no reliable country-specific measurements are carried out, the default emission factor for Western Europe from the Revised 1996 IPCC Guidelines (745 kg CH₄/PJ) is applied. Fugitive emissions of CO₂ from transport of crude oil are not estimated (NE) as no country-specific measurements have been carried out and no default IPCC emission factor for tanker ships is available.

3.3.2.2.3 *Refining/Storage, CRF 1.B.2.A.4*

Sweden estimates both CO₂ and non CO₂ emissions by using the Tier 2 method. The Tier 2 method requires data at plant level and Sweden uses data provided by the refineries in their annual environmental reports. However,

for CH₄ emissions, the IPCC Tier 2 equals a mass-balance estimate, whereas Sweden uses a country specific (CS) method (see below). Emissions are reported from catalytic cracking (CO, SO₂, NOX), desulphurisation (SO₂) and from the storage and handling of oil (NMVOC, CH₄). Catalytic cracking occurs at one plant in Sweden. CO emissions from catalytic cracking are calculated as:

$$CO = \left(\frac{\text{Batched cracker in the material raw of amount}}{\text{Total plant in the material raw of amount batched}} \right) \times \text{Total plant for the emission CO}$$

Due to some operational problems at the plant the total emissions of CO were high for 1997 and 1998 compared to other years.

The emissions of SO₂ from desulphurisation increased in year 2006 compared to previous years due to operational disturbances at one facility.

Fugitive emissions of NMVOC and CH₄ from refineries include emissions from the process area as well as emissions from the refinery harbours when loading tankers. The estimates of NMVOC are mainly based on reported data from the facilities' environmental reports and older reports from the Swedish EPA^{77, 78, 79, 80} and Statistics Sweden⁸¹. The activity data, as crude oil throughput, is known for almost all years. Implied emission factors have been developed, based on reported emissions and known activity data. Reported data for years for which either activity data or emission data is missing have been calculated using the implied emission factors thus developed. In table 3-31, reported NMVOC emissions as well as activity data can be seen. The estimate of fugitive CH₄ emissions are for two refineries based on reported data in the facilities' environmental reports. For the remaining three refineries the fugitive CH₄ emissions are estimated as 5% of the total fugitive VOC emission. This estimate has been provided by one refinery that refines about 50 % of the crude oil in Sweden. Since no information from the two remaining refineries was obtained the same percentage has been used to estimate the fugitive CH₄ emissions even from these plants. The reported emissions of CH₄ are very uncertain due to limited measurements. In Table 3-31, the reported emissions of CH₄ and also activity data can be seen.

The trend of hydrocarbon emission does not follow the fluctuations of the crude oil throughput very well. This is most likely due to the uncertainties in the method used by the refineries to estimate the emissions.

⁷⁷ Swedish EPA, 1990.

⁷⁸ Swedish EPA, 1994a.

⁷⁹ Swedish EPA, 1994b.

⁸⁰ Swedish EPA. 1995.

⁸¹ Statistics Sweden. 1996 Emissions to air in Sweden of volatile organic compounds (VOC) 1988 and 1994.

Table 3-31. Throughput of crude oil in refineries and estimated fugitive emissions of NMVOC and CH₄ (Mg) reported in CRF 1.B.2.A.4.

Year	Throughput of crude oil Mg	Total emissions of NMVOC Mg	Total emissions of CH ₄ Mg
1990	17,330,000	14,408	460
1995	19,430,000	7,643	400
2000	20,253,120	11,568	577
2005	19,919,968	7,691	399
2006	20,050,576	8,269	425
2007	17,706,518	8,877	417
2008	20,420,061	8,578	447
2009	19,669,472	8,779	438
2010	20,278,888	8,924	469
2011	19,034,115	8,938	431

In submission 2009, emissions from combustion of cracker coke in refineries earlier reported in CRF 1A1b were reallocated to CRF 1.B.2.A.4 to be in line with the IPCC guidelines. This is based on a recent study performed by SMED⁸². The cracking reactions produce some carbonaceous material (referred to as *coke*) that deposits on the catalyst and very quickly reduces the catalyst reactivity. The catalyst is regenerated by burning off the deposited coke. Hence the combustion is not carried out for energy purposes and thus the emissions should not be reported in CRF 1.A.

3.3.2.2.4 Gasoline handling and distribution, CRF 1.B.2.A.5

Calculated fugitive emissions of NMVOC from the storage of oil products have been obtained from the environmental reports of the oil depots. The calculations are based on the amount of product handled in the depots. The calculations cover 1990 – 2011 and are based on methods given by Concawe 85/54⁸³ for the years 1990-2006 and on Concawe 03/07⁸⁴ for 2007 and onwards. More than 30 depots have been considered during later years. Gas recovery systems and the recovered amount of gas have been considered in the calculations. For five depot areas the reported NMVOC emissions are based on emission measurements in the depot areas and not on calculations based on the amount product handled in the depots. For years with missing data, interpolation between years with available data has been used for estimates. Handled amounts of gasoline and fugitive emissions of NMVOC from depots and gasoline stations are presented in Table 3-2.

⁸² Skårman, T., Danielsson, H., Kindbom, K., Jernström, M., Nyström, A-K. 2008. Fortsättning av riktad kvalitetskontrollstudie av utsläpp från industrin i Sveriges internationella rapportering. SMED Report 2008

⁸³ Concawe, 1986, Hydrocarbon emissions from gasoline storage and distribution systems, Report No 85/54.

⁸⁴ Concawe Report No. 3/07, Air pollutant emission estimation methods for E-PRTR reporting by refineries

The calculation of fugitive NMVOC emissions from gasoline distribution, 1990-2011 (Table 3-32), is based on methods given by Conca⁸⁵, including annual national gasoline consumption and assumptions on the share of gasoline evaporated at different stages of the handling procedure, as well as effects of applied abatement technology at gasoline stations⁸⁶. The basic assumptions are presented in Table 3-33. Ethanol both for use in blends with gasoline and for use unblended are included in the reported gasoline volume.

Table 3-32. Handled and distributed amount of gasoline and ethanol, estimated fugitive emissions of NMVOC (Gg) from storage at depots and estimated fugitive emissions of NMVOC (Gg) from gasoline stations reported in CRF 1.B.2.a.5.

Year	Volume of gasoline and ethanol m ³	Fugitive emissions of NMVOC at depots Gg	Fugitive emissions of NMVOC at gasoline stations Gg
1990	5,201,225	2.48	13.3
1995	5,257,786	1.93	3.06
2000	5,131,814	2.07	3.08
2005	5,252,212	2.31	3.15
2006	5,136,312	2.47	3.08
2007	5,020,203	2.35	3.01
2008	4,917,095	2.53	2.95
2009	4,638,389	2.41	2.78
2010	4,409,828	2.21	2.65
2011	4,151,038	2.43	2.49

Table 3-33. Assumptions for calculating fugitive emissions from the handling and distribution of gasoline.

Parameter	Assumption	
Density of gasoline	730 kg/m ³ 1990 - 1996 750 kg/m ³ 1997 -	
Distribution of gasoline to gas stations	0.16 %	of distributed volume
Spill	0.01 %	of distributed volume
Filling of car tanks	0.18 %	of filled volume
Measures at distribution to gas station	90 %	Efficiency of measures
Measures at filling cars	70 %	Efficiency of measures

The measures at distribution and filling were introduced over a period of time from 1991-1994, to the extent presented in Table 3-34. The amount of gasoline and ethanol sold at large and small gas stations, respectively, was assumed to be 50/50 for the years 1990-1994. Data on the distributed amounts of gasoline (Table 3-32) is taken from the HBEFA3.1 road emission model. The model is based on a bottom-up approach considered to be Tier 2. The model is described in detail in Annex 2.

⁸⁵ Conca, 1986, Hydrocarbon emissions from gasoline storage and distribution systems, Report No 85/54.

⁸⁶ Andersson, 2000.

Table 3-34. Fraction of gasoline stations with technical measures installed.

Year	Large gas stations > 2,000 m ³	Small gas stations
1990	0%	0%
1991	50%	0%
1992	75%	25%
1993	100%	75%
1994 -	100%	100%

3.3.2.2.5 TRANSFER LOSSES OF GAS WORKS GAS, CRF 1.B.2.A.5

Transmission losses of gas works gas are reported from the producers of gas works gas to Statistics Sweden and published in Statistics on the delivery of gas products (and also reported as “losses” in the annual energy balances). At present, the emissions from these losses are calculated with the same emission factors that are used for stationary combustion, although they are reported as fugitive emissions. The allocation to CRF 1B2A5 is according to the 2006 IPCC Guidelines.

3.3.2.2.6 NATURAL GAS TRANSMISSION, CRF 1.B.2.B.3

Sweden estimates fugitive emissions and emissions due to venting activities from natural gas transmission pipelines in a manner consistent with the IPCC Good Practice Guidance. Default emission factors for fugitives of 2.90×10^{-3} Gg methane per year and per kilometre of transmission pipeline and 1.6×10^{-5} Gg carbon dioxide per year and per kilometre of transmission pipeline from Table 2.16 in the IPCC Good Practice Guidance are used. The methane emission factor is the largest in the range given in Table 2.16 and reflects the use of mostly reciprocating compressors. Since at least one reciprocating compressor is used in the Swedish gas transmission network this emission factor has been used. For estimates of emissions from venting activities, emission factors from the same table in IPCC Good Practice Guidance are used. Emission factors of 1.2×10^{-3} Gg methane per year and per kilometre of transmission pipeline and 8.5×10^{-6} Gg carbon dioxide per year and per kilometre of transmission pipeline are used.

Activity data as kilometre of transmission pipeline has been obtained only for 1996⁸⁷ (320 km) and for 2008^{88, 89}, (620 km). For years before 1996 the pipeline length has been estimated to be of the same length as in 1996 and for 2009 to 2011 the length has been estimated to be as in 2008. The latter assumption is based on information from one of the operators⁹⁰ of the network saying that there has been no extension of the transmission network

⁸⁷ Technical description of the Swedish natural gas distribution system, Swedish gas center, Report SGC 088, June 1997

⁸⁸ Distributionsformer för biogas och naturgas i Sverige, Grontmij – för Svenska Gasföreningen, november 2009

⁸⁹ Tredje inre marknadspaketet för el och naturgas – Fortsatt europeisk harmonisering – betänkande av Nya el- och gasmarknadsutredningen, Statens offentliga utredningar, SOU 2010:30, 2010

⁹⁰ Phone conversation 2011-10-10. Sigvard Trönell (E on)

between 2008 and 2011. For years between 1996 and 2008 the activity data has been estimated by interpolation.

Methane emissions from storage are also reported in 1.B.2.B.3 and are based on national estimates from the operators of the transmission system. There is one storage facility for commercial use in Sweden and it has been in use since 2006.

Figure 3-5 shows total emissions of CH₄ and CO₂ as CO₂ equivalents reported in CRF 1.B.2.B.3.

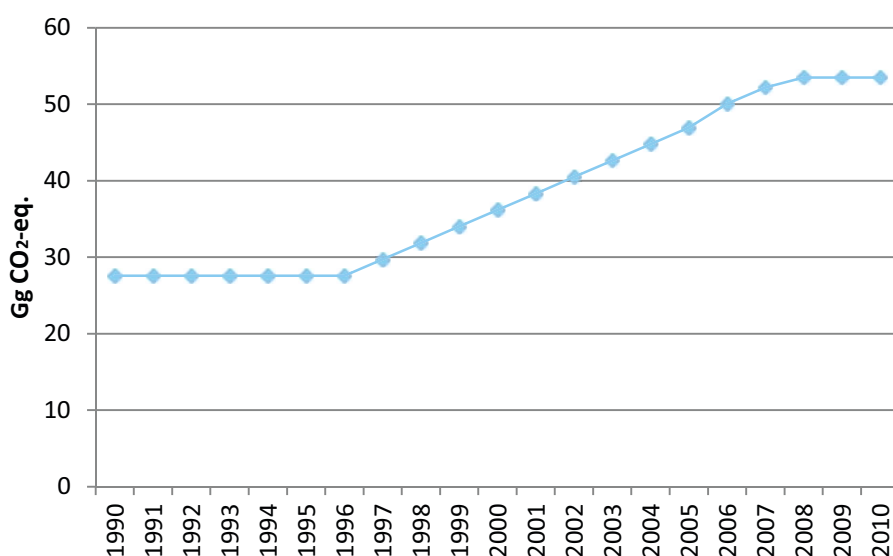


Figure 3-5 Total emissions of CH₄ and CO₂ as CO₂-equivalents reported in CRF 1.B.2.B.3.

3.3.2.2.7 Natural gas distribution, CRF 1.B.2.B.4

Sweden estimates fugitive methane emissions from natural gas distribution in a manner consistent with the IPCC Good Practice Guidance. A default emission factor of 6.15×10^{-4} Gg per year and per kilometer of distribution mains has been used. The used emission factor is an average of the upper and lower default emission factors given in the IPCC Good Practice Guidance, Table 2.16.

The main extension phase of the Swedish natural gas distribution network ended in 1990. Since then there has been a reduced expansion in stages. Activity data as kilometer of distribution mains has been obtained only for 1996⁸⁷ (2,050 km) and for 2008^{88, 89} (2,600 km). For years before 1996 the pipeline length has been estimated to be of the same length as in 1996. For years between 1996 and 2008 the activity data has been estimated by interpolation. The network length in 2009 has been estimated to be 2,610 km and 2010 to 2011 it has been estimated to 2,620 km. The latter estimates are based on information from one of the operators⁹⁰ of the distribution network saying that the network has been extended with approximately 20 km between the end of 2008 and the end of 2010. Since the operator could not estimate the extension per year the expansion has been estimated as 10 km per year.

Figure 3-6 shows total emissions of CH₄ and CO₂ as CO₂ equivalents for the years reported in CRF 1.B.2.B.4.

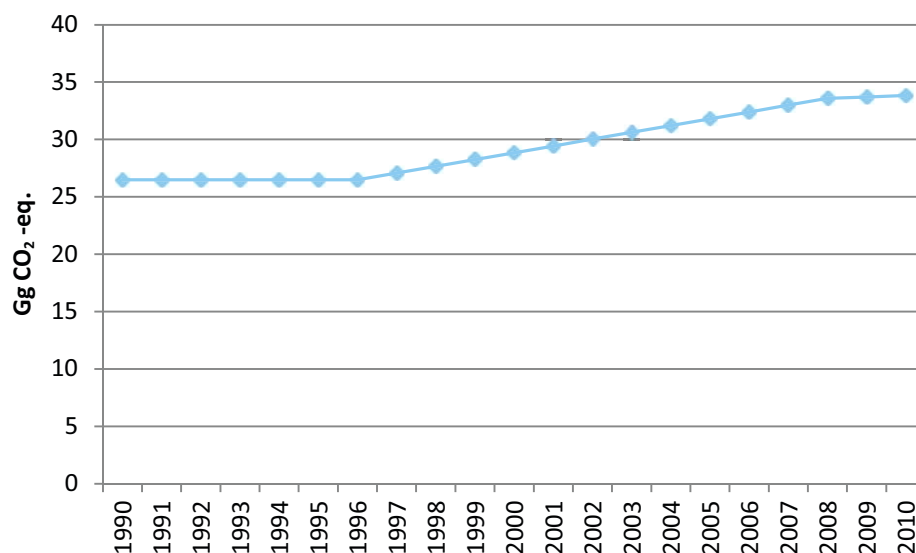


Figure 3-6 Total emissions of CH₄ and CO₂ as CO₂-equivalents reported in CRF 1.B.2.B.4.

3.3.2.2.8 Venting (CRF 1.B.2.C.1)

In submission 2010, emissions from venting at oil refineries were reported using the Tier 1 approach and default emissions factors from the 2000 Good Practice Guidance. For emissions of CH₄ and CO₂, the maximum values in the default EF range were chosen to avoid any risk of underestimating the emissions. In submission 2011, an analysis was carried out with the aim to investigate if vented emissions from refineries already were included in reported emissions in other CRF categories. The conclusion from this study was that the emissions from venting at refineries most probably are included in other categories of fugitive emissions; mainly in CRF 1.B.2.A.4 but possibly partly in 1.B.2.C.2. Hence, it was concluded that the emissions reported in 1.B.2.C.1, in submission 2010 were double counted, and in submission 2011 and later, emissions in CRF 1.B.2.C.1 are reported as IE (in 1.B.2.A.4 and 1.B.2.C.2.) The fugitive CH₄ emissions from oil refineries reported in CRF 1.B.2.A.4 are based on measurements of total hydrocarbon emissions from the refinery areas. These emissions include leakages but also emissions from venting activities. It is therefore not possible to report fugitive emissions and emissions from venting separately. However the hydrocarbon emissions from venting activities at refineries are assumed to be very small, since during normal operation conditions the vented gases enters the gas flare systems. Venting of CH₄ and CO₂ from natural gas transmission pipelines are included in CRF 1.B.2.B.3.

3.3.2.2.9 *Flaring (CRF 1.B.2.C.2)*

Flaring of liquid fuels was estimated and reported for the first time in the Swedish inventory in submission 2005. Data includes flaring of refinery gases at four refineries and one chemical industry, and flaring of LPG at three iron and steel plants and one pulp industrial plant. Emissions in this CRF category varies quite widely between years due to large variations in the amount of refinery gases that needs to be flared each year. Data has been collected directly from the plant operators. For the years 2005 and later, data from the EU ETS system has been used when possible. Data from the EU ETS system are verified against data from environmental reports and vice versa. In submission 2010 EU ETS data was analyzed carefully. It was concluded that the notation key for flaring of natural gas (NE in earlier submissions) could be changed, since no such flaring could be found in the EU ETS data and all plants that might be flaring are included in the EU ETS. There is a slight possibility that some flaring of natural gas is reported included in the flaring of liquid fuels. Because of this the notation key IE is used rather than NO, referring to emissions reported under CRF 1.B.2.C.2.1 Oil.

3.3.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty analysis tables are presented in Annex 7 and a general description of the uncertainties is presented in section 1.7. The extent of flaring is by nature very variable between years, and the uncertainties in activity data and emission factors are high compared to other activities.

3.3.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

The coherence between environmental reports and ETS data is checked when possible, and when differences occur, the facilities are contacted for verification. For a few plants that flare small amounts of gas, activity data as amount of flared gas is shown neither in the environmental reports, nor in the ETS data. Flaring at these plants was investigated in 2005, and the same values are used for later years. Every year, these facilities are asked to verify that the default value is still valid.

3.3.2.5 SOURCE-SPECIFIC RECALCULATIONS

- Since submission 2012 there has been a smaller revision of the distributed amount of gasoline reported from the used road transport emission model HBEFA3.1. The revision resulted in a decrease of NMVOC emissions in CRF 1.B.2.A.5 of 0.2% (0.03 Gg) in 1990 and 4% (0.12 Gg) in 2011, all compared to submission 2012.
- CH₄ and CO₂ emissions from venting activities from natural gas transmission pipelines have been estimated by using the default method in the IPCC Good Practice Guidance. Emissions are allocated to CRF 1.B.2.B.3. In earlier submissions these emissions were not estimated at all.
- CH₄ emissions from storage of natural gas have been estimated by applying estimates from the operators of the transmission system. Emissions are allocated to CRF 1.B.2.B.3. In earlier submissions these emissions were not estimated at all.
- SO₂ emission corrected for one oil refinery in CRF 1.B.2.A.4. Elementary sulphur was reported instead of SO₂.

3.3.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4 Industrial processes (CRF sector 2)

4.1 Overview of sector

For Sweden the most important industries within the industrial sector has historically been base industries such as mining, iron and steel industry and pulp and paper industry. Other important industries when considering emissions of greenhouse gases from industrial processes include the cement industry, primary aluminium production, consumption of fluorinated greenhouse gases and some processes in the chemical industry.

Greenhouse gas emissions from the industrial processes sector have increased 331 Gg CO₂ equivalents from 6,330 Gg CO₂ equivalents in 1990 to 6,661 Gg CO₂ equivalents in 2011, an increase of 5.2% (Figure 4-1). The trend is mainly affected by increased emissions of HFCs (+809 Gg CO₂ equivalents) and CO₂ (+534 Gg CO₂ equivalents), but also in decreased emissions of N₂O (-772 Gg CO₂ equivalents) and PFCs (-194 Gg). In Figure 4-1 it can be seen that in 2011, CO₂ is by far the largest contributor among the greenhouse gases in this sector, accounting for 82.0 % of the GHG emissions. Emissions of HFCs are the second largest greenhouse gas in 2011, accounting for 12.2 % of the sector emissions.

Among the industries in this sector, metal production (CRF 2.C) is the largest contributor in 2011, accounting for 3,458 Gg CO₂ equivalents or 51.9% (Figure 4-2). In Figure 4-2 it can be seen that there was a sharp decrease in greenhouse gas emissions from metal production (CRF 2.C) in 2009. This was mainly due to the economic recession in 2009 which largely affected the production volumes of iron and steel in Sweden and thus the emissions are significantly reduced in 2009.

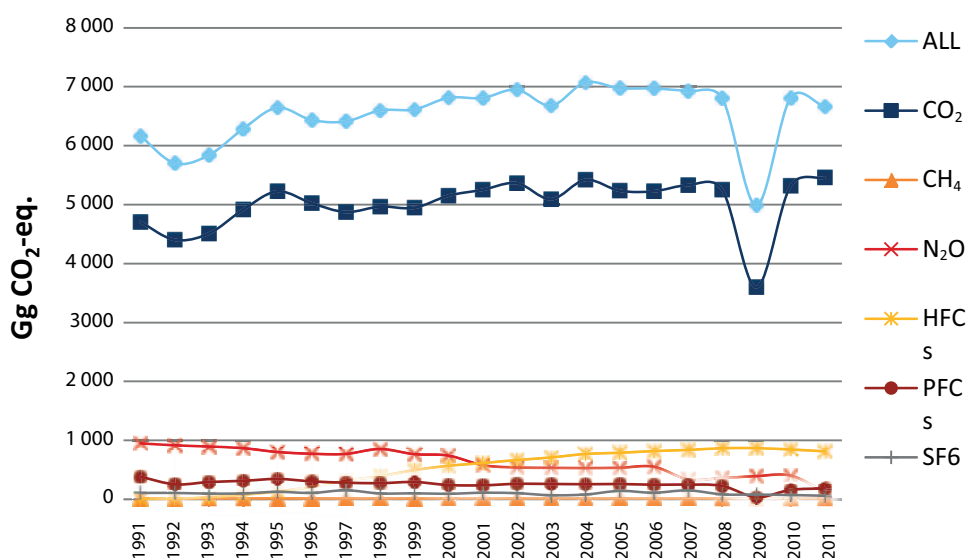


Figure 4-1. Total emissions of all greenhouse gases calculated as CO₂ equivalents from CRF 2 Industrial processes.

The second largest contributor of greenhouse gases to this sector 2011 is mineral products (CRF 2.A) with 2,072 Gg CO₂ equivalents, or 31.1 % of the sector emissions. Compared to 1990 there is an increase in greenhouse gas emissions from mineral products of about 20.4 % (Figure 4-2), mainly due to increased production of lime and clinker.

For chemical industry (CRF 2.B), greenhouse gas emissions have decreased with 776 Gg CO₂ equivalents since 1990 and amounted to 193 Gg CO₂ equivalents in 2011. The reduction is closely linked to N₂O emissions from nitric acid production.

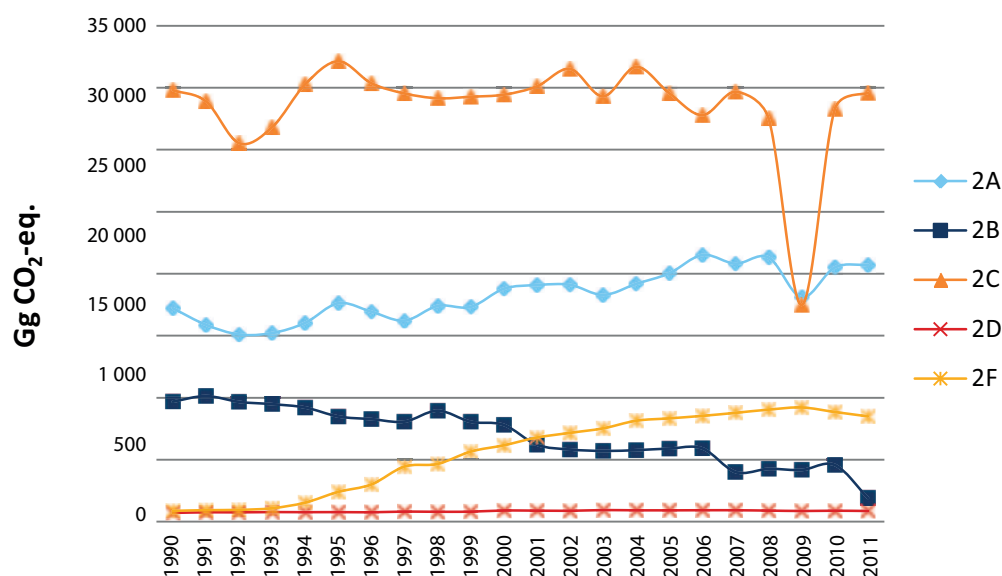


Figure 4-2. Total emissions of all greenhouse gases calculated as CO₂ equivalents from the different Industrial processes sub-sectors.

2A Mineral products. 2B Chemical industry. 2C Metal production. 2D Other production. 2F Consumption of Halocarbons and SF₆.

The estimated emissions of fluorinated greenhouse gases consist of emissions from the use of these in various applications, as well as PFC emissions from the primary aluminium production process. No production of halocarbons or SF₆ (CRF 2.E) occurs in Sweden. The consumption of fluorinated greenhouse gases (CRF 2.F) has increased substantially, 762 Gg CO₂ equivalents, since 1990 (Figure 4-2). The use of HFCs as refrigerants in refrigerators, freezers and air-conditioning equipment has contributed to the larger share in later years.

Process emissions from pulp and paper in other production (CRF 2.D) do not contribute significantly to the emissions of greenhouse gases in Sweden. Table 4-1 shows the impact of recalculations reported in submission 2013 for GHG emissions by sector and sub-sectors for 1990, 1995, 2000 and 2005-2010. The recalculations are mainly due to exclusion of previously double-counted CO₂ emissions from organic carbon in the raw material in cement production (CRF 2.A.1) 2005 – 2010.

Table 4-1. Impact of recalculations of GHG emissions submission 2013 in the industrial processes sector.

Impact of recalculations submission 2013 (Gg CO ₂ eq.)							
CRF	2A	2B	2C	2D	2F	Total CRF 2	% CRF 2
1990	NA	NA	NA	NA	NA	0	0.0%
1995	NA	NA	NA	NA	NA	0	0.0%
2000	NA	NA	NA	NA	0	0	0.0%
2005	-28	NA	NA	NA	0	-28	-0.4%
2006	-30	NA	NA	NA	0	-30	-0.4%
2007	-28	NA	1	NA	0	-27	-0.4%
2008	-30	NA	1	NA	0	-28	-0.4%
2009	-26	NA	1	NA	0	-25	-0.5%
2010	-27	NA	1	NA	-5	-30	-0.4%

0: value less than 0.5. NA: no recalculation is performed.

4.2 Mineral products (CRF 2.A)

Reported emissions include estimates for cement production (2.A.1), lime production (2.A.2), limestone and dolomite use (2.A.3) soda ash use (2.A.4), asphalt roofing (2.A.5), road paving with asphalt (2.A.6), and other (2.A.7). In the source category other (2.A.7), glass production (2.A7.1), non-iron ore mining and dressing plants, glass and mineral wool production, LECA production and production of roofing tiles, bricks and ceramics are included. Until 1998 also emissions from battery manufacturing are included in code 2.A.7.

4.2.1 Cement production (CRF 2.A.1)

4.2.1.1 SOURCE CATEGORY DESCRIPTION

Cement production occurs at three facilities in Sweden (owned by one company), with one being dominant. Annual production of cement in Sweden is about 2,000-3,000 ktonnes. Emissions from cement production stem both from combustion of fuels and from raw materials used in the processes. Emissions arising from fuel combustion are, with the exception of NO_x, reported in the energy sector (CRF 1.A.2.f).

For process-related emissions, facility data are obtained from environmental reports, EU ETS (European Union Emission Trading Scheme) and by direct contacts with the facilities. Process related CO₂ emissions from cement production arise as a by-product during the production of clinker as limestone is heated to produce lime. Process related CH₄ and N₂O emissions from cement production are not occurring according to the IPCC Guidelines and thus reported as not applicable (NA).

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-2.

Table 4-2. Summary of source category description, CRF 2.A.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.1	CO ₂	X			T2	PS	Yes
	CH ₄				NA	NA	NA
	N ₂ O				NA	NA	NA

T2 Tier 2. PS Plant-specific.

4.2.1.2 METHODOLOGICAL ISSUES

In line with the Good Practice Guidance Tier 2 methodology, plant-specific CO₂ emission estimations in Sweden are based on clinker production and include emissions from by-pass dust and cement kiln dust (CKD) as well as emissions from organic carbon of raw meal.

For 1990-2004, information from the company on CO₂ emissions is based on clinker production and default EF from GHG protocol, CKD correction factor and organic carbon in raw meal.

CO₂ = Production of cement clinker (Gg) * 0.525 (Gg CO₂/ Gg clinker, i.e. default value in the GHG-protocol) * CKD correction factor+ CO₂ from organic carbon content of raw meal

The emission estimates were made on initiative by the WRI (World Resources Institute) for the WBCSD (Working Group Cement CO₂ Emissions Inventory Protocol, Version 1.6.), see Facts about the GHG protocol below and on their website⁹¹. The protocol tool calculates CO₂ emissions from raw material converted to clinker, by-pass dust and CKD discarded, and has been used for all years except 1991-1994 and 1996, for which insufficient information was provided from the plants. Instead the cement company has reported production and CO₂ emissions 1991-1994 and 1996 based on mean values from adjacent years.

Facts about the GHG protocol

The GHG protocol has been developed to enable companies to uniformly report their emissions of greenhouse gases. Emissions from stationary combustion and from processes are included.

Over 500 experts have developed the protocol and it is used by over 150 companies including industry associations representing pulp and paper, aluminium and cement. The protocol for CO₂ emissions from the production of cement (WBCSD CSI, version 2.0) can be found on: <http://www.ghgprotocol.org>

From 2005, the company reports plant-specific data on CO₂ emissions to the EU ETS. The CO₂ emissions are based on production of clinker and CaO content of clinker, but also include CO₂ contained in released non-recycled dust (CKD and by-pass) as prescribed by the national guidelines for reporting to the EU ETS⁹². Also CO₂ emissions from organic carbon of raw meal are included in the CO₂ emissions reported in the EU ETS.

⁹¹ <http://www.ghgprotocol.org>. 2005-10-20.

⁹² Lyberg, A., Cementa, Personal communication, September 2011

Table 4-3 shows information on clinker production and total CO₂ emissions from clinker production. For the years prior to 2005 the table shows the calculated emissions from CKD and the resulting CKD correction factor as well as CO₂ emissions from organic carbon content of raw meal.

Table 4-3. Amount of produced clinker and associated CO₂ from specific sources.

Year	Clinker Production	Total CO ₂ emissions	CO ₂ from Clinker (from 2005 incl. CKD and organic carbon content)	CO ₂ from CKD	CKD correction factor	CO ₂ from organic carbon content of raw meal
	Gg	Gg	Gg	Gg		Gg
1990	2,348	1,272	1,233	13	1.010	27
1995	2,405	1,296	1,263	6	1.005	27
2000	2,389	1,288	1,254	6	1.005	27
2005	2,457	1,313	1,313	IE	NA	IE
2006	2,660	1,439	1,439	IE	NA	IE
2007	2,493	1,337	1,337	IE	NA	IE
2008	2,644	1,395	1,395	IE	NA	IE
2009	2,305	1,260	1,260	IE	NA	IE
2010	2,454	1,322	1,322	IE	NA	IE
2011	2,603	1,359	1,359	IE	NA	IE

IE - Included elsewhere. NA – Not applicable.

Total emissions of NO_x by facility are found in the environmental reports or have been obtained directly from the company. They include emissions from fuel combustion as well as from industrial processes. Due to the use of a large variety (within and between years) of waste fuels there is a lack of correct emission factors for the energy sector (1.A.2.f), and thus all emissions are reported in this source category as process emissions. Reported emissions are decreasing over time since 1990.

Data on SO₂ emissions from cement production has been obtained directly from the company or from the environmental reports. Reported emissions are decreasing over time since 1990.

4.2.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Activity data and CO₂ emissions are reported to the EU ETS and have thus been verified by an accredited verification body. The uncertainty for activity data is judged to be ± 2 % and the uncertainty of the emission factor for CO₂ is judged to be ± 5 %.

All three cement producing facilities in Sweden are covered in the reported estimates and the time-series are considered complete, accurate and more or less consistent. As described above, for 1990-2004 constant CO₂ EF (0.525 Gg CO₂/Gg clinker produced) is used together with CKD correction factor and CO₂ emissions from organic carbon of raw meal. Since 2005, CO₂ emissions are retrieved from EU ETS (including CKD and emissions from organic carbon in the raw meal). This means that different methods are used over

time. In Figure 4-3 it can be seen that CO₂ IEF for total emissions from clinker production show larger variations after the introduction of EU ETS data as data source. The varying IEF may depend on the variation of the composition of the raw material. A larger proportion of marl gives lower CO₂ emissions.

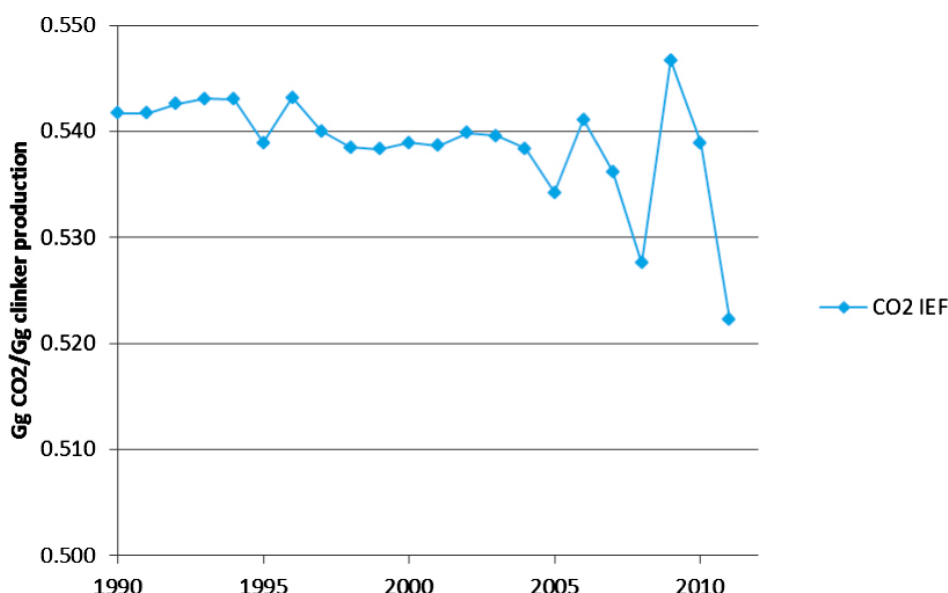


Figure 4-3. CO₂ IEF for total emissions from clinker production 1990-2010

4.2.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In response to previous review recommendations, discussions with the cement producers have been finalized and it has been concluded that CO₂ emissions from dust and from carbon content in the raw material are included in the estimations for the whole time-series (see methodological issues above). In Table 4-3 above, information on clinker production, emissions from production, the calculated emissions from CKD before 2005 and the corresponding CKD correction factors are presented. Compared to the IPCC default value (1.02) the presented CKD correction factor is generally lower which is in line with the conception that dust emission in Sweden is low or nearly non-existent.

Moreover, the implied emission factor for total CO₂ emissions 2005-2011 (average 0.5435 Gg CO₂/Gg clinker produced) is somewhat higher than the IPCC Guidelines default value (0.5071 Gg CO₂/Gg produced clinker) and among the highest of the reporting Parties of the Convention. This is due to higher CaO content in clinker (data for 2008 and 2009 shows a variation in CaO content between 63.9 to 67.6 %) compared to IPCC Guidelines default value (64.6 %), but also because emissions from the organic carbon content of raw meal and CKD are included in the Swedish estimates whereas these sources are not included in the IPCC Guidelines default value.

To follow the Good Practice Guidance Tier 2 method, information shall also include the CaO content of the clinker and data on non-carbonate feeds to kilns. The cement production company reports the CaO content of the clinker to be approximately 65 %.

4.2.1.5 SOURCE-SPECIFIC RECALCULATIONS

In the previous submission, CO₂ emissions from organic carbon in the raw material were double-counted for the years 2005 – 2010. For these years CO₂ emissions from organic carbon are included in emissions reported in EU ETS. In submission 2103 this has been corrected. Due to this recalculation reported CO₂ emissions 2005 - 2010 were reduced with around 2% per year, representing between 26 and 30 Gg CO₂.

4.2.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.2.2 Lime production (CRF 2.A.2)

4.2.2.1 SOURCE CATEGORY DESCRIPTION

In Sweden, quicklime, hydraulic lime and dolomite lime is produced at a number of facilities, owned by a few companies. Produced lime is, for instance, used in blast furnaces, in sugar and carbide production and in the pulp and paper industry to bind impurities and purify the produced material. The production of lime has increased steadily since 1990 from about 400 Gg to about 700 Gg in 2011. In 2009 there was a large decrease in lime production due to an economic recession in the EU. CO₂ is emitted during lime production through calcination of the calcium carbonate (CaCO₃) in limestone, or through the decomposition of dolomite (CaCO₃·MgCO₃). Process related CH₄ and N₂O are not emitted during lime production and thus report as not applicable (NA). Lime contains sulphur which is released as SO₂ during the production process.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-4.

Table 4-4. Summary of source category description, CRF 2.A.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.2	CO ₂				D	D	Yes
	CH ₄				NA	NA	NA
	N ₂ O				NA	NA	NA

D Default.

4.2.2.2 METHODOLOGICAL ISSUES

4.2.2.2.1 CO₂ (Gg)

The emissions of CO₂ from the production of lime are based on activity data by type of lime, i.e. on produced amounts of quicklime, hydraulic lime and dolomitic lime, and corresponding EFs and purity of the limestone from the 2006 IPCC Guidelines. As CO₂ emissions also depend on the production process, the methods for collecting activity data and estimating CO₂ emissions are described by data source (i.e. the sugar industry, the pulp and paper industry and other production of lime) below.

Sugar industry

For determining activity data and emissions of CO₂ within the sugar industry, the amounts of limestone for the production of quicklime are used. The quantities are obtained directly from the sugar producing company from 1999. For years prior to 1999 no data on used amounts of limestone are available. For those years the amounts of limestone used for production of quicklime are estimated using the quantity of coke used for lime production 1990 – 1998, together with the average ratio coke/limestone for the years 1999 to 2002. According to the company the used limestone consists to 97 % of CaCO₃.

In the production of sugar, lime is used for purification of the juice. Lime is added to the raw juice and some impurities are precipitated. In the carbonisation step CO₂ is bubbled through the juice and most of the remaining lime is precipitated as CaCO₃. The precipitated “limestone” is sold and used within agricultural activities. Information from the company gives that around 88% of the lime used was precipitated as CaCO₃ for the years before 2005. For later years this share has increased and varies between 90 and 94 %.

In earlier submissions the whole amount of lime produced and used within the sugar industry was reported as activity data without taking into account that a large amount of the produced lime is precipitated as CaCO₃ in the carbonation process. Since submission 2010, only the part of CaO which is not recovered as CaCO₃ is reported as activity data.

In Table 4-5 the used amounts of limestone, the amounts of produced lime and emitted CO₂, the precipitated CaCO₃, and the reported activity data and CO₂ emissions from lime production within the sugar industry is presented.

Table 4-5. Limestone used, amount of produced lime and emitted CO₂, precipitated CaCO₃ and reported activity data and CO₂ emissions from lime production within the sugar industry

Year	Used amounts of limestone Gg	Amount of lime produced Gg	CO ₂ from lime production Gg	Precipitated share of lime %	Precipitated amount of lime Gg	Reported Activity Data (lime) Gg	Reported CO ₂ emissions Gg
1990	94.7	51.4	40.4	87.5%	45.0	6.4	5.0
1995	76.4	41.5	32.6	87.5%	36.3	5.2	4.1
2000	70.0	38.0	29.9	87.5%	33.3	4.8	3.7
2005	60.9	33.1	26.0	92.0%	30.4	2.6	2.1
2006	68.1	37.0	29.1	92.0%	34.0	3.0	2.3
2007	48.6	26.4	20.7	91.3%	24.1	2.3	1.8
2008	57.3	31.1	24.4	94.4%	29.4	1.7	1.4
2009	55.8	30.3	23.8	94.1%	28.5	1.8	1.4
2010	43.3	23.5	18.5	92.7%	21.8	1.7	1.3
2011	59.8	32.5	25.5	90.4%	29.4	3.1	2.4

Pulp and paper industry

In response to previous review recommendations, since the 2011 submission, detailed data on the quantities of lime used as make-up lime in the pulp and paper industry, and quantities of limestone and dolomite used for production of make-up lime, have been obtained from the Swedish Lime Association and The Swedish Lime Industry from 1995⁹³.

Based on 2006 IPCC Guidelines, the purity of the limestone is set to 95 % for the production of lime within the pulp and paper industry. The corresponding figure for dolomite is 100 %. For the years before 1995, the amounts of make-up lime consumed are estimated using the average ratio between the quantity of make-up lime used and kraft pulp produced for the period 1995 – 2009 and corresponding production data for 1990 – 1994. Earlier information on the need for make-up lime has indicated that it would be less than 20 kg per Mg pulp. New information from a small number of Swedish pulp and paper industries shows that the need may vary considerably, from less than 10 kg per Mg to over 30 kg per Mg⁹⁴. The data used in submission 2011 gives an average need (1995 – 2008) of 20 kg make-up lime per Mg kraft pulp (Table 4-6) and can therefore be considered reliable to use to estimate the need for make-up lime to the pulp industry for years before the 1995. Similarly, the amount of CO₂ emitted is estimated for 1990 – 1994 by using the average ratio between emitted CO₂ and used amounts of make-up lime for the period 1995 – 2008. The used amount of Make-up lime was very low in 2009 which led to a reduced amount of emitted CO₂ in 2009 compared to previous years.

⁹³ Swedish Lime Association and The Swedish Lime Industry, Svenska Kalkföreningen, personal communication

⁹⁴ Håkan Sbttriple, IVL Swedish Environmental research Institute, personal communication

Table 4-6. Produced amounts of kraft pulp, IEF (Make-up lime used per produced amounts of kraft pulp), IEF (CO₂ emitted per produced make-up lime) and reported activity data and CO₂ emissions from make-up lime production for the pulp and paper industry

Year	Produced amounts of kraft pulp Gg	Reported Activity Data (Make-up lime) Gg	IEF (Make-up lime/kraft pulp) Gg/Gg	Reported CO ₂ emissions Gg	IEF (CO ₂ /Make-up lime) Gg/Gg
1990	5,944	118.7*	0.020**	88.5*	0.7457**
1995	6,377	119.4	0.019	89.0	0.7458
2000	7,557	138.0	0.018	103.0	0.7459
2005	7,784	171.5	0.022	128.4	0.7463
2006	7,828	156.0	0.020	116.8	0.7463
2007	7,835	188.4	0.024	140.7	0.7460
2008	7,635	164.7	0.022	122.8	0.7458
2009	7,299	116.9	0.016	87.2	0.7458
2010	7,462	169.4	0.023	126.4	0.7458
2011	7,279	172.1	0.024	128.4	0.7458

* estimated

** average ratio for 1995–2009

Other production of lime

For all other production of quicklime, hydraulic lime and dolomite (mainly used in iron and steel production), detailed data from 1990 are obtained from the Swedish Lime Association⁹⁵. To avoid double counting of emissions, activity data for produced quicklime, hydraulic lime and dolomite lime in the sugar industry and the pulp and paper industry has been deducted.

Based on 2006 IPCC Guidelines, the purity of the limestone is set to 95 % for the production of lime in conventional lime mills. The corresponding figure for dolomite is 100 %. The produced amounts of quick lime and dolomitic lime in conventional lime mills was very low in 2009 which led to a reduced amount of emitted CO₂ in 2009 compared to previous years.

Table 4-7. Produced amounts of quick lime and dolomitic lime, emitted CO₂ and IEF (CO₂ emitted per produced quick lime and dolomitic lime) in conventional lime mills.

Year	Reported Activity Data (quick lime and dolomitic lime, excluding lime in sugar and pulp industry) Gg	Reported CO ₂ emissions (excluding emissions in sugar and pulp industry) Gg	IEF (CO ₂ /quick lime + dolomitic lime) Gg/Gg
1990	264.3	201.1	0.7609
1995	258.4	196.4	0.7599
2000	403.9	306.0	0.7576
2005	495.6	375.6	0.7578
2006	543.7	412.1	0.7581
2007	530.5	402.6	0.7590
2008	541.4	409.6	0.7566
2009	400.6	301.6	0.7528
2010	530.6	399.4	0.7527
2011	509.8	382.7	0.7507

⁹⁵ Swedish Lime Association, Svenska Kalkföreningen, personal communication

4.2.2.2.2 SO₂ (Gg)

The emissions of SO₂ from 1990 have been estimated for production of quick lime. The estimations from quick lime production were calculated using emission factors presented in environmental reports by one of the producers⁹⁶. The emission factor provided by the lime producer is substantially higher for 2008 than for earlier years. This resulted in an increase of reported SO₂ emissions for 2008 compared to earlier years. However in 2009 the reported SO₂ emissions were again on the same level as before 2008 due to less use of lime. For 2009, 2010 and 2011 the emission factor for 2008 has been used for the estimation of emissions of SO₂ due to lack of newer information in the environmental reports.

Emissions of SO₂ from quick lime production intended for the pulp and paper industry are not included in the estimates reported in CRF 2.A.2.

4.2.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty for activity data is $\pm 2\%$ and the uncertainty of the emission factor for CO₂ is $\pm 5\%$. The time-series are considered accurate, consistent and complete.

4.2.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

The activity data reported in CRF 2.A.2 has been compared with national statistics from Statistics Sweden in line with the Good Practice Guidance Tier 2⁹⁷.

The comparison (Figure 4-4) shows that national statistics are more irregular but for most years the coherence is good. The differences are especially high in 1998, 1999 and from 2003 and onwards. The national statistics are based on national surveys mainly aiming at collecting data for economic statistics. In these surveys not all facilities are included and for those the produced amounts are estimated, which might lead to over- or underestimations of, in this case, produced amounts of lime. This leads to larger fluctuations and higher uncertainties in the national statistics from Statistics Sweden compared to data from the Swedish Lime Association and the Swedish Lime Industry⁹⁸ is used as data source.

Quick lime is also produced and used within carbide production. According to the IPCC Guidelines, CO₂ emissions arising from this lime should be reported under CRF 2.B.4, together with other CO₂ emissions from carbide production. It is not known whether this lime is included in the national statistics in Figure 4-4, but it is most likely not.

⁹⁶ Nordkalk, <http://www.nordkalk.com>

⁹⁷ Statistics Sweden. Data from the Industrial production database: www.scb.se

⁹⁸ Swedish Lime Association and The Swedish Lime Industry, Svenska Kalkföreningen, personal communication

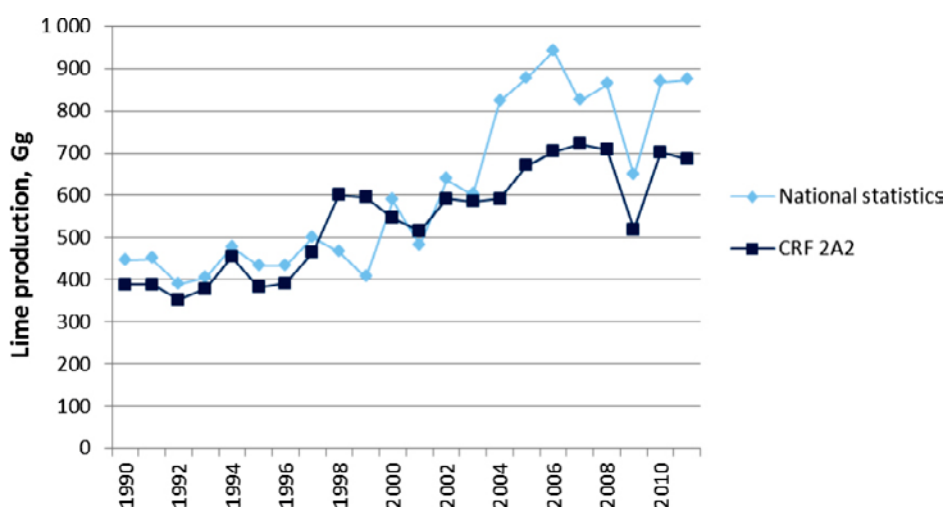


Figure 4-4. National total on produced amount of lime according to data from Statistics Sweden and reported data in CRF 2.A.2.

4.2.2.5 SOURCE-SPECIFIC RECALCULATIONS

Emissions of CO₂ and SO₂ have been recalculated for 2010 due to a minor correction of activity data. The recalculation resulted in an increase of reported CO₂ emissions for 2010 of 1 Gg. For SO₂ the recalculation led to an increase of less than 0.001 Gg.

4.2.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.2.3 Limestone and dolomite use (CRF 2.A.3)

4.2.3.1 SOURCE CATEGORY DESCRIPTION

Limestone and dolomite are used in various processes in Sweden such as cement production, lime production, carbide production, iron sinter production (further described in 2.C.1.3), steel and other metal production, production of clay-based products, glass wool and mineral wool production, glass production, flue gas purification in energy industries and production of chemical products. The use of limestone and dolomite in these processes gives rise to emissions of CO₂. However, not all CO₂ emissions from the use of limestone and dolomite are reported in this source category (see methodological issues below). Process-related CH₄ and N₂O are not emitted during limestone and dolomite use and thus reported as not applicable (NA). The largest contributor of CO₂ emissions in this source category is the iron sinter production. Three facilities (owned by one company) in the north of Sweden produce iron sinter. The use of dolomite in iron sinter production is on average about 80-130 Gg, while the use of limestone was introduced 1999 and has since increased over time to reach the highest use of about 100 Gg in 2010.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-8.

Table 4-8. Summary of source category description, CRF 2.A.3.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.3	CO ₂				CS	D	Yes
	CH ₄				NA	NA	NA
	N ₂ O				NA	NA	NA

D Default. CS Country Specific

4.2.3.2 METHODOLOGICAL ISSUES

This source category comprises of activity data, CO₂ emissions from the use of limestone and dolomite within facilities producing iron sinter, glass wool and mineral wool, chemical products, but also use of limestone and dolomite for flue gas purification.

Process-related CO₂ emissions from the use of limestone and dolomite in the production of cement (2.A.1), lime (2.A.2) and carbide (2.B.4) are reported in corresponding CRF source categories in accordance with the IPCC Guidelines. According to the IPCC Guidelines, all other emissions of CO₂ from the use of limestone and dolomite should be reported as process emissions from limestone and dolomite use in CRF 2.A.3. Since the Centralized review of submission 2004 the ERT has repeatedly recommended Sweden to follow the guidelines. Since the CO₂ emissions from limestone and dolomite are small in some source categories it is not considered to be good practice to spend resources obtaining underlying data to separate these emissions.

Sweden has chosen to not include in 2.A.3 (but in corresponding categories):

- CO₂ emissions from the use of limestone and dolomite in primary and secondary production of steel (2.C.1.1, 2.C.1.2),
- CO₂ emissions from the use of limestone and dolomite in other metal production (2.C.5),
- CO₂ emissions from the use of limestone and dolomite in production of clay based products (2.A.7) and
- CO₂ emissions from the use of limestone and dolomite in glass production (2.A.7.1).

In the case of limestone and dolomite use within the production of glass, the reallocation of CO₂ emissions from 2.A.3 to 2.A.7.1 is due to recommendations from the EC (European Commission) Internal review in 2009. Emissions of CO₂ from use of limestone and dolomite for the production of clay based products are reported in 2.A.7. This is due to the fact that emissions originating from the use of limestone and dolomite seldom are separately reported in the ETS, but rather reported together with other carbon containing raw materials.

Data on the use of limestone and dolomite in this source category has been acquired from environmental reports, the ETS and through direct contacts with the companies. The calculations are made by applying the IPCC Guidelines default emission factors for limestone and dolomite⁹⁹.

Formulas for CO₂ emissions from limestone and dolomite:

$$CO_2 \text{ (Gg)} = \frac{44.0098}{100.0892} \times f \times \text{limestone (Gg)}$$

$$CO_2 \text{ (Gg)} = \frac{88.02}{184.4} \times f \times \text{dolomite (Gg)}$$

where *f* is the purity of the limestone and dolomite, set to 97% and 100% respectively.

In Table 4-9 the use of limestone and dolomite, and corresponding CO₂ emissions, for glass production (2A7.1), primary (2C1.2) iron and steel production and other metal production (2.C.5) are presented for 2005-2011. In relation to the amounts reported in 2.A.3 (Table 4-10), the yearly amounts not included in 2.A.3 represents around 35 % of the total use of limestone and dolomite in Sweden 2005 – 2011.

Table 4-9. Used amounts of limestone and dolomite and estimates of corresponding CO₂ emissions for glass production, primary steel production and other metal production, 2005–2011.

Year	2A7.1		2A7-clay based products		2C1.1		2C1.2		2C5	
	AD, Gg	CO ₂ , Gg	AD, Gg	CO ₂ , Gg	AD, Gg	CO ₂ , Gg	AD, Gg	CO ₂ , Gg	AD, Gg	CO ₂ , Gg
2005	68	31.7	*	*	*	*	94	40.7	5	2.1
2006	73	33.7	*	*	*	*	92	39.7	4	1.8
2007	72	33.6	*	*	*	*	80	34.7	5	2.3
2008	73	33.7	*	*	*	*	96	41.3	4	1.8
2009	55	25.5	*	*	*	*	61	26.4	4	1.7
2010	66	30.8	*	*	*	*	87	37.4	6	2.5
2011	66	30.7	*	*	*	*	84	36.2	4	1.6

* not possible to separate CO₂ from limestone/dolomite for included facilities

⁹⁹ IPCC. Revised 1996 Guidelines for National Greenhouse Gas Inventories: Reference Manual section 2.5.2

Table 4-10. Used amounts of limestone and dolomite and corresponding CO₂ emissions reported i 2A3.

Year	Total 2A3 Activity data	Total 2A3 CO ₂	Iron sinter production % of total CO ₂	Flue gas desulphurisation % of total CO ₂	Glass and mineral wool % of total CO ₂	Other chemical industry % of total CO ₂
1990	194	90	42.2	11.5	26.1	20.1
1995	214	100	54.1	14.2	17.9	13.8
2000	245	113	67.4	9.5	15.1	8.0
2005	254	116	70.2	15.8	9.6	4.5
2006	242	109	64.9	18.6	10.9	5.6
2007	268	122	70.1	15.0	9.8	5.1
2008	287	131	73.6	11.7	10.0	4.7
2009	230	105	69.9	13.8	9.3	6.9
2010	298	135	75.3	10.6	9.2	5.0
2011	298	136	75.3	12.1	7.7	4.9

The emissions have increased during the reporting period due to higher limestone and dolomite use in the production of ore-based iron pellets. This increase is however partly compensated by a decrease in the use within the mineral and glass wool industry and the chemical industry. Decreased emissions from the glass wool industry are partly due to an increased use of recycled materials and thereby less need for limestone and dolomite for raw glass wool production. During 2009 the used amounts of limestone and dolomite was lower compared to previous year due to the economic recession.

Data on the use of limestone and dolomite have been acquired from environmental reports, the ETS and through direct contacts with the companies.

4.2.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty for activity data is $\pm 7\%$ and the uncertainty of the emission factor for CO₂ is $\pm 5\%$. The time series are considered accurate, consistent and complete. It is however possible that there are small facilities using (insignificant amounts of) limestone and dolomite which perhaps are not included in the Swedish inventory.

4.2.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed. For facilities part of the EU ETS, data on CO₂ emissions should however be used for verification of calculated CO₂ emissions using the IPCC default values.

4.2.3.5 SOURCE-SPECIFIC RECALCULATIONS

CO₂ and activity data were added for one new facility using limestone for flue gas purification (2004 – 2010). Activity data and corresponding CO₂ emissions were corrected for two facilities (2009 and 2010). Activity data not included in submission 2012 was added for one facility (2005 – 2010). The recalculations resulted in yearly increases in CO₂ emissions of about 0.2 to 0.3 Gg (2004 – 2010).

4.2.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.2.4 Soda ash use (CRF 2.A.4)

4.2.4.1 SOURCE CATEGORY DESCRIPTION

Soda ash is used in the production of glass wool, moist snuff and chemicals i.e. detergents, and until 2004 also in flue gas desulphurisation at energy plants. Soda ash is also used in production of glass (2A7.1). Soda ash is not produced in Sweden. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-11.

Table 4-11. Summary of source category description, CRF 2.A.4.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.4	CO ₂				CS	D	Yes
	CH ₄				NA	NA	NA
	N ₂ O				NA	NA	NA

D Default. CS Country Specific

4.2.4.2 METHODOLOGICAL ISSUES

In 2004 a study was carried out to collect data on soda ash use and calculate CO₂ emissions.¹⁰⁰ From this study it became clear that no production of soda ash occur in Sweden, and is hence reported as NO in the CRF. Activity data consists of soda ash use from ten plants within several areas:

- production of glass wool, moist snuff and chemicals
- until 2004, in flue gas desulphurisation at energy plants

As for the use of limestone and dolomite, the emissions and activity data concerning use of soda ash within the glass industries have been reallocated to 2.A.7.1 due to recommendations from the EC Internal review in 2009. This reallocation reduces the reported CO₂ emissions by approximately 50 % in the early 1990s and by over 90 % for the last years. The reason for the large effect on reported emissions in later years is due to large changes in the use of soda ash in one chemical industry. This industry spent during the early 1990s considerable amounts of soda ash, and has since 1997 sharply reduced their consumption. In the beginning of the new millennium the soda ash used for manufacturing at this industry is bound in products, and thus no CO₂ is emitted.

¹⁰⁰ Nyström. 2004. SMED-report: CO₂ from the use of soda ash.

Activity data for the use of soda within water treatment and moist snuff production, by others than the dominant manufacturer, has been estimated based on information from expert organisations¹⁰¹ and the dominant snuff manufacturer. The emissions are calculated by applying the IPCC Guidelines default emission factors for soda ash for all activity data:

$$CO_2 \text{ (Gg)} = \frac{44.0098}{105.9884} \times \text{sodaash(Gg)}$$

Data on the use of soda ash have been acquired from the ETS and through direct contacts with the reporting companies.

The data used for national GHG estimations from soda ash use is believed to be more consistent and complete, compared with the data from national statistics, since the data for the inventory is collected from the ETS, from the environmental reports of the facilities or by direct contact with the plants.

4.2.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty for activity data is $\pm 7\%$ and the uncertainty of the emission factor for CO_2 is $\pm 5\%$. The time series is consistent and complete for the major plants, but it has to be noted that some facilities using small amounts of soda ash might be missing in the inventory.

4.2.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.2.4.5 SOURCE-SPECIFIC RECALCULATIONS

In current submission corrections have been made for the amounts of soda ash used within one chemical industry (2007 – 2010) and in moist snuff production by others than the dominant manufacturer (2008 – 2010). This recalculation for the years 2007 - 2010 led to minor changes in reported CO_2 (from a reduction of 0.3 Mg to an increase of 3.3 Mg).

4.2.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.2.5 Asphalt roofing (CRF 2.A.5)

4.2.5.1 SOURCE CATEGORY DESCRIPTION

Since the end of the 1990's there have only been two companies in Sweden producing asphalt-saturated felt. Production and emission data provided by the manufacturers have been used for developing emission factors for estimations of the NMVOC emissions. No measurements or estimations on CO emissions have been performed by the industry and are consequently reported

¹⁰¹ The Swedish Chemicals Agency (KemI), www.kemi.se

NE, not estimated, for the whole time-series. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-12.

Table 4-12. Summary of source category description, CRF 2.A.5.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.5	CO ₂				NA	NA	NA

D Default. CS Country Specific. T2 Tier 2. PS Plant-specific.

4.2.5.2 METHODOLOGICAL ISSUES

Data on the total Swedish production of asphalt-saturated felt was provided by the producing companies. Emission factors for asphalt roofing manufacture are presented in EMEP/CORINAIR Emission Inventory Guidebook.¹⁰² These are based on studies performed during the 1970s in the USA and presented by EPA.¹⁰³ As stated in the guidebook, the level of uncertainty regarding the suggested emission factors is high, and it is recommended that better factors should be developed and used.

After contact with the industry, emission factors based on measurements and calculations made by the manufacturers were developed before submission 2005 for estimating the NMVOC emissions from the Swedish production of asphalt-saturated felt (Table 4-13)¹⁰⁴.

Table 4-13. Estimated emissions of NMVOC from manufacturing of asphalt-saturated felt (CRF 2A5) in Sweden 1990 – 2011.

Year	NMVOC emissions from asphalt roofing, 2A5 Mg
1990	77.7
1995	98.6
2000	111.1
2001	112.9
2002	109.2
2003	101.1
2004	113.7
2005	139.7
2006	132.7
2007	142.4
2008	138.6
2009	103.1
2010	105.4
2011	109.6

¹⁰² EMEP/CORINAIR Emission Inventory Guidebook: <http://reports.eea.eu.int/EMEP/CORINAIR4/en>

¹⁰³ Shrager, Brian and Marinshaw, Richard. 1994. Emission Factor Documentation for AP-42, Section 11.2, Asphalt Roofing, Final Report. For U.S. Environmental Protection Agency, Office for Air Quality Planning and Standards, Emission Inventory Branch. MRI Project No. 4601-01.

¹⁰⁴ Danielsson, H. 2004. SMED report: Investigation on the occurrence of emissions from asphalt roofing in Sweden.

The NMVOC emissions from the production of asphalt-saturated felt originate from the felt saturation and coating processes and from leakage from the asphalt storage tanks, the latter being the dominating source. For the calculation of the NMVOC emissions, separate emission factors were used, 0.068 kg/Mg and 1.56 kg/Mg, respectively. The emission factors are based on measurements/estimations from 2003 and 1997. Previously reported notation keys for activity data have been changed from NE to C.

4.2.5.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The time-series are consistent

4.2.5.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.2.5.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.2.5.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.2.6 Road paving with asphalt (CRF 2.A.6)

4.2.6.1 SOURCE CATEGORY DESCRIPTION

Large changes have occurred in asphalt paving technology over the last decade, with a gradual change towards use of water-based emulsions instead of solvent-containing bitumen solutions. Industry representatives estimated that the naphtha content in the solutions used for road paving was on average 23 % in 2002 and 46 % in 2011. In this inventory, only NMVOC emitted in the process of paving the roads is included. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-14.

Table 4-14. Summary of source category description, CRF 2.A.6.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.6	CO ₂				NA	NA	NA

4.2.6.2 METHODOLOGICAL ISSUES

Estimates for the early 1990s are taken from investigations and inventories made in the early 1990s. Data for the years 2002 – 2011 has been calculated based on information from the asphalt producers on the average amount of solvent (naphtha) in the mixtures used for road paving. The producers have also provided figures on the total amount of road paving mixtures delivered in Sweden. It is assumed that all solvents in the solvent-based bitumen are emit-

ted when used. Emissions of NMVOC reported for the years in mid- and late 1990s were interpolated (Table 4-15). In the calculations no emissions from imported solvent-based bitumen are used. The amount of imported solvent-based bitumen is most likely very small. In 2005 the emission of NMVOC was very high due to the fact that a heavy storm ruined many roads in southern Sweden. These roads needed to be restored quickly and solvent-based bitumen was used for this purpose.

Table 4-15. Emissions of NMVOC 1990–2011 from road paving with asphalt.

Year	NMVOC from road paving with asphalt Mg
1990	6,200
1995	3,800
2000	1,170
2005	1,230
2006	750
2007	935
2008	855
2009	341
2010	256
2011	216

4.2.6.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The time-series is consistent.

4.2.6.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.2.6.5 Source-specific recalculations

No source-specific recalculations have been performed.

4.2.6.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.2.7 Other (CRF 2.A.7)

4.2.7.1 SOURCE CATEGORY DESCRIPTION

Specified sub-categories under this heading are “Glass production (2.A.7.1)”, “Non-Iron ore mining and dressing”, “Glass and mineral wool production”, “Battery manufacture” and “Light expanded clay aggregate (LECA), roofing tile, brick, and ceramics production”. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-16.

Table 4-16. Summary of source category description, CRF 2.A.7.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.A.7	CO ₂				CS	CS, D	No, see Annex 5
	CH ₄				NA	NA	NA
	N ₂ O				NA	NA	NA

D Default. CS Country Specific. T2 Tier 2. PS Plant-specific.

4.2.7.1.1 Glass production, CRF 2.A.7.1

In Sweden there is one facility for float glass production, one for container glass and several small facilities for manual glass production. Emissions of CO₂ from the use of limestone and soda ash in glass production are from submission 2010 and onwards reported in 2.A.7.1. In earlier submissions CO₂ from the use of limestone and of soda ash in glass productions were reported under 2.A.3 and 2.A.4, respectively. The CO₂-emissions in 2009 are lower than in adjacent years due to the fact that the demand for glass was low in 2009. Also less amount of glass was manufactured from raw material that year - instead recycled glass was used. From the float glass production, the total emissions of SO₂ and NO_x from the glass furnace are allocated to 2.A.7.1 since a separation into energy-related and process-related emissions is not possible. From the container glass production, SO₂ emissions originating from the raw material and small amounts of NMVOC are reported. All other emissions from the glass production facilities are from combustion for energy purposes, and are allocated to the Energy sector (CRF 1).

4.2.7.1.2 Non-Iron ore mining and dressing, CRF 2.A.7

The only emissions reported for the non-iron ore mining and dressing are, in this submission, NO_x released from use of explosives. Also CO is emitted but no data concerning the CO emissions are available and the time series 1990 – 2011 is thus reported NE. Estimated emissions from combustion of fuels are included in the Energy sector (CRF 1).

4.2.7.1.3 Glass and mineral wool, CRF 2.A.7

Glass and mineral wool production occurs at three facilities run by two companies. Before 2004 there were four facilities but one closed down during 2003.

4.2.7.1.4 Battery manufacturing, CRF 2.A.7

One battery producer of NiCd-batteries previously used iso-propanol in their processes, which resulted in emissions of NMVOC. The process was changed in 1998 and, since then, no NMVOC emissions occur from this source.

4.2.7.1.5 *Light expanded clay aggregate (LECA), roofing tile, brick and ceramics production, CRF 2.A.7*

In this sub-code under 2.A.7 Sweden report CO₂ emissions from production of clay based materials such as LECA, roofing tiles, bricks and ceramics. During the production CO₂ is emitted from the burning of fuels, reported in CRF 1.A.2.f, but CO₂ originating from the clay, the limestone and from other carbon containing material is also emitted. Reported CO₂ emissions represent the emissions from totally six facilities during the years 1990-2008 and from totally five facilities from 2009 and onwards since one facility closed down during 2008. One of the facilities is dominating in CO₂ emissions. All CO₂ emissions from raw material used are reported in 2.A.7.

4.2.7.2 METHODOLOGICAL ISSUES

Specified sub-categories under this heading are, “Non-Iron ore mining and dressing”, “Glass and mineral wool production”, “Glass production”, “Battery manufacture” and “Light expanded clay aggregate (LECA), roofing tile, brick, and ceramics production”.

4.2.7.2.1 *Glass production, CRF 2.A.7.1*

Emissions of CO₂ from the use of limestone and from the use of soda ash in glass production are reported in CRF 2.A.7.1 together with CO₂ emissions from other carbon containing raw material. Of the reported total CO₂ emissions in 2.A.7.1, approximately 44 % is caused by the use of soda ash and 55 % on the use of limestone and dolomite. The remaining CO₂ is emitted as a result of use of other carbon containing raw materials.

Activity data and emissions are mainly collected from the ETS or from the facilities yearly environmental reports. For small glass production plants a constant amount of 0.9 Gg CO₂ per year, and corresponding amount of limestone, is added. This estimate is based on information from a survey made in the late 1990s by the Swedish EPA on small glass production facilities and represents data from 1997. Two different estimates were made, one based on the consumption of carbonates for the production of glass and crystal, and the other based on the knowledge on the percentage weight loss depending on emitted CO₂, from weight of raw material to produced amount of glass or crystal. Both estimates result in CO₂ emissions of around 0.9 Gg, yearly.

The process-related SO₂ emissions from container and float glass production are reported for the period 1990 – 2011 in CRF 2.A.7. The reported NO_x emissions originate from the production of float glass. Data has been provided directly by the companies or collected from their environmental reports.

4.2.7.2.2 *Non-Iron ore mining and dressing, 2.A.7*

Data on NO_x emissions from use of explosives within the non-iron ore mining industry are reported 2002 – 2011, but for the years 1990 – 2001 no information is presently available. Data on NO_x emissions are collected from the companies' environmental reports to the authorities.

4.2.7.2.3 *Glass and mineral wool production, 2.A.7*

Within mineral wool production, the limestone and dolomite used cause process emissions of CO₂ which are allocated to CRF 2.A.3 according to the IPCC Guidelines. For some years however (1990-1995 and 1998-1999), blast furnace slag was used in the process causing CO₂ emissions as well. These emissions are reported in CRF 2.A.7. Activity data on the slag consumption has been obtained for the mentioned years from the mineral wool producers. The emission factor is 0.04 Gg CO₂ /Gg slag based on that the slag contains 1 % carbon and the CO₂ emissions are calculated by using the formula:

$$\text{Emissions of CO}_2 \text{ (Mg) from use of slag} = \text{Slag (Mg)} * 0.01 * (\text{C content}) * 44/12$$

For glass and mineral wool production, the time series of NMVOC emissions is based on data received from the companies directly or as reported in environmental reports together with earlier total estimates. The emissions of NMVOC consist of formaldehyde and phenol.

4.2.7.2.4 *Battery manufacture, 2.A.7*

NMVOC emissions from battery manufacture for the period 1990-1998 are compiled from data presented in the companies' environmental reports. The process has changed and no emissions of NMVOC occur after 1998.

