Informative Inventory Report
Sweden 2018

Submitted under
the Convention on Long-Range Transboundary Air Pollution
# Contents

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EXECUTIVE SUMMARY

ES.1 Background information on the air pollutant emission inventory

Sweden has carried out inventories on air pollutants since the 1980's to meet the obligations of the United Nations Economic Commission for Europe Convention on Long-Range Transboundary Air Pollution (UNECE CLRTAP).

The inventory reports of air pollutants for the year 2018 is prepared in accordance with the 2013 Reporting Guidelines and according the CLRTAP’s revised Gothenburg Protocol as agreed by the parties to the Convention in Geneva, 2012. The inventory reports are submitted to the UNECE Secretariat and to the EEA annually.

This report constitutes Sweden’s IIR 2018 (inventory data 2016) for anthropogenic emissions of air pollutants: NOX, CO, NMVOC, SO2, NH3, TSP, PM10 and PM2.5, BC, heavy metals, dioxins, HCB, PCBs and PAH-4. The report contains information on Sweden’s inventories of air pollutants for all years from 1980 to 2016, including descriptions of methods, data sources, uncertainties, the quality assurance and quality control (QA/QC) activities carried out and a trend analysis. Data on estimated emissions and corresponding activity data are provided in NFR tables. Thermal values and emission factors are provided in Annex 2. The report also shows how Sweden follows the guidelines for estimating and reporting of emission data to ensure the transparency, consistency, accuracy, comparability and completeness of the reported emission data.

Emission estimates are mainly based on official Swedish statistics, e.g. energy statistics, agricultural statistics, environmental reports from industry and emission factors (nationally developed factors as well as internationally recommended ones). Sweden uses the Guidelines for Estimating and Reporting Emission Data for reporting to the Convention on Long-Range Transboundary Air Pollution (CLRTAP) and the EMEP/EEA Air Pollutant Emission Inventory Guidebook as methodological guidance.

Sweden also uses methodologies in accordance with the IPCC 2006 Guidelines for National Greenhouse Gas Inventories and methods that are in general in line with Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories IPCC-NGGIP (Good Practice Guidance). 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-2013. In a few cases, as indicated in IIR, methodologies from the EMEP/EEA Air Pollutant Emission Inventory Guidebook-2016 have been implemented.

2 The Good Practice Guidance can be found at: http://www.ipcc-nggip.iges.or.jp/public/gp/english/
Due to changes in the routines for handling data it was discovered that a higher degree of confidentiality is required for some activity data. This has affected some sub-sectors in the energy sector (NFR1A and 1B) which have been classified with the notation key Classified (C). Sweden works continuously on limiting the extent of confidentiality in inventory data.

**ES.2 Overview of source category emission estimates and trends**

The main sources of air pollutants have been divided into the following sectors: energy, industrial processes and product use, agriculture and waste. No air pollutant emissions have been estimated for the land use, land use change and forestry sector.

Emissions of pollutants regulated in the amended Gothenburg Protocol (SO₂, NOₓ, NH₃, NMVOCs, PM₁₀, PM₂.₅ and BC) have been reduced significantly since 1990. Other air pollutants, such as CO, poly aromatic hydrocarbons (PAH-4), dioxins, HCB and priority heavy metals cadmium namely, (Cd), mercury (Hg) and lead (Pb) have all been also reduced extensively since 1990.

The decline in emissions of the main air pollutants and heavy metals has been achieved through regulatory controls and application of better technologies in industry, energy, and transport sector. Examples include switching from higher sulphur fuels to lower sulphur fuels, phasing out the use of leaded gasoline, catalytic converters on vehicles and other instruments such as NOₓ-fee.

**Nitrogen oxides**

The estimated emissions of nitrogen oxides (NOₓ) were about 131 kt in 2016. The total emissions of NOₓ have declined by more than a half (53 %) since 1990 and by 3 % compared to the previous year.

The energy sector (NFR1) accounted for most of the NOₓ emission (81 %) of which transport was responsible for 40 % of the national total. The industrial processes and product use sector (NFR2) and the agriculture sector (NFR3) were responsible for about 9 and 10 %, respectively. NOₓ -emissions from the waste sector (NFR5) are very limited.

Emissions from the transport sector (NFR1A3) have declined by more than two thirds since 1990 and by about 7 % compared to 2015. Emissions from diesel passenger cars continued to increase (6 %) between 2015 and 2016, while emissions of NOₓ from heavy-duty vehicles declined by about 18 % compared to 2015. The decline in emissions from heavy duty vehicles is mainly due to the tightening of the EU road vehicle emission regulation standards.

Other important reasons for the general decline of NOₓ emission is the increased use of district heating and the introduction of NOₓ-fee in 1992 which have resulted in a reduction of emissions from the manufacturing industries and construction (NFR1A2) and the energy industries sectors (NFR1A1).
**Sulphur dioxide**

Emissions of sulphur dioxide (SO$_2$) decreased from 104 kt in 1990 to 19 kt in 2016, a reduction of 84%. However, the emissions in 2016 increased by 6% compared to 2015. About a half of the total SO$_2$ emission comes from the energy sector. The remaining emissions (51%) arise from the industrial processes and product use sector. SO$_2$-emission from the waste sector is very limited. Among the largest sources is of SO$_2$ emission are metal industry (2C), 23%, pulp and paper industry (2H1), 22%, manufacturing industries and construction (1A2), 20% and public electricity and heat production (1A1a), 17%. Transport (1A3) was a major source of SO$_2$ in early 90s but now is responsible for just 2% of the total emission. The general reduction in these sectors including the transport sector is mainly due to a transfer from fuels with high sulphur content to low-sulphur fuels.

**Ammonia**

The total emissions of ammonia (NH$_3$) amounted to 53 kt in 2016. Compared to emission levels in 1990, the emissions were 12% lower in 2015. The agriculture sector was the dominant source of NH$_3$ in 2016, accounting for 88% of total emissions. NH$_3$ is emitted from farm animals dung and urine and the use of inorganic fertilizers. The rest of the emission comes mainly from pulp and paper industry and transport, mainly from urea in vehicles filters that release HN$_3$. The main drivers for the reduced emission within the agriculture sector are a decline in number of animals, reduced use of inorganic fertilizers, measures and a more effective production.

**Non-methane volatile organic compounds (NMVOCs)**

In 2016, a total of 159 kt of NMVOCs were emitted in Sweden. Less than a half (47%) of the NMVOC emissions come from the industrial processes and product use sector (NFR2). The energy sector and the agriculture sector contributed with 34 and 19%, respectively. The remaining emissions arise from the waste sector.

The total emissions of NMVOCs have declined by 55% since 1990. The aggregated emission increased by 2% compared to 2015. The decline is sharp in the energy sector (mainly transport) and is clearly visible in the industrial processes and product use sector, amounting to about 74% and 30%, respectively compared to 1990. The main drivers for the sharp decrease in the energy sector in the last two decades are the increased energy efficiency in cars and the introduction of stricter emission standards in the EU-regulations for road vehicles.

**Carbon monoxide**

The aggregated emissions of carbon monoxide (CO) have decreased from about one million ton in 1990 to less than 430 kt in 2016, a decline of 60%. In 2016, the energy sector (NFR1) accounted for most of the CO emission (93%). The rest (7%) comes from the industrial processes and product use sector. CO-emission from the waste sector is very limited. Emissions of CO from transport sector decreased by 90% between 1990 and 2016. The introduction of catalytic converters in cars is the main driver for this reduction.
**Particles Matter (PM$_{10}$ and PM$_{2.5}$)**

In 2016, the total emission of PM$_{10}$ and PM$_{2.5}$ in Sweden were about 38 and 18 kt, respectively. Compared to 1990, the emissions have been reduced by 35 % and 51 %, respectively. The emission of PM$_{10}$ and PM$_{2.5}$ has increased by 2 % compared to previous year. Road transport is a large source for the emissions of PM. The amount of emissions depends on total traffic work and the use of studded tires. The main reason for the overall reductions in transport sector since 1990 is stricter European emission standards, resulting in lower emissions from heavy duty vehicles and busses. The combined emissions of PM from road abrasion and tires- and brake wear show an increasing trend. This is because the total volume of traffic has increased since 1990.

**Black Carbon**

Emissions of back carbon (BC) were about less than 3 kt in 2016. The emissions were about 40 % lower in 2016 than in 2000 and decreased by 2 % since 2015. The largest single source is other sectors (NFR1A4) accounting for 46 % of the total BC emissions. The transport sector (NFR1A3) accounted for about one-fourth of total BC emissions in 2016 of which road traffic is the major contributor of the emissions in the sector. Emissions from road transport have declined by 65 since 2000 due to stricter exhaust requirements.

**Poly Aromatic Hydrocarbons**

The total emissions of poly aromatic hydrocarbons (PAH-4) were 12.6 tons in 2016. The emissions have been reduced by 26 % since 1990. Compared to 2015 the emissions in 2016 have increased by 2 %. Wood combustion for residential heating is the dominant emission source for PAH, accounting for about two-third of the total emissions. The largest reduction of PAH-4 emissions comes from the metal industry (NFR2C), 81 %, compared to 1990, and has been achieved through the application of new technologies.

**Hexachlorobenzene**

The total emissions of hexachlorobenzene (HCB) were 2.7 kg in 2016. Emissions have decreased by 84 % since 1990. Compared to 2015 the emissions in 2016 have decreased by 30%. The largest single source of HCB emissions in 2016 was the electricity- and heat production (NFR1A1), 37 % of the total emissions. However, emissions from electricity- and heat production have doubled since 1990, mainly due to increased use of biomass as a fuel. The increased emission of HCB from iron and steel industry since 1990 is due to increased production volume. The largest increase of HCB in relative terms comes from the incineration and open burning in the waste sector, which has increased by 340 % times since 1990, due to increased combustion of hazardous waste.

**Priority Heavy Metals (Cd, Hg and Pb)**

In 2016, the total emissions of cadmium (Cd) in Sweden were 540 kg, a decrease of 77 % since 1990. The largest sources of Cd emissions in 2016 were electricity- and heat production and stationary combustion in the residential sector, accounting for about 30 % each. The industrial processes and product use was responsible for 13 % of the total emissions. Emissions from the industrial processes and product
use have decreased by more than 96% since 1990, mainly due to better technologies applied in the metal industry.

In 2016, the total emissions of mercury (Hg) in Sweden were about 412 kg, a decrease of about 73% since 1990. The largest single source of Hg emissions in 2016 was electricity- and heat production (NFR1A1) which was responsible for about one-third of the total emissions. Other large sources were metal industry (NFR2C) and manufacturing industries and construction (NFR1A2) accounting for 17% and 14%, respectively. The waste sector is also a significant contributor (8%) of the total emission. Emissions from industrial processes and product use have decreased by more than 85% since 1990, mainly due to better technologies applied in the metal industry.