4.2.7.2.5 *Light expanded clay aggregate- (LECA), roofing tile, brick and ceramics production, 2.A.7*

Activity and emissions data for LECA production 1990 - 2004 is retrieved directly from the production plant, split into emissions from clay and emissions from additives (limestone and other carbon containing material). From 2005 and onwards, the equivalent data is acquired through the ETS and the Swedish LECA producer's annual report.

For roofing tile, brick and ceramics production, activity and emission data from 2005 and onwards is acquired through the ETS. The data in the ETS does not always separate between emissions from limestone/dolomite use and CO₂ emissions from other carbon containing raw material (i.e. from the clay and other carbonates used) needed for the production. In order to as far as possible report an accurate total process-related CO₂ emission for the facilities included in this 2.A.7 sub-code, Sweden have chosen to report all CO₂ emissions in 2.A.7.

As there is a lack of data before 2005, the reported emissions for 2005 are extrapolated for 1990-2004.

As activity data reported in this 2.A.7 sub-code produced amounts of LECA is reported due to lack of activity data for remaining facilities. The implied emission factor may vary somewhat from one year to another because of the specific composition of limestone, clay and additives with different carbon contents. In 2007, the C-content in one of the additives for LECA pro-

duction was unusually high which has resulted in comparatively high CO₂-emissions for that year. The use of limestone and other additives in LECA production has declined in favour of clay which today contributes to about 88 % of all process related CO₂ emissions from LECA production. The facility producing LECA corresponds to around 75 % of yearly reported CO₂ emissions in this 2.A.7 sub-code.

4.2.7.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainties of the direct CO₂ emissions in 2A7 are considered to be ± 7 % based on expert judgements.

4.2.7.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.2.7.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed except for non-iron ore mining and dressing, 2.A.7. Data on NO_x emissions from the use of explosives in 2010 was added. The recalculations resulted in an increase in NO_x emissions of about 0.076 to 0.079 Gg yearly (2010).

4.2.7.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.3 Chemical industry (CRF 2.B)

Sources covered in the reporting are nitric acid production (2.B.2), carbide production (2B4) and other (2B5), which include a large variety of processes in the chemical industry. No ammonia production (2.B.1) or adipic acid production (2B3) occurs in Sweden.

4.3.1 Ammonia production (CRF 2.B.1)

4.3.1.1 SOURCE CATEGORY DESCRIPTION

There is an annual production of about 5 Gg of ammonia in Sweden, according to United Nation statistics¹⁰⁵. This ammonia is however not intentionally produced, but is a by-product in one chemical industry producing various chelates and chelating agents, such as EDTA, DTPA and NTA¹⁰⁶. Emissions from this industry are included in CRF code 2B5. Ammonia production, 2.B.1, is thus reported as NO in the CRF-tables.

¹⁰⁵ UN. Commodity Production Statistica Database. Department of Economic and Social Affairs, Statistics Division,. As referred in FCCC Synthesis and Assessment report 2002 Part I.

¹⁰⁶ Kindbom, 2004. SMED report: Investigation on the occurrence of ammonia production in Sweden. 2004-05-11.

4.3.2 Nitric acid production (CRF 2.B.2)

4.3.2.1 SOURCE CATEGORY DESCRIPTION

Production of nitric acid has taken place at three facilities in Sweden during 1990-2000. One of these was shut down at the end of 2000, and a second one was shut down during 2001. Therefore, there is currently only one facility producing nitric acid in Sweden. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-17.

Table 4-17. Summary of source category description, CRF 2.B.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.B.2	CO ₂				NA	NA	NA
	CH ₄				NA	NA	NA
	N ₂ O		X		T2	PS	Yes

T2 Tier 2. PS Plant-specific.

4.3.2.2 METHODOLOGICAL ISSUES

Activity data, such as the produced amount of nitric acid, has been obtained from the facilities and from official statistics. Emission estimates of N₂O have been reported in the companies' environmental reports or have been provided by the facilities directly. Emission data is not available for all facilities for 1991-1993. Since two plants have been shut down, it is no longer possible to acquire this information. Calculations have therefore been made based on production statistics and an assumed emission factor (Table 4-18). The assumed emission factor of 7 kg/Mg for 1991 - 1993 is based on the calculated emission factors for 1990 and 1994 and is in line with the default factors for nitric acid production presented in Table 4.7 in IPCC Good Practice Guidance.

Documentation has been received from the facility concerning production data, production capacity and abatement measures, used emission factors and the method used for estimating emissions as well as uncertainty in emission estimates. However, this information is confidential.

Table 4-18. Activity data, emission factors and emissions for N₂O for nitric acid production.

Year	Production of nitric acid Gg	Calculated EF (1990 and 1994-2011),kg/Mg	Emissions of N ₂ O, Gg
1990	374	7.02	2.63
1991	395	7.00*	2.77
1992	380	7.00*	2.66
1993	369	7.00*	2.58
1994	377	6.62	2.50
1995	417	5.48	2.29
1996	400	5.48	2.19
1997	390	5.56	2.17
1998	400	6.10	2.44
1999	383	5.58	2.14
2000	430	4.80	2.06
2001	282	5.48	1.55
2002	263	5.41	1.42
2003	258	5.39	1.39
2004	257	5.37	1.38
2005	264	5.37	1.42
2006	272	5.42	1.47
2007	249	3.16	0.788
2008	266	3.26	0.866
2009	243	4.05	0.983
2010	257	3.92	1.01
2011	263	0.50	0.132

*Emission factors have been assumed

4.3.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty for activity data is ± 2 % and the uncertainty of the emission factor for N₂O is ± 5 %. The time-series is consistent. The fluctuations in the calculated total EF for N₂O 1994 – 2000 (Table 4-18) are mainly due to fluctuations in one of the facilities. The IEFs are within the IPCC default interval (2-19 kg N₂O/Mg). Activity data and reported emissions have been acquired from reporting in e.g. environmental reports from the facility, but since the facility has shut down, it is no longer possible to check previously reported estimates. Beside emissions of N₂O also emissions of NO_x are reported.

The lower level of N₂O emissions from 2001 and onward compared to earlier years is a result of one facility being shut down in late 2000 and a second one during 2001. Emissions for all years, except 1991 - 1993, are as reported from the facilities. The higher level of NO_x emissions in year 2004 is a result of a long lasting leakage of NO_x from one of the production units at the active facility. During 2007 catalytic abatement was installed at one of the production units at the active facility and as a result the emissions of N₂O and NO_x were reduced compared to previous years. The used abatement system

is described in the BREF document for large volume inorganic chemicals¹⁰⁷. During 2009 the production of nitric acid was lower compared to previous years and also lower than later years. The higher N₂O emission factor in 2009 is due to that the N₂O reduction catalysts were not used during 2009. This was because 2009 was set as base year in a joint implementation project with the aim to reduce N₂O emissions. For some months in 2010 N₂O-reducing catalysts were used again, now in both production units at the facility. In one of the production units the catalyst was used from March and in the other unit from December. The fact that the catalysts were not used not during all months of the year is the reason for the higher emission factor in 2010 compared to 2007 and 2008. In 2011 the catalysts in both production units were used the whole year with a significant decrease of N₂O emissions as result.

4.3.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.3.2.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.3.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.3.3 Carbide production (CRF 2.B.4)

4.3.3.1 SOURCE CATEGORY DESCRIPTION

Silicium carbide production does not occur in Sweden but calcium carbide is produced at one facility. All process-related CO₂ emissions from the industry are included in the code 2.B.4.2. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-19.

Table 4-19. Summary of source category description, CRF 2.B.4.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.B.4	CO ₂				D	PS	Yes

D Default. PS Plant-specific.

¹⁰⁷ European Commission, 2007

4.3.3.2 METHODOLOGICAL ISSUES

To cover all sources of CO₂ from the production of calcium carbide, estimates of emissions from the production of quick lime, from the reduction of quick lime to calcium carbide and CO₂ from use of calcium carbide have been made. In the tables and text below the estimated CO₂ emissions to be reported in CRF 2B4.2 are presented.

4.3.3.2.1 CO₂ emissions from quick lime production

In order to estimate the CO₂ emissions from the first step of the production of calcium carbide, the amount of limestone used for quick lime production is used as activity data together with the default emission factor from Revised 1996 IPCC Guidelines, 0.44 Mg CO₂/Mg limestone used. During the time period 1990 – 2011 the facility has for some years produced all quick lime that has been used in the production while in other years large amounts of quick lime has been bought from other producers of quick lime. Thus, the CO₂ emission varies between years as can be seen in table 4-20.

Table 4-20. Limestone used for quick lime production and associated CO₂ emissions to be reported in CRF 2.B.4.2.

Year	Limestone used for quick lime production, Gg	CO ₂ from quick lime production, Gg
1990	96	42
1995	79	35
2000	64	28
2005	75	33
2006	23	10
2007	69	30
2008	61	27
2009	9	4
2010	51	22
2011	51	22

4.3.3.2.2 CO₂ emissions from calcium carbide production

Calcium carbide is produced in an electric arc furnace at high temperature, 2,000 – 3,000 °C. Quick lime, CaO, is reduced with coke and forms CaC₂. In this process an energy rich gas is produced as a by-product. This gas is used as fuel within the facility and to some extent in other nearby plants and thus only a minor part of the gas is flared. To calculate the CO₂ emissions from the reduction of quick lime to calcium carbide, data on produced amounts of calcium carbide, share of gas flared and default emission factor in IPCC Guidelines are used. Since there only is one producer of calcium carbide in Sweden the produced amounts are reported as confidential, C.

Table 4-21. Share of flared carbide oven gas and associated CO₂ emissions to be reported in CRF 2.B.4.2.

Year	Carbide oven running time/ Flaring time, %	CO ₂ from the reduction of CaO to CaC ₂ , Gg
1990	6%	4
1995	7%	4
2000	14%	6
2005	15%	7
2006	13%	6
2007	10%	4
2008	8%	4
2009	10%	3
2010	9%	3
2011	11%	4

4.3.3.2.3 CO₂ emissions from use of calcium carbide

In Revised 1996 IPCC Guidelines it is stated that in addition to reporting CO₂ emissions from calcium carbide production, also CO₂ originating from the use of calcium carbide has to be reported. To be able to estimate the CO₂ emission from use of calcium carbide only the amount of calcium carbide for acetylene production and the use within the country has to be taken into account. Information from the calcium carbide producer in Sweden indicates that one third of the calcium carbide is used for acetylene production. Assuming that imported and exported amounts of acetylene have the same utilisation it is possible to reasonably well estimate the CO₂ emissions originating from acetylene use. Yearly statistics on imported and exported amounts from 1998 and onwards are available from Statistics Sweden¹⁰⁸. Amounts used for acetylene production for earlier years are estimated. The default emission factor presented in the Revised 1996 IPCC Guidelines, 1.1 Mg CO₂/Mg calcium carbide use, has been used for the estimations.

Table 4-22. Amount of calcium carbide used for acetylene production, and CO₂ emissions from acetylene use reported in CRF 2.B.4.2.

Year	Amount of calcium carbide for acetylene production, Gg	CO ₂ from use of acetylene, Gg
1990	7	8
1995	6	7
2000	6	7
2005	9	10
2006	8	8
2007	8	9
2008	9	9
2009	4	5
2010	5	6
2011	5	6

¹⁰⁸ www.scb.se

4.3.3.2.4 Time series reported in CRF 2.B.4. 2

In Table 4-23, the total CO₂ emission for some years in the time series reported in submission 2013 is presented. Since there is only one producer of calcium carbide in Sweden production statistics are reported as confidential, C. The total reported CO₂ emissions in CRF 2.B.4.2 are based on:

- produced amounts of quick lime and emission factors from Revised 1996 IPCC Guidelines
- produced amounts of calcium carbide, share of gas flared and the default emission factor according to the Revised 1996 IPCC Guidelines
- amount of calcium carbide used for acetylene production within the country and the default emission factor presented in the Revised 1996 IPCC Guidelines.

Table 4-23. Time series reported in CRF 2.B.4 2.

Year	Produced calcium carbide, Gg	CO ₂ emissions from production and use of calcium carbide, Gg
1990	C	54.2
1995	C	45.5
2000	C	40.8
2005	C	49.6
2006	C	24.2
2007	C	43.4
2008	C	40.2
2009	C	11.7
2010	C	31.6
2011	C	32.5

4.3.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As can be seen in Table 4.23, CO₂ emissions for 2006 and 2009 are much lower compared to surrounding years. For 2006 a minor part of the quick lime needed for production of calcium carbide were produced at the facility, the remaining part was bought from external producers. The sharp decrease of the CO₂ emission in 2009 is due to that the lime kiln was only operating during the last quarter of the year. Consequently, most of the quick lime used for calcium carbide production in 2009 was bought from external lime producers.

4.3.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.3.3.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.3.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.3.4 Other (CRF 2.B.5)

4.3.4.1 SOURCE CATEGORY DESCRIPTION

This sub-category includes various chemical industries, such as sulphuric acid production, the pharmaceutical industry, production of base chemicals for plastic industry, various organic and inorganic chemical productions and other non-specified chemical production, which are not covered elsewhere. Approximately 70 larger industrial facilities are included in the emission estimates. Emissions of CO₂, CH₄, N₂O, NO_x, CO, NMVOC and SO₂ are reported in this sub-category. It is possible though that some emissions of NMVOC reported in CRF 2.B.5 should be reported in CRF 3C (e.g. pharmaceutical industries), but as it has been difficult to make the distinction clear between process emissions and solvent use, all NMVOC emissions from these facilities have been included in CRF 2B5.

Emission time-series for GHG are relatively stable. There is a slight drop in emissions of GHG in 2009 compared to 2008 e.g. due to lower production of carbon black. In addition, CH₄-emissions decreased in 1999 due to a much lower production at one facility and N₂O-emissions increased in 1999 due to the fact that one facility within "Pharmaceutical industry" reported higher emissions that year.

The SO₂ emissions reported in 2B5 decreased dramatically in 2004 in comparison to earlier years. This is due to that in December 2004 one facility for production of viscose staple fibre was shut down. The yearly SO₂ emissions from this facility represented between 8 and 20 % of the totally reported SO₂ emission in CRF 2 – Industrial Processes, 1990 - 2003.

CO-emission from "Other inorganic chemical production" increased from below 200 Mg in 2005 to 500 Mg in 2006. This increase is due to unusually high CO emission in 2006 from one facility producing PVC. In 2007 the CO-emissions were very low from one facility producing PVC.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-24.

Table 4-24. Summary of source category description, CRF 2.B.5.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.B.5	CO ₂				CS	PS	Yes
	CH ₄				CS, D	PS, D	Yes
	N ₂ O				CS	PS	Yes

D Default. CS Country Specific. PS Plant-specific.

4.3.4.2 METHODOLOGICAL ISSUES

The primary information on emissions of CO₂, CH₄, N₂O, NO_x, CO, NMVOC and SO₂ is as reported by the companies in their environmental reports. A total of approximately 70 facilities are included. In the IPCC Guidelines, methods for estimating CH₄ emissions for several chemical products are presented and consequently the CRF Reporter is divided on those products (2.B.5.1-5). Since several plants in Sweden produce several chemicals products each but report emissions aggregated by plant, it is not possible to report emissions in accordance with the suggested split in the CRF Reporter. In Sweden, since submission 2006 the emissions are thus presented allocated to six separate branch categories: sulphuric acid production, pharmaceutical industry, production of base chemicals for plastic industry, organic chemical production, inorganic chemical production and other non-specified chemical production.

In Sweden there is one company producing carbon black. CH₄ emissions are included from 1990 and onwards based on production data from the company's environmental reports and IPCC Guidelines default EF (11 g CH₄/kg production). Due to data confidentiality, emissions are included under 2.B.5 (Other inorganic chemical production).

4.3.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Based on expert judgements, the uncertainties of collected emissions of CO₂, CH₄ and N₂O are as follows: ±50 %, ±100 % and ±125 %, respectively.

The time-series for GHG have been reviewed and are considered to be consistent.

4.3.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Emissions reported in the plant-specific environmental reports are carefully studied annually to retrieve the most appropriate data for the GHG inventory.

In addition, emissions in this sub-category were reviewed as part of a quality control SMED project, financed by the Swedish EPA, during 2010. The project aimed at increasing the quality and reducing the uncertainties in the most important air emissions substances from chemicals industries in Sweden¹⁰⁹. Emissions reported in the environmental reports were compared to plant-specific data in the GHG inventory, significant discrepancies were investigated, and recommendations were provided on feasible improvements for submission 2011 as well as recommendations on further investigations¹¹⁰.

Overall, the QC-project showed that total reported GHG emissions from the chemical industries in the Swedish inventory are in coherence with the plant emission data.

¹⁰⁹ Gustafsson, T., Nyström, A-K., Gerner, A. Riktad kvalitetskontrollstudie av utsläpp från kemiindustrin i Sveriges internationella rapporter. SMED report 2010.

¹¹⁰ Most recommendations on further investigations refer to the energy sector

4.3.4.5 SOURCE-SPECIFIC RECALCULATIONS

- Calculation errors regarding the emission of NO_x for the years 2009 and 2010 have been corrected.
- An error in the calculation sheet of 2.B.5, Base chemicals for plastic industry, regarding year 2006 was found and corrected for. This correction resulted in a minor increase in emissions of NO_x and NMVOC for year 2006.
- Subsector 2B5/Other organic chemical prod: Emissions of NMVOC for year 2010 has been updated.

4.3.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.4 Metal production (CRF 2.C)

All sub-categories are covered in the estimates, i.e. iron and steel production (2.C.1), ferroalloy production (2.C.2), aluminium production (2.C.3), SF₆ used in magnesium foundries (2.C.4) and other (2.C.5), which consists of estimates for one large non-ferrous smelter plant and one metal recycling plant.

4.4.1 Iron and steel production (CRF 2.C.1)

4.4.1.1 SOURCE CATEGORY DESCRIPTION

In Sweden, there are three primary iron and steel facilities and about ten secondary steel plants equipped with electric arc furnaces. In total, there are approximately 20 different facilities included in the different estimates. Processes occurring besides the primary processes and secondary steel production are rolling mills, pickling and other steel-related processes. From submission 2009 and onwards, emissions from two major iron ore mines and three facilities producing pellets in Sweden are reported in 2C1.3 (reallocated from previous reporting in 2.A.7). Emissions from a sinter producing facility are also included until 1995, when the production closed down.

Process emissions arising from reducing agents in the primary steel works and secondary iron and steel works are reported in CRF 2.C.1. As the plants also generate emissions from fuel combustion (CRF 1.A.1c and CRF 1.A.2.a) and fugitive emissions (CRF 1.B.1.c), the text in this section is closely connected to the text in the corresponding section in the energy chapter.

In the Swedish inventory, emissions from primary iron and steel production and secondary steel production are reported separately and fed into the CRF Reporter under 2.C.1.2 Pig iron and 2.C.1.1 Steel, respectively. This enables process emissions from the two integrated iron and steel production plants in Sweden to be reported together (2.C.1.2 Pig iron), and thus not introducing further sources of uncertainty due to additional data handling.

The GHG emission trend 1990-2008 is rather stable with some minor inter-annual variations. However, the economic recession in 2009 had a great effect on the production volumes of iron and steel in Sweden and thus the emissions 2009 are significantly reduced. In 2010 the emissions are back at the same level as before 2009.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-25.

Table 4-25. Summary of source category description, CRF 2.C.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.C.1	CO ₂	X	X		CS, T2	PS	Yes
	CH ₄				CS	PS	No, see Annex 5
	N ₂ O				NA	NA	NA

CS Country Specific. T2 Tier 2. PS Plant-specific.

4.4.1.1.1 Secondary steel production, CRF 2.C.1.1

The reported CO₂ emissions include emissions from reducing agents such as coke, coal and electrodes in electric arc furnaces in secondary steel plants. These emissions are not primarily a result of combustion, but are necessary for the process and should hence be reported in CRF 2.C.1.1. Reported CO₂ emissions also include emissions from the use of limestone and dolomite in secondary steel industry.

In submissions prior to submission 2010, the reported CO₂ emissions in CRF 2.C.1.1 included data from nine plants in 1990-2003 and eight plants from 2004, since one plant was shut down in 2004. From submission 2010 another two secondary steel industries are included in the reported CO₂ time series. Also another four plants with process related NO_x and/or NMVOC emissions are included in this sector. These plants do not produce steel, and hence do not emit CO₂.

Production and consequently emissions have increased slowly since 1990 due to higher demand of these products. The high production and emissions level in 1990 compared to 1991 is explained by the fact that one plant closed its production in 1991.

4.4.1.1.2 Primary iron and steel production, CRF 2.C.1.2

In Sweden there are three producers of primary iron and steel, i.e. the basis of their production is iron ore pellets. Two plants produce pig iron and steel as part of their integrated coke ovens, blast furnaces and steel converters. The primary purpose of the use of coal and coke in the blast furnace is to secure oxidation and act as reducing agents, and the associated emissions are thus to be reported as industrial processes from iron and steel production in CRF 2.C.1, according to the IPCC Guidelines and Good Practice Guidance. The third plant produces iron sponge and iron powder.

4.4.1.1.3 Iron ore mining, dressing, sintering and iron ore pellets production, CRF 2.C.1.3

Emissions of CO₂ from the use of limestone and dolomite within the production of ore based iron pellets are reported in CRF 2.A.3. Estimated emissions from combustion of fuels are included in the Energy sector (CRF 1).

Emissions considered in CRF 2.C.1.3 are CO₂ from the use of bentonite and organic binder, SO₂ from the sulphur content in the ore and NO_x emitted as a result of the use of explosives. The use of mining explosives also causes emissions of carbon monoxide, CO¹¹¹.

4.4.1.1.4 CO₂ emissions reported in Coke, CRF 2.C.1.4

Emissions of CO₂ from the production of coke are reported in CRF 1.A.1.c and emissions of CO₂ from the use of coke in blast furnaces are reported in CRF 2.C.1.2 in line with the IPCC Guidelines.

4.4.1.1.5 CO₂ emissions reported in Other, CRF 2.C.1.5

No emissions of CO₂ reported in this sector.

4.4.1.2 METHODOLOGICAL ISSUES

4.4.1.2.1 Secondary steel production, CRF 2.C.1.1

In most cases, data from the Swedish enquiry for the Swedish national allocation plan (NAP) for the EU ETS could be used for the years 1998-2002. Data for 1990-1997 and 2003-2004 has been collected directly from the plants. From 2005, the equivalent data are acquired from the ETS, from the facilities environmental reports and through contacts with the companies.

Data in the ETS includes information concerning carbon bound in products, slag, etc, but also other sources for process related CO₂ emissions. Prior to submission 2010, these other emissions were not included for all facilities. Estimates of these missing CO₂ emissions were performed using ETS data for 2005 – 2008 and production data for years before 2005. All CO₂ emissions presented for the facilities in ETS 2005 – 2011 are included in 2C1.1 in submission 2013.

Reported CO₂ emissions until year 2008 are for all facilities, except the one which closed down in 2004, based on data in the ETS. Reported CO₂ emissions can therefore be classified to follow the Good Practice Guidance method Tier 2 since, according to the ETS guidelines; reported emissions shall be based on all carbon input to and carbon output from the process. For the years from 2009 background data needed for estimation of process-related CO₂ emissions for one facility, earlier included in the ETS, was collected from the facility's environmental report since this facility is not included in ETS. Before the facility shut down in 2004, plant specific methods were applied. For non-CO₂ emissions, the companies' environmental reports are the main source of information. NO_x, NMVOC and SO₂ emissions emitted from elec-

¹¹¹ Wieland, M.S. 2004.

tric arc furnaces are reported in 2.C.1.1. NO_x emissions may also arise from pickling and NMVOC emissions from rolling mills. These sources are also included in the estimates.

4.4.1.2.2 *Primary iron and steel production, CRF 2.C.1.2*

Production of iron powder

The emissions of CO₂ are calculated using the Good Practice Guidance method Tier 2. Plant-specific data on emissions from carbon-containing input materials such as coke and anthracite and also specific carbon-contents of output iron and by-products are used for all years. From 2005, ETS data is used and 1990-2004, information has been acquired from the plant. The emissions are verified using national statistics from Statistics Sweden on amounts of coke, anthracite and out-put material. CO₂ emissions from natural gas used for production of reduction gas used in the process are considered to be process-related and thus reported in 2.C.1.2. The remaining amounts of natural gas used by the facility are considered as energy-related and the corresponding emissions are reported in the Energy sector (CRF 1.A.2.a). To be consistent with calculations of emissions from production of pig iron, limestone used in the production is included in the emissions from the production of iron powder in CRF 2.C.1.2. Activity data reported is produced amount of direct-reduced iron (iron sponge).

Production of primary pig iron and steel

As a response to recommendations from UNFCCC expert review teams, since submission 2010, Sweden uses the recommended Tier 2 method according to the IPCC Guidelines, to base the calculations of CO₂ emissions on carbon mass-balances in order to reduce the risk of double counting or omitting CO₂ emissions.

The carbon contents of external input materials such as coking coal, coke, injection coal, limestone, etc., are balanced against final output materials; coke¹¹², pig iron¹¹², steel, tar, sludge, slag, etc. The remaining carbon contents are accounted for as CO₂ emissions:

$$CO_2 \text{ emissions}_{TotalCRF1and2} = \left[\sum_i (MI_i * C_i) - \sum_p (MO_p * C_p) \right] * 44/12$$

where,

MI_i = External carbon material input *i* fed into any part of the integrated processes (t).

MO_p = Final carbon material output *p* (t).

C_x = Carbon content of material input or output *x* (t C/t material *x*).

Figure 4-5 gives an overview of the input and output materials, the carbon flows between the different processes (plant stations), and the CO₂-emitting sources.

¹¹² If put in stock or sold externally

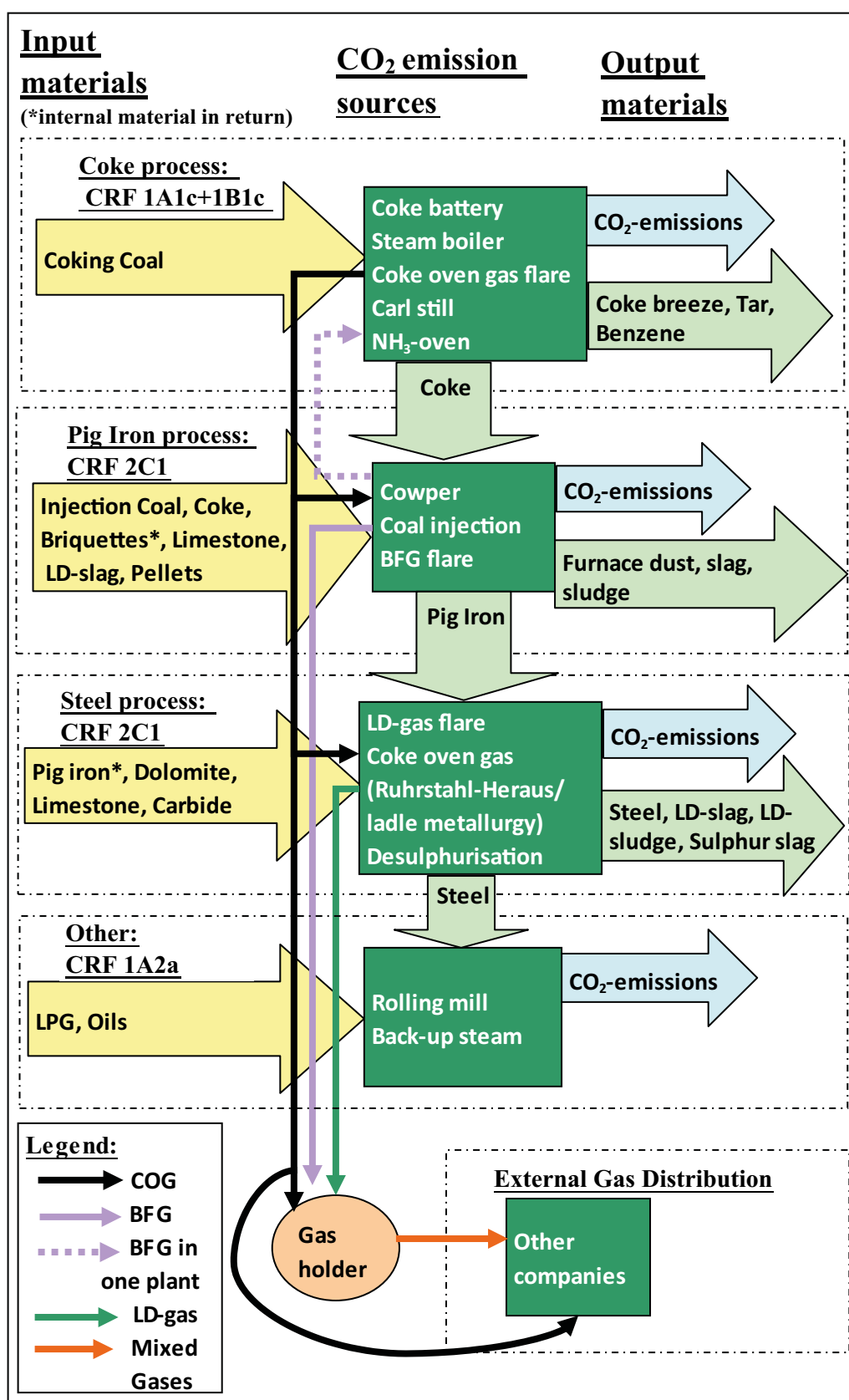


Figure 4-5. Carbon flow chart of integrated primary iron and steel plants in Sweden

In the coke ovens (battery), coking coal is turned into coke through dry distillation. During the process, coke oven gas (COG) and by-products are formed. The coke oven gas is purified through several procedures and used as fuel in other plant stations, but smaller amounts are also flared. Produced amounts of coke are fed into the blast furnace together with injection coal to act as reduction agent when pig iron is produced from iron ore pellets. Limestone is added to extract slag and other by-products from the pig iron. Besides pig iron and by-products, blast furnace gas (BFG) is produced in the process. The main use for the blast furnace gas is to heat up the cowpers (and in one plant used in the coke oven), but some excess gas is released through flaring.

In the steelworks, pig iron is transformed into various qualities of steel depending on the demand. Dolomite, pig iron, carbide, etc., are added depending on the different metallurgic processes. LD-gas is produced in the steel converter and used as fuel or flared. Some steel is treated in the rolling mills where LPG and different oils are used as fuel.

Considerable amounts of energy gases (coke oven gas, blast furnace gas and LD-gas) from the different processes are collected in a gas holder and sold to external consumers (mainly in CRF 1.A.1.a electricity and heat production). These amounts of gases and their associated emissions are allocated to the source category where they are consumed and thus not accounted for in the iron and steel production. This is not in accordance with the 1996 Guidelines as they fall under the category “Auto-producers¹¹³”, but in line with the 2006 IPCC Guidelines¹¹⁴ where allocation of emissions from delivered gases is described. Sweden has chosen to follow the 2006 IPCC Guidelines in this case as they are more in line with the emission reporting for the annual environmental reports and the EU ETS reporting. Detailed carbon mass balances are compiled for plants included in the reporting according to EU ETS, but due to confidentiality reasons the mass balances cannot be included in the NIR. However, they can be delivered to the UNFCCC Expert Review Team upon request.

During the whole process from raw material to final product, emissions of CO₂ are released. The allocation of both plants total CO₂ emissions on plant stations and consequently CRF sub-sector is based on detailed mass-balances (Table 4-26).

¹¹³ See IPCC Guidelines: Reporting instructions 1.3

¹¹⁴ See 2006 IPCC Guidelines: Volume 3: Industrial Processes and Product Use, Box 1.1 (page 1.8)

Table 4-26. CO₂ emission allocation 2011 in integrated primary iron and steel production (excluding external gas distribution).

CRF	Plant station	CO ₂ emissions 2011 (Gg)
1.A.1.c	Coke Oven	345
1.A.2.a	Combustion in Rolling Mills + Power and Heat Production	726
1.B.1.c	Flare in Coke Oven (COG)	6
2.C.1.2	Blast Furnace + Steelworks (including Flaring of BFG and LD-gas)	2,436
Total		3,512

According to the IPCC Guidelines, emissions of CO₂ from the use of limestone should be reported separately as process emissions from limestone and dolomite use in CRF 2.A.3. Since the Centralized review from submission 2004 the ERT has repeatedly recommended Sweden to follow the guidelines. As the CO₂ emissions from limestone and dolomite are small (<1 per cent of the plants total CO₂ emissions) it is not considered to be good practice to spend resources obtaining underlying data to separate these emissions. Hence Sweden choose to include these CO₂ emissions in CRF 2.C.1.2.

Activity data (amount of pig iron produced) on integrated pig iron and steel production along with CO₂ emissions and consumed amounts of energy gases (coke oven gas, blast furnace gas and LD-gas) and other fuels, are reported by the plants in the environmental reports since 2003. Mass-carbon balances and associated CO₂ emissions are also reported to the EU ETS since 2005. For some years, CO₂ emissions to the EU ETS did not include all plant stations (rolling mills), and additional information from the plants was obtained in order to ensure that no omissions occurred. Since 2008 annual CO₂ emissions reported by the plants in their environmental reports are equal to those reported to the EU ETS. For 2003 onwards, information on activity data and emissions for all plants (CRF 1.A.1.c, 1.A.2.a, 1.B.1.c and 2.C.1.2) are taken from the environmental reports. Amounts of pig iron produced 1990-2002 were obtained directly from both plants, together with total CO₂ emissions 1990-2002 for one of the plants. For the other plant, CO₂ emissions 1990-2002 are calculated using its pig iron production 1990-2002 and an average CO₂ IEF 2003-2007. Allocation of CO₂ emissions on different sub-categories (CRF 1.A.1.c, 1.A.2.a, 1.B.1.c and 2.C.1.2) are based on the plant specific average distributions 2003-2007.

Consumed amounts of different energy gases and other fuels 1990-2002 are derived by applying the Good Practice Guidance surrogate method using the average values 2003-2007 and the CO₂ emissions as the surrogate parameter. Activity data reported in CRF Reporter in CRF 2C1.2 is produced amount of primary pig iron.

Emissions of CH₄, N₂O, NMVOC and CO are not reported in the plants' environmental reports. In the Swedish inventory these emissions are instead estimated from consumed amounts (including flared amounts) of energy gases multiplied by country-specific emission factors (see Annex 2). Emissions of

CH₄, NMVOC and CO from coke oven gas, blast furnace gas and LD-gas in the blast furnace and steel converter are allocated to CRF 2.C.1.2, whereas emissions of N₂O are assumed to be not applicable (NA) in this sub-category, in accordance with the IPCC Guidelines. Emissions of NO_x and SO₂ are based on detailed plant information from the environmental reports.

4.4.1.2.3 *Iron ore mining, dressing, sintering and iron ore pellets production, CRF 2.C.1.3*

Data on production statistics as well as on SO₂ emissions have been supplied by the facilities for the entire time-series. Amounts of bentonite and organic binder used for the production of iron ore pellets and the corresponding CO₂ emissions are for later years collected from the EU ETS. For earlier years the amounts of bentonite and organic binder were provided by the company and EFs for bentonite and organic binder from the EU ETS were used for the calculations. No data concerning the CO emissions is available and the time series is thus reported NE.

4.4.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The largest implication on the national total uncertainties from this category stems from uncertainties in CO₂ emissions in primary iron and steel production (CRF 2.C.1.2); based on expert judgement by SMED expertise the estimated uncertainty is $\pm 5\%$. It should be noted however, that total emissions of CO₂ from iron and steel production, including energy related emissions, are likely to deem lower uncertainty estimates.

There is an obvious decrease in CO₂ IEFs since 1990 (Table 4-27) for primary pig iron and steel production, from 0.80 Gg CO₂/kton iron in 1990 to 0.55 Gg CO₂/kton pig iron in 2009. This is due to the undertaking of several energy efficiency measures, e.g. increased temperature in the blast furnaces and increased recycling of energy gases and by-products¹¹⁵, leading to decoupling between CO₂ emissions and primary pig iron production in Sweden.

Table 4-27 shows a significant increase in the IEF for year 2011. This is due to the fact that the reported CO₂ emissions are overestimated for one plant. During 2011 one of the two blast furnaces at the plant was out of operation from July until December and consequently the production of pig iron decreased compared to previous year. At the same time the production rate at the coke plant were kept under normal conditions. This resulted in an increased intermediate stock of coke at the plant. After consulting the operator it was concluded that the operator does not take into account any intermediate stock change of produced coke in the carbon mass balance used when calculating the CO₂ emissions. The same method has been used for all years since emission year 2005, i.e. the first reporting year to EU ETS. The exclusion of the change in storage of coke in the carbon mass balance is more pro-

¹¹⁵ ENET-Steel, 2007.

nounced for years when for example the operation of the blast furnaces has been restricted. During 2012 the operator applied to the county administrative board to change their monitoring methodology for CO₂. However, the method change will not apply until emission year 2012. In late 2012 the Swedish EPA will initiate a development project in order to correct the overestimated time series for CO₂ for the plant, and consequently the time series will be revised in submission 2014.

The CO₂ IEF is overall significantly lower than the Tier 1 default emission factor (1.35 Gg CO₂/kton pig iron produced) presented in 2006 IPCC Guidelines Table 4.1. This is partly due to that a large share of the energy gases produced at one of the plants is distributed to other companies (as described above) and thus not accounted for in iron and steel production. Adding CO₂ from all the external gas distribution (though some gas is COG) would lead to an IEF in 2008 of 1.21 Gg CO₂/kton pig iron produced and thus considered to be reasonable with regard to the IPCC Guidelines default.

Table 4-27. CO₂ implied emission factors (IEF) for primary pig iron production

Year	CO ₂ IEF (Gg CO ₂ /kt primary pig iron produced)
1990	0.80
1995	0.78
2000	0.70
2005	0.58
2006	0.57
2007	0.56
2008	0.58
2009	0.55
2010	0.68
2011	0.75

4.4.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All plants in this category report their emissions in environmental reports. For plants included in the EU ETS the report data is scrutinized and compared to EU ETS data. EU ETS data is applied wherever it is judged to be appropriate in line with the Good Practice Guidance. Detailed carbon mass balances are compiled for plants included in the reporting according to EU ETS, but due to confidentiality reasons the mass balances cannot be included in the NIR. However, they can be provided to the UNFCCC Expert Review Team upon request. More information on QC activities related to EU ETS is included in Annex 8.1.

For primary iron and steel production, activity data is compared to production statistics from the Swedish Steel Producers' Association and only minor differences are detected for the time-series.

4.4.1.5 SOURCE-SPECIFIC RECALCULATIONS

2.C.1.1

Reported SO₂ emissions were corrected for 2009 and 2010.

2.C.1.2

No recalculations have been performed in CRF 2.C.1.2 in submission 2013.

2.C.1.3

No recalculations have been performed in CRF 2.C.1.3 in submission 2013.

4.4.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.4.2 Ferroalloy production (CRF 2.C.2)

4.4.2.1 SOURCE CATEGORY DESCRIPTION

Ferroalloy production is reported for only one facility in Sweden. There is also ferroalloy production at one more plant, but since the main production at this facility is iron and steel, the emissions are reported in CRF 2.C.1- Iron and steel production. The production of iron silicide has decreased sharply since 2005, and since 2008 there is no production at all. This leads to ending of the emissions of CH₄ from 2008. Production of ferrosilicon leads to larger emissions of SO₂ compared to production of ferrochromium. From 2005 the production of ferrosilicon has been much reduced and from 2008 no ferrosilicon is produced. This leads to a distinct decrease in SO₂ emissions in later years. The economic recession in 2009 had a great effect on the production volumes of ferroalloys in Sweden and thus the emissions 2009 are significantly reduced.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-28.

Table 4-28. Summary of source category description, CRF 2.C.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.C.2	CO ₂				T2	PS	Yes
	CH ₄				D	D	Yes
	N ₂ O				NA	NA	NA

D Default. T2 Tier 2. PS Plant-specific.

4.4.2.2 METHODOLOGICAL ISSUES

CO₂ emissions within the production of ferroalloys are plant specific (in line with Tier 2). The CO₂ emissions reported by the plant are calculated from consumed amount of reducing agents (Tier 1a¹¹⁶), i.e. electrodes and coke (and in 2003 coal) and their specific carbon contents. Input data is also the amount of carbon bound in produced ferroalloys. The common distribution of carbon in the incoming and outgoing materials is:

Coke	+	Electrodes	→	Ferroalloys	+	Emissions	+	Particles
95%	+	5%	→	10%	+	89.5%	+	0.5%

To verify the emissions reported by the plant, emissions are calculated from activity data on coal, coke, electrodes and the amount of carbon in produced ferroalloys and:

- Emission factors and thermal values used for stationary combustion for coke and coal and information from the company that the electrodes contain 90 % carbon.
- IPCC default factors for coal, coke and electrodes¹¹⁷.

The used formula is:

$$CO_2(Mg) = Coke(Mg) \times EF \times Thermal\ value + Coal(Mg) \times EF \times Thermal\ value + Electrode(Mg) \times C\text{-}content \times \frac{44}{12} - CO_2\ in\ produced\ ferroalloys(Mg, plant\ data)$$

where 44/12 are the molecule weights of CO₂ and carbon. As can be seen in Table 4-29, there are differences in the plant specific data and emissions based on Swedish default EF and emissions estimated with IPCC Guidelines default values. The differences are due to the fact that - according to the company - the carbon content of the coke may vary from one year to another.

The total amount of carbon in the produced ferroalloys is presented in Table 4-30, and is calculated based on the carbon content in coke, coal, electrodes and dust by the company. The amount of carbon in the produced ferroalloys varies between 0.1 % and 7 %. This carbon is reported under CRF 1.AD.10 - coke and coal. CH₄ emissions from production of FeSi alloys are reported from submission 2010 and calculated based on FeSi alloy production (Tier 2¹¹⁸)

Data on non-CO₂ emissions has been obtained directly from the company for the whole time series. The reported emissions include NO_x and SO₂ from the process.

¹¹⁶ <http://www.ipcc-nggip.iges.or.jp/public/gl/guidelin/ch2wb2.pdf>

¹¹⁷ IPCC. Revised 1996 Guidelines for National Greenhouse Gas Inventories: Reference Manual, Table 2.12.

¹¹⁸ 2006 IPCC Guidelines for National Greenhouse Gas Inventories, Table 4.8

Table 4-29. Total emissions of CO₂ based on plant specific data (reported in the CRF), data based on Swedish EF and thermal values, and based on IPCC Guidelines default values.

Year	Plant specific data, Gg CO ₂	Swedish values, Gg CO ₂	IPCC default values, Gg CO ₂
1990	243	244	263
1995	265	274	295
2000	240	266	287
2005	225	214	231
2006	220	209	225
2007	220	188	203
2008	194	164	177
2009	48	48	52
2010	107	96	104
2011	117	122	132

Table 4-30. Total amount of carbon bound in produced ferroalloys.

Year	Carbon in ferroalloys, Gg
1990	8.4
1995	8.7
2000	9.5
2005	8.0
2006	8.3
2007	8.4
2008	7.4
2009	1.8
2010	4.0
2011	4.7

4.4.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties in this category have little impact on the estimated national total emission uncertainty. Emission uncertainties of CO₂ are judged by SMED expertise to be low at $\pm 5\%$ as plant-specific values and Swedish default values give similar results.

Time-series are considered to be consistent.

4.4.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

As presented in Table 4-29 verification of CO₂ emissions reported by the plant is obtained as calculated Swedish default values give similar results.

4.4.2.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.4.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.4.3 Aluminium production (CRF 2.C.3)

4.4.3.1 SOURCE CATEGORY DESCRIPTION

There is one facility that produces primary aluminium in Sweden. The facility consists of two plants. One of the potlines (plant 1) includes 56 closed prebake cells (CWPB), each of 150 kA. The other plant (plant 2) consisted of 262 cells and, until the beginning of 2008, operated three prebake cells and 259 open cells with Söderberg anodes (VSS). The Söderberg anodes were produced in an electrode pulp factory at the facility.

In 2008 a project was started to convert the Söderberg ovens to ovens with prebake cells. All pot-lines operating the Söderberg technology were shut-down by December 2008. The anode effect (AE min/oven day) were for the Prebake process higher in 2008 compared to 2007 due to start-up problems for the new Prebake cells.

By the end of December 2009, 120 of a total of 262 cells in plant 2 had been converted to the prebake technology and the conversion to prebake cells continued under 2010. In the beginning of December 2010 242 prebake cells in plant 2 were in operation. At the end of December 2010 a power outage lead to major disturbances in plant 2 leading to both increased emissions and major production problems. On January 7, 120 prebake cells were shut down as a direct result of the power outage. At the end of June 2011 all prebake cells in plant 2 were restarted and in operation.

The shutdown of Söderberg ovens explains the very large decline in PFC emissions in 2009 (-85% compared to 2008) (Figure 4-6). Also the reported CO₂ has declined in 2009 relative to previous years. The relatively stable implied emission factor (Table 4-32) provides the explanation that the reduced CO₂ emissions are due to reduced aluminium production in 2008 and 2009. The cold winter in 2010 resulted in high power input to the anodes, thus leading to high emissions of PFCs. There were also problems with power outages which affected the production and led to increased number of AE minutes. During the start-up period in 2011, emissions to air increased but later in 2011 the emissions decreased to expected levels.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-31.

Table 4-31. Summary of source category description, CRF 2.C.3.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.C.3	CO ₂				T2	PS	Yes
	CH ₄				NA	NA	No, see Annex 5
	PFCs		X		T2	D	Yes

D Default. T2 Tier 2. PS Plant-specific.

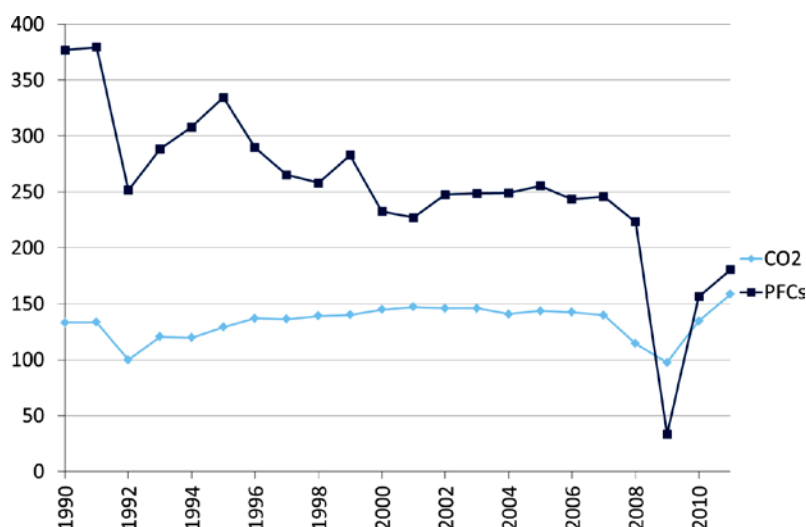


Figure 4-6. Time series for CO₂ and PCF emissions from aluminium production, CRF 2C3.

4.4.3.2 METHODOLOGICAL ISSUES

Primary aluminium is in Sweden produced in one facility, where the prebaked process is used. The time series of emissions compiled for primary aluminium production include emissions of CO₂, PFCs, NO_x, CO, NMVOC and SO₂. Reported production statistics and emissions data are based on information in the environmental reports or received directly from the company.

Reported emissions of NO_x are calculated from production statistics using emission factors defined by Swedish EPA¹¹⁹. NMVOC emissions are calculated from reported emissions of tar, assuming that 70 % of the tar is emitted as NMVOC¹¹⁹. CO emissions were for the first time reported in submission 2008 and are for 2002 - 2011 as reported in the company's environmental reports. For the period 1990 – 2001, the CO emissions are calculated based on production statistics and emission factors provided by the company as also for the SO₂ emissions during 1990 - 2005. For later years SO₂ emissions data are based on environmental reports published by the company.

Emission data for CO₂ from the production of primary aluminium 2002 - 2011 are derived through measurements and reported directly by the plants, whereas the emissions for 1990-2001 are calculated based on the mass of coal elements (anodes) such as electrodes, coke etc. and the amount of carbon that is bound in soot. The formula used for CO₂ (Mg) for 1990-2001 is:

$$\text{Mass anodes (100\% C)} \times \frac{44}{12} \times (1 - 0.257^*)$$

* Mass CO₂ bound in soot and rest anodes in 2002

¹¹⁹ Ahmadzai, H. Swedish EPA. Personal communication. 2000.

The value for carbon bound in soot and rest anodes (0.257) is based on the reported value for 2002. For subsequent years the amounts bound in soot and rest anodes vary between 0.181 and 0.298.

For the years from 2002 and onwards the emissions reported by the plant have been verified by collecting data on the amount of coal elements used and by calculating the emissions based on the equation above. The results are very comparable.

Due to the fact that the carbon bound in soot is not included in the reported CO₂ emissions in 2C3, the IEF (implied emission factor) values in the Swedish inventory (given as Mg CO₂/ Mg Al produced) are lower than the IPCC Guidelines default emission factors for prebaked and Söderberg (1.8 and 1.5 Gg CO₂/Gg produced Al) (Table 4-32).

Table 4-32. Implied emission factor for CO₂ for the production of aluminium.

Year	Aluminium production Gg	Emissions of CO ₂ Gg	IEF Gg CO ₂ /Gg Al
1990	96	133	1.4
1995	94	129	1.4
2000	101	145	1.4
2005	103	144	1.4
2006	102	142	1.4
2007	100	140	1.4
2008	82	114	1.4
2009	70	98	1.4
2010	96	135	1.4
2011	113	159	1.4

The two different processes for aluminium production, prebaked (CWPB) and Söderberg (VSS), have substantially different emission factors for PFCs. Estimates of emissions are based on the number of ovens and the number and duration of anode effects. This activity data is considered to be of good quality.

Activity data used for the PFC emission calculations, anode effects in min/oven day and production statistics, were provided by the company, and specified for the prebaked and Söderberg processes. The reported emissions and calculated Implied Emission Factors are presented in Table 4-33.

Table 4-33. Activity data, emissions of C₂F₆, CF₄ and calculated IEF for aluminium production.

Year	Al production, CWPB, Gg	Al production, VSS, Gg	Total emissions, C ₂ F ₆ Mg	Total emissions, CF ₄ Mg	Calculated IEF			
					CWPB kg C ₂ F ₆ /Mg	VSS kg C ₂ F ₆ /Mg	CWPB kg CF ₄ /Mg	VSS kg CF ₄ /Mg
1990	23.4	72.9	3.05	53.66	0.0443	0.0276	0.3444	0.6255
1995	22.8	71.2	2.29	48.25	0.0106	0.0287	0.0827	0.6510
2000	23.0	78.1	1.57	33.58	0.0059	0.0184	0.0460	0.4165
2005	23.6	78.9	1.66	36.93	0.0022	0.0204	0.0171	0.4629
2006	23.6	78.1	1.59	35.21	0.0024	0.0196	0.0188	0.4453
2007	23.3	76.5	1.61	35.54	0.0026	0.0202	0.0205	0.4583
2008	29.6	52.0	1.83	31.74	0.0223	0.0226	0.1737	0.5113
2009	69.7	-	0.56	4.36	0.0080	-	0.0625	-
2010	96.1	-	2.62	20.36	0.0272	-	0.2118	-
2011	113.3	-	3.02	23.49	0.0267	-	0.2073	-

4.4.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As can be seen in table 4-33 the IEFs show a downward trend from 1990 to 2007, especially so for CF₄. This reflects the company's on-going work aiming to reduce the time and frequency of the anode minutes. For 2008 it can be seen that the IEF for C₂F₆ as well as for CF₄ from the prebake ovens are higher compared to 2007. According to the company the reason for this is due to initial start-up problems with the new prebake ovens. The IEF for 2009 indicates less start-up problems of new prebake ovens in 2009. For both 2010 and 2011 the high IEFs reflect start-up problems and the major disturbances in plant 2 in 2010.

As described earlier (Figure 4-6) is the sharp decline in PFC emissions in 2009 caused by the closure of all Söderberg ovens in 2008.

The reported time series are considered to be consistent.

4.4.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.4.3.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.4.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.4.4 SF₆ used in aluminium and magnesium foundries (CRF 2.C.4)

4.4.4.1 SOURCE CATEGORY DESCRIPTION

In Sweden, four magnesium foundries use SF₆ as a cover gas. No SF₆ is used in aluminium foundries (CRF 2.C.4.1) as far as known, and thus reported as not occurring (NO). The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in table 4-34.

Table 4-34. Summary of source category description, CRF 2.C.4.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.C.4	SF ₆				D	D	Yes

D Default.

4.4.4.2 METHODOLOGICAL ISSUES

The total amount of SF₆ used annually in the magnesium foundries (CRF 2.C.4.2) is reported as emissions, according to the IPCC Guidelines and Good Practice Guidance. Data is obtained from companies using SF₆.

4.4.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Estimated uncertainty in SF₆ emissions is judge by SMED to be ±40%. Time series are considered to be consistent.

4.4.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In response to questions raised during the 2011 submission review, data for 2009 has been checked with information from the Swedish Chemicals Agency's Products Register and the data was found to be consistent.

4.4.4.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.4.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.4.5 Other metal production (CRF 2.C.5)

4.4.5.1 SOURCE CATEGORY DESCRIPTION

This sub-category includes CO₂, NO_x and SO₂ emissions from one large smelter producing various non-ferrous metals; copper, lead, zinc etc, and from one metal recycling company mainly producing lead. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-35.

Table 4-35. Summary of source category description, CRF 2.C.5.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.C.5	CO ₂				D	PS	Yes

D Default. CS Country Specific.

4.4.5.2 METHODOLOGICAL ISSUES

Emissions of CO₂ originate from one plant producing copper, lead and zinc, and one metal recycling plant mainly producing lead by melting used batteries and recover the lead.

CO₂ emissions from the smelter are calculated based on the amounts of coke, coal, limestone, plastics and other raw material used in the production. The company directly reports these activity as well as carbon content in slag products. The emissions from coal and coke are calculated based on national thermal values (TV) and emission factors (EF). IPCC default value is used for CO₂ emissions from limestone. The equation used for the smelter is:

$$\begin{aligned}
 \text{CO}_2 \text{ (Mg)} &= \text{Coke (Mg)} \times \text{EF} \times \text{Thermal value} + \text{Coal (Mg)} \times \text{EF} \times \text{Thermal value} \\
 &+ \text{Limestone (Mg)} \times 0.97 \times \frac{44.0098}{100.0892} + \text{C in raw material and plastics (Mg)} \times \frac{44}{12} \\
 &- \text{Slag (Mg)} \times 0.0002 \times \frac{44}{12}
 \end{aligned}$$

The metal recycling plant emits CO₂ from the melting of lead batteries composed of carbon containing plastics (polypropylene). The total CO₂ emissions from the plant are reported by the company for all years from 1990. For the years 1990 to 2003 the reported total CO₂ emissions also include energy related emissions. From 2004 the amount of plastics, their carbon content, as well as the CO₂ emission from plastics are known. This information for 2004 is used for estimating the process related CO₂ part of the total CO₂ emissions from the plant for the years 1990 until 2003. Also CO₂ originating from the limestone used is included. For the years 1990 – 2003 the yearly amounts of limestone used are estimated using activity data for 2004.

The reported emissions of SO₂ originate from the sulphur content in the raw materials used.

4.4.5.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Time-series are considered to be consistent.

4.4.5.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Both plants in this category report their emissions in yearly environmental reports. For the one plant included in the EU-ETS the reported activity data and emissions are analysed and compared to EU-ETS data. Where EU ETS data is judged to be appropriate and in line with the Good Practice Guidance, it is applied. More information on QC activities related to EU ETS is included in Annex 8.1.

4.4.5.5 SOURCE-SPECIFIC RECALCULATIONS

Slightly increased CO₂ emissions due to an additional raw material CO₂ source. The CO₂ emissions have been recalculated for the years 2007 to 2010.

4.4.5.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.5 Other production (CRF 2.D)

Other production covers emissions from the pulp and paper industry (2.D.1) as well as estimates from the production of food and drink (2.D.2). Emissions of fossil CO₂ are not estimated for this sector. According to the IPCC Guidelines Reference Manual, emissions of fossil CO₂ from this sector are not likely.

4.5.1 Pulp and paper (CRF 2.D.1)

4.5.1.1 SOURCE CATEGORY DESCRIPTION

The pulp and paper industry in Sweden is an important source of industrial process emissions. 42 individual pulp and paper facilities are included in the reported emissions, as well as two manufacturers of cardboard. One of these facilities shut down during 2008 and during 2009 another two plants closed down their pulp and paper production. The Kraft process (sulphate) dominates in Sweden but there are also emissions from four sulphite and 16 CTMP (Chemo Thermo Mechanical Pulp) or TMP (Thermo Mechanical Pulp) facilities reported in CRF 2.D, 1990 - 2011.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-36.

Table 4-36. Summary of source category description, CRF 2.D.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.D.1	CO ₂				NA	NA	NA
	CH ₄				CS	CS	Yes
	N ₂ O				CS	CS	Yes

CS Country Specific.

4.5.1.2 METHODOLOGICAL ISSUES

Reported emissions from the pulp and paper industry are primarily based on information about production and emissions in the companies' environmental reports. The industrial organisation within this sector has, for several years, cooperated closely with its members in developing sector-specific methods of measuring and calculating emissions, which have resulted in high quality emissions data. The reported emissions of NMVOC do not include terpenes.

The Swedish definition of process emissions includes the combustion of spent cooking liquor which gives rise to emissions of N_2O and CH_4 . The cooking liquor contains organic compounds and chemicals and is combusted to recover Na and S, but also to utilise the energy in the cooking liquor. The recovered Na and S (as Na_2CO_3 and Na_2S) are recycled and used in the process again. In submission 2008 and earlier, due to technical reasons, these emissions were reported in CRF 2.G. From submission 2008 and onwards, N_2O and CH_4 are reported in 2D1.

The estimated process emissions of CO_2 from quick lime production within this industry are allocated in CRF 2.A.2.

4.5.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainty in activity data is $\pm 5\%$ and uncertainty in emission factors (CH_4 and N_2O) are $\pm 20\%$.

4.5.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.5.1.5 SOURCE-SPECIFIC RECALCULATIONS

Minor corrections affecting NO_x in 2010, SO_x in 2009 and 2010 and NMVOC in 2010.

4.5.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.5.2 Food and drink (CRF 2.D.2)

4.5.2.1 SOURCE CATEGORY DESCRIPTION

The food and drink industry is a moderate source of NMVOC in Sweden. The industry consists of beer, wine and liquor producers, bread, sugar, yeast and margarine and solid cooking fat producers, coffee roasters and animal feed producers. Greenhouse gas emissions have not been estimated due to the lack of available methodology and data. Emissions of greenhouse gases are however considered to be insignificant.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-37.

Table 4-37. Summary of source category description, CRF 2.D.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.D.2	CO ₂				NA	NA	No, see Annex 5
	CH ₄				NA	NA	NA
	N ₂ O				NA	NA	NA

4.5.2.2 METHODOLOGICAL ISSUES

Estimates of NMVOC emissions are based on activity data from different official statistics. For wine the estimation of NMVOC emissions is based on data on sold amount¹²⁰ together with figures on import and export¹²¹. NMVOC emissions from beer production are based on the Swedish annual total production of beer^{122 123}. NMVOC emissions originating from the production of liquors, bread, sugar, yeast, margarine and solid cooking fat, coffee roasters and animal feeds are all based on statistics available at Statistics Sweden's website. For the NMVOC emission estimates, emission factors presented in Table 4-38 were used. Emissions of CO₂ are not estimated but are believed to be minor or of biogenic origin.

Table 4-38. NMVOC emission factors for the reported production activities in CRF 2.D.2 - Food and drink.

Production activity	Emission factor	Unit	Reference
Wine	0.8	kg/1,000 litres	120
Beer	0.35	kg/1,000 litres	123
Liquors	0.6	kg/1,000 litres	EF based on emission and activity data from one producer, 2001
Bread (sponge dough)	8	kg/Mg	121
Bread (white)	4.5	kg/Mg	121
Bread (whole meal and light rye)	3	kg/Mg	121
Bread (dark rye)	0	kg/Mg	121
Cakes	0.1	kg/Mg	121
Biscuits	0.1	kg/Mg	121
Breakfast cereals	0.1	kg/Mg	121
Sugar	10	kg/Mg	121
Yeast	18	kg/Mg	121
Margarine and solid cooking fats	10	kg/Mg	121
Coffee roasting	0.55	kg/Mg	121
Animal feed	0.1	kg/Mg	121

¹²⁰ Systembolaget. Försäljningsstatistik. <http://www.systembolaget.se/>

¹²¹ Statistics Sweden. <http://www.scb.se/>

¹²² Carlsberg Sweden. <http://www.carlsberg.se>

¹²³ Bryggeriföreningen. <http://sverigesbryggerier.se>

4.5.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The time series is consistent.

4.5.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.5.2.5 SOURCE-SPECIFIC RECALCULATIONS

Activity data, thus affecting reported NMVOC emissions, have been updated for:

- Wine: Import 2009, Export 2010
- Cider: Produced amount 2008-2010
- Beer: Produced amount 2009, 2010
- Liquors: Import and export 2010
- Bread (sponge dough): Produced amount 2008-2010
- Bread (white): Produced amount 2007-2010
- Bread (whole meal and light rye): Produced amount 2008-2010
- Cakes: Produced amounts 2003, 2004, 2008-2010
- Biscuits: Produced amounts 2002-2004, 2008-2010
- Breakfast cereals: Produced amounts 2002-2004, 2008-2010
- Sugar: Produced amounts 2002-2004, 2008-2010
- Margarine and solid cooking fats: Produced amounts 2003, 2004, 2008-2010
- Animal feed: Produced amounts for 2002, 2008-2010
- Coffee roasting: Produced amounts 2002-2004, 2008, 2010
- Yeast: Produced amounts 2002, 2004, 2010.

The recalculations resulted in changes in NMVOC emissions of about ± 0.2 Gg 2004-2010.

4.5.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.6 Production of Halocarbons and SF₆ (CRF 2.E)

Production of halocarbons and SF₆ does not occur in Sweden.

4.7 Consumption of Halocarbons and SF₆ (CRF 2.F)

Use and emissions of halocarbons have increased since 1990, especially in refrigeration and air-conditioning equipment, which is the major source of halocarbon emissions in Sweden in later years. The second largest source in 2010 is foam blowing (XPS-foam), followed by electrical equipment and aerosols. All remaining sources are comparatively small emitters of fluorinated greenhouse gases.

All sub-categories are covered in the estimates except solvents (2.F.5) According to the information available, solvents containing HFCs or PFC are not used in Sweden.

An overview of actual reported emissions in CRF 2F are shown in Table 4-39.

Table 4-39. Overview of submitted actual emissions data, Gg CO₂ equivalents.

CRF	Category	1990	1995	2000	2005	2006	2007	2008	2009	2010	2011
F1	Refrigeration and air conditioning equipment	3	125	430	668	714	752	782	798	780	745
F2	Foam blowing	NA	NA	111	87	74	54	51	40	32	37
F3	Fire extinguishers	NA	NA	5	6	6	6	8	6	6	6
F4	Aerosols/Metered dose inhalers	1	7	22	29	24	26	26	25	27	28
F5	Solvents	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO
F6	Other use of ODS substitutes	NO	NO	NO	NO	NO	NO	NO	NO	NO	NO
F7	Semiconductor manufacture	NA	11	8	NO	NO	NO	NO	NO	NO	NO
F8	Electrical equipment	81	95	32	28	22	29	28	44	31	30
F9	Other	2	3	8	14	12	9	8	8	8	5

In estimating the actual emissions in all subcategories, as far as possible, a national model has been used, corresponding to the IPCC Tier 2 approach. The basis for the emission estimates are the annual bulk import and export statistics of fluorinated greenhouse gases recorded in the Swedish Chemicals Agency's Products Register. However, the register does not cover all chemicals already included in products imported to or exported from Sweden (e.g. air-air heat pumps). In order to make a complete reporting of fluorinated greenhouse gas emissions and, as far as possible, to facilitate allocation of emissions onto the IPCC source categories, additional information from various trade associations and companies are collected annually. Based on an earlier inventory model on actual emissions of fluorinated greenhouse gases in Sweden covering the time

period 1990-1999¹²⁴, in 2005, the model was updated and refined e.g. concerning the calculations from the accumulated bank ¹²⁵. The model takes into consideration changes in accumulated amounts each year resulting from additional amounts of HFC, PFC and SF₆ imported and used within the country, as well as the decline in accumulated stock caused by exports or emissions from operating systems. In 2011, a SMED study¹²⁶ has been carried out to analyze the model's flexibility to adapt to the newly introduced international and national legislations on fluorinated greenhouse gases. In addition, the study aimed at updating model factors using available information, but also to analyze the accuracy of the estimates of e.g. emissions from disposal. The study resulted in several recalculations for the 2012 submission, but also suggestions on future improvements. The model is described in more details in Annex 3:1.

Due to a recurring one year lag of updating of the data from the Products Register from the Swedish Chemicals Agency, data on bulk import and export in 2010 are updated. This results in revised data on actual emission estimates from stationary refrigeration and air-conditioning equipment (2.F.1) and from electrical equipment (2.F.8) for 2010 due to the calculation system (as described in Annex 3:1).

4.7.1 Refrigeration and air conditioning equipment (2.F.1)

4.7.1.1 SOURCE CATEGORY DESCRIPTION

Emissions of HFCs and PFCs from heat pumps, stationary air-conditioning, mobile air-conditioning, refrigeration and freezing equipment are included in this category. Emissions of SF₆ from refrigeration and air conditioning equipment are not applicable (NA) in Sweden. The most important source of greenhouse gases to the category is emissions of HFC-134a from air-conditioning in cars. It can be seen in Table 4-39 that emissions of HFCs and PFCs from this source category has increased from 3 Gg CO₂ equivalents 1990 to 745 Gg CO₂ equivalents 2011. The use of HFCs as refrigerants in refrigerators, freezers, heat pumps and air-conditioning equipment in vehicles (MAC) is the main reason for the large increase in emissions. In 2010 and 2011, however, the emissions of HFCs are lower compared to 2009, mainly due to reduced use of HFC-134a in other stationary refrigeration equipment.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-40a.

¹²⁴ Kindbom, K., Haeger Eugensson, M. and Persson, K. 2001. Kartläggning och beräkning av potentiella och faktiska utsläpp HFC, FC och SF₆ i Sverige. IVL B-1428.

¹²⁵ Kindbom, K. 2005. Revision of Methodology and Estimated Emissions of Fluorinated greenhouse Gases in Sweden. Report Series SMED Nr 16 2005, www.smed.se

¹²⁶ Gustafsson, T. 2011. Fluorinated Greenhouse Gases in Sweden. Review of Methodology and Estimated Emissions Reported to the UNFCCC and the EU monitoring Mechanism. SMED report 2011.

Table 4-40a. Summary of source category description, CRF 2.F.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.F.1	HFCs	X	X		CS, T2	CS, D	Yes
	PFCs				CS	CS, D	Yes
	SF ₆				NA	NA	NA

D Default. CS Country Specific. T2 Tier 2.

4.7.1.2 METHODOLOGICAL ISSUES

Input data for the calculation of actual emissions consists of information from various sources; the Swedish Chemicals Agency, equipment producers and importers. Table 4-40b presents values for chemical charge, lifetime and emission factors for the applications used in the Swedish inventory. They are based on information from the equipment producers and IPCC default values. When data from equipment producers is used it has been compared against IPCC default data and been judged as reasonable. In the recent SMED study¹²⁷, based on contacts with the Swedish road vehicles manufacturers, several factors were modified for MAC for 2010 onwards to be more in line with the present status of the Swedish road vehicle fleet.

Table 4-40b. Typical values on equipment lifetimes, amounts of chemical per unit and emission factors for different applications of HFCs or PFCs used in calculations of actual emissions in Sweden. Intervals given indicate changes between 1990 and the last inventory year used in the calculations.

Application	Fluorinated substances	Lifetime**	Amount installed /unit, kg	Emissions at manufacturing	Emissions per year during use	Remained in product at disposal	Emissions at disposal
Household fridges and freezers	HFCs	20	0.1	2%***	1%	90%	5%
Heat pumps	HFCs	20→15	5→1	1%	10→1%	90%	5%
Other refrigeration and air conditioning equipment	HFCs PFC-218	15	*	3.5	7→3.6%	90%	5%
Refrigerated transport	HFCs	10	10→6	4.5%	30→7%	90%	15%
Mobile air-conditioning, lorries	HFCs	6	1.2	1%→0.5%	15→10%	90%	15%
Mobile air-conditioning, cars	HFCs	11	0.8→0.7	1%→0.5%	15→7.5%	90%	15%
Mobile air-conditioning, buses	HFCs	12	7	1%→0.5%	10%	90%	15%

* Top-down calculations

** Lifetime means the average expected lifetime of a product, not the designed technical lifetime from its first commissioning.

*** From Revised 1996 IPCC Guidelines

¹²⁷ Gustafsson, T. 2011. Fluorinated Greenhouse Gases in Sweden. Review of Methodology and Estimated Emissions Reported to the UNFCCC and the EU monitoring Mechanism. SMED report 2011.

The information on refrigerant-related imported amounts of fluorinated gases from the Swedish Chemicals Agency's Products Register is compared to calculations made in the model, based on assumptions and information from other sources. Since not all sources are possible to trace separately in the inventory, the amounts imported to the country according to the products register is larger than calculated from the individual sources covered in the model. In order to account for the total volumes of refrigerant-related fluorinated substances, the amount of imported chemical to Sweden, derived from the Products Register, is assumed to be the correct data. From these data, the amounts of chemicals already accounted for in other applications, treated separately in the calculations, are subtracted. The resulting remainder of all refrigerant-related HFCs and PFCs from the Products Register is allocated as input data in the sub source "other stationary refrigeration". The chemicals concerned are HFC-23, HFC-32, HFC-125, HFC-134a, HFC-143a, HFC-152a and PFC-218 (C_3F_8). In addition, it was discovered in the recent SMED study that imported air-air heat pumps were prefilled with the refrigerant (R410A), and thus not included in the bulk import statistics from the Swedish Chemicals Agency.

Due to that data are derived from source-independent national statistics in the Product Register, as well as from some end users, it is currently impossible to correctly fill in the CRF background data table asking for domestic, commercial and industrial applications. Consequently, industrial refrigeration as well as stationary air-conditioning have been reported as included elsewhere (IE) and the emissions are included in commercial refrigeration in CRF table 2(II) F.

4.7.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The largest contribution to the total national emission uncertainty from this category stem from HFCs from mobile air conditioner and other refrigeration. Based on SMED expert judgement AD and EF uncertainty are ± 10 % and ± 40 % for mobile air conditioner, and ± 25 % and ± 50 % for other refrigeration.

Data in the category is of varying quality, but generally considered, by expert judgment, to be of medium quality and is usually better for the later years than for the earlier years of the inventory. The time-series are calculated using the same methodology for all years and are thus considered to be consistent.

4.7.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Estimates have been checked with trade associations KYS (Kylbranschens Samarbetsstiftelse) and SVEP (Svenska Värmepumpföreningen) and with experts at the Swedish EPA¹²⁸. The information on refrigerant-related imported amounts of fluorinated gases from the Products Register is compared to calculations made in the model, based on assumptions and information from other sources.

¹²⁸ Swedish EPA . Ujfalusi, Bernekorn , and Björnell. Personal communication.

As HFCs from mobile air-conditioning in cars is the most influential sub-source in the category, its underlying factors are compared to IPCC default values and differences are analysed (Table 4-41). The values for car air-conditioner lifetime, charge and annual leakage were chosen based on information in IPCC Guidelines and Good Practice Guidance. The values for EF for production, remaining at decommissioning and share recovered are attained from the Swedish car manufacture Volvo and in cooperation with experts at the Swedish EPA.

Table 4-41. Comparison of IPCC default factors and Swedish factors for MAC in cars.

Parameter	1996 IPCC/ update GPG	Swedish factors	Comment
Lifetime (y)	12/12	11	OK
Charge (kg)	0.8	0.8	OK
Annual leakage (%/year)	10-30/10-20	15-10	OK
EF _{production} (%)	4-5/0.5	1-0.5	OK
Remaining at decommissioning (%)	75/40	90	High; We assume that there is continuous maintenance and refilling of the equipment
Share recovered (%)	0/0	85	OK according to experts at Swedish EPA

4.7.1.5 SOURCE-SPECIFIC RECALCULATIONS

General: Due to a recurring one year lag in the updating of the data from the Products Register from the Swedish Chemicals Agency, data on bulk import and export in 2010 were updated. It mainly affected emissions of HFC-134a and HFC-152 in commercial refrigeration and in total, emissions was reduced about 10 Gg CO₂ equivalents.

2.IIA.F.1.2 Commercial Refrigeration:

Calculation errors regarding the emission of HFC143a for the years 1999 to 2010 have been corrected. For most years this results in an increase of less than 1 %. For 2004 the decrease was 1.3 %. For 2008-2009 the correction lead to a decrease less than 1 %.

4.7.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.7.2 Foam blowing (2.F.2)

4.7.2.1 SOURCE CATEGORY DESCRIPTION

This category consists of HFCs emissions from production and use of XPS foam in Sweden. Emissions of PFCs and SF₆ from foam blowing are reported as not applicable (NA). Emissions of HFCs peaked in year 2000 and has since then decreased due to reduced leakage during manufacturing, according to data from the manufacturer.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-42.

Table 4-42. Summary of source category description, CRF 2.F.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.F.2	HFCs				CS	PS	Yes
	PFCs				NA	NA	NA
	SF ₆				NA	NA	NA

CS Country Specific. PS Plant-specific.

4.7.2.2 METHODOLOGICAL ISSUES

Data is obtained from the producer on the used amount of HFC-134a and HFC-152a, emissions at production as well as the exported amount of chemicals in products each year. The use of HFCs in this application started in 1996. The company has also provided algorithms to calculate leakage of HFC-134a and HFC-152a during the product lifetime (Table 4-43).

Table 4-43. Typical values on equipment lifetimes, amounts of chemical per unit and emission factors for different applications of HFCs used in calculations of actual emissions in Sweden.

Application	Fluorinated substances	Lifetime**	Amount installed /unit, kg	Emissions at manufacturing	Emissions per year during use	Remained in product at disposal	Emissions at disposal
Foam blowing (XPS)	HFCs	> 12	*	35%	Declining	\$	<76%***

* Top-down calculations

** Lifetime means the average expected lifetime of a product, not the designed technical lifetime from its first commissioning.

*** Based on remaining HFC in products at disposal after 12 years. 2008 is the first year for emissions at disposal in Sweden.

\$ Calculated according to a declining curve, different for HFC-134a and HFC-152a.

The basis for the calculation is the amount of HFC-134a and HFC-152a that is introduced into products used in Sweden, and subsequently leached from the products. Beside annual losses from products over time, the reported Swedish emissions in the CRF tables contain emissions from manufacturing.

More detailed information on the methodological issues for foam blowing is presented in Annex 3:4.

4.7.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The quality of activity data, such as amount of chemical used in applications, is usually better for the later years than for the earlier years of the inventory. Data from the manufacturers is considered to be complete and cover all sources of HFC emissions in Sweden. The time series are calculated using the same methodology for all years and are thus consistent.

4.7.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

The current calculation method provided by the company, used for reporting of emissions, has been compared to the Tier 2 method given in the Good Practice Guidance, see Annex 3:4.

4.7.2.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.7.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.7.3 Fire extinguishers (2.F.3)

4.7.3.1 SOURCE CATEGORY DESCRIPTION

HFC may be used as extinguishing medium in fixed fire extinguishing systems. In Sweden, emissions of HFCs from fire extinguishers are reported since 1997. Emissions of PFCs and SF₆ for the category are not applicable (NA).

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-44.

Table 4-44. Summary of source category description, CRF 2F3.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2F3	HFCs				CS	CS	Yes
	PFCs				NA	NA	NA
	SF ₆				NA	NA	NA

CS Country Specific.

4.7.3.2 METHODOLOGICAL ISSUES

All imports of HFCs to be installed in fire extinguishers are registered at the Swedish Chemicals Agency. From 2001, the use of HFC-227ea in fire extinguishers has been introduced in Sweden. Data has been obtained from the companies supplying such systems (Table 4-45).

Table 4-45. Typical values on equipment lifetimes, amounts of chemical per unit and emission factors for different applications of HFCs used in calculations of actual emissions in Sweden.

Application	Fluorinated substances	Lifetime**	Amount installed /unit, kg	Emissions at manufacturing	Emissions per year during use	Remained in product at disposal	Emissions at disposal
Fire extinguishing	HFCs	30	*	0.5%	2%/0.1%***	95%	1%

* Top-down calculations

** Lifetime means the average expected lifetime of a product, not the designed technical lifetime from its first commissioning.

*** HFC-227ea 0.1 %, other HFCs 2 %.

4.7.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties are mainly associated with the exported amounts, which are relatively large.

The time series are calculated using the same methodology for all years and are thus consistent.

4.7.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.7.3.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.7.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.7.4 Aerosols/metered dose inhalers (2.F.4)

4.7.4.1 SOURCE CATEGORY DESCRIPTION

HFC may be used as propellant gas in aerosols, but also as the actual product e.g. in cleaning sprays. In asthma medication inhalers, HFC-134a (norflurane) is sometimes used as propellant gas. Emissions of PFCs and SF₆ for the category are not applicable (NA).

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-46.

Table 4-46. Summary of source category description, CRF 2.F.4.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.F.4	HFCs				CS, T2	D	Yes
	PFCs				NA	NA	NA
	SF ₆				NA	NA	NA

D Default. CS Country Specific. T2 Tier 2.

4.7.4.2 METHODOLOGICAL ISSUES

Emission estimates cover technical aerosols as well as metered dose inhalers. The estimates consist of emissions from production of technical aerosols at one facility, and emissions from the use of imported technical aerosols and metered dose inhalers containing HFCs. The contribution from metered dose inhalers is relatively small, but has increased in later years.

The aerosol manufacturer provided information on the used amount of HFC-134a as well as emissions from production, and exported amounts of HFC-134a in products. Table 4-47 present the assumptions on product life-time, emissions at manufacturing and disposal as well as remaining HFC in product at disposal.

For metered dose inhalers, statistics on the numbers of sold inhalers was received from the Swedish retailer for medical products, Apoteket. Information concerning the content of HFC in the inhalers was provided by the Swedish Medical Products Agency.

Table 4-47. Typical values on equipment lifetimes, amounts of chemical per unit and emission factors for different applications of HFCs used in calculations of actual emissions in Sweden.

Application	Fluorinated substances	Lifetime**	Amount installed /unit, kg	Emissions at manufacturing	Emissions per year during use	Remained in product at disposal	Emissions at disposal
Aerosols/ MDI	HFCs	2	*	NA	50%	50%	100%

* Top-down calculations

** Lifetime means the average expected lifetime of a product, not the designed technical lifetime from its first commissioning.

4.7.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The largest uncertainty in this source concerns the amount of HFC-134a imported in technical aerosols for which there are no statistics available. In 2000, a survey was sent to approximately 10 importers of technical aerosol products. The majority of the importers responded to the survey, and provided estimates on the amount of HFC imported each year in technical aerosols. In 2004 an update on estimated import was made for the whole time series, in cooperation with the Swedish Aerosol Association (Svenska Aerosolföreningen). The information from this survey was used to update the time series up to year 2003 at that time. The activity data also includes estimates of e.g. Novelty aerosols.

The quality of activity data, such as figures of estimated emissions or amount of fluid used in different applications, is usually better for the later years than for the earlier years of the inventory. The time series are calculated using the same methodology for all years and are thus consistent.

4.7.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Data and information from the Products Register, hosted by the Swedish Chemicals Agency, could not be used directly for validation and reporting purposes due to confidentiality.

4.7.4.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.7.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.7.5 Solvents (2.F.5)

Efforts have been made to find national information concerning this sub-category. For instance potential users of solvents containing HFCs or PFCs were contacted. No information indicating that these kinds of solvents are used in Sweden was found. Emissions from solvents are consequently reported as NO, not occurring.

4.7.6 Other applications using ODS substitutes (2.F.6)

No other applications are covered in the Swedish inventory.

4.7.7 Semiconductor manufacture (2.F.7)

4.7.7.1 SOURCE CATEGORY DESCRIPTION

HFC, PFC and SF₆ are used in the semiconductor manufacturing process. Semiconductor manufacture has in recent years occurred on a commercial scale at only one facility in Sweden. Previously one more facility was located in Sweden, but production was moved abroad. During 2004 the production in the only facility left was also closed down.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-48.

Table 4-48. Summary of source category description, CRF 2.F.7.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.F.7	HFCs				T1	D	Yes
	PFCs				T1	D	Yes
	SF ₆				T1	D	Yes

D Default. T1 Tier 1.

4.7.7.2 METHODOLOGICAL ISSUES

Information concerning the annually used amounts of various fluorinated substances has been provided by the company, and as far as possible been compared to information from the Products Register at the Swedish Chemicals Agency. Emissions are calculated by using the Good Practice Guidance Tier 1 method (top-down calculations) using an average expected lifetime of 1 year.

4.7.7.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Emission estimates are judged to be of good quality. The quality of activity data is usually better for the later years than for the earlier years of the inventory. The time series are calculated using the same methodology for all years and are thus consistent.

4.7.7.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Cross-references with the Products Register at the Swedish Chemicals Agency could not be made for later years, since the level of detail in the Products Register was insufficient.

4.7.7.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.7.7.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.7.8 Electrical equipment (2.F.8)

4.7.8.1 SOURCE CATEGORY DESCRIPTION

In Sweden, emissions of SF₆ from electrical equipment consist of two different parts, emissions from the production of gas-insulated switchgear (GIS), and emissions from SF₆ installed in distribution systems. Emissions of HFCs and PFCs are not applicable (NA) for this category.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-49.

Table 4-49. Summary of source category description, CRF 2.F.8.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2F8	HFCs				NA	NA	NA
	PFCs				NA	NA	NA
	SF ₆				CS, D	CS, PS	Yes

D Default. CS Country Specific. PS Plant-specific.

4.7.8.2 METHODOLOGICAL ISSUES

The larger part of annual SF₆ emissions in earlier years originated from the manufacture of GIS (Table 4-50), where emissions in 1995 and 1997 peak due to a leaking valve in 1995 and to rebuilding and accidental leakages in 1997. The SF₆ emissions from production have decreased in later years due to measures taken at the production facility. These estimates, obtained from industry, are of medium to high quality, with better quality in later years. For the early 1990s, assumptions on the emitted amounts of SF₆ from GIS manufacture were made in cooperation with industry. Industry has also provided information concerning the used amount of SF₆ for GIS manufacture (Table 4-50), as well as the share of products that are exported from the country, which exceeds 90 % of the production.

Table 4-50. Typical values on equipment lifetimes, amounts of chemical per unit and emission factors for different applications of SF₆, used in calculations of actual emissions in Sweden. Intervals given indicate changes between 1990 and the last inventory year used in the calculations.

Application	Fluorinated substances	Lifetime**	Amount installed /unit, kg	Emissions at manufacturing	Emissions per year during use	Remained in product at disposal	Emissions at disposal
Electrical insulation and GIS manufacture	SF ₆	30	*	12→1.5%	0.6→0.5%	#	NA

* Top-down calculations

** Lifetime means the average expected lifetime of a product, not the designed technical lifetime from its first commissioning.

Estimated lifetime at least 30 years, NE.

Emissions from installed amounts of SF₆ for insulation purposes in operating systems have previously contributed less to the actual annual emissions. In 2001-2002, a questionnaire was sent out to power companies from the trade association Swedenergy¹²⁹ (Svensk Energi) asking for the installed amounts of SF₆ in operating equipment, and the replaced amounts of SF₆ during service. The results showed an installed accumulated amount of approximately 80 Mg and an annual leakage rate of 0.6 % (equals the amount replaced from the questionnaire) and these were used as input data in the inventory. For later years, data on replaced amounts of SF₆ in operating systems results in a calculated annual leakage rate of 0.5 % (Swedenergy and power distribution companies).

Table 4-51. Calculated emissions and accumulated stock of SF₆ for electrical equipment.

Year	Emissions from GIS manufacture SF ₆ Mg	Annual losses SF ₆ Mg	Accumulated stock Mg	Total emissions SF ₆ Mg
1990	3.0	0.4	65.7	3.4
1995	3.5	0.5	76.0	4.0
2000	0.7	0.6	107.9	1.3
2005	0.5	0.7	143.0	1.2
2006	0.2	0.7	148.7	0.9
2007	0.4	0.8	162.9	1.2
2008	0.3	0.9	173.4	1.2
2009	0.9	0.9	185.2	1.8
2010	0.3	1.0	196.6	1.3
2011	0.2	1.0	207.8	1.3

In accordance with the methodology described for deriving amounts of refrigerant chemicals not accounted for, the same procedure was adopted for SF₆. When comparing the amounts of SF₆ accounted for in various applications

¹²⁹ Swedenergy. Matz Tapper. Personal communication. 2005.

with data from the Products Register, a rather large annual volume of SF₆ remains unallocated (about 15-21 %). Sources of SF₆ emissions that are covered in the calculations are the use in semi-conductor manufacture, in production of sound-proof windows, in magnesium foundries, in the production of gas-insulated switchgear and as insulation in electrical equipment. Information from the Products Register did not indicate that any areas of use have not been covered and are missing from the calculations.

For all sources, except as insulation in electrical equipment, the levels of annual SF₆ consumption is comparatively easy to estimate with some confidence since there are few end-users. It was thus concluded that the amounts of SF₆ not already accounted for elsewhere, most reasonably should be allocated to the electrical equipment source. However, even though information concerning SF₆ in electrical equipment is more difficult to judge concerning completeness, indications from end-users are that the difference between imported amounts according to the Products Register and those already accounted for in the calculations seem too large to annually be consumed for electrical insulation. One explanation to the difference could be that there is an underreporting of exported SF₆ from the Products Register, where no export at all of SF₆ is registered.

As the question of the remaining amount of SF₆ at present could not be unambiguously solved, the unaccounted SF₆ from the Products Register was allocated to be used as electrical insulation (accumulated stock).

4.7.8.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The quality of activity data is usually better for the later years than for the earlier years of the inventory. The time series are calculated using the same methodology for all years and are thus consistent.

4.7.8.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.7.8.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.7.8.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.7.9 Other (2.F.9)

4.7.9.1 SOURCE CATEGORY DESCRIPTION

The estimated emissions from the use of SF₆ in jogging shoes and in sound-proof windows are reported in CRF 2.F.9. No production of SF₆-containing shoes occurs in Sweden. Since 2008 SF₆ has not been used in production of sound-proof windows.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 4-52.

Table 4-52. Summary of source category description, CRF 2.F.9.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
2.F.9	HFCs				NA	NA	NA
	PFCs				NA	NA	NA
	SF ₆				CS	CS, D, PS	Yes

D Default. CS Country Specific. PS Plant-specific.

4.7.9.2 METHODOLOGICAL ISSUES

For jogging shoes, a more or less rough estimate has been made. It has not been possible to obtain any national data, so a Norwegian estimate was scaled to the Swedish population.¹³⁰ According to the results from a study performed in early 2004¹³¹ a phasing out of SF₆ and replacement with PFC-218 was started in 2003. The lifetime for shoes is set to 8 years in the national model (Table 4-53).

Manufacturers of windows have provided data on the amount of SF₆ used in the manufacture of barrier gas windows. The manufacturers have also provided estimates of the share of SF₆ emitted in production (Table 4-53). These estimates vary considerably between manufacturers, from 5-50 %. The reason for the increase in emissions in later years is the lifetime and the associated time lag for emissions originating from disposal. Calculating a weighted average of the emission factor at production results in a national figure in the order of 30 %, which is in line with the point estimate of 33 % given in the Good Practice Guidance.

Table 4-53. Typical values on equipment lifetimes, amounts of chemical per unit and emission factors for different applications of PFCs or SF₆, used in calculations of actual emissions in Sweden. Intervals given indicate changes between 1990 and the last inventory year used in the calculations.

Application	Fluorinated substances	Lifetime**	Amount installed /unit, kg	Emissions at manufacturing	Emissions per year during use	Remained in product at disposal	Emissions at disposal
Sound proof windows	SF ₆	30	*	5-50%##	1%	#	NA
Jogging shoes	SF ₆ PFC-218	8	*	NA	NA	100%	25%

* Top-down calculations

** Lifetime means the average expected lifetime of a product, not the designed technical lifetime from its first commissioning.

Estimated lifetime at least 30 years, NE.

Different emissions at different production units.

¹³⁰ Weholt, Ø. 1999. Materialstrømsanalyse av SF₆. Beregning av potensielt og faktisk utslipp over tid

¹³¹ Kindbom, K. and Skårman, T. 2004. Nya scenarier för fluorerade växthusgaser. U952, Swedish EPA.

4.7.9.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The quality of activity data is usually better for the later years than for the earlier years of the inventory. The time series are calculated using the same methodology for all years and are thus consistent.

4.7.9.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.7.9.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations have been performed.

4.7.9.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

4.8 Consumption of Halocarbons and SF₆ Potential Emissions (CRF 2.F.P)

4.8.1 Potential emissions

Data on bulk imports and exports are obtained from the Products Register hosted by the Swedish Chemicals Agency, which did not register these substances until 1995. Estimates of potential emissions for imports and exports were, however, made for all years in the time series, 1990-2004 in a special study in 2005¹³². The method of estimating potential emissions for the following years was made accordingly.

Due to the recurring one year lag of updating of the data from the Products Register from the Swedish Chemicals Agency, data on bulk import and export in 2009 are updated. This results in revised data on potential emissions for 2009.

4.9 Other, CRF 2G

Not applicable for Sweden.

¹³² Kindbom, K. 2005. Revision of Methodology and Estimated Emissions of Fluorinated Greenhouse Gases in Sweden. Report Series SMED Nr 16 2005.

5 Solvent and other product use (CRF sector 3)

5.1 Overview of sector

This chapter describes emissions from solvents and other product use. Use of solvents and products containing solvents result in emissions of non-methane volatile organic compounds (NMVOC), which is regarded as an indirect greenhouse gas as it over a period of time will oxidise to CO₂ when emitted to the atmosphere.

Estimates reported in this sector include emissions from paint application (CRF 3.A), degreasing and dry-cleaning (CRF 3.B), chemical products, manufacture and processing (CRF 3.C) and other solvent use (CRF 3.D.5). Other use of N₂O (CRF 3.D.4) includes evaporative emissions of N₂O arising from other types of product use. This includes N₂O emissions from anaesthesia and aerosol cans.

Emissions of total greenhouse gases from the solvent and other product use sector (CRF 3) have decreased by 13 % from 332 Gg CO₂ equivalents in 1990 to 289 Gg CO₂ equivalents in 2011 (see Figure 5-1). The decline can largely be explained by a reduction in the use of solvents in CRF 3.A (paint application) due to a shift to water-based paints, which contain a smaller fraction of solvents compared to solvent-based paints.

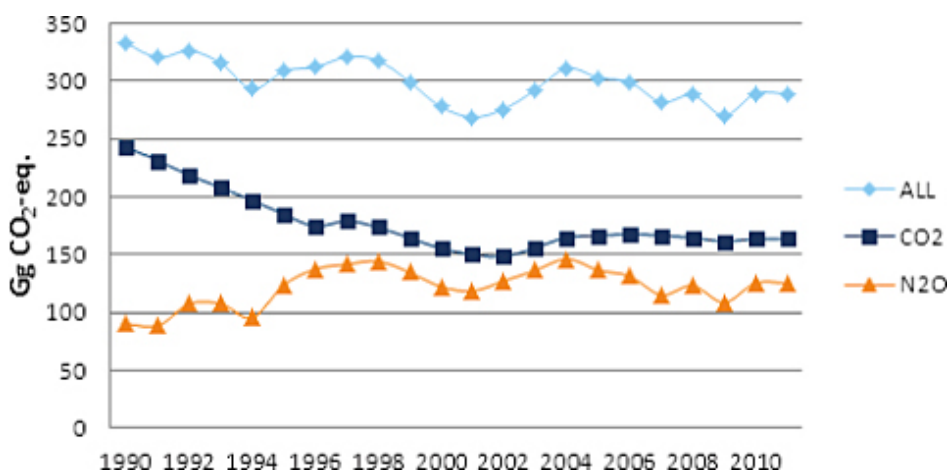


Figure 5-1. Total emissions of all greenhouse gases calculated as CO₂ equivalents from CRF 3 Solvent and Other product use.

CO₂ emissions in CRF 3.A (paint application) have decreased by 71 % from 94 Gg CO₂ equivalents in 1990 to 27 Gg CO₂ equivalents in 2011 (Figure 5-2). The largest source of greenhouse gas emissions from solvents in CRF sector 3 is CRF 3.D (other). For emissions of CO₂ in CRF 3.D we can see a minor decrease by less than 1 % from 138 Gg CO₂ in 1990 to 135 Gg CO₂ in 2011. CRF 3.D (other) also consists of N₂O emissions from CRF 3.D.4 (other use of N₂O). The use of N₂O has increased with 39 % from 90 Gg CO₂ equivalents in 1990 to 125 Gg CO₂ equivalents in 2011. In total, greenhouse gas emissions in CRF 3.D have increased by 14 % from 229 Gg CO₂ equivalents to 261 Gg equivalents in 2011.

Table 5-1 shows the impact of recalculations reported in submission 2012 for GHG emissions by sector and sub-sectors for 1990, 1995, 2000 and 2005-2010. Detailed descriptions of the recalculations are found under sector specific sections below.

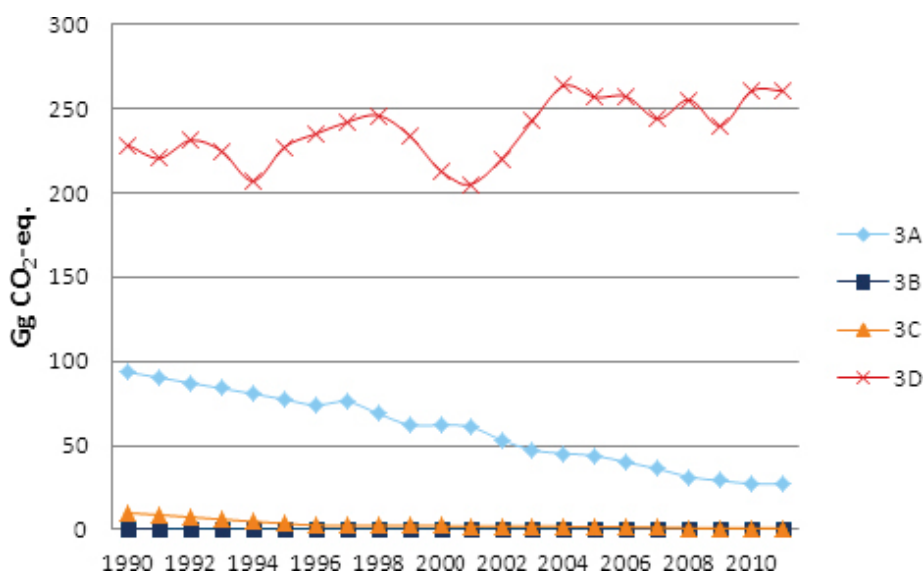


Figure 5-2. Total emissions of all greenhouse gases calculated as CO₂ equivalents from the different Solvent and Other product use sub-sectors. 3A - Paint application. 3B - Degreasing and dry-cleaning. 3C - Chemical products, manufacture and processing. 3D - Other.

Table 5-1. Impact of recalculations of GHG emissions submission 2013 in the Solvent and Other product use sector.

Impact of recalculations submission 2013 (Gg CO ₂ eq.)						
CRF	3A	3B	3C	3D	Total CRF 3	% CRF 3
1990	NA	NA	NA	NA	NA	NA
1995	NA	NA	NA	NA	NA	NA
2000	NA	NA	NA	NA	NA	NA
2005	NA	0	NA	0	0	0.0%
2006	0	0	0	-1	-1	-0.2%
2007	0	0	0	-15	-15	-5.1%
2008	-2	0	0	-21	-23	-7.5%
2009	-2	0	0	-39	-41	-13.2%
2010	-4	0	0	-18	-22	-7.1%

0: value less than 0.5 Gg. NA: no recalculation is performed.

The model used for estimating the CO₂ and NMVOC emissions reported in sector 3 is described in detail in Annex 3.3.

5.2 Paint application (CRF 3.A)

5.2.1 Source category description

Includes paints sold for “industrial use” and for “consumer and other professional use”.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 5-2.

Table 5-2. Summary of source category description, CRF 3A.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
3.A	CO ₂				CS	CS	Yes

CS Country Specific.

5.2.2 Methodological issues

All activity data from 1995 has been obtained from the Products register at the Swedish Chemicals Agency. Emissions from 1988 are taken from a time series that was compiled in a special study concerning NMVOC emissions, carried out by SMED in 2002¹³³. The emissions for 1990-1994 have been interpolated based on the information from the late 1980's and known data for 1995.

5.2.3 Uncertainties and time-series consistency

Reported time series are considered to be consistent, except for last year (2011) where data for previous year (2010) has been reported. This practice has been questioned by the ERT several times. The reason for Sweden to report activity data and emissions in CRF 3 with a delay of one year is due to the fact that activity data from the Product Register is not provided in sufficient time data to be able to perform the calculations and report in a timely manner.

5.2.4 Source-specific QA/QC and verification

No source-specific QA/QC or verification is performed.

5.2.5 Source-specific recalculations

Due to the recurring one year lag of updating of the data from the Product Register from the Swedish Chemicals Agency and the use of moving average for compiling the NMVOC and CO₂ time series in subsector 3A, the reported emissions for 2008 - 2010 were updated in submission 2013.

By mistake a substance with a vapour pressure of less than 0.01 kPa have been included in activity data derived from the Product Register. In submission 2013 this substance has been excluded which leads to minor recalculations in CRF 3.A for the years 2006 – 2010 (Table 5-1).

A correction has been performed in the calculation model for CRF 3.A for emission year 2008, which slightly affects reported emissions and activity data for 2005 - 2010 (Table 5-1).

5.2.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

¹³³ Kindbom, K., Boström, C-Å., Skårman, T., Gustafsson, T. and Talonpoika, M. 2003. Estimated Emissions of NMVOC in Sweden 1988-2001.

5.3 Degreasing and Dry cleaning (CRF 3.B)

5.3.1 Source category description

Includes solvents sold to the laundry and dry cleaning industry. Degreasing is included in CRF 3D.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 5-3.

Table 5-3. Summary of source category description, CRF 3B.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
3.B	CO ₂				CS	CS	Yes

CS Country Specific.

5.3.2 Methodological issues

All activity data from 1995 has been obtained from the Products Register at the Swedish Chemicals Agency. Emission data for 1988 is based on reported quantities of tetrachloroethylene from the Swedish Chemicals Agency. After 1995 also other substances for degreasing and dry cleaning are included. Of the total amount of NMVOC used within CRF 3B these “non tetrachloroethylene” substances contribute approximately 30%. As not only tetrachloroethylene is included in the time series after 1995, the NMVOC emissions reported 1988 is recalculated using a correction factor based on the proportion of other NMVOCs of the total NMVOC for 1995 (tetrachloroethylene plus 30 %). Emissions between 1990 and 1994 have been interpolated based on the information from the late 1980’s and known data for 1995. The solvents used within CRF 3B includes a lower carbon share compared to the solvents used in the other sub-codes within CRF 3.

5.3.3 Uncertainties and time-series consistency

Reported time series are considered to be consistent, except for last year (2011) where data for previous year (2010) has been reported. This practice has been questioned by the ERT several times. The reason for Sweden to report activity data and emissions in CRF 3 with a delay of one year is due to the fact that activity data from the Product Register is not provided in sufficient time data to be able to perform the calculations and report in a timely manner.

5.3.4 Source-specific QA/QC and verification

No source-specific QA/QC or verification is performed.

5.3.5 Source-specific recalculations

Due to the recurring one year lag of updating of the data from the Product Register from the Swedish Chemicals Agency and the use of moving average for compiling the NMVOC and CO₂ time series in subsector 3B, the reported emissions for 2008 - 2010 were updated in submission 2013.

A few corrections for CRF 3.B have been performed in the calculation model for emission years 2004 and 2007. These corrections affect slightly reported emissions and activity data for 2002 - 2010 (Table 5-1).

5.3.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

5.4 Chemical products, Manufacture and Processing (CRF 3.C)

5.4.1 Source category description

Includes solvents sold for car manufacturing, paint industry and rubber industry.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 5-4.

Table 5-4. Summary of source category description, CRF 3C.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
3.C	CO ₂				CS	CS	Yes

CS Country Specific.

5.4.2 Methodological issues

The category includes emissions from car manufacturing, paint industry and from rubber industry. Emissions from car manufacturing contributed in 2005 by approximately 50%, paint industry by 35 % and rubber industry by 15 % of the reported emissions in CRF 3C. The corresponding figures for 2011 are 35 %, 45 % and 20 %, respectively. Emission data for car manufacturing has been compiled from environmental reports for 1990 and data for 1991-1994 has been interpolated. For paint industry emission data for 1990-1994 has been taken from the old time series given in a special study concerning NMVOC emissions, carried out by SMED in 2002¹³³. Emission data for the rubber industry is known for 1988¹³³ and data for 1990-1994 have been interpolated based on the information from the late 1980's and known data for 1995.

5.4.3 Uncertainties and time-series consistency

Reported time series are considered to be consistent, except for last year (2011) where data for previous year (2010) has been reported. This practice has been questioned by the ERT several times. The reason for Sweden to report activity data and emissions in CRF 3 with a delay of one year is due to the fact that activity data from the Product Register is not provided in sufficient time data to be able to perform the calculations and report in a timely manner.

5.4.4 Source-specific QA/QC and verification

No source-specific QA/QC or verification is performed.

5.4.5 Source-specific recalculations

Due to the recurring one year lag of updating of the data from the Product Register from the Swedish Chemicals Agency and the use of moving average for compiling the NMVOC and CO₂ time series in subsector 3C, the reported emissions for 2008 - 2010 were updated in submission 2013.

By mistake a substance with a vapour pressure of less than 0.01 kPa has been included in activity data derived from the Product Register. In submission 2013 this substance have been excluded which leads to minor recalculations in CRF 3.C for the years 2006 – 2010 (Table 5-1).

5.4.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

5.5 Other (CRF 3.D)

5.5.1 Source category description

All data concerning solvents, NMVOC and CO₂, are reported in CRF 3.D.5. CRF 3.D.5 includes solvents sold to the printing industry, for preservation of wood, to leather industry and to textile industry. The code also includes solvents used by other industries not reported separately, and solvents for domestic use. In CRF 3.D.4 sold amounts and use of N₂O are reported. Due to confidentiality, data for 3.D.1 - Use of N₂O for Anaesthesia and 3.D.3 – N₂O from Aerosol cans cannot be reported separately.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 5-5.

Table 5-5. Summary of source category description, CRF 3D.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
3.D	CO ₂				CS	CS	Yes
3.D	N ₂ O				CS	CS	Yes

CS Country Specific.

5.5.2 Methodological issues

Solvents used in printing industry, for preservation of wood, in leather industry and in textile industry have been estimated separately. The code also includes solvents used by other industries not reported separately, and also solvents for domestic use. The printing industry contributes to totally reported CO₂ and NMVOC in CRF 3.D by around 8 %. The corresponding figure for preservation of wood and leather and textile industry is below 1%, while general solvent use represents over 90% of the total reported emissions in CRF 3D. Emission data for 1988 is known for most industries included in CRF 3D and in most cases the emissions for 1990-1994 have been interpolated based on information from the late 1980's and known data for 1995.

There are two companies in Sweden selling N₂O in gas cylinders. Information on sold amounts was obtained from one of the companies (1990 - 1991) and from the Products Register at the Swedish Chemicals Agency (1992 - 2010). The time series of use of N₂O in Sweden are reported in Other use of N₂O (3.D.4) since no background data is available to separate between the source categories Use of N₂O for Anaesthesia (3.D.1) and N₂O from Aerosol cans (3.D.3). Consequently CRF codes 3.D.1 and 3.D.3 are both reported as IE. Activity data for the latest year, 2011, is not yet official and hence Sweden has chosen to report data from 2010 also for 2011. Data for 2011 will be updated in the next submission.

5.5.3 Uncertainties and time-series consistency

Reported time series are considered to be consistent, except for last year (2011) where data for previous year (2010) has been reported. This practice has been questioned by the ERT several times. The reason for Sweden to report activity data and emissions in CRF 3 with a delay of one year is due to the fact that activity data from the Product Register is not provided in sufficient time to be able to perform the calculations and report in a timely manner.

5.5.4 Source-specific QA/QC and verification

No source-specific QA/QC or verification is performed.

5.5.5 Source-specific recalculations

Due to the recurring one year lag of updating the data from the Product Register from the Swedish Chemicals Agency, the reported emissions of N₂O in CRF 3D4 for 2010 is updated in submission 2012.

Due to the recurring one year lag of updating of the data from the Product Register from the Swedish Chemicals Agency and the use of moving average for compiling the NMVOC and CO₂ time series in subsector 3D, the reported emissions for 2008 - 2010 were updated in submission 2013.

By mistake a substance with a vapour pressure of less than 0.01 kPa has been included in activity data derived from the Product Register. In submission 2013 this substance have been excluded which leads to recalculations in CRF 3.D for the years 2006 – 2010 (Table 5-1). This correction affects mainly emissions and activity data for printing industry included in CRF.3.D.5.

Also corrections have been made in the calculation model for CRF 3.D.5 Other, other non-specified use of solvents for emission years 2004, 2005, 2007, 2008 and 2009. These corrections affect slightly reported emissions and activity data for 2002 - 2010 (Table 5-1).

5.5.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

6 Agriculture (CRF sector 4)

6.1 Overview of sector

Only emissions of nitrous oxide (N₂O) and methane (CH₄) are reported in the agricultural sector. Carbon dioxide emissions from agriculture are reported under other sectors. Carbon dioxide from working vehicles and other energy use is reported in the energy sector and carbon dioxide from liming of agricultural land is reported in the LULUCF sector. Sweden's inventory includes emissions from enteric fermentation, manure management and agricultural soils. Rice cultivation, burning of savannas or burning of agricultural residues does not occur in Sweden. The agriculture in Sweden has undergone radical structural changes and rationalisations over the past 50 years. One fifth of the Swedish arable land cultivated in the 1950s is no longer farmed. Closures have mainly affected small holdings and those remaining are growing larger. Livestock farmers predominately engage in milk production and the main crops grown in Sweden are grain and fodder crops.¹³⁴ The decrease of agricultural land area has continued since Sweden joined the European Union in 1995 and the acreages of land for hay and silage has increased. Organic farming has increased from 6 % of the arable land area in 1996 to 17 % in 2010.¹³⁵

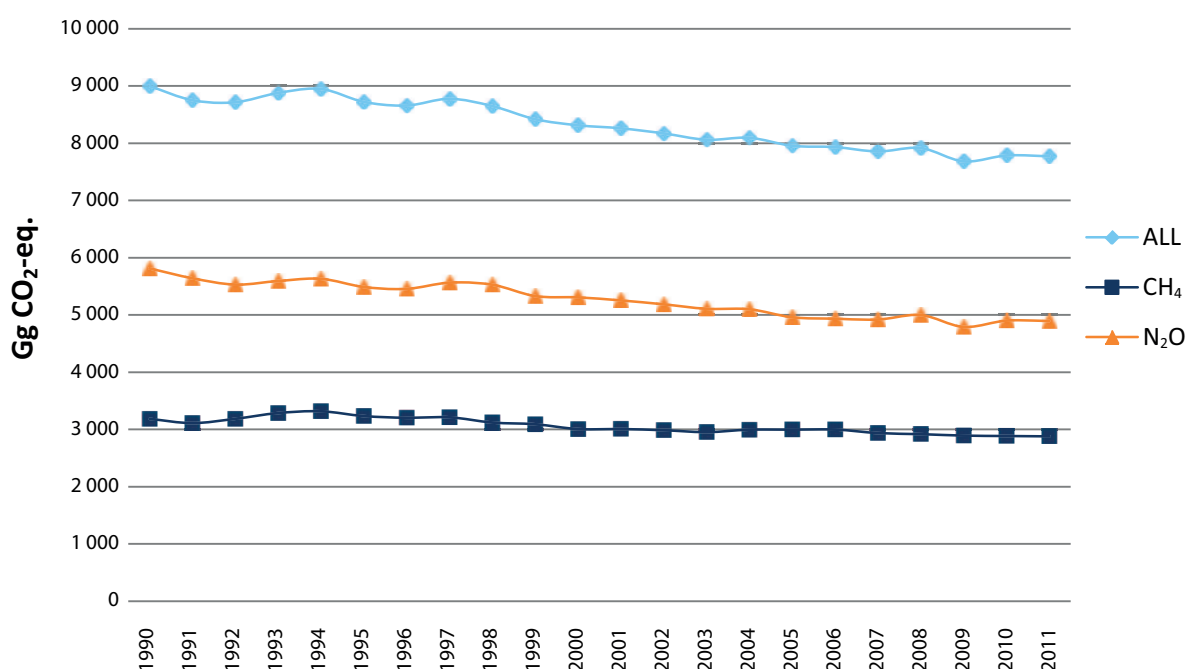


Figure 6-1. Total emissions of all greenhouse gases calculated as CO₂ equivalents from CRF 4 Agriculture.

¹³⁴ Ministry of the Environment, 2001.

¹³⁵ Swedish Board of Agriculture, www.jordbruksverket.se, <http://miljomal.nu>

The total greenhouse gas (GHG) emissions from sector four have decreased by 14 % since 1990, from 8,997 Gg CO₂ equivalents to 7,772 Gg CO₂ equivalents (Figure 6-1). The largest emissions in this sector are nitrous oxide from nitrogen circulation in agricultural land (Figure 6-2).

Recalculations are made in submission 2013 (Table 6-1). Detailed descriptions of the recalculations are found under the sector specific sections.

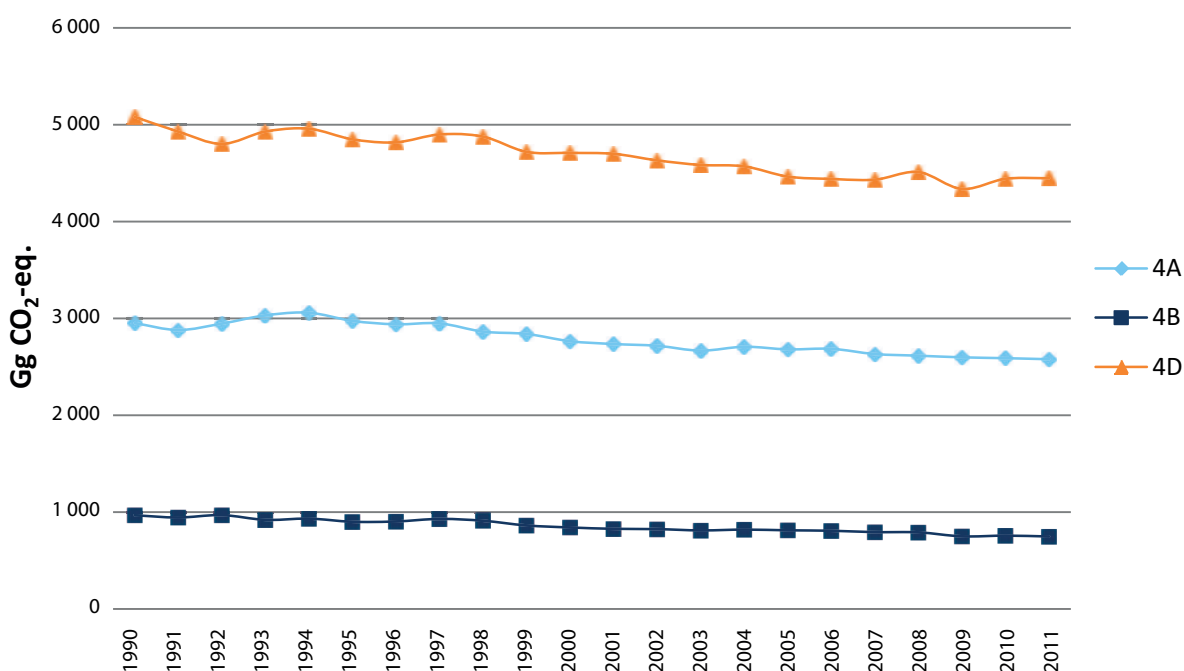


Figure 6-2 Total emissions of all greenhouse gases calculated as CO₂ equivalents from the different agricultural sub-sectors. There are no emissions from 4.C or 4.E-G.

Table 6-1. Impact of recalculations of GHG emissions submission 2013 in the agricultural sector.

Impact of recalculations submission 2013 (Gg CO ₂ eq.)								
CRF	4.A	4.B	4.D.1	4.D.2	4.D.3	4.D.4	Total CRF 4	% CRF 4
1990	-120	0	0	0	1	1	-118	-1.3
1995	-134	1	0	0	0	0	-132	-1.5
2000	-126	1	1	0	2	1	-121	-1.4
2005	-129	1	1	0	1	1	-125	-1.5
2006	-127	1	1	0	1	1	-122	-1.5
2007	-126	1	1	0	1	1	-121	-1.5
2008	-128	1	0	0	0	0	-126	-1.6
2009	-127	1	0	0	0	0	-126	-1.6
2010	-125	1	-1	0	-1	-1	-127	-1.6

0: value less than 0.5.

Livestock is the main contributor to greenhouse gas emissions from agriculture. In Table 6-2 all the livestock subgroups used in the calculations are presented. Mink and foxes are minor contributors to greenhouse gas emissions and are not included in the inventory due to a lack of well-founded emission factors.

Table 6-2. Livestock subgroups used in the calculations.

Categories according to IPCC Guidelines	Sub-categories Enteric Fermentation	Sub-categories Methane from manure management	Sub-categories N ₂ O from manure management	Sub-categories N ₂ O from grazing animals
Dairy Cattle (**)	Dairy cows	Dairy cows	Dairy cows	Dairy cows
Non-Dairy Cattle (**)	Beef cows	Beef cows	Beef cows	Beef cows
	Other cattle	Growing animals (12-24 months)	Growing animals (12-24 months)	Growing animals (12-24 months)
		Calves > 6 months	Calves > 6 months	Calves > 6 months(*)
		Calves < 6 months	Calves < 6 months	Calves < 6 months(*)
Swine	Swine	Sows	Sows	NO
		Boars	Boars	
		Pigs for meat production	Pigs for meat production	
		Piglets	Piglets	
Sheep	Sheep	Sheep	Sheep	Sheep
Goats	Goats	Goats	Goats	Goats
Horses (***)	Horses	Horses	Horses	Horses
Poultry	Poultry	Poultry	Laying hens (**)	NO
			Chickens (**)	
			Slaughter Chickens (****)	
Other (*****)	Reindeer	NO	NO	Reindeer

(*) The age distribution of calves is accomplished by using standard values.

(**) Farm Register. (***) Statistics Sweden. (****) Swedish Poultry Meat Association. (*****) Sametinget (The Sami Parliament of Sweden).

The Farm Register provides the main basis for agricultural statistics in Sweden. The Register is administered by the Swedish Board of Agriculture and Statistics Sweden and provides annual information on the total number of animals of different categories on Swedish farms¹³⁶. The information on livestock refers to the situation prevailing in mid-June of that year and thus is considered to be equivalent to a one-year average. Most of the information on livestock numbers comes from the Farm Register, but the distribution of calves (older and younger than 6 months respectively) is model-assisted: 60 % are assumed to be younger than 6 months and the rest are assumed to be over 6 months old.

Concerning horses, the Farm Register gives an underestimation of the number of horses because only horses on farms are included (i.e. not horses for leisure activities). However, a separate survey¹³⁷ estimated total number of horses in Sweden in 2010 and 2004, these estimates are used in the calculations instead. The number of slaughter chickens (mean number of chickens kept during the year) is provided by the Swedish Poultry Meat Association. This estimate is generally higher than the estimate given by the Farm Register, which on the other hand is considered to be too low.

¹³⁶ Swedish Board of Agriculture, JO 20-series.

¹³⁷ Swedish Board of Agriculture, 2011

6.2 Enteric Fermentation (CRF 4.A)

6.2.1 Source category description

The animal husbandry sub-sector is an important sector influencing GHG emissions from agriculture in Sweden. Livestock farming, including farmyard manure management, is the major source of CH₄ emissions. From the total emission of CH₄ about 85 % derives from enteric fermentation from cattle. The total numbers of livestock in Sweden in 1990-2011 are presented in Table 6-6 and Figure 6-3.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 6-3.

Table 6-3. Summary of source category description, CRF 4.A.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
4.A	CH ₄	X			CS, T1, T2	CS, D	Yes

CS Country Specific. T1 Tier 1. T2 Tier 2. D Default.

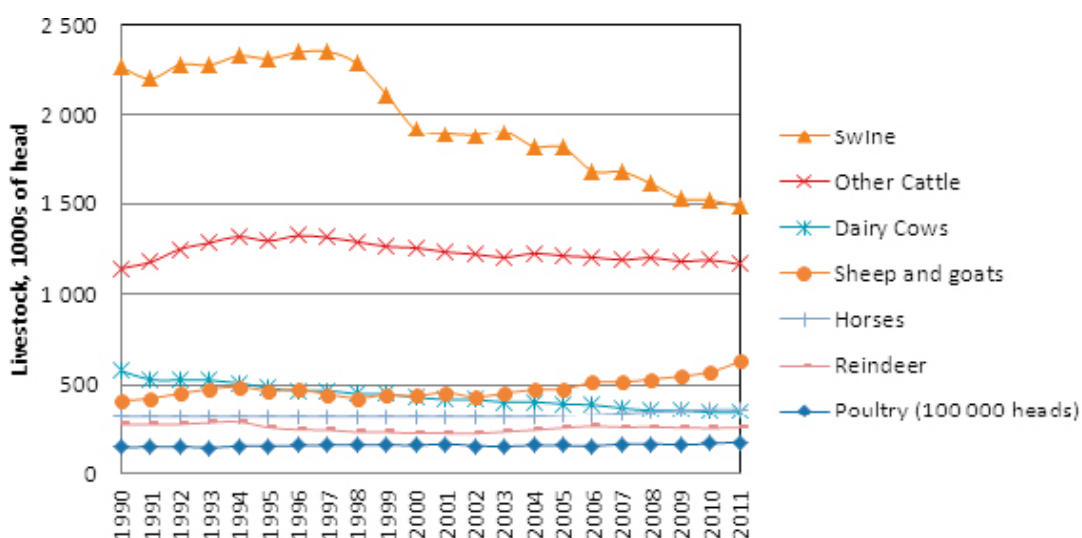


Figure 6-3 Livestock in Sweden 1990-2011, 1,000s of head

6.2.2 Methodological issues

6.2.2.1 EMISSION FACTORS, METHANE

The livestock population (Table 6.6) in each category is multiplied by an emission factor and the total emission is calculated as:

$$emissions = \sum_i population_i * EF_i$$

Emission factors for the significant cattle subgroups are national. For reindeer, where the IPCC Guidelines do not provide default values, an emission factor is calculated according to the IPCC Guidelines methodology using a Finnish

value of gross energy requirements.¹³⁸ For emissions from swine, sheep, goats and horses the IPCC default values are used. Statistics on livestock categories are presented in Table 6-. A national methodology based on feed energy requirements expressed as metabolisable energy¹³⁹ is used in the Swedish inventory to estimate emission factors for dairy cows, beef cows and other cattle. For dairy cows during the lactation we first calculate the metabolisable energy (MJ/day) from the energy requirements for maintenance and lactation. Data on milk yield is from the trade organisation Swedish Milk and built on the production evaluation tool “Kokontrollen”. It is a voluntary tool but because the farmers can use the results for optimisation and follow-up on productivity, about 80 % of the farmers are connected. For the cows not connected Swedish Milk assumes a lower productivity. The data used to calculate average milk production are given in Table 6-6. The metabolisable energy is then used to estimate digestible energy, and from this the emission of methane is calculated using the methane conversion rate.

The calculations for dairy cows were revised some years ago¹⁴⁰. The emission factors for other cattle groups were also re-evaluated, using the same methodology¹⁴⁰. The conclusion led to a decision to use the emission factor, 50 kg CH₄/head and year for the sub-category “Other Cattle”, a value close to the Good Practice Guidance default value for non-dairy cattle (48 kg CH₄/head and year). Due to the recommendation of the ERT during the in-country visit in Sweden in 2007, CH₄ emission factors for beef cows and reindeers were revised to 78.0 kg CH₄/head and year and 19.9 kg CH₄/head and year, respectively. Good Practise Guidelines (GPG) recommends that the methane conversion rate should be zero for calves feeding only milk. Personal communication with Swedish experts resulted in that this period is assumed to be two months for calves of dairy breed and three months for other calves.

Below follows a review of the different stages in the calculation for dairy cows. Metabolisable energy (MJ/day) is calculated using the formula from Spörndly 1999:

Lactation period:

$$\text{Metabolisable energy (MJ/day)} = 1.11 \times (62 + \text{Milk Production/day} \times 5) - 13.6$$

Dry period:

$$\text{Metabolisable energy (MJ/day)} = 62 + 13$$

Metabolisable energy is then converted to digestible energy using the formula¹⁴¹:

$$\text{Metabolisable energy (\% of digestible energy)} = 83.2 + 2.53 \times L - 0.045 \times G - 0.184 \times R_p$$

¹³⁸ Statistics Finland, 2007

¹³⁹ Lindgren, 1980; Murphy, 1992; Bertilsson, 2002.

¹⁴⁰ Bertilsson, 2001.

¹⁴¹ Lindgren, 1980.

Methane conversion rate of digestible energy is calculated using the formula¹⁴²:

$$\text{Methane conversion rate (\% methane of digestible energy)} = 15.7 - 0.030 \times \text{SK} - 1.4 \times \text{L}$$

Where L is the total feed intake expressed as multiples of maintenance energy. G is the share of coarse feed and Rp is the crude protein content in the food expressed as percentage of total food consumption in dry matter (dm). SK is the digestibility of the feed (% of gross energy). All used constant for the different years are given in Table 6-4. Finally, the actual emission of methane is estimated with the formula:

$$\text{Emission of methane (Kg/day)} = (\text{Digestible energy} \times \text{Methane conversion rate}) / 55.65$$

The Swedish method resembles the one used by IPCC. The main difference is that Sweden uses metabolisable energy in the calculations as opposed to gross energy intake. Furthermore, the energy loss caused by methane emissions is calculated as a fraction of digestible energy. This fraction is in turn determined by total feed intake and digestibility of the feed. IPCC instead express energy in methane as a constant fraction of gross energy in feed.

Table 6-4. Constants used for estimating methane from dairy cattle.

Year	Dry cows				During lactation			
	L	G	Rp	SK	L	G	Rp	SK
1990	1.2	80.1	14	69	3.0	55	16	69
1995	1.2	80.1	14	69	3.2	55	16	69
2000	1.2	80.1	14	69	3.5	55	16	69
2005	1.2	80.1	14	69	3.5	55	16	69
2006	1.2	80.1	14	69	3.5	55	16	69
2007	1.2	80.1	14	69	3.5	55	16	69
2008	1.2	80.1	14	69	3.5	55	16	69
2009	1.2	80.1	14	69	3.5	55	16	69
2010	1.2	80.1	14	69	3.5	50	17	69
2011	1.2	80.1	14	69	3.5	50	17	69

For the year 2011 this resulted in an emission of 397 g of methane per day for milk cows during the lactation period. The lactation period is estimated to 305 days per year. For cows during the non-lactation period the value does not change between years and is estimated to 201 g of methane per day. The final emission from milk cow per head and year is then calculated from the emission during the lactation and the non-lactation period.

From these variables it is possible to calculate gross energy intake (GE) and the methane conversion rate for gross energy (Y_m) for dairy cows that are reported in the CRF-tables despite that we do not actually use them in the calculation of the emission.

¹⁴² Lindgren, 1980.

For this we use the formulas:

Gross energy intake (GE) = ((Digestible energy during lactation × 305 + Digestible energy for dry cows × 60)/365) / SK

Methane conversion rate (Y_m) = CH_4 /head/year × 55.65 / (GE × 365)

The default values in the IPCC Guidelines are used for the less significant animal groups¹⁴³ and for these groups the development of a national emission factor has not been given priority. The emission factors used for dairy cattle and other animal groups are collected in Table 6-7.

Table 6-5. Number of dairy cows and average milk production.

Year	Dairy cows in country, number of head(*)	Dairy cows in the official control activity, number of head (**)	Produced milk per head in official control activity, kg/head/yr (**)	Produced milk per head, not in official control activity (**)	Average milk production per head, kg/yr (***)
1990	576,000	421,780	7,319	5,330	6,786
1991	528,000	388,860	7,376	5,280	6,824
1992	526,000	367,452	7,376	5,400	6,780
1993	525,000	376,126	7,740	5,600	7,133
1994	509,000	383,124	8,011	6,100	7,538
1995	482,000	390,146	8,083	6,200	7,724
1996	466,000	382,511	8,033	6,150	7,696
1997	468,000	380,760	8,209	6,250	7,844
1998	449,000	380,567	8,298	6,258	7,987
1999	449,000	378,623	8,377	6,300	8,051
2000	428,000	368,350	8,537	6,430	8,243
2001	418,000	360,364	8,742	6,627	8,450
2002	417,000	354,801	8,784	6,665	8,468
2003	403,000	346,133	8,794	6,750	8,506
2004	404,000	332,367	8,994	6,750	8,596
2005	393,000	332,367	8,994	6,750	8,648
2006	388,000	318,986	9,283	6,750	8,832
2007	369,646	298,865	9,412	6,750	8,902
2008	357,194	293,939	9,322	6,750	8,867
2009	356,776	285,246	9,486	6,750	8,937
2010	348,095	275,715	9,468	6,750	8,903
2011	346,495	287,606	9,480	6,750	9,016

(*) Farm Register, (**) Swedish Dairy Association. (***) Calculated value.

¹⁴³ According to current estimations, "other animal groups" produce less than 10 % of the total methane that results from enteric fermentation.

Table 6-7. Methane from animals, used emission factors.

Livestock subgroups	Kg CH ₄ / head/year	Method
Dairy cows in 1990, average milk production 6,786 kg/yr/head	120.3	1
Dairy cows in 2011, average milk production 9,016 kg/yr/head	133.1	1
Beef cows	78	4
Growing animals (12-24 months)	50	4
Calves (excluding calves on milk)	50	4
Swine	1.5	2
Sheep	8	2
Goats	5	2
Horses	18	2
Poultry	No fermentation assumed	
Reindeer	19.9	3

(1) The emission factor is related to milk production and calculated from Spörndly, 1999 and Bertilsson, 2001.

(2) IPCC Guidelines. (3) Statistics Finland, 2007, Tier 2. (4) Bertilsson, 2001.

6.2.3 Uncertainties and time-series consistency

Between 1995 and 1996 there was an increase in the number of sows by 13 %. The reason for this sudden increase is that as from this year also uncovered gilts are included in this group. When no estimate on number of horses exists before 2004, this value is used for all preceding years. From 2010 there was a minor change in the values of the share of coarse feed and the crude protein content in dairy cow food (see Table 6-4).

6.2.4 Source-specific QA/QC and verification

The time series for the different populations and milk production is checked for consistency. Recalculations are identified and verified. Emission estimates and methods are reviewed yearly by external national experts.

6.2.5 Source-specific recalculations

Due to comments received from the EU ESD review, the emissions from calves are recalculated, now assuming zero emissions from calves feeding only milk.

6.2.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

Table 6-6. Population size of different animal groups (1,000s heads).

Year	Dairy cows	Non-Dairy Cattle		Swine		Sheep		Horses		Goats		Other	Poultry	
	Dairy cows	Beef Cows	Growing animals (12-24 months)	Calve	Sow (***)	Pig for meat pro-duction	Piglet	Boar	Sheep	Lamb	Horse (*)	Goat (**)	Kid (***)	Slaughterer Chicken (**)
1990	576	75	543	524	221	1,276	758	8.6	162	244	316	2.9	1.4	5,500
1991	528	98	543	537	219	1,239	736	8.3	168	251	316	3.2	1.6	5,833
1992	526	136	565	548	225	1,283	763	8.3	180	267	316	3.5	1.8	6,333
1993	525	154	549	581	241	1,272	756	7.9	189	282	316	3.5	1.8	6,333
1994	509	165	561	592	241	1,264	815	8.2	196	288	316	3.5	1.8	6,833
1995	482	157	596	542	237	1,300	768	7.6	195	266	316	3.5	1.8	7,083
1996	466	164	617	543	273	1,303	765	6.9	203	266	316	3.5	1.8	7,250
1997	468	169	614	530	269	1,313	764	5.8	195	247	316	3.5	1.8	7,833
1998	449	170	611	509	255	1,293	733	4.8	187	234	316	3.5	1.8	7,833
1999	449	165	600	499	220	1,239	651	4.2	194	244	316	3.5	1.8	7,833
2000	428	167	589	500	202	1,146	566	4.2	198	234	316	3.5	1.8	7,917
2001	418	166	573	49 4	212	1,089	586	3.9	208	244	316	3.5	1.8	8,708
2002	417	169	553	499	208	1,096	574	3.4	197	229	316	3.5	1.8	8,833
2003	403	165	527	512	204	1,127	567	3.9	210	238	316	3.7	1.8	8,668
2004	404	172	539	514	192	1,095	528	3.1	220	246	316	3.7	1.8	8,752
2005	393	177	527	508	185	1,085	538	2.7	222	249	323	3.7	1.8	8,387
2006	388	178	530	496	184	1,002	492	2.6	244	262	331	3.7	1.8	8,892
2007	370	186	516	489	179	1,015	480	2.5	242	267	339	3.7	1.8	8,925
2008	357	196	513	492	167	974	465	2.4	251	273	347	3.7	1.8	8,975
2009	357	192	502	488	158	943	426	2.4	254	287	355	3.7	1.8	8,533
2010	348	197	513	479	154	937	427	2.3	273	292	363	3.7	1.8	8,565
2011	346	196	495	475	151	901	429	1.9	297	326	363	3.7	1.8	8,965

Most data from the Farm register, Swedish Board of Agriculture and Statistics Sweden. (*) Estimated number of horses 2010, by Statistics Sweden. (**) Swedish Poultry Meat Association. (***) Data on goats were available until 1992, this data have been extrapolated. (****) Between 1995 and 1996 there was an increase in number of sows by 13 %. The reason for this sudden increase is that as from this year also uncovered gilts are included in this group. (*****) Including a small fraction of turkeys.

6.3 Manure Management (CRF 4.B)

6.3.1 Source category description

This category includes emissions of methane and nitrous oxide from manure management. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 6-8.

Table 6-8. Summary of source category description, CRF 4.B.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
4.B	CH ₄				T1, T2	CS, D	Yes
	N ₂ O	X	X		T1, T2	CS, D	Yes

CS Country Specific. T1 Tier 1. T2 Tier 2. D Default.

6.3.2 Methodological issues

Statistics on manure management and the use of manure and fertilisers are collected biannually by Statistics Sweden¹⁴⁴. Data on stable periods (Table 6-9) and manure management systems (Table 6-10, Table 6-11 and Table 6-12) originate from this survey. Since dairy cows are often stabled at night and spend time in the stables during milking, the data on stable periods for this animal category is combined with an assumption that 38 % of its manure was produced in the stable during the grazing period. The Swedish Board of Agriculture publishes data on manure production from cattle and swine as well as on nitrogen production from most of the animal subgroups included in the inventory¹⁴⁵. Much of the data are from a model called STANK in the MIND. It is primarily a tool for environmental fertiliser and manure guidance to farmers. The program can be used for nutrient balances and contains a database with e.g. nitrogen production. The data is based on various sources, mostly from public reports made by the Swedish Board of Agriculture, but expert judgements also occur. A complete reference list is found in the manual to the program. As milk production has increased during the reporting period nitrogen excretion is calculated from milk data. How manure and nitrogen vary with milk production is showed in Table 6-13. The values for manure production and nitrogen production in each of the other animal groups are given in Table 6-14 and Table 6-15, respectively. The data for manure and nitrogen production are estimated values per head, year and box. That is, the production if one hypothetically assumes that an animal of a specific animal category would stay in the box for a year. Total yearly production per animal category is then estimated by multiplying these values with number of animal in mid-June according to the farm register. Naturally the number of animals

¹⁴⁴ Statistics Sweden, MI 30-series.

¹⁴⁵ Swedish Board of Agriculture, 1993; and Swedish Board of Agriculture, 2001; Swedish Board of Agriculture, 1995; Swedish Board of Agriculture, 2000. The given values are calculated according to the model STANK – the official model for input/output accounting on farm level in Sweden (Linder, 2001).

in the different categories/age classes will vary somewhat during a year. i.e. piglets will grow and enter the next growth class. However, we use the value from the farm register as an approximation of the annual average. Due to more intense swine production, the yearly production for sows and pigs for meat production were updated in 2001. When no default emission factor exists for reindeer, we use the same value as for sheep. All emission factors used in the calculations are presented in Table 6-16.

Table 6-9. Stable periods for cattle, months.

Year	Dairy cows	Beef cows	Steers and bulls	Heifers	Calves	Sheep, horses, goats	Reindeer	Poultry, Swine
1990	7.2	6.2	7.6	6.5	7.8	6	0	12
1995	7.2	6.2	7.6	6.5	7.8	6	0	12
2000	7.2	5.8	7.6	6.1	7.6	6	0	12
2005	(*) 6.9	(*) 5.4	(*) 7.9	(*) 5.5	(*) 7.3	6	0	12
2006	6.9	5.4	7.9	5.5	7.3	6	0	12
2007	(*) 7.2	(*) 5.2	(*) 8.3	(*) 5.7	(*) 8.0	6	0	12
2008	7.2	5.2	8.3	5.7	8.0	6	0	12
2009	(*) 7.1	(*) 5.5	(*) 8.7	(*) 5.8	(*) 8.1	6	0	12
2010	7.1	5.5	8.7	5.8	8.1	6	0	12
2011	7.3	5.5	8.6	5.9	8.0	6	0	12

(*) Statistics Sweden. Other values are standard values, or extrapolated.

Table 6-10. Waste management systems, fraction of liquid systems.

Year	Dairy cattle	Other cattle	Pigs for meat production	Other swine	Sheep, goats, horses, reindeer	Poultry
1990	0.23	0.17	0.44	0.44	0	0.25
1991	0.23	0.17	0.44	0.44	0	0.25
1992	0.23	0.17	0.44	0.44	0	0.25
1993	0.29	0.22	0.58	0.58	0	0.25
1994	0.29	0.22	0.58	0.58	0	0.25
1995	0.31	0.23	0.63	0.63	0	0.25
1996	0.31	0.23	0.63	0.63	0	0.25
1997	(*) 0.33	(*) 0.16	(*) 0.8	(*) 0.24	0	0.25
1998	0.33	0.16	0.8	0.24	0	0.25
1999	(**) 0.39	(**) 0.14	(**) 0.82	(**) 0.26	0	0.25
2000	0.39	0.14	0.82	0.26	0	0.25
2001	(***) 0.44	(***) 0.15	(***) 0.86	(***) 0.31	0	0.25
2002	0.44	0.15	0.86	0.31	0	0.25
2003	(****) 0.46	(****) 0.14	(****) 0.88	(****) 0.38	0	0.25
2004	0.46	0.14	0.88	0.38	0	0.25
2005	(*****) 0.50	(*****) 0.15	(*****) 0.87	(*****) 0.33	0	0.25
2006	0.50	0.15	0.87	0.33	0	0.25
2007	(*****) 0.55	(*****) 0.14	(*****) 0.94	(*****) 0.47	0	0.25
2008	0.55	0.14	0.94	0.47	0	0.25
2009	(*****) 0.58	(*****) 0.18	(*****) 0.94	(*****) 0.61	0	0.25
2010	0.58	0.18	0.94	0.61	0	0.25
2011	(*****) 0.62	(*****) 0.18	(*****) 0.95	(*****) 0.47	0	0.25

(*) Statistics Sweden, 1998. (**) Statistics Sweden, 2000b. (***) Statistics Sweden, 2002b. (****) Statistics Sweden, 2004. (*****) Statistics Sweden, 2006. (*****) Statistics Sweden 2008, (*****) Statistics Sweden 2010, (*****) Statistics Sweden 2012. Other values are standard values, or interpolated /extrapolated.

Table 6-11. Waste Management Systems, fraction of solid systems.

Year	Dairy cattle	Other cattle	Pigs for meat production	Other swine	Sheep, goats	Reindeer	Horses	Poultry
1990	0.52	0.32	0.54	0.45	0.5	0	0.48	0.55
1991	0.52	0.32	0.54	0.45	0.5	0	0.48	0.55
1992	0.52	0.32	0.54	0.45	0.5	0	0.48	0.55
1993	0.46	0.27	0.40	0.31	0.5	0	0.48	0.55
1994	0.46	0.27	0.40	0.31	0.5	0	0.48	0.55
1995	0.44	0.26	0.35	0.26	0.5	0	0.48	0.55
1996	0.44	0.26	0.35	0.26	0.5	0	0.48	0.55
1997	0.41	0.33	0.17	0.65	0.5	0	0.48	0.55
1998	0.41	0.33	0.17	0.65	0.5	0	0.48	0.55
1999	0.35	0.31	0.18	0.67	0.5	0	0.48	0.55
2000	0.35	0.31	0.18	0.67	0.5	0	0.48	0.55
2001	0.31	0.26	0.13	0.58	0.5	0	0.48	0.55
2002	0.31	0.26	0.13	0.58	0.5	0	0.48	0.55
2003	0.27	0.24	0.11	0.51	0.5	0	0.48	0.55
2004	0.27	0.24	0.11	0.51	0.5	0	0.48	0.55
2005	0.23	0.20	0.05	0.44	0.5	0	0.48	0.55
2006	0.23	0.20	0.05	0.44	0.5	0	0.48	0.55
2007	0.20	0.22	0.06	0.41	0.5	0	0.48	0.55
2008	0.20	0.22	0.06	0.41	0.5	0	0.48	0.55
2009	0.16	0.20	0.05	0.32	0.5	0	0.48	0.55
2010	0.16	0.20	0.05	0.32	0.5	0	0.48	0.55
2011	0.13	0.19	0.05	0.41	0.5	0	0.48	0.55

Table 6-12. Waste management systems, fraction of deep litter systems (categorised as “other” in the CRF-tables).

Year	Dairy cattle	Other cattle	Pigs for meat production	Other swine	Sheep, goats, reindeer	Horses	Poultry
1990	0.01	0.08	0.02	0.11	0	0.02	0.20
1991	0.01	0.08	0.02	0.11	0	0.02	0.20
1992	0.01	0.08	0.02	0.11	0	0.02	0.20
1993	0.01	0.08	0.02	0.11	0	0.02	0.20
1994	0.01	0.08	0.02	0.11	0	0.02	0.20
1995	0.01	0.08	0.02	0.11	0	0.02	0.20
1996	0.01	0.08	0.02	0.11	0	0.02	0.20
1997	0.01	0.08	0.02	0.11	0	0.02	0.20
1998	0.01	0.08	0.02	0.11	0	0.02	0.20
1999	0.01	0.09	0.01	0.07	0	0.02	0.20
2000	0.01	0.09	0.01	0.07	0	0.02	0.20
2001	0.00	0.12	0.01	0.12	0	0.02	0.20
2002	0.00	0.12	0.01	0.12	0	0.02	0.20
2003	0.01	0.14	0.01	0.11	0	0.02	0.20
2004	0.01	0.14	0.01	0.11	0	0.02	0.20
2005	0.01	0.16	0.01	0.22	0	0.02	0.20
2006	0.01	0.16	0.01	0.22	0	0.02	0.20
2007	0.01	0.17	0.00	0.12	0	0.02	0.20
2008	0.01	0.17	0.00	0.12	0	0.02	0.20
2009	0.01	0.16	0.01	0.07	0	0.02	0.20
2010	0.01	0.16	0.01	0.07	0	0.02	0.20
2011	0.01	0.17	0.00	0.12	0	0.02	0.20

Table 6-13. Manure and nitrogen production from dairy cows.

Animal groups	Manure kg dm/day/head	Nitrogen kg/year/head
Dairy Cows (Milk production 6,000 kg/yr)	5.75	97
Dairy Cows (Milk production 8,000 kg/yr)	6.07	117
Dairy Cows (Milk production 10,000 kg/yr)	6.19	137

Table 6-14. Manure production from other animal groups.