Total emissions of lead (Pb) in Sweden were less than 12 ton in 2016 and have decreased by 97% since 1990. The largest single source of Pb emissions in 2016 was the metal industry (NFR2C) followed by the transport sector (NFR1A3) which accounted for about one-third and 29%, of the total emission, respectively. Emissions of Pb from the transport sector have decreased by more than 99% since 1990 due to the phasing out the use of leaded gasoline, while the decrease of emissions from the industrial processes and product use (about 95%) is mainly attributed to better technologies applied in the metal industry.
1 Introduction

Reporting of emission data to the Executive Body of the Convention on Long-range Trans-boundary Air Pollution (CLRTAP) is required in order to fulfil obligations regarding strategies and policies in compliance with the implementation of Protocols under the Convention. Parties should use the reporting procedures and are required to submit annual national emissions of SO₂, NOₓ, NMVOC, CO and NH₃, particulate matter, black carbon (BC), various heavy metals and POPs using the revised 2014 Guidelines for Estimating and Reporting Emission Data under the Convention on Long-range Transboundary Air Pollution⁵.

This report constitutes Sweden’s Informative Inventory Report (IIR) due by March 15 2018. The report contains information on Sweden’s inventories for all years from 1980 to 2016 including descriptions of methods, data sources, QA/QC activities carried out, and a trend analysis. The inventory accounts for anthropogenic emissions of SO₂, NOₓ, NH₃, NMVOC, CO, BC, TSP, PM₁₀, PM₂.₅, Pb, Cd, Hg, As, Cr, Cu, Ni, Se, Zn, PAH, HCB and dioxins.

Emission estimates are mainly based on official Swedish statistics, e.g. energy statistics, agricultural statistics, environmental reports from industry and emission factors (nationally developed factors as well as internationally recommended ones).

Sweden uses the revised 2014 Guidelines for Estimating and Reporting Emission Data for reporting to the Convention on Long-Range Transboundary Air Pollution (CLRTAP) and the 2013 EMEP/EEA Air Pollutant Emission Inventory Guidebook as methodological guidance⁶. Data are also reported under the EU National Emissions Ceiling Directive on emission of air pollutants to the European Commission. Sweden also uses methodologies in accordance with the IPCC 2006 Guidelines for National Greenhouse Gas Inventories⁷ and methods that are in general in line with Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories IPCC-NGGIP (Good Practice Guidance⁸). Some parts of the methodologies are taken directly from the IPCC Guidelines, the Good Practice Guidance and the 2013 EMEP/EEA Air Pollutant Emission Inventory Guidebook.

1.1 Institutional arrangements

The national system is 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.

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⁸ http://www.ipcc-nggip.iges.or.jp/public/gp/english/
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 fulfils all quality criteria, i.e. is transparent, accurate, consistent, comparable and complete.

The national system is, where applicable, used also for the reporting to the Convention on Long-Range Transboundary Air Pollution (CLRTAP) under the United Nations Economic Commission for Europe (UNECE) and under the EU National Emissions Ceiling Directive on emission of air pollutants to the European Commission.

### 1.1.1 Legal arrangements

The ordinance (2005:626) concerning climate reporting has been updated and enlarged to fulfil all the reporting requirements under the EU Monitoring Mechanism Regulation 525/2013/EC. The new ordinance 2014/1434 concerning climate reporting came into force and replaced the old ordinance the 29th of December 2014, and have been operational since the preparation of submission 2015.

The ordinance on climate reporting (OCR) describes the roles and responsibilities of the relevant government agencies in this area. The ordinance ensures that sufficient capacity is available for reporting. It also includes other improvements needed on the national level.

Supplemental to the new ordinance, formal agreements between the Swedish Environmental Protection Agency (Swedish EPA) and other concerned national agencies have been signed, listing in detail what is required regarding content and timetable from each agency.

Sweden also has legislation indirectly supporting climate reporting efforts by providing a basis for estimating greenhouse gas emissions and removals. Environmental reports are submitted under the Environmental Code (SFS 1998:808), and the Official Statistics Act (SFS 2001:99) imposes an obligation for large industries to submit annual data. In addition, government agencies in Sweden must comply by the Information and Secrecy Act (offentlighets- och sekretesslag) (SFS 2009:400).
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.

There is legislation in Sweden that indirectly supports the work by providing a basis for the estimation of air pollutants. 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 air pollutants 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 air pollutant inventory.

1.1.2 Institutional arrangements

Where applicable, the same institutional arrangements are used as for the Greenhouse gas inventory:

The illustration in Figure 1-1 and Table 1-1 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 Convention on Long-Range Transboundary Air Pollution (CLRTAP) under the United Nations Economic Commission for Europe (UNECE) and under the EU National Emissions Ceiling Directive on emission of air pollutants to the European Commission.

Depending on the role of the government agencies in 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 (in red). In addition to what is described in the OCR, the Swedish EPA engages the SMED consortium as consultants with expert skills to conduct the inventory and reporting in the area of air pollutants.
1.1.2.1 RESPONSIBILITIES OF THE SWEDISH ENVIRONMENTAL PROTECTION AGENCY

The Swedish Environmental Protection Agency (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 the Environment and Energy – submits the inventory to the NEC directive/EU and to the CLRTAP/UNECE. Finally, the Swedish EPA is responsible for national publication of the air pollutants inventory.

1.1.2.2 RESPONSIBILITIES OF NATIONAL AGENCIES

Table 1-1 below shows the responsibilities of the Swedish agencies according to the Ordinance concerning climate reporting.
Table 1. Responsibilities according to the ordinance concerning climate reporting

<table>
<thead>
<tr>
<th>Sector</th>
<th>Data and documentation provided by</th>
<th>Peer review conducted by</th>
</tr>
</thead>
<tbody>
<tr>
<td>Energy</td>
<td>Swedish Energy Agency, the Swedish Transport Administration, the Swedish Transport Agency, Transport Analysis, the Swedish Armed Forces.</td>
<td>Swedish Energy Agency (energy sector excluding transports) Transport Analysis (transports)</td>
</tr>
<tr>
<td>Industrial Processes and Product Use</td>
<td>Swedish Chemicals Agency, Medical Products Agency.</td>
<td>The Swedish EPA (CO₂, CH₄ and N₂O)</td>
</tr>
<tr>
<td>Agriculture</td>
<td>Swedish Board of Agriculture, Statistics Sweden (SCB).</td>
<td>The Swedish Board of Agriculture</td>
</tr>
<tr>
<td>Land Use, Land-Use Change and Forestry Sector</td>
<td>Swedish University of Agricultural Sciences (SLU), Statistics Sweden (SCB), the Swedish Forest Agency, the Swedish Meteorological and Hydrological Institute (SMHI), the Swedish Board of Agriculture, Swedish Civil Contingencies Agency (MSB), the Geological Survey of Sweden (SGU).</td>
<td>Swedish Forest Agency The Swedish Board of Agriculture (agriculture related parts)</td>
</tr>
<tr>
<td>Waste</td>
<td></td>
<td>The Swedish EPA</td>
</tr>
</tbody>
</table>

1.1.2.3 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)⁹, 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 covered the whole first commitment period under the Kyoto Protocol. During 2014 the contract with the consortium SMED was prolonged for another period (2015 – 2022). The structure of the consortium for the prolonged contract is a little bit different from since it is based on an agency agreement for the national agencies (SMHI, SCB and SLU) and a negotiated procurement of services under the terms of the Public Procurement Act for the IVL, this to be able to have the same setting for the consortium as during the former period.

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 SMHI is main responsible for production of gridded emission data. SCB 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 SLU is responsible for the LULUCF sector. The IVL is main responsible for the industrial process and product use sector and also parts of the waste sector and energy sector.

⁹ http://www.smed.se/
On behalf of the Swedish EPA, SMED also conducts specific projects necessary for improving the inventory.

1.2 Inventory planning, preparation and management

The present Swedish air pollutant inventory was compiled according to the recommendations for inventories set out in the revised 2014 Guidelines for Estimating and Reporting Emission Data for reporting to the Convention on Long-Range Transboundary Air Pollution (CLRTAP\textsuperscript{10}) and the 2013 EMEP/EEA Air Pollutant Emission Inventory Guidebook as methodological guidance\textsuperscript{11} and also the UNFCCC reporting guidelines in accordance with the Doha Amendment to the Kyoto Protocol (1/CMP.8), the IPCC 2006 Guidelines for National Greenhouse Gas Inventories, the IPCC Good Practice Guidance and Uncertainty Management in National Greenhouse Gas Inventories (IPCC GPG, 2000\textsuperscript{12}). Data are reported in the updated NFR format.

The inventory of air pollutants for reporting to the UNECE (CLRTAP) is integrated with the greenhouse gas inventory (for reporting to the UNFCCC and EU). This assures effective use of resources and consistency between the reporting to the UNFCCC and to the CLRTAP.

1.2.1 Quality system

In order to fulfil the obligations of reporting, 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.

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 plan\textsuperscript{13}. 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.


\textsuperscript{11} http://www.eea.europa.eu/publications/emep-eea-guidebook-2013

\textsuperscript{12} http://www.ipcc-nggip.iges.or.jp/public/gp/english/

\textsuperscript{13} Swedish EPA, National Greenhouse Gas and Air Pollutants Inventory System in Sweden
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 inventory work have to ensure that the methodologies applied in the reporting and inventories of emissions attain the quality required for it to be possible for Swedish air pollutant reporting to be done in the correct manner and with correct information.

The governments 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 SMED. The SMED quality system is described in a detailed manual including several appendices. 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 NFR category and documented procedures for uncertainty and key source analyses, as well as procedures for handling and responding to UNECE’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 for greenhouse gases which is largely applicable to the air pollutant inventory.

Figure 1-2. Structure of the quality system.
<table>
<thead>
<tr>
<th>Date</th>
<th>Event</th>
</tr>
</thead>
<tbody>
<tr>
<td>Jan</td>
<td>Swedish Energy Agency, Swedish Transport Administration, Swedish</td>
</tr>
<tr>
<td></td>
<td>Maritime Administration, Swedish Armed Forces</td>
</tr>
<tr>
<td>Feb</td>
<td>Annual Environmental Reports</td>
</tr>
<tr>
<td>Mar</td>
<td>Trade associations and private companies</td>
</tr>
<tr>
<td>Apr</td>
<td>Swedish Chemicals Agency</td>
</tr>
<tr>
<td>May</td>
<td>Swedish Board of Agriculture</td>
</tr>
<tr>
<td>Jun</td>
<td>Statistics Sweden</td>
</tr>
<tr>
<td>Jul</td>
<td>Trade associations</td>
</tr>
<tr>
<td>Aug</td>
<td>Swedish EPA, Statistics Sweden</td>
</tr>
<tr>
<td>Sep</td>
<td>SMED</td>
</tr>
<tr>
<td>Oct</td>
<td>Swedish EPA, Statistics Sweden</td>
</tr>
<tr>
<td>Nov</td>
<td>Swedish EPA, Transport Analysis, The Swedish EPA, Swedish Chemicals</td>
</tr>
<tr>
<td>Dec</td>
<td>Agency, Swedish Board of Agriculture, National Board of Forestry</td>
</tr>
</tbody>
</table>

**Figure 1-3. Overview of inventory planning, preparation and management.**
1.2.2 Training, awareness and skills

Training, awareness and skills in air pollutant reporting are essential to maintain the level of quality required according to specified requirements. Skills are ensured for the Swedish EPA and the majority of the government agencies involved in the work by the government agency being the sector government agency with staff that have particular skills in different specialist areas.

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.