Animal groups	Manure production, kg dm/day
Beef cows (*)	2.64 (in stable); 3.64 (during grazing)
Growing animals (12-24 months)	2.6
Calves > 6 months	1.12
Calves < 6 months	0.69
Sows	0.74 (1990-2001); 0.793 (as from 2002)
Boars	0.52
Pigs for meat production	0.42
Piglets	0.05

Swedish Board of Agriculture, 1993. Swedish Board of Agriculture, 1995. Swedish Board of Agriculture, 2001.

Table 6-15. Nitrogen production from other animal groups.

Animal groups	Nitrogen kg/ year/head, 1990-2001	Comment	Updated values on nitrogen prod. used from 2002, kg/ year/ head	Comment
Beef cows	63			
Growing animals 12-24 months	47			
Calves > 6 months	28			
Calves < 6 months	28			
Sows	18.5		22.5	
Boars	13			
Pigs for meat production	9.5	2.5 prod. cycles/ year	10.8	3 prod. cycles / year
Piglets	0.5		0.5	
Sheep	13	Ewes incl. 1.5 lambs		
Lambs	0			
Goats	13	Does incl. 1.5 kids		
Kids	0			
Horses	50	Mean value for all animals		
Laying hens and turkeys	0.64			
Chickens	0.28	2.5 prod. cycles/ year		
Slaughter Chickens	0.29	6.5 prod. cycles/ year		
Reindeer*	10			

Values are calculated according to the STANK IN MIND model (Swedish Board of Agriculture)

* Data from Statistics Finland

Table 6-16. Emission factor manure management

Manure management	Emission factor for CH ₄	Note
MFC Solid manure (*)	1 % of B ₀	1
MFC Liquid manure (*)	3.5 % of B ₀	3
MFC Deep litter (*)	39 % of B ₀	1
MFC Pasture/Range/Paddock (*)	1 % of B ₀	1
Dairy Cattle - volatile solid (VS)	1,947 kg VS/animal/yr	2
Dairy Cattle - B ₀ (**)	0.24 m ³ CH ₄ /kg VS	1
Dairy Cattle - Emission per animal	8.9 kg CH ₄ /animal/yr	2
Non-Dairy Cattle – volatile solid (VS)	528 kg VS/animal/yr	2
Non-Dairy Cattle – B ₀	0.17 m ³ CH ₄ /kg VS	1
Non-Dairy Cattle – emission/animal (***)	5.6 kg CH ₄ /animal/yr	2
Swine – volatile solids (VS)	112 kg VS/animal/yr	2
Swine - B ₀ **	0.45 m ³ CH ₄ /kg VS	1
Swine – emission per animal (***)	1.44 kg CH ₄ /animal/yr	2
Sheep – emission	0.19 kg CH ₄ /animal/yr	1
Goats – emission	0.12 kg CH ₄ /animal/yr	1
Horses – emission	1.40 kg CH ₄ /animal/yr	1
Poultry – emission	0.08 kg CH ₄ /animal/yr	1
Reindeer – emission (assuming same as sheep)	0.19 kg CH ₄ /animal/yr	1
Manure management	Emission factor for N ₂ O	Note
Waste Management System	% N ₂ O-N of N-supply	
Liquid manure	0.1	1
Solid manure	2	1
Deep litter	2	1

(*)MCF = Methane Conversion Factor. (**) B₀ = maximum methane producing capacity for manure. (***) Weighted value – more than one animal category. (1) IPCC Guidelines (2) Calculated. (3) National, Rodhe et al. 2008.

6.3.2.1 4.B(A). METHANE (INCLUDING EXCRETION FROM GRAZING ANIMALS)

The Good Practice Guidance Tier 2 methodology for estimating methane from manure management, including excretions from grazing animals, is applied for cattle and swine, and the corresponding Tier 1 methodology is used for other animal groups¹⁴⁶. The formula for the emission factor for livestock group “i”, according to the Good Practice Guidance Tier 2 methodology is:

$$emissionfactor_i = VS_i * B_{0i} * 0.67 * \sum_{jk} MCF_{jk} * MS_{ijk}$$

where VS_i is the volatile substance excreted per year, B_{0i} is the maximum methane producing capacity for manure produced by an animal within the livestock group, MCF_{jk} is a conversion factor for methane production, given

¹⁴⁶ According to current estimations, cattle and swine produce about 85-90% of the total methane emissions from manure management.

a manure management system j (where grazing animals are considered as one of the systems), and a climate region k . MS_{ijk} is the fraction of animal manure handled using manure system j in climate region k .

The B_{0i} and MCF factors used are the default values in the Good Practice Guidance, except for the revised MCF for liquid manure, where the value of 3.5 % is used. This value was developed by Rodhe et al., 2008 and is considered to be more appropriate for Swedish conditions. The values reported in the CRF tables are sometimes aggregated after the calculation has been carried out for more specific animal groups. Hence the implied emission factor for “other cattle” will depend not only on different manure management systems and stable periods over the years, but also on the relative composition of the different subgroups. The implied emission factor therefore varies between the reported years.

The Swedish Board of Agriculture provides data from a national database on manure production from cattle and swine¹⁴⁷.

Information on waste management systems is collected from the surveys published in the biannual statistical report on the use of fertilisers and animal manure in agriculture¹⁴⁸ and the interpolated values are used for the intermediate years. Three manure management systems are considered apart from grazing animals: liquid systems (Table 6-10), solid storage (Table 6-11) and deep litter (Table 6-12) (sometimes categorised as “other” in the national inventory). National estimates of stable periods for cattle are collected from the statistical report on use of fertilisers and animal manure in agriculture¹⁴⁹. This information has been available biannually since 1997. Before 1997, the data are extrapolated to 1990.

When looking on the trends for the implied emission factor (IEF) there is a distinct increase for both non-dairy and dairy cattle. For non-dairy cattle this is caused by a decreased use of solid manure systems and an increased use of deep litter systems, which has a higher methane conversion factor. For dairy cattle the use of solid systems has also decreased but here the use of liquid systems has increased instead.

6.3.2.2 4.B(B). NITROUS OXIDE

The methodology for estimating N_2O from manure management is in accordance with the IPCC Guidelines Tier 2 methodology; it is based on emission factors from the IPCC Guidelines in combination with national activity data. The emission of N_2O from manure management is calculated as:

$$emission = \sum_S \left(\sum_T N_T \times Nex_T \times MS_{(T,S)} \right) \times EF_s \times 44 / 28$$

¹⁴⁷ Swedish Board of Agriculture, 1993. Swedish Board of Agriculture 1995. Swedish Board of Agriculture 2001. The given values are calculated according to the model STANK – “Stallgödselnäring i kretslopp” the official model for input/output accounting on farm level in Sweden (Linder, 2001). STANK is currently being evaluated in a study launched by The European Commission.

¹⁴⁸ Statistics Sweden, MI 30-series.

¹⁴⁹ Statistics Sweden, MI 30-series.

where is the number of heads of livestock in category T in the country, is the annual average excretion of N per head of category T in the country, $MS_{(T,S)}$ is the fraction of total annual excretion for each livestock category T managed in manure management system S in the country (including pasture, range and paddock manure). Even though dairy cattle generally spends the summer on pasture, part of the manure is still produced in the stable during milking and when spending the nights in the stable. This manure is excluded from the pasture, range and paddock manure and instead allocated to a manure system. Data on nitrogen and manure production has been derived by the Swedish Board of Agriculture (Table 6-13, Table 6-14 and Table 6-15). Stable period and manure management systems are the same as used in the methane calculations (Table 6-9, Table 6-10, Table 6-11 and Table 6-12).

The emission factors are described in Table 6-17. In the CRF tables, where some animal subgroups are aggregated, the implied emission factors (IEFs) may change over the years, depending on the relative size of the respective subgroups aggregated.

6.3.3 Uncertainties and time-series consistency

Due to more intense swine production, the nitrogen production for sows and pigs for meat production were updated in 2002.

6.3.4 Source-specific QA/QC and verification

Times series and activity data are checked for consistency. Recalculations are identified and verified. Emission estimates and methods are reviewed yearly by external national experts.

6.3.5 Source-specific recalculations

A smaller error in the manure production rate for grazing beef cows was corrected.

6.3.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

6.4 Agricultural Soils (CRF 4.D)

Emissions of nitrous oxide from agricultural soils are presented under CRF 4.D.1 to CRF 4.D.4. When the subsectors represent relatively different processes they are divided in separate paragraphs and also treated independently in the key categories analyses. Table 6-17 gives an overview of all emission factors used in this sector.

Table 6-17. Emission factor for N₂O from soils

Direct emissions from soils	Emission factor % N ₂ O-N of N-supply	Note
Mineral fertiliser	0.8	1
Manure	2.5	1
Crop residue	1.25	2
N-fixing Crops	1.25	2
Manure during grazing	2	2
Background emission due to cultivation	Kg N₂O-N/ha/yr	
Cultivation of Histosols	8	2
Cultivation of Mineral Soils	0.5	1
Indirect emissions from soils		
Atmospheric Deposition	1 % of emitted N	2
Nitrogen Leaching and run-off	2.5 % of N lost from leaching	2

(1) National, Klemedtsson, 2001. (2) IPCC Guidelines.

6.4.1 Direct Soil Emissions (CRF 4.D.1)

6.4.1.1 SOURCE CATEGORY DESCRIPTION

The category includes the direct emission of nitrous oxide from soils. In terms of magnitude the most important emissions are from animal manure applied to soils followed by cultivation of histosols and from the use of synthetic fertilisers, respectively. Also included in this category are emissions from nitrogen fixing crops and crop residues. In this category Sweden also includes emission from sewage sludge used as fertilisers. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 6-18.

Table 6-18. Summary of source category description for the entire category CRF 4.D.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
4.D.1	N ₂ O	X	X		CS, T1, T1a, T1b, T2	CS, D	Yes

CS Country Specific. T1 Tier 1. T2 Tier 2. D Default.

6.4.1.2 METHODOLOGICAL ISSUES

6.4.1.2.1 Emission factors

For estimating direct soil emissions, the Good Practice Guidance encourages parties to use country-specific emission factors for N₂O from agricultural fields, where possible. A suggested alternative is to use factors from other countries with comparable management and climatic conditions. In order to update the information from research, a literature study was carried out, requested by the Swedish EPA¹⁵⁰. The study includes documented N₂O emis-

¹⁵⁰ Klemedtsson, 2001.

sion measurements carried out in Sweden and in other countries in northern Europe and in Canada. The results show that there is limited data available. It was for example not possible to develop different emission factors for added nitrogen to mineral or organic soils. Best data availability was found for the use of synthetic fertilisers on mineral soils. Here the emissions of N_2O -N were between 0 and 0.8 % of added nitrogen. However, when the emissions are caused by other processes than newly added fertiliser, a correlation between added nitrogen and emitted nitrogen was not always apparent. For example, the amount of fertiliser used on cereals did not affect the magnitude of emitted N_2O . Thus, a method was suggested where the emissions are divided into two parts, one is dependent on fertiliser use and the other is a constant background emission. When the use of synthetic fertiliser on grasslands increased the emissions with some 0.8 % of added nitrogen an emission factor of 0.8 % was suggested for the use of synthetic fertiliser on all agricultural land. A lower value than the IPCC Good Practice Guidance default value of 1.25 % for mineral fertiliser nitrogen has also been suggested in a synthesis of literature data¹⁵¹. The background emission from mineral soils was suggested to be 0.5 kg N_2O -N/ha/year (section 6.4.4). For the use of animal manure the study concluded that there are even less data available and the emissions according to different surveys varied between 0.6 % and 8 % of added nitrogen. Using a regression model an emission factor of 2.5 % was suggested. The study also suggested country specific emission factors for background emission from organic soils and from indirect emissions of N_2O from atmospheric deposition and nitrogen leaching. These emission factors were however not adopted in the inventory as they were considered to be too unreliable. Instead IPCC default factors were used. A summary of all emission factors applied for 4.D is given in Table 6-17.

6.4.1.2.2 *Emission of ammonia*

The calculations of ammonia emissions from the agricultural sector are mainly built on data collected through Statistics Sweden's field investigation among farmers. The calculation methods have been developed by the Swedish EPA and Statistics Sweden in collaboration with the Swedish Board of Agriculture and the Swedish Institute of Agricultural and Environmental Engineering¹⁵². The calculations have been made by Statistics Sweden since 1990 at national and regional levels. Regional results are published from 2005 at the web-site of Statistics Sweden¹⁵³. However, due to a change in a questionnaire in Statistics Sweden's field investigation among farmers and updated methods for calculating the emissions, the results from 1990-1994 are not compatible with these from 1995-2007. For that reason we use the value from 1995 for 1990 through 1994.

¹⁵¹ Lægveid and Aastveit, 2002.

¹⁵² Swedish Environmental Protection Agency 1997

¹⁵³ Statistics Sweden 2007

In short the calculations are made as follows:

$$A = (V + L + S)$$

$$V = D \times N \times P \times F(v)$$

$$L = D \times N \times P \times (1 - F(v)) \times F(l)$$

$$S = D \times N \times P \times (1 - F(v)) \times (1 - F(l)) \times F(s)$$

A = emission of nitrogen in ammonia

V = emission of nitrogen through stable ventilation (depending on type of handling, type of animal and type of manure)

L = emission of nitrogen during storing (depending on type of manure, storing method and type of animal)

S = emission of nitrogen during spreading (depending on type of manure, time of spreading, method of spreading and time period between spreading and mulching)

D = number of animal¹⁵⁴

N = production of nitrogen, kg, per type of animal, year and handling¹⁵⁵

P = stable period¹⁵⁶

F(v) = emission of nitrogen through stable ventilation, % of total nitrogen content in stable manure¹⁵⁷.

F(l) = emission of nitrogen during storing, % of total nitrogen content in stable manure after ventilation losses¹⁵⁸.

F(s) = emission of nitrogen during spreading, % of ammonium nitrogen content in stable manure after ventilation and storing losses¹⁵⁹.

The calculated data is differentiated by type of animal, type and handling of manure, milk production, time and method of spreading and time period between spreading and mulching. Type of manure, way of storing and time of spreading etc. are estimated from the field investigation among farmers¹⁶⁰. Ventilation-, storage- and spreading-losses originate from a data calculating program (called STANK IN MIND) from Swedish Board of Agriculture and from Swedish Institute of Agricultural and Environmental Engineering, Table 6-20 and Table 6-21 give an overview of the emission factors used in the calculations.

¹⁵⁴ Swedish Board of Agriculture 2008 and other sources

¹⁵⁵ Swedish Board of Agriculture 1995; Swedish Board of Agriculture 2000; Swedish Board of Agriculture 2001

¹⁵⁶ Statistics Sweden 2008

¹⁵⁷ Swedish Board of Agriculture 2005

¹⁵⁸ Swedish Institute of Agricultural and Environmental Engineering 2002

¹⁵⁹ Swedish Institute of Agricultural and Environmental Engineering 2002

¹⁶⁰ Statistics Sweden, 2008

Table 6-19. Nitrogen losses caused by ventilation in stables, % of Total-N.

Type of animal	Solid-manure	Deep litter	Liquid manure	Semisolid manure	Urine
Cattle	4	20	4	4	4
Swine	10	25	14	10	10
Laying hens	10	35	10		
Chickens	10	20	10		
Slaughter chickens			10		
Horses	4	15			
Sheep	4	15			

Table 6-20. Nitrogen losses caused by ammonia emission during storage of manure, % of total-N.

Type of manure, handling	Type of animal					
	Cattle	Swine	Laying hens/ chickens	Slaughter chickens	Horses	Sheep
Solid manure	20	20	12		25	25
Semisolid manure	10	10				
Liquid manure, uncovered						
Filled from underneath	6	8	8			
Filled from above	7	9	9			
Liquid manure, covered						
Filled from underneath:						
Roof	1	1	1			
floating crust	3	4	4			
other	2	2	2			
Filled from above:						
roof	1	1	1			
floating crust	4	5	5			
other	3	3	3			
Urine, uncovered						
Filled from underneath	37	37				
Filled from above	40	40				
Urine, with cover						
Filled from underneath:						
roof	5	5				
floating crust	17	17				
other	10	10				
Filled from above:						
roof	5	5				
floating crust	20	20				
other	12	12				
Deep litter manure	30	30	20	5		33

Table 6-21. Nitrogen losses caused by ammonia emission during spreading of manure (% of total-N).

Season/ Spreading method	Spreading strategy and tillage timing	Solid manure	Urine	Liquid manure
Early spring/late winter				
Broadcast	Spread on frozen ground	20	40	30
Trailing hoses			30	20
Spring				
Broadcast	Immediately	15	8	10
	Mulching within 4 h	33	14	15
	Mulching within 5-24 h	50	20	20
	Spread on pasture	70	35	40
	Spread on grain		11	20
Trailing hoses	Immediately		7	5
	Mulching within 4 h		14	8
	Mulching within 5-24 h		20	10
	Spread on pasture		25	30
	Spread on grain		10	15
Shallow injection	Spread on pasture		8	15
Early summer, summer				
Broadcast	Spread on pasture	90	60	70
	Spread on grain		10	20
Trailing hoses	Spread on pasture		40	50
	Spread on grain		10	7
Shallow injection	Spread on pasture		15	30
Early autumn				
Broadcast	Immediately	20	15	5
	Mulching within 4 h	35	23	18
	Mulching within 5-24 h	50	30	30
	No mulching	70	45	70
Trailing hoses	Immediately		10	3
	Mulching within 4 h		18	9
	Mulching within 5-24 h		25	15
	No mulching		30	40
Late autumn				
Broadcast	Immediately	10	10	5
	Mulching within 4 h	15	15	8
	Mulching within 5-24 h	20	20	10
	No mulching	30	25	30
Trailing hoses	Immediately		4	3
	Mulching within 4 h		11	4
	Mulching within 5-24 h		18	5
	No mulching		25	15

6.4.1.2.3 4.D.1.1 Nitrous oxide from synthetic fertilisers

Emissions from fertilisers are calculated as:

$$emissions = N_{FERT} \times (1 - Frac_{GASF}) \times EF \times 44 / 28$$

where N_{FERT} is the total amount of fertiliser nitrogen consumed annually, and $Frac_{GASF}$ is the fraction that volatilises as ammonia. Statistics on sales of fertilisers, recalculated into nitrogen quantities, are published annually by Statistics Sweden¹⁴⁹. The estimated emissions are based on amount of nitrogen in mineral fertilisers sold in Sweden excluding the nitrogen lost as ammonia (Table 6-22). The proportion of nitrogen lost as ammonia ($Frac_{GASF}$) differs between different types of fertilisers. The values used are from the EMEP/EEA emission inventory guidebook 2009 and calculated using the mean spring temperature of 5.9 degrees centigrade (table 6-23). In Table 6-22 sold quantities of ammonia-emitting products are shown, which directly explains variations in the $Frac_{GASF}$ between different years.

Table 6-22. Sold quantity of ammonia emitting fertilisers and nitrogen in sludge used as fertilizers.

Year	N in sold fertilisers, tonnes	Ammonium Nitrate, AXAN, N26, N27, N28, tonnes of product	N-solution, tonnes of product	Urea, tonnes of product	NPK, tonnes of N	NP, tonnes of N	NK, tonnes of N	Proportion of emitted fertiliser-N ($Frac_{GASF}$)	Sludge, tonnes of N.
1990	224,500	225,387	10,089	5,932	64,600	11,000	0	0.0,084	1,180
1991	208,600	237,612	6,498	4,683	52,100	11,000	3,700	0.0,082	1,180
1992	178,400	179,234	8,837	2,980	45,400	8,500	3,000	0.0,083	1,180
1993	207,200	200,004	5,257	3,501	46,100	9,800	3,300	0.0,083	1,180
1994	216,400	167,150	7,820	3,061	55,900	12,300	3,000	0.0,087	1,433
1995	198,300	182,486	11,193	1,955	51,050	13,451	2,912	0.0,089	2,304
1996	192,300	158,613	5,949	1,474	48,000	14,000	2,500	0.0,078	2,304
1997	204,600	175,558	4,399	1,104	51,500	15,900	2,300	0.0,076	2,304
1998	205,600	209,463	2,631	889	53,723	14,286	2,033	0.0,076	2,027
1999	179,200	166,077	3,111	745	50,092	14,619	1,746	0.0,079	2,027
2000	189,400	205,869	3,772	655	51,600	11,400	2,200	0.0,075	1,758
2001	196,900	235,495	2,036	553	54,000	11,300	3,000	0.0,075	1,171
2002	184,800	198,981	1,641	484	54,900	10,900	2,300	0.0,078	593
2003	180,100	238,828	1,083	382	53,900	10,600	2,200	0.0,077	692
2004	176,800	240,553	4,928	475	54,500	11,900	1,800	0.0,083	796
2005	161,500	273,036	3,364	519	59,000	8,400	1,600	0.0,092	1,053
2006	160,300	267,754	3,164	225	57,800	8,500	1,800	0.0,091	1,322
2007	167,100	285,064	0	271	61,100	5,300	2,000	0.0,087	1,322
2008	186,500	360,415	13	235	71,029	3,931	1,805	0.0,091	2,481
2009	142,400	298,882	67	1,088	43,600	2,000	1,100	0.0,094	2,205
2010	168,000	348,691	0	1,002	52,800	2,600	1,400	0.0,091	2,224
2011	169,800	314,225	0	1,090	56,600	1,300	900	0.0,088	2,224

Statistics on fertilisers from Swedish Board of Agriculture, 2011 and Statistics Sweden, 2011.

(*) Statistics Sweden 1997b and Statistics Sweden 2001. (**) from Statistics Sweden 1992 and Statistics Sweden 1995. Other values are expert judgements. (***) The decrease in 2009 is due to an overconsumption in 2008 due to a dropped tax on fertilisers

Table 6-23. Fraction of nitrogen for different fertiliser types that is lost as ammonia

Fertiliser	Lost as ammonia (% of N)
Ammonium Nitrate, AXAN, N26, N27, N28	0.9
N-solution	6.3
Urea	12.7
NPK	0.9
NP	1.4
NK	0.9
Sludge	30

6.4.1.2.4 4.D.1.2 Nitrous oxide from animal manure applied to soils

To calculate the N₂O from animal manure, the default methodology according to the IPCC Guidelines is used combined with national estimates of N content in manure (section 6.3.2) and a national estimation of ammonia-N emissions. The formula is:

$$emissions = \sum_T N_T \times Nex_T \times (1 - Frac_{GASM}) \times (1 - Frac_{PRP}) \times EF \times 44 / 28$$

Where T is a category of livestock. Frac_{GASM} is a national value for the fractions nitrogen that is lost as ammonia-N (Table 6-24). Frac_{GASM} is estimated by Statistics Sweden and the Swedish EPA¹⁶¹. The estimates are model-based and take into account many factors that influence gas emissions (see 6.4.1.2.2).

Table 6-24. Ammonia-N emissions from manure, fraction.

	1995	1997	1999	2001	2003	2005	2007	2009	2010	2011
Stable manure (FracGASM)	0.33	0.33	0.33	0.33	0.33	0.32	0.33	0.33	0.33	0.33
Manure from grazing animals ("FracGASG")	0.12	0.08	0.08	0.08	0.08	0.08	0.09	0.09	0.09	0.09

Statistics Sweden, MI 37-series.

6.4.1.2.5 4.D.1.3 N₂O from N-fixing crops in pure stands and in temporary grass

Nitrogen fixation crops are pasture grounds with features of clover, leguminous crops (cooking and fodder peas, preserved peas, vetches, field beans etc). This nitrogen fixation by leguminous plants is a part of the nitrogen circulation in agricultural soils and the corresponding N₂O emissions are included in the inventory. Data derives from national estimates of nitrogen fixation, which account for regional differences, in combination with the Good Practice Guidance's default emission factor for direct N₂O emissions.

The formula is:

$$emissions = \sum_{crop, county} Area_{crop, county} \times NfixingAmount_{crop, county} \times EF \times 44 / 28,$$

¹⁶¹ Statistics Sweden, MI 37-series.

The total production of the respective crops is given by multiplying the cultivated area, according to the Farm Register, by standard yield. The reason for using standard yields instead of actual yields in the calculations is that the time series becomes more consistent and not drastically effected by stochastic events like extreme weather conditions. Estimated standard yields for different crops are published annually by the Swedish Board of Agriculture/Statistics Sweden and are a function of crop yields estimated by surveys conducted over the last 15 years¹⁶² (Table 6-29 and Table 6-30).

Areas are given in Table 6-26, Table 6-27 and Table 6-28. To estimate nitrogen fixation from the atmosphere, a model according to Høgh-Jensen has been used since submission 2006¹⁶³. The model covers fixation from root and stubble as well as transmission to other plants. It has been adapted to account for Swedish conditions¹⁶⁴ and has also been used by others such as the Swedish Board of Agriculture. According to the model the amount of fixed nitrogen is estimated as a part of the total amount of nitrogen in the plant's biomass. This part varies depending on the kind of leguminous plant, the age of the pasture, the number of harvests and, to some extent, the fertilised amount of fertiliser applied.

6.4.1.2.6 4.D.1.4 N₂O from crop residue

For the estimation of N₂O from crop residues we also use standard yields instead of actual yields in the calculations. To estimate N₂O from nitrogen circulation in crop residues, the methodology recommended in the Good Practice Guidance is used combining national activity data on removed residues and other parameters, such as nitrogen content, at crop level with the Good Practice Guidance's default emission factor for direct N₂O emissions. The data on crop residues builds on a one-time study from 1997¹⁶⁵ on how straw and tops from different crops are used. The formula used for the calculations is:

$$emission = \sum_{crop} yield_{crop} \times area_{crop} \times Fracresidues_{crop} \times FracN_{crop} (1 - Fracresiduesremoved_{crop}) \times EF \times 44 / 28,$$

Yield is the standard yield, Fracresidues are the crop residues as a fraction of the harvest, FracN is the fraction of nitrogen in crop residues and Fracresiduesremoved is the fraction of crop residues that is removed according to a field survey¹⁶⁶ from 1997. When calculating N-circulation in residues from cereal crops, national factors for recalculation from harvest to crop residue and the corresponding N-content based on national measurement data are used¹⁶⁷. For other crops, a combination of national factors and IPCC default values was used¹⁶⁸. All factors used for calculating N input with crop residues

¹⁶² Statistics Sweden, 2002e.

¹⁶³ Høgh-Jensen et al. 2004.

¹⁶⁴ Frankow-Lindberg, 2005.

¹⁶⁵ Statistics Sweden, 1999.

¹⁶⁶ Statistics Sweden, 1999.

¹⁶⁷ Mattson, 2005.

¹⁶⁸ Adolfsson, R. 2005.

are given in Table 6-25. Areas of different crops used in the calculations are stated in Table 6-26, Table 6-27 and Table 6-28. Standard yield¹⁶⁹ of different crops used in the calculations is presented in Table 6-29 and table 6-30.

Table 6-25. Data used for calculating nitrogen input in crop residues.

Crop	Fraction of crop residues removed (ResiduesRemoved)	Fraction of N in crop residues in dry matter (FracN)	Fraction residues in relation to harvest, (FracResidues)	Dry matter content, fraction
Winter wheat	0.06	0.0051	0.87	0.85/0.86
Spring wheat	0.06	0.0044	0.96	0.85/0.86
Winter rye	0.09	0.0060	1.08	0.85/0.86
Winter barley	0.23	0.0051	0.87	0.85/0.86
Spring barley	0.12	0.0077	0.83	0.85/0.86
Oats	0.12	0.0073	0.89	0.85/0.86
Mixed grain	0.18	0.0067	0.98	0.85/0.86
Triticale	0.06	0.0060	1.08	0.85/0.86
Sugar beets	0.09	0.0225	0.66	0.85
Winter rape	0.02	0.0107	0.47	0.91
Spring rape	0.02	0.0107	0.47	0.91
Winter turnip rape	0.02	0.0107	0.47	0.91
Spring turnip rape	0.02	0.0107	0.47	0.91
Table potatoes and Potatoes for starch prod.	0	0.0110	0.40	0.20
Temporary grass	0	0.0130	0.25	0.84
Temporary grass for seed	0.49	0.0130	0.94	0.84
Green fodder	0	0.0130	0.25	0.84
Pasture ground	0	0.0130	0.40	0.67
Peas, Peas for fodder and brown beans	0.02	0.0142	1.50	0.85
Peas for conservation	0	0.0142	1.50	0.85

Adolfsson, R. 2005.

¹⁶⁹ Statistics Sweden, 2002e.

Table 6-26. Areas of different crops used in the calculations (hectares).

Year	Winter wheat	Spring wheat	Winter rye	Winter barley(*)	Spring barley	Oats	Mixed grain	Triticale(**)
1990	320,120	29,595	73,460	-	492,027	387,823	32,628	-
1991	225,330	33,387	43,239	-	490,896	364,272	40,337	-
1992	233,678	36,647	34,597	-	454,097	360,859	47,420	-
1993	271,818	32,581	46,390	-	420,437	321,961	35,330	35,330
1994	212,095	39,722	38,957	29,536	443,489	341,415	25,421	42,526
1995	222,304	39,076	39,693	26,220	427,115	278,322	27,124	44,577
1996	292,170	42,392	33,558	22,061	446,503	283,588	34,230	61,694
1997	299,594	44,588	29,416	15,272	467,628	315,465	30,247	66,473
1998	359,024	39,021	34,617	15,949	429,011	311,467	26,972	66,751
1999	209,641	65,777	24,507	11,883	470,104	305,658	33,022	32,586
2000	353,201	48,364	34,533	12,997	398,227	295,544	45,328	40,728
2001	354,495	44,670	34,403	9,577	387,922	278,174	25,370	39,642
2002	285,249	54,350	24,395	6,386	410,456	295,002	22,623	30,809
2003	364,058	47,290	24,366	6,345	362,127	279,808	25,235	44,661
2004	349,823	53,585	24,402	5,268	392,006	229,696	18,697	52,195
2005	295,325	59,430	21,386	5,356	373,208	200,122	18,857	50,292
2006	317,603	43,333	23,454	5,691	309,444	206,055	17,430	55,406
2007	323,182	38,367	24,716	8,274	318,407	207,909	15,317	53,914
2008	311,632	49,915	27,581	10,396	395,367	227,588	15,955	49,287
2009	326,838	48,297	36,633	18,278	351,878	196,038	17,050	53,571
2010	331,805	68,184	24,228	17,928	300,847	164,386	19,150	36,231
2011	349,793	66,986	24,094	14,370	313,459	181,170	18,676	24,215

Statistics from the Farm Register. (*) Before 1994, statistics on winter barley and spring barley revised as one crop. (**) Before 1993, statistics on Triticale was included in mixed grain.

Table 6-27. Areas of different crops used in the calculations (hectares).

Year	Sugar beets	Winter rape	Spring rape	Winter turnip rape	Spring turnip rape	Table potatoes	Potatoes starch prod.
1990	38,502	84,598	44,203	9,068	30,035	27,305	8,866
1991	47,963	75,724	41,046	8,089	26,362	28,269	8,807
1992	51,287	51,364	56,519	3,145	26,366	30,414	8,791
1993	51,287	74,460	46,203	2,455	22,370	27,815	8,469
1994	53,353	46,035	53,033	1,746	27,647	25,449	7,539
1995	57,518	56,084	23,311	1,587	23,661	27,630	7,371
1996	59,223	21,737	18,976	811	23,869	27,577	9,060
1997	60,459	22,888	19,475	1,787	19,432	26,732	9,081
1998	58,737	23,159	16,705	1,470	13,238	25,133	8,567
1999	59,881	19,626	31,273	1,206	23,784	24,422	8,391
2000	55,484	24,870	12,112	1,395	9,791	23,610	9,293
2001	54,834	19,900	13,591	857	10,425	23,776	8,460
2002	54,820	31,219	21,943	1,899	12,408	23,142	8,589
2003	50,100	23,352	26,670	817	7,734	21,923	8,617
2004	47,625	37,496	36,715	1,244	8,343	23,015	8,656
2005	49,182	34,997	38,578	1,460	7,116	22,081	8,372
2006	44,184	47,638	35,148	1,138	6,270	20,212	7,966
2007	40,682	50,341	33,044	1,117	3,341	20,330	8,032
2008	36,778	61,860	24,359	834	2,453	19,590	7,293
2009	39,782	67,841	29,245	282	2,150	19,706	7,252
2010	37,950	71,836	35,695	496	2,207	19,842	7,361
2011	39,638	56,600	36,112	395	1,781	20,046	7,616

Table 6-28. Areas of different crops used in the calculations (hectares).

Year	Temporary grass	Temporary grass for seed	Green forage	Pasture ground	Peas incl fodder	Peas for conser- vation (*)	Brown beans (*)	Total area arable land (**)
1990	727,590	10,753	39,698	190,503	32,742	-	-	3,103 141
1991	696,069	10,418	33,509	239,818	23,327	-	-	3,094 792
1992	708,384	8,791	2,896	292,825	14,059	-	-	3,087 260
1993	748,094	7,863	23,137	314,458	8,720	-	-	3,076 156
1994	757,000	8,241	23,000	314,666	6,598	-	-	3,071 416
1995	766,776	7,907	23,695	276,927	11,959	8,578	709	3,066 027
1996	750,085	7,854	22,268	247,369	17,713	8,821	690	3,060 329
1997	746,832	8,470	24,443	234,677	32,742	9,028	921	3,056 259
1998	742,068	9,013	21,935	221,418	49,150	8,524	938	3,051 159
1999	760,227	8,165	21,867	198,091	30,053	8,752	872	3,036 358
2000	760,227	8,465	21,867	198,091	27,892	8,525	835	3,032 121
2001	750,200	10,300	26,400	179,400	29,928	8,862	756	3,025 401
2002	759,419	12,439	32,387	181,604	31,959	8,909	717	3,015 788
2003	769,200	12,306	31,748	164,100	28,942	9,121	767	3,013 381
2004	770,412	12,329	35,715	164,359	33,116	9,318	767	3,005 827
2005	803,920	12,847	39,628	192,670	31,285	8,874	707	2,985 171
2006	816,400	15,151	42,463	206,270	26,180	8,954	646	2,974 183
2007	831,390	14,276	46,482	190,400	19,198	8,824	535	2,969 457
2008	870,740	14,260	44,619	183,380	17,414	7,343	498	2,965 985
2009	888,800	13,969	54,442	178,210	24,705	8,791	521	2,966 181
2010	894,470	14,818	57,069	172,780	36,084	9,368	658	2,958 992
2011	917,620	14,743	55,562	152,740	32,506	8,472	712	2,960 184

(*) Before 1995 statistics on peas & beans were aggregated. (**) Data from the national forest inventory

Table 6-29. Standard yield of different crops used in the calculations, total weight (including water), kg/hectare.

Year	Winter wheat	Spring wheat	Winter rye	Winter barley (*)	Spring barley	Oats	Mixed grain	Triticale	Sugar beets	Winter rape
1990	5,818	4,918	4,195	-	3,911	3,866	3,305	5,818	44,843	2,748
1991	5,929	4,948	4,242	-	3,947	3,872	3,323	5,929	45,272	2,758
1992	6,040	4,979	4,288	-	3,982	3,879	3,341	6,040	45,701	2,767
1993	6,151	5,009	4,335	-	4,018	3,885	3,359	6,151	46,130	2,776
1994	6,207	5,012	4,398	-	4,036	3,869	3,359	6,207	46,446	2,777
1995	6,262	5,014	4,461	-	4,053	3,853	3,360	6,262	46,762	2,777
1996	6,393	5,078	4,600	-	4,103	3,882	3,394	5,434	46,985	2,752
1997	6,477	5,151	4,705	-	4,153	3,897	3,421	5,505	46,838	2,735
1998	6,592	5,021	5,010	-	4,136	3,714	3,336	5,603	46,686	2,681
1999	6,503	5,017	5,103	-	4,153	3,710	3,342	5,528	46,637	2,638
2000	6,446	5,059	5,204	-	4,137	3,658	4,431	6,446	46,300	2,609
2001	6,408	5,134	5,348	-	4,168	3,685	4,517	6,408	46,249	2,607
2002	6,351	5,176	5,448	-	4,204	3,747	3,976	6,351	46,416	2,634
2003	6,376	5,265	5,561	-	4,266	3,823	4,045	6,376	46,626	2,717
2004	6,231	5,227	5,526	-	4,245	3,853	4,049	6,231	46,661	2,789
2005	6,196	5,282	5,567	-	4,248	3,880	4,064	6,196	46,389	2,887
2006	6,169	5,201	5,515	-	4,201	3,870	4,036	6,196	47,193	3,027
2007	6,128	5,042	5,561	-	4,184	3,869	4,027	6,128	47,990	3,147
2008	6,184	4,966	5,580	5,177	4,280	3,997	3,245	4,849	49,129	3,214
2009	6,210	4,804	5,626	5,408	4,266	3,967	3,292	5,044	51,703	3,300
2010	6,268	4,585	5,711	5,460	4,320	4,016	3,305	5,060	52,920	3,423
2011	6,262	4,387	5,695	5,438	4,316	3,996	3,261	5,027	53,921	3,496

Swedish Board of Agriculture, Statistics Sweden, JO 15-series. (*) Until 2008 no standard yields was produced. The value for winter rye was used as an approximation.

Table 6-30. Standard yield of different crops used in the calculations, total weight (inc. water), kg/hectare.

Year	Spring rape	Winter turnip rape	Spring turnip rape	Table potatoes	Potatoes for starch prod.	Temporary grass	Green fodder
1990	1,777	1,821	1,587	29,194	36,045	7,490	5,000
1991	1,762	1,804	1,578	29,769	36,502	7,120	5,000
1992	1,746	1,787	1,570	30,343	36,958	6,010	5,000
1993	1,731	1,770	1,562	30,918	37,415	5,933	5,000
1994	1,715	1,747	1,558	31,409	37,585	5,856	5,000
1995	1,699	1,724	1,555	31,900	37,754	5,779	5,000
1996	1,679	1,682	1,542	31,817	37,651	5,702	5,000
1997	1,680	1,622	1,533	31,832	37,613	5,625	5,000
1998	1,607	1,523	1,420	34,910	39,706	5,548	5,000
1999	1,657	1,474	1,431	35,598	40,665	5,471	5,000
2000	1,720	1,471	1,451	35,146	40,401	5,394	5,000
2001	1,809	1,444	1,483	34,608	40,268	5,317	5,000
2002	1,910	1,390	1,511	33,866	39,812	5,240	5,000
2003	2,008	1,415	1,553	33,436	39,368	5,010	5,000
2004	2,062	1,440	1,573	32,461	38,530	5,010	5,000
2005	2,141	1,496	1,596	31,536	38,426	4,840	5,000
2006	2,175	1,586	1,590	30,976	38,367	4,450	5,000
2007	2,214	1,655	1,583	30,493	37,982	5,030	5,000
2008	2,217	1,709	1,534	29,857	37,315	4,863	5,000
2009	2,184	1,755	1,479	29,470	36,985	4,807	5,000
2010	2,144	1,811	1,428	29,374	37,020	4,889	5,000
2011	2,068	1,884	1,366	29,336	36,644	5,044	5,000

Swedish Board of Agriculture, Statistics Sweden, JO 15-series

6.4.1.2.7 4.D.1.5 Background emissions of N₂O from cultivation of organic soils

The background emissions from organic soils vary with different crops¹⁷⁰. They are considered to be higher from ploughed soils than from pasture or temporary grass lands¹⁷¹ and the suggested emission factors are 1 and 6 kg N₂O-N ha⁻¹, respectively. The IPCC Guidelines' default value is however implemented in the inventory since a Swedish/Finnish research group concluded that not enough data exists to generate different emission factors for different management and soil types¹⁷². The area of organic soils has only been estimated intermittently. The latest survey in 2009 concluded that approximately 5 % of the total area of arable land consists of organic soils¹⁷³. That fraction has then been used for all years, assuming that the area of organic soils relative the total area of arable land stays constant over time.

¹⁷⁰ Klemedtsson, 2001.

¹⁷¹ Klemedtsson, 2001.

¹⁷² Klemedtsson et al., 1999.

¹⁷³ Berglund, Berglund & Sohlenius, 2009

6.4.1.2.8 4.D.1.6 N₂O from sludge used as fertiliser

N₂O from sewage sludge used as fertiliser is a part of the N₂O emissions from agricultural soils and may be reported, according to the Good Practice Guidance, if sufficient information is available. This emission was included for the first time in the inventory for submission 2006. The activity data used is found in Table 6-22. Out of the total amount of nitrogen emitted, 70 % is assumed to emanate from direct emissions and 30 % from indirect emissions. The emission factor used for the direct emissions is 1.25 % of the nitrogen in the sewage sludge. The corresponding value for the indirect emissions is 1 %. Statistics on the use of sludge have been collected intermittently by Statistics Sweden and the Swedish EPA from sewage treatment plants (table 6-22). The emissions are calculated as:

$$emissions = \sum_i F_{Ri} \times N_{SEWAGESLUDGE} \times EF_i \times 44 / 28$$

F_{Ri} is the fraction of N emitted as direct/indirect emissions and EF_i is the corresponding emission factor. The direct emissions from sewage sludge have been reported as an optional category in the CRF and the indirect emission is reported under CRF category 4.D.3.1.

6.4.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Two related parameters are the amount of nitrogen in sold fertiliser, estimated by the sales statistics, and the nitrogen in used fertilisers, estimated from interviews with farmers. Sales statistics are collected annually by the Swedish board of agriculture and Statistics Sweden¹⁷⁴. Data has been collected in the same way from the larger producers and retailers since the early 1960s. Statistics on the use of fertiliser and manure have been collected biannually since the end of the 1980s¹⁷⁴. The estimated nitrogen content in sold products has for most years been somewhat higher. The two estimates should be about the same, at least in the long run. The difference may be due to storage and/or the fact that estimation methods are affected by different error types. The sales statistics also contain quantities sold for use outside the agricultural sector and are therefore expected to result in a higher figure. The user statistics provide valuable information about the use of fertilisers in different crops and regions, but the sales statistics are considered to give a more accurate estimate of total use. Therefore, the latter have been used in the GHG inventory. Another advantage of the sales statistics is that they are updated annually.

Statistics on the use of sewage sludge have been published irregularly and in different reports, but a time series has been created through interpolation/extrapolation and certain assumptions. The quality of data has increased over time and data for the latest years is of satisfactory quality.

¹⁷⁴ Statistics Sweden, MI 30-series.

6.4.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Times series and activity data are checked for consistency. Recalculations are identified and verified. Emission estimates and methods are reviewed yearly by external national experts.

6.4.1.5 SOURCE-SPECIFIC RECALCULATIONS

The total area of agricultural land has changed for all years. This is a consequence of the method used by SLU for calculating cropland (see 7.2.2.1). The largest changes are for the most recent years. Data for sludge use has been updated for 2010. Due to a late publication it was not possible to update this year in submission 2012.

6.4.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

6.4.2 Pasture, Range and Paddock Manure (CRF 4.D.2)

6.4.2.1 SOURCE CATEGORY DESCRIPTION

The category includes N₂O emissions from nitrogen excreted during grazing. Calculations are carried out according to the methodology in the IPCC Guidelines. The summary of the latest key category assessment, methods and EF used, and information on completeness, are presented in Table 6-31.

Table 6-31. Summary of source category description for the entire category CRF 4.D.2.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
4.D.2	N ₂ O	X	X		T2	CS, D	Yes

CS Country Specific. T2 Tier 2. D Default.

6.4.2.2 METHODOLOGICAL ISSUES

For N₂O emissions from N excreted on permanent pastures the default emission factor of 2 % N₂O-N/kg nitrogen excreted is used for all animal groups. This is probably an overestimation of the emission when the nitrogen lost as N₂O is likely to be lower in cold climates as in Sweden. However, very scarce information is available and until better empiric data is available the IPCC default EF will be used. The emissions are calculated as:

$$emissions = \sum_T N_T \times Nex_T \times Frac_{prp} \times EF \times 44 / 28$$

N_T is the number of animals of type _T in the country, Nex_T is the N-excretion of animals of type _T, Frac_{prp} is the fraction of the manure allocated to pasture, range and paddock (see Table 6-32) and EF is the emission factor. The nitrogen production for the different animal groups is presented in Table 6-13 and table 6-15.

Table 6-32. Waste management systems, fraction of manure deposited on Pasture, Range and Paddock.

Year	Dairy cattle	Other cattle	Pigs for meat production	Other swine	Sheep, Goats	Reindeer	Horses	Poultry
1990	0.25	0.42	NO	NO	0.50	1.00	0.50	NO
1995	0.25	0.44	NO	NO	0.50	1.00	0.50	NO
2000	0.25	0.46	NO	NO	0.50	1.00	0.50	NO
2005	0.26	0.49	NO	NO	0.50	1.00	0.50	NO
2006	0.26	0.49	NO	NO	0.50	1.00	0.50	NO
2007	0.25	0.47	NO	NO	0.50	1.00	0.50	NO
2008	0.25	0.47	NO	NO	0.50	1.00	0.50	NO
2009	0.25	0.46	NO	NO	0.50	1.00	0.50	NO
2010	0.25	0.46	NO	NO	0.50	1.00	0.50	NO
2011	0.24	0.46	NO	NO	0.50	1.00	0.50	NO

6.4.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The time-series is consistent.

6.4.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Times series and activity data are checked for consistency. Recalculations are identified and verified. Emission estimates and methods are reviewed yearly by external national experts.

6.4.2.5 SOURCE-SPECIFIC RECALCULATIONS

In submission 2012 data for nitrogen lost as ammonia was subtracted before the EF for N₂O emissions was applied. However, owing to comments from the UNFCCC centralized review and a “Saturday Paper” on this issue, data was resubmitted without the subtraction of ammonia. However, compared to the resubmission there are no recalculations for this source.

6.4.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

6.4.3 Indirect Emissions (CRF 4.D.3)

6.4.3.1 SOURCE CATEGORY DESCRIPTION

This category includes indirect emissions from soils. In general the Good Practice Guidance default emission factors are used. The Good Practice Guidance stresses the lack of knowledge on a global scale and the extreme variability in the suggested emission factors and parties are not encouraged to use national values unless rigorously documented and previewed country-specific values have been developed. However, values for losses of nitrogen as ammonia (see 6.4.1.2.2) and nitrogen leakage are national. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 6-33.

Table 6-33. Summary of source category description for the entire category CRF 4.D.3.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
4.D.3	N ₂ O	X	X		CS, T1a	CS, D	Yes

CS Country Specific. T1 Tier 1. D Default.

6.4.3.2 METHODOLOGICAL ISSUES

6.4.3.2.1 4.D.3.1 *Atmospheric deposition*

The formula for estimating the emissions is:

$$N_{Tot} = N_{Fert} \times Frac_{GASF} + N_{Sludge} \times 0.3 + N \times Nex \times Frac_{GASM} + N \times Nex \times Frac_{GASG}$$

$$N_2O = N_{Tot} \times EF \times 44 / 28$$

N_{fert} is the nitrogen supply by mineral fertiliser and $Frac_{GASF}$ is the corresponding N fraction emitted as ammonia, calculated from sold quantities of different fertilisers and with emission factors from the EMEP/EEA air pollutant emission inventory guidebook 2009. $N \times Nex$ is the total amount of nitrogen excreted from animals, combined with national estimates of $Frac_{GASM}$, the fraction of nitrogen from animal manure emitted as ammonia, and $Frac_{GASG}$, the fraction of manure from grazing animals emitted as ammonia (see 6.4.1.2 for explanations on the calculations of $Frac_{GASM}$ and $Frac_{GASG}$)

6.4.3.2.2 4.D.3.2 *Nitrogen Leaching and Run-off*

The national estimate of nitrogen leaching is estimated by the SLU and calculated from the SOILNDB model¹⁷⁵, which is a part of the SOIL/SOILN model¹⁷⁵. This simulation model was developed during the 1980s in order to describe nitrogen processes in agricultural soils¹⁷⁶. Since then the model has been elaborated and tested on data from controlled leaching experiments, and these tests show that the model estimates leaching from soil with good precision¹⁷⁷. By using national data on crops, yields, soil, use of fertiliser/manure and spreading time, the leaching is estimated for 22 regions. These regions are based on similarities in agricultural production areas. On average data from this model has been published every five years, intermittent years have been interpolated (Table 6-34).

For calculating nitrogen leaching in the inventory, the average N leaching per hectare, calculated by the SOILNDB model, is multiplied by the total Swedish area of agricultural soil.

¹⁷⁵ Johnsson, 1990; Swedish EPA, 2002.

¹⁷⁶ Johnsson et al., 1987.

¹⁷⁷ Swedish EPA, 2002b.

Table 6-34. Nitrogen leaching estimated by the SOIL/SOILN model.

Year	Nitrogen leaching (kg/ha)	Year	Nitrogen leaching (kg/ha)
1990	24.6 *	2001	20.0
1991	23.9	2002	19.5
1992	23.2	2003	19.0
1993	22.4	2004	18.5
1994	21.7	2005	18.0 **
1995	21.0 **	2006	18.0
1996	21.0	2007	18.0
1997	21.0	2008	18.0
1998	21.0	2009	18.0 **
1999	21.0 **	2010	18.0
2000	20.5	2011	18.0

* Interpolated using results from 1985 and 1994.

** Estimated with the SOIL/SOILN model.

Other years are interpolated/extrapolated.

The indirect N₂O emission is stated:

$$emissions = area \times leachfactor \times EF \times 44 / 28$$

To estimate the implied FracLEACH, which is required as additional information in CRF 4D for each reporting year, the leached nitrogen, according to the national model, is divided by the sum of nitrogen in fertilisers and animal production. This quotient varies between 0.20 and 0.25, which is relatively close to the IPCC Guidelines' default value of FracLEACH (0.3).

6.4.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The average nitrogen leaching from agricultural soils, the leach factor, has been estimated by Swedish University of Agricultural Sciences. The value for 1990 is calculated from an investigation that estimated nitrogen leaching for the years 1985 and 1994, however, only for the southern part of Sweden. So the value used is corrected to apply to the whole of Sweden. The reason for the decrease between 1999 and 2005 is believed to mainly be dependent on an increase in the area of catch crops. But an increased awareness of the eutrophication problem also lead to changed fertilising patterns. This model is considered to be the best available in Sweden, taking many relevant factors into account. Since statistics on the use of fertilisers and manure are produced every other year,¹⁷⁸ the estimates can be updated at most every second year. However, due to economic reasons, the data has been published intermittently.

¹⁷⁸ Statistics Sweden, NA 30-series; Statistics Sweden, MI 30-series.

6.4.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Times series and activity data are checked for consistency. Recalculations are identified and verified. Emission estimates and methods are reviewed yearly by external national experts.

6.4.3.5 SOURCE-SPECIFIC RECALCULATIONS

The total area of agricultural land has changed for all years. This is a consequence of the method used by SLU for calculating cropland (see 6.4.4.3 and 7.2.2.1). The largest changes are for the most recent years.

6.4.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

6.4.4 Other (CRF 4.D.4)

6.4.4.1 SOURCE CATEGORY DESCRIPTION

Under CRF 4.D.4 Sweden report a background emission from agricultural soils. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 6-35.

Table 6-35. Summary of source category description for the entire category CRF 4.D.4.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
4.D.4	N ₂ O	X	X		CS	CS	Yes

CS Country Specific.

6.4.4.2 METHODOLOGICAL ISSUES

Based on a study of national emission factors¹⁷⁹, a background emission from the cultivation of mineral soils have been included with the national emission factor of 0.5 kg N₂O-N ha⁻¹. The total area of mineral soils is calculated as total area of arable land minus the area of organic soils.

6.4.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The estimated emission is strongly dependent on estimated total area of agricultural land. This classification is performed by the Swedish National Forest Inventory (NFI¹⁸⁰). In total the sample consists of approximately 30,000 sample plots. However, only one fifth are investigated yearly. As a consequence of this the accuracy will increase retroactively until the whole five year cycle is completed. The estimate for the last year will only be based on 6,000 sample

¹⁷⁹ Klemedtsson, 2001.

¹⁸⁰ Ranney et al., 1987

plots, and then the accuracy gradually increases until the whole five year cycle is completed. That is, the number of sample plots for one, two, three, four and five years old data is 6,000, 12,000, 18,000, 24,000 and 30,000, respectively. However, to reduce the variation that sometimes arise for the most recent years due to sampling errors the estimates of the areas are calculated as a running average for the last five years (see 7.2.2.2).

6.4.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Times series and activity data are checked for consistency. Recalculations are identified and verified. Emission estimates and methods are reviewed yearly by external national experts.

6.4.4.5 SOURCE-SPECIFIC RECALCULATIONS

The total area of agricultural land has changed for all years. This is a consequence of the method used by SLU for calculating cropland (see 6.4.4.3 and 7.2.2.1). The largest changes are for the most recent years.

6.4.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

7 Land Use, Land-Use Change and Forestry (CRF sector 5)

7.1 Overview of LULUCF

Sweden reports carbon stock changes and greenhouse gas emissions from Forest land, Cropland, Grassland and Settlements and associated land-use transfers¹⁸¹. These land use categories are considered managed. Except for a small area used for peat extraction, Wetlands and Other land are considered unmanaged and consequently only areas of these categories are reported. The reporting also includes N₂O emissions associated with nitrogen fertilization of Forest land, N₂O emissions due to land conversions to cropland, CO₂ emissions associated with liming and N₂O as well as CH₄ emissions from biomass burning.

In 2011 the net removal from the Land Use, Land-Use Change and Forestry LULUCF-sector was estimated to ca 35,217 Gg CO₂. The net removal increased slightly from 2010 to 2011 by ca 4,538 Gg CO₂.

The long-term trend with decreasing net removals in living biomass is mainly due to an increase in harvest rates (Figure 7-1), while the increase in gross removal originating from tree growth has not been equally large. Gross removal (growth) in Sweden shows an increasing trend and lies currently around 120 Mm³sk (approx. 160 Mtons CO₂ per year). The harvest is also increasing but the annual fluctuations are large. In 2011 the gross harvest was approximately 90 Mm³sk (Figure 7-1). Since the harvest level is well below the growth this results in a steady increase in the carbon stock in living biomass which has prevailed since the beginning of the 20th century (Figure 7-1).

Since Submission 2010 Sweden reports data and supplementary information for the Kyoto Protocol including emission/removals for activities under article 3.3 (Afforestation, Reforestation and Deforestation) and for Forest management under article 3.4. In 2011 changes in carbon pools for afforestation, reforestation and deforestation together constituted an emission of 1,660 Gg CO₂ and Forest management represented a removal of 37,632 Gg CO₂. The supplementary information required for the reporting under the Kyoto Protocol is found in the NIR part II (Chapter 11).

¹⁸¹ Sweden uses random sampling methodology to estimate land-use and land-use transfers. The reporting is based on 30,000 permanent sample plots inventoried by the Swedish National Inventory of Forests (RIS) in five cycles each based on around 6,000 plots per year. The permanent sample plots have been re-inventoried at intervals of 5-10 years, from 2003 every five year, and the land-use of each is described from the year of the first inventory and every year thereafter. The land-use of years between inventories has been interpolated. This means that a full record of plots comprise 30,000 plots whereas the latest year is only represented by 6,000 measured plots. As the inventory cycle continues in subsequent years, more data covering the latest years are made available. Estimates of the five most recent years are thus re-calculated using re-measured plots in each submission. From submission 2012 and onwards, results for the latest years (those years without a full record of plots per cycle) are still calculated on a full record of plots using extrapolated data per cycle as approximate values for the not-yet-inventoried plots, see Figure 7-6. Both the extrapolation and the re-calculation is made to improve the accuracy of estimates.

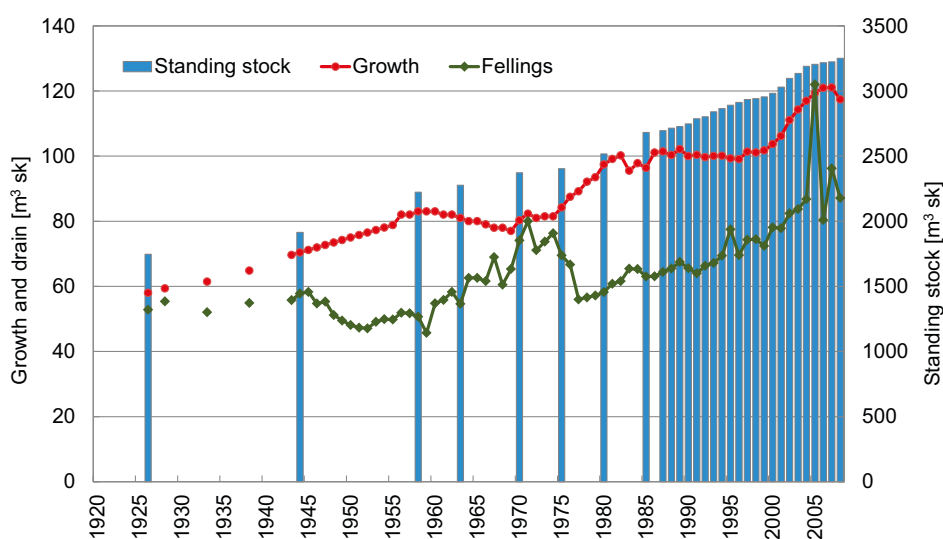


Figure 7-1 Annual growth and fellings and the standing stock (Source: Official Forest Statistics (<http://www.slu.se/en/webbtjanster-miljoanalys/forest-statistics/>))

Forest is the major land-use category in Sweden. The total forest area (FAO definition) is about 28 million hectares. The productive forests (where annual stem wood production per hectare and year is larger than 1 m³), on which most of the reported changes in carbon pools occur, is 23 million hectares¹⁸².

Harvest of trees is more or less restricted to productive forests and the unrestricted productive forest area has decreased since 1990, as a result of the establishment of nature reserves. Increased demand for forest products has led to a continuous increase in felling during the reported period peaking at 2005 (due to wind throws originating from a severe storm), while the growth rate only increased moderately. However, harvest statistics indicate a high demand of forest products in recent years with quite large fluctuations between years. It should be noted that from the base year and onwards, the reported growth is larger than the drain – defined as harvest and self-mortality.

The land use and land-use change matrices (Table 7-1a, 7-1b) is based on about 6,000 (1990-2011) and 30,000 sample plots (1990-2007), respectively¹⁸³. Due to a five-year inventory cycle we can only provide a full record of data 1990-2007. Note that only the area in Table 7.1b corresponds to the reported area in the CRF-tables (year 2007) while Table 7.1a might deviate somewhat because it is based on a smaller sample¹. Forest land is the most important land-use category. The gross and net conversions indicate that conversions from Forest land to Settlements are frequent. The reason for reporting two land use matrixes is based on a request from reviewers.

¹⁸² Swedish University of Agricultural Sciences, 2011

¹⁸³ The reason for reporting two land use matrixes is mainly because it is mandatory to report a land use matrix 1990-2011 and because the accuracy increases when using a full record of sample plots.

Table 7-1a¹⁸⁴ Land Use Categories 1990, 2011 and gross and net land use transfers 1990-2011 (based on about 6,000 permanent sample plots inventoried 1986-2011). The carbon stock of Forest land in the mountain area¹⁸⁵ (915,000 ha) is not monitored in the field and changes in the carbon pools for this area are not reported.

Area [1,000 ha]	"From"	"To" Year 2011					
	Year 1990	Forestland	Cropland	Grassland	Wetland	Settlements	Other land
Forest land	28,118	27,575	2	35	276	188	43
Cropland	3,283	74	3,066	46	4	94	0
Grassland	561	65	47	439	0	9	0
Wetlands	7,098	257	23	8	6,679	18	114
Settlements	1,683	84	20	12	0	1,562	5
Other land	4,279	59	0	6	15	0	4,200
Sum after transfers		28,114	3,157	546	6,974	1,870	4,362

Table 7-1b Land Use Categories 1990, 2007 and gross and net land use transfers¹⁸⁶ 1990-2007 (based on about 30 000 permanent sample plots inventoried 1983-2011).

Area [1 000 ha]	"From"	"To" Year 2006					
	Year 1990	Forestland	Cropland	Grassland	Wetland	Settlements	Other land
Forest land	28,220	27,901	5	26	103	146	41
Cropland	3,102	82	2,903	46	4	67	0
Grassland	508	43	41	405	3	12	4
Wetlands	7,303	226	2	3	6,963	5	104
Settlements	1,705	76	18	5	8	1,588	10
Other land	4,341	47	0	2	41	1	4,250
Sum after transfers		28,375	2,969	487	7,121	1,819	4,409

The largest carbon stocks are found in the living biomass and soil organic carbon pools on Forest land, and the largest annual stock change is the change in the living biomass pool (Figure 7-2 and 7-3). A net removal of CO₂ due to increases in the living biomass pool is reported for every year during the period.

The dead organic matter pool resulted in net removals during most years of the reported period, whereas the soil organic carbon resulted in net emissions during most years of the reported period. Some soils act as sources whereas others act as sinks. The major source is the emissions from drained organic soils (Histosols) on Forest land and on Cropland. An area of about 4.2 Mha of the Forest land was considered as Histosols 2011 and 1.2 M ha of the Histosols are drained. The Cropland area on Histosols is estimated to approx. 145 kha (2011) and all of that area is drained. The area is decreasing since the total Cropland area (the average trend) in Sweden is decreasing, leading to a trend towards decreased emissions from Cropland. The decrease in area is mainly due to conversion to Forest land and Settlements. Conversion from and to Grassland do also occur.

¹⁸⁴ Table 7.1a is put together based on a request from the ERT in the centralized review (Subm 2009).

¹⁸⁵ Löfgren, 1998

¹⁸⁶ The carbon stock of Forest land in the mountain area (915 000 ha) is not monitored in the field and changes in the carbon pools for this area are not reported.

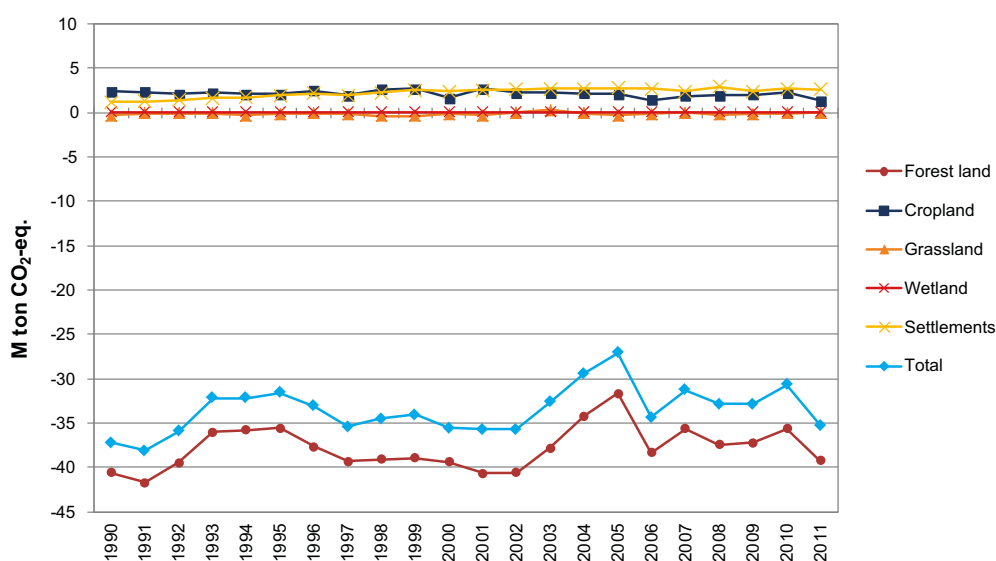


Figure 7-2 Net emissions/removals of GHG in the LULUCF sector from different land-use categories and total net removals for the LULUCF-sector

There has been considerable variation between submissions for specific years in the soil organic carbon pool on mineral soils. These variations are partly caused by random variation in the sample. Since the total pool is huge and the changes in the pool comparatively small the numbers are sensitive to random variation when small changes are multiplied by large areas. It should be noted that a change of 0.1% in the pool is equivalent to more than 3,000 Gg CO₂. The variation between years has been reduced after the introduction of a new method for extrapolation of data on plot basis (first time used and described as a recalculation in Submission 2011). Variation between submissions may still be substantial. We expect that this variation will decrease with time when more plots are re-inventoried.

Emissions of CO₂, N₂O and CH₄ from i) direct N₂O emissions from nitrogen fertilization, ii) N₂O emissions from disturbance associated with land-use conversion to Cropland, iii) CO₂ emissions from agricultural lime application, and iv) GHG-emissions from biomass burning are quite limited in Sweden. The total emission shows no obvious trend but instead a quite stable emission less than 0.3 Mton CO₂-equivalents every year during the period 1990-2011. Among these categories, the largest emissions originate from liming. Sweden does not report N₂O emissions from drainage of soils. A summary of emissions/ removals is found in Tables 7-2a and 7-2b.

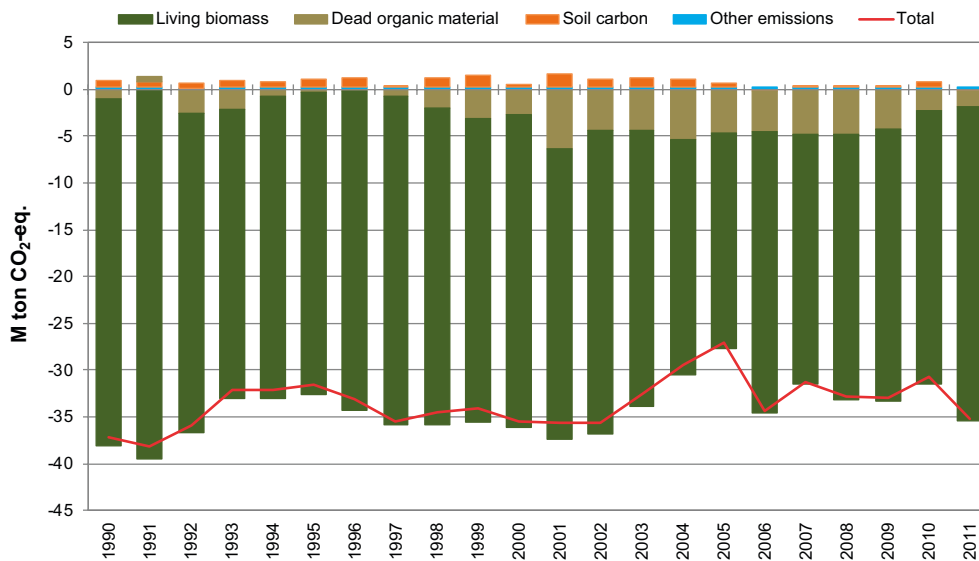


Figure 7-3 Net removals/emissions of GHG in the LULUCF sector from different carbon pools.

Table 7-2a. Summary of net removals (-)/emissions (+) in living biomass (LB), dead organic matter (DOM) and soil organic carbon (SOC) per land use category.

7.2a	Net removals (minus=removal) [Mt CO ₂]															
	Forest land				Cropland				Grassland				Wetland	Settlement		
	LB	DOM	SOC	Min. Org.	LB	DOM	SOC	Min. Org.	LB	DOM	SOC	Min. Org.	SOC	LB	DOM	SOC
1990	-36.6	-1.0	-12.0	9.0	0.1	0.0	0.1	2.1	-0.6	-0.4	0.3	0.3	0.0	0.0	0.3	0.8
1991	-39.1	0.5	-12.1	9.0	-0.1	0.0	0.2	2.1	-0.3	-0.4	0.3	0.3	0.0	0.0	0.4	0.9
1992	-33.6	-2.7	-12.2	9.0	-0.1	0.0	0.1	2.1	-0.3	-0.3	0.3	0.3	0.0	0.0	0.4	1.0
1993	-30.6	-2.2	-12.2	9.0	-0.1	0.0	0.1	2.1	-0.4	-0.3	0.3	0.3	0.0	0.1	0.5	1.1
1994	-31.6	-1.0	-12.2	9.0	-0.1	0.0	-0.1	2.0	-0.5	-0.3	0.3	0.3	0.0	0.0	0.5	1.2
1995	-31.8	-0.6	-12.2	9.0	-0.2	0.0	0.1	2.0	-0.4	-0.3	0.3	0.3	0.0	0.1	0.6	1.3
1996	-33.9	-0.4	-12.3	9.1	-0.1	0.0	0.3	2.1	-0.3	-0.3	0.3	0.3	0.0	0.1	0.6	1.4
1997	-34.5	-1.0	-12.8	9.1	-0.1	0.0	-0.1	2.0	-0.3	-0.3	0.2	0.3	0.0	-0.2	0.7	1.5
1998	-33.0	-2.4	-12.7	9.1	-0.1	0.0	0.5	2.1	-0.6	-0.3	0.3	0.3	0.0	0.0	0.7	1.5
1999	-31.9	-3.5	-12.7	9.2	-0.1	0.0	0.5	2.1	-0.7	-0.3	0.3	0.3	0.1	0.2	0.7	1.6
2000	-32.4	-3.2	-13.0	9.2	-0.4	0.0	-0.2	2.0	-0.4	-0.3	0.3	0.3	0.1	-0.1	0.8	1.7
2001	-30.0	-6.9	-13.0	9.2	-0.3	0.0	0.7	2.1	-0.6	-0.3	0.3	0.3	0.1	-0.1	0.8	1.8
2002	-31.8	-5.0	-13.0	9.2	-0.1	0.0	0.1	2.0	-0.3	-0.3	0.3	0.3	0.1	-0.1	0.9	1.9
2003	-29.1	-5.0	-13.0	9.2	-0.2	0.0	0.2	2.0	0.0	-0.3	0.3	0.3	0.1	-0.2	0.9	2.0
2004	-24.5	-6.0	-13.0	9.2	-0.3	0.0	0.3	2.0	-0.3	-0.3	0.2	0.3	0.0	-0.1	0.9	1.9
2005	-22.3	-5.3	-13.4	9.2	-0.2	0.0	0.1	2.0	-0.6	-0.3	0.2	0.3	0.1	0.1	0.8	1.8
2006	-29.3	-4.8	-13.6	9.3	-0.5	0.0	-0.2	2.0	-0.4	-0.2	0.2	0.3	0.0	0.2	0.8	1.7
2007	-26.1	-5.3	-13.6	9.3	-0.4	0.0	0.1	2.0	-0.3	-0.2	0.2	0.3	0.1	0.0	0.7	1.7
2008	-28.0	-5.3	-13.6	9.3	-0.4	0.0	0.1	2.0	-0.5	-0.3	0.2	0.3	0.1	0.4	0.7	1.7
2009	-28.2	-4.8	-13.5	9.3	-0.3	0.0	0.2	2.0	-0.4	-0.2	0.2	0.3	0.1	-0.1	0.7	1.8
2010	-28.7	-2.8	-13.5	9.3	-0.4	0.0	0.3	2.0	-0.4	-0.2	0.2	0.3	0.1	0.2	0.7	1.8
2011	-32.8	-2.3	-13.5	9.3	-0.6	0.0	-0.2	1.9	-0.3	-0.2	0.2	0.3	0.1	0.1	0.7	1.8

Table 7-2b. Summary of net emissions (+)/removals (-) in living biomass (LB), dead organic matter (DOM) and soil organic carbon (SOC) and other sources. The total LULUCF removals are expressed as CO₂-equivalents.