1.2.3 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

- new international and national requirements, decisions and guidelines
- recommendations from international review not yet implemented in the inventory
- recommendations from national peer review not yet implemented in the inventory
- key source analysis (focus on major sources)
- uncertainty analysis (focus on sources 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

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 1.2.8 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 initiated.

From January to 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.
1.2.4 Inventory preparation (DO)
SMED gather data and information from various government agencies, organisations 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 3-7 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.

1.2.4.1 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 is 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. Default emission factors from the EMEP/EEA 2013 Guidebook are used to complement the national estimates.

1.2.4.2 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. Default emission factors from the EMEP/EEA 2013 Guidebook are used to complement the national estimates.

1.2.4.3 ENERGY – FUGITIVE EMISSIONS
For handling of solid fuels, activity data from Statistical Sweden is used, together with national emission factors for coal and peat.
Emissions from coke production are partly compiled from the facilities’ environmental reports, partly calculated via facility-specific activity data and default emission factors from the EMEP/EEA 2013 Guidebook.

For flaring in refineries and chemical industries, activity data from ETS are used for 2005 and later. In earlier years, data was collected through personal contacts with the facilities. Activity data from hydrogen production in oil refineries are taken from ETS. Regular emission factors for stationary combustion are used.

Activity data for transfer losses of gasworks gas are taken from the environmental reports provided by the facilities. Data on venting and flaring of gas in the national gas transmission network (natural gas and biogas) is reported by the operator. Emission factors for stationary combustion are used for flaring. Losses of gas during transmission, storage, venting and distribution are estimated using a national method and national data on typical gas compositions.

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.

### 1.2.4.4 INDUSTRIAL PROCESSES AND PRODUCT USE

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. Default emission factors from the EMEP/EEA 2013 Guidebook are used to complement the national estimates.

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.
1.2.4.5 AGRICULTURE
Data on livestock population, crop areas, crop yields, sales of manure, manure management systems 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, SLU and the Swedish Institute of Agricultural and Environmental Engineering. Default emission factors from the EMEP/EEA 2013 Guidebook are used to complement the national estimates.

1.2.4.6 WASTE
Emissions reported for waste incineration are compiled from the facilities' annual environmental reports. Other reported data are mainly based on models and uses statistical sources as activity data and default emission factors from EMEP/EEA 2013 Guidebook.

1.2.5 QA/QC procedures and extensive review of emission inventory (CHECK)

1.2.5.1 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 1-2).

All QC measures performed are documented by SMED in QC checklists for each CRF code or group of codes. When the reporting tables and the NIR are completed by SMED, a project quality control team (QC-team) performs checks before internal delivery to the SMED quality coordinator. The QC-team consists of one inventory compiler from each SMED organization (IVL, SCB and SLU) and the review
is performed so that each member of the team checks data and documentation from the inventory that it has not been involved in developing. In addition, the QC-team performs data checks in terms of the functionality of the CRF Reporter (i.e. checks of completeness, time-series consistency and recalculation explanations).

Before delivery of the inventory to the Swedish EPA, the SMED quality coordinator performs the final quality control. The QC-team and SMED quality coordinator checks serve as both quality control and quality assurance in accordance with the 2006 IPCC guidelines.

In addition, the project management team performs checks of submission data using the REPDAB tool\(^\text{15}\).

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

<table>
<thead>
<tr>
<th>CRF</th>
<th>Action</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A, 1B and parts of 2</td>
<td>Energy amounts</td>
</tr>
<tr>
<td>2C1</td>
<td>Iron and steel production</td>
</tr>
</tbody>
</table>

1.2.5.2 QUALITY ASSURANCE

The Swedish QA/QC system includes several QA activities outside the SMED QA/QC procedures. At the final stages of completion of the inventory, the Swedish EPA performs a peer review for each sector. In the 2018 submission, the Swedish EPA inventory compiler team consisted of seven members.

The Swedish QA/QC system also includes national peer reviews by sectoral authorities prior to inventory submission. The peer review is defined in the Ordinance on Climate Reporting (2014:1434) and thus includes only review of greenhouse gases. However, most underlying data is the same for the estimation of greenhouse gases and air pollutions, and thus, the national reviews serve as quality improvements also for the air pollution reporting. The national reviews include all sectors and are conducted by a person who has not taken part in the inventory preparation. The Swedish EPA is responsible for coordinating the peer reviews. From the 2016 submission, the national peer review is conducted in two steps:

- **Annual national review.** The aim of the review is to check the robustness of the national system and to guarantee that politically independent emissions data is reported. The review is performed by sectoral authorities prior to submission.

\(^{15}\) [http://www.ceip.at/check-your-inventory-repdab/](http://www.ceip.at/check-your-inventory-repdab/)
• *In-depth expert peer review*. Each year there is also an in depth peer review of one sector or part of a sector. The choice of sector depends on the outcome of the results from the EU and UNFCCC reviews and if the national review has identified problems or other needs discovered by SMED or the Swedish EPA. The aim of the in-depth expert peer review is to improve the inventory data quality. The review is performed by sectoral authorities and other national and international experts.

The annual national review is organised as a yearly meeting. Before the meeting the sectoral authorities have reviewed NIR in terms of the functionality of the national system and a general overview of methodology and statistics used (chapter 1), emission trends (chapter 2) and changes in methods, if changed (chapter 9). Thereafter the reviewers will provide feedback on whether they find the inventory reliable and independent, the trends are correct and the national system functional. Any recommendations for improvements are recorded in the list of suggested improvements described in section 1.2.8.

The in-depth expert peer review includes methodologies, models, activity data and emissions factors. 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 1.2.8.

The UNECE secretariat administers an international peer review of Swedish reporting after submission approximately every fourth year. Recommendations from the review reports are collected to the list of suggested improvements described in section 1.2.8.

### 1.2.6 Finalization, publication and submission of the inventory

The Swedish Environmental Protection Agency informs the Ministry of the Environment and Energy about the air pollutants inventory report in mid-December. At the same time, the inventory is published nationally.\(^\text{16}\)

The Swedish EPA submits the inventory to the CLRTAP/UNECE and NEC Directive/European Commission on February 15th. Reported data in the submission of year X relates to emissions year X-2, in other words emissions which took place during 2016 are reported in early 2018.

\(^{16}\) www.naturvardsverket.se
1.2.7 Data storage

A system for handling emission data, entitled Technical Production System (TPS)\(^\text{17}\), 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 peer reviewers to view data, plot time series and make comparisons between different years and submissions. For all NFR categories and sub-categories, time series from 1990 (sometimes 1980) 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 NFR tables or to plain MS Excel. Finally, TPS is used for data archiving of each submission. For access to the TPS, login with password is requested.

In addition to TPS, documentation, data and all calculations for each submission are stored at each organization’s servers and, for collective use and archiving, at two projects at Projectplace\(^\text{18}\). 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 major part of 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.

1.2.8 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 1.2.3. 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.

\(^\text{17}\) https://tps.naturvardsverket.se/

\(^\text{18}\) www.projectplace.com
1.3 Key source categories

Key source categories are sources that together contribute with either 95% of the level or 95% of the overall trend of reported emissions in Sweden. In this inventory, level and trend assessments are carried out for the following pollutants: As, Cd, CO, Cr, Cu, dioxins/furans, Hg, NH₃, Ni, NMVOC, NOₓ, PAH 1-4, Pb, PM₂.₅, PM₁₀, Se, SO₂, TSP and Zn. The level and the trend assessment are done with both the approach 1 and the approach 2 methods. The results of the Swedish key source analysis for 2015 are presented in Annex 1 together with the methodology.

1.4 General uncertainty evaluation

Since submission 2012, the general uncertainty evaluation is updated every submission. The uncertainties in the Swedish emission inventory reported to the CLRTAP were for the first time evaluated in 2003¹⁹, covering the emissions in 1990 and 2001. In order to prioritise efforts and resources in subsequent years, expert judgments mainly by the inventory staff together with IPCC references on uncertainties in activity data and emission factors have been the basis for the IPCC Tier 1 uncertainty evaluation.

In 2009, SMED performed a study to provide transparent uncertainty estimates of national emissions for the Swedish reporting to the CLRTAP of the submission 2010 in accordance with the Tier 1 methodology described in the EMEP/EEA Guidebook 2009 (Table 6-1)²⁰.

The complete results of the Swedish uncertainty analysis for 2016 are presented in Annex 1 together with the methodology. The summary table below (Table 1-3) show the uncertainty for the total emissions together with the uncertainty for the trend for all substances. The tables A.1.1 to A.1.19 in the annex show for all pollutants; estimated emissions 1990 and 2016, the uncertainty for the trend 1990-2016 and the uncertainty in national emissions 2016 together with the estimated uncertainty for the emission factor and for the activity data. For several of the substances the majority of the total variance derives from only a limited number of sources. For example, 72% of the variance in total NOₓ emissions derives from the application of mineral fertilisers. In general, the emission factors are more uncertain than the activity data.

¹⁹ Kindbom, 2004
²⁰ Gustafsson, 2009
1.5 General assessment of the completeness

The Swedish inventory covers all air pollutants required and most relevant sources with some exceptions. The general completeness for each sector is discussed below.

Sources where no pollutants not yet have been estimated, but may occur, include: 1A3ei Pipeline transport; 2C7d, Storage handling and transport of metal products; 2J, Production of POPs; 2K, Consumption of POPs and heavy metals; 3Da4, Crop residues applied to soils; 3Db, Indirect emissions from managed soils; 3Dd, Off-farm storage handling and transport of agricultural products; and 5C1biii, Clinical waste incineration.

For sources where PAH-4 is estimated, usually benzo(a)pyrene is estimated separately but not always the other 3 specified PAH-substances due to lack of information. As a consequence, national totals for the 4 specified PAH species in relation to PAH-4 may be misleading.

As a consequence of the in-depth review of the 2013 submission of the Swedish inventory under the CLRTAP and EU NEC directive, Sweden has implemented emission estimates for all missing sources and pollutants in the Swedish inventory where there are default emissions factors available and they can be considered to be representative of the national circumstances (first time in Submission 2016). In Table 1-4a and 1-4b those sources are listed that are not estimated in the Swedish
emission inventory and where there are default emission factors available in the EMEP/EEA Guidebook 2013. For each of these sources, an explanation is given as to why emissions have not been estimated.

Table 1-4a. Sources in the Swedish air pollutant inventory for which emissions have not been estimated – Main pollutants and particulates. Explanations are given below.

<table>
<thead>
<tr>
<th>NFR Code</th>
<th>NOx</th>
<th>NMVOC</th>
<th>SOx</th>
<th>NH3</th>
<th>PM2.5</th>
<th>PM10</th>
<th>TSP</th>
<th>BC</th>
<th>CO</th>
</tr>
</thead>
<tbody>
<tr>
<td>2B10b</td>
<td>NE (2)</td>
<td>NE (2)</td>
<td>NE (2)</td>
<td>NE (2)</td>
<td>NE (2)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1B1c</td>
<td>NE (5)</td>
<td>NE (5)</td>
<td>NE (5)</td>
<td>NE (5)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2C1b</td>
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Table 1-4b. Sources in the Swedish air pollutant inventory for which no emissions have been estimated – Heavy metals and POPs. Explanations are given below.

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(1) Notation key should be IE (included in NFR2C1). Will be corrected in submission 2018
(2) Notation key should be IE (included in NFR2B10a). Will be corrected in submission 2018
(3) Notation key should be IE (included in NFR2C7c). Will be corrected in submission 2018
(4) Default EF results in unreasonable emissions and is not applicable to Swedish conditions
(5) Not possible to estimate with the country-specific methodology currently used.
(6) No default method exists
(7) Notation key should be IE (included in NFR 5C1bii). Will be corrected in submission 2018
(8) Source is not mandatory
For the 1980’s no other data than the National total is reported in the NFR-tables for most of the pollutants and NE is inserted for sources where emissions could not be reported but would be expected. Data which are detailed enough to correctly report emissions by source for the 1980’s are not available. Particles, PAH and dioxin are however reported for individual sources from 1980. From 1988 this also applies for NMVOC.

1.5.1 Energy
Estimated emissions are considered to be complete for most sources. There might still be some incompleteness as regards in-house generated fuels in the chemical industry and in smaller companies.

Fugitive emissions, i.e. venting and flaring of liquid and gaseous fuels, are most likely not complete for smaller companies. However, all Swedish plants that flare gas and that are included in the European trading scheme for CO₂ from 2005, are included. For smaller plants, data might be reported in NFR1A instead of NFR1B. Hence lack of data on emissions from flaring is considered to be insignificant.

1.5.2 Industrial Processes and Product Use
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 EMEP/EEA air pollutant emission inventory guidebook – 2013. However, some exceptions do exist, mainly in terms of some heavy metal emissions and POPs from product use and consumption of POPs and heavy metals, where the default guidebook emission factors were judged inappropriate for Swedish conditions.

1.5.3 Agriculture
Emissions of NOₓ and NH₃ from crop residues are currently not estimated, because neither a country specific method nor a default method in the guidebook exists. Particulate matter from fur-bearing animals planned to be added to the inventory in submission 2018. Also emissions of NH₃ from application of sewage sludge are planned to be implemented in submission 2018. Beyond that all relevant agricultural emissions and sources are considered to be included in the inventory. For example does the majority of the country’s horses not belong to a farm, but are despite that included in the agricultural sector of the inventory. All sales of fertilizers are included, also quantities used in other sectors.

1.5.4 Waste
Emissions from incineration of Municipal Solid Waste (MSW) are included in IA1a, as MSW is used for energy production. In NFR 5, emissions of some pollutants from hazardous waste incineration, cremation, landfill fires and garden burning/bonfires are included, but not complete. For hazardous waste incineration, emissions from one large plant are included, and there may be emissions from smaller plants that are not covered. The overall completeness for the waste sector is unknown, but the inventory can be considered as complete in terms of using the suggested methods in EMEP/EEA Guidebook 2013.
2 Explanation of key trends

2.1 Emissions of pollutants regulated in the amended Gothenburg Protocol

Emissions of pollutants regulated in the amended Gothenburg Protocol (SO₂, NOₓ, NH₃, NMVOCs, PM₂.₅, and BC) have been reduced since 1990.

Figure 2-1. Trends in emissions 1990-2016 for NOₓ, SO₂, NH₃, NMVOC, PM₂.₅, and BC. Index year 1990=100 and BC index year 2000=100.
2.1.1 Nitrogen oxides (NO\textsubscript{X})

Swedish emissions of NO\textsubscript{X} amounted to 131.5 kt in 2015. Total emissions have decreased by 53 % since 1990 and by 3 % since 2015. The different sectors’ share of the NO\textsubscript{X} emissions in Sweden 2016 is shown in Figure 2-2.

![Distribution of NO\textsubscript{X} emissions among major contributing sectors and subsectors in 2016.](image)

The largest source of NO\textsubscript{X} emissions in 2016 was the transport sector (NFR1A3), where emissions amounted to 52.9 kt, corresponding to 40 % of total emissions. Emissions from the transport sector have been reduced by 66 % since 1990 and by 7 % since 2015. The main reason for the reduction since 1990 is stricter emission requirements for new vehicles during the period.

NO\textsubscript{X} emissions from passenger cars (NFR1A3b) were 68 % lower in 2016 compared to 1990 but since 2011 the emissions are increasing. Since 1990 there is more than a tenfold increase in NO\textsubscript{X} emissions from diesel passenger cars and between 2015 and 2016 the increase was 6 %. The combination of increased use of diesel due to the political ambition to reduce CO\textsubscript{2}-emissions from cars and the problems with the large discrepancies between Euro standards and real driving emissions (RDE) from diesel cars are the main reasons for this development. Emissions of NO\textsubscript{X} from gasoline cars and heavy-duty vehicles continue to decrease. Emissions from heavy duty vehicles were reduced by 18 % between 2015 and 2016 and by 73 % since 1990.
Figure 2-3. Trend in NOx emissions from transport (1A3b) 1990-2016.

The second largest source of NOX emissions in 2015 was combustion in manufacturing industries and construction (NFR1A2) in which emissions were 23.1 kt corresponding to 18% of total emissions. NOX emissions in the sector have been reduced by 42% since 1990 and increased by 2% since 2015.

Emissions of NOX from electricity and heat generation (NFR1A1a) amounted to 11.1 kt in 2016 corresponding to 9.5% of total emissions. Emissions have varied during the period with decreasing emissions during the 90’s and increasing emissions between 2000 and 2010. Energy supply for electricity and heat generation has increased by 60% between 1990 and 2016 while emissions were 17% lower in 2016 compared to 1990. Emissions to some extent depend on temperature and precipitation resulting in higher emissions during 1996 and 2010 which were cold and dry years.

A NOX fee, introduced in 1992, includes combustion plants (NFR1A1a and 1A2) with an output of more than 25 GWh per year. Since the introduction of the fee NOX emissions per unit produced energy have been reduced to less than half. This is an important reason why emissions of NOX from combustion in manufacturing industries and construction and electricity and heat production not have increased despite increased energy production.
Emissions of NO$_X$ from other sectors (NFR1A4) were 14.8 kt corresponding to 11 % of total emissions in 2016. Emissions have been reduced by 57 % since 1990. The reduction since 1990 is due to expanded district heating and increased use of heat pumps that have replaced oil heaters.

Agriculture contributed with 9.9 % of total emissions of NO$_X$ in 2016. The main sources are inorganic fertilizers and animal manure applied to soils. This post is new in the Swedish inventory as from submission 2016 and is considered to have high uncertainty.

Industrial processes (2) accounted for 9.4 % or 12.3 kt NO$_X$ emissions in 2016. Emissions have decreased by 16 % since 1990 and increased slightly, by 1 % since 2015. The main part, 87 %, of the emissions originates from the pulp- and paper industry (NFR2H1).
2.1.2  Sulphur dioxide (SO₂)

Total emissions of SO₂ have decreased by 87 % since 1990 and increased by 6 % since 2015 and amounted to 19.1 kt in 2016. The different sectors share of the SO₂ emissions in Sweden 2016 is shown in Figure 2-5.

![Distribution of SO₂ emissions among major contributing sectors and subsectors in 2016.](image)

Industrial processes (NFR2) was the sector with largest emissions of SO₂ in 2015. Processes within the metal industry (NFR2C) and pulp and paper industry (NFR2H1) generated 4.5 and 4.2 kt of SO₂ respectively in 2016, corresponding to 23 and 22 % respectively. Emissions have been reduced by 54 % and 67 % since 1990 and emissions from the metal industry (NFR2C) were reduced by 4 % while emissions from the pulp and paper industry (NFR2H1) increased by 3 % since 2015.

Emissions from combustion in manufacturing industries and construction (NFR1A2) amounted to 3.8 kt in 2016, corresponding to 20 % of total emissions. Emissions have been reduced by 81 % since 1990. The main reasons for the large reduction since 1990 are a reduced use of oil in the sector and better pollution control. Between 2015 and 2016 emissions increased by 29 % mainly due to increased combustion of biomass within pulp, paper and print (1A2d).

Emissions from electricity and heat generation (NFR1A1a) amounted to 3.0 kt in 2016. Emissions from the sector have been reduced by 81 % since. The two main reasons for the reduction since 1990 are reduced use of coal and oil and pollution control. Between 2015 and 2016 emissions increased by 10% mainly due to increased use of biomass but to some extent also increased use of oil.
Transport (NFR1A3) and other sectors (NFR1A4) amounted to 0.3 and 0.9 kt respectively in 2016, each corresponding to roughly 2 and 5%. Emissions in these sectors have been reduced by over 90% since 1990 due to reduced use of oil and reduced sulphur content in the oil used.

Figure 2-6. Trends in SO$_2$ emissions from major sectors and subsectors 1990-2016.
2.1.3 Ammonia (NH$_3$)

Total emissions of NH$_3$ in Sweden were 12% lower in 2016 than in 1990 and amounted to 53.1 kt. Between 2015 and 2016 emissions decreased by 2%. The different sectors’ share of the NH$_3$ emissions in Sweden 2016 is shown in Figure 2-7.

![Figure 2-7. Distribution of NH3 emissions among major contributing sectors and subsectors in 2016.](image)

Agriculture (NFR3) was the main source for emissions of NH$_3$ in 2016 and emissions were 46.7 kt. Ammonia is emitted from farm animals’ dung and urine and during the spreading of animal manure and the use of inorganic fertilizers.

Emissions from agricultural soils 3D were 24.9 kt in 2016 contributing with 47% to total emissions. The main contributions coming from animal manure and inorganic fertilizers applied to soils. Emissions were slightly lower in 2016 compared to 1990 but since 2012 a slight increase can be discerned.

Emissions from manure management (NFR3B) were 21.9 kt in 2016, contributing with 41% of total emissions in 2016. The main contribution came from dairy cattle, 3.5 kt, and non-dairy cattle, 9.3 kt. Emissions from manure management were 17% lower in 2016 compared to 1990. The reasons for the reduction of emissions are a decline in number of animals, a reduced use of inorganic fertilizers and a more effective production. There is a direct relationship between emissions of NH$_3$ and the number of animals.
Figure 2-8. Trend in NH₃ emissions from manure management (NFR3B) and Agricultural Soils (3D) 1990-2016.
2.1.4 NMVOC

Emissions of NMVOC which amounted to 158.9 kt in 2016 have decreased by 55 % since 1990 and by 2 % since 2015. The different sectors’ share of the NMVOC emissions in Sweden 2016 is shown in Figure 2-9.

Figure 2-9. Distribution of NMVOC emissions among major contributing sectors and subsectors in 2016.

Emissions of NMVOC from solvent use (NFR2D3) were 61.5 kt in 2016 and it was the dominant source contributing with 39 % to total emissions. Emissions from solvent use have decreased by 32 % since 1990. Important reductions occurred from coating applications (NFR2D3d) where emissions decreased by 70 % since 1990. Emissions from other non-specified (NFR2D3i) have almost tripled since 1990.

Emissions from the transport sector (NFR1A3) which accounted for 11.6 kt, corresponding to 7 % of national totals in 2016, have decreased by 92 % since 1990 and by 8 % since 2015. The main part of the reduction since 1990 (57 %) comes from passenger cars (NFR1A3bi) with the main reason being more energy efficient cars and the introduction of new exhaust requirements. Emissions from gasoline evaporation (NFR1A3bv) have been reduced by 99 % since 1990 due to emission control measures.
Emissions from other sectors (NFR1A4) amounted to 23.8 kt in 2016, corresponding to 15%. Since 1990 emissions have increased by 10%. The main contribution, 15.8 kt or 10% of total emissions, comes from the residential sector (NFR1A4b) where the sources are combustion of biomass and the use of gasoline in gardening equipment.