7.2b	Total carbon pool changes [Mt CO ₂]			Other emissions [Mt substance]						Total	
	LB	DOM	SOC	Fert.	To CL	Liming	Biomass burning			LULUCF	Total LULUCF
							5 (I) N ₂ O	5 (III) N ₂ O	5 (IV) CO ₂	5 (V) CO ₂ CH ₄ N ₂ O	
Year											[Mt CO ₂ -eq]
1990	-37.0	-1.0	0.6	1.9E-04	7.0E-05	1.7E-01	IE	8.2E-05	5.6E-07	5.6E-07	-37.2
1991	-39.5	0.5	0.6	1.1E-04	8.8E-05	1.3E-01	IE	7.6E-05	5.3E-07	5.3E-07	-38.1
1992	-34.1	-2.6	0.6	7.6E-05	9.2E-05	1.1E-01	IE	7.7E-05	5.3E-07	5.3E-07	-35.9
1993	-30.9	-2.1	0.7	6.7E-05	1.1E-04	1.3E-01	IE	8.0E-05	5.5E-07	5.5E-07	-32.2
1994	-32.2	-0.8	0.6	5.9E-05	1.2E-04	1.6E-01	IE	7.6E-05	5.3E-07	5.3E-07	-32.2
1995	-32.3	-0.3	0.8	6.9E-05	1.3E-04	1.7E-01	IE	7.7E-05	5.3E-07	5.3E-07	-31.6
1996	-34.2	-0.1	1.0	6.2E-05	1.4E-04	1.9E-01	IE	8.2E-05	5.6E-07	5.6E-07	-33.0
1997	-35.2	-0.7	0.2	4.9E-05	1.5E-04	1.7E-01	IE	4.2E-04	2.9E-06	2.9E-06	-35.4
1998	-33.7	-2.0	1.0	5.0E-05	1.5E-04	1.3E-01	IE	2.2E-05	1.5E-07	1.5E-07	-34.5
1999	-32.5	-3.1	1.3	6.5E-05	1.5E-04	1.6E-01	IE	1.4E-04	9.7E-07	9.7E-07	-34.0
2000	-33.4	-2.7	0.3	6.4E-05	1.7E-04	1.6E-01	IE	1.4E-04	9.7E-07	9.7E-07	-35.5
2001	-31.0	-6.4	1.4	5.4E-05	1.8E-04	1.4E-01	IE	1.4E-04	9.9E-07	9.9E-07	-35.7
2002	-32.4	-4.4	0.9	3.7E-05	1.9E-04	1.3E-01	IE	2.3E-04	1.6E-06	1.6E-06	-35.7
2003	-29.5	-4.3	1.0	4.4E-05	2.0E-04	1.3E-01	IE	2.9E-04	2.0E-06	2.0E-06	-32.6
2004	-25.1	-5.4	0.9	5.5E-05	2.1E-04	1.2E-01	IE	2.6E-04	1.8E-06	1.8E-06	-29.4
2005	-23.0	-4.7	0.4	8.2E-05	2.1E-04	1.2E-01	IE	2.4E-04	1.6E-06	1.6E-06	-27.1
2006	-30.0	-4.3	-0.2	8.8E-05	2.1E-04	9.1E-02	IE	5.8E-04	4.0E-06	4.0E-06	-34.3
2007	-26.7	-4.8	0.0	1.2E-04	2.1E-04	1.2E-01	IE	1.1E-04	7.9E-07	7.9E-07	-31.3
2008	-28.4	-4.8	0.1	1.6E-04	2.3E-04	1.0E-01	IE	6.3E-04	4.3E-06	4.3E-06	-32.8
2009	-29.0	-4.3	0.2	1.5E-04	2.3E-04	9.8E-02	IE	1.3E-04	8.6E-07	8.6E-07	-32.9
2010	-29.2	-2.2	0.5	2.1E-04	2.3E-04	9.1E-02	IE	3.4E-05	2.3E-07	2.3E-07	-30.7
2011	-33.5	-1.7	-0.1	1.4E-04	2.3E-04	7.9E-02	IE	1.0E-04	7.0E-07	7.0E-07	-35.2

7.2 Description of categories 5A, 5B, 5C, 5D, 5E and 5F

7.2.1 Characteristics of categories

A summary of the key categories under the LULUCF-sector is found in Table 7-3. In the LULUCF-sector CO₂ emissions/removals for Forest land, Cropland, Grass-land and Settlements are considered key-categories for reporting of carbon stock changes in the CRF-categories 5A, 5B, 5C and 5E. Sweden uses Tier 2 and 3 methodologies and country-specific emissions factors. Land under 5D (except for a small area used for peat extraction) and 5F are considered unmanaged and not reported. The reported land under 5D refers to quite limited emissions from peat extraction. Emissions from categories 5I, 5III, 5IV and 5V are reported. For some of these categories it is not possible to separate emissions into land use categories since emissions are based on the total amount used (nitrogen fertilisation) or sold (liming). Category 5III is considered key-category (included in 5B Cropland N₂O). Category 5II, Non-CO₂ emissions from drainage of soils and wetlands, is not reported.

7.2.2 Information on approaches used for representing land areas and on land-use databases used for the inventory preparation

7.2.2.1 THE SWEDISH NATIONAL INVENTORY OF FORESTS

The Swedish National Inventory of Forests (RIS¹⁸⁷) consists of the Swedish National Forest Inventory (NFI¹⁸⁸) and The Swedish Forest Soil Inventory (MI¹⁸⁹). The NFI and the MI are integrated in the same sample design, using the same permanent sample plots. However, the sampling interval of the soil inventory is longer since processes in the soil are much slower than in the living biomass.

The plots associated to the NFI are re-inventoried every fifth year and the plots associated to the MI are re-inventoried every tenth year. Moreover, top soil cores are only taken at every second sample plot and deeper soil horizons are only sampled on every fourth sample plot. The reported data of changes in the living biomass and dead wood pools are based on the NFI-measurements and changes in the litter and soil organic carbon pools are based on the MI-measurements.

¹⁸⁷ Swedish University of Agricultural Sciences, 2011

¹⁸⁸ Ranneby et al., 1987

¹⁸⁹ Swedish University of Agricultural Sciences, <http://www-markinventeringen.slu.se/>

The NFI is an annual, systematic, cluster-sample inventory of Sweden's forests (Figure 7-4 and 7-5). Each year roughly a thousand survey sample clusters are inventoried in the field. One third of the clusters are temporary and two thirds are permanent. Only permanent sample plots are used for the UNFCCC reporting. The clusters are distributed all over the country in a pattern that is denser in the southern part of than in the northern part of the country. The clusters (tracts) are square-shaped with sample plots along each side. Each cluster consists of four to eight sample plots, depending on geographical region. Each year, about 6,000 permanent survey sample plots are inventoried in the field. On each circular sample plot, with a radius usually of 10 or 20 m, information is collected about the trees, the stand and the site. The main focus of the NFI is on monitoring forests for timber production and environmental protection.

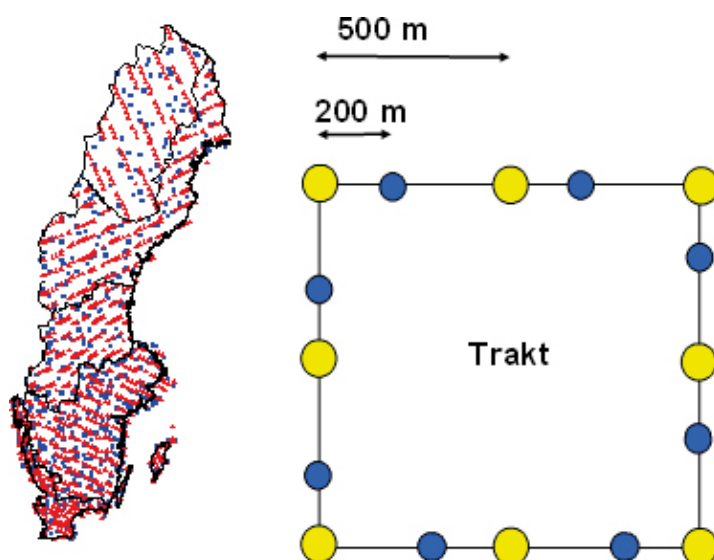


Figure 7-4 Covering the whole country of Sweden, each year a permanent sample grid (red) is re-inventoried and a temporary sample grid (blue) is inventoried. To be able to trace both gross and net land use transfers, only permanent sample plots are used in the reporting. When estimating changes of e.g. C, the accuracy is also higher using permanent than when using temporary sampling plots. Each red dot represents a cluster of sample plots (Trakt) and within Trakt the yellow plots are used for the inventory while the blue plots are used for validation of harvests (estimates on up to one year old stumps).

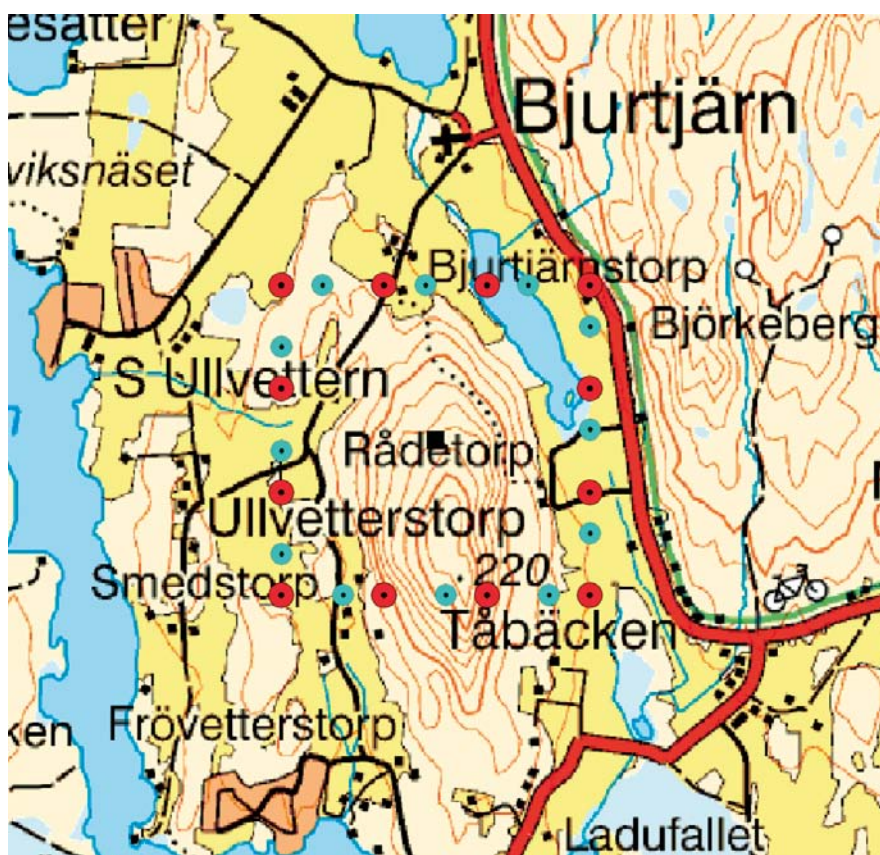


Figure 7-5 The sample plots (red) are covering all relevant land use. On the example above, plots are located on e.g. Forest land, Cropland, Wetlands and some plots are divided into more than one land use category. On the plots, measurements are made to estimate standing biomass of trees. If at the next re-inventory, the trees remains and has been growing the plot represents a net sink but if they have been harvested the plot represents a source (stock change method). Volume of dead wood per decay classes are also measured on the plot. Soil samples from different soil horizons are sampled and analyzed for C concentration and other properties. Litter is partly estimated using data from the plot and partly modeled. Observe that the shape of tracts differs by county. An additional sample (blue) is used for estimates of harvests.

The soil inventory uses the 10-m radius sampling plot. A number of variables are recorded including general site variables, soil and humus type. The litter and different soil layers are sampled for further laboratory analysis. The O, H or A horizon are sampled using an augur. The mineral soil is sampled in different layers according to the distance from the soil surface and to some extent depending on the soil type. From 2003 and onwards the soil sampling has been harmonized with an ongoing European inventory, i.e. Biosoil¹⁹⁰ and soil samples are taken at fixed depths.

¹⁹⁰ <http://biosoil.jrc.it/>

7.2.2.2 SAMPLE BASED ESTIMATIONS

The sample frame consists of a map covering the whole land and fresh water area of Sweden. A sea archipelago zone where islands covered by vegetation might occur is also included in the frame (but no sea area is reported). The frame is divided into 31 strata (i.e. representing counties) and a specific number of sample units are sampled per stratum. Each cluster (tract) of sample plots is assumed to be the sample unit. The inventoried area of a tract is given a specific area weight and will consequently represent a larger area. The weighing is generated so that the sum of all represented areas will be equal to the total county area.

The land use of whole plots or parts of plots may change by time but the total tract area will always represent the same area. At the county level, the reported value of a change in a carbon pool (for example a change in the living biomass pool for the land use category Forest land remaining Forest land) will be estimated by a ratio estimator¹⁹¹. Finally the reported value on national level is estimated as the sum of the county values (For further information, see Annex 3.2).

A five year inventory cycle is used and five different samples were randomly distributed (using a systematic grid) 1983, 1984, 1985, 1986 and 1987, respectively. Each of these samples consists of around 6,000 sample plots. The expected value of an estimator is theoretically the same for any given sample but to reduce sample randomness all five samples are merged. Full sets of samples are currently only available for years until 2007 and consequently only 24,000, 18,000, 12,000 and 6,000 sample plots are available for the estimates of 2008, 2009, 2010 and 2011 respectively. When five years have proceeded since any reporting year, all samples have been re-inventoried covering that particular year and the full set of data can be used to produce the estimate. Therefore, the five most recent reporting years are re-calculated and revised in each submission.

Since the random variation of the samples largely affects the estimates of areas and carbon stock changes for the four most recent years (as also noted in the annual review reports), Sweden now extrapolates each of the five sample series (cycles) using the running average for the five years prior to the year of the latest actual re-measurement, to enlarge the data set for each reported year. In Figure 7-6, the extrapolation of each sample series and its consequences on the estimates are illustrated. The effect of the extrapolation levels out “strange” area and carbon stock variations evolving from the randomness of the sampling as exemplified in Figure 7-6¹⁹².

¹⁹¹ Thompson, 1992

¹⁹² This improvement and the information provided is a response to reviewers (ARR 2011)

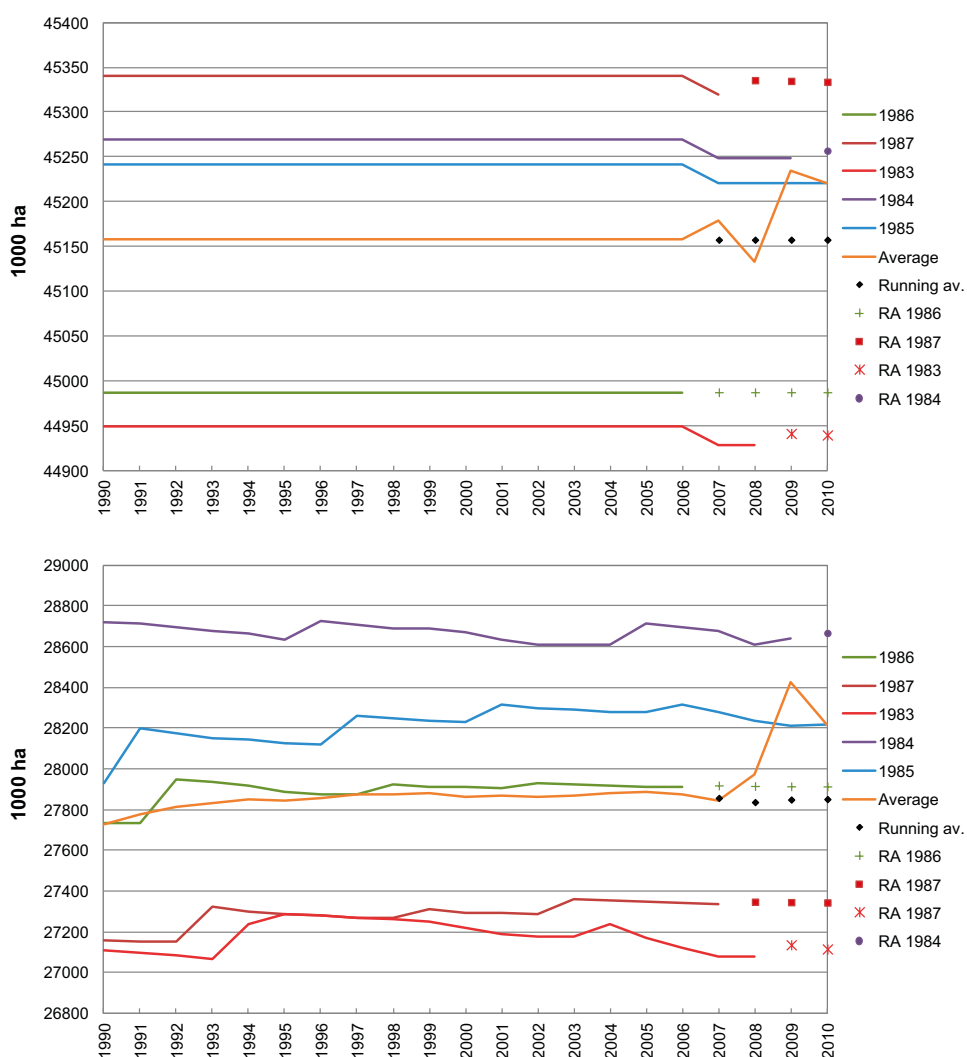


Figure 7-6. The upper panel shows the total area of Sweden with (R average) and without (average) the extrapolated numbers, continuous lines represents measured and interpolated values and dots represents extrapolated values. The lower panel shows the area of Forest land remaining Forest land with (R average) and without (average) the extrapolated numbers, continuous lines represents measured and interpolated values and dots represents extrapolated values.

Sweden reports “human induced” carbon changes only, where “human induced” has the interpretation of “managed”, i.e. the biomass stock change on unmanaged land are set to zero. However, the “actual” stock on unmanaged land is considered when calculating stock changes after conversions between unmanaged and managed land and vice versa. This is possible since trees are inventoried on almost any land. All areas, managed or unmanaged, are reported.

7.2.2.3 THE LULUCF-REPORTING DATABASE

Around 30000 permanent sample plots inventoried by RIS are used for the area based sampling. The plots were distributed during the period 1983-1987, representing all land and fresh-water areas have been re-inventoried at intervals of 5-10 years (from 2003 in a five-year cycle). Land-use of each plot (or sub-plot for plots divided in two or more land use classes) is described from the year of the first inventory and every year thereafter. The land-use of years between inventories has been interpolated by linear interpolation (see Table A.3.2.1 in Annex 3:2). Biomass pools for years between inventories are also linearly interpolated. If final felling is identified on a plot, the year of harvest between inventories is assessed in field. If both harvest and a land use conversion are identified, then these two events are matched to the same year (the year of land use conversion between inventories is assessed in field since 2006, see next section).

7.2.2.4 LAND USE TRANSFERS CRF-TABLES 5A, 5B, 5C, 5D, 5E AND 5F

Until 2005 information of the year of a land use change was not recorded by the NFI. Therefore, land-use transfers between 1990 and 2005 are assumed to occur at a random year between two consecutive inventories. From the inventory year 2006 and onwards the year of conversion is assessed in field. Every plot that is converted to another land-use category is reported for 20 years in the land-use transfer class. After 20 years the plot will be reported in the class to which it was transferred. If a second land-use conversion occurs within the 20 years, the counting starts all over again and the second transfer is reported for 20 years in the land-use transfer class as in the first example. In the reporting database it is possible to trace some of the land-use transfers that occurred up to 20 years before 1983 and consequently it is possible to decide how many years a sample plot has belonged to a certain land-use category and what land-use category it was converted from already at the start of the reporting period (1990). Consequently, several land use transfer categories include areas already before 1990. The IPCCs “20 year default rule” has vast influence on reported figures. One example is the fact that accumulation of carbon in the Living biomass pool in the conversion category Cropland converted to Forest starts already before the base year. After twenty years, parts of these land areas are reported as Forest remaining Forest. Consequently a loss of carbon may be reported under Cropland to Forest for and the corresponding removal is instead reported under Forest remaining Forest. To some extent, the continuously transfer of land use between categories and the five-year inventory cycle average out such events.

The FAO definition of Forest land was introduced in the Swedish NFI in 1998. Until 1998 Forest land was assessed based on the national definition of Forest land. Therefore, land-use categories have to be re-determined for the period 1990-1997. There are two main types of redetermination cases which are handled as follows:

1. If the land-use category for a sample plot was registered as Forest land at the first inventory after 1997 and the national land-use category (see Table 7-4) had been the same at all earlier inventories since 1990, the plot are assumed to have always belonged to the land-use category Forest land.
2. If the land-use category was registered as Forest land at the first inventory after 1997 and the national land-use category (see Table 7-4) had changed since 1990, the first land-use category are assumed to remain until the year of conversion. If at consecutive inventories after that, the land-use category belonged to the same land-use category, the plot is assumed to belong to the category Forest land all years after the year of conversion.

Two types of inconsistently classified land-use transfers have been identified and corrected:

1. Inconsistency over time in applying land-use category definitions.
2. Inconsistency in delineating borders between plots divided into more than one land-use category.

One example of the first type is when at different inventories, the land-use category of a sample plot has been classified as Forest land at the first inventory, as Wetland at the next inventory and then again as Forest land at the third inventory without traces of human activities. A case like this is corrected so that the land-use category is assumed to be Forest land on all three occasions. Another example of the first type is when a recreation forest close to a city has been converted from Settlements (section 7.2.3.1, national land-use category 13, "Urban land") to Forest land and the new land-use category consists of old trees. This has been corrected so the land-use is assumed as Forest land on both occasions. One example of the second type is when the delineation of a divided plot, representing more than one land-use category, has been changed at the re-inventory due to personal judgments rather than due to actual changes. These land-use changes should not be registered as land use changes and have been corrected by keeping the newer delineation, usually if the assumed incorrect new delineation deviates approximately less than 0.75 m² from the old delineation. If the affected area is larger, the new delineation is assumed to be correct. Rules for automatic and manual corrections of inconsistencies and the actual corrections are saved and could be verified on request.

7.2.3 Land-use definitions and the classification systems used and their correspondence to the LULUCF categories

Sweden has defined Forest land according to the Global Forest Resources Assessment (FAO/FRA) 2005¹⁹³. Forest land is land with a tree crown cover (or equivalent stocking level) of more than 10 percent at maturity, with a minimum area of 0.50 hectare and the trees should be able to reach a minimum height of 5 m at maturity *in situ*. However, there are two small discrepancies between the FRA 2005 definition and the definition implemented in the Swedish inventory.

Table 7-3. Summary of key category description, CRF 5. If no emissions/ removals are associated with a category, this category is not shown. Emissions/ removals from CRF-tables 5I - 5V are very small and included in CRF 5A – 5F. All pools and emissions are reported except 5II.

CRF	Gas	Key Category Assessment 2011			Method	EF
		Level	Trend	Qualitative		
5A	CO ₂ ^{1,3}	X	X		T1,T2, T3	CS
	N ₂ O				T1 (5I, 5V)	D (5I, 5V)
	CH ₄				T1 (5V)	D (5V)
5B	CO ₂ ²	X	X		T1,T2,T3, T1 (5IV)	CS, D (5IV)
	N ₂ O		X		T1 (5III)	D (5III)
5C	CO ₂ ³	X			T1, T2, T3	CS
	N ₂ O				T1 (5V)	D (5V)
	CH ₄				T1 (5V)	D (5V)
5D	CO ₂				T2	CS
5E	CO ₂	X	X		T1,T2, T3	CS

¹=5I included, no key category, T1, CS; ²=5III included, no key category, T1, D; ³=5V included, no key category, T1, CS

In the Swedish inventory permanent forest roads (width>5m) are not considered Forest land, and no minimum width to constitute Forest land is considered (note that the strict terms of Decision 11/CP.7 does neither mention the minimum width nor the forest roads¹⁹⁴. All Forest land is considered managed, i.e. even protection of forests in reserves is considered as management. Cropland is defined as regularly tilled agricultural land and all Cropland is assumed managed. Grassland is defined as agricultural land that is not regularly tilled and all Grassland is assumed managed. Generally, Wetlands is assumed unmanaged and is defined as mires and areas saturated by fresh water. However, approx. 10,000 ha of the Wetland area is used for peat extraction and therefore assumed managed. Settlements are defined as infrastructure components such as roads and railways, power lines within forests, municipality areas, gardens and gravel pits. All Settlements are assumed managed. Other land is defined as impediments (waste land) and includes

¹⁹³ Food and Agriculture Organization of the United Nations, 2004

¹⁹⁴ FCCC/CP/2001/13/Add.1, p 58

most of the mountain area in northwest Sweden. All Other land is assumed unmanaged. Land-use categories are monitored within the Swedish National Forest Inventory (NFI¹⁹⁵).

7.2.3.1 THE CONNECTION BETWEEN NATIONAL AND REPORTED LAND USE CATEGORIES

The reported land use categories are based on 16 national land use categories monitored by the Swedish National Inventory of Forests (RIS). For example in year 2000 the area of Forest land, according to the definition of forest described above, was estimated to 27,414,000 ha. This corresponds to 22,749,000 ha Productive Forest land (national category 01), 1,678,000 ha Mire (04), 520,000 ha Rock Surface (05), 268,000 ha Sub alpine Coniferous Woodland (06), 376,000 ha High Mountain (07), 1,615,000 ha Protected Area, Nature Reserve (11) and 208,000 ha to other categories, using the national land use categories (Table 7-4). Note that the international land use category (FRA 2005) Forest land is superior to all other land use categories.

7.2.3.2 CONSISTENCY IN REPORTING LAND USE CATEGORIES

The NFI has monitored land-use categories in a reasonably consistent way since 1983. Based on permanent sample plots, it is possible to trace both gross and net land-use transfers from 1983 and onwards. On Forest land, it is also possible to determine former land-use (i.e. Cropland or Grassland) before the base year (1990). All land areas are included in the field inventory except high mountains and urban land. These latter land-use categories are only inventoried for area by remote sensing. It is assumed that their relative importance for the Swedish carbon budget is negligible.

A few historical inconsistencies in the land-use category assessment have been identified and corrected. Before year 2003, protected areas ("Protected Area, Nature Reserve"; section 7.2.3.1) were not regularly inventoried. From 2003 and onwards this areas are included in other land-use categories. Usually there are data from at least one field inventory of "protected areas" before 1990, but sometimes no data are available. If no data are available, the change in carbon pools in former "protected areas" is assumed to be zero from 1990 to 2002. From 2003 potential changes will be reported based on field inventory data. The FRA 2005 definition of Forest land was introduced in the field inventory in 1998 and therefore land-use categories in earlier inventories has been re-determined. A description on the treatment of former protected areas, re-determination of land-use categories and the methodology for correcting inconsistencies in the land-use category assessment are described in more detail in the methodology section.

¹⁹⁵ Ranneby et al., 1987

Table 7-4 National Land Use Categories, their connection to the UNFCCC Land Use Categories and their potential importance for carbon reporting. A=all land is considered FAO Forest land, B=large areas are considered FAO Forest land. Observe that this example is based on both temporary and permanent sample plots. Thus, the total area is not comparable with reported areas.

National Land Use Category	UNFCCC/KP-Land Use Category	Carbon Stock In Living Biomass of Trees [T gram] Year 2000	Area [1,000 ha] Year 2000	Additional explanation
Productive Forest land (01)	F	1,047	22,749	Land which hosts a potential yield of stem-wood exceeding one cubic metre per hectare and year (A).
Grazing Land (02)	G	5.7	494	Not regularly cultivated.
Arable Land (03)	C	1.5	3,052	Regularly cultivated
Mire (04)	W	35.6	4,588	Land which hosts a potential yield of stem-wood lower than one cubic metre per hectare and year (B).
Rock Surface (05)	O	16.4	896	Rocky or stony areas. (B)
Sub alpine Coniferous Woodland (06)	F	8.2	307	Land-zone usually located between (01) and (07). (A)
High Mountain (07)	O	Low	3,010	Usually unstocked or sparsely stocked. (B)
Climatic Impediment (08)	O	0.7	48	Usually located in flat terrain in northern Sweden. (B)
Road and Railroad (09)	S	0.5	445	For permanent use. Not only roadway and rail but also other connected areas as embankments and ditches.
Power line Within Forest (10)	S	0.2	145	Minimum width 5 m, otherwise Productive Forest land (01)
Protected Area, Nature Reserve (11)	(F)	Medium	3,967	This land use category was left out 2003 and is thereafter included in the remaining land use categories.
Military Impediment (12)	S	Low	69	Could not be inventoried for security or safety reasons.
Urban Land (13)	S	Low	1,185	Settlements of many different kinds.
Other land (14)	S	2.1	115	Different kinds of land that is not covered by Other land use categories. Examples: gravel pits, halting places and slalom slopes
Water (not sea) (15)	W	0	4,009	Lakes, rivers, creeks, canals, pounds etc. Minimum width of 2 m.
Sea (16)	-	-	-	To check if the total land area is constant.
Total		1,118	45,080	

7.2.4 Definition of carbon Pools, CRF 5A, 5B, 5C, 5D, 5E and 5F

7.2.4.1 LIVING BIOMASS

The reported carbon pool changes refer to the biomass of all living trees with a height of at least 1.3 m. Thus, small trees, shrubs and other vegetation, such as herbs are not included in the figures. Both above-ground and below-ground biomasses are reported. Above-ground biomass is defined as living biomass above stump height (1 % of tree height). Scots pine (*Pinus sylvestris*), Norway spruce (*Picea abies*) and birch (*Betula pendula* and *Betula pubescens*) constitute about 92 % of the standing volume¹⁹⁶. Broad-leaved species constitute most of the remaining 8 %. Below-ground biomass is defined as living biomass below stump height (1 % of tree height) down to a root diameter of 2 mm (fine roots, <2 mm, are operationally defined as belonging to the dead organic matter pool or in the soil organic carbon pool). The living biomass pool is reported for all land-use categories assessed in the field inventory.

7.2.4.2 DEAD ORGANIC MATTER

The dead organic matter pool includes the carbon pools dead wood and litter. Dead wood is defined as fallen dead wood, snags or stumps including coarse and smaller roots down to a minimum “root diameter” of 2 mm. Dead wood of fallen dead wood or snags should have a minimum “stem diameter” of 100 mm and a length of at least 1.3 m. Dead wood of stumps with corresponding roots are reported for Forest land remaining Forest land (and Forest management under the Kyoto Protocol), while fallen dead wood and snags are reported for all relevant land-use categories. Litter includes all non-living biomass not classified as dead wood, lying dead, in various states of decomposition above the mineral or organic soil. This includes the litter, fomic, and humic layers. Live fine roots (<2 mm), are included in litter if found in the O horizon since they cannot be separated during sampling. Coarse litter is defined as dead organic material with a “stem diameter” between 10-100 mm and originating from dead trees. Fine litter from the previous season or earlier is regarded as part of the O horizon.

7.2.4.3 SOIL ORGANIC CARBON

The soil organic carbon pool on forest land and grassland includes all carbon in the mineral soil below the litter, fomic and humic layers in mineral soils and all organic carbon in soils classified as Histosols¹⁹⁷. The carbon pool considered is soil organic carbon down to a depth of 0.5 m measured from top of the mineral soil or, alternatively, from the soil surface when the soil is classified as a Histosol.

¹⁹⁶ Swedish University of Agricultural Sciences, 2011

¹⁹⁷ Food and Agriculture Organization of the United Nations, 1994.

7.2.5 Emissions of N₂O, CO₂ and CH₄, CRF 5(I), 5(II), 5(III), 5(IV) and 5(V)

7.2.5.1 DIRECT N₂O EMISSIONS FROM N FERTILIZATION, CRF 5(I)

To increase the forest production, some middle aged or older forest stands on mineral soils are occasionally fertilized. In 1990, the fertilized forest area was estimated to 69,200 ha¹⁹⁸. Since then, the annual fertilized area has decreased to about 15,000 - 33,000 ha in 1992-2006. In recent years, this area has increased to about 53,000 ha in 2011. The underlying data (areas) are based on an annual questionnaire sent to approximately 150 large-scale forest companies and constitute Official Statistics of Sweden collected by the Swedish Forest Agency. Large-scale forestry are defined as forest companies with more than 10 employees or owners of more than 5,000 ha Forest land., contributes with 98.5 % of fertilizer related emissions of N₂O. Consequently, small-scale forestry is assumed to contribute with approximately 1.5 % of the emissions. To estimate the total annual emission, area figures are multiplied with normal average amount of fertilizer N spread per hectare. The normal average amount N spread per hectare is obtained from companies that are carrying out the fertilization in practice (there are only a few companies in this business).

7.2.5.2 N₂O EMISSION FROM DRAINAGE OF SOILS, CRF 5(II)

According to UNFCCC (decision 13/CP.9), reporting emissions of nitrous oxide from drainage (N₂O-direct $N_{drainage}$) is optional. One reason for that is the limited understanding of the processes controlling the emissions. No N₂O emissions from drainage of soils will be reported this year, but some preliminary studies indicate that reliable methods may be available in a few years.

7.2.5.3 N₂O EMISSIONS FROM DISTURBANCE ASSOCIATED WITH LAND-USE CONVERSION TO CROPLAND, CRF 5(III)

Due to more intensive soil management on Cropland, the transfer of other land-use to Cropland is usually associated with a temporary increase in the mineralization of organic matter. Part of the released N may be converted to N₂O through denitrification. Land converted to Cropland is reported as belonging to the conversion class for twenty years (if no secondary conversion occurs). The accumulated area converted from Forest land to Cropland during 20 years has been up to 5,000 ha and the area converted from Grassland to Cropland is now around 55,000 ha. For the first time, from 2007, we have identified land converted from Wetlands to Cropland. The accumulated area in this conversion class is between 2,000 and 5,000 ha (2007-2011).

¹⁹⁸ Swedsih Forest Agency, 2012

7.2.5.4 CARBON FROM AGRICULTURAL LIME APPLICATION, CRF 5(IV)

Lime is used for soil improvement in both agriculture and horticulture to mitigate acidification that is caused by the export of biomass, acidifying fertilizers and acid rain. The reported figures are based on quantities sold for agricultural and horticultural purposes plus lime from sugar mills and steel production. The quantities are separated into dolomite ($\text{CaMg}(\text{CO}_3)_2$) and limestone (CaCO_3), where dolomite and Mg-lime are reported as dolomite and all other categories are reported as lime-stone. All categories are supposed to contain 100 % dolomite/limestone except residual lime from sugar production which is assumed to contain 65 % limestone due to a water content of approximately 35 %. The accuracy of estimates of the sold quantities is assumed to be high and constitutes Official Statistics of Sweden¹⁹⁹. Separate default IPCC emission factors are used for limestone and dolomite, respectively.

7.2.5.5 N_2O , CH_4 AND CO_2 FROM BIOMASS BURNING, CRF 5(V)

Forest fires are very rare in Sweden. Wildfires have been monitored by the Swedish Rescue Services Agency since 1996²⁰⁰ and the area of wildfires has varied from 400 to 6,400 ha yr^{-1} . Controlled burning after clear-cutting to improve regeneration of trees is monitored by a full record from 1990 and onwards (Swedish Forest Agency). Controlled burning for nature conservation is monitored from 2006. In recent years, an area of approximately 300-3,000 ha is annually burned after clear cutting and 100-2,000 ha is now annually burnt for nature conservation. The Swedish Rescue Services Agency reports the annual area of wildfires for three different land categories: "Forest", "Sparsely covered by trees" and "No tree cover". The definition of "Forest" almost corresponds to the national definition of productive forest. "Sparsely covered by trees" are areas sparsely covered by trees such as mires, forest in the mountain area and parks. "No tree cover" is land with no trees such as agricultural land, open areas but also some mires. The assumed former stock on burned areas is based on estimates of above-ground living and dead biomass inventoried by the NFI by matching national definitions to the definition by the Swedish Rescue Services Agency. The area of wildfires is probably slightly underestimated since the reported numbers only include actual turn-outs by the fire brigade. The accuracy of the burned amount of carbon per land category is probably low. This is due to a lack of knowledge about the burned stock in typically burned forests.

¹⁹⁹ Statistics Sweden, 2004

²⁰⁰ Swedish Rescue Services Agency, 2004

7.3 Methodological issues

7.3.1 CRF-tables 5A, 5B, 5C, 5D, 5E and 5F

7.3.1.1 BASE METHODOLOGY

Sweden reports emission/removals from carbon pools mainly according to the IPCC stock change method. The stock change method is combined with a sample-based inventory design which makes it possible to estimate errors of the estimates. The Swedish National Inventory of Forests (RIS²⁰¹) has monitored the most relevant carbon pools since 1983. A particular advantage with the Swedish NFI is that it has been undertaken using permanent sample plots, on all land use categories, which makes it possible to monitor both gross and net land-use conversions for the six land-use categories in a consistent and transparent manner (for further details, see Annex 3:2).

7.3.1.2 METHODOLOGY LIVING BIOMASS CRF 5A, 5B, 5C, 5D, 5E AND 5F

A national methodology (Tier 3) is used. The above-ground biomass per fractions is estimated by applying Marklund's²⁰² biomass functions to trees on permanent sample plots of the NFI²⁰³. The below-ground biomass is estimated by using Peterssons and Ståhl's²⁰⁴ biomass functions on biomass data from the same trees as for the above-ground biomass. The conversion factor 0.49 is used to convert biomass to carbon²⁰⁵. Estimates of the annual change in the carbon pools are based on repeated measurements. Consequently, the stock change of for example year 2000 is calculated as the difference in stock between year 2000 and year 1999. Since the estimates are based on representative allometric single tree regression functions or on direct measurements, a low risk of bias is assumed.

7.3.1.3 METHODOLOGY DEAD ORGANIC MATTER CRF-TABLES 5A, 5B, 5C, 5D, 5E AND 5F

A national methodology (Tier 3) is used to estimate the dead organic matter pool. The pool includes different sub-pools (dead wood, litter and the organic soil horizon) that are estimated slightly differently.

The inventory of dead wood began in 1994 for northern Sweden and from 1995 for the whole country. The carbon content in dead wood was calculated using conversion factors from volume per decay class to biomass for the species Norway spruce and Scots pine. The volume is measured by the NFI. Below-ground dead wood originating from stump and root systems of harvested trees is reported based on indirect measurements of harvest.

²⁰¹ Swedish University of Agricultural Sciences, 2011

²⁰² Marklund, 1987 and 1988

²⁰³ Ranney et al., 1987

²⁰⁴ Petersson and Ståhl, 2006

²⁰⁵ Sandström et al., 2007

The harvest is estimated based on estimates of growth (stem volume, from measurements of increment bore cores of sample trees) converted to carbon dioxide equivalents minus the net change in the living biomass carbon pool. Growth is estimated by the National Forest Inventory and represents “productive forest land” while the net change in the living biomass pool represents all Forest land (FAO-definition)– consequently, the annual inflow to the stump carbon pool might be slightly underestimated. The harvest of stems is converted by conversion factors to stump and root biomass, and the conversion factors are calculated on estimates of stem volume²⁰⁶ and stump and root biomass²⁰⁷ applied to sample trees representing the standing stock of Swedish forests. The decay of stump systems is modelled²⁰⁸ by simple decomposition functions. The described methodology is consistently used during the reported period. Emissions from stump systems before 1990 are considered by similar methodology (1853-1989). Sweden is discussing options to improve the accuracy of estimates in future reporting²⁰⁹.

The carbon in the litter pool is estimated based on three different sources: (i) coarse litter (ii) annual litter fall and (iii) litter < 2 mm. Coarse litter is defined as dead organic material with a “stem diameter” between 10-100 mm and originating from dead trees. Coarse litter is not inventoried but calculated as 15 % of the above-ground dead wood. Litter fall is calculated using empirical functions based on tree stand properties and litter fall for deciduous species by biomass functions based on leaf biomass. This fraction of litter is regarded as an annual pool. The remaining part of this pool after one year is included in the O horizon and thus measured by the soil inventory. The fine litter (< 2 mm) is estimated by sampling the O or H horizon sample which is taken on an area basis, weighed and analysed for carbon content (for further details, see , see Annex 3:2).

7.3.1.4 METHODOLOGY SOIL ORGANIC CARBON CRF 5A, 5B, 5C, 5D, 5E AND 5F

The soil organic carbon pool is estimated using different approaches depending on the land use. For Forest land and Grassland on mineral soils, estimates are based on repeated soil sampling in combination with pedotransfer functions. For organic forest and grassland soils the changes are based on emission factors and area estimates of different sub-categories. For Cropland the ICBM model^{210, 211} is used to predict changes in the soil organic carbon stock on mineral soils and to calculate the change on organic soils an estimation of the subsidence is added.

²⁰⁶ Näslund 1947

²⁰⁷ Petersson and Ståhl, 2006

²⁰⁸ Melin et al., 2009

²⁰⁹ Petersson and Melin, 2010; Melin et al., 2010

²¹⁰ the Introductory Carbon Balance Model

²¹¹ Andrén & Kätterer, 2001

7.3.1.4.1 *Forest land and Grassland on mineral soils CRF 5A and 5C*

The method is a Tier 3 method. The estimates are based on repeated measurements on the NFI plots of several variables. The basic function used to determine the amount of carbon in a soil layer is based on the amount of carbon in a certain soil layer and the fraction of fine earth. The amount of fine earth is dependent on the bulk density and amount of gravel, stones and boulders in the soil (for further details, see Annex 3:2).

7.3.1.4.2 *Forest land and Grassland on organic soils CRF 5A and 5C*

The method is a Tier 2 method. Changes in the organic carbon pool are calculated as the difference between annual below ground litter input and the heterotrophic respiration. Annual litter production is derived from the National Forest Inventory and the emission factors for drained and undisturbed organic forest soils are based on studies from Sweden and Finland (for further details, see Annex 3:2).

7.3.1.4.3 *Cropland on mineral soils CRF 5B*

The method to estimate the carbon balance of agricultural soils is a Tier 3 method. The carbon changes in the mineral soil are calculated based on data from eight agricultural production regions using the model ICBM-region. The ICBM model is described in Andrén & Kätterer²¹². The calculations are based on daily weather data, annual crop harvest statistics, the use of manure in each region and the results from a nationwide survey of agricultural soils including data on carbon content and texture²¹³(for further details see Annex 3:2).

7.3.1.4.4 *Cropland on organic soils CRF 5B*

The method to estimate the carbon balance of organic agricultural soils is a Tier 2 method. A national emission factor for cropland on organic soils is used to calculate the mean annual carbon loss per cm soil subsidence. The emission factor is modified according to crop type. The relative area proportion of the different crop types and the total area of organic soils under agricultural production were estimated in a national survey²¹⁴ Compared to earlier estimated and reported areas the area is now essentially smaller (for further details see Annex 3:2). The area has been linked to the changes in total cropland area so that decreasing cropland area proportionally affects the area of cropland on organic soils²¹⁵ (for further details see Annex 3:2).

²¹² Andrén & Kätterer, 2001.

²¹³ Eriksson 1997,1999

²¹⁴ Berglund and Berglund, 2009

²¹⁵ In previous submissions a small discrepancy in the area of organic soils reported for cropland remaining cropland in the LULUCF sector and the area of cultivated organic soils reported in the agriculture sector was detected (ERT, centralised review Subm 2009). In this submission the consistency has been improved and areas have been estimated as described in the text.

7.3.1.5 METHODOLOGY FOR DEAD ORGANIC MATTER AND SOIL ORGANIC CARBON FOR CONVERSION BETWEEN LAND-USE CLASSES CRF-TABLES 5A.2.1-5, 5B.2.1-5, 5C.2.1-5, 5D.2.1-5, 5E.2.1-5 AND 5F.2.1-5 CROPLAND ON ORGANIC SOILS CRF 5B

The method to estimate the emission/removals in the DOM – and the SOC pools associated with land use changes is a Tier 2 method. In general (except for dead wood and coarse litter) the carbon stock changes associated with conversion of lands is estimated using an emission/removal factor is used in combination with the areal change in land-use for further details see Annex 3:2).

7.3.1.6 CO₂ EMISSION FROM MINERALIZATION WHEN EXTRACTING PEAT CRF 5D

The method used to estimate CO₂ emission from peat extraction areas is a Tier 1 approach. A limited area of Wetlands (10,000 ha) used for peat extraction is considered managed and reported under Wetlands remaining Wetlands. The reported CO₂ emissions refer to mineralization when extracting peat for fuel and agricultural purposes. The emitted CO₂ [Mton•yr⁻¹] is calculated as the product of the extracted area and an emission factor (for further details see Annex 3:2).

Peat extraction is only ongoing on part of the production area. The peat extraction is usually proceeding many years on the same production area until this area is closed down and restored. Former managed peat land is usually restored by saturation by water or by conversion to Forest land. The water saturation will probably stop most carbon mineralization and Wetlands converted to Forest land is reported under Wetlands converted to Forest land (for further details see Annex 3:2). There are no new activity data for years 2010 and 2011 and the area for year 2009 is assumed also for years 2010 and 2011.

7.3.2 CRF 5(I), 5(II), 5(III), 5(IV) and 5(V)

7.3.2.1 DIRECT N₂O EMISSIONS FROM N FERTILIZATION, CRF 5(I)

A Tier 1 methodology is used and the reported figures refer to $N_2O_{direct\ fertilizer}$ (of N). All fertilization is assumed to occur on Forest land remaining Forest land²¹⁶. In year 1990 nitrate of lime (Ca(NO₃)₂) was the dominant fertilizer but thereafter the fertilizer have been based on 50 % NO₃-N and 50 % NH₄-N. The reported annual $N_2O_{direct\ fertilizer}$ [Gg•yr⁻¹] is calculated as the product of the applied amount and the emission factor (for further details see Annex 3:2).

²¹⁶ ERT (centralized review submission 2009) recommended Sweden to report emissions from organic and mineral soils separately. The methodology is based on the total retailed amount and there is no appropriate statistics available on where the fertilizer is applied.

7.3.2.2 N₂O EMISSIONS FROM DRAINAGE OF SOILS, CRF 5(II)

Not reported (optional).

7.3.2.3 N₂O EMISSIONS FROM DISTURBANCE ASSOCIATED WITH LAND USE CONVERSION TO CROPLAND, CRF 5(III)

A Tier 1 methodology is used. The reported annual N₂O emission from disturbance associated with land use conversion to Cropland (N_2O_{conv} [Gg•yr⁻¹]) is calculated according to equation 3.3.15 in IPCC GPG for LULUCF (IPCC²¹⁷) (for further details see Annex 3:2).

7.3.2.4 CARBON FROM AGRICULTURAL LIME APPLICATION, CRF 5(IV)

Methodology level Tier 1-2 is used for reporting carbon emissions from liming. The reporting is based on consumption studies²¹⁸ and all liming is assumed to occur on Cropland remaining Cropland. The reported annual carbon emission from agricultural lime application (C_{lime} ; [Gg•yr⁻¹]) is calculated as the product of the applied lime and the emission factors (see Annex 3:2 for further details).

7.3.2.5 EMISSIONS FROM BIOMASS BURNING, CRF 5(V)

A Tier 1 methodology and IPCC default emission factors are used. All land categories are monitored but the reported emission is assumed to occur only on Forest land remaining Forest land and on Grassland remaining Grassland. Calculations are based on the amount of biomass per area, burned area and emission factors. Observe that, to avoid double-counting, CO₂ emissions from wildfires and controlled burning are included in carbon stock changes in living biomass (for further details see Annex 3:2).

7.4 Uncertainties and time series consistency

7.4.1 Uncertainties

Since the Swedish reporting system of the LULUCF-sector mainly is based on sampling, a national method is used to estimate the overall uncertainty. Uncertainties in the reported estimates arise from random and systematic errors. Random errors dominate the uncertainty for the part of the living biomass, dead organic matter and soil organic pools that are calculated based on sampling data whereas systematic errors dominate the uncertainty for other emissions/removals. A summary of uncertainties per greenhouse gas is found in Table 7-5.

²¹⁷ Intergovernmental Panel on Climate Change, 2003

²¹⁸ Statistics Sweden, 2004

Random errors could be estimated by straight forward statistical theory but systematic errors are often hard to quantify. Generally for Sweden, the systematic error induced by activity data is small compared to the error due to use of incorrect emission factors. Systematic errors could also arise from missing or overlapping pools. Systematic errors are subjectively judged with help from experts and from default error values according to IPCC²¹⁹.

Table 7-5 Estimated annual net emissions/removals and their corresponding uncertainty (2•relative “standard error”). For categories Living biomass, Dead organic matter and Soil organic carbon, standard errors are based on random sampling. For other categories, standard errors refer to biases that are assumed. Assuming GWP=1 for CO₂, 310 for N₂O and 21 for CH₄, the uncertainty level for the total net removal is estimated to 31 %. Combined uncertainties are calculated according to IPCC, minus=removal

Category	Emission/Removal [Gg•yr ⁻¹]			2•Relative Standard Error [%]		
	CO ₂	N ₂ O	CH ₄	CO ₂	N ₂ O	CH ₄
Living biomass	-33,544	-	-	30	-	-
Dead organic matter	-1,725	-	-	50	-	-
Soil organic carbon	-159	-	-	35	-	-
Direct N fertilization, 5 (I)	-	0.140	-	-	50	-
Drainage of soils, 5 (II)	-	NE	-	-	NE	-
Conversion Cropland, 5 (III)	-	0.230	-	-	100	-
Agricultural lime application, 5 (IV)	79	-	-	50	-	-
Biomass burning, 5 (V)	-	0.001	0.101	-	75	75
All	-35,349	0.371	0.101	31	85	75

7.4.2 Living biomass, CRF 5A, 5B, 5C, 5D, 5E and 5F

The estimated accuracy of the living biomass pool depends mainly on the sample design of the NFI. Results from the control inventory of the NFI indicate that measurement errors, registration errors and errors caused by the instruments (callipers) could be assumed to be close to zero. Potential bias induced by incorrectly specified models and an unrepresentative derivation data are ignored. Unpublished research indicates that the influence of model errors is less than 10%. Estimates for reporting years 1990-2007 are based on approximately 30,000 sample plots and with a corresponding estimated relative standard error of 12 %. Estimates for reporting years 2008, 2009, 2010 and 2011 are based on measurements on approximately 24,000, 18,000, 12,000 and 6,000 sample plots, respectively, combined with extrapolated data. We believe that this extrapolation increases the accuracy substantially but to be on the sure side, to avoid a potentially risk of systematic errors, we gradually update extrapolated data using data from re-measured sample plots (see Annex 3:2 for further details).

²¹⁹ Intergovernmental Panel on Climate Change, 2003

7.4.3 Dead organic matter, CRF 5A, 5B, 5C, 5D, 5E and 5F

Estimates of dead organic matter are based on sampled data from the litter pool and dead wood pool from the NFI and the MI. The sample error for the entire dead organic matter pool is calculated similarly to the living biomass calculation and is given in Table 7-5. There is probably a small error in the estimates of dead wood due to incorrect measured volumes and due to errors connected to the conversion from volume to carbon. Coarse litter is calculated as 15 % of the dead wood. The error of this proportion might be large since the knowledge of the relation between the amount of dead wood and coarse litter is poor. Compared to submission 2012 accuracy has improved since the reported figures now are based on more repeated measurements of permanent sample plots. For changes in carbon in the O-horizon the measurements are based on samples from 1993-1999 (first inventory) and from 2003-2009 (second inventory), while dead wood measurements are from the period 1995 to 2011. We are now also basing the estimate on interpolated values for years between inventories. The accuracy will increase in the future when more data from repeated measurements will be accessible.

One of the major difficulties in reporting changes in DOM and SOC is that the pool is very large and the changes small in comparison to the pool. As seen in Figure 7-7 the reported changes are considerable in terms of carbon and they do have an impact on the national carbon budget. However, the annual changes are still only in the order of a few % of the pool and can hardly be detected in the right panel. When tested statistically the changes are not significant at $p=0.05$ and the system is sensitive to systematic errors like small changes in data collection between inventories.

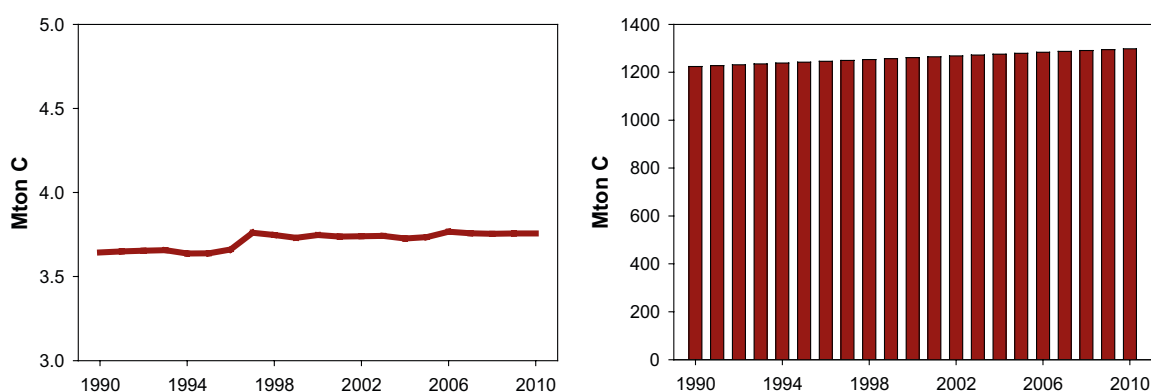


Figure 7-7 The reported change in soil carbon on mineral soils (left panel) and the corresponding stock (right panel) for Forest remaining Forest in the current submission.

7.4.4 Soil organic carbon, CRF 5A, 5B, 5C, 5D, 5E and 5F

The sample error for the soil organic carbon pool is calculated similarly to the living biomass calculation and is given in Table 7-5. The problems associated to the estimates of changes in the soil carbon pool is of the same nature as the ones described for the dead organic matter pool above, and significant improvements are expected when the proportion of repeated measurements will increase. Another problem associated to our methodology is the risk of systematic errors in the sampling and analysis of data. Since there are rather small changes in large pools even a small systematic error may cause a trend in the material. From 2003 the sampling methods of soil samples have been changed compared to earlier inventories in order to avoid subjective judgments in sampling, e.g. regarding determination of soil horizon boundaries. This might give rise to problems of comparability between inventories, but should improve the quality of the data by reducing future risks of systematic errors.

Significant efforts are made to check data and to remove possible sources of error in the field data collection. The uncertainty in activity data (area) for CO₂ emission from drained forest land is judged to 25 % and errors in the emission factor to 25 % as well. The uncertainty in activity data (area) for CO₂ emission from mineralization when extracting peat is judged to 25 % and the uncertainty due to errors in the emission factor chosen is judged to 300 %. The high error of the EF is based on the fact that i) the variation between different emission factors is significant²²⁰, and ii) the underlying data of the EF does not perfectly match the target population^{221, 222}.

7.4.5 Other CO₂ emissions, CRF 5(IV) and 5(V)

The reported CO₂ emission from agricultural lime application is based on consumption studies and the design is regarded as a total inventory with no random error. The error due to activity data is probably small and the reason for this is high quality data on quantities of limestone and dolomite sold. It is assumed that the error due to the use of incorrect emission factors used might be quite large. The reported uncertainty is based on a default error coefficient from IPCC²²³.

Uncertainties from biomass burning arise from the errors in the estimated area that is burned and in the emission factors used. The emitted amounts per area unit depend on the biomass stock before the fire and the proportion of this biomass that actually is burned. The error of the estimated burned area is probably quite small but the knowledge of emitted amount per area is quite poor. The reported uncertainty is based on a default error coefficient from IPCC.

²²⁰ Statistics Sweden, 2002

²²¹ Kasimir-Klemedtsson et al., 2000

²²² Sund et al., 2000

²²³ Intergovernmental Panel on Climate Change, 2003

7.4.6 N₂O and CH₄ emissions, CRF 5(I), 5(III) and 5(V)

Generally for all N₂O and CH₄ emissions, the error in activity data is small compared to the error due to errors associated to the emission factors.

For N₂O emissions from N-fertilization, the error due to activity data is judged to 3 % (the Swedish Forest Agency) and the default total error to 25 %. However, a recommendation is that emission factors chosen should be within the range 0.25 % to 6 % and the interpretation is that a badly chosen emission factor could lead to an error that is much larger than 25 %. Based on this information a total error of 50 % for N₂O emissions from N-fertilization is suggested.

The accuracy of estimates of N₂O emissions from disturbance associated with land-use conversion to Cropland is assumed to be lower than for N₂O emissions from N-fertilization. This is because it is assumed that the error of the activity data (ΔC from mineralization) is higher and due to a large potential error in the selected C:N-ratio. The uncertainty level is based on this reasoning and on IPCC default values (IPCC).

According to the points raised in the discussion above on uncertainties in CO₂ emissions from biomass burning, the uncertainty of N₂O and CH₄ emissions from biomass burning are assumed to be 75 %, (Managing uncertainties: A.1.4).

7.4.7 Completeness

Each category has been reported only once. This is ensured by using only one source of information for the land area representation of carbon pools.

Sweden reports carbon stock changes in all carbon pools and all other emissions for mandatory categories considered managed (Forest land, Cropland, Grassland, Settlements and a small area of Wetland used for peat extraction). The notation key “NO” is used when there is no observed occurrence for a certain category (i.e. uncommon land use changes). The notation key “IE” is used when it is not possible to separate emissions/removals on relevant land use categories and according to the use of the stock change methods. In the latter case either gains or losses are reported “IE”. The notation key “NA” is used when the reported activity does not result in emissions/removals. This notation key is also used for emissions/removals from unmanaged land. The notation key “NE” is only used for categories not estimated because they are currently optional to report. Harvested wood products are not reported (optional) and factoring out has not been considered.

7.4.8 Time series consistency and verification

The time series of change in carbon stock for the living biomass pool is consistently measured from 1990 and onwards. The trend has been validated and confirmed by the default method (growth minus drain) but the level of the annual net removals could not be verified. We assume that most of the discrepancy could be explained by the basic biomass expansion factors applied using the default method. The time series for the dead wood pool extrapolates data in the beginning of the period and this because the inventory did not begin until

the mid-1990th. Due to a relative high sampling error, a trend is reported and it is quite difficult to match emissions/removals from dead wood to the correct year. The dead wood pool constitutes a net removal and this could partly be explained by the fact that an increasing amounts of dead wood and snags have been left after harvest, however, no proper validation has been made.

The time series of the dead organic matter pool is consistently measured since 1994 with only minor changes in sampling methodology. The soil organic carbon has been sampled annually since 1993. In 2003 a revision of sampling methodology was made to harmonize sampling with international monitoring programs. Studies on the effects of these changes in sampling have not revealed any systematic differences with respect to soil carbon pool estimates. The time series for dead organic matter and soil organic carbon in forests have been compared to results from two process-oriented models. Models and measurements agreed well in estimation of the soil carbon pool and in the direction of change, but there were small changes with respect to the rate of change between the models and the measurements²²⁴.

7.5 QA/QC

7.5.1 Quality assurance

The quality assurance system of the data collection within RIS used for the UNFCCC and Kyoto reporting has been described by the Swedish University of Agricultural Sciences²²⁵. These routines were improved during 2006 cooperating with SLU (Swedish University of Agricultural Sciences). SLU also works closely with the Swedish EPA to enhance the QA/QC. For this submission, quality assurance has been carried out in an internal review by experts at SLU. A national independent review has been carried out by representatives for the Swedish Forest Agency and Swedish Board of Agriculture.

7.5.2 Quality control

An internal quality control has been performed following level Tier 1, (Table 5.5.1 in Good Practice Guidance 2003).

For reported activity data, descriptions of definitions, description of underlying models, description of sampling design and emission factors used were studied and no errors were found. This was also valid for descriptions of land areas, eventual transcription errors and references. Both calculations and units of estimates were cross checked and judged as reasonable. Original data from the NFI constitute official statistics of Sweden and were not checked. All data (and methodologies used) is archived by the SLU.

²²⁴ Ortiz C. et. al. 2009.

²²⁵ Karlton, E. et. al.. 2005.

7.6 Source-specific Recalculations

Recalculations can be divided into four categories of which the two first ones can be considered “ordinary” recalculations due to the applied methodology using random sampling.

The first category is recalculations due to updated NFI-data which mainly affects the estimates for the previous four years as described in section 7.2.2.2. Small corrections of historical land use changes may affect estimates for earlier years, especially for categories using area as activity data.

The second category is recalculations related to extended datasets for litter and soil from the MI. Since the whole dataset is included using extrapolation and interpolation techniques this may generate updated data for the entire time series.

The third category is when new activity data (not related to NFI or MI) or emission factors have become available (i.e. better sales statistics, information on biomass burning or emission factors related to land-use change).

The fourth category is when the methods have been improved.

In the current submission, the living biomass pool (also valid for areas per land use category), land use areas and areas subject to land use transfers have been recalculated for the years 2007-2011 to improve accuracy and each estimate are now based on 6,000 more sample plots and incomplete inventory cycles have been extrapolated to 2011, see also section 7.2.2.2. Minor corrections of single plots have been made and that is why also small deviations from former submissions occur also for years 1990-2007. The effect of this annual recalculation on Living biomass on Forest land remaining forest land is illustrated in figure 7-8.

The pools dead organic matter and soil organic carbon on mineral soils on Forest land remaining forest land and Grassland-remaining-Grassland have been recalculated for the whole time series from 1990 to 2011 due to introduction of more re-inventoried sample plots.

To increase transparency, peat extraction areas are now reported separately from other Wetlands under Wetland remaining wetland²²⁶. There are no new activity data for peat extraction areas for years 2010 and 2011 and the area for year 2009 is used also for years 2010 and 2011.

Recalculations occur also for non-carbon pools due to slightly adjustments in activity data (areas), but these re-calculations are very small. No recalculations have been made for Nitrogen emissions from Nitrogen fertilization (5I) but data for 2010 and 2011 are preliminary. Due to recalculated estimates of areas, emissions from disturbance associated with land use conversion to Cropland (5III) have been updated and emissions for land converted from Wetlands to Cropland, occur for the first time.

Some rounding errors have been corrected (general for both carbon and non-carbon pools). Inconsistencies in the reporting of direct N₂O emissions for N fertilization and N₂O emissions from disturbance associated with

²²⁶ This disaggregation of land was made as a response to reviewers (ARR 2011)

land-use conversion to cropland between the UNFCCC-reporting and the KP-LULUCF reporting have been taken care of²²⁷. The recalculations are summarized in Table 7.6.

Table 7-6a and b. Recalculations of carbon stock changes and other emissions between submission 2012 and submission 2013 in the LULUCF-sector. Positive numbers indicate an increase in emissions and negative numbers indicate an increase in removals (or a decrease in emissions for categories 5I to 5V).

7-6 a Difference between Submission 2012 and 2012 [Mt CO₂]					
Year	Forest land	Cropland	Grassland	Wetland	Settlement
1990	3.5	0.0	0.6	0.0	0.0
1995	3.3	0.0	0.6	0.0	0.0
2000	2.8	0.0	0.6	0.0	-0.1
2006	2.0	0.0	0.6	0.0	-0.1
2007	5.4	0.0	0.7	0.0	-0.3
2008	3.8	0.0	0.5	0.0	-0.3
2009	2.6	0.0	0.6	0.0	-0.3
2010	2.5	0.3	0.7	0.0	-0.2

7-6 b	Total carbon pool changes [M ton CO ₂]			Other emissions [M ton substance]						Total	
				Fert.	To CL	Liming	Biomass burning				
	LB	DOM	SOC	5 (I)	5 (III)	5 (IV)	5 (V)			[M ton CO2-eq]	[%]
Year				N ₂ O	N ₂ O	CO ₂	CO ₂	N ₂ O	CH ₄		
1990	0.1	2.6	1.3	0.0	0.0	0.0	NA	0.0	0.0	4.1	-10
1995	-0.1	2.7	1.3	0.0	0.0	0.0	NA	0.0	0.0	3.9	-11
2000	-0.2	2.6	1.0	0.0	0.0	0.0	NA	0.0	0.0	3.4	-9
2006	-0.9	2.9	0.5	0.0	0.0	0.0	NA	0.0	0.0	2.5	-7
2007	3.5	1.7	0.4	0.0	0.0	0.0	NA	0.0	0.0	5.7	-15
2008	1.8	1.6	0.5	0.0	0.0	0.0	NA	0.0	0.0	4.0	-11
2009	0.8	1.6	0.4	0.0	0.0	0.0	NA	0.0	0.0	2.9	-8
2010	0.8	1.7	0.9	0.0	0.0	0.0	NA	0.0	0.0	3.4	-10

0 equals value less than 0.5.

²²⁷ The inconsistency now taken care of was observed by reviewers (ARR 2011)

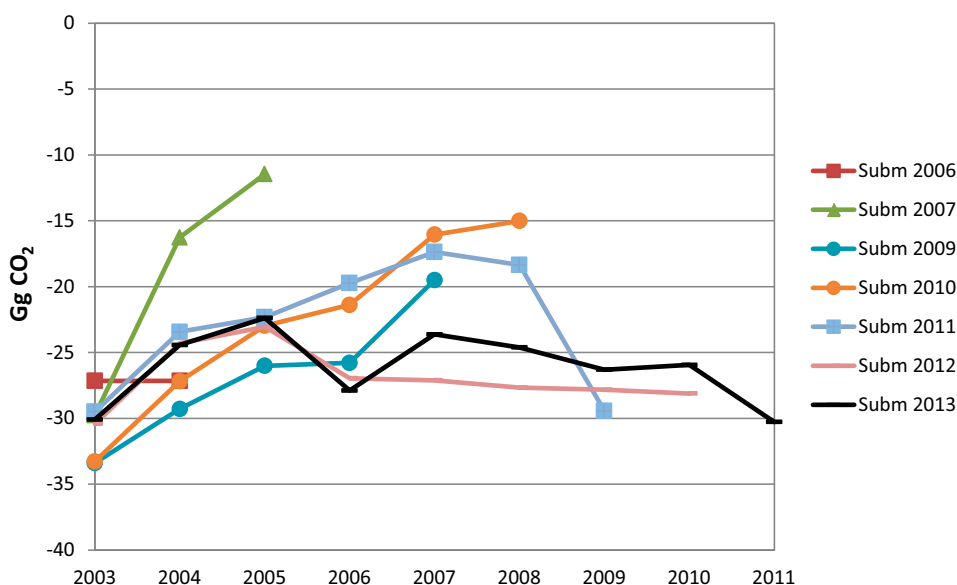


Figure 7-8 Reported living biomass on Forest land remaining forest land (5A1) according to different submissions. The values (the five latest reported years) are continuously recalculated (due to a resubmission, submission 2008 correspond to submission 2009).

7.7 Coming improvements

7.7.1 Informal reporting of HWP

On an informal basis, Sweden reports emissions/removals from Harvested Wood Products (HWP) using the Production Approach (PA) according to decisions taken in Durban, decision 2/CMP.7. Thus, emissions from HWP are estimated as changes in the carbon pool of HWP in use originating from Swedish forests. A Tier 3 model was used for the calculations.

Input data originated from national data sources; Swedish Forest Agency, SDC (Forest industry information association), Statistics Sweden, and Swedish Forest Industries Federation. The data used concerned production and trade of raw material and primary products and covered different length of time. Data about harvested round wood and sawn wood for instance covered 1850-2011.

The model calculates net-emissions of CO₂ by estimating differences in the size of the HWP-carbon pool between years. Equation 12.1 in IPCC 2006 Guidelines was used. Separate calculations were made for different product categories: sawn wood, wood based panels, and paper products. A HWP-carbon pool for a certain year was estimated by adding an inflow of new products to, and subtracting an outflow (decay) of consumed products from the pool of previous year. The difference between the years was translated into CO₂ emissions or CO₂ uptake.

Inflow was calculated by adding primary products made from exported raw material of domestic origin such as round wood, chips and pulp, to domestically produced products of domestic origin. Domestic production from imported raw material was excluded at each step along the refinement chain. Production of primary paper products based on recovered paper is

excluded since domestically collected recovered paper might originate from imported paper. Excluding recovered paper also ensures that carbon in paper is not calculated as inflow more than once.

A first-order decay was assumed, i.e. it is proportional to the size of the carbon pool, and is calculated using half-life (number of years until 50 % is consumed) as input variable.

Half-times for sawn wood, wood based panels and paper products were set to 35, 25 and 2 years respectively, according to decision 2/CMP.7. The net-emission of CO₂ varied between -8.47 and -2.47 M ton during 1990-2011 (Figure 7-9 and Table 7-7).

Table 7-7 Net removals (-) from the HWP-pool originating from Swedish forests 1990-2011.

Year	1990	1995	2000	2001	2005	2006	2007	2008	2009	2010	2011	average
M ton CO ₂	-4.15	-5.02	-6.44	-4.59	-7.5	-8.47	-8.34	-5.32	-3.88	-4.09	-3.69	-5.14

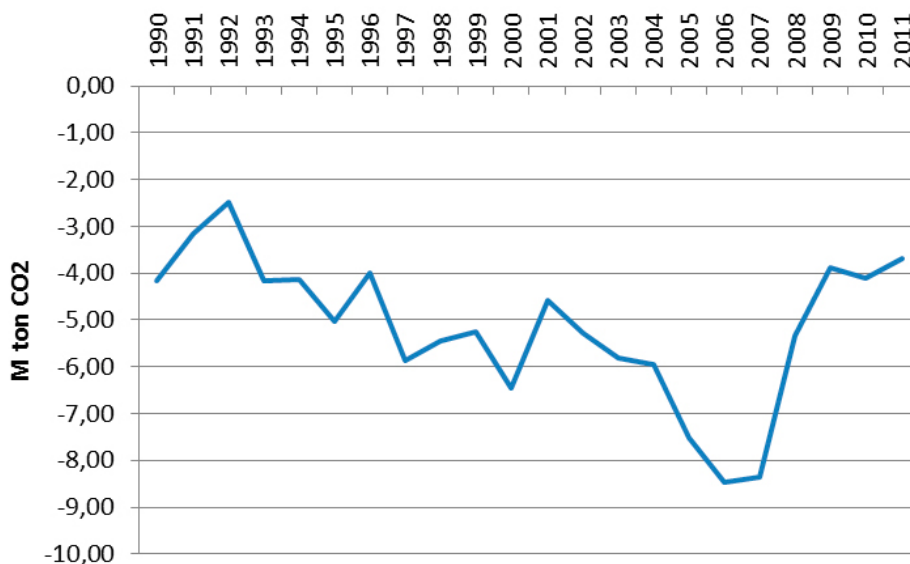


Figure 7-9 Net emissions of CO₂ from the HWP-carbon pool originating from Swedish forests during 1990-2011.