Emissions of NMVOC from manure management (NFR3B) were 20.7 kt in 2016, corresponding to 13%. Emissions are related to the number of animals and have as a result been reduced by 24% since 1990.

Fugitive emissions (NFR1B) of NMVOC were 11.8 kt in 2016 corresponding to 7% of total emissions of NMVOC. The emissions have decreased by 63% since 1990 but increased by 11% since 2015. The main part (close to 70%) of fugitive emissions originates from refineries and the main reduction can be seen in the emissions from gasoline distribution.
2.1.5 Particulate matter (PM$_{2.5}$)

Emissions of PM$_{2.5}$ were 18.4 kt in 2016. Emissions have been reduced by 51% since 1990 but increased by 2% since 2015. The different sectors’ share of the PM$_{2.5}$ emissions in Sweden 2016 is shown in figure 2-11.

![Distribution of PM$_{2.5}$ emissions among major contributing sectors and subsectors in 2016.](image)

Other sectors (NFR1A4) were the largest source of PM$_{2.5}$ in 2016 accounting for 6.7 kt or 37% of total emissions in 2016. Emissions were 15% lower in 2016 compared to 1990. The main part of the emissions in the sector (72%) came from stationary biomass combustion and biomass combustion in the residential sector (NFR1A4b). Emissions from biomass combustion in the residential sector, which have varied over the period, were 10% lower in 2016 compared to 1990.

Emissions from road transport (NFR1A3b), the second largest source, amounted to 4.1 kt in 2016. Emissions from road transport have decreased by 42% since 1990. The main reason for reductions since 1990 is stricter standards resulting in lower emissions from heavy duty vehicles (88% reduction since 1990) and buses (94% reduction since 1990). Specific emissions of PM$_{2.5}$ from diesel passenger cars have been reduced by over 90% since 1990 but overall emissions have not decreased to the same extent due to the sharp increase in diesel consumption which resulted in emissions being 56% lower in 2016 compared to 1990. The emissions from road abrasion, which depend on the total traffic work and the use of studded tires, increased with 28% between 1990 and 2016.

Emissions from combustion in manufacturing industries and construction (NFR1A2) were 1.4 kt in 2016. Emissions from the sector have been reduced by 71
% since 1990 but increased by 5% since 2016. About one third of the total emissions in the sector originated from biomass combustion within pulp- and paper (NFR1A2d) and other (NFR1A2g) including combustion in industries for wood and wood products. Emissions from biomass combustion in the sector have been halved since 1990. About half of emissions in the sector originate from off-road vehicles and other machinery in manufacturing industries and construction (NFR1A2g), mainly from the mining industry.

Emissions from electricity and heat generation (NFR1A1a) amounted to 0.9 kt in 2016. Emissions have decreased by 56% since while energy supply for electricity and heat generation has increased by 60% between 1990 and 2016. The reason for this decrease is better pollution control. Emissions in 2016 were 8% higher compared to 2015.

The pulp and paper (NFR2H1) and metal industries (NFR2C) are the two most important subsectors within industrial processes and product use (NFR2) where emissions were 1.6 and 1.0 kt respectively in 2016. In the metal industries there was a large reduction in emissions between 2013 and 2014 much of which can be explained by a new electro filter at a large pellet plant. Between 2015 and 2016 emissions were increased by 33%. There has been a significant reduction (74%) in emissions from the pulp- and paper industry between 1990 and 2016.

![Figure 2-12. Trends in PM$_{2.5}$ emissions from major sectors and subsectors 1990-2016.](image-url)
2.1.6 Black carbon (BC)

Total emissions of BC in Sweden were 2.9 kt in 2016. Emissions have been reduced by 38 % since 2000 and by 2 % since 2016. The different sectors’ share of the BC emissions in Sweden 2016 is shown in fig 2-13.

![Distribution of BC emissions among major contributing sectors and subsectors in 2016.](image)

The largest source of BC emissions with over one third of emissions in 2016 is other sectors (NFR1A4) where emissions amounted to 1.4 kt. The main part, 1.0 kt, originates from stationary biomass combustion. Emissions of BC from stationary biomass combustion were 32 % higher in 2016 than in 2000.

The second largest source is the transport sector (NFR1A3) in which emissions amounted to 0.7 kt in 2016. The most important source was road transport (NFR1A3b), contributing with 0.6 kt or 85 % of the emissions in the sector. Emissions from road transport have been reduced by 64 % since 2000 and the most important contribution to the reduction comes from heavy duty vehicles (80 %) and passenger cars (70 %) since 2000. The reductions are a result of stricter exhaust requirements.

Emissions from combustion in manufacturing industries and construction (NFR1A2) were 0.6 kt in 2015. Emissions were 46 % lower in 2016 compared to 2000. About 56 % originates from off-road vehicles and other machinery in (NFR1A2g), mainly from the mining industry.
Figure 2-14. Trends in BC emissions from major sectors and subsectors 1990-2016.
2.2 CO, PM$_{10}$, PAH-4, HCB & Dioxins

Emissions of CO, PM$_{10}$, PAH-4, HCB and dioxins all show decreasing trends over the period (1990-2016). The reduction of emission of the above-mentioned pollutants varies between 26 to 84 % but emissions have varied over the period.

Figure 2-15. Trends in emissions 1990-2016 for CO, PM$_{10}$, PAH-4, HCB and dioxins. Index 1990=100.
2.2.1 Carbon monoxide (CO)

The aggregated emissions of carbon monoxide (CO) have decreased from 1.1 Mt in 1990 to about 0.4 Mt in 2016, a decline of 60%. Between 2015 and 2016, emissions increased by 1%. The different sectors’ share of the CO emissions in Sweden 2016 is shown in figure 2-16.

![Distribution of CO emissions among major contributing sectors and subsectors in 2016.](image)

Other sectors (NFR1A4) was the largest source for CO emission, accounting for about two-thirds of the total emissions in 2016. Most of the emission is derived from biomass combustion in residential stationary plants, (NFR1A4bi). The rest were derived from other stationary combustion (NFR1A4ai), off-road vehicles and other machinery (NFR1A4aii and 1A4bii) and agriculture, forestry and fishing (NFR1A4c). Carbon monoxide emissions from other sectors (NFR1A4) were 19% higher in 2016 compared to 1990, mainly due to increased biomass combustion (Figure 2-17).

In 2016, the transport sector (1A3) was responsible for 19% of the total emissions of CO. Emissions from the transport sector have decreased by 90% since 1990, the main reason being the introduction of catalytic converters in passenger cars and light duty vehicles. Emissions of CO from passenger cars have decreased by 93% and from light duty vehicles by 95% since 1990.
Figure 2-17. Trends in CO emissions from other sectors (NFR1A4) and transport (NFR1A3) 1990-2016.

Carbon monoxide emissions from the industrial processes and product use sector (NFR2) were 28 kt in 2016 contributing with 7% of the total emissions and derived largely from aluminum (NFR2C3) and iron and steel production in the metal industry (NFR2C) as well as from pulp and paper industry (NFR2H1). Emissions from industrial processes and product use have increased by about 29% compared to 1990.

Emissions from electricity- and heat production (NFR1A1a) and combustion within manufacturing industries and construction (NFR1A2) amounted to 6 and 15 kt CO, respectively. Most of the emission in electricity- and heat production is derived from biomass combustion and the emissions from the sector have increased by more than 80% since 1990 due to increased combustion of biomass. In the manufacturing industries (NFR1A2) the emissions originate mainly from Off-road Vehicles and Other Machinery (1A2gvii) contributing with 2% of the national total emissions.
2.2.2 Particulate matter (PM$_{10}$)

Emissions of PM$_{10}$ were about 38 kt in 2016 and have decreased by 35% since 1990. The aggregated emission in 2016 increased by 2% compared to 2015. The different sectors’ share of the PM$_{10}$ emissions in Sweden 2016 is shown in figure 2-18.

![Figure 2-18. Distribution of PM$_{10}$ emissions among major contributing sectors and subsectors in 2016.](image)

The main sources of PM$_{10}$ emissions in 2016 were road abrasion, tyre and brake wear (NFR1A3bvi & NFR1A3bvii) accounting for 16.0 kt in which the major part (95%) comes from road abrasion. Emissions from road abrasion, tyre and brake wear were 28% higher in 2016 than in 1990. The magnitude of PM$_{10}$ emission depends on total traffic work and the use of studded tires.

The second largest source of PM$_{10}$ emissions was other sectors (NFR1A4) and amounted to 6.8 kt or 18% of total emissions in 2016. Most of the emission is derived from biomass combustion in residential stationary plants, (NFR1A4bi) with a total of 4.8 kt. Emissions from residential stationary plants varied during the period (1990-2016) and were 18% lower in 2016 than in 1990. The aggregated emissions from 1A4 in 2016 increased by 2% compared to 2015. (Figure 2-19).

Emissions from agriculture sector (NFR3) were 4.6 kt in 2016 or 12% of the national total. Manure management (NFR3B) and agricultural soils (NFR3D) contributed with about 2.5 and 2.1 kt, respectively. Emissions from manure management (NFR3B) (mainly poultry) have increased since 1990 due to increased numbers of slaughter chickens while emissions from agricultural soils (NFR3D) have decreased by about 17% during the same period due to decline of the area of cultivated cropland.
The aggregated emissions of PM$_{10}$ from industry (NFR2) were about 5 kt or about 13% of the national total. The pulp and paper industry (NFR2H1) accounts for the largest part of the emissions on the sector and contributes with 2.0 kt or 6% of the national total in 2016. Emissions in the sector have been reduced by two thirds since 1990. Large reductions in emissions have occurred in metal industry (NFR2C), 74% since 1990, mainly due to installation of new flue gas treatment in pellets production (NFR2C1e).

Emissions of PM$_{10}$ from combustion in manufacturing industries and construction (NFR1A2) amounted to 1.7 kt in 2016 and have been reduced by 72% since 1990. Biomass combustion in pulp, paper and print (NFR1A2d) contributed with about 0.5 kt in 2016. Diesel combustion in off-road vehicles (NFR1A2gvii) and biomass combustion in industry (NFR1A2gviii) contributed with about 1 kt.

Emissions from electricity- and heat production (NFR1A1a) were about 1.3 kt or about 3% of the total emission in 2016 and most of the emission is derived from biomass combustion. Emissions have decreased by about a half since 1990 mainly due to improved technology in large combustion plants (Figure 2-19).

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**Figure 2-19.** Trends in PM$_{10}$ emissions from major sectors and subsectors 1990-2016.
2.2.3 Poly Aromatic Hydrocarbons (PAH-4)

Emissions of poly aromatic hydrocarbons (PAH-4) were 12.6 t in 2016. Emissions have been reduced by 26% since 1990 but increased by 2% between 2015 and 2016. The different sectors’ share of the emissions of PAH1-4 in Sweden 2016 is shown in figure 2-20.

Figure 2-20. Distribution of PAH-4 emissions among major contributing sectors and subsectors in 2016.

Other sectors (NFR1A4) was the largest source of PAH-4 and contributed with about 9.9 t or 78% of the total emissions in 2016. Most of the emission is derived from biomass combustion in stationary plants (NFR1A4b). Emissions were slightly higher in 2016 compared to 1990 (Figure 2-21). The largest source of PAH-4 emissions within other sectors is combustion of biomass in residential stationary plants (NFR1A4bi), about 8.1 t in 2016. The emission trend for residential biomass combustion varied during the period and were about 14% lower in 2016 compared to 1990. The rest of the emission, about 1.7 t, comes from stationary biomass combustion in agriculture, forestry and fishing (NFR1A4ci) and commercial (NFR1A4ai).

PAH-4 emissions from solid fuel transformation (NFR1B1b) amounted to 1.1 t in 2016, equivalent to 9% of the total emissions.

Metal industry (NFR2C) is also a significant source of PAH-4 emissions which was responsible for 0.9 t in 2016. Between 1990 and 2016 emissions have been reduced by about 80% due to application of new technologies. Aluminium production (NFR2C3) was a key source of PAH-4 emission in Sweden until 2008 and decreased strongly since then as all pot-lines operating the Söderberg technology were shut-down in 2008. This has resulted in emission reduction by more than 99
% between 2008 and 2009 (Figure 2-21). In 2016, the emission from aluminium production was about 15 kg.

Figure 2-21. Trends in PAH-4 emissions from major sectors and subsectors 1990-2016
2.2.4 Hexachlorobenzene (HCB)

The total emissions of HCB in Sweden were 2.7 kg in 2016. Emissions have decreased by 84 % since 1990. Compared to previous year, the emissions in 2016 decreased by 30 %. The different sectors’ share of the HCB emissions in Sweden in 2016 is shown in figure 2-22.

![Figure 2-22. Distribution of HCB emissions among major contributing sectors and subsectors in 2016.](image)

Emissions of HCB from electricity- and heat production (NFR1A1a) amounted to about 1 kg in 2016 of which about half was derived from biomass combustion. Emissions from NFR1A1a have more than doubled (130 %) since 1990 mainly due to increased biomass combustion. HCB emission from this source depends, to some extent, on temperature and precipitation conditions in Sweden. For example, the observed high emission peaks in 1996 and 2010 were particularly cold years (Figure 2-23).

Emissions of HCB from metal industry (NFR2C) were about 0.7 kg in or 27 % of the national total in 2016. Emissions derived mostly from pellets production (NFR2C1e) and have more doubled since 1990 due to increased production. The economic recession in 2009 is clearly visible in form of a dip in the emission trend (Figure 2-23).

In 2016, about 0.4 kg or 15 % of HCB was derived from combustion within the manufacturing industry and construction (NFR1A2), mainly due to biomass combustion in pulp, paper and print (NFR1A2d) and other (NFR1A2g). Emissions have varied since 1990 and were 14 % higher in 2016 compared to 1990.
The waste sector (NFR5) is also a significant source of HCB emission and was responsible for about 0.3 kg or 10% of the total emission in 2016. The emission was mostly derived from incineration and open burning of waste (NFR5C). The emissions have increased by about 300% since 1990 due to increased burning of hazardous waste.

Biomass combustion in residential stationary plants (NFR1A4bi) accounted for 8% of the national emission of HCB in 2016. Emissions have increased by 7% since 1990. Emissions from heating of public and other facilities have also increased.

Figure 2-23. Trends in HCB emissions from major sectors and subsectors 1990-2016.
2.2.5 Dioxins - Polychlorinated dibenzodioxins and furans (PCDD/F)

The aggregated emissions of dioxins in Sweden were 24 g-Teq in 2016. Emissions have decreased by 64% since 1990 but increased by about 5% compared to 2015. The share of different sectors’ emissions of dioxins in Sweden in 2016 is shown in Figure 2-24.

![Distribution of dioxins emissions among major contributing sectors and subsectors in 2016.](image)

The waste sector (NFR5E) is the sector with the highest contribution to the emissions of dioxins to air with 7 g-Teq in 2016 and accounted for 30% in 2016. Emissions have increased by about 8% since 1990. The main source within the waste sector is House and car fires (NFR 5E) contributing with 25% of the total national emission of dioxins. A small amount of dioxin emissions to air come from combustion of hazardous waste. Since 1990 the emissions doubled due to increased combustion of hazardous waste.

Emissions of dioxins from electricity- and heat production (NFR1A1a) amounted to about 7 g-Teq or 29% of the national emission in 2016. Most of the emission was derived from biomass combustion. Emissions from electricity and heat production have decreased by 56% since 1990 due to technological improvement in combustion plants. The amount of dioxins emission from (NFR1A1a) depends to some extent on temperature and precipitation conditions in Sweden. The observed elevated emissions (jumps) in 1996 and 2010 were particularly cold years (Figure 2-25).

In 2016, about 3.1 g-Teq of dioxins emission to air were derived from manufacturing industry and construction (NFR1A2), mostly due to biomass combustion in pulp, paper and print (NFR1A2d). Emissions from manufacturing industries and
construction (NFR1A2) decreased by 65 % since 1990, due to improved technologies that have been adopted in various industries especially in pulp, paper and print. Emission of dioxin from the metal industry (NFR2C) were about 2 g-Teq or 8 % of the national total. The emission decreased by more than 90 % mainly from iron and steel production due to use of better technologies.

Stationary combustion plants (NFR1A4), i.e. residential plants (NFR1A4bi) accounted for 12 % of the national emission of dioxins in 2016. The emission from residential plants has increased by 7 % since 1990. Emissions resulted from commercial and institutional (NFR1A4ai) stationary combustion have also increased due to increased combustion of biomass.

Figure 2-25.  Trends in dioxins emissions from major sectors and subsectors 1990-2016.
2.3  Emissions of priority heavy metals

Emissions of cadmium (Cd), mercury (Hg) and lead (Pb) have all been reduced significantly since 1990. The most drastic decrease in emissions came from Pb being phased out from gasoline blends in the early 1990’s. Pb emissions have since 1990 been reduced by 97 %. In the early 1990’s Cd emissions were also heavily reduced, mainly due to efficiency improvements in metal production. Hg emissions were reduced by nearly half from 1990 until 1999 and most of the reduction came from improvements in metal processing and the waste sector (mainly cremation).

Figure 2-26.  Trends in emissions 1990-2016 for Cd, Hg and Pb. Index 1990=100.
2.3.1 Cd

Total emissions of Cd in Sweden were 540 kg in 2015, a total decrease of 77% compared to emissions in 1990. Emissions of Cd in 2016 are dominated by five sectors, as seen in Figure 2-27.

![Figure 2-27. Distribution of Cd emissions among major contributing sectors and subsectors in 2016.](image)

The largest source of Cd emissions in 2016 was electricity and heat production (NFR1A1a) where emissions amounted to 158 kg, corresponding to 29% of total emissions. Emissions of Cd from electricity and heat production have increased by 65% from 1990, much due to intensified activity over the years. Most of these emissions occurred at plants for combined heat and power generation where emissions from incineration of biomass have almost doubled since 1990. To some extent the emissions depend on temperature and corresponding demand on electricity and heat production. Year 1996 and 2010 were particularly cold which resulted in especially high emissions for these years.
Stationary combustion in the residential sector (NFR1A4bi) was the second largest source of Cd emissions with a total of 127 kg in 2015, corresponding to 23 % of total emissions. These emissions have decreased by 6 % compared to levels in 1990.

Combustion in manufacturing industries and construction (NFR1A2) represent 15 %, 83 kg, of total Cd emissions in 2016, mostly due to use of biofuels. Most of these emissions, 8 % of total Cd emissions, occurred in the pulp, paper and print industries (NFR1A2d). Cd emissions in the pulp, paper and print industries have decreased by 32 % since 1990. Emissions from combustion in manufacturing industries and construction have been relatively stable over the past ten-year period. From 2015 to 2016 the emissions increased by 1 %.

Industrial processes and product use (NFR2) amounted to 73 kg in 2016, corresponding to 13 % of total Cd emissions. Most of these emissions occurred in the metal industry (NFR2C) and pulp and paper industry (NFR2H1), with 8 % and 5 %, respectively, of total Cd emissions. Emissions in each industry are seen in Figure 2-29. Emissions from industrial processes and product use have decreased by 96 % from 1990. The emissions were drastically reduced during the early 1990’s and levels have been relatively stable since the early 2000’s.
Emissions from road traffic in the transport sector (NFR1A3b) accounted for 11\%, 61 kg, of the Cd emissions in 2016. The major part of these emissions came from gasoline use in passenger cars, with a total of 22 kg Cd emitted in 2016, corresponding to a decrease by 46\% since 1990. During the same period emissions from use of diesel fuels have increased, both in passenger cars and heavy goods vehicles. The increase from 1990 to 2016 was almost 800\% for passenger cars, 15\% for heavy duty vehicles and over 700\% for light duty vehicles. Cd emissions from use of diesel fuel amounted to 25 kg in 2016, with approximately 45\% coming from passenger cars.
2.3.2 Hg

Total emissions of Hg in Sweden were 412 kg in 2016, a total decrease of 73 % compared to emissions in 1990. Distribution of emissions of Cd between different sectors is shown in Figure 2-30.

In 2016, roughly one third of Hg emissions were generated through electricity and heat production (NFR1A1a) where emissions amounted to 148 kg. Most of these emissions, 115 kg, were generated from combined heat and power generation, where Hg emissions have decreased by 64 % since 1990. Much of these emissions were related to the use of biomass as fuel. The emissions mainly decreased between 1990 and 2000, after which emission levels increased for three consecutive years. For the past ten years the emissions from electricity and heat generation have been relatively stable.

Figure 2-30. Distribution of Hg emissions among major contributing sectors and subsectors in 2016.
Processes within the metal industry (NFR2C) accounted for 17 % of total Hg emissions in 2016. Emissions have been reduced by 86 % since 1990 and emissions in 2016 amounted to 68 kg.

In 2016, 14 % of total emissions occurred due to combustion in manufacturing industries and construction (NFR1A2), which corresponded to 56 kg of total emissions. Emissions have been reduced by 33 % since 1990, much due to less fuel being used in sectors of non-metallic minerals (NFR1A2f), pulp, paper and print (NFR1A2d) and other (NFR1A2g).

Emissions from road traffic in the transport sector (NFR1A3b) accounted for 10 % of the Hg emissions in 2016. Almost half of these emissions came from gasoline use in passenger cars, with a total emission of nearly 20 kg in 2016, while the other half came from use of diesel and other fuels. Emissions from road transportation have decreased by five percent since 1990 although emission levels remained nearly unchanged from 2015 to 2016. While emissions from combustion of gasoline have decreased since 1990 the emissions from combustion of the increasingly popular diesel have increased during the same period.

Emissions from incineration and open burning of waste in the waste sector (NFR5C) accounted for 8 % of the Hg emissions in 2016. The emissions in 2016 were 82 % lower than in 1990. During this period more waste have been handled by the energy sector where electricity and district heating is produced from waste incineration, resulting in less material being accounted for in the waste sector.
2.3.3  \textbf{Pb}

Total emissions of lead (Pb) in Sweden were 12 t in 2016, correlating to a total decrease of 97\% compared to emissions in 1990. Emissions of Pb are dominated by five sectors, as seen in Figure 2-32.