8 Waste (CRF sector 6)

8.1 Overview of sector

In this sector, the most important emissions of greenhouse gases are those of CH₄ from solid waste landfills, CRF 6.A. Minor categories are the subcategories of wastewater handling, CRF 6.B, from where N₂O and CH₄ are reported. Emissions of CO₂, NO_x, SO₂ and NMVOC are reported from waste incineration, CRF 6.C. No emissions are reported in CRF 6.D.

For all greenhouse gases together, the trend over the last ten years has been a constant reduction of emissions (Figure 8-1). For CH₄, the trend can be explained by decreasing quantities of organic waste deposited at landfills. Also, the quantities of recovered landfill gas were increasing until from 1990 until 2003. For N₂O there has been a reduction in the quantity of nitrogen discharged from municipal wastewater treatment plants from the mid-1990s when nitrogen treatment in wastewater treatment plants in Sweden was developed. Emission of greenhouse gases from waste incineration is small about 4 % of the emission of CRF 6 in 2011.

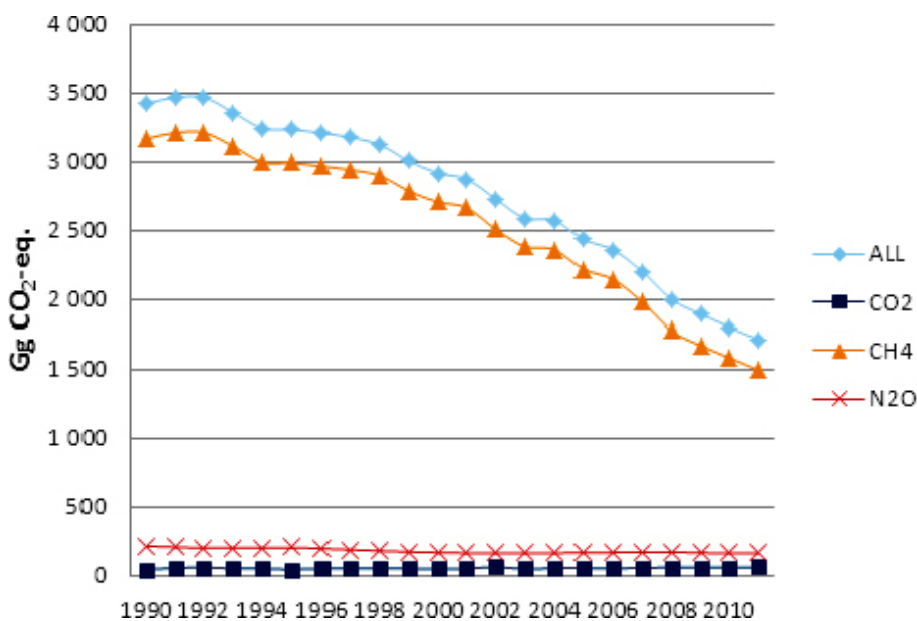


Figure 8-1. Total emissions of all greenhouse gases calculated as CO₂ equivalents from CRF 6 Waste.

Figure 8-2 shows that greenhouse gas emissions from the Waste sector (CRF 6) largely come from solid waste disposal on land (CRF 6.A). CH₄ in sub-sector 6.A represents between 84.2 % and 67.4 % of totally reported greenhouse gases in the Wastewater sector during the period 1990 – 2010.

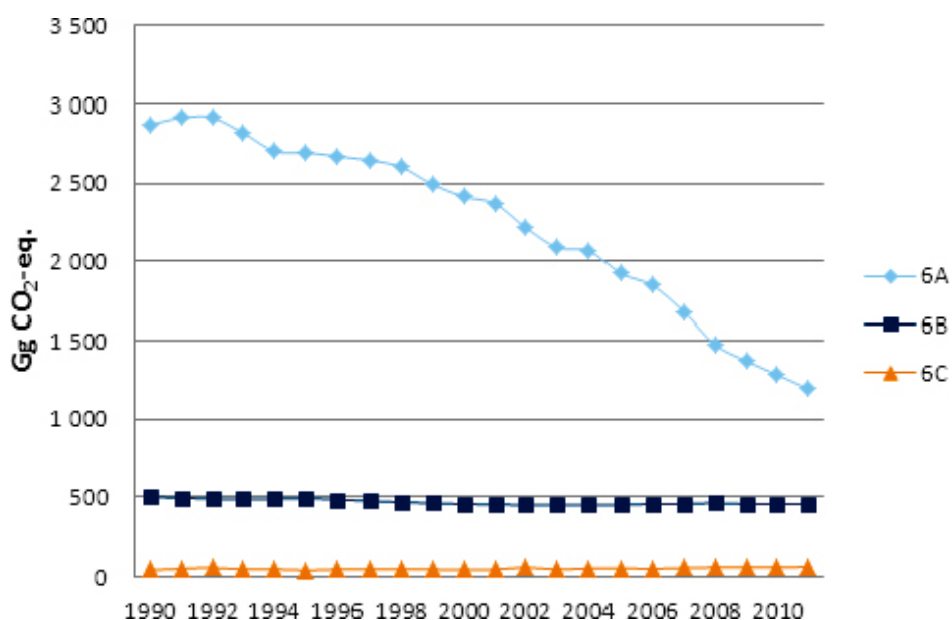


Figure 8-2. Total emissions of all greenhouse gases calculated as CO₂ equivalents from the different Waste sub-sectors.

Table 8-1 shows the impact of recalculations reported in submission 2012 for GHG emissions by sector and sub-sectors for 1990, 1995, 2000 and 2005-2009. The recalculations are mainly due to new data for protein consumption in human sewage (CRF 6.B.2.2). More detailed descriptions of the recalculations are found under sector specific sections below.

Table 8-1. Impact of recalculations of GHG emissions submission 2013 in the waste sector.

Impact of recalculations submission 2013 (Gg CO ₂ eq.)					
CRF	6.A	6.B	6.C	Total CRF 6	% CRF 6
1990	NA	NA	0	0	0.0%
1995	NA	NA	0	0	0.0%
2000	NA	NA	0	0	0.0%
2005	NA	NA	-39	-39	-1.6%
2006	0	NA	-22	-22	-0.9%
2007	1	NA	-49	-48	-2.1%
2008	4	NA	-66	-62	-3.0%
2009	5	-3	-50	-48	-2.5%
2010	4	-6	-48	-50	-2.7%

0: value less than 0.5. NA: no recalculation is performed.

8.2 Solid waste disposal on land (CRF 6.A)

Waste management in Sweden has been developed considerably over the past twenty years. Legislation, such as the implementation of EU directives and national tax policies in the waste management field, has forced and encouraged investments in new technical solutions and waste treatment methods. There has been a comprehensive extension of the treatment capacity of Swedish incineration plants for household waste (with energy recovery) and development of waste management practices other than solid waste disposal on land (landfilling).

Since Sweden is a country with a developed mining and quarrying industry, mining waste is by far the most dominating single waste category in generation of waste and landfilling. In year 2008, 93 % of the landfilled waste (or 43.9 Mt of 47.2 Mt) was mining waste.

In the 1990s, the amount of deposited waste (other than mining waste) decreased significantly. This is especially notable for household waste (in Sweden also referred as “Municipal waste”), which is the largest contributor of greenhouse gases of all waste categories. Only 0.9 % of the generated household waste was deposited in 2011²²⁸ which can be compared with 43.8 % in 1990. The remaining part of the generated household waste in 2010 was either incinerated (51.4 %), recycled (32.8 %) or treated biologically (14.9 %).

²²⁸ Avfall Sverige / Swedish Waste Management 2012

Depositing has become an expensive waste management solution for disposal of waste. Since January 1st 2000, there is taxation on depositing, currently 435 SEK²²⁹ per ton of waste liable to taxation. Another important change is the implementation of the national prohibitions on the landfilling of burnable and organic wastes in the 9-10 §§ of the Landfill Ordinance (2001:512). The landfilling of combustible wastes has been prohibited since 2002, and in 2005 the ban was extended to organic wastes. These prohibitions are regulated in more detail through regulation NFS 2004:4 from the Swedish EPA.

In the end of 2008 a new EU regulation for deposition came into force and almost 50 % of landfills for municipal waste were closed, according to the trade association Avfall Sverige – Swedish Waste Management. Landfill was conducted at 79 sites during 2011.

In year 2011, landfill gas was extracted at 57 landfills²³⁰ whereof 46 were active landfills. According to a survey²³¹, the production of biogas in Sweden in 2010 was totally 1,387 GWh (or 99.5 Gg in methane). 21.5 % of the produced biogas was produced at landfills. The biogas production (collected gas) on landfills decreased by 46.9 % from 2003 to 2011, since the amounts of deposited organic waste has decreased significantly the past years, due to the implementation of waste treatment policies.

Biogas from landfills is mainly used for heating but also for production of electricity. Currently, none of this gas is used as vehicle fuel because of the difficulties to upgrade the gas to sufficient quality. About 11 % of the biogas produced (collected gas) at landfills was flared in 2011.

Sweden is reporting emissions of methane for CRF 6.A.1 (managed waste disposal sites). For CRF 6.A.2 (unmanaged waste disposal sites), Sweden is reporting NO (not occurring), since there are no known unmanaged disposal sites for organic waste or municipal solid waste in use²³².

Practises regarding landfills were regulated in 1969. Since this, the unmanaged (or illegal) landfills are very uncommon in Sweden.

Sweden has some concerns about the unmanaged waste: *littering*. This occurs in particular around recycling stations. Other kinds of littering of organic waste are the disposal of smaller amounts of garden waste from households in nature or that residuals from the hunted animals are disposed in situ. When littering is discovered however, the clean-up is performed or the cost for the clean-up is paid by the responsible operator. If the responsible operator cannot be found, the relevant municipality is responsible to perform the clean-up of the site.

²²⁹ SFS 1999:673

²³⁰ Avfall Sverige / Swedish Waste Management 2011

²³¹ Swedish Energy Agency, 2011

²³² Nygren, 2010

8.2.1 Managed waste disposal on land (CRF 6.A.1)

8.2.1.1 SOURCE CATEGORY DESCRIPTION

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 8-2.

Table 8-2. Summary of source category description, CRF 6.A.1.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
6.A	CO ₂				NA	NA	NA
	CH ₄	X	X		T2	CS, D	Yes
	N ₂ O				NA	NA	NA

CS Country Specific. T2 Tier 2. D Default.

8.2.1.2 METHODOLOGICAL ISSUES

The decrease in deposited waste quantities reduces the potential of methane emissions from landfills. Figure 8-3 shows the methane emissions calculated by the IPCC default model and the IPCC First Order Decay (FOD) model respectively.

The two methods are not really comparable. According to the default model, there is a rapid decrease that immediately follows the decrease in deposited waste. By using this model, the annual landfill gas potential is calculated, rather than the actual gas emissions. The gas emission value for 2011 is negative (-3.9 Gg) since the quantity of recovered gas exceeds the landfill gas potential for waste deposited the same year.

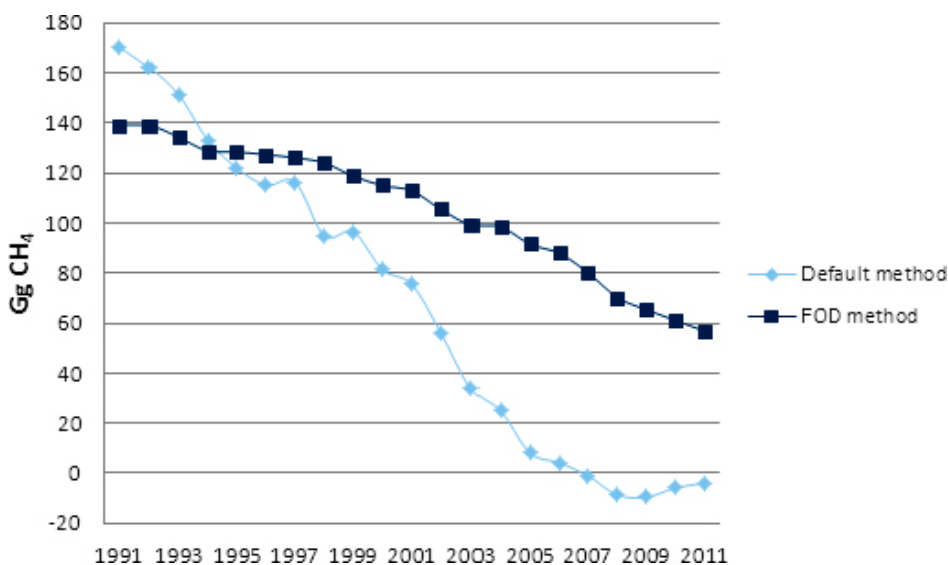


Figure 8-3 Emission of methane from Swedish landfills 1990-2011, estimated by the two IPCC methodologies, (Gg CH₄).

The FOD model, on the other hand, uses a time factor representing the delay in methane production, which results in a slower decrease of emitted methane. The estimates of the FOD model are used in the Swedish National GHG Inventory. In Table 8-3, and Table 8-4 the estimates from the Default model, the FOD model and the deposited amount of municipal solid waste (MSW) are presented.

Table 8-3 Methane emission from Swedish landfills according to IPCC Default and FOD methods. Deposited MSW*, sludges and total (excl. mining waste), 1990-2005

Year	Gas emissions Default method Gg CH ₄	Gas emissions FOD method Gg CH ₄	Deposited MSW* in 1,000 tonnes	Deposited sludge from wastewater handling and pulp industry in 1,000 tonnes	Total deposited waste (excl. mining waste)** in 1,000 tonnes
1990	185	137	2,323	1,400	5,563
1991	170	139	2,223	1,262	5,161
1992	162	139	2,203	1,174	4,977
1993	151	134	2,199	1,086	4,824
1994	133	129	2,166	860	4,547
1995	122	128	1,974	850	4,330
1996	115	127	1,856	880	4,145
1997	116	126	1,842	975	4,203
1998	95	124	1,678	700	3,868
1999	96	119	1,756	620	3,853
2000	81	115	1,529	587	3,720
2001	76	113	1,488	514	3,488
2002	56	105	1,338	341	3,006
2003	34	99	1,034	223	2,688
2004	25	98	810	113	2,380
2005	8	92	541	58	2,067

* Includes household and similar waste, park and garden waste, industry- and non-industry specific waste (organic fractions), construction and demolition waste (organic fraction).

** Includes household and similar waste, park and garden waste, industry- and non-industry specific waste (organic and inorganic fractions), construction and demolition waste (organic and inorganic fractions) and sludge from wastewater handling and pulp industry.

Table 8-4 Methane emission from Swedish landfills according to IPCC Default and FOD methods, deposited solid waste (containing Degradable Organic Carbon), sludges (containing DOC) and total (incl. mining waste), 2006-2011

Year	Gas emissions Default method Gg CH ₄	Gas emissions FOD method Gg CH ₄	Deposited solid waste (containing DOC)* in 1,000 tonnes	Deposited industrial effluent sludges and common sludge* in 1,000 tonnes	Total deposited waste (excl. mining waste)* in 1,000 tonnes	Total deposited mining waste* in 1,000 tonnes
2006	4	88	1,249	180	4,143	61,820
2007	-1	80	1,144	144	4,260	60,450
2008	-9	70	1,039	108	4,376	59,080
2009	-10	65	871	141	4,376	59,080
2010	-6	61	648	164	3,300	47,200
2011	-4	57	648	164	3,300	47,200

* Activity data and statistics for 2006, 2008 and 2010 are from Swedens reporting to the Commission according to the Waste Statistic Regulation. Activity data for 2007, 2009 and 2011 are interpolated/extrapolated values.

Table 8-3 presents emissions and waste data used for the years before 2006. The waste data are from various sources and uses national waste categories. These waste categories are different from the ones used currently in Sweden.

Table 8-4 is presenting emissions and waste data used from 2006. The waste data are from Sweden's reporting to the Commission according to the Waste Statistic Regulation and uses waste categories as defined in the regulation.

8.2.1.2.1 *National application to IPCC First Order Decay (FOD)*

The method used for estimating methane emissions from municipal solid waste is the Tier 2 methodology, the IPCC First Order Decay model, with a slightly different time factor and with some estimates on the national gas potentials. The time factor year i , is calculated as:

$$\begin{cases} 1 - e^{-0,5k}, & i = 0 \\ e^{-k(i-0,5)} \cdot (1 - e^{-k}), & i = 1, 2, \dots \end{cases}, \text{ where } k \text{ is the generation rate constant.}$$

This model corresponds to the assumption²³³ that all waste is deposited on 1 July, which is approximately equivalent to a uniformly distributed deposition.

Comparisons between the suggested IPCC gas potentials and Swedish estimates show that the IPCC values tend to be higher, but considering the large methodological uncertainties, which is the same in both cases, the difference should be within a reasonable interval.

Historical data has been extrapolated five half-life periods back in time, which means that, for the calculations of 1990, all deposited gas potentials since 1952 are considered. All available historical information on national deposited quantities is used in the calculation. The quality of data on household waste is high since 1980, but data on organic industrial waste is scarce. The consequence is that many assumptions on historical deposited waste quantities have been made, which have greater impact on the calculated emissions in 1990 than in 2011.

Methane potentials

IPCC values for gas potentials are used for the different fractions of household waste, as well as garden waste. As noted above, these values are somewhat higher than Swedish estimates, but lie within a reasonable interval.

The IPCC gives no gas potential for deposited sludge (already treated, for example, by rotting) from wastewater treatment. The content of Degradable Organic Carbon (DOC) in sludge from wastewater treatment is approximately 7 per cent.²³⁴ The gas potential of the sludge is reduced by 50 % because it is treated.²³⁵ By using formulas given in Good Practice Guidance the gas potential can be calculated to 24 kg/tonnes of sludge.

²³³ Statistics Sweden, 2005b

²³⁴ Recounted from RVF, 1996.

²³⁵ Sweco Viak, 2000.

For wastewater sludge from the pulp industry, a national value of 45 kg methane /tonnes of waste is used.²³⁶

Recovered gas

Since gas recovery can be of importance for the final emissions of methane, Good Practice Guidance recommends formulas that subtract the recovered gas from the produced gas. In Sweden the first plant for biogas extraction from landfills was started in 1983. The business has increased until 2003 when gas was recovered in 72 plants. Since 2009, about 57 gas plants are in operation, and the amount of recovered gas is now decreasing because of the dramatic reduction of deposition of organic waste. Information on recovered gas (in energy units) is provided by Avfall Sverige and converted to quantity (in tonnes) by Statistics Sweden (Table 8-5).

Table 8-5 Recovered methane from landfill gas, tonnes.

Year	Recovered gas
1982	0 ¹
1983	NE ²
1990	12,000 ³
1991	12,210 ³
1992	14,430 ³
1993	20,800 ⁴
1994	27,500 ⁴
1995	30,000 ⁴
1996	30,000 ⁵
1997	30,000 ⁵
1998	30,000 ⁵
1999	33,000 ⁵
2000	34,000 ⁵
2001	32,400 ⁵
2002	35,947 ⁵
2003	36,449 ⁵
2004	30 1,3 ⁵⁵
2005	29 4,1 ⁸⁵
2006	24 5,6 ⁷⁶
2007	24 5,5 ³⁶
2008	26 9,7 ⁹⁶
2009	24 2,4 ⁰⁶
2010	21 4,3 ⁹⁶
2011	19 3,4 ⁴⁶

1) No gas recovery. 2) 1st plants started. 3) Swedish EPA/RVF. 4) RVF, 1996c. 5) RVF, 1997-2006. 6) Avfall Sverige (Swedish Waste Management), 2007-2011

²³⁶ Swedish EPA, 1993.

Table 8-6 shows quantities of produced energy from landfill gas and how much that is flared in Sweden. The energy is used for production of electricity and for heating.

Table 8-6 Energy recovery and flaring at landfills in Sweden, MWh²³⁷.

Year	2005	2006	2007	2008	2009	2010	2011
Energy recovery	340,000	282,200	290,100	310,800	294,240	262,200	237,400
of which is used for production of electricity	20,000	20,800	22,600	23,700	17,400	20,400	16,000
Flaring	70,000	60,200	52,100	65,100	43,600	36,600	32,200
Total	410,000	342,400	342,200	375,900	337,840	298,800	285,600

Other parameters

The Methane Correction Factor for modern Swedish landfills is equal to one (1.0) (Table 8-7). Waste management was centralised during the 1970s. Before 1980, landfills were smaller and presumably less compact. Information that helps establish the MCF (Methane Conversion Factor) (cover material, mechanical compacting and levelling of waste) is missing. For calculations before 1980 the IPCC default value 0.6 was used.

The IPCC default value 50 % is used for the methane content in landfill gas (F) (Table 8-7). The value of DOC_F 0.5 has been chosen according to IPCC methodology.

The oxidation factor is estimated to be 10 %, and the half-life of the methanogenesis is 7.5 years.²³⁸ The choice of the half-life factor has also been motivated by the rather wet climate conditions in Sweden (MAP/PET>1), and that the 2006 IPCC Guidelines recommends the default value of 7 for such climate conditions.

Table 8-7 Other used parameters in the methane emission calculations.

Parameter	Value	Motivation
MCF -1979	0.6	IPCC Default
MCF 1980-	1	Well managed(*)
F	50 %	IPCC Default
DOC_F	0.5	IPCC Default
OX	10 %	National(**)
$t_{1/2}$	7.5 years	National(***)

(*) Swedish EPA, 1999b, (**) Swedish EPA, 1997b, (***) Swedish EPA, 1993b.

²³⁷ Avfall Sverige (Swedish Waste Management)

²³⁸ Börjesson, 2000

Until about 1975, waste burning at landfills was a common waste treatment method, but it ceased about five years later. There is no information on the waste fraction that was burned, except that burning was practiced at 311 of the 847 landfills in 1975.²³⁹ An assumption is therefore made that before 1976, 37 % of all deposited household waste was burned.

8.2.1.2.2 *Waste statistics in sweden, 1980 - 2005*

The Swedish EPA made the first national survey in Sweden in 1980, collecting data on deposited waste (only for household waste and similar). Statistics Sweden collected similar data in 1985, 1990 and 1994. Since 1994, the Swedish Waste Management (former RVF) has carried out an annual survey on deposited waste. Thus, household waste is the best documented waste category, with high quality data available since 1980. Household waste is also the most important category for methane production in landfills. Statistics on deposited sludge from households and park and garden waste are available since 1990. Standard values on fractions of deposited household waste from 1970 and 1975 are also available at the Swedish Waste Management.

Statistics on organic waste from industries are much scarcer. There is information on industrial waste from the 1980s but organic fractions were not specified. The official statistics from 1993 and 1998 on waste from manufacturing do not emphasize generation and treatment of organic waste. Dedicated studies on quantities and treatment of biological waste from industry were carried out in 1993 and 1996 by the Swedish EPA. According to these studies, deposited sludge from the pulp industry has previously been the most important organic deposited industrial waste category. This waste category is also documented by surveys, carried out regularly until 2000 by the Swedish EPA and later by Swedish Forest Industries Federation. Today, sludge from the pulp industry is incinerated and composted.

There are no time series of data available on landfilled organic industrial waste (except from data on sludge from pulp industry).

8.2.1.2.3 *Waste statistics in sweden, 2006 and onward*

The Regulation of the European Parliament and the Council No 2150/2002 of 25 November 2002 on waste statistics (hereafter referred to as “the Waste Statistics Regulation” or “WStatR”) contains rules for the reporting of waste statistics to the EU. Reporting in accordance with the regulation is to take place every second year. Reporting shall be submitted each time 18 months after the end of the reporting period. The first round of reporting by all member states was completed by 30 June 2006 and concerned waste generation and recovery and disposal of waste for the year 2004. The treatment of waste is to be reported by treatment method for the different types of waste according to the Waste Statistical Nomenclature (EWC-Stat). The method

²³⁹ Swedish EPA, 1983.

of treatment relates to various recovery and disposal operations (“R and D codes”) are compiled into 6 different groups. Group 4, “Disposal operations: Land filling, deep injection, surface impoundment, permanent storage and others”, is relevant for “Solid waste disposal on land, CRF 6A”.

The Swedish EPA is responsible for the reporting in accordance with the regulation. So far, waste data has been reported for the reference years 2004, 2006, 2008 and 2010. No waste statistics on landfilling are compiled for the intermediate years by SEPA.

In 2010, a study²⁴⁰ was carried out in order to analyze possibilities to use the reported waste data to WStatR for the calculations of CH₄ from solid waste landfills. The study recommended implementation of WStatR-data from reference year 2006 and onwards. The advantages of WStatR-data in relation to waste statistics for 1980-2005 are mainly that:

- WStatR-data uses more specific and better developed descriptions of waste classifications.
- It is produced regularly (every second year). Therefore it is to a less extent based on extrapolations of old waste data and expert judgments. This means it is more sensitive for rapid changes in amounts of waste and DOC content.
- WStatR-data has per definition 100 % coverage (completeness).

Relevant waste categories (those who is containing Degradable Organic Carbon) were chosen, and the DOC content of the chosen waste categories was investigated by analyzing the statistical source material in cooperation with waste experts. Interpolations and extrapolations have been made for the intermediate years.

8.2.1.2.4 *Waste categories, 1980-2005*

Household waste, sludge and garden waste

Table 8-8 summarizes the available statistics on household waste, sludge from waste water treatment and garden waste. Interpolation is used for the intermediate years. Before 1990, park/garden waste and sludge from households are assumed to be directly proportional to the population, with the same proportion as in 1990.

²⁴⁰ Edborg, Stenmarck, Sundquist & Szudy, 2010

Table 8-8 Deposited household waste, sludge and garden waste (1,000 tonnes).

Year	Household waste (and similar)	Sludge from waste water treatment, wet weight	Garden waste
1980 ¹	1,450
1985 ²	1,040
1986 ³	1,020
1988 ⁴	1,080
1990 ⁵	1,400	900	70
1994 ⁶	1,380	610	80
1995 ⁷	1,200	540	60
1996 ⁸	1,110	470	70
1997 ⁸	1,150	455	50
1998 ⁹	1,020	490	45
1999 ¹⁰	972.5	490	45
2000 ¹¹	869.5	345	53
2001 ¹²	880	330	44
2002 ¹³	820	215	40
2003 ¹⁴	575	155	33
2004 ¹⁵	380	102	0*
2005 ¹⁶	210	58	0*

1) Swedish EPA, 1983. 2) Statistics Sweden, 1988; RVF. 3) RVF, 1988. 4) RVF, 1990. 5) Statistics Sweden, 1992. 6-16) RVF, 1996-2006.

* Included in household waste from reference year 2004. ** Estimate

The composition of household waste has been investigated in many studies over the years. Ohlsson²⁴¹ presents a historic overview of Swedish investigations, the first of which was carried out in 1977. The time series indicates a rather constant composition of components, except the paper content, which declines during the 1990s. The chosen composition²⁴² for 1990 and 1995 are presented in Table 8-3. The composition in the years between the surveys is interpolated. It should be pointed out that this type of analysis contains an unknown variation, and the source of error may be large. Ohlsson also shows that different studies may differ greatly in methods and results.

In 2005, another overview of household waste composition was published.²⁴³ Different fractions of household waste from southern Sweden have been analysed with the same methodology in 3 different years (1997, 2000 and 2004), see further in Table 8-9.

²⁴¹ Ohlsson, 1998 and REFORSK, 1998

²⁴² Ohlsson, 1998

²⁴³ RVF, 2005

Table 8-9. Content of Swedish household waste, %.

	1990	1995	1997	2000	2004
A, Paper and textiles	33	28	23	25	18
B, Garden/park waste, and diapers	14	14	14	11	13
C, Food waste	40	40	41	39	43
D, Wood	1	1	1	1	1

In Sweden the section of the Ordinance prohibiting the deposition of organic waste as landfill was implemented on January 1st 2005. The waste treatment plants need permissions in order to deposit organic waste.

The impact of the new legislation on the DOC content of deposited household waste has not been investigated and documented, but the waste composition and DOC content of the deposited household waste has probably changed since the analysis from 2004. Separation of organic fractions made by the households should lead to a decrease of the DOC content. The organic fractions are treated by composting and anaerobic digestion. Organic fractions (and other fractions) from the mixed waste generated by households and companies are also separated at waste treatment plants before landfilling.

Industrial waste

As noted above, statistics on deposited industrial waste are not divided into organic waste categories. Special studies of organic waste are considered to be the most important information sources of industrial waste categories. In 2004 a study on deposition of organic waste was carried out by Profu and financed by the Swedish EPA.²⁴⁴ The estimates have been made with information from many different sources, such as national statistics, screening inspections of waste content, information on capacity of energy recovery from waste and extrapolation back in time using the industries part of Gross National Product (GNP). The study shows that great amounts of paper and wood have been deposited in construction and demolition waste, as well as in the category of “non specific” industrial waste.

The first study on “specific” organic industrial waste was published in 1993;²⁴⁵ the waste groups found to generate methane in landfills are presented in Table 8-10. The most important subgroup here is sludge from the pulp industry and the other subgroups are mainly from the food industry. The gas potentials stated in the report are based on literature studies and rotting experiments. The gas potentials are used in the methane calculations for 1990.

²⁴⁴ Profu, 2004.

²⁴⁵ Swedish EPA, 1993

Table 8-10 Organic industrial waste, early 1990s (Swedish EPA, 1993).

Waste category	Produced quantity, 1,000 tonnes/yr	Deposited fraction, %	Deposited quantity, 1,000 tonnes/yr	Gas potential, Mm ₃ CH ₄ /yr
Sludge from pulp industry	1,000	50	500	31.5
Carcasses	8	35	2.8	0.63
Waste from slaughter houses	40	5	2	0.45
Sludge from slaughterhouses	45	8	3.6	0.28
Entrails	30	5	1.5	0.09
Manure from slaughterhouses	10	5	0.5	0.03
Draff	5.5	0.5	0.0275	0.03
Waste from sugar beet industry	100	0.5	0.5	0.02
Waste from potato industry	46	0.5	0.23	0.01
Returned bread	13	3	0.39	0.11
Mycelia waste	2	1	0.02	0.01
Scrows waste	5.5	100	5.5	0.8
Waste from fishing industry		50	0	0.5
Whey	1,000	0	0	0
Tinned foods industry	53	50	26.5	1.55
Total:				
Sludge from pulp industry			500	31.5
Other			43.6	4.5

Data on deposited sludge from the pulp industry is available from a survey carried out annually from 1994 up to year 2000 by the Swedish EPA. In 2004, data on deposited sludge from the pulp industry is taken from the Swedish Forest Industries Federation. Data for the intermediate years have been interpolated. The reports contain detailed information on waste and waste treatment for each pulp and paper producer. Intermediate values (1991-1993) have been interpolated (Table 8-11).

Table 8-11 Values of deposited wastewater sludge from the pulp industry, wet weight.

Year	Quantity 1,000 tonnes/year
1990	500 ¹
1994	250 ²
1995	310 ³
1997	520 ⁴
1998	210 ⁵
1999	130 ⁶
2000	242 ⁷
2001	184 ⁸
2002	126 ⁸
2003	68 ⁸
2004	10.5 ⁹
2005	0 ⁹

1) Swedish EPA, 1993. 2) Swedish EPA, 1995. 3) Swedish EPA, 1996b. 4) Swedish EPA, 1998b. 5) Swedish EPA, 1999. 6) Swedish EPA, 2000. 7) Swedish EPA, 2001. 8) Value interpolated no similar survey carried out. 9) Swedish Forest Industries Federation.

A study on organic industry-specific waste was published in 1996²⁴⁶. In accordance with the report, the deposited waste categories are presented in Table 8-12. The gas potentials were calculated by Sweco Viak.

Table 8-12 Organic Industrial Waste 1996.

Waste category	Deposited quantity, 1,000 tonnes/yr	Gas potential, Mm3 CH₄/yr
Waste from slaughter houses	22.5	0.88
Waste from potato and vegetable industries	11.5	0.64
Total:	34	1.52

Swedish EPA, 1996

The final gas potential is used as gas potentials in the methane calculations for 1996 and later. By using the two reports, values are interpolated between 1990 and 1996.

In addition to the gas potentials from these industries, the gas potentials for paper and cardboard waste from industries, which is not included in the referred reports, have to be added. Information on these gas potentials is extracted from a survey ("Waste from the manufacturing and minerals extraction industries in 1998") made by the Swedish EPA and Statistics Sweden.²⁴⁷ In 1998, about 6,000 tonnes of paper and wrapping material were deposited. This quantity is added each year to the industrial waste already noted.

²⁴⁶ Swedish EPA, 1996

²⁴⁷ Statistics Sweden, 2000

Composition of deposited waste

Table 8-13 illustrates the estimated composition of deposited waste (excl. mining waste) 1990-2005.

Table 8-13 Composition of deposited waste, in per cent.

Year	Paper	Food	Plastic	Glass	Textile	Napkins	Sludge from waste water	Sludge from pulp industry	Wood	Other inert	Other organic
1990	7.1	13.5	2.1	0.6	0.7	1.3	16.2	9.0	0.3	34.9	14.3
1991	7.4	14.6	2.2	0.7	0.8	1.5	15.5	9.0	0.3	34.5	13.6
1992	7.5	15.4	2.3	0.7	0.8	1.5	15.1	8.5	0.3	34.2	13.7
1993	7.5	16.1	2.4	0.7	0.8	1.6	14.5	8.0	0.4	34.1	14.0
1994	7.7	17.2	2.6	0.8	0.9	1.7	13.4	5.5	0.4	35.8	14.2
1995	6.8	15.8	2.4	0.7	0.8	1.6	12.5	7.2	0.3	36.9	15.1
1996	6.3	15.9	2.3	0.7	0.8	1.5	11.3	9.9	0.3	36.1	14.8
1997	5.6	16.0	2.5	0.7	0.8	1.6	10.8	12.4	0.3	35.5	13.8
1998	5.4	15.6	2.4	0.7	0.8	1.5	12.7	5.4	0.3	41.0	14.2
1999	5.2	15.0	2.3	0.7	0.8	1.5	12.7	3.4	0.3	40.7	17.5
2000	5.4	13.5	2.5	0.8	0.7	1.2	9.3	6.5	0.2	45.5	14.6
2001	5.8	14.2	2.7	0.8	0.8	1.2	9.5	5.3	0.2	45.2	14.4
2002	6.3	15.5	2.9	0.9	0.9	1.3	7.2	4.2	0.2	46.9	13.8
2003	5.0	13.0	2.3	0.7	0.7	1.1	5.8	2.5	0.1	55.4	13.5
2004	2.8	10.4	1.8	0.4	0.4	0.9	4.3	0.4	0.1	63.1	15.5
2005	1.9	7.8	1.2	0.2	0.2	0.6	2.8	0.0	0.1	72.2	13.0

8.2.1.2.5 Used statistics on deposited waste, 1952-2011

Used statistics 1952-2005

Table 8-14 shows the activity data 1952-2005 used in the calculations of methane emissions from solid waste disposal on land.

Table 8-14 Overview over used statistics on deposited waste and interpolated/-extrapolated values: Solid waste.

Year	Standard value: Household waste/citizen (kg)	Fraction deposited household waste	Fraction of burned household waste on landfills	Deposited household waste and similar, 1,000 tonnes	Deposited park and garden waste, 1,000 tonnes	Deposited organic industrial waste(**), 1,000 tonnes	Deposited industrial waste (not industry specific), organic fraction(**), 1,000 tonnes	Deposited construction and demolition waste, organic fraction(**), 1,000 tonnes
1952	290	76%	37%	992	58	56	207	63
1953	290	76%	37%	998	59	56	211	64
1954	290	76%	37%	1,005	59	56	215	66
1955	290	76%	37%	1,012	59	56	220	68
1956	290	76%	37%	1,018	60	56	226	70
1957	290	76%	37%	1,024	60	56	232	71
1958	290	76%	37%	1,030	60	56	234	73
1959	290	76%	37%	1,035	61	56	239	75
1960	290	76%	37%	1,041	61	56	250	77
1961	290	76%	37%	1,049	62	56	260	78
1962	290	76%	37%	1,056	62	56	272	80

Year	Standard value: Household waste/citizen (kg)	Fraction deposited household waste	Fraction of burned household waste on landfills	Deposited household waste and similar, 1,000 tonnes	Deposited park and garden waste, 1,000 tonnes	Deposited organic industrial waste(**), 1,000 tonnes	Deposited industrial waste (not industry specific), organic fraction(**), 1,000 tonnes	Deposited construction and demolition waste, organic fraction(**), 1,000 tonnes
1963	290	76%	37%	1,064	62	56	280	82
1964	290	76%	37%	1,072	63	56	301	83
1965	290	76%	37%	1,079	63	56	316	85
1966	290	76%	37%	1,088	64	56	325	87
1967	290	76%	37%	1,096	64	56	330	89
1968	290	76%	37%	1,105	65	56	345	90
1969	290	76%	37%	1,114	65	56	349	92
1970	290	76%(*)	37%	1,122	66	56	364	94
1971	290	76%	37%	1,126	66	56	369	96
1972	290	76%	37%	1,129	66	56	372	97
1973	290	66%	37%	984	66	56	391	99
1974	290	66%	37%	987	67	56	406	101
1975	290	66%(*)	37%(*)	990	67	56	409	103
1976	290	66%	30%	1,109	67	56	452	116
1977	290	66%	22%	1,229	67	56	483	131
1978	290	58%	15%	1,186	67	56	517	145
1979	290	58%	7%	1,292	68	56	593	162
1980			0%	1,450(*)	68	56	628	177
1981				1,400	68	56	632	179
1982				1,300	68	56	627	182
1983				1,200	68	56	551	158
1984				1,100	68	56	579	161
1985				1,040(*)	68	56	595	163
1986				1,020(*)	68	56	602	165
1987				1,050	69	56	615	168
1988				1,080(*)	69	56	624	170
1989				1,240	70	56	630	172
1990				1,400(*)	70(*)	56	622	175
1991				1,390	72	57.1	567	137
1992				1,390	75	58.2	554	126
1993				1,390	77	59.3	558	115
1994				1,380(*)	80(*)	60.3	564	82
1995				1,200(*)	60(*)	61.4	571	82
1996				1,110(*)	70(*)	62.5	536	78
1997				1,150(*)	50(*)	62.5	495	85
1998				1,020(*)	45(*)	62.5	477	73
1999				972.5(*)	45(*)	62.5	580	96
2000				869.5(*)	53(*)	62.5	473	71
2001				880(*)	44(*)	62.5	439	62
2002				820(*)	40(*)	62.5	370	45
2003				575(*)	33(*)	62.5	323	40
2004				380(*)	0(***)	62.5	321	47
2005				210(*)	0(***)	62.5	231	37

(*) Taken from statistical sources. Other values are interpolated or extrapolated.
(**) Estimate. (***) Included in household waste from reference year 2004.

**Table 8-15 Overview over used statistics on deposited waste and interpolated/extrapolated values:
Sludge, wet weight.**

Year	Deposited sludge from waste water treatment, 1,000 tonnes	Deposited sludge from pulp industry, 1,000 tonnes
1952	748	500
1953	753	500
1954	759	500
1955	764	500
1956	768	500
1957	772	500
1958	777	500
1959	781	500
1960	786	500
1961	791	500
1962	797	500
1963	803	500
1964	809	500
1965	814	500
1966	821	500
1967	827	500
1968	834	500
1969	840	500
1970	847	500
1971	849	500
1972	852	500
1973	855	500
1974	857	500
1975	860	500
1976	862	500
1977	865	500
1978	867	500
1979	869	500
1980	871	500
1981	872	500
1982	873	500
1983	874	500
1984	875	500
1985	876	500
1986	881	500
1987	885	500
1988	890	500
1989	895	500
1990	900(*)	500(*)
1991	800	462
1992	750	424
1993	700	386
1994	610(*)	250(*)
1995	540(*)	310(*)
1996	470(*)	410(*)
1997	455(*)	520(*)

Year	Deposited sludge from waste water treatment, 1,000 tonnes	Deposited sludge from pulp industry, 1,000 tonnes
1998	490(*)	210(*)
1999	490(*)	130(*)
2000	345(*)	242(*)
2001	330(*)	184
2002	215(*)	126.3
2003	155(*)	68
2004	102(*)	10.5(*)
2005	58(*)	0(*)
2005	58(*)	0(*)

(*) Taken from statistical sources. Other values are interpolated or extrapolated.

Used statistics 2006-2009

Table 8-16 shows waste statistics for 2006-2009 used in the calculations of methane emissions from solid waste disposal on land. It also shows estimated DOC content for each waste category.

Table 8-16. Overview over used statistics* 2006-2009 on deposited waste and interpolated/extrapolated values, 1,000 tonnes, and estimated DOC content, percent.

EW-Stat code	Description of waste categories	2006(*)	2007	2008(*)	2009	DOC content
03.1	Chemical deposits and residues	C	C	176.946	176.946	2
03.2	Industrial effluent sludges: <i>Dry matter</i>	11.914	11.247	10.580	10.580	9
05.	Health care and biological wastes: <i>Hazardous</i>	C	C	0.004	0.004	8
05.	Health care and biological wastes	C	C	0.010	0.005	8
07.2	Paper and cardboard wastes	38.977	20.637	2.296	1.427	36
07.5	Wood wastes	C	C	1.840	0.949	40
07.6	Textile wastes	0.228	0.600	0.972	0.486	24
09A	Animal and vegetal wastes (<i>excl. 09.11 & 09.3</i>)	11.548	8.803	6.058	6.058	15
09.11	Animal waste of food preparation and products	0.303	0.323	0.343	0.343	15
09.3	Animal faeces, urine and manure	0.372	0.224	0.075	0.038	9
10.1	Household and similar wastes	203.821	161.904	119.986	68.500	18
10.2	Mixed and undifferentiated materials	482.743	352.593	222.442	222.442	3.1
10.3	Sorting residues	311.483	409.541	507.599	393.591	2.5
11A	Common sludges (<i>excl. dredging spoils</i>): <i>Dry matter</i>	26.383	19.763	13.142	19.685	28
Total, wet weight		1,428.879	1,287.740	1,146.601	1,011.530	

* Waste statistics for 2006 and 2008 are from Sweden's reporting to the Commission in accordance to the Waste Statistic Regulation. Waste statistics for 2007 and 2009 are interpolated/extrapolated values.

C: Confidential

Used statistics 2010-2011

Table 8-17 shows waste statistics for 2010-2011 used in the calculations of methane emissions from solid waste disposal on land. The EWC-stat codes as well as the DOC content differs somewhat compared to those in Table 8-16. This is due to changes in the EWC-stat codes implemented in the 2010 year data, which is also mentioned in section 8.2.1.5. Due to the changes in EWC-codes there was also an investigation on the DOC-contents regarding the codes changed²⁴⁸.

Table 8-17. Overview over used statistics* 2010-2011 on deposited waste and extrapolated values, 1,000 tonnes, and estimated DOC content, percent.

EWC-Stat code	Description of waste categories	2010 ^(*)	2011	DOC content
02A	Chemical wastes	85.323	85.323	5
03.2	Industrial effluent sludges: <i>Dry matter</i>	1.282	1.282	12.5
03.2	Industrial effluent sludges: <i>Dry matter Hazardous</i>	7.000	7.000	2
05.	Health care and biological wastes: <i>Hazardous</i>	0	0	8
05.	Health care and biological wastes	0	0	8
07.2	Paper and cardboard wastes	0.577	0.577	36
07.5	Wood wastes	0.057	0.057	40
07.6	Textile wastes	0	0	24
09.1	Animal and mixed food waste	1.183	1.183	13
09.2	Vegetal wastes	2.304	2.304	20
09.3	Animal faeces, urine and manure	0	0	9
10.1	Household and similar wastes	17.013	17.013	18
10.2	Mixed and undifferentiated materials	262.226	262.226	8.5
10.3	Sorting residues	279.583	279.583	2.5
11A	Common sludges (<i>excl. dredging spoils</i>): <i>Dry matter</i>	26.228	26.228	28
Total, wet weight		812.514	812.514	

* Waste statistics for 2010 are from Sweden's reporting to the Commission in accordance to the Waste Statistic Regulation. Waste statistics for 2011 is extrapolated values.

8.2.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Because of the simplifications in the used top-down model and the difficulties in estimating many of the parameters, the estimated emissions in the waste sector are uncertain. The time dependency in methane production makes the model estimate further dependent on assumptions of waste management from

²⁴⁸ Sundqvist & Szudy, 2012

earlier years. The uncertainty is highest in 1990 and then decreases, mainly due to better and more frequent activity data on household waste during the 1990s. The section of the Ordinance prohibiting deposition of organic waste as landfill was implemented on January 1st 2005. It has led to higher uncertainties since the data on DOC has not been updated during the last years to cover the changes. The DOC from the year 2005 is probably overestimated.

IPCC Guidelines suggest that the error in estimated methane generation potential may be about 15 % given high quality data and 50 % given poor data on methane generation, per ton of waste. The uncertainty in statistics on deposited waste may be 10 %, if the waste is weighted, or more than 200 % if the data quality is poor. The errors in estimated methane recovery will probably be small, according to the Good Practice Guidance. Given these standard uncertainty ranges and applying the simple error propagation formula, a total error of estimated methane emissions of about 20 % would be achievable, in the best case, given high quality data.

According to Good Practice Guidance there is some extra uncertainty in the methane generation rate constant [-40 %, 300 %], and in the oxidation factor, if oxidation is assumed. An assessment of the confidence interval for the Swedish methane estimate from landfills would be around 50-60 % for 2005. Swedish waste statistics 2005 on household waste, in particular, are of high quality, but the estimates are still dependent on lower quality data and extrapolations from earlier years. Furthermore, statistics on different waste fractions in household waste, and especially industrial waste, are still of lower quality. The quality of parameters based on IPCC default values may also be low, since they rely on older research, and data from Swedish on-site measurements is not yet extensive enough for verification.

The time series in the waste sector are calculated consistently and in line with the Good Practice Guidance. When statistics are not produced annually, interpolation and extrapolation have been necessary tools for imputation.

In submission 2013 the uncertainties were recalculated since errors in the data sheets were discovered. This affected mainly the calculated uncertainties for the base year 1990.

8.2.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

8.2.1.4.1 *Quality Assurance and Quality Control*

All quality procedures according to the Swedish QA/QC plan (Manual for SMED's Quality System in the Air Emission Inventories) have been implemented during the work with this submission.

8.2.1.4.2 *Verification of data and reducing compiling errors*

Statistics Sweden and the IVL has on behalf of the Swedish EPA scrutinized the activity data (quantities of deposited; household waste, park and garden waste, sludge from waste water treatment) used for calculations. The accuracy in these activity data is judged to be good.

8.2.1.5 SOURCE-SPECIFIC RECALCULATIONS

Recalculations have been made for CH₄ from 6.A.1 Managed waste disposal on land, mainly due to availability of new data on landfilled waste for the year 2010 and the changes in the EWC-stat codes implemented in the 2010 year data. These changes have an impact on emission years 2009-2010. Also the data for EWC-stat 10.3 (Sorting residues) has been changed for 2006 and 2008 due to typing error. These changes have an impact on emissions years 2006-2009. The recalculations have led to a slight increase of emissions of CH₄ (0 % - 0.35 %) from category 6.A.1 Managed waste disposal on land in submission 2013.

8.2.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

8.3 Waste water handling (CRF 6.B)

Sweden is reporting data on emissions of nitrous oxide (N₂O) and methane (CH₄) for the sectors Industrial Wastewater, CRF 6.B.1 and Domestic and Commercial Wastewater, CRF 6.B.2.

8.3.1 Industrial, domestic and commercial wastewater (CRF 6.B.1 and CRF 6.B.2)

8.3.1.1 SOURCE CATEGORY DESCRIPTION

In Sweden, internal wastewater treatment is practised in some industries (see further in section 8.3.1.2. Industries with internal wastewater treatment). These plants are situated both by the coast and in the inland.

There are almost 500 municipal wastewater treatment plants in Sweden with treatment capacity for more than 2,000 personal equivalents. 95 % of the wastewater is treated mechanically, chemically and biologically. In some larger plants, or plants with sensitive recipients, special nitrogen treatment is performed. These wastewater treatment plants also receive wastewater from industries without internal wastewater treatment.

In addition, there are also a number of smaller plants or private plants of varying standard.²⁴⁹ There are also approximately 1.3 million people in Sweden not connected to any wastewater treatment plant.

Considerable quantities of heat and bioenergy are recovered from sewage and wastewater.²⁵⁰ Anaerobic wastewater treatment and anaerobic digestion of sludge is practised in Sweden and generates methane for production of electricity, heating, vehicle fuel and for local gas distribution networks. Some of the methane is flared.

²⁴⁹ Swedish EPA & SMED, 2003

²⁵⁰ Ministry of the Environment, 2001.

The estimations of emissions (leakage) of nitrous oxide and methane from the wastewater treatment processes and sludge treatment processes need further improvements. However, some improvements have been made in submission 2011 by calculating and reporting emissions of CH₄ from wastewater treatment (Industrial Wastewater, CRF 6B1a, and Domestic and Commercial Wastewater, CRF 6B2a). Also emissions of CH₄ from sludge treatment (Domestic and Commercial Wastewater, CRF 6B2b) are calculated and reported.

According to a survey²⁵¹ by the Swedish Energy Agency on biogas production and utilization, the production of biogas in Sweden in 2010 was 1,387 GWh (or 99.5 Gg in methane) to be compared with 1,285 GWh (or 92.2 Gg in methane) in 2005²⁵². In 2010, 44 % of the produced energy from biogas was produced at wastewater treatment plants (anaerobic digestion of sludge). The biogas production at wastewater treatment plants increased by 9.8 % from 2005 to 2010. Approximately 9.4 % of the biogas produced at wastewater treatment plants was flared in 2010.

The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), is presented in Table 8-18.

Table 8-18. Summary of source category description, CRF 6.B.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
6.B	CO ₂				NA	NA	NA
	CH ₄	X	X		CS, D, T1	CS, D	Yes
	N ₂ O				CS	D	Yes

CS Country Specific. T1 Tier 1. D Default.

8.3.1.2 METHODOLOGICAL ISSUES

8.3.1.2.1 Nitrous oxide (N₂O)

National activity data on nitrogen in discharged wastewater from municipal wastewater treatment plants and industries are used, in combination with a model estimating nitrogen in human sewage from people not connected to municipal wastewater treatment plants.

The general formula to calculate the emissions is:

$$(N_{Industry} + N_{WastewaterTreatmentPlants} + PROTEIN * Nr_{People} * 0.16) * EF * 44 / 28$$

where N_{WastewaterTreatmentPlants} and N_{Industry} are the nitrogen in discharged wastewater from municipal wastewater treatment plants (including industries without internal wastewater treatment) and other industries (with internal wastewater treatment) respectively. IPCC's default emission factor, 1 % N₂O-N/N, is used

²⁵¹ Swedish Energy Agency, 2011

²⁵² Swedish Energy Agency, 2007

as emission factor (EF) for the discharges from all three sources (Wastewater treatment plants, Industries and unconnected households).

IPCC Guidelines suggest an emission factor of 1 % ($\text{N}_2\text{O-N/kg}$ sewage N discharged sewage effluent) for sewage nitrogen that enters rivers and estuaries (Good Practice Guidance, Table 4.18), while the N_2O emissions associated with sewage treatment and land disposal are considered to be negligible. There is no Swedish research that will motivate a national emission factor concerning discharged sewage nitrogen.

Wastewater treatment statistics and activity data related to nitrous oxide emissions in Sweden

According to Swedish environmental protection law, all municipal wastewater treatment plants designed for more than 2,000 person equivalents, including industry, need to report their discharges in legal environmental reports delivered to their supervision agency. Statistics are published every other year by the Swedish EPA.²⁵³

For industrial wastewater handling, Sweden has better data on discharges in tonnes than in cubic metres and therefore has chosen to publish data of discharges of nitrogen in tonnes. Discharges in tonnes are more relevant than in cubic meters because large quantities of the waste water output reported in the environmental reports are actually cooling water and not process water. It is less confusing for the reporting companies to report the quantity/quantities of their discharges of various substances in *tonnes* instead of in *cubic metres*.

The statistics on discharges of nitrogen exclude municipal wastewater treatment plants designed for less than 2,000 person equivalents. These were surveyed in 1999, and were found to represent about 6 % of the total discharged nitrogen, which is compensated for using a “1.1 factor” in the above formula.

The statistics also exclude approximately 1.3 million people in rural areas, who are not connected to municipal wastewater treatment. Until submission 2009, Sweden estimated this population to almost 1 million people. The new estimate is based on new data²⁵⁴ for 1995, 2000 and 2005. The mean of these data is approx. 1,264,000 people, which is rounded up to 1.3 million people in the calculations to compensate for suspected underestimation for year 1995. Sweden cannot see any national trend or variations over the years that are significant enough to apply in the calculations, since the data for 1995 (1,205,686 people) is likely an underestimation, and 2000 (1,296,757 people) and 2005 (1,291,299 people) are very similar. However, the nitrogen from these people is accounted for in the formula as well, through the model estimate of nitrogen production.

National values for protein consumption are used and are available in time series. Previously a constant value was used.

²⁵³ Statistics Sweden, MI 22 SM, Swedish EPA and SMED.

²⁵⁴ Statistics Sweden MI 11 SM 0701, Korrigerad version

Industries with internal wastewater treatment

The formula is: $N_{Industry} * EF * 44 / 28$

The sector covers; pulp and paper industry, oil refineries, chemical industry, iron and steel industry, food manufacturing industry, manufacturing of wood products and mining and quarrying industry.

Table 8-19 Discharges of nitrogen from mining and quarrying and manufacturing industries: Pulp and paper industry (total), Oil refineries (total), Chemical industry (inland and coastal), Iron and steel industry (inland and coastal), Food manufacturing industry (inland and coastal), Manufacturing of wood products (inland and coastal) and Mining and quarrying (total), tonnes.

Year	Pulp and paper (tot.)	Oil ref. (tot.)	Chemical (inl.)	Chemical (coast.)	Iron and steel (inl.)	Iron and steel (coast.)	Food (inl.)	Food (coast.)	Wood prod. (inl.)	Wood prod. (coast.)	Mining (tot.)
1990	5,500
1992	3,630
1994	3,200
1995	3,844	80	..	385	..	70	..	0
1997	3,433
1998	3,307	78	..	423	..	230	..	1
1999	3,042
2000	3,241	38	..	361	..	114	..	109
2001	3,014
2002	3,169	68	..	268	..	72	..	3
2003	3,162
2004	3,039	30	..	224	..	54	..	11	2	6	451
2005	3,222
2006	3,200	39	..	144	..	74	..	17	2	3	496
2007	2,825
2008	2,830	26	256	139	807	68	89	27	2	2	480
2009	2,600
2010	2,590	45	205	140	769	84	96	25	0	4	321
2011	2,500

Source: NV 4657, NV 4434, NV 4657, NV 4924, NV 4987, NV 5114, Swedish Forest Industries Federation, .MI 22 SM, Swedish EPA and SMED

Municipal wastewater treatment plants

The formula is: $N_{WastewaterTreatmentPlants} * EF * 44 / 28$

$N_{WastewaterTreatmentPlants}$ is magnified by 10 %, in order to compensate for wastewater from small treatment plants, not included in the statistics.

Table 8-20 Discharges of nitrogen from large municipal wastewater treatment plants (from treatment of domestic, commercial and industrial waste water), tonnes.

Year	Municipal wastewater treatment plants
1990	26,200
1992	25,310
1995	25,940
1998	21,376
2000	18,977
2002	18,036
2004	17,779
2006	18,347
2008	18,433
2010	17,419

Source: MI 22 SM, Swedish EPA and SMED

Households not connected to municipal wastewater treatment plants

The formula is: $(PROTEIN * Nr_{People} * 0.16) * EF * 44 / 28$

PROTEIN is the annual per capita consumption per person/year, Nr_{People} is the number of people not connected to municipal wastewater treatment plants, and 0.16 is the fraction of nitrogen in proteins (Table 8-21).

Table8-21. Protein consumption in Sweden, g/person/day

Year	Protein consumption g/person/day
1980	87
1985	86
1990	89
1995	89
2000	97
2003	101
2004	102
2005	102
2008	111
2009	110
2010	112

Source: The Swedish yearbook of agricultural statistics 2007, 2011 & 2012.

8.3.1.2.2 Methane (CH₄)

Methane emissions from wastewater treatment

CRF 6B1a, Industrial wastewater, Wastewater

The majority of the facilities in Sweden are using aerobic processes, where no CH₄ is supposed to be generated because of the use of aeration in the wastewater treatment process. In 2010, there were only five (5) facilities using anaerobic wastewater treatment processes in Sweden. These facilities were in the pulp industry and food industry.

For methane emissions from industries with internal wastewater treatment, Sweden has chosen a national method to estimate the emissions based on data availability. According to wastewater treatment expertise²⁵⁵, the loss of CH₄ in the energy recovery process should be within the range of 2 - 5 %. This factor can be combined with data on energy recovery from the anaerobic processes.

In 2010, 114 GWh²⁵⁶ (or 8.2 Gg CH₄) was recovered. By using the upper value (5 %) of the leakage factor to ensure a conservative estimate, the emission of CH₄ is calculated to 0.43 Gg for 2010.

No plant-specific data has been used in the calculations for any year. At the moment only statistical data sources are available and used. Statistical data on energy recovery from anaerobic processes are available for 2005-2010. For 1990-2004 no statistical data are available. Therefore the activity data for 2005 was extrapolated for 1990-2004. In other words, the value for 2005 was also used for the years 1990-2004. There is no additional statistical information available on upgrades or modifications of the recovery system in general. The estimate of the loss of CH₄ in the energy recovery process (5 %) is used for all years.

CRF 6B2a, Domestic and Commercial Wastewater, Wastewater

When analyzing the sector CRF 6B2a in Sweden it is necessary to divide the sector into three sections:

- a) Large wastewater treatment plant (treatment capacity: more than 2,000 pe)
- b) Small wastewater treatment plants (treatment capacity: 25 - 2,000 pe)
- c) Population not connected to wastewater discharge system
 - a) In Sweden, all large wastewater treatment plants are using aerobic wastewater treatment processes. No CH₄ is supposed to be generated because of the use of aeration in the wastewater treatment process.
 - b) For small wastewater treatment plants, the situation is at the moment not well enough investigated and therefore Sweden is using the IPCC Good Practice Guidance method (Page 5.15 Box 5.1 Check method):

²⁵⁵ Ek, 2010

²⁵⁶ Swedish Energy Agency, 2011

$$WM = P \times D \times SBF \times EF \times FTA \times 365 \times 10^{-12}$$

where

WM = Annual CH₄ emission per country, from domestic wastewater (Tg)

P = Population of country or urban population for some developing countries (person)

D = Organic load in biochemical oxygen demand per person (g BOD/person/day),
overall default = 60 g BOD/person/day

SBF = Fraction of BOD that readily settles, default = 0.5

EF = Emission factor (g CH₄/g BOD), default = 0.6

FTA = Fraction of BOD in sludge that degrades anaerobically,
default = 0.8

Activity data on population connected to small wastewater treatment plants (700,000 people) is derived of background data from a survey from 2010 on treatment methods and sewage networks in Swedish municipal waste water treatment plants. A report²⁵⁷ was published in 2011.

c) For population not connected to wastewater discharge system, the following applies:

- 1) The sludge in the wastewater (SBF = Fraction of BOD that readily settles) is collected and transported to anaerobic digestion plants located at larger wastewater treatment plants²⁵⁸. It is covered and reported in section CRF 6B2b (sludge treatment).
- 2) CH₄ emissions from the remaining wastewater are likely to be NO (not occurring) or negligible. The waste water is rich in oxygen, and for biological processes to occur the water must not be too cold.²⁵⁹ Sweden has a rather cold climate with an average annual temperature of 4.8 (°C) 1991-2005. At the moment Sweden does not²⁶⁰ have any written references to support this statement and therefore emissions are estimated by using the default Check method (as for 'b') above).

The activity data (population not connected to wastewater discharge system) is the estimate based on data²⁶¹ for 1995, 2000 and 2005 (1,300,000 people).

²⁵⁷ Brånvall & Svanström, 2011

²⁵⁸ Ek, 2010

²⁵⁹ Ibid

²⁶⁰ Ek & Szudy, 2011

²⁶¹ Statistics Sweden MI 11 SM 0701, Korrigerad version

Methane emissions from sludge treatment

CRF 6.B.1.b, Industrial wastewater, Sludge

The European Union Waste Statistics Regulations (EC 2150/2002), also called WStatR, came into force in November 2002. It requires all Member States to provide data to the European Commission every two years on the generation and treatment of waste and on the number and capacities of waste management facilities. In this context, waste treatment covers incineration, recovery and disposal. Sweden has so far, as a Member State of the European Union, reported official national statistics on waste to the Commission for the years 2004, 2006, 2008 and 2010.

The inventory work and the data collection for WStatR have provided Sweden's national waste experts with valuable information on existing waste management practices in Sweden. Since the WStatR by definition shall cover 100 % of the generated waste in all sectors and incineration, recovery and disposal of waste, it is considered a reliable data source.

Both environmental reports and questionnaires have been used as data sources. All companies which perform waste treatment activities need permits and also to compile an annual environmental report. These environmental reports have been read by a team of waste experts for the purpose of finding waste data for WStatR.

The conclusions²⁶² on the subject are that no activities such as anaerobic digestion of sludge from industrial wastewater treatment have been found during the inventory. Therefore Sweden is reporting "NO" (not occurring) for *emissions* and *recovery* for 6B1: *sludge treatment*.

CRF 6.B.2.b, Domestic and commercial wastewater, Sludge

A study²⁶³ conducted by SWECO (Swedish Consultants) on behalf of Stockholm Water Authority (Stockholm Vatten) on two wastewater treatment plants in Stockholm shows that this leakage is between 4 % and 7 % of the gas production for these plants. If these plants are representative for all 135 wastewater treatment plants that apply anaerobic digestion in Sweden year 2010, then a national methane emission estimates can be calculated by combining this factor with data on energy recovery from the anaerobic sludge treatment.

In 2010, 614 GWh²⁶⁴ (or 44.0 Gg CH₄) was recovered. By using the upper value (7 %) of the leakage factor to ensure a conservative estimate the emission of CH₄ is calculated to 3.32 Gg for 2010.

Data on energy recovery from anaerobic sludge treatment are available for 2005-2010. For 1990-2004 no data are available and therefore the data for 2005 was extrapolated for these years. The estimate of the loss of CH₄ in the energy recovery process (7 %) is used for all years.

²⁶² Memo "Occurrence of treatment of sludge by anaerobic digestion in Swedish industries", Statistics Sweden, 2011 "

²⁶³ " Metanförluster vid avloppsreningsverken i Henriksdal och Bromma", Stockholm Vatten, 2004

²⁶⁴ Swedish Energy Agency, 2011

8.3.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Because of the simplifications in the used top-down model and the difficulties in estimating many of the parameters, the estimated emissions in the waste sector are uncertain.

The statistics of discharges from municipal wastewater treatment plants are biased from sources of inaccuracy such as under coverage, non-response or no observations and sample errors “within” the treatment plants. No objective methods of calculating accuracy measures have been developed, but data on nitrogen is considered to have a margin of inaccuracy of well under 10 % at national level. The inaccuracy in the emission factor is estimated to be at least 50 %, according to Good Practice Guidance. This results in an overall inaccuracy exceeding 50 % annually, and more for years where activity data have been extrapolated.

For methane emissions, both national methods and default methods are used. For the activity data “energy recovery” for the early years with no observations, a 50 % uncertainty is assumed, while for the recent years with actual observations a 10 % uncertainty is assumed. The uncertainty of the emission factor “leakage” is assumed to be the same for all years (40 %). All of these estimates are expert judgements.

For the activity data “population connected to small wastewater treatment plants” and “population not connected to wastewater treatment plants” the uncertainty for the early years is judged to be higher (10 %, expert judgement) than for the recent years (5 %, IPCC default). The uncertainty of the emission factor (Equation 5.6 in Good Practice Guidance) “Check method” is based on expert judgements and assumed to be high (65 %).

The time series in the waste sector are calculated consistently and in line with the Good Practice Guidance. When statistics are not produced annually, interpolation and extrapolation have been necessary tools for imputation.

In submission 2013 the uncertainties were recalculated since errors in the data sheets were discovered. This affected mainly the calculated uncertainties for the base year 1990.

8.3.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All quality procedures according to the Swedish QA/QC plan (Manual for SMED's Quality System in the Air Emission Inventories) have been implemented during the work with this submission.

8.3.1.5 SOURCE-SPECIFIC RECALCULATIONS

Recalculations have been made for N₂O from 6.B.1 Industrial Wastewater and 6.B.2 Domestic and Commercial Wastewater. New data has been available on discharges from wastewater treatment plants as well as protein consumption for year 2010, which in turn affects the interpolated data for 2009. The recalculations have led to a decrease of emissions of N₂O from category 6.B Waste water handling. The reduction amounts to -1.93 % for 2009 and -3.44 % for 2010.

8.3.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

8.4 Waste incineration (CRF 6.C)

8.4.1 Source category description

Emissions from incineration of hazardous waste, and in later years also MSW and industrial waste, from one large plant are reported in CRF 6.C. Emissions from other MSW incineration plants combusting waste for energy purposes are included in CRF 1. The summary of the latest key category assessment, methods and EF used, and information on completeness, i.e. if any sources are not estimated (NE), are presented in Table 8-22.

Table 8-22. Summary of source category description, CRF 6.C.

CRF	Gas	Key Category Assessment 2011			Method	EF	All sources estimated
		Level	Trend	Qualitative			
6.C	CO ₂				T3	PS	Yes
	CH ₄				T2	PS	Yes
	N ₂ O				T2	PS	Yes

PS Plant Specific. T2 Tier 2. T3 Tier 3.

8.4.2 Methodological issues

For the whole waste category, the methodology and time series consistency are in line with the Good Practice Guidance.

Emissions from incineration of hazardous waste, and in later years also MSW and industrial waste, from one large plant are reported in CRF 6.C. Reported emissions are for the whole time series obtained from the facility's Environmental report or directly from the facility on request. CO₂, SO₂ and NO_x are measured continuously in the fumes at the plant. In 2003 capacity was increased substantially at the plant by taking one new incinerator into operation. The new incinerator incinerates a mixture of MSW, industrial waste and hazardous waste. As a consequence of increased capacity, the emissions from 2003 are increased compared to earlier years. Only a minor part (1 – 2%) of the total amount of waste incinerated for energy purposes in Sweden are incinerated in the facility included in 6.C. All other emissions from incineration of MSW are reported in CRF 1.

Emissions reported are CO₂, CH₄, N₂O, NO_x, SO₂ and NMVOC.

In previous submissions, only the amount of CO₂ emissions of biogenic origin of the MSW fraction of the waste, have been estimated using information

from a report published by the Swedish Waste Management²⁶⁵. In this report the information is given that approximately 70 % of the emitted CO₂ from incineration of MSW have biogenic origin. All other incinerated waste fractions have been assumed to be of fossil origin. The amounts of incinerated MSW have been obtained by the company for the years 2003 until 2009. For 2010 this information was considered to be market-related, not official, information. Reported MSW for 2010 was in submission 2012 estimated applying the same proportion of MSW of the total amount of incinerated waste as in 2009.

In current submission the time series of reported biogenic and fossil CO₂ are revised. For the years 2008, 2009, 2010 and 2011 the company has, besides total emissions of CO₂, also reported CO₂ with respect to biogenic or fossil origin. The company has based their reporting of biogenic and fossil CO₂ on a detailed study performed in 2008. In this study they found that 63% of the totally emitted CO₂ had biogenic origin. This finding is in good agreement with newly published results from a Swedish study showing that about one third of the carbon in solid waste is of fossil origin²⁶⁶. As the mixture of incinerated wastes has been almost the same for all years from 2003, when the new incinerator was taken into operation, the company considered this biogenic percentage of the totally emitted CO₂ to be valid also for the years 2003 to 2008. For the period before 2003 the company considers reported CO₂ emissions to be almost 100% fossil.²⁶⁷

Before 2008 occasional measurements of CH₄ in the flue gas was performed. The company reported CH₄ emission around 1.1 Mg for 2008. This information, together with information of incinerated amounts of waste 1990 until 2007, has been used for estimating a time series 1990 – 2008 for emissions of CH₄ in CRF 6C. For 2008, 2009, 2010 and 2011 reported CH₄ emissions are based on continuous measurements in the flue gas. Also N₂O from waste incineration is reported for the whole time series. The estimates are based on occasional measurements of the N₂O concentrations in the flue gas together with information on yearly flue gas volumes 2003 - 2011. For 1990 until 2002 the volumes are not known and for these years the flue gas volumes have been estimated using the average of the ratios between volumes and incinerated amounts of waste for 2003 to 2008. Activity data and emission factors used for the CH₄ and N₂O estimates are presented in Table 8-23.

²⁶⁵ Swedish Waste Management. RVF rapport 2003:12 (in Swedish). Förbränning av avfall. Utsläpp av växthusgaser jämfört med annan avfallsbehandling och annan energiproduktion.

²⁶⁶ Swedish Waste Management. RAPPORT U2012:05. Determination of the fossil carbon content in combustible municipal solid waste in Sweden.

²⁶⁷ Personal communication, Hanna Eriksen, Hanna.Eriksen@sakab.se, 2012-08-23

Table 8-23. Activity data and emission factors used for estimations of CH₄ and N₂O emissions in CRF 6.C.