![Figure 2-32. Distribution of Pb emissions among major contributing sectors and subsectors in 2016.](image)

The most important sector, with one third of emissions, was processes and product use in the metal industry (NFR2C). In 2016 these emissions amounted to 3.7 t, and have been reduced by 95\% since 1990. Emissions from this sector were reduced drastically between 1990 and 1996 and since 2002 emissions have been leveled out with only slight variation between years.
The transport sector (NFR1A3) was the second most important source of emissions with 3.4 t of Pb in 2016 corresponding to 29% of the national totals. The main source comes from gasoline use for road transportation (NFR1A3b). As a consequence of lead being phased out from gasoline blends in the early 1990’s these emissions have been reduced by 99% compared to 1990. In 1990 the emissions from passenger cars amounted to 246 t, and in 2016 these emissions were 2 t.

Figure 2-33. Trend in emissions of Pb from Metal industry (2C) 1990-2016.

Figure 2-34. Trend in emissions of Pb from transport (1A3) 1990-2016.
Almost one fifth of the total emissions comes from public electricity and heat production (NFR1A1a) and is mainly related to combustion of biofuels. The emissions in this sector have been reduced by 23 % compared to emission levels in 1990. Emissions from manufacturing industries and construction (NFR1A2) accounted for 13 % of the Pb emissions in 2016. Almost half of these emissions come from the use of biofuels in pulp, paper and print industries (NFR1A2d). Since 1990 the emissions from manufacturing industries and construction has reduced by 74 %.

Five percent of total emissions, 0.6 t, occurred in the residential sector (NFR1A4b) where combustion of biomass was the major source. Compared to levels in 1990 total emissions from the residential sector has decreased by 24 % at the same time as emissions from biomass combustion in this sector have increased by 7 %.

Figure 2-35. Trends in emissions from Public electricity (1A1a) and heat production, Pulp, paper and print (1A2d) and residential stationary (1A4bi) 1990-2016.
3 Energy (NFR sector 1)

3.1 Overview

The energy sector includes emissions from fuel combustion (NFR1A) and fugitive emissions from fuel production and handling (NFR1B). 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 DIOX-ide dominate overwhelmingly in this sector.

3.2 Fuel combustion, NFR1A

Emissions from fuel combustion, NFR1A, are allocated to a number of subsectors.

NFR1A1 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.

NFR1A2 manufacturing industries, combustion-related emissions in manufacturing industries and construction and working machinery within the construction sector allocated to this subsector. Emissions from working machinery within the construction sector are allocated to NFR1A2, but apart from that, NFR1A2 includes only stationary combustion.

NFR1A3, emissions from domestic transport include aviation, road traffic, railways and navigation.

NFR1A4, emissions from other sectors, include stationary and mobile sources in households, service, agriculture, forestry and fisheries.

NFR1A5, emissions from other combustion include domestic military operations.

In addition, emissions from International aviation and international navigation (international bunkers) and multilateral operations, NFR1D, are not included in the national total.

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:

\[ \text{Emission}_{\text{fuel}} = \text{Fuel consumption}_{\text{fuel}} \times \text{Thermal value}_{\text{fuel}} \times \text{EF}_{\text{fuel}} \]
Different tier methods are used for different sub-sectors as discussed in sections below. Activity data sources, thermal values and emission factors are described in detail in Annex 2.

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 NFR2 and methods used are described in section 4.

3.2.1 Public electricity and heat production, NFR1A1a

3.2.1.1 SOURCE CATEGORY DESCRIPTION

Since 1980 the Swedish energy system has changed substantially. The dependence on fossil fuels (oil and coal), both for heating purposes and in industry, has decreased. During the period 1980-1990, the production of electricity from nuclear power plants increased from 26 TWh to 68 TWh. Another factor behind the decrease in fossil fuel use is the increased use of district heating. Use of biofuel (wood chips, bark) and incineration of municipal waste is common in district heating plants. The use of heat pumps both in district heating plants and in residential houses has increased since 1985. In the manufacturing industry the combustion of oil products has decreased from 1980 and has to a large extent been replaced by biofuels.

Swedish electricity production 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, mainly due to variations in ambient wintertime temperatures. 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. 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. Electricity is an important energy source in the manufacturing industry, which is dominated by the pulp and paper industry and the steel industry.

Production of district heating is currently to a large extent based on biomass and waste. There has been a shift from fossil fuels towards biomass since 1990. In 1990, 25% of fuels used were biomass including biogenic waste, and 6% was fos-
sil waste. In 2016, 73 % of all fuels used for district heating were biomass (including the biogenic fraction of waste), while waste (fossil fraction) accounted for 11 % \textsuperscript{21}. 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 176 PJ (2015) \textsuperscript{22} but, due to the more frequent use of biomass, greenhouse gas emissions from district heating were lower in 2016 than in 1990.

The number and distribution of Swedish power stations in 2015 are presented in Table 3.1 \textsuperscript{23}. Changes in number of plants and their installed effect have been minor in the production of district heating, but due to growing wind power the number of plants in the electricity sector have increased.

<table>
<thead>
<tr>
<th>Type of plants</th>
<th>Number of plants</th>
<th>Gross Production GWh</th>
<th>Gross Production TJ</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total power stations</td>
<td>4 332</td>
<td>161 959</td>
<td>583 052</td>
</tr>
<tr>
<td>Power generation not based on fuels</td>
<td>4 163</td>
<td>91 707</td>
<td>330 145</td>
</tr>
<tr>
<td>Wind power</td>
<td>3 174</td>
<td>16 268</td>
<td>58 565</td>
</tr>
<tr>
<td>Hydropower</td>
<td>989</td>
<td>75 439</td>
<td>271 580</td>
</tr>
<tr>
<td>Power generation based on fuels</td>
<td>169</td>
<td>70 252</td>
<td>252 907</td>
</tr>
<tr>
<td>Nuclear power</td>
<td>3</td>
<td>56 348</td>
<td>202 853</td>
</tr>
<tr>
<td>Conv. thermal power</td>
<td>166</td>
<td>13 906</td>
<td>50 062</td>
</tr>
</tbody>
</table>

The trend in fuel consumption in this sector varies depending on the production of hydropower and climate variables. The greatest changes in fuel consumption are for biomass fuels, where the consumption has increased significantly due to, for instance, increased district heating. There was a significant increase in the use of natural gas in 2009 due to an increase in the number of gas fuelled facilities. In 2010, the production and use of district heating was unusually high due to the cold weather with unusually low temperatures in the beginning and the end of the year.

A summary of the latest key category assessment is presented in Table 3-2.

\textsuperscript{21} All numbers are according to data used in the greenhouse gas inventory this submission. The proportions given are calculated for heat production, and may include plants in both 1.A1.A.ii and 1.A1.A.iii

\textsuperscript{22} Statistics Sweden/Swedish Energy Agency EN11SM 1601 (Electricity supply, district heating and supply of natural and gasworks gas 2015.). Data for 2015 currently not available.

\textsuperscript{23} Data for 2016 currently not available. Statistics Sweden /Swedish Energy Agency EN11SM 1601 (Electricity supply, district heating and supply of natural and gasworks gas 2015).
Table 3-2. Summary of key source analysis, NFR1A1a, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>Level</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1A1a</td>
<td>Biomass – As, Pb, Cd, Hg, DIOX, Zn, Ni, Se, Cu, Cr, NOx, PM1.5, NMVOC, PM10, SO2, TSP, PAH 1-4, CO</td>
<td>Biomass – As, Pb, Cd, Hg, DIOX, Zn, Ni, Se, Cu, Cr, NOx, PM1.5, NMVOC, PM10, SO2, TSP, PAH 1-4, NH3</td>
</tr>
<tr>
<td></td>
<td>Gaseous –</td>
<td>Gaseous –</td>
</tr>
<tr>
<td></td>
<td>Liquid – Ni, SO2, NOx</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Solid – SO2, Hg, As, Cr, Cd, Ni</td>
<td>Solid – Se, PM2.5, SO2, PM10, TSP, Hg, As, DIOX, NOx</td>
</tr>
<tr>
<td></td>
<td>Other – Hg, As, Se, Pb, DIOX, SO2, Cr, Cd, Ni, PM2.5, NOx</td>
<td>Other – Hg, As, Pb, DIOX, SO2, Cr, Cd, PM2.5, NOx</td>
</tr>
<tr>
<td></td>
<td>Peat – SO2, As, Hg</td>
<td>Peat – PM10, TSP, SO2, DIOX, Hg, Ni, NOx, PM2.5, Se</td>
</tr>
</tbody>
</table>

3.2.1.2 METHODOLOGICAL ISSUES
A combined Tier 2 and 3 methods is used. Activity data for emissions in NFR1A1a are taken from quarterly fuel statistics. For this sector, the quarterly fuel statistics is sent to all companies registered as ID 40 according to databases used by Statistics Sweden and the response rate is almost 100 %. This gives very good data to the inventory, accurate, complete and consistent and with very low uncertainties.

No emissions from the integrated iron and steel industry are allocated to NFR1A1a. However, emissions from steelwork gases sold to and combusted by ISIC 40 facilities are still allocated to NFR1A1a.

3.2.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
For the energy sector, the largest uncertainties come from activity data for the 1980's and from emission factors.

Due to problems with data files on energy consumption in energy industries and manufacturing industries 1980-1989, it has not been possible to recalculate emissions as has been done for different sectors 1990 and later years (described below). Because of this, time series 1980-89 and 1990-later are not directly comparable. The differences are largest for NFR1A1b, 1A1c, 1A2a and 1A2c.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for all fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As, diox, HCB, PCB and Zn
- Other fuels for As, Cr, Cu, Diox, Ni, Se, Zn, Cd
- Other fuels for HCB, PAH, PCB
- Peat for PCB
- Solid fuels for Cd, Cu, HCB, PCB, Se
Due to the final revision of emission factors for NFR 1A1a some IEF time series are in general lower than in submission 2017. These are in particular Zn, PM10 and the indirect GHGs SO$_2$ and NO$_X$.

- Liquid fuels: Lower for NO$_X$, SO$_2$
- Biomass fuels: Lower for NO$_X$ and Zn
- Peat: Lower for PM10

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Experts at the Swedish EPA conduct a review of the inventory estimates, methodologies and emissions factors used. The experts also identify areas of improvement, which constitute part of the basis for improvements in coming submissions.

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 explanations. IEFs for all reported substances are calculated per fuel, substance and NFR-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 reports from the UNFCCC and CLRTAP/NEC reviews have been carefully read and taken into account.

As part of the inventory procedure for submission 2007, a separate study$^{24}$ was performed to verify the quality of all fossil fuel combustion-related activity data from the largest plants (in terms of CO$_2$-emissions) in Sweden in 2005. The verification consisted of a comparison of plant-specific SMED-data (energy statistics from the quarterly fuel statistics) with data from the EU Emission Trading System (ETS). The results showed that for 21 plants, accounting for about 50% of the fossil fuel

$^{24}$ Backman & Gustafsson, 2006
consumption of the 63 plants included in the study, no significant differences between the two data sources were identified. For a number of plants, large differences occurred between the two data sources. In 2007, 19 of these plants were further surveyed in another study. Again, energy statistics (the quarterly fuel statistics) and ETS data by plant were compared and analysed.