Year	Total amounts of incinerated waste	Flue gas volume	N ₂ O	CH ₄
	Gg	1,000 m ³	EF, g/1,000 m ³	IEF, kg/Gg
1990	30	220,674	15.00	7.73**
1995	33	240,637	15.00	7.73**
2000	28	205,778	15.00	7.73**
2001	31	228,934	15.00	7.73**
2002	33	240,887	15.00	7.73**
2003	122	789,438	15.00	7.73**
2004	130	881,100	15.00	7.73**
2005	126	1,099 338	15.00	7.73**
2006	122	902,039	15.00	7.73**
2007	140	915,032	15.00	7.73**
2008	146	1,189 691	15.00	7.73
2009	162	1,107 410	15.00	5.32
2010	115	1,007 061	15.00	7.84
2011	163	1,229 605	15.00	6.09

* = estimated volume

** = IEF for 2008 used for estimations 1990 - 2007

8.4.3 Uncertainties and time-series consistency

In Revised 1996 IPCC Guidelines no information concerning uncertainties for CO₂, CH₄ and N₂O can be found. In 2006 IPCC Guidelines is stated that if a default value for emission factor is used the uncertainty has been estimated to be ± 100 per cent or more and the uncertainty for plant specific activity data is ± 5 %. In this case the activity data referred to is amount of waste incinerated. The Swedish reporting of N₂O is based on an emission factor and measured yearly amounts of flue gas and the uncertainty for emission factor is set to ± 100 % and the uncertainty for activity data is set to 5 %.

In 2006 IPCC Guidelines it is not easy to find information concerning uncertainties for measured amounts of emitted CO₂ but corresponding information for measured amounts of CH₄ is likely to be in order of ± 10 %. Due to lack of other information the emissions data uncertainty for CO₂ and CH₄ are set to ± 10 %.

As can be seen in Table 8-23 the implied emission factor (IEF) for CH₄ varies slightly for later years, especially so between 2008 and 2009. Reported emissions for 2008 – 2011 are based on continuous measurements and the reason for the variation between 2008 and 2009 may be explained by variations in the composition of the incinerated waste.

8.4.4 Source-specific QA/QC and verification

No source-specific QA/QC or verification is performed.

8.4.5 Source-specific recalculations

In the previous submission, CH₄ emissions for the years before 2008 were calculated using a wrong IEF for 2008. In this submission the emissions have been correctly estimated using the accurate IEF for 2008. This recalculation of reported CH₄ emissions 1990 - 2007 led to minor changes in reported CH₄ (between a reduction of 70 kg to an increase of 214 kg).

Total CO₂ emissions for 2006 were corrected. In current submission the total CO₂ emission for 2006 is 23 Gg higher compared to the previous submission. Also the allocation of CO₂ with respect to fossil or biogenic origin has been changed for the period 2003 to 2010.

8.4.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.

9 Other

Not applicable for Sweden.

10 Recalculations and improvements

Since the last submission, recalculations of GHG emissions for several years have been carried out throughout the inventory. The recalculations are due to comments and implemented recommendations from the national and international review teams in the on-going progress to make the inventory be fully in line with the IPCC Guidelines and the Good Practice Guidance. The recalculations include new methods, emission factors, thermal values and activity data. Some recalculations are due to errors in earlier inventories detected during the work with the present inventory.

10.1 Explanations and justifications for recalculations

The explanations and justifications for the recalculations made in this submission since submission 2012, together with descriptions on their implications for the emission levels, are given in the sector specific chapters.

10.2 Implications for emission levels

This section provides a general description for each sector of the major recalculations made. The implications for emission levels of GHG emissions by sector are presented in Table 10-1. In section 10.3 the implications for emission trends are presented.

10.2.1 Energy, CRF 1

10.2.1.1 STATIONARY COMBUSTION

Combustion of coke for production of heat used in carbide manufacturing has been included in CRF 1A2c in submission 2013. Emissions are reported for every year 1990-2011. In earlier submissions, activity data for this combustion was collected from energy statistics, but in submission 2013 it was discovered that this data was not complete. Because of this, activity data from EU ETS is used for this particular activity.

Activity data for stationary combustion of all fuels used in the other sector, e.g. CRF 1A4 and parts of CRF 1A2f (small industrial enterprises, the construction sector) 2009-2010 has been revised due to revisions of the annual energy balances.

Minor errors in the emission factor for CO₂ from natural gas in 2007-2010 have been corrected.

10.2.1.2 MOBILE COMBUSTION

The ratio of aviation gasoline for national aviation was underestimated in submission 2012 and the ratio for aviation kerosene was as a result overestimated. This has been adjusted in submission 2013.

The database used by the Swedish Defence Research Agency (FOI), in their calculations of fuel consumption and emissions from aviation, has been updated for the years 2008-2011.

The road transport emission model HBEFA 3.1 is updated yearly with new information, but in submission 2013 there has been an extended review of traffic work and the driving distances were updated for all vehicle types all the way back to 1999. The HBEFA model has also been updated with information regarding the use of air conditioning in passenger cars for all years and data regarding the number of passenger cars equipped with particle filters has also been updated.

In submission 2012 the emissions of N_2O and CH_4 from ethanol used by road traffic were overestimated and the emissions of CH_4 and N_2O from low blended gasoline in E85 vehicles were not included. This is corrected in submission 2013.

As from submission 2013, CH_4 and N_2O emissions from HBEFA are no longer adjusted to be in line with fuel delivery statistics, as these emissions are based on other factors such as mileage, type of vehicle, speed limits etc. These factors are considered in the road traffic model HBEFA.

The consumption of ethanol for road traffic by the military has been added as from 2007.

In submission 2013 activity data on residual fuel oil used for domestic navigation, was interpolated for 2009 and 2010 based on data for 2008 and 2011 due to discovered errors in activity data.

10.2.1.3 FUGITIVE EMISSIONS

CH_4 and CO_2 emissions from venting activities from natural gas transmission pipelines have been estimated for the first time in submission 2013 and are reported in CRF 1.B.2.B.3.

CH_4 emissions from storage of natural gas have been estimated for the first time in submission 2013 and are reported in CRF 1.B.2.B.3.

10.2.2 Industrial processes, CRF 2

In the previous submission, CO_2 emissions from organic carbon in the raw material were double-counted for the years 2005 – 2010. (For these years CO_2 emissions from organic carbon are included in emissions reported in EU ETS.) In submission 2013 this has been corrected.

10.2.3 Solvents and other products use, CRF 3

By mistake a substance with a vapour pressure of less than 0.01 kPa has been included in activity data derived from the Product Register. In the 2013 submission, this has been corrected. This correction affects mainly emissions and activity data for printing industry included in CRF.3.D.5 2006-2010.

10.2.4 Agriculture, CRF 4

Due to comments received from the EU ESD review, the emissions from calves are recalculated, now assuming zero emissions from calves feeding on only milk. This affected the emission estimate for CH₄ from 4.A.

In 4.B a small error in the manure production rate for grazing beef cows was corrected which affected the estimate for the CH₄ emissions.

A new data for sludge use 2010 have been published by Statistics Sweden. This data was published too late to be included in the 2012 submission. This affected the emission estimate for 4.D.1.6 Use of sewage sludge as fertilizers.

In submission 2012 data for nitrogen lost as ammonia was subtracted before the EF for N₂O emissions was applied. However, following comments from the UNFCCC centralized review and a “Saturday Paper” on this issue, data for submission 2012 was resubmitted without the subtraction of ammonia. This affected the estimate of N₂O from 4.D.2. However, compared to the 2012 resubmission there are no recalculations for this source.

There are some minor differences in the estimate of total area of agricultural land through the whole time series. The reason is the method used by SLU for estimating agricultural (see 7.2.2.1). This lead to difference in the emission estimates from the following CRF-codes, 4.D.1.5, 4.D.3.2 and 4.D.4\ Cultivation of mineral soils.

10.2.5 LULUCF, CRF 5

The major part of the differences in total removals 1990-2010 is due to updated AD from the NFI, mainly affecting carbon stock changes in soil carbon and litter. From 2007 and onwards the difference is also affected by the annual update of the reporting database, which results in recalculated data in Living biomass from 2007 to 2010. The difference is also due to the recalculation of individual sample series using extrapolation as described in section 7.2.2.2.-

10.2.6 Waste, CRF 6

Recalculations have been made for CH₄ from 6.A.1 Managed waste disposal on land, mainly due to availability of new data on landfilled waste for the year 2010 and the changes in the EWC-stat codes implemented in the 2010 year data. These changes have an impact on emission years 2009-2010. Also the data for EWC-stat 10.3 (Sorting residues) has been changed for 2006 and 2008 due to typing errors. These changes have an impact on emissions years 2006-2009. The recalculations have led to a slight increase of emissions of CH₄ (0 % - 0.35 %) from category 6.A.1 Managed waste disposal on land in submission 2013.

Recalculations have been made for N₂O from 6.B.1 Industrial Wastewater and 6.B.2 Domestic and Commercial Wastewater. New data has been available on discharges from wastewater treatment plants as well as protein consumption for year 2010, which in turn affects the interpolated data for 2009. The recalculations have led to a decrease of emissions of N₂O from category 6.B Waste water handling. The reduction amounts to -1.93 % for 2009 and -3.44 % for 2010.

In the previous submission, CH₄ emissions from 6.C for the years before 2008 were calculated using a wrong IEF for 2008. In submission 2013 the emissions have been correctly estimated using the accurate IEF for 2008. This recalculation of reported CH₄ emissions 1990 - 2007 led to minor changes in reported CH₄ (between a reduction of 70 kg to an increase of 214 kg).

Total CO₂ emissions for 2006 were corrected for 6.C. In current submission the total CO₂ emission for 2006 is 23 Gg higher compared to previous submission. Also the allocation of CO₂ with respect to fossil or biogenic origin has been changed for the period 2003 to 2010.

Table 10-1. Recalculations of GHG emissions between submission 2013 and submission 2012 by CRF sector.

Recalculation difference														
Year	Total (excl LULUCF)	CRF 1		CRF 2		CRF 3		CRF 4		CRF 5		CRF 6		
	Gg CO ₂ eq.	%	Gg CO ₂ eq.	%	Gg CO ₂ eq.	%	Gg CO ₂ eq.	%	Gg CO ₂ eq.	%	Gg CO ₂ eq.	%		
1990	-55	-0.1%	63	0.1%	0	0.0%	0	0.0%	-118	-1.3%	4,075	-9.9%	0	0.0%
1991	-60	-0.1%	64	0.1%	0	0.0%	0	0.0%	-124	-1.4%	3,343	-8.1%	0	0.0%
1992	-85	-0.1%	45	0.1%	0	0.0%	0	0.0%	-130	-1.5%	4,198	-10.5%	0	0.0%
1993	-124	-0.2%	16	0.0%	0	0.0%	0	0.0%	-140	-1.6%	3,842	-10.7%	0	0.0%
1994	-124	-0.2%	20	0.0%	0	0.0%	0	0.0%	-144	-1.6%	3,911	-10.8%	0	0.0%
1995	-110	-0.1%	22	0.0%	0	0.0%	0	0.0%	-132	-1.5%	3,893	-11.0%	0	0.0%
1996	-113	-0.1%	20	0.0%	0	0.0%	0	0.0%	-134	-1.5%	3,887	-10.5%	0	0.0%
1997	-112	-0.2%	19	0.0%	0	0.0%	0	0.0%	-131	-1.5%	3,530	-9.1%	0	0.0%
1998	-98	-0.1%	25	0.0%	0	0.0%	0	0.0%	-123	-1.4%	3,677	-9.6%	0	0.0%
1999	-93	-0.1%	28	0.1%	0	0.0%	0	0.0%	-120	-1.4%	4,114	-10.8%	0	0.0%
2000	-94	-0.1%	27	0.1%	0	0.0%	0	0.0%	-121	-1.4%	3,370	-8.7%	0	0.0%
2001	-90	-0.1%	29	0.1%	0	0.0%	0	0.0%	-119	-1.4%	3,232	-8.3%	0	0.0%
2002	-95	-0.1%	26	0.0%	0	0.0%	0	0.0%	-121	-1.5%	3,288	-8.4%	0	0.0%
2003	-128	-0.2%	28	0.1%	0	0.0%	0	0.0%	-124	-1.5%	3,744	-10.3%	-32	-1.2%
2004	-128	-0.2%	33	0.1%	0	0.0%	0	0.0%	-124	-1.5%	3,398	-10.4%	-37	-1.4%
2005	-153	-0.2%	39	0.1%	-28	-0.4%	0	0.0%	-124	-1.5%	3,799	-12.3%	-39	-1.6%
2006	-146	-0.2%	28	0.1%	-30	-0.4%	-1	-0.2%	-122	-1.5%	2,516	-6.8%	-22	-0.9%
2007	-131	-0.2%	80	0.2%	-27	-0.4%	-15	-5.1%	-121	-1.5%	5,671	-15.4%	-48	-2.1%
2008	-230	-0.4%	9	0.0%	-28	-0.4%	-23	-7.5%	-126	-1.6%	3,952	-10.7%	-62	-3.0%
2009	-372	-0.6%	-132	-0.3%	-25	-0.5%	-41	-13.2%	-126	-1.6%	2,856	-8.0%	-48	-2.5%
2010	-784	-1.2%	-554	-1.1%	-30	-0.4%	-22	-7.1%	-127	-1.6%	3,355	-9.9%	-50	-2.7%

10.3 Implications for emission trends

The total emissions of GHG have changed for all inventory years due to the recalculations. Below a more detailed description is presented of implications for emission trends due to recalculations of the base year emissions and the last recalculated year's emissions. Note that this section does not include implications for emission trends in the LULUCF sector. In Table 10-2 it can be seen that compared to the estimated assigned amounts, the base year emissions in submission 2013 are about 713 Gg CO₂ equivalents higher.

Table 10-2. Difference between Assigned Amount and Base Year emissions submission 2013 by GHG, excluding LULUCF

GHG	Assigned Amount (Gg CO ₂ eq.)	Base Year* emissions Submission 2013 (Gg CO ₂ eq.)	Difference between Base Year emissions Submission 2013 and Assigned Amount (Gg CO ₂ eq.)
CO ₂	56,301.08	56,954.05	652.97
CH ₄	6,719.22	6,938.23	219.01
N ₂ O	8,534.73	8,369.65	-165.09
F-gases	596.61	602.23	5.62
Total	72,151.65	72,864.16	712.51

*1995 for F-gases and 1990 for other GHG emissions (excluding LULUCF)

Based on submission 2013, the estimated GHG emissions in Sweden decreased by 15.7% between the base year (72,864 Gg CO₂ equivalents) and 2011 (61,449 Gg CO₂ equivalents). In Table 10-3 it can be seen that in submission 2012 the trend from the base year to 2010 shows a 9.1% decrease. It can also be seen that the recalculation of GHG emissions in submission 2013 increased the downward trend between the base year and 2010 by 729 Gg CO₂ equivalents or 1.0 % points compared to submission 2012.

Table 10-3. Impact on emission trends (base year to 2010) due to recalculations of GHG emissions between submission 2013 and submission 2012 by GHG, excluding LULUCF

Trend Base Year* to 2010						
GHG	Submission 2012		Submission 2013		Difference between submission 2013 and submission 2012	
	Gg CO ₂ eq.	%	Gg CO ₂ eq.	%	Gg CO ₂ eq.	% points
CO ₂	-4,007	-7.0%	-4,652	-8.2%	-645	-1.1
CH ₄	-1,795	-25.5%	-1,862	-26.8%	-67	-1.4
N ₂ O	-1,324	-15.8%	-1,337	-16.0%	-13	-0.2
F-gases	479	79.5%	474	78.7%	-5	-0.8
Total	-6,647	-9.1%	-7,377	-10.1%	-729	-1.0

*1995 for F-gases and 1990 for other GHG emissions (excluding LULUCF)

10.4 Recalculations and other changes made in response to the UNFCCC review process

As the inventory time cycle in Sweden is planned for a national independent review of the inventory, submission 2013 is already compiled in mid-October 2012. The preliminary result of the centralized review in 2012, taking place in September, can thus only be taken into account as minor recalculations and changes in response to the review process. In Table 10-4 the recalculations and other changes in data and in the NIR made in response to the UNFCCC review process are described briefly and referenced to relevant sections to the NIR. Table 10-3 describes the Expert Review team recommendations for submission 2011 and earlier not yet implemented in the Swedish inventory and the reasons for that together with possible implementation plans.

Table 10-4. Recalculations and other changes made in response to the UNFCCC review process.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2006	General	12. In the NIR provided more methodological detail so that the relationship between activity data (AD), emission factors (EFs) and equivalent parameters and emission estimates was clear, and if the reasons for apparent outliers or anomalies in implied emission factors (IEFs) had been easier to understand. This would have reduced the number of questions and requests for background material during the review. The ERT recommends that the accessible style of the NIR be retained, but that more use is made of tabular and graphic material, and annexes to convey the methodological detail.	Several measures are in place to enable further review of data. E.g. IEFs are compared to IPCC defaults in CRF 2C1.2 and CRF 2F.
submission 2007/2008, (Submission 2010)	General	10, (24). The LULUCF sector requires improved descriptions of country circumstances and the approaches to estimating emissions and removals. One area where transparency could be improved is the use of EU ETS data in the national GHG inventory, in the description of the national system.	The description of the LULUCF sector has continuously been improved since submission 2008 with information concerning the national circumstances. Information on the use of EU ETS data in the Swedish GHG inventory has continuously been improved since submission 2008. See section 1 and 3 and 4 and Annex 2 (1.1.10), and Annex 8:1. Further information will be added as needed in future submissions.
Submission 2007/2008	General	83. Include information on the commitment period reserve.	Information is included in section 12.5.
Submission 2009, (Submission 2010)	General	8, (6, 13). Provide information in CRF table 7 as presented in Annex 1 to the NIR for the purpose of completeness.	Included in submission 2011.
Submission 2009, (Submission 2010)	General	19, (18). Extend the information on its national system to include the specific responsibilities of the organizations participating in SMED and consultants who assist the Swedish EPA in the inventory preparation.	Information on the national system is revised and updated in submission 2011 and in submission 2012, see section 1.2 and 1.3
Submission 2009, (Submission 2010)	General	20, (19). Total emission estimates used for the key category analysis were not the same as those reported in the NIR, CRF tables and the background tables (appendix 20B to the NIR). Perform the key category analysis correctly and report it in the next annual submission.	The key category analysis is subject to QA and QC activities before delivered to the UNFCCC but they were not fully applied in the resubmission of 2009 due to limited time.
Submission 2009	General	21. Extend its overall uncertainty analysis to include the LULUCF sector. Perform uncertainty analysis taking correlations between gases or categories into account in the next annual submission.	Uncertainty estimates for LULUCF are included in submission 2011. IPCC Tier 1 does not allow correction of correlations between activity data when used for estimating several GHG (e.g. CO ₂ , CH ₄ and N ₂ O based on fuel combustion). The activity data uncertainty is thus applied in several cells in the IPCC Table 6.1 calculations leading to underestimations of total emissions.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2009	General	23. Provide information on QA applied to data from EU ETS.	Information on the use of EU ETS data in the Swedish GHG inventory has continuously been improved since submission 2008. See section 1 and 3 and 4 and Annex 2 (1.1.10) and Annex 8:1. Further information will be added as needed in future submissions.
Submission 2009	General	24. Improve transparency by providing more precise and detailed explanations of methodologies, AD and EFs used as well as relevant category-specific QA/QC activities in cases Sweden uses AD from different sources for a single category, country-specific EFs, or methods that are not explicitly explained in the Revised 1996 IPCC Guidelines or the IPCC good practice guidance.	More information is included in submission 2011.
Submission 2009	General	28. (a) If recommendations identified during the previous review cannot be implemented, the Party should clearly explain the reasons;	Fulfilled through table 10.5
Submission 2009	General	28 (d) The implementation of QA/QC procedures needs to be improved to avoid calculation errors and inconsistency between the CRF tables and the NIR;	Sweden's internal handling plan for deliverables between contracting agency SMED and Swedish EPA have been revised in order to enable more time for internal QA/QC routines. Further efforts will be continuously made in future submissions in order to improve QA/QC procedures. Improvements will be reported in coming NIRs.
Submission 2009	General	88. In the SIAR enhance the availability of the required public information mentioned above and ensure that rejected transactions are terminated, and should report, in its next annual submission, on these two identified issues.	See complete response in Annex 6
Submission 2009	General	89. Include correct information on its commitment period reserve in its next annual submission.	Correct information on the commitment period reserve is included in section 12.5.
Submission 2009	General	90. Report any changes in its national system in accordance with section I.F of the annex to decision 15/CMP.1.	The information on the national system is revised and updated in submission 2011 and submission 2012, see section 1.2 and 1.3
Submission 2009	General	91. Report in its next annual submission any changes in its national registry in accordance with section I.G of the annex to decision 15/CMP.1.	Appropriate information is provided in submission 2011
Submission 2010	General	11, 14. Continue to include emission estimates from civil aviation, industrial wastewater and domestic and commercial wastewater raised by the ERT in the list of potential problems.	Addressed in submission 2012, see section 3.2.15 and section 8.3.1.
Submission 2010	General	21. Use the results of uncertainty analysis to prioritize improvements in the inventory for its next annual submission.	Since the 2011 submission, a Tier 2 key category analysis is performed, taking into account uncertainty estimates. The Tier 2 analysis is used when prioritizing resources and efforts for improving the inventory (see section 1.2.3) together with other input such as international reviews and national peer review.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2011	General	25. The ERT recommends that, in the NIR of its next annual submission, Sweden improve the explanation of which inventory improvements lead to recalculated inventory estimates and improved uncertainty estimates, and how these improved uncertainties are considered in the uncertainty analysis.	The Swedish inventory planning is described under 1.3.3 of the NIR. Information on improved uncertainties due to development projects is included under 1.7.1 of the NIR.
Submission 2011	General	26. There are several instances where the uncertainty of AD is indicated as "0" in table A.7.2. in annex 7 to the NIR. The ERT recommends that, in the NIR of its next annual submission, Sweden revise these uncertainty estimates.	In line with GPG (2000) uncertainties associated with measured emissions (e.g. for CH ₄ emissions from oil and natural gas) are allocated to either AD or EF depending on its likelihood of correlation over time. However, for clarity we now write "-" instead of "0" in table A.7.2 and A.7.3 in the NIR annex.
Submission 2006	Energy	23, 37. Categories not estimated	Fugitive emissions of CH ₄ from transport of crude oil are estimated in submission 2010 and added to CRF 1B2a iii Transport, whereas CO ₂ emissions in the same category is considered to be not estimated (NE) in Sweden (see NIR 3.3.2).
Submission 2007/2008	Energy, Industrial processes, Waste	19, 32, 80. Categories not estimated	1B2c Flaring: In submission 2010, all emissions are estimated. All plants where flaring occurs are part of the EU ETS system, and according to this data no flaring of natural gas occurs. If natural gas is included in the "burning gas" sometimes reported to the EU ETS, the emissions are estimated but reported as IE (in flaring of liquid fuels). 1B2c Flaring: In submission 2010, all emissions are estimated. All plants where flaring occurs are part of the EU ETS, and according to this data no flaring of natural gas occurs. If natural gas is included in the "burning gas" sometimes reported to the EU ETS, the emissions are estimated but reported as IE (in flaring of liquid fuels). CH ₄ from Carbon Black added in CRF 2B5 all years (see NIR 4.3.4)
Submission 2006	Energy	24. Institutionalize system-level checks to minimize the risk of missing plants or data in its future submissions.	The emissions from stationary combustion are calculated with activity data from a sample survey (thoroughly described in Annex 2). SMED internal QC procedures are in place to minimize error in compilation and data handling. The procedures will be continuously improved also in future submission.
Submission 2006	Energy	32 Methodology for Iron and steel industry	Addressed in submission 2010. Emissions have been revised for all years and reported in accordance with IPCC Guidelines (see NIR section 4.4.1)
Submission 2006	Energy	34. Improve explanations on fluctuating trend for fugitive emissions of refinery gas in Petroleum refining	Text in NIR section 3.3.2 clarified in submission 2010
Submission 2006	Energy	36. The allocation of fuel between civil aviation and aviation bunkers is not transparently described in the NIR, especially for the period 1990–1994.	A more detailed description is provided in NIR submission 2010 section 3.2.15

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
submission 2007/2008	Energy	27. The CO ₂ IEF for diesel decreased due to the change in the mix of the different types of diesel in Sweden. The ERT recommends that Sweden provide explanations in its next NIR, together with the specific carbon content values of Swedish MK1 and MK3 diesel.	In submission 2010, these specific carbon content values are shown in Appendix 20.
submission 2007/2008	Energy	30. The ERT recommends that Sweden provide specific information in its next NIR on how technology improvements influence the CH ₄ EFs from biomass burning.	In submission 2010, extensive information on this issue is included in NIR Annex 2. Very detailed information is provided in Paulrud et al, 2005: Methane emissions from residential biomass combustion. This report can be provided to the ERT if requested.
Submission 2009	Energy	32. Report emissions from some categories were reported as "NE", such as CO ₂ , CH ₄ and N ₂ O from venting of oil and gas and flaring of gas for all years.	Emissions from venting are considered to be included in other subsectors in 1.B.2 (see section 3.3.2.2). Flaring of natural gas is probably not occurring, but if occurring, it is included in flaring of oil.
Submission 2009	Energy	44. Clearly describe recalculations for off road vehicles and machinery made with the logic for making the revisions in the NIR in its next annual submission.	In submission 2011 there are no recalculations on this category. All recalculations made in other categories are clearly described in the NIR and in the CRF.
submission 2007/2008	Energy and Industrial processes	28, 29, 33, 35, 36. Sweden uses a CS-method to estimate and allocate CO ₂ emissions from pig iron production, not in line with the good practice guidance as this method allocates all CO ₂ emissions to the output (i.e. the blast furnace), rather than using an input based CO ₂ calculation method.	Addressed in submission 2010. Emissions have been revised for all years and reported in accordance with IPCC Guidelines (see NIR section 4.4.1)
Submission 2009, (Submission 2010)	Energy	38, (42). Investigate how distribution of marine distillate fuels and residual fuel oils between domestic and international navigation data corresponds to the definition of international and domestic marine transport in IPCC and explain large inter-annual variations.	Fuel data in the Monthly fuel, gas and inventory statistics, which is used as activity data for estimating emissions for national navigation and international maritime bunkers, has been analyzed in a SMED study (Eklund et al. 2011. Emissions from navigation and fishing including international bunkers). It has been found to be of good quality and fuels used for domestic and international navigation have been separated correctly and in line with IPCC Guidelines.
Submission 2010	Energy	33. Provide information and improve the transparency on the amount of fuel consumed in construction.	A table showing fuel consumption in the construction sector has been included in section 3.2.14.2
Submission 2010	Energy	34. For the largest iron and steel plants include explanatory information on the accuracy of AD and the consistency of the time series of data, and ensure the accuracy of its reporting without compromising the consistency of the time series of data.	Information is included in sections 3.2.9.2.1 and 0
Submission 2010	Energy	35. Investigate the reliability of the different sources of information, use appropriate and consistent sources of data for NCVs for its next submission, and provide the justification and reasoning for revising the previously used NCVs.	Explanations are provided in Annex 2.
Submission 2010	Energy	37. Provide explanatory information on the rationale for recalculations.	Explanations are provided in the section "Source specific recalculations" for those CRF categories where recalculations have been made in the current submission.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2010	Energy	43. Report correct notation key for CO2 emissions from solid fuels use in agriculture/forestry/fisheries	The notation key is "NO" for 2001 and onwards which is correct since no solid fuels have been used in this category since 2000.
Submission 2010	Energy	44. Provide information in the NIR on the fluctuating trend (e.g. large inter-annual changes) in the CO2 IEFs of other fuels for public electricity and heat production.	Information is provided in the section "Uncertainties and time-series consistency" (section 3.2.6.3)
Submission 2010	Energy	46. For civil aviation explain in more detail the method used to estimate the AD for 2008.	The estimation of CO2 from civil aviation is based on the data on supply and delivery of petroleum products from Statistics Sweden. Non-CO2 emissions are based on information from the Swedish Transport Agency and adjusted to match the delivered amount of aviation fuels (see NIR section 3.2.15)
Submission 2010	Energy	48. Explain the approach and method used to estimate fugitive emissions from distribution of oil products.	Explanations are provided in section 3.3.2.2
Submission 2009, (Submission 2010)	Energy	31, 36, 39, (39). The ERT reiterates the recommendation made by the previous review that Sweden investigates the cause of the difference between the data reported to the IEA and that reported to the UNFCCC.	An study ² was carried out in 2010. The recommendations from that study have been implemented when possible, as described in Annex 4 (Reference approach). The reference approach was revised in submission 2013.
Submission 2011	Energy	43. The ERT recommends that the Party continue to use this framework of verification and QC when considering the implementation of additional EU ETS data into the national inventory.	The verification and QC routines mentioned are used in submission 2013
Submission 2011	Energy	47. The ERT noted that in the CRF tables, small inconsistencies occur between tables 1.C and 1.A(b) for jet kerosene used in international aviation bunkers for 2009, and for gas/diesel oil and residual fuel oil used in international marine bunkers for all years. The ERT recommends that Sweden correct these discrepancies in its next annual submission.	This has been corrected in submission 2012. The same activity data and thermal values are now used for jet kerosene in CRF 1C and 1Ab.
Submission 2011	Energy	48. The EFs for coke oven gas and steel converter gas used in public electricity and heat production is based on measurements from one plant conducted in 2001. The ERT considers that EFs could vary over time and between plants. The ERT recommends that Sweden describe any recalculations and changes clearly in the NIR of its next annual submission.	Since submission 2012, facility specific emission factors are used.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2011	Energy	50. Sweden reports the CO ₂ emissions from the combustion of some CH ₄ and CH ₄ -based gas mixtures under liquid fuels in chemicals (NIR, page 95). The ERT recommends that Sweden use fuel-specific and year-specific CO ₂ EFs will be used in its next annual submission, review the allocation and clearly explain any recalculations in its next annual submission, because depending on the mixture it may be more appropriate to report these emissions under gaseous fuels.	Since submission 2012, year specific emission factors based on data from the plants using the fuel used for 2001 and later years, and this is described in NIR Section 3. It has not been possible to determine how much of the gas that is of gaseous origin, but since the coherence between the reference and sectoral approaches is good for gaseous fuels, it is probably correct to assume that most of the by product gas is of liquid origin.
Submission 2011	Energy	53. In its 2011 annual submission, Sweden has reported CO ₂ , CH ₄ and N ₂ O emissions from the transfer losses of gas-works gas under the distribution of oil products category. The ERT recommends that Sweden revise the allocation of these emissions, possibly allocating them to solid fuel transformation or other (fugitive emissions from solid fuels). The ERT reiterates the recommendation from the previous review report that Sweden describe the method to estimate emissions from the transfer losses of gas-works gas.	The description in NIR has been improved. Allocation and calculation methods will be reviewed in 2013.
Submission 2011	Energy	51. In CRF table 1.B.2, Sweden has reported CO ₂ and CH ₄ emissions from venting of oil, gas and combined as included elsewhere ("IE"). The ERT recommends that Sweden allocate these emissions correctly or, if this is not possible, include additional information indicating where they are allocated in its next annual submission.	Additional information has been added to the NIR, section 3.3.2.2.8 Venting (CRF 1.B.2.C.1), to improve transparency.
Submission 2011	Energy	52. The ERT agrees with these estimates and strongly recommends that Sweden report estimates for CO ₂ emissions from natural gas transmission and CH ₄ emissions from natural gas transmission and distribution and in its next annual submission.	As ERT recommends Sweden now reports estimates for CO ₂ from natural gas transmission and CH ₄ estimates from natural gas transmission and distribution in its annual submissions.
Submission 2011	Energy	54. The ERT noted that Sweden in its 2011 annual submission did not estimate CH ₄ and N ₂ O emissions from the combustion of ethanol. The ERT considers the potential problem to be solved and recommends Sweden to continue the reporting of these estimates in its next annual submission.	See NIR 3.2.16.2
Submission 2009	Energy	35. The ERT reiterates the recommendation made by the previous review that Sweden reconciles the differences between the reference approach and the sectoral approach.	Data and methodology for the reference approach have been reviewed and revised in the three last submissions. Although much effort has been made, there are still quite large differences between the reference approach and the sectoral approach for some years. Detailed information that explains the differences are provided in NIR and Annex 4. Further explanations are given in a SMED report from 2012. ³

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2010	Energy	38. ... the documentation box of CRF table 1.A referred to section 3.3.6 of the NIR, which relates to the chemicals category of the energy sector and does not address the issue raised above. The ERT recommends that Sweden correct this reference in its next annual submission. ERT noted that the difference between the reference and sectoral approaches is largely associated with solid fuels (24.9 per cent) and other fuels (2.2 per cent). The ERT recommends that Sweden investigate the ways to properly separate fugitive emissions and emissions from fuel combustion in the industrial processes sector.	Documentation box: corrected in submission 2013 Differences: Data and methodology for the reference approach have been reviewed and revised in the three last submissions. Although much effort has been made, there are still quite large differences between the reference approach and the sectoral approach for some years. Detailed information that explains the differences are provided in NIR and Annex 4. Further explanations are given in a SMED report from 2012. ⁴
submission 2011	Energy	44. the ERT noted that the documentation box of CRF table 1.A(c) referred to section 3.3.6 of the NIR, which is not correct. The ERT reiterates the recommendation of the previous review report that Sweden correct this reference in its next annual submission.	The documentation box is corrected in submission 2013
submission 2011	Energy	45. The ERT noted several differences between the data reported in Sweden's annual submission and those reported to the International Energy Agency (IEA), including peat production figures, imports and exports of lubricants and liquid fuel stock changes. In its NIR Sweden notes that these differences were studied and analyzed during 2010. However, due to budget and time constraints, the recommendations from the study have not been fully implemented in the current submission. The ERT recommends that Sweden act on the recommendations of the study to improve alignment between the two data sets and report on this in its next annual submission.	Data for peat and lubricants were revised in submission 2012. Other issues were dealt with in submission 2013.
submission 2007/2008	Industrial processes	37. Inconsistency in IEF since 2005 is not explained in the NIR. Sweden is encouraged to provide the time series for the content of calcium oxide (CaO) in clinker to validate the single average value (65%) for the whole period.	Addressed in submission 2010. Before 2005 the company used the default EF 0.525 for the estimates. From 2005, CO ₂ emissions are based on analysis on the CaO content in the clinker. Data from 2008 and 2009 show a CaO content variation between 63.9 to 67.6%. See NIR 4.2.1.
submission 2007/2008	Industrial processes	38. The ERT recommends that Sweden improves the transparency of the reporting of the methodology used to estimate the CO ₂ removals in the pulp and paper industry.	Addressed in submission 2010. Reported data is revised in submission 2010. The revised time series affects reported activity data as well as CO ₂ emission data. See NIR section 4.5.1
submission 2007/2008	Industrial processes	40. The ERT encourages Sweden to correct and improve its reporting in the NIR, and to improve the transparency of the applied approach by adding an allocation table of the annual amounts of limestone used and emissions for each category.	Reported data in 2A3 is revised in submission 2010. An allocation table is added in the NIR and activity and emission data is provided for 2005 - 2008. See NIR section 4.2.3

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2009	Industrial processes	46. Correct the discrepancy between figures in CRF and NIR.	Figures are correctly reported in submission 2011 as better SMED internal QC procedures are in place to minimize discrepancy between figures in CRF and NIR.
Submission 2009, (submission 2010)	Industrial processes	48, (54) Continue the discussion with the cement producing company and improve the CO ₂ estimates as appropriate.	Addressed in submission 2012, see NIR section 4.2.1.4
Submission 2009	Industrial processes	49. Complete the planned revision of methods and explain the new methods for pulp and paper industry in a transparent manner in the next annual submission.	Addressed in submission 2010, see NIR section 4.2.2
Submission 2009, (submission 2010)	Industrial processes	50, (62) Implement planned improvements in Limestone and dolomite use – CO ₂ with regards to transparency and allocation of emissions.	Addressed in submission 2010, see NIR section 4.2.3
Submission 2009	Industrial processes	52. Implement planned improvements in Iron and steel production and provides a transparent explanation of the revised estimation methods as well as the reallocation of emissions from the energy sector to this category.	Addressed in submission 2010. Emissions have been revised for all years and reported in accordance with IPCC Guidelines (see NIR section 4.4.1).
Submission 2009	Industrial processes	Include in the NIR a brief discussion on the results of the carbon balance checks.	
Submission 2009	Industrial processes	53. Clearly explain this in more detail in the NIR if it continues reporting CO ₂ from limestone use in iron and steel production.	Addressed in submission 2010, see NIR section 4.4.1
Submission 2009	Industrial processes	54. Make efforts to estimate the emissions from foam blowing that are not estimated currently, and include them in the next annual submission.	Data from the manufacturers is considered to be complete, see NIR section 4.7.2.
Submission 2009	Industrial processes	55. Calculate and include CO ₂ emissions from the use of calcium carbide using the default EF presented in the Revised 1996 IPCC Guidelines unless there is evidence showing that the calcium carbide produced is not used in the country.	Addressed in submission 2011, see NIR section 4.3.3
Submission 2011	Industrial processes	61. report on the progress of these discussions with the facilities about the accuracy of CO ₂ emissions from CKD reported by the facilities by clarifying the occurrence of CO ₂ emissions from CKD in its next annual submission.	After contact with the company we now know that CO ₂ emissions from CKD are included in ETS data (from 2005 and onwards). Recalculation performed to submission 2012.
Submission 2011	Industrial processes	62. include additional relevant information on how the CO ₂ EF for cement production is obtained in its next annual submission.	The EF 0.525 was used until 2004. From 2005 Plant-specific analysis data is used.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2011	Industrial processes	63. The ERT also noted that there is an inconsistency between the CO ₂ IEF for cement production reported for 2009 in CRF table 2(I)A-G (0.5593 t/t) and that reported in the NIR (0.5425 t/t).	Sweden has improved their QC procedures to minimize the risk of similar errors to occur.
Submission 2011	Industrial processes	69. Sweden has reported HFC and PFC emissions from the subcategory solvents as "NE" in its NIR (table 4.39) but as "NO" in CRF table 2(I). In its NIR (page 189).	The notation key for solvents in Sweden's NIR Table 4.39 has been changed to NO. The notations in the NIR and the CRF tables are now consistent. An explanation of why NO is used is given in NIR section 4.7.5 Solvents (2.F.5).
submission 2006	Agriculture	52. Break in time series 1995	Addressed in submission 2010. In the footnote to Table 6.6 an explanation to this is added.
submission 2006	Agriculture	54. Background data on enteric fermentation for cattle in CRF tables not consistent with calculations.	Addressed in submission 2010. The Background data this is referring to is not used in the actual calculations. The paragraph concerning the method for the calculations of enteric fermentation has been revised.
submission 2006	Agriculture	61. Sweden does not provide sufficient information in the NIR about the volatilization ratios of ammonia (NH ₃) and nitrogen oxide (NO _x) from the use of synthetic fertilizers and the application of animal manure.	A new paragraph named "Emission of ammonia" was added to NIR in submission 2011 to clarify this issue.
Submission 2007/2008	Agriculture	44. Provide detailed information in its NIR on the assumptions and national conditions supporting the calculation/selection of EFs (e.g. N ₂ O emissions from manure management and from agricultural soils).	The NIR has been updated with additional information about this.
submission 2007/2008	Agriculture	46. The ERT recommends that Sweden re-examine the preparation of table 4.B(b) to ensure that it accounts for all N excretion for the estimation of N ₂ O from manure management and for the quantification of N input for manure applied to soils and excretion on pasture range and paddock (table 4.Ds1). The ERT also recommends that Sweden ensure that its QA/QC procedures provide for accurate and correct completion of CRF tables in the agricultural sector.	This was due to a miscount that resulted in some incorrect activity data in submission 2009. It did, however, not affect the estimate of the actual emissions. This was corrected in submission 2010.
Submission 2007/2008	Agriculture	47. Revise this N-excretion rate in accordance with the conditions it is reporting for the number of pigs, that is, number of animals produced (including rotations) or number of average livestock at a given time.	The reference day for the estimation of number of pigs is the first of June. This should approximately be the average number of pigs for an arbitrary time of year.
Submission 2007/2008	Agriculture	48. Encourage Sweden to clarify the decrease in the IEF for solid storage from 0.0197 kg N ₂ O-N/kg N in 1990 to 0.0192 kg N ₂ O-N/kg N in 2006 and indicate which management systems underlie the IEF of 0.02 kg N ₂ O-N/kg N in the category other.	In CRF this value is constantly 0.02 for all reporting years.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
submission 2007/2008	Agriculture	51. The ERT recommends that Sweden ensure the consistency of information between CRF tables 4.B(b) and 4.D	In table 4.B (b) the total amount of nitrogen excreted is given. In table 4.D the amount of nitrogen directly lost as ammonia is subtracted from the total amount and the remaining amount is used as activity data for release of nitrous oxide.
submission 2007/2008	Agriculture	52. Sweden does not provide sufficient information in the NIR on the volatilization ratios of ammonia (NH ₃) and nitrogen oxide (NO _x) from the use of synthetic fertilizers, and the application of animal manure.	A new paragraph named "Emission of ammonia" was added to NIR in submission 2011 to clarify this issue.
Submission 2009	Agriculture	58. Following the recommendations from previous reviews, the ERT recommends that Sweden improve transparency by providing additional information on how the EFs are calculated for tier 2 methods.	Additional information about the country specific EF has been included in the NIR for submission 2010.
Submission 2009	Agriculture	59. Further details on how the EF for dairy cattle is developed.	This has been clarified in submission 2011.
Submission 2009	Agriculture	60. Provide further documentation to explain the trend in emissions for manure management and the changing IEF.	The trend is explained in the beginning of section 6, Agriculture.
Submission 2009	Agriculture	63. Provide further information in the NIR on the appropriateness for Swedish conditions of the factors: country-specific EFs of 0.8 and 2.5 per cent kg N ₂ O-N/kgN for nitrogen from synthetic fertilizer and nitrogen from manure applied to soils, respectively.	This has been clarified in submission 2011.
Submission 2009	Agriculture	64. Further explanation be provided in the NIR of the appropriateness of the use of this country-specific EF (pasture, range and paddock manure is 0.016 kg N ₂ O-N/kg N and unfertilized pastures/grasslands is based on a value in the range of 0.002–0.01 kg N ₂ O-N/kg N).	Sweden has now changed the EF for pasture, range and paddock manure to the IPCC default value of 2%.
Submission 2010	Agriculture	66. Use the notation key "NO" in accordance with the IPCC good practice guidance in particular in CRF table 4.D.	This has been corrected in submission 2011
Submission 2010	Agriculture	67. Improve the transparency of its reporting and include all relevant information on the country-specific EFs and methodologies used in the NIR, in particular for estimating N ₂ O emissions from agricultural soils.	The NIR has been updated with additional information in submission 2011.
Submission 2010	Agriculture	69. Include in the NIR and in the CRF tables additional information e.g. on milk yield per cow, digestibility of feed and gross energy intake, in order to improve the transparency.	This has been adjusted in submission 2011
Submission 2010	Agriculture	70. For CH ₄ from manure management include explanations for the fluctuations and the increasing IEF trends.	This information is included in the NIR since submission 2011

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2010	Agriculture	72. Account for the average livestock population of all growing animal species, namely piglets	This information can be found in Table 6.6
Submission 2010	Agriculture	73. For direct soil N ₂ O emissions, include further explanations regarding country-specific EFs.	This information can now be found in the paragraph "Emission factors" under "Direct Soil Emissions (CRF 4.D.1)"
Submission 2010	Agriculture	75. For N ₂ O from pasture, range and paddock manure, provide consistent information in the NIR and in the CRF tables, and transparent explanations of the relevant corrections and changes made.	The paragraph in the NIR has been updated and a new table included in submission 2012
Submission 2010	Agriculture	76. For N ₂ O emissions from atmospheric deposition, include all relevant information on country-specific methodologies in the NIR, and provide well-documented explanations with regard to all country-specific EFs and parameters used.	A new paragraph regarding the emissions of ammonia from fertilisers and manure (and by that, $Frac_{GASM}$ and $Frac_{GASG}$) has been included in the NIR.
Submission 2011	Agriculture	73. including further background information on the calculation of average milk yield, on the N flow model (STANK) and on the CH ₄ IEF trends for manure management.	Additional information about the calculation of milk yield is included in NIR, paragraph 6.2.2.1. Information on STANK in paragraph 6.3.2 and information of IEF for emission of CH ₄ in paragraph 6.3.2.1.
Submission 2011	Agriculture	74. report a time series for gross energy intake and Ym in both its NIR and CRF in its next annual submission.	Time series for GE and Ym are now provided in the CRF tables.
Submission 2011	Agriculture	75. include additional information on the calculation of annual average milk yield in the NIR of its next annual submission.	Additional information on the calculation of average milk yield is now included in NIR, paragraph 6.2.2.1.
Submission 2011	Agriculture	76. improve its use of notation keys in the additional information table for CRF table 4.A in its next annual submission.	The ERT recommended notation keys are now used in the CRF tables.
Submission 2011	Agriculture	77. apply consistent definitions of animal waste management systems in the NIR for both the CH ₄ and the N ₂ O calculations, in line with the IPCC good practice guidance equations 4.17 and 4.18.	This is a misunderstanding from the ERT. Our definition of PRP-manure is in agreement with GPG. $MS = \text{"fraction of animal species i's manure handled using manure system j"} (GPG \text{ eqv } 4.17)$. This is also our definition of PRP-manure. However, we realise that the equation in paragraph 6.3.2.2 in NIR was confusing. The equation has now been altered to more clearly reflect how we calculate emissions from AWMS (including PRP-manure).
Submission 2011	Agriculture	78. include this information on the value of piglets presented in the NIR assumes that piglets do not grow during their first year and that the number of piglets listed in the table takes this assumption into consideration.	This information is now included in paragraph 6.3.2.
Submission 2011	Agriculture	79. report the N fraction of legumes ($Frac_{NCRBF}$) and that of other crops ($Frac_{NCRO}$) separately in the additional information table of CRF table 4.D, in its next annual submission.	This has been done since submission 2012.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2011	Agriculture	80. apply the default EF and equation 4.18, in line with the IPCC good practice guidance, in its next annual submission.	This has been revised in the Swedish 2012 resubmission.
Submission 2011	Agriculture	81. provide more information on the STANK model in the NIR of in its next annual submission, especially showing whether underlying studies reflect field data, expert judgement or studies reported in the scientific literature.	Information on this is now included in paragraph 6.3.2.
Submission 2010	LULUCF	20. Perform key category analysis at a more disaggregated level of categories, and as a result revise its key categories for activities under Article 3, paragraphs 3 and 4, of the Kyoto Protocol, for its next annual submission and explain in the NIR how it plans to use the key category analysis to prioritize improvements in the inventory.	Sweden performed the analysis on the level suggested by the ERT.
Submission 2010	LULUCF	82. Consider using the category other land to adjust the annual fluctuations in its total land area.	Sweden has started investigating how the estimates could be improved using complementary methods for the last reporting years. In this submission the inconsistency in total area as well as other categories has been solved using extrapolation for sample series ending earlier than 2010. See section 7.2.2.2.
Submission 2010	LULUCF	86. The ERT noted that there were some errors in the NIR, especially in the references related to the figures and tables provided in the NIR and recommended Sweden to improve the consistency of the reporting within the NIR, enhance its QC procedures and correct such errors in its next annual submission.	This has been checked and corrected in submission 2011.
Submission 2010	LULUCF	87. The ERT recommended that Sweden provide in its NIR more information on the drivers of the emission trends and their impact on the annual carbon stock changes, in order to improve the transparency of the reporting and facilitate the review of the inventory of the Party's next annual submission.	This was implemented in Submission 2011.
Submission 2010	LULUCF	88. The ERT noted that there is an inconsistency between the reported area of land converted to forest land and the area subject to the afforestation/reforestation activities reported under Article 3, paragraph 3, of the Kyoto Protocol and recommended that Sweden revise the method used to identify the relevant area and ensure the consistency of its reporting under the Convention and under the Kyoto Protocol, thus improving the transparency and accuracy of its reporting.	Sweden has started investigating how the estimates could be improved using complementary methods for the last reporting years. In this submission the inconsistency has been solved using extrapolation for sample series ending earlier than 2010. See section 7.2.2.2. However since AR and D areas are aggregated since 1990 and the area accumulation of land converted to Forest land starts earlier these are not comparable per se.
Submission 2009	LULUCF	59. Improve its approach for determining land-use change in order to report a consistent time series of annual land-use change matrices, as is suggested in the IPCC good practice guidance for LULUCF.	Sweden has increased the transparency in the description of methods used in the inventory in accordance with IPCC GPG.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2009	LULUCF	55. Improve the transparency of its inventory by providing all the necessary documentation and information in its future submissions, in accordance with the IPCC good practice guidance for LULUCF.	Sweden has increased the transparency in the description of methods used in the inventory in accordance with IPCC GPG.
Submission 2007/2008	LULUCF	70. Clarify in the NIR whether annual land-use change data are used to produce the estimates and report a consistent time series of these annual land-use change data in accordance with the IPCC good practice guidance for LULUCF.	Now clearly addressed in NIR
Submission 2007/2008	LULUCF	54. The ERT encourages the Party to improve the completeness of its reporting in its future annual submissions by providing estimates and relevant information for categories that are not estimated.	Sweden's reporting of LULUCF is now complete except for categories that are currently not mandatory to report.
submission 2007/2008	LULUCF	55. The ERT recommends that Sweden improve the transparency of its inventory by providing all the necessary documentation and information in its future submissions, in accordance with the IPCC good practice guidance for LULUCF.	Figures have been introduced to make it easier for the reviewer to follow the quite complicated sample design use. The text is continuously improved. See NIR section 7.3.1. Estimators are moved to Annex to NIR.
submission 2007/2008	LULUCF	56. The ERT recommends that Sweden consider the use of notation keys NO or IE either for gains or losses when the stock change method is applied.	Sweden is following this advice from submission 2010.
Submission 2007/2008	LULUCF	60. The total area of organic soil reported in the LULUCF sector does not match the area of cultivated organic soils reported in the agriculture sector (CRF table 4.Ds1).	In submission 2010, the reporting of total cropland areas in the agriculture sector (CRF 4) has been revised to be consistent with the reporting in the LULUCF sector. The area of cultivated organic soils still differ due to differences in data sources. The full correction is introduced in submission 2011.
Submission 2007/2008	LULUCF	63. Sweden reports a net carbon increase for the living biomass (except 1991) and dead organic matter pools for all years. The ERT recommends that Sweden provide an explanation for this trend in its next NIR in order to improve the transparency of its reporting.	The transparency in describing this issue has improved in submission 2010. See NIR section 7.1.
Submission 2007/2008	LULUCF	66. With the exception of the years 2002, 2003 and 2005 Sweden reports a net carbon increase in living biomass associated with land-use change from forest land to grassland. The ERT recommends that Sweden provide an explanation for the outlined trend in its next NIR.	The transparency in describing this issue has improved in submission 2010. See NIR section 7.1.
Submission 2007/2008	LULUCF	67. With the exception of the years 1992 and 2006, Sweden reports a net carbon increase in living biomass associated with land-use change from forest land to settlements. To improve the transparency, the ERT recommends that Sweden provide an explanation for the outlined trend in its next NIR.	The transparency in describing this issue has improved in submission 2010.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
submission 2007/2008	LULUCF	68. Sweden has not been able to separate emissions from organic and mineral soils (CRF table 5 (III)). The ERT recommends that Sweden improve its methodology in order to be able to report the two soil categories separately.	Categories are reported separately in submission 2010.
Submission 2006	LULUCF	69. Improve consistency in the CRF tables by correcting the errors.	Corrected.
Submission 2006	LULUCF	70. Estimate CO ₂ emissions from land converted to wetland.	Addressed in submission 2010. No longer reported unmanaged.
Submission 2011	LULUCF	84. Sweden indicated that all reported years in its 2012 annual submission will be based on five different inventory cycles to maximize accuracy, and that all emissions and removals for the last years in the time series will be recalculated. To increase transparency, the ERT recommends that Sweden include additional information on these recalculations in the NIR of its next annual submission.	84. The information related to this issue are described in the Annex to NIR Table A 3:2.1. The new method of extrapolating data is in detail described in section 7.2.2.2, and consequences compared to not introducing extrapolation in Figure 7-6. See also the footnotes on page 267 and on page 271.
Submission 2011	LULUCF	87. provide estimates for land converted to wetland or otherwise revise its use of notation keys for carbon stock changes in wetlands in its next annual submission.	87. As a response to this issue Sweden introduced "Peatland" as a separate sub-category to Wetlands remaining Wetlands. See section 7.3.1.6 and 7.6 (recalculations).
Submission 2011	LULUCF	90. The ERT recommends that Sweden address the discrepancies between the areas reported for the different land-use categories (NIR tables 7.1a and 7.4) and also for the total area of the country (NIR tables 7.1a, 7.1b and 7.4 report 45,250 kha, 45,158 kha and 45,080 kha, respectively, for total area).	Sweden reports also a land use matrix 1990-2011 based on one cycle (estimates based on approximately 20% of the plots). Using such approach the total land and fresh water area deviates only 0.3%. See footnote on page 421
Submission 2011	LULUCF	92. The ERT noted that there are large differences in the areas of all land-use categories reported in the 2011 annual submission compared with the 2010 annual submission, especially in forest land area. The ERT recommends that Sweden demonstrate that this discrepancy is reduced with decreasing uncertainties for each year that adds to the full fiveyear inventory cycle in its next annual submission.	92. The introduced extrapolation (see question 84) has completely removed the problem with random variations for the most recent years.
Submission 2011	LULUCF	93. The ERT recommends that Sweden include additional information on the drivers of the emission/removal trends and their impacts on the annual carbon stock change and improve the transparency of the information in its next annual submission.	93. See the overview section in NIR chapter 7. On page 267 and 268 we are trying to verify estimates by the IPCC default method (studying growth and harvests). In chapter 7.4 we refer to a study (2003), validating estimates based on measurements by using different models.
Submission 2006	Waste	75. The ERT recommends that Sweden provide further information on the utilization of gas recovery in its next NIR.	From submission 2008, the use of recovered gas is described in NIR.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2006	Waste	79. The ERT recommends that Sweden use the notation key "not estimated" ("NE") for CH ₄ emissions from wastewater treatment, instead of "included elsewhere" ("IE"), in CRF table 6.B.	From submission 2010, the notation key NE is reported for CH ₄ emissions from wastewater treatment (sludge). In submission 2011, by following the recommendations of ERTs Saturday paper, emission estimates has been reported for CH ₄ emissions from wastewater treatment (wastewater). This is further described in the NIR section 8.3.1.2
Submission 2006	Waste	76. Measure CH ₄ and N ₂ O emissions from hazardous waste incineration periodically on-site.	Addressed in submission 2010, see NIR section 8.4
Submission 2007/2008	Waste	73. CO ₂ emissions from solid waste disposal on land could be better reported using the notation key NO.	From submission 2010, the notation key NO is reported for CO ₂ emissions.
Submission 2007/2008	Waste	74. CH ₄ emissions from wastewater treatment have been reported as NE	In submission 2011, by following the recommendations of the ERT, emission estimates has been reported for CH ₄ emissions from wastewater treatment (wastewater). This is further described in the NIR section 8.3.1.2
submission 2007/2008	Waste	75. The ERT recommends that Sweden change the notation key from IE to NE for the emissions from sludge resulting from treatment during the wastewater handling process.	In submission 2011, emission estimates has been reported for CH ₄ emissions from wastewater treatment (sludge).
Submission 2007/2008	Waste	74. Attempt to estimate CH ₄ emissions from wastewater treatment as part of the general assessment for "NE" categories and to provide the background data on industrial wastewater in CRF table 6.B.	CH ₄ emissions from wastewater treatment (wastewater and sludge) are estimated in submission 2011. This is further described in the NIR section 8.3.1.2
Submission 2007/2008	Waste	76. Clarifications on methodology Wastewater handling needed in NIR	In submission 2010, justifications have been made in NIR section 8.3.2 (See section 8.3.1.2.1 in NIR submission 2011)
Submission 2007/2008	Waste	80. CH ₄ and N ₂ O emissions from waste incineration are reported as 'NE'. The ERT recommends that Sweden calculate these emissions (however insignificant) using IPCC default EFs, rather than wait for the outcome of actual measurements.	Estimates of N ₂ O and CH ₄ emissions are included in submission 2010. See NIR section 8.4
Submission 2009	Waste	81. Clarify about unmanaged landfill sites throughout the time series from 1990 to 2007 and provide more information on managed and unmanaged landfill sites in Sweden.	In NIR submission 2011, more information is provided on the matter. This is further described in the NIR section 8.2
Submission 2009	Waste	82. Confirm which Tier is used in CH ₄ from Solid waste disposal on land and if necessary correct this information.	Tier 3 was changed to Tier 2 in submission 2010.
Submission 2009	Waste	83. Give more and updated information on the amount of landfill gas recovered that was used for energy and was flared.	More information was provided in NIR 2011. This is further described in the NIR section 8.2.1.2
Submission 2009	Waste	85. Provide some additional information about the lower value in 2006 for CO ₂ emissions from waste incineration.	Information is provided in section 8.4. Data are direct measurements from the facilities flue gases and are considered to be correct.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2010	Waste	99. Continue to estimate and report the CH ₄ emissions from industrial, domestic and commercial wastewater and sludge, ensure consistency between the information reported in the NIR and that in the CRF tables and include all necessary information (e.g. category description, methodological issues, uncertainty and time-series consistency, recalculations and planned improvements) in order to improve the transparency. Provide information on the rationale for using the notation keys to report on wastewater handling	Sweden has improved the reporting of CH ₄ emissions from industrial, domestic and commercial wastewater and sludge in submission 2010 (resubmission) and further more in submission 2011. Emissions and methodologies are presented and described in the CRF-tables and in NIR.
Submission 2010	Waste	100. Consider using time-dependent AD on protein consumption (e.g. statistics from the Food and Agriculture Organization of the United Nations) to further improve the accuracy.	Since submission 2011, Sweden is using existing time series data on protein consumption from the Swedish yearbook of agricultural statistics.
Submission 2011	Waste	108. justify why the country-specific fossil carbon fraction for all waste incinerated at that plant can be assumed to be the same as the value obtained from a study on MSW incineration. The ERT also recommends that Sweden provide more transparent information on the carbon content used in the estimation in its next annual submission.	In submission 2011 the fossil carbon fraction of 30% is only applied to the incinerated amounts of MSW. All other wastes (industrial and hazardous) incinerated at the plant are assumed to be 100% fossil. In submission 2013 the time series of reported biogenic and fossil CO ₂ are revised. For the years 2008, 2009, 2010 and 2011 the company has, besides total emissions of CO ₂ , also reported CO ₂ with respect to biogenic or fossil origin. The company has based their reporting of biogenic and fossil CO ₂ on a detailed study performed in 2008. In submission the allocation between fossil and biogenic CO ₂ (see NIR, 8.4 Waste incineration (CRF 6.C)).
Submission 2011	Waste	105. Sweden has reported CH ₄ emissions from wastewater handling for the whole time series. Data for the period 1990–2004 were obtained by extrapolating backwards the data for 2005 because data on energy recovery is only available for the period 2005–2009. The ERT recommends that Sweden provide additional information on how this extrapolation was done, including how the potential changes, upgrades or modifications of the recovery system in each plant in the period 1990–2005 was considered, in its next annual submission.	See information in NIR 8.3.1.2.
Submission 2011	Waste	107. CH ₄ emissions from industrial sludge treatment in anaerobic plants are estimated based on a country-specific CH ₄ leakage factor during energy recovery. However, the ERT noted that this leakage factor is obtained from a study on two plants, which are assumed to be representative of all 138 plants that engage in the anaerobic treatment of sludge in Sweden (NIR, page 302). The ERT recommends that Sweden include additional information on how the Party ensures that the leakage factor obtained from two plants is representative of the other 136 plants and does not lead to an underestimation of emissions.	Note that this issue is NOT regarding industrial sludge treatment, but municipal waste water treatment (sludge). This is addressed in NIR section 8.3.1.2. To ensure a conservative estimate the upper value (7 %) of the leakage factor is used. We believe this is the most appropriate way of addressing this issue – good practice – as this is a very small source.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Actions as a result of ERT recommendations
Submission 2011	KP	111. The ERT noted that N2O emissions from the drainage of soils under forest management is reported as "NE" in table NIR-1 but as "NA" in table 5(KP-II)2. The ERT recommends that Sweden address this inconsistency in its next annual submission.	111. Sweden has now changed notation keys from "NA" to "NE" in table 5(KP-II)2.
Submission 2011	KP	112. In the NIR (page 340), Sweden has indicated that the estimates for carbon stock changes for afforestation, reforestation and deforestation are "very uncertain" and, for forest management, "quite uncertain". The ERT noted that, although Sweden has chosen to account for KP-LULUCF activities at the end of the commitment period, reporting on KP-LULUCF activities is mandatory since the 2010 annual submission, and therefore recommends that Sweden make every possible effort to reduce the uncertainty of these estimates in its next annual submission.	112. From submission 2012, the accuracy of estimates of areas and living biomass for AR, D and FM has been improved substantially using a similar extrapolation as for the UNFCCC reporting (question 84), see chapter 11.1.
Submission 2011	KP	113. The ERT recommends that Sweden justify how 2–3 sample plots can be representative of an activity or, if not possible, address this sampling issue, in its next annual submission.	113. See section 11.3.1.4 and 11.3.1.5 in the NIR.
Submission 2011	KP	116. In response to a question raised by the ERT during the review Sweden explained why it is not necessary that afforestation and reforestation areas and land converted to forest land areas are exactly comparable. To improve transparency, the ERT recommends that Sweden include this information in its next annual submission.	116. See NIR section 11.1.
Submission 2011	KP	117. The ERT noted that Sweden considers its estimates of carbon stock changes for deforestation as "very uncertain" and therefore reiterates the recommendation made in paragraph 112 above that Sweden make every possible effort to reduce the uncertainty of these estimates in its next annual submission.	117. This question refers to uncertainty and the reviewer refers back to question 112. We believe that we could answer this question by repeating our answers to questions 112 and 113.
Submission 2011	KP	119. The table NIR-2 shows that, for 2009, the area for forest management has increased by 720.2 kha while at the same time the area for afforestation and reforestation has not changed and the deforestation area has increased by 34.1 kha. The ERT recommends that Sweden address the inconsistencies in the Party's reporting of areas for KP-LULUCF activities in its next annual submission.	119. This question about NIR 2 (the land use matrix) is answered by our reply to question 116.

¹ FCCC/IRR/2006/SWE, FCCC/ARR/2008/SWE and FCCC/ARR/2009/SWE and FCCC/ARR/2010/SWE. The draft ARR for the 2011 submission has not been available in time for implementation in submission 2012.

² Hedlund & Lidén, 2010

³ Anderson, Eklund, Gerner & Gustafsson, 2012

⁴ Anderson, Eklund, Gerner & Gustafsson, 2012

Table 10-5. Recommendations from the UNFCCC review process not yet implemented in the Swedish inventory.

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Rationale for not yet adapting ERT recommendation and possible improvement plan
Submission 2011	General	19. The ERT noted that the areas identified by Sweden as areas for further improvement could be better linked with the pending recommendations from previous review reports, and the ERT therefore recommends that Sweden include any recommendations from the previous review report not yet addressed in the specific sections on category-specific planned improvements in the NIR of its next annual submission	The information will be kept in table 10.6 to avoid duplication error and double work. We believe this recommendation should have been expressed as an encouragement by the ERT.
Submission 2007/2008	Energy	79. The ERT recommends that only one value (e.g. 25 kg CO ₂ /GJ) be used for all years for incinerated municipal waste in order to maintain time-series consistency and consistency with reporting on incineration in category 1.A.	A study performed in 2009 ² addressed this issue. It was concluded that there is not enough evidence to revise the emission factor for years prior to 2003, and SEPA decided not to revise the emission factor at all. We are aware of this issue and our ambition is to improve this emission factor in future submissions.
Submission 2009	Energy	32. Report emissions from some categories were reported as "NE", such as the CH ₄ and N ₂ O emissions from mobile military use of biomass (FAME) for the years 1999 to 2001, CO ₂ from oil transport for all years,	In submission 2011 all categories for which IPCC default methodologies exist are estimated. There are no emission factors available to estimate CH ₄ and N ₂ O emissions from mobile military use of biomass for the years 1999 to 2001 but emissions are expected to be minor. Sweden lack information on CO ₂ emissions from oil transport and there is no IPCC default methodology for CO ₂ from oil transport using tanker ships.
submission 2011	Energy	49. The ERT noted that Sweden's IEF for N ₂ O (8.41–15.97 kg/TJ) is the highest across all Parties for all years for the public electricity and heat production category (0.06–15.97 kg/TJ). The high IEFs are driven largely by high N ₂ O EFs for coal and coke. The N ₂ O EFs for these fuels are sourced from a country-specific study conducted in 2004.6 The ERT recommends that Sweden review the N ₂ O EFs and report on any updates or provide further justification for the existing factors in the NIR of its next annual submission.	This will be addressed in the coming submissions.
submission 2007/2008	Industrial processes	41. For CO ₂ and N ₂ O from solvent and other product use, Sweden reported identical emission estimates for 2005 and 2006 in its 2008 submission. The ERT recommends that Sweden improve its data collection procedures in order to estimate final emissions in a timely manner.	Not implemented but addressed in NIR Annex 3.3. We intend to make special arrangements for data collection for submission 2014 in order to avoid using preliminary data in that submission.
Submission 2010 (Submission 2011)	Industrial processes	59. (65) Allocate CO ₂ emissions from limestone and dolomite used in iron and steel production to limestone and dolomite use category.	Not implemented, but addressed in NIR section 4.4.1.2.
Submission 2009, (Submission 2010)	Industrial processes	56. (63). 68. Report CH ₄ and N ₂ O from combustion of cooking liquor in the pulp, paper and print category under the energy sector, in accordance with the Revised 1996 IPCC Guidelines.	This will be addressed in the coming submissions. Addressed in NIR section 4.5.1.
Submission 2011			

Review ¹	Sector	Paragraph and recommendation in report (shortened)	Rationale for not yet adapting ERT recommendation and possible improvement plan
Submission 2011	Industrial processes	66. The ERT recommends that Sweden report emissions from auto-producers in the iron and steel processes in accordance with the Revised 1996 IPCC Guidelines. To increase transparency in the allocation of emissions, the ERT encourages Sweden to provide a detailed carbon mass balance for all the iron and steel processes.	Sweden will continue to report emissions allocated as prescribed in the 2006 IPCC Guidelines. Due to confidentiality reasons the mass balances cannot be included in the NIR. However, they can be delivered to the UNFCCC Expert Review Team upon request (see NIR section 4.4.1.2).
Submission 2007/2008	LULUCF	75. The ERT reiterates the recommendation from the previous review that Sweden improve its methodology (to separate emissions from organic and mineral soils (CRF table 5 (III))) in order to be able to report the two soil categories separately in future submissions.	Currently we do not have appropriate information on exactly on which land these changes occur. Therefore the emissions are reported aggregated.
Submission 2007/2008	LULUCF	69. The ERT noted that Sweden may be underestimating the C stock increase in living biomass and recommends that Sweden verify differences in IEF compared to Finland and Norway and make revisions if necessary.	Comparing IEF may not tell the whole story. The annual harvest may differ significantly between years in and between countries. Therefore the difference may not be seen as exceptional.
Submission 2009	LULUCF	57. Report the subcategory mire under grassland, further distinguishing between managed and unmanaged land subcategories.	Grasscovered mires are always saturated with water and therefore reported under Wetlands.
Submission 2006	Waste	77. Account for CO ₂ emission only from non-biogenic waste incineration sources according to the IPCC Good Practice Guidelines.	This will be addressed in the coming submissions.
Submission 2007/2008	Waste	72. Include information on time series for industrial organic waste in order to provide a more complete picture of municipal solid waste AD.	No data are available, which has been further described in NIR section 8.2.1.2.2.
Submission 2007/2008	Waste	78. Include information on biogenic fraction of incinerated municipal waste in its next NIR, and update and validate this value regularly as the biogenic fraction of incinerated municipal solid waste varies over time. It is good practice to assume that the composition of incinerated municipal solid waste is similar to that of generated municipal solid waste (IPCC good practice guidance, page 5.28).	This will be addressed in the coming submissions.
Submission 2011	Waste	103. Sweden has used the IPCC first order decay method with mostly default parameters and country-specific AD for estimating CH ₄ emissions from solid waste disposal sites. The ERT recommends that Sweden develops country-specific parameters to estimate the emissions from this category in its next annual submission.	This will be addressed in the coming submissions.
Submission 2011	Waste	106. For estimating CH ₄ emissions from domestic and commercial wastewater treatment apply the method from the Revised 1996 IPCC Guidelines in order to improve the accuracy of emission estimates or justify why the check method gives more accurate estimates in its next annual submission.	This will be addressed in the coming submissions.

¹ FCCC/IRR/2006/SWE, FCCC/ARR/2008/SWE, FCCC/ARR/2009/SWE and FCCC/ARR/2010/SWE. The draft Annual Review Report for the 2011 submission has not been available in time for implementation in submission 2012.

² Paulrud, Fridell Strippel, 2009

10.5 Major changes in methodological descriptions

Table 10-6. Documentation of major changes in methodological descriptions compared to previous year NIR.

Greenhouse gas source and sink categories	Description of methods	Recalculations	Reference
Total (Net Emissions)			
1. Energy			
A. Fuel Combustion (Sectoral Approach)			
1. Energy Industries			
2. Manufacturing Industries & Construction			
3. Transport	X	X	See NIR 3.2.16
4. Other Sectors			
5. Other			
B. Fugitive Emissions from Fuels			
1. Solid Fuels			
2. Oil and Natural Gas	X	X	See NIR 3.3.2
2. Industrial Processes			
A. Mineral Products			
B. Chemical Industry			
C. Metal Production			
D. Other Production			
E. Production of Halocarbons and SF6			
F. Consumption of Halocarbons and SF6			
G. Other			
3. Solvent and Other Product Use			
4. Agriculture			
A. Enteric Fermentation	X	X	See NIR 6.2.5
B. Manure Management			
C. Rice Cultivation			
D. Agricultural Soils	X	X	See NIR 6.4.2.5
E. Prescribed Burning of Savannas			
F. Field Burning of Agricultural Residues			

Greenhouse gas source and sink categories	Description of methods	Recalculations	Reference
	Please tick where the latest NIR includes major changes in methodological descriptions compared to the previous year NIR	Please tick where this is also reflected in recalculations compared to the previous year CRF	If ticked please provide some more detailed information for example related to sub-category, gas, reference to pages in the NIR, etc
G. Other			
5. Land Use, Land-Use Change and Forestry			
A. Forest Land			
B. Cropland			
C. Grassland			
D. Wetlands			
E. Settlements			
F. Other Land			
G. Other			
6. Waste			
A. Solid Waste Disposal on Land			
B. Waste-water Handling			
C. Waste Incineration	X	X	See NIR 8.4
D. Other			
7. Other (as specified in Summary 1.A)			
Memo Items:			
International Bunkers			
Aviation			
Marine			
Multilateral Operations			
CO2 Emissions from Biomass			
NIR Chapter	Description		Reference
	Please tick where the latest NIR includes major changes in descriptions compared to the previous year NIR		If ticked please provide some more detailed information for example reference to pages in the NIR

PART 2: SUPPLEMENTARY INFORMATION REQUIRED UNDER ARTICLE 7, PARAGRAPH 1

11 KP-LULUCF

11.1 General information

Sweden provides supplementary information under Article 7 of the Kyoto Protocol (KP) for the Land Use, Land-Use Change and Forestry sector. The requested information is further specified in Decision 15/CMP.1, 16/CMP.1 and IPCC GPG for LULUCF (IPCC²⁶⁸).

The activities Afforestation and Reforestation (AR) and Deforestation (D) are quite uncommon in Sweden, each representing around 10 kha annually in average (Table 11.1 and 11.2). The accumulated AR and D-areas steadily increase by time when estimates are based on a full set of inventory plots (30,000 plots). However, if based only on the most recently re-measured plots, this might not always be the case. This is due to the five-year inventory cycle, where the estimates for the five recent years are based on a decreasing number of plots. To avoid a risk of an incorrect decrease in AR and D area and to improve the accuracy of the estimates, inventory cycles without a full record to 2011 are extrapolated (see 7.2.2.2 and Figure 7-6). Since areas under ARD are accumulated, their extrapolations are based on the trend of the five previous years to the actual year, while extrapolation for area under Forest management (FM) and living biomass for all Art. 3.3 and 3.4 activities are based on a running average of the five years previous to the actual year. There are several options to make this extrapolation but the method chosen has the advantage that the total land and fresh water area is constant over time.”Five years” is chosen as a trade off between being enough to reduce random variation and be reasonably up to date. Each submission, data for the five recent years are re-calculated to limit a potential small risk of bias induced by the extrapolation (a pilot study by historical data indicated that the result of re-calculated data using measured full records of sample plots does not deviate significantly from extrapolated data). IPCC recommends a five-year inventory cycle and to re-calculate data when the intention is to improve the accuracy. Chapter six in IPCC 2003 GPG describes that extrapolation is a valid approach to improve estimates for years with missing data. Comments by previous reviewers to report a constant total area and consistent areas under AR and D, initiated the introduction of extrapolation. The methodology has also the advantage to improve the accuracy of estimates. The only potential disadvantage is the challenge to trace back extrapolated carbon of living biomass to land use from 1990 and onwards.

²⁶⁸ Intergovernmental Panel on Climate Change, 2003

Sweden has elected the activity Forest management (FM) under Article 3.4 of the Kyoto Protocol (KP). The KP-reporting of FM and AR harmonize (areas) almost with the UNFCCC-reporting of Forest land and land converted to Forest land. However, according to IPCC GPG for LULUCF (e.g. decision tree 4.1.1), land that during the commitment period is converted from Forest land should either be reported under D or under FM. Thus Forest land converted to Wetland or Other land (both unmanaged) are reported under FM (KP) and under Forest converted to Wetland/Other land (UNFCCC). This explains the insignificant difference in area between FM+AR and Forest land+land converted to Forest land for years 2010 and 2011 (see also NIR 2). Due to a slow growth rate in boreal forests, land under AR will not be considered harvested during the first commitment period²⁶⁹ and these juvenile forests are not fertilized. Therefore, direct N₂O emissions from N fertilization and emissions from forest fires are reported only under FM. Forest fires –both natural and wild-fires– are uncommon and, this far, has not been registered on ARD-land. N₂O emissions from disturbance associated with land use conversion from Forest land to Cropland are reported under D. N₂O emissions from drainage of soils are not reported (voluntary). Liming is assumed only to occur on agricultural land and is assumed to occur only on Cropland remaining Cropland. Thus no liming is reported under the KP (In 2011, 6,300 ha is estimated to be D to Cropland, which constitute only 0.21 % of the total Cropland area.).

The KP-reporting uses the same institutional arrangements, national system and corresponding QA/QC procedures as for the UNFCCC reporting. Emissions reported under Article 3, paragraph 3 and 4 are not overlapping with those emissions reported under KP Annex A. The section below focuses on differences in aggregating underlying data between the UNFCCC- and the KP-reporting.

The same underlying methodology is used for the reporting under the KP of the LULUCF-sector as described for the UNFCCC reporting of LULUCF (chapter 7). The estimates of emissions/ removals and areas are based on permanent sample plots inventoried by the Swedish National Inventory of Forests covering all land and fresh water areas. A major difference from the UNFCCC reporting is that the carbon pool living biomass is separated into above ground and below ground living biomass and that the dead organic matter pool is separated into dead wood and litter in the reporting under the KP. Only emissions/ removals on land under the activities AR, D and FM are reported under the KP. ARD land is accumulated from 1990 using permanent sample plots covering all land and fresh water areas. Changes in carbon pools on ARD-land are reported for years 2008-2011.

Land under FM is accumulated from 1990 and changes in carbon pools are reported on such land 2008-2011.

Sweden has elected commitment period accounting for LULUCF for the first commitment period.

²⁶⁹ The concept “harvest” is important when a party claims to offset emissions from land under harvested AR-land (e.g. FCCC/KP/CMP/2005/8/Add.3 p.6) but no definition of “harvest” has been found. So Sweden assumes that “harvest” refers to emissions at final felling and such AR-land is not expected to exist in Sweden during the first commitment period.

Table 11.1 The accumulated area under activities AR, D and FM .

[M ha]	1990	.	.	.	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
AR	0.02				0.12	0.13	0.14	0.15	0.17	0.19	0.21	0.22	0.24	0.25	0.26	0.26
D	0.01				0.13	0.14	0.15	0.16	0.16	0.17	0.18	0.19	0.20	0.21	0.22	0.22
FM	28.20				28.09	28.08	28.07	28.07	28.07	28.08	28.11	28.15	28.18	28.18	28.17	28.17

Table 11.2 Emissions / removals (minus), CO₂ [Mton] from reported carbon pools in AR, D and FM for the fourth year in the commitment period (2011).

[Mton]	Above ground biomass	Below ground biomass	Dead wood	Litter	Soil organic carbon
AR	-0.60	-0.20	-0.01	-0.27	0.17
D	0.48	0.16	0.00	0.95	0.96
FM	-23.90	-8.10	-6.87	5.67	-4.44

11.1.1 Definitions of forest and any other criteria

For reporting purposes under the Kyoto Protocol, Forest land is defined, according to the FAO definition, as land with a tree crown cover (or equivalent stocking level) of more than 10 %, an area of more than 0.5 ha and a minimum height of 5 m. Both crown cover and height refers to maturity *in situ*, and consequently, Forest land could temporary be unstocked due to human intervention such as final felling. Normally such land is regenerated within a few years and Forest land is not considered deforested if not confirmed in field. Assessed land that meets the forest criteria above but where other land-use is predominating is not considered Forest land. For example, agriculture land normally fulfils the forest criteria except for the predominant land use and is not considered Forest land. Tree-rows narrower than 10 m are not considered forests. Roads and power-line routes within forests are considered forest only if they are narrower than 5 m. Tree covered areas less than 0.5 ha does not fulfil the forest criteria and is reported as belonging to the neighbouring land use category – this implies that carbon stock changes in living biomass may be reported under any land use category.

All Forest land is assumed managed. Thus, the definition of Forest land and the assumption that all Forest land is managed are consistent with reporting under the UNFCCC. The underlying data are also consistent for the whole reporting period. In fact, the area of Forest land under the UNFCCC reporting should nearly be equal the sum of the areas subject to activities Forest management and Afforestation/ Reforestation under the Kyoto Protocol (this far all land under Afforestation/ Reforestation has secondary classification Forest management and none under Deforestation).

The definition of Forest land is consistent with former reporting under the UNFCCC and to other international bodies such as the FAO. However, to be able to trace both gross and net land use transfers, only permanent sample plots are used in the reporting under the UNFCCC and the KP while both temporary (only visited once) and permanent (fixed position and re-inventoried) sample plots are normally used for most assessments and reporting of the Swedish forest situation to other bodies. In both cases the expected values of estimates are the same but estimates might vary from randomness of the sample.

Under the Kyoto Protocol it is central to distinguish between definitions of land use categories, activities and spatial assessment units (Figure 11.1). The definition of Forest land has a minimum area but this is not the case for activities. For Sweden the spatial assessment unit is a permanent sample plot (radius 10 m) and since this plot could be delineated into more than one land use category, deforestation close to 0 m² could be detected. Area-based sampling is used and each sample plot represents a certain area in the estimation algorithm so that all sample plots together represent the total land and fresh water area of Sweden. The Swedish NFI has the advantage that the sample frame covers all land- categories required for the UNFCCC-reporting. This is essential when both gross and net land use transfers over time have to be traced.

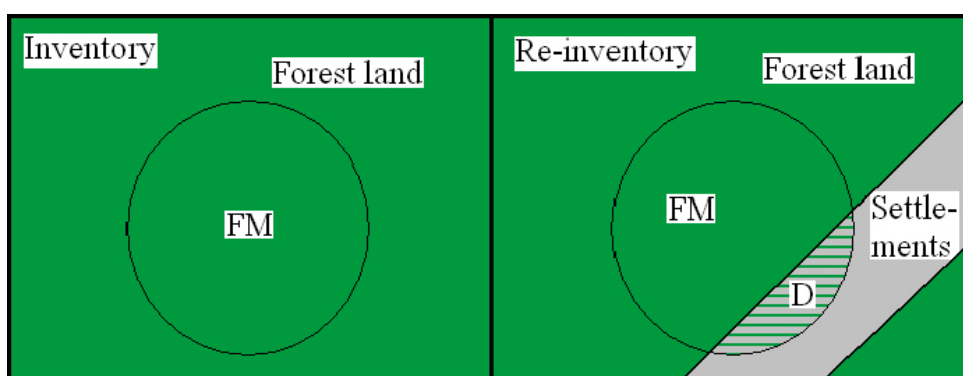


Figure 11.1 Example distinguishing the concepts of land use category, activities under the Kyoto Protocol, and spatial assessment unit in the Swedish sample based inventory: At the first inventory, only the land use category Forest land exist in an area but at the re-inventory part of the Forest land has been deforested to the land use category Settlements. Activities under the Kyoto Protocol are estimated using area based sampling by circular sampling plots (the spatial assessment unit). At the first inventory, the whole plot represents the activity Forest management (FM) but at the re-inventory the plot represent the activities FM and Deforestation (D), respectively. Observe that both land use categories and activities have definitions but Sweden has no minimum area limit set for estimating activities.

11.1.2 Elected activities under Article 3, paragraph 4, of the Kyoto Protocol

For the accounting of LULUCF-activities under article 3.4 during the first commitment period, Sweden has elected Forest management (FM). FM is defined as activities on Forest land.

11.1.3 Description of how the definitions of each activity under Article 3.3 and each elected activity under Article 3.4 have been implemented and applied consistently over time

Sweden defines Deforestation (D) as land use conversions from Forest land (all forest land area is regarded managed) to Cropland, Grasslands or Settlements (all land under these three categories are assumed managed). Afforestation/ Reforestation (AR) is defined as land use conversions in the opposite direction (Figure 11.2). Land use categories are strictly defined (see NIR chapter 7.2.3) and land use conversions are observed in field using a five-year inventory cycle. The approximately 30,000 permanent sample plots were laid out between 1983 and 1987 and have thereafter been re-inventoried in a consistent way (Figure 11.3). If the land use of a sample plot or part of a sample plot is assessed as converted between consecutive inventories the exact year of the conversion is estimated from properties on the sample plot (site, stand and vegetation properties). This method has been used since 2006. For the years 1990-2006, the conversion year between consecutive inventories is randomly assumed. AR on former Cropland, Grasslands and Settlements are connected with an active human decision. Normally regeneration is following shortly after the land conversion. All AR land is by national legislation considered as Forest land and the same definition of Forest land is used in the Forestry act

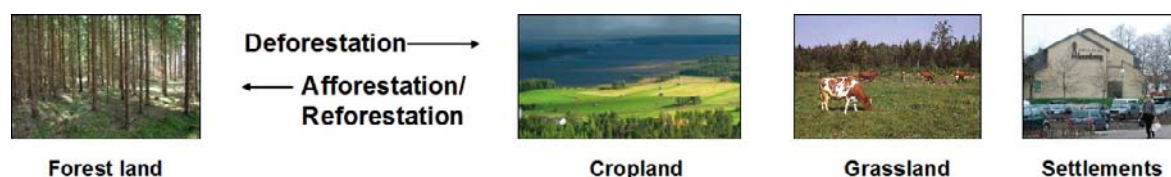


Figure 11.2 D is defined as land use conversions from Forest land (managed) to another managed land use class (all Cropland, Grasslands and Settlements are assumed managed). AR are defined as land use conversions in the opposite direction (C, G or S to F).

(1979:429 2 § 1.) as for the UNFCCC reporting. The activity Forest management (FM) is assumed occurring on all land fulfilling the forest definition (see 11.1.1). If land is subject to AR (or D), this land may have secondary classification FM. Land could only be reported under one activity or none (to avoid double counting).

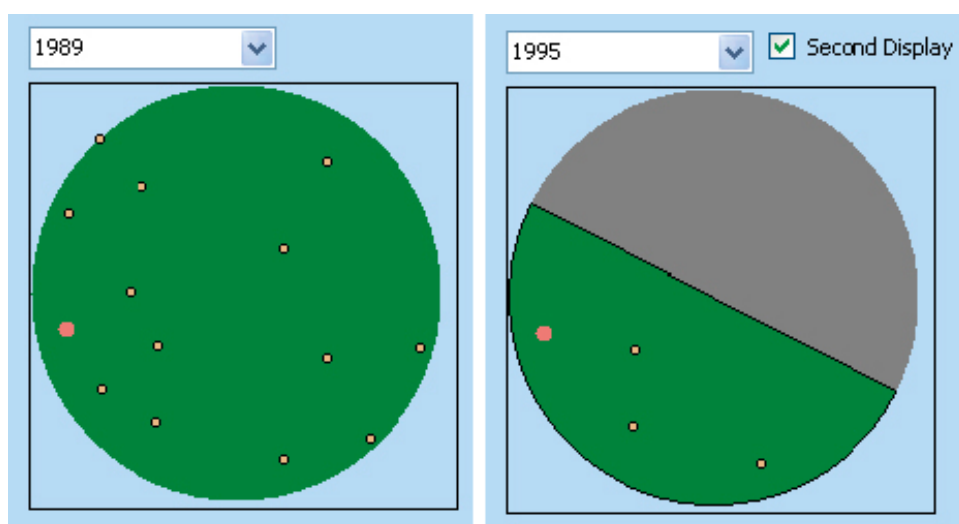


Figure 11.3 The figure shows data for a specific sample plot in the Swedish LULUCF-database. The individual tree biomass on approximately 30,000 permanent sample plots are matched to land use and traced back to before the base year in a consistent way. Applying area based sampling all 30,000 permanent sample plots represents the whole land and fresh water area in Sweden and carbon stock changes are estimated using the stock change method on these plots. Part of this specific plot was Deforested between 1991 and 1992. The positioning of trees is central when matching carbon stock changes in living biomass to activity (about 75% of Deforested plots are divided into more than one land use category). The position and biomass of the marked tree (right panel) is identified at both inventories and demonstrates the possibility to match individual trees to activities over time.

11.1.4 Descriptions of precedence conditions and/or hierarchy among Article 3.4 activities, and how they have been consistently applied in determining how land was classified.

Precedence conditions are: D, AR and FM since FM is the only activity elected under article 3.4. Land under Deforestation (D) cannot leave this category. This is basically also valid for land under AR – except after D. Land under Afforestation/ Reforestation (AR) usually has secondary classification FM (always if reported under Forest land remaining Forest land or conversion to Forest land under the UNFCCC). Theoretically, land under Deforestation can have secondary classification FM (if reported under Forest land remaining Forest land or conversion to Forest land under the UNFCCC) but this far such land does not exist. Land areas under FM that are naturally degraded can leave the category until 2008 (but not during the commitment period) and is not reported under the KP (usually reported as Forest land converted to Wetland or Other land under the UNFCCC). Deforested land can leave FM at any time from 1990 and onwards and these land areas are then reported under D. Land associated with the activities reported (AR, D and FM) is accumulated from end of 1989 and onwards, and changes in carbon pools and other emissions on these land areas are reported during the commitment period.

11.2 Land-related information

11.2.1 Spatial assessment unit used for determining the area of the units of land under Article 3.3

The “Spatial assessment unit”, which is the same as for the UNFCCC-reporting, is used to determine the area of accounting for ARD. The “Spatial assessment unit” is defined as the minimum area used to detect a land use conversion.

Sweden monitors land use transfers based on field measurements using circular sample plots (radius 10 m). If any part of a plot is converted from one land use category to another, it can be detected. Thus, the “Spatial assessment unit” will be a sample plot part and activities down to an area of 0 m² could be detected. The same “Spatial assessment unit” has consistently been used in both the UNFCCC and the KP-reporting (Figure 11.1).

11.2.2 Methodology used to develop the land use matrix

Data from the Swedish National Forest Inventory (NFI) have been used for developing the land use matrix. The underlying data are consistent with the data used for developing the land use matrix under the UNFCCC-reporting. The main difference is that activities are reported under the KP while land use categories are reported under the UNFCCC.

The Swedish National Forest Inventory covers all land and fresh water areas before the base year and onwards on sample plots with a fixed position (permanent sample plots). This makes it possible to consistently trace both gross and net land use transfers over time.

11.2.3 Maps and/or database to identify the geographical locations, and the system of identification codes for the geographical locations

Sweden uses a geographical boundary encompassing units of land (Reporting method 1) and has adopted approach 3 (Table 4.2.2 in GPG LULUCF 2003) for reporting emissions/ removals under article 3 of the KP (Figure 11.4). In practice a sample frame of approximately 30,000 permanent sample plots is covering all relevant managed land in Sweden (see chapter 7). The sample frame is divided into about 30 strata and the distance between sample units within stratum is based on autocorrelation. A five-year inventory cycle is used and each year about 6,000 sample plots are inventoried over the whole country. Each sample plot has an identification code and a registered geographical position. This information is confidential due to sampling reasons. However, on request (i.e. in connection with an in country review) it is possible to visit any plot. A certain year, each sample plot (or a part of a sample plot) could only represent one activity (D, AR or FM) or none. The status of activities on sample plots could be traced back from the current year to the base year (1990; Figure 11.4).

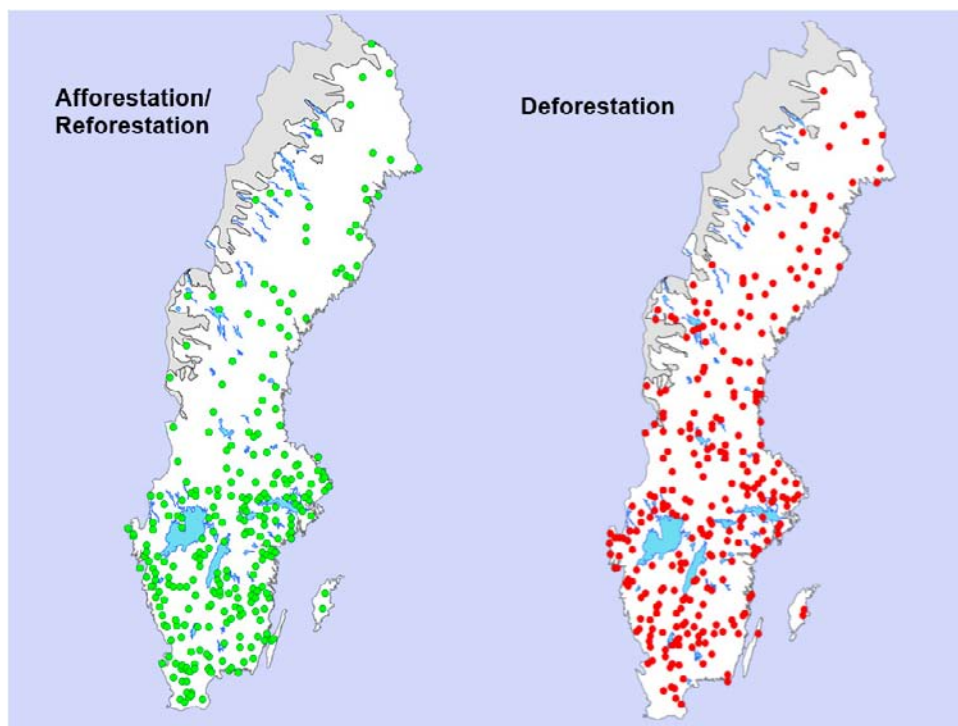


Figure 11.4 The location of sample plots partly or completely reported under ARD in Sweden (1990-2011).

11.3 Activity-specific information

11.3.1 Methods for carbon stock change and GHG emission and removal estimates

In most cases, methodologies, models and assumptions under the KP-reporting are consistent with the UNFCCC-reporting. This chapter focuses on discrepancies.

11.3.1.1 DESCRIPTION OF THE METHODOLOGIES AND THE UNDERLYING ASSUMPTIONS USED

11.3.1.1.1 *Carbon pools*

The living biomass pool changes is estimated in exactly the same way as under the UNFCCC reporting using the stock change method and area based sampling (See 7.3.1.2 + NIR Annex 3:2). However, the living biomass is reported separately for above-ground and below-ground biomass, respectively.

The dead wood, litter and soil organic pools are calculated using the same methods as for the UNFCCC-reporting (See 7.3.1.3-4 + Annex 3:2) using the area distribution associated with the reported activities under the Kyoto protocol (ARD and FM).

All methods used for FM is Tier 3 whereas methods for Litter, Dead wood and Soil organic carbon for ARD is Tier 2.

11.3.1.1.2 *Other emissions*

Emissions associated with direct N₂O emissions from N fertilization in forests (TABLE 5(KP-II)1) are estimated in the same way as under the UNFCCC (see 7.2.5.1). The estimates are based on activity data and emission factors with no information of the actual geographical distribution of fertilizer used. The fertilization is strictly regulated by the Forestry act and Sweden assumes that no fertilizer is applied in young forests. Therefore all emissions are assumed to occur under the activity FM and none under AR and the reported figure under “Forest Land remaining Forest Land” (UNFCCC, TABLE 5(I)) should correspond to the reported figure under FM (TABLE 5(KP-II)1)). It should be noted that fertilization is very restricted in Sweden. The annual fertilized area is expected to increase in the coming years but to cover less than 0.5 % of the total area of Forest land.

In line with the UNFCCC-reporting (TABLE 5(II)), N₂O emissions from drainage of soils (TABLE 5(KP-II)2) are not reported. The justification for omitting the emission is found in section 11.3.1.2.

The reporting of N₂O emissions from disturbance associated with land-use conversion to Cropland (TABLE 5(KP-II)3) are only relevant for the activity D and is reported. The reported figure should be similar to the figure reported under “2.1 Forest land converted to Cropland” (UNFCCC, TABLE 5(III) and discrepancies arise only from a different accumulations of land between the two reporting's (see activities).

All forest fires (TABLE 5(KP-II)5) are reported under FM and this figure should correspond to the figure reported under Forest land (UNFCCC, TABLE 5(V)). Forest fires may occur in all kinds of forests but no fires have been registered by the National Forest Inventory on land reported under activities AR. As mentioned under chapter 11.1, only 0.21% of area of Cropland is D and liming does probably only occur on Cropland remaining Cropland. Thus no emissions from liming are reported under the KP (this far).

11.3.1.1.3 Activities

Kyoto Protocol activity areas are accumulated from the base year and onwards and, normally, do not leave the class. For the UNFCCC-reporting converted land stays in the conversion class for twenty years and is thereafter reported under the land use category it was converted to. The twenty-year accumulation of land under the UNFCCC-reporting may begin long before the base year and is therefore not suitable to, for example, compare D under the Kyoto Protocol with Forest land converted to Cropland, Grassland or Settlements. Using conversions from Forest land as a “proxy” for D has led to several misunderstandings when assessing the outcome of the UNFCCC and the KP reporting.

11.3.1.2 JUSTIFICATION WHEN OMITTING ANY CARBON POOL OR GHG EMISSIONS/REMOVALS FROM ACTIVITIES UNDER ARTICLE 3.3 AND ELECTED ACTIVITIES UNDER ARTICLE 3.4

Sweden accounts for all carbon pools (above-ground biomass, below-ground biomass, litter, dead wood and soil organic carbon). This is also valid for all non-carbon pool emissions except nitrous-oxide emissions from drainage of soils under FM (Table 5(KP-II)2). These emissions are optional to report since the available methods to estimate the emissions are not accurate enough (IPCC 2003 GPG, Appendix 3a.2).

11.3.1.3 INFORMATION ON WHETHER OR NOT INDIRECT AND NATURAL GHG EMISSIONS AND REMOVALS HAVE BEEN FACTORED OUT

Sweden argues that the issue of “factoring out” was solved during negotiations with the cap for FM. A footnote of par. 7 of decision 16/CMP.1 “recognizes that the intent of the appendix to the annex to decision 16/CMP.1 is to factor out the effects described in paragraph 7 (a)–(c) of these guidelines for the first commitment period”). So Sweden has indirectly “factored out” 7 (a)–(c) by the cap for FM but no direct “factoring out” has been made. Moreover, sound science for a direct “factoring out” does not exist and no methodology has been adopted.

11.3.1.4 CHANGES IN DATA AND METHODS SINCE PREVIOUS SUBMISSIONS (RECALCULATIONS)

The uncertainty of estimates increases by decreasing number of sample plots and Table 11.3 illustrate the need of annual recalculations of the most recent years to increase the accuracy. In the current submission, the living biomass pool and activity areas have been recalculated for the most recent years to improve accuracy and each estimate are now based on 6,000 more measured sample plots. To avoid an increasing uncertainty of estimates by decreasing number of sample plots Sweden has now introduced extrapolation for inventory cycles without a full record of sample plots until 2011. Inconsistencies in the reporting of direct N₂O emissions for N fertilization and N₂O emissions from disturbance associated with land-use conversion to cropland between the UNFCCC-reporting and the KP-LULUCF reporting have been taken care of.²⁷⁰

Based on 30,000 sample plots the accuracy of estimates of carbon stock changes in living biomass for ARD activities are certain in absolute but uncertain in relative terms. The estimated accuracy (Standard Error) for AR and D is around 0.1 and 0.3 Mton CO₂ per year, respectively. However, since ARD is quite uncommon in Sweden and quite close to zero, the relative error might be large. The corresponding estimated accuracy for FM is 3 Mton CO₂ per year.

The uncertainty will most probably not influence the accounting since the net removal from FM is much larger than the net emission from ARD, thus it will be possible to offset net emissions from ARD by the net removal from FM and further claim credits from FM limited by the cap (2.13 Mtons CO₂ per year).

²⁷⁰ The inconsistency now taken care of was observed by reviewers (ARR 2011)

Table 11.3 The accumulated area under activities AR, D and FM and the approximate number of sample plots each estimate is based on (partly extrapolated plots inside brackets), presented by submission. Last three rows express the difference between submissions due to recalculations.

Submission	1990	.	.	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011
AR [M ha]	0.02	.	.	0.13	0.14	0.15	0.17	0.19	0.21	0.22	0.23	0.23	0.24	-
2012 D [M ha]	0.01	.	.	0.15	0.16	0.16	0.17	0.18	0.19	0.19	0.20	0.21	0.21	-
FM [M ha]	28.23	.	.	28.10	28.09	28.09	28.09	28.10	28.13	28.18	28.24	28.28	28.31	-
No. of plots (10 ³)	30	.	.	30	30	30	30	30	30	(30)	(30)	(30)	(30)	-
AR [M ha]	0.02	.	.	0.13	0.14	0.15	0.17	0.19	0.21	0.22	0.24	0.25	0.26	0.26
2013 D [M ha]	0.01	.	.	0.14	0.15	0.16	0.16	0.17	0.18	0.19	0.20	0.21	0.22	0.22
FM [M ha]	28.20	.	.	28.08	28.07	28.07	28.07	28.08	28.11	28.15	28.18	28.18	28.17	28.17
No. of plots (10 ³)	30	.	.	30	30	30	30	30	30	30	(30)	(30)	(30)	(30)
Difference between Submission 2013 and 2012														
AR [M ha]	0.00	.	.	0.00	0.00	0.00	0.00	0.00	0.00	0.00	-0.01	-0.02	-0.02	-
D [M ha]	0.00	.	.	0.01	0.01	0.00	0.01	0.01	0.01	0.00	0.00	0.00	-0.01	-
FM [M ha]	0.03	.	.	0.02	0.02	0.02	0.02	0.02	0.02	0.03	0.06	0.10	0.14	-

Table 11.4 Emissions / removals (minus), [Mton CO₂] from reported carbon pools in AR, D and FM per submission

Activity	Carbon pool	2008 Subm. 2012	2009 Submi. 2012	2010 Submi. 2012	2008 Subm. 2013	2009 Subm. 2013	2010 Subm. 2013	2011 Subm. 2013
AR	Above ground biomass	-0.54	-0.53	-0.53	-0.58	-0.60	-0.60	-0.60
	Below ground biomass	-0.18	-0.18	-0.18	-0.19	-0.20	-0.20	-0.20
	Dead wood	-0.02	-0.02	-0.01	-0.02	-0.02	-0.01	-0.01
	Litter	-0.25	-0.26	-0.26	-0.26	-0.28	-0.28	-0.27
	Soil organic carbon	0.17	0.17	0.18	0.17	0.18	0.19	0.17
	Total	-0.83	-0.82	<u>-0.80</u>	-0.88	-0.92	<u>-0.90</u>	-0.90
D	Above ground biomass	1.03	0.86	0.60	0.84	0.63	0.40	0.48
	Below ground biomass	0.35	0.28	0.20	0.29	0.20	0.14	0.16
	Dead wood	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	Litter	0.92	0.95	0.98	0.92	0.96	0.99	0.95
	Soil organic carbon	0.94	0.96	0.99	0.94	0.98	1.00	0.96
	Total	3.24	3.05	<u>2.76</u>	2.99	2.77	<u>2.53</u>	2.55
FM	Above ground biomass	-21.7	-19.8	-21.4	-20.3	-20.4	-20.9	-23.9
	Below ground biomass	-7.57	-8.51	-7.24	-6.91	-6.99	-7.01	-8.10
	Dead wood	-9.15	-8.75	-6.92	-9.48	-9.06	-7.02	-6.87
	Litter	2.88	2.90	3.13	5.38	5.40	5.66	5.67
	Soil organic carbon	-4.55	-4.56	-4.57	-4.46	-4.45	-4.45	-4.44
	Total	-40.1	-38.7	<u>-37.0</u>	-35.8	-35.5	-33.7	-37.6

11.3.1.5 UNCERTAINTY ESTIMATES

Estimates of carbon stock changes are based on the same underlying data as the reporting under the UNFCCC. These estimates originate mainly from a sampling design with the intention to keep systematic errors as low as possible. The systematic error is reduced by using representative functions, by direct measurements in field and at laboratory. We assume that the major source of uncertainty arise from random variation due to sampling. The sampling error is estimated using statistical theory for living biomass and partly for other carbon pools (all Tier 3). A consistent methodology for estimating carbon pools has been used from 1990 and onwards. Therefore, we expect the uncertainty to be the same for all years where all sample units are used to estimate the annual change. The uncertainties for other categories are assumed by expert judgment. For the first time we can provide separate formal estimates of uncertainty of AR and D, respectively. The estimated uncertainty was larger than expected for D, while the uncertainty for AR was as expected.

Based on 30,000 sample plots the accuracy of estimates of carbon stock changes for ARD activities are certain in absolute but uncertain in relative terms. The estimated accuracy (Standard Error) for living biomass for AR and D is around 0.1 and 0.3 Mton CO₂ per year, respectively. This is valid when estimates are based on all 30,000 sample plots. However, when based on one year sample (6,000 plots), the estimated accuracy (Standard Error) for living biomass for AR and D is around 0.2 and 0.7 Mton CO₂ per year, respectively. To increase the accuracy Sweden uses extrapolated data for the most recent

years. Since ARD is quite uncommon in Sweden and quite close to zero the relative error might be large. The corresponding estimated accuracy for FM is 3 Mton CO₂ per year (when based on 6,000 sample plots, 7 Mton CO₂ per year). Three Mton CO₂ per year should be compared with a total stock of more than 4,000 Mton CO₂ (relative error 0.07 %). For other carbon pools than living biomass, the uncertainty is based on assumptions²⁷¹.

Table 11.5 Estimated and assumed uncertainty for KP-activities. (Uncertainty=2•relative “standard error”).

Activity	Category	2•Relative Standard Error [%]		
		CO ₂	N ₂ O	CH ₄
FM	Living biomass	30	-	-
	Dead organic matter	50	-	-
	Soil organic carbon	35	-	-
	Direct N fertilization, 5 (I)	-	50	-
	Biomass burning, 5 (V)	-	75	75
AR	Living biomass	29	-	-
	Dead organic matter	70	-	-
	Soil organic carbon	35	-	-
D	Living biomass	82	-	-
	Dead organic matter	70	-	-
	Soil organic carbon	35	-	-
	Conversion Cropland, 5 (III)	-	100	-

11.3.1.6 INFORMATION ON OTHER METHODOLOGICAL ISSUES

There are currently no methods identified that needs further clarification than those already explained.

11.3.1.7 THE YEAR OF THE ONSET OF AN ACTIVITY, IF AFTER 2008

The onset of activities follows IPCC GPG for LULUCF (IPCC²⁷²) and no activity has been set on after 2008.

²⁷¹ This section is an amendment due to a request from reviews (ARR 2011) to improve the information on uncertainties in estimates of ARD.

²⁷² Intergovernmental Panel on Climate Change, 2003

11.4 Article 3.3

11.4.1 Information that demonstrates that activities under Article 3.3 began on or after 1 January 1990 and before 31 December 2012 and are direct human-induced

Sweden defines D as land use conversions from Forest land (all managed) to Cropland, Grasslands or Settlements (all land under these three categories are assumed managed). AR is defined as land use conversions in the opposite direction (Figure 11.2). Land use categories are strictly defined (see NIR chapter 7.2.3) and land use conversions are confirmed in field at consecutive inventories. The estimates are based on area sampling using the approximately 30,000 permanent sample (see chapter 7 for further details on the NFI). The inventory has been consistent since 1983.

This implies that Sweden uses the broad interpretation of “direct human induced” and an active human removal of trees followed by a land use conversion from Forest land to a managed non-forest land use category is considered direct human induced deforestation. This is also valid for the choice to actively abandon managed land in favour for the management of forests (afforestation, reforestation). The management of Forest land on abandoned former managed non-forest land is regulated by the Forestry act (1979:429). The intention of a human induced land use conversion should be permanent. If, for example, a land owner decides to convert former Cropland to Forest land by planting trees, this action is considered AR, but if the land owner in the future decides to cultivate this land back to Cropland, then the land will be reported under D. No such reversed-conversions have been identified (this far).

The NFI is used to confirm that activities under Article 3.3 began on or after 1 January 1990 and before 31 December 2012 and are direct human-induced. If the land use of a sample plot or part of a sample plot is considered converted between consecutive inventories the exact year of conversion is estimated from properties on the sample plot (site, stand and vegetation properties). This is valid from 2006. Until 2006, the conversion year between consecutive inventories is randomly distributed.

11.4.2 Information on how harvesting or forest disturbance that is followed by the re-establishment of forest is distinguished from deforestation

Final felling is a natural step in the rotation cycle of forestry. Also storms may result in large areas of felled trees (wind-throws). If final felling or disturbances as storms have been identified between two consecutive inventories this is not enough to classify the plot as D. However, if for instance a new road, a power line or other land use preceding the definition of forest is located on the former Forest land, then the plot is considered D. The emission from “loss of biomass” is matched to the conversion year. If final felling has occurred on a plot between two consecutive inventories with no sign of D, but D is confirmed at the next re-inventory, then the year of D is “re-calculated” to match the “loss of biomass” to the conversion year.

11.4.3 Information on the size and geographical location of forest areas that have lost forest cover but which are not yet classified as deforested

About 200,000 ha²⁷³ Forest land is annually losing its forest cover as a natural step in the forest rotation cycle. The position and status of every sample plot that has lost forest cover is known but D is not reported until confirmed (see 11.4.2). The geographical position and area of all final fellings on Forest land are monitored by detection of changes using the remote sensing system ENFORMA²⁷⁴. Each land owner has to apply to the Forestry Agency before harvest and in that regard, state whether the removal of trees is a natural step in the forest rotation cycle or harvest followed by a permanent land use change (D).

11.5 Article 3.4

11.5.1 Information that demonstrates that activities under Article 3.4 have occurred since 1 January 1990 and are human induced

The activity FM is assumed to occur on all Forest land and land areas with first classification FM is reported under Article 3.4 FM. Land reported under AR usually has secondary classification FM. Land under activity FM is accumulated from 1990 but could leave this category for D at any time. Before 2008 (but not during the commitment period), land under FM could leave this category by natural degradation to Wetland or Other land and is then not reported at all. This IPCC-rule has no practical significance for reported removals/ emissions. The area under FM is quite stable and all land use categories, including Forest land, are consistently monitored in field since 1983. Therefore it is possible to trace back all land use categories and land use conversions to at least 1990. "Human induced" is assumed equivalent with "managed" and all Forest land is assumed managed. Most forest biomass is actively managed for timber and pulp production and remaining forest biomass is managed for nature conservation. The definition used coincides with definition of Forest land according to the Forestry act (1979:429).

11.5.2 Information relating to Cropland Management, Grazing Land Management, and Revegetation, if elected, for the base year

Sweden has not elected these activities.

²⁷³ Swedish University of Agricultural Sciences, 2010

²⁷⁴ Olsson et al, 2005

11.5.3 Information relating to Forest Management

The net removal from living biomass is important for the total net removal reported under FM. The net removal from living biomass is the result from growth and drain and is sensitive to the demand of forest products from the forest industry.

11.6 Other information

11.6.1 Key category analysis for Article 3.3 activities and any elected activities under Article 3.4

A qualitative key category analysis has been made (IPCC GPG for LULUCF (IPCC²⁷⁵), p 5.38-5.40). Carbon dioxide emissions/ removals from Activities Forest management, Afforestation/ Reforestation and Deforestation were considered key-categories (CO₂). Every key-category is estimated using Tier 2 and 3. Emissions from non-carbon pools are very restricted under the KP. These are not considered as key categories and reported using Tier 1. Under the UNFCCC, part of 5III N₂O emissions from land use conversions to Cropland corresponds to D from Forest to Cropland. However, this category was not identified as key-category because most of the emissions under the UNFCCC refer to land use conversions from Grasslands to Cropland and due to a “conservative” high assumed uncertainty (100%). Sweden will follow up this emission carefully in the future (Tier 1) and we will compare the probably overestimated uncertainty with neighbouring countries using similar methodology. Sweden base the key-category analysis on a higher disaggregation level than suggested (GPG 2003, Table 5.4.4), because the key-category analysis is made per land use and gas (as suggested by reviewers). Since Sweden uses higher tier for all key-categories under the KP, Sweden has no immediately plan to improve the inventory (LULUCF). However, the key-category analysis is useful for monitoring potential changes in emission/removal trends.

11.7 Information relating to Article 6

Information relating to Article 6 is provided in Annex 6:1 and 6:3.

²⁷⁵ Intergovernmental Panel on Climate Change, 2003

12 Information on accounting of Kyoto units

12.1 Background information

Each Party must include information on its aggregate holdings and transactions of Kyoto Protocol units in its annual report. The reporting will be submitted according to the special report standard, the Standard Electronic Format (SEF) with the annual inventory on 15 April. Sweden began the annual reporting in 2009.

Sweden's Standard Electronic Format report for 2012 containing the information required in paragraph 11 of the annex to decision 15/CMP.1 The SEF will be submitted to the UNFCCC Secretariat electronically.

The ETS operates in 30 countries: the 27 EU Member States plus Iceland, Liechtenstein and Norway. It covers CO₂ emissions from installations such as power stations, combustion plants, oil refineries and iron and steel works, as well as factories making cement, glass, lime, bricks, ceramics, pulp, paper and board.

The revised EU ETS Directive that was adopted in 2009 provides for the centralization of the EU ETS operations into a single European Union registry, operated by the Commission as well as the inclusion of the aviation sector into the system at the start of 2012. At the same time and with a view to increase efficiency in the operations of their National Registry, the EU Member States who are also Parties to the Kyoto Protocol (25) plus Iceland, Liechtenstein and Norway decided to operate their registries in a consolidated manner.

For this purpose, in 2012, the EU registry has undergone a major redevelopment with a view to comply with the new requirements of Commission Regulation 920/2010 and Commission Regulation 1193/2011 in addition to implementing the Consolidated System of EU registries (CSEUR).

Considering the importance of the change, the consolidated registry was subject to the formal recertification process by the UNFCCC, the process included the following steps:

Item	Date passed
Common readiness documentation review	15/12/2011
Specific readiness documentation	31/05/2012
Connectivity reliability test	30/05/2012
Distinctness test	9/12/2011
Interoperability test	30/05/2012 (the SEF report must still be tested)

A complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. This description notably includes:

- Readiness questionnaire
- Application logging plan
- Change management procedure
- Disaster recovery
- Manual Intervention
- Operational Plan
- Roles and responsibilities
- Security Plan
- Time Validation Plan
- Version change Management

The Consolidated System of EU registries that was certified on 1/6/2012 and which went to production on 20 June 2012 (called V3) constitutes the new baseline for reporting changes in the national registries of the Parties concerned. It is to be noted that the full activation included the migration of all data hosted by the national registries of the Member States as well as switching all connections to the UNFCCC ITL (one per MS) to the consolidated system. A new central service desk was also set up to support the registry administrators of the consolidated system. The new service desk acts as 2nd level of support to the local support provided by the Parties. It also plays a key communication role with the ITL Service Desk with regards notably to connectivity or reconciliation issues.

Referring to paragraph 22 of the annex to Decision 15/CMP.1, the following changes have occurred in the Consolidated System of EU registries since its recertification:

On 2 October 2012 a new software release (called V4) including functionalities enabling the auctioning of phase 3 and aviation allowances, a new EU ETS account type (trading account) and a trusted account list went into Production. The trusted account list adds to the set of security measures available in the single registry. This measure prevents any transfer from a holding account to an account that is not trusted. Those changes were all relating to EU ETS processes and had no impact on Kyoto Protocol processes.

A release V5 was being implemented to allow the allocation of EU allowances in phase III early 2013. Delivered early December 2012, release V5 will be activated in January 2013.

12.2 Summary of information reported in the SEF tables

Annual Submission Item	Party provided content
15/CMP.1 annex I.E paragraph 11: Standard electronic format (SEF)	Sweden's Standard Electronic Format report for 2012 will contain the information required in paragraph 11 of the annex to decision 15/CMP.1. The SEF will be submitted to the UNFCCC Secretariat electronically.

12.3 Discrepancies and notifications

Annual Submission Item	Party provided content
15/CMP.1 annex I.E paragraph 12: List of discrepant transactions	This section will be finalized in the S-IAR reporting to the UNFCCC in April 2013. Refer to Separate Electronic Attachment "SIAR Reports 2012-SE v 1.0.xls" Worksheet R2.
15/CMP.1 annex I.E paragraph 13 & 14: List of CDM notifications	No CDM notifications were received by the National Registry during the 2012 reporting period, pursuant of 15/CMP.1 annex I.E paragraphs 13 & 14. Refer to Separate Electronic Attachment "SIAR Reports 2012-SE v 1.0.xls" Worksheet R3.
15/CMP.1 annex I.E paragraph 15: List of non-replacements	No non-replacements occurred during the 2012 reporting period, pursuant of 15/CMP.1 annex I.E paragraph 15. Refer to Separate Electronic Attachment "SIAR Reports 2012-SE v 1.0.xls" Worksheet R4.
15/CMP.1 annex I.E paragraph 16: List of invalid units	No invalid units exist as at 31 December 2012, pursuant of 15/CMP.1 annex I.E paragraph 16. Refer to Separate Electronic Attachment "SIAR Reports 2012-SE v 1.0.xls" Worksheet R5.
15/CMP.1 annex I.E paragraph 17: Actions and changes to address discrepancies	The overall change to a Consolidated System of EU Registries also triggered changes to discrepancies procedures, as reflected in the updated manual intervention document and the operational plan . The complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries.

12.4 Publicly accessible information

Annual Submission Item	Party provided content
15/CMP.1 annex I.E Publicly accessible information	<p>The following information is now deemed publicly accessible and as such is available via the homepage of the SE registry and Swedish Energy Agency – http://www.energimyndigheten.se/en/Sustainability/EU-ETS---Implementation-in-Sweden/EUETS/Reports/</p> <p>In accordance with the requirements of Annex E to Decision 13/CMP.1, all required information for a Party with an active Kyoto registry is provided with the exceptions as outlined below.</p> <p>Account Information (Paragraph 45) and Account holders authorised to hold Kyoto units in their account (Paragraph 48)</p> <p>In light of the amendments introduced by Article 78 of the revised Registries Regulation that came into force in October 2010 and for security reasons, it is considered that the representative identification information as required in paragraph 45 and paragraph 48 is held as confidential.</p> <p>Since there are no provisions in Swedish law on which kyoto unit types legal entities are authorised to hold in the Swedish National Registry, It is difficult to provide a list of legal entities authorized to hold party holding accounts. All legal entities (person or organisation) authorized to participate in the Swedish national registry under the Kyoto mechanisms, must have a separate holding account for each legal entity according to the Data Exchange Standards (DES). The list of legal entities that currently have party holding accounts in the Swedish registry can be found through a report tool on the following public website:</p> <p>http://www.energimyndigheten.se/en/Sustainability/EU-ETS---Implementation-in-Sweden/EUETS/Reports/</p> <p>JI projects in Sweden (Paragraph 46)</p> <p>Two Article 6 (Joint Implementation) projects have been reported for conversion to ERU under an Article 6 project, and the conversion occurred in the specified period. The list of the conversion that occurred the Swedish registry can be found through a reports on the following public website:</p> <p>http://www.energimyndigheten.se/en/Sustainability/EU-ETS---Implementation-in-Sweden/EUETS/Reports/Kyoto-units-in-SUS/</p> <p>Holding and transaction information of units (Paragraph 47)</p> <p>Holding and transaction information is provided on a holding type level, due to more detailed information being declared confidential by EU Regulation.</p> <p>Article 10 of EU Regulation 2216/2004/EC, provides that “All information, including the holdings of all accounts and all transactions made, held in the registries and the Community independent transaction log shall be considered confidential for any purpose other than the implementation of the requirements of this Regulation, Directive 2003/87/EC or national law.”</p> <p><u>Paragraph 47c</u></p> <p>The total quantity of ERUs issued and converted on the basis of Article 6 projects (Joint Implementation), are displayed in the public accessible information on the web site.</p>
15/CMP.1 annex I.E Publicly accessible information	<p><u>Paragraph 47e</u></p> <p>Sweden does not perform LULUCF activities and therefore does not issue RMUs</p> <p><u>Paragraph 47g</u></p> <p>No ERUs, CERs, AAUs and RMUs have been cancelled on the basis of activities under Article 3, paragraphs 3 and 4 to date.</p> <p><u>Paragraph 47h</u></p> <p>No ERUs, CERs, AAUs and RMUs have been cancelled following determination by the Compliance Committee that the Party is not in compliance with its commitment under Article 3, paragraph 1 to date.</p> <p><u>Paragraph 47j</u></p> <p>No ERUs, CERs, AAUs and RMUs have been retired, other than retirements of CERs and AAUs as an outcome of the EU compliance retirement within EU Emission Trading Scheme (EU ETS) and not through retirement under article 3, paragraphs 3 and 4. Retired CERs and AAUs are presented in the SEF report.</p> <p><u>Paragraph 47k</u></p> <p>There is no previous commitment period to carry ERUs, CERs, and AAUs over from.</p>

12.5 Calculation of the commitment period reserve (CPR)

12.5.1 Assigned Amount

The assigned amount pursuant to Article 3, paragraphs 7 and 8, has been calculated in accordance with the annex to decision 13/CMP.1. Sweden's base year is 1990 and the Party has chosen 1995 as the base year for HFCs, PFCs and SF₆. Sweden's quantified emission limitation is 92 per cent as included in Annex B to the Kyoto Protocol. As Sweden is part of the European Community, whose Member States will meet their reduction commitment jointly in accordance with Article 4 of the Kyoto Protocol, Sweden's quantified emission limitation is 104 per cent. Sweden's assigned amount is calculated based on the Party's Article 4 commitment.

In response to inventory issues identified during the review of the Initial Report, Sweden submitted revised estimates of its base year inventory, which resulted in a recalculation of the assigned amount. Based on the revised estimates for Sweden's base year emissions – 72,152 Gg CO₂ eq. equal to 72,151,646 tonnes CO₂ eq. – the assigned amount is calculated to be 375,189 Gg CO₂ eq. equal to 375,188,559 tonnes CO₂ eq.

12.5.2 Commitment Period Reserve (CPR)

According to the annex to decision 11/CMP.1 (paragraph 6), "Each Party included in Annex I shall maintain, in its national registry, a commitment period reserve which should not drop below 90 per cent of the Party's assigned amount calculated pursuant to Article 3, paragraphs 7 and 8, of the Kyoto Protocol, or 100 per cent of five times its most recently reviewed inventory, whichever is lowest."

Sweden's original commitment period reserve was based on 90 per cent of assigned amount. However in the centralized review of submission 2009 the ERT pointed out that the option "100 per cent of five times its most recently reviewed inventory" gave a lower CPR. The CPR for submission 2009 was recalculated and since then the CPR has been calculated as the amount of 100 per cent of five times the national total not including LULUCF.

In submission 2013 the CPR is calculated from "100 per cent of five times its most recently reviewed inventory". The national total not including LULUCF (in 2011 reported in submission 2013) is 61,449 Gg CO₂ eq. equal to 66,448,940 tonnes CO₂ eq. The CPR is then 307,245 Gg CO₂ eq. equal to 307,244,703 tonnes CO₂ eq.

12.6 KP-LULUCF accounting

Sweden reports and accounts for activities under article 3.3 and the activity Forest management under article 3.4 of the Kyoto protocol. Detailed descriptions on definitions of activities and carbon pools as well as methods for the quantification of emissions and removals related to these activities can be found in chapter 11 of the NIR. For 2010 the activities under article 3.3 constituted a net source of 2.0 Mtons CO₂. Forest management under article 3.4 constituted a removal of 37 Mtons CO₂. After offsetting the article 3.3 source the remaining removal from Forest management constitutes 35 Mtons CO₂. However, final accounting quantity for 2010 is limited by the cap to 2.13 Mtons CO₂. It may be noted that Sweden has elected commitment period accounting. The referred figures represent only 2010.

13 Information on changes in national system

There have been no changes to the national system since last submission.

14 Information on changes in national registry

Directive 2009/29/EC adopted in 2009, provides for the centralization of the EU ETS operations into a single European Union registry operated by the European Commission as well as for the inclusion of the aviation sector. At the same time, and with a view to increasing efficiency in the operations of their respective national registries, the EU Member States who are also Parties to the Kyoto Protocol (25) plus Iceland, Liechtenstein and Norway decided to operate their registries in a consolidated manner in accordance with all relevant decisions applicable to the establishment of Party registries - in particular Decision 13/CMP.1 and decision 24/CP.8.

With a view to complying with the new requirements of Commission Regulation 920/2010 and Commission Regulation 1193/2011, in addition to implementing the platform shared by the consolidating Parties, the registry of EU has undergone a major re-development. The consolidated platform which implements the national registries in a consolidated manner (including the registry of EU) is called Consolidated System of EU registries (CSEUR) and was developed together with the new EU registry on the basis the following modalities:

- 1) Each Party retains its organization designated as its registry administrator to maintain the national registry of that Party and remains responsible for all the obligations of Parties that are to be fulfilled through registries;
- 2) Each Kyoto unit issued by the Parties in such a consolidated system is issued by one of the constituent Parties and continues to carry the Party of origin identifier in its unique serial number;
- 3) Each Party retains its own set of national accounts as required by paragraph 21 of the Annex to Decision 15/CMP.1. Each account within a national registry keeps a unique account number comprising the identifier of the Party and a unique number within the Party where the account is maintained;
- 4) Kyoto transactions continue to be forwarded to and checked by the UNFCCC Independent Transaction Log (ITL), which remains responsible for verifying the accuracy and validity of those transactions;
- 5) The transaction log and registries continue to reconcile their data with each other in order to ensure data consistency and facilitate the automated checks of the ITL;
- 6) The requirements of paragraphs 44 to 48 of the Annex to Decision 13/CMP.1 concerning making non-confidential information accessible to the public would be fulfilled by each Party individually;

- 7) All registries reside on a consolidated IT platform sharing the same infrastructure technologies. The chosen architecture implements modalities to ensure that the consolidated national registries are uniquely identifiable, protected and distinguishable from each other, notably:
- a) With regards to the data exchange, each national registry connects to the ITL directly and establishes a distinct and secure communication link through a consolidated communication channel (VPN tunnel);
 - b) The ITL remains responsible for authenticating the national registries and takes the full and final record of all transactions involving Kyoto units and other administrative processes such that those actions cannot be disputed or repudiated;
 - c) With regards to the data storage, the consolidated platform continues to guarantee that data is kept confidential and protected against unauthorized manipulation;
 - d) The data storage architecture also ensures that the data pertaining to a national registry are distinguishable and uniquely identifiable from the data pertaining to other consolidated national registries;
 - e) In addition, each consolidated national registry keeps a distinct user access entry point (URL) and a distinct set of authorisation and configuration rules.

Following the successful implementation of the CSEUR platform, the 28 national registries concerned were re-certified in June 2012 and switched over to their new national registry on 20 June 2012. During the go-live process, all relevant transaction and holdings data were migrated to the CSEUR platform and the individual connections to and from the ITL were re-established for each Party.

The following changes to the national registry of Sweden have therefore occurred in 2012, as a consequence of the transition to the CSEUR platform:

Reporting Item	Description
15/CMP.1 annex II.E paragraph 32.(a) Change of name or contact	No changes of name or contact regarding the RSA.
15/CMP.1 annex II.E paragraph 32.(b) Change regarding cooperation arrangement	<p>The EU Member States who are also Parties to the Kyoto Protocol (25) plus Iceland, Liechtenstein and Norway have decided to operate their registries in a consolidated manner. The Consolidated System of EU registries were certified on 1 June 2012 and went to production on 20 June 2012.</p> <p>A complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. This description includes:</p> <ul style="list-style-type: none"> • Readiness questionnaire • Application logging • Change management procedure • Disaster recovery • Manual Intervention • Operational Plan • Roles and responsibilities • Security Plan • Time Validation Plan • Version change Management <p>The documents above are provided as an appendix to this document.</p> <p>A new central service desk was also set up to support the registry administrators of the consolidated system. The new service desk acts as 2nd level of support to the local support provided by the Parties. It also plays a key communication role with the ITL Service Desk with regards notably to connectivity or reconciliation issues.</p>
15/CMP.1 annex II.E paragraph 32.(c) Change to database structure or the capacity of national registry	<p>In 2012, the EU registry has undergone a major redevelopment with a view to comply with the new requirements of Commission Regulation 920/2010 and Commission Regulation 1193/2011 in addition to implementing the Consolidated System of EU registries (CSEUR).</p> <p>The complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. The documentation is annexed to this submission.</p> <p>During certification, the consolidated registry was notably subject to connectivity testing, connectivity reliability testing, distinctness testing and interoperability testing to demonstrate capacity and conformance to the Data Exchange Standard (DES). All tests were executed successfully and lead to successful certification on 1 June 2012.</p>
15/CMP.1 annex II.E paragraph 32.(d) Change regarding conformance to technical standards	<p>The overall change to a Consolidated System of EU Registries triggered changes the registry software and required new conformance testing. The complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. The documentation is annexed to this submission.</p> <p>During certification, the consolidated registry was notably subject to connectivity testing, connectivity reliability testing, distinctness testing and interoperability testing to demonstrate capacity and conformance to the DES. All tests were executed successfully and lead to successful certification on 1 June 2012,</p>

Reporting Item	Description
15/CMP.1 annex II.E paragraph 32.(e) Change to discrepancies procedures	The overall change to a Consolidated System of EU Registries also triggered changes to discrepancies procedures, as reflected in the updated manual intervention document and the operational plan . The complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. The documentation is annexed to this submission.
15/CMP.1 annex II.E paragraph 32.(f) Change regarding security	The overall change to a Consolidated System of EU Registries also triggered changes to security, as reflected in the updated security plan . The complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. The documentation is annexed to this submission.
15/CMP.1 annex II.E paragraph 32.(g) Change to list of publicly available information	No major changes in publicly available information occurred during the reporting period. Only the URL to the publicly available information has been changes, http://www.energimyndigheten.se/en/Sustainability/EU-ETS---Implementation-in-Sweden/EUETS/Reports/ .
15/CMP.1 annex II.E paragraph 32.(h) Change of Internet address	The new internet address of the Swedish registry is: https://ets-registry.webgate.ec.europa.eu/euregistry/SE/index.xhtml
15/CMP.1 annex II.E paragraph 32.(i) Change regarding data integrity measures	The overall change to a Consolidated System of EU Registries also triggered changes to data integrity measures, as reflected in the updated disaster recovery plan . The complete description of the consolidated registry was provided in the common readiness documentation and specific readiness documentation for the national registry of EU and all consolidating national registries. The documentation is annexed to this submission.
15/CMP.1 annex II.E paragraph 32.(j) Change regarding test results	On 2 October 2012 a new software release (called V4) including functionalities enabling the auctioning of phase 3 and aviation allowances, a new EU ETS account type (trading account) and a trusted account list went into Production. The trusted account list adds to the set of security measures available in the CSEUR. This measure prevents any transfer from a holding account to an account that is not trusted.
Previous Annual Review recommendations for the National Registry	FCCC/ARR/2011/SWE
<i>ERT recommendation:</i>	No recommendations to address

15 Information on minimization of adverse impacts in accordance with Article 3, paragraph 14

The Swedish reporting of information regarding minimizations of adverse impacts in accordance with Article 3, paragraph 14 of the Kyoto Protocol is presented below. The outline follows that of CMP 15: § 23 and § 24.

Paragraph 23

Each Party included in Annex I shall provide information relating to how it is striving, under Article 3, paragraph 14, of the Kyoto Protocol, to implement its commitments mentioned in Article 3, paragraph 1, of the Kyoto Protocol in such a way as to minimize adverse social, environmental and economic impacts on developing country Parties, particularly those identified in Article 4, paragraphs 8 and 9, of the Convention

According to the provisions of Article 2 of the Kyoto Protocol, each party with quantified commitments under the Protocol is to introduce policies and measures to achieve the emission reductions to which it has made a commitment. The measures implemented are to be compatible with overarching objectives of sustainable development. Measures which would mean that all greenhouse gases regulated by the Protocol can decrease and cover all sectors of society are emphasised. The parties to the Kyoto Protocol are to aim to introduce policies and measures so that adverse effects are minimised. Such effects include adverse effects of a changed climate, effects on international trade and social, environmental and economic effects on other parties, particularly on developing countries.

In connection with the implementation of policies and measures in Sweden, an impact assessment is carried out, including an environmental impact assessment as a basis for decision-making. Such an analysis as far possible also includes assessing the risk of adverse effects in other countries. Formulation of proposals for changes of policy instruments is undertaken in a consultation procedure that makes it possible for operators concerned to give their comments on the proposals. In consultations that include suggestions for new rules or guidelines that may affect trade with other country shall be notified within the EU and to be alerted under the WTO's rules. This process makes it possible for other countries to influence the design of proposals for changed policy instruments and highlight any negative side effects that may arise.

The Swedish research activities, as indicated in Chapter 8 of Sweden's National Communication Report 5 (NC 5), among other things contribute to a sustainable global development. There are several examples of interdisciplinary research efforts focused on improving knowledge of effects globally (socially, economically and ecologically) of large-scale introduction of measures to reduce greenhouse gas emissions. Sweden's focus on increased use

of bioenergy, both through increased domestic production but also through increased imports in particular from developing countries, has meant that this area has been specially prioritised in systems-science research in the country.

Results from research have already influenced, and will in future influence, the development of policy. The special sustainability criteria devised for vehicle biofuels under the EU Renewables Directive is one such example.

Both positive and negative effects must be taken into account. Sweden contributes to a number of measures that may have positive effects on the prospects of developing countries adapting to climate change and implementing their own measures to reduce their greenhouse gas emissions. A description is given in Chapter 7 of NC5 of such efforts in the areas of technology transfer, knowledge building and support for adaptation measures. In addition, Sweden contributed multilateral aid in addition to our commitments by additional financial support to the special climate change Fund and least developed countries fund.

Finally Sweden wishes to emphasise that its climate strategy with its broad focus on many different types of measures covering the majority of sectors of society (both in and outside the country) and all greenhouse gases governed by the Kyoto Protocol has a form which fundamentally limits (minimises) the risk of adverse effects.

Paragraph 24 (a)

Annex II Parties shall incorporate information on how they give priority to the following actions:

(a) The progressive reduction or phasing out of market imperfections, fiscal incentives, tax and duty exemptions and subsidies in all greenhouse-gas-emitting sectors, taking into account the need for energy price reforms to reflect market prices and externalities

Sweden has to a large extent reformed the energy markets and phased out any market imperfections. The market price on electricity is deregulated and governed by the balance between demand and supply on a cross-border electricity market. In Sweden fossil fuels used outside the EU emissions trading scheme (ETS) is subject to a carbon dioxide tax to reflect the external cost. In EU ETS it is mainly the price of allowances that reflect the external effect of carbon dioxide emissions and the market failure.

Paragraph 24 (b)

Removing subsidies associated with the use of environmentally unsound and unsafe technologies.

Sweden does not extract oil, natural gas or coal, and therefore, has no subsidies on these fuels. With the introduction of the EU ETS for CO₂ emissions a cost have been imposed on environmentally harmful technologies such as fossil fuel based heat- and electricity production and industries.

Paragraph 24 (c)

Cooperating in the technological development of non-energy uses of fossil fuels, and supporting developing country Parties to this end

The chemical industry including refineries contributes to a fairly small share of the overall Swedish industrial production. This technological field is not a high priority in the Swedish research policy.

Paragraph 24 (d)

Cooperating in the development, diffusion, and transfer of less-greenhouse-gas-emitting advanced fossil-fuel technologies, and/or technologies, relating to fossil fuels, that capture and store greenhouse gases, and encouraging their wider use; and facilitating the participation of the least developed countries and other non-Annex I Parties in this effort

Sweden has an almost fossil free heat- and power production and therefore don't give priority to research and technology development in the field of advanced fossil based techniques for electricity and heat production technology. Since there is an automotive industry in Sweden, research programmes in the areas of hybrid technologies, automatic control systems for more energy-efficient internal combustion engines and the use of diesel oil for hydrogen production have been carried out over a long time period. The programmes are designed in particular to contribute to reduced fuel consumption for road vehicles. A development which is also of value for more fuel efficient passenger- and goods transport in non-Annex 1 countries, particularly those who are dependent on imports of oil, diesel and petrol.

Carbon Capture and Storage technology has in recent years been given priority in the Swedish research and climate policy and Sweden is keen on launching a demonstration project in this area. In the long term Sweden have the ambition to participate in the field of multilateral research collaborations

Paragraph 24 (e)

Strengthening the capacity of developing country Parties identified in Article 4, paragraphs 8 and 9, of the Convention for improving efficiency in upstream and downstream activities relating to fossil fuels, taking into consideration the need to improve the environmental efficiency of these activities

Sweden contribute to technology development in developing countries through development assistance and CDM projects, see chapter 7 of NC 5. The focus on transfer of technologies is primarily on energy efficiency technologies and on the introduction of renewable energy, but also to contribute to capacity-building. By providing knowledge about how CDM projects evolve, are administered and implemented for approval, which Sweden has made in African countries, the ability to *inter alia* obtain technology that enhances the efficiency of fossil fuel-intensive activities as well as other climate-related environmental technology projects improves.

Paragraph 24 (f)

Assisting developing country Parties which are highly dependent on the export and consumption of fossil fuels in diversifying their economies

Sweden assists developing countries which are dependent on imports for its fossil fuel consumption with the transfer of more energy-efficient technologies, renewable energy technologies and capacity-building which enhances diversification of the economy in these countries (see chapter 7 of NC 5). *Inter alia*, through;

- support for research programmes on renewable energy technologies coordinated by Asian Institute of technology,
- education on sustainable energy technology in partnership with universities in Uganda, Mozambique, Ethiopia and Tanzania,
- support for photovoltaic technologies for energy services to rural areas in Zambia.

In addition to development cooperation projects, Sweden is engaged in CDM projects in biomass based electricity generation, wind energy, biogas production, hydro-electric power production and energy efficiency projects which contribute to economic development and diversification of the economy in fossil fuel dependent developing countries. Capacity-building about how CDM projects evolve, are administered and implemented for approval, which Sweden has made in African countries, support a greater diversification of the economy in the countries concerned.

16 Other information

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16.2 Units and Abbreviations

t	1 (metric) tonne = 1 megagram (Mg) = 10^6 g
toe	tonne oil equivalent 1 toe = 41.87 GJ
Mg	1 megagram = 10^6 g = 1 tonne
Gg	1 gigagram = 10^9 g = 1 kilotonne (kt)
Tg	1 teragram = 10^{12} g = 1 megatonne (Mt)
TJ	1 terajoule
A	
AR	Afforestation and Reforestation
ARTEMIS	Assessment and Reliability of Transport Emission Models and Inventory Systems
AWMS	Animal Waste Management System
C	Carbon or Confidential
CH ₄	Methane
EMIR	Emissions database of the county administrative boards
ERT	Expert Review Team
CFCs	Freons
CKD	Cement kiln dust
CO	Carbon monoxide
CO ₂	Carbon dioxide
COP	Conference Of the Parties
CORINAIR	EMEP/CORINAIR Emission Inventory Guidebook
CRF	Common Reporting Format
D	Deforestation
DOM	Dead organic matter
SOC	Soil organic carbon
EC	Environmental Class
EAA	European Aluminium Association
EEA	European Environment Agency
EF	Emission Factors
EU	European Union
EMV	Emission Model for Road Traffic
ETS	Emission Trading Scheme
FAME	Fatty Acid Methyl Ester (earlier called RME)
F-gases	Fluorinated gases (HFCs, PFCs, SF ₆)
FM	Forest management
FMV	Swedish Defence Material Administration
FAO	Food and Agriculture Organisation of the UN
FOD model	IPCC First Order Decay model
FOI	Swedish Defence Research Agency
FORTV	Swedish Fortification Department
FRA	Forest Resource Assessment
FRA	National Defence Radio Institute

FTP	Federal Test Procedure
GHG	Greenhouse gases
Good Practice Guidance	IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories IPCC NGGIP
GWP	Global Warming Potential
Halocarbons	Organic compounds containing one or more halogens
HBEFA	Handbook Emission Factors for Road Transport
HWP	Harvested Wood products
HFCs	Hydrofluorocarbons
IE	Included Elsewhere
IEA	International Energy Agency
IEF	Implied Emission Factors
Industrial statistics	Industrial energy statistics
IPCC	Intergovernmental Panel on Climate Change
IPCC Guidelines	Revised 1996 Guidelines for National Greenhouse Gas Inventories
IPCC EFDB	IPCC Emission factor data base
ISIC	International Standard Industrial Classification of All Economic Activities
IVL	IVL Swedish Environmental Research Institute AB
Jernkontoret	Swedish Steel Producers' Association
KemI	The Swedish Chemicals Agency
KP	the Kyoto protocol
LPG	Liquefied Petroleum Gas
LTO	Landing and Take-Off
LUCF	Land-use change and forestry
LULUCF	Land-use, land-use change and forestry
MI	Markinventeringen (Swedish soil inventory)
MSW	Municipal solid waste
N ₂ O	Nitrous oxide
NAP	Swedish national allocation plan
NA	Not Applicable
NBF	National Board of Forestry
NCV	Net Calorific Value
NE	Not Estimated
NFI	National Forest Inventory
NIR	National Inventory Report
NMVOC	Non Methane Volatile Organic Compounds
NO	Not Occuring
NO _x	Nitrogen oxides
NSFSV	National Survey of Forest Soils and Vegetation
MTC	Motor Test Center
O ₃	Ozone
PA	Production approach
PAH	Polycyclic aromatic hydrocarbons

PDCA	Plan, Do, Check, Act
PFCs	Perfluorocarbons
QA/QC	Quality assurance and Quality control
Quarterly statistics	Quarterly fuel statistics
RIS	Riksinventeringen av skog (national forest inventory)
RME	Rapeseed Methyl Ester fuel
RVF	Swedish Association of Waste Management
SF ₆	Sulphur hexafluoride
SDC	Forest industry information association
SGU	Geological Survey of Sweden
SJV	Swedish Board of Agriculture
SLU	Swedish University of Agricultural Sciences
SMED	Swedish Environmental Emissions Data
SMHI	Swedish Meteorological and Hydrological Institute
STA	Swedish Transport Administration
STrA	Swedish Transport Agency
SO ₂	Sulphur dioxide
SPBI	Swedish Petroleum and Biofuel Institute
Swedish EPA	Swedish Environmental Protection Agency
TSP	Total amount of suspended particles
TPS	Technical Production System
UNFCCC	United Nations Convention on Climate Change
VBA	Visual Basic for Applications
VETO	Mechanistic model for simulations on road traffic
VTI	Swedish Road- and Transport Research Institute
WBCSD	World Business Council for Sustainable Development
WRI	World Resource Institute



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