The results show that the reported fuel amounts differ slightly between the data sets and since ETS data are verified, they are likely to be more correct. Another deficiency in the quarterly fuel statistics is that unconventional fuels are often grouped and the emission factors of these fuels are associated with very large uncertainties, since they are not specific for the current fuel and plant. Finally, another problem is that some of those unconventional fuels are incorrectly classified. According to data reported to ETS, some of these fuels are often partly biogenic and should hence be classified as "Other biomass".

3.2.1.5 SOURCE-SPECIFIC RECALCULATIONS
In submission 2017, a major revision of the emission factors was initiated and finished in submission 2018. The revision is described in more detail in Annex 2. Briefly, the completion of the revision in NFR 1A1a includes several fuel types and recalculations are in general made for certain periods or years and not whole time series.

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

3.2.2 Refineries, NFR1A1B
3.2.2.1 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. There are five refineries in Sweden. Three of these produce fuel products such as gasoline, diesel and heating oils. The other two 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 is mainly based on refinery gas, which is a by-product in the refining process. The use has increased due to higher demand of refined products.

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25 Nyström, 2007
Table 3. Summary of key source analysis, NFR1A1b, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A1b</td>
<td>Liquid Fuels - PM$_{2.5}$, Ni, SO$_2$, NO$_x$</td>
<td>Liquid Fuels - PM$<em>{2.5}$, PM$</em>{10}$, TSP, Ni, SO$_2$.</td>
</tr>
</tbody>
</table>

3.2.2.2 METHODOLOGICAL ISSUES

The Tier 2 method is used.

The statistics for NFR1A1b are based on a total of seven plants with the Swedish Standard Industrial Classification 192, petroleum refining. Five of these companies are real refineries which use more than 99% of the energy within the sector and thereby cause most of the emissions. The other two plants are oil companies, mainly involved in production of lubricating grease.

Activity data for the five refineries has been 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 for 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.

Data from ETS are used for the refinery plants for 2005 and later years. In 2008 and later years, the quality of ETS data is considered to be very high for all the five refineries, and thus this is the primary source of activity data for this sector.

The fuel consumption in this sector is mainly based on liquid fuels and the use has increased compared to the early 1990’s due to higher demand of refined products. Since the early 2000’s, however, there is no obvious trend in fuel consumption in this sector.

3.2.2.3 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 in earlier years has been analysed and no (significant) signs of inconsistency have been found. In recent years, environmental reports are used for verification.

The assigned uncertainties are based on information directly from the facilities. These are updated regularly but not annually. The uncertainty of the activity data is around 1.5%, but the uncertainty of the NCV is unknown, so the total uncertainty for the activity data was judged to 10%. Activity data uncertainty for the 1990’s is also estimated to 10%.

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26 Backman & Gustafsson, 2006.
Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for Gaseous fuels and higher for Liquid fuels. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty is Liquid fuels for all emissions often larger than 100%. EF for As, CO and NMVOC are larger than 100%. See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for NFR 1A1b as for 1A1a described above. For each of the five refineries, ETS data for the latest year are verified against the refineries’ legal environmental reports. 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 NFR 1A1b 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.

In submission 2018, a development project was carried out with the specific purpose to improve emission allocation between the energy sector and IPPU and to establish a procedure for annual cross-sectoral control of reported emissions. Within this project, a new quality control tool has been developed to ensure that comparisons of emissions in the energy and IPPU sector are done on a more regular basis, for the same range of facilities, and using a unified procedure. In the tool, for each of the relevant facilities, emissions reported in the energy sector (NFR 1A) are summed up with the emissions reported in the NFR 2, and the sum is compared to both EU ETS data (if available) and environmental reporting provided by facilities. Quality control is therefore being conducted on a facility level. In case of discrepancies, they are easily identified and further investigated regarding potential gaps or double-counting. This work is ongoing, and feasible reallocations will be done in the submission 2019.

3.2.2.5 SOURCE-SPECIFIC RECALCULATIONS

During submission 2018, a major revision of emission factors for whole industry sector was implemented, including 1A1b. The revision concerned various fuel types and for all emission factors except for CO₂. A more in depth description of the new emission factors used are found in Annex 2. In addition due to revision of
allocation of fuel consumption between NFR 1 and 2 refinery gas was reallocated between the NFR codes 1A1b. and 1B. according to the results of the study.²⁷

3.2.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.3 Manufacture of solid fuels and other energy industries, NFR1A1c

3.2.3.1 SOURCE CATEGORY DESCRIPTION
This category includes emissions 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.²⁸ The trend is related to the amounts of iron and steel produced, and hence there was a dip in 2009. Since 2009, the production and the emissions have increased gradually, and in 2013 the emissions were about the same level as in the early 2000’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 3-4.

Table 3-4. Summary of key source analysis, NFR1A1c, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source</th>
<th>Assessment 2016</th>
<th>Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A1c</td>
<td>Solid fules – SO₂</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Solid fules – TSP</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

3.2.3.2 METHODOLOGICAL ISSUES
The Tier 2 method is used. Emissions from fuel combustion in the manufacturing of solid fuels are reported under NFR1A1c, in line with IPCC Guidelines. This includes emissions from combustion in coke ovens in the iron and steel industry.

Activity data on coke production is taken from environmental reports. Emissions of NMVOC and CO are estimated with the Tier 2 methodology with national emission factors. Estimates of emissions of SO₂ and NOₓ are available from environmental reports on an aggregate level, and these emissions are distributed over the different NFR codes (1A1c, 1A2a, 1B1c and 2C1, SO₂ also 2B5 and 1B1b) according to the activity data distribution.

For 1980-1989, activity data is taken from the industrial statistics and quarterly statistics. For more details on the surveys see Annex 2. Activity data on combustion of coke oven gas and blast furnace gas in coke ovens is discussed in connection

²⁸ Fuel combustion in manufacturing of nuclear fuels was included in NFR 1A1c in previous submissions, but for confidentiality reasons the very small emissions from these facilities have been included in NFR 1A1aiii instead.
with other emissions from the iron- and steel industry in section 4.4 Metal production, NFR2C.

Since 1990, solid fuel consumption has increased slightly due to higher production of coke caused by higher demand of primary iron and steel. In 2009, however, solid fuel consumption decreased considerably due to lower production of coke, caused by a lower demand of primary iron and steel.

For confidentiality reasons all emissions in 1A1b are reported as C.

3.2.3.3 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.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Liquid fuels for Cd, Cu, Hg, Pb, Se, Zn
- All fuels for Cr, Ni, PAH, SO2, TSP and PM10 and PM2.5

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.3.4 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.

The improvements in methodology and allocation of emissions from the integrated iron and steel industry in submission 2010 were made based on a study carried out in 2008 looking at emissions from several industrial plants, including the two largest iron and steel plants in Sweden, where inventory data from submission 2008 was compared with data from environmental reports. In 2010, activity data

and emission factors for the chemical industry and the most important metal foundries were verified against data from environmental reports in a similar study\textsuperscript{30}.

3.2.3.5 SOURCE-SPECIFIC RECALCULATIONS
In submission, 2018 there were several recalculations for the sector. The whole particle emissions time series was revised for the two largest plants.

3.2.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.4 Iron and steel, NFR1A2a

3.2.4.1 SOURCE CATEGORY DESCRIPTION
A limited number of industries account for the majority of industrial energy use, i.e. the pulp and paper industry, iron and steel works and the chemical industry together account for about 65\% of the fuel used. 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.

In Sweden, there are three primary steel works that base their production on iron ore pellets producing either steel or iron powder. There are also 10 secondary steel plants producing steel based on scrap metal. In 2009, fuel consumption in the iron and steel industry fell sharply as a consequence of decreased production (2.8 Mt of steel) due to the global recession. In 2016, the production was 4.6 Mt\textsuperscript{31}, an increase of almost 6 percent compared to 2015. Emissions from iron and steel companies with less than 10 employees are allocated to NFR 1A2g because the model estimate of fuel consumption for these small companies is produced on an aggregate level and not separated by ISIC code.

The trend of the fuel combustion is increasing slightly since 1990 due to higher production of iron and steel products. In 2009 this trend was broken due to decreasing demand of iron and steel. In 2010, production and fuel consumption recovered to more “normal” levels.

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.

\textsuperscript{30} Gustafsson, T., Nyström, A-K., Gerner, A., 2010
\textsuperscript{31} The Swedish Steel Producers’ Association, 2017-11-08.
Table 3-5. Summary of key source analysis, NFR1A2a, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2a</td>
<td>Liquid – Ni, SO₂, NOₓ</td>
<td>Liquid – Ni, Se,</td>
</tr>
<tr>
<td></td>
<td>Solid – SO₂</td>
<td>Solid – SO₂c</td>
</tr>
</tbody>
</table>

3.2.4.2 METHODOLOGICAL ISSUES

The Tier 2 method is used.

During 2009, a new methodology was implemented for the two largest primary iron and steel works. Activity data for all other facilities 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 NFR1A1c, 1A2a, 1B1b, 1B1c and 2C1 since some emissions arise from fuel combustion and some from reducing agents in the process. The text in this section is hence closely connected to the text in section 4.4 Metal production, NFR2C. NFR2C1 (iron and steel production). Fuel combustion has increased slightly since 1990 due to higher production of iron and steel products. However, there was a significant decrease in solid fuel consumption in 2009 due to lower production of coke, caused by a lower demand of primary iron and steel.

Due to confidentiality reasons liquid and biomass fuels are reported as C for energy consumption.

3.2.4.2.1 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 Figure 4- 1 in section 4.4 Metal industry (NFR2C). Emissions from fuel combustion (oils, LPG 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 are allocated to this sub-sector in accordance with the IPCC Guidelines.

3.2.4.2.2 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. For these facilities, data on fuel consumption for energy purposes is from the quarterly fuel statistics. National NCVs and emission factors are used.
3.2.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

For the two largest facilities, the time series is considered to be very consistent since the time series developed in 2009 was compiled in close cooperation with the facilities. For NFR1A2a 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 annual 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.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Liquid fuels for As, Cd, Cr, Cu, NH3, Ni, Pb, Se, Zn
- Biomass for As
- All fuels for diox, PAH, PM10, PM2.5 SO2

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for NFR1A2a as for 1A1a 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.

In submission 2018, a development project was carried out with the specific purpose to improve emission allocation between the energy sector and IPPU and to establish a procedure for annual cross-sectoral control of reported emissions. Within this project, a new quality control tool has been developed to ensure that comparisons of emissions in the energy and IPPU sector are done on a more regular basis, for the same range of facilities, and using a unified procedure. In the tool, for each of the relevant facilities, emissions reported in the energy sector (NFR 1A) are summed up with the emissions reported in the NFR 2, and the sum is compared to both EU ETS data (if available) and environmental reporting provided by facilities. Quality control is therefore being conducted on a facility level. In case of discrepancies, they are easily identified and further investigated regarding potential gaps or double-counting. This work is ongoing, and feasible reallocations will be done in the submission 2019.
3.2.4.5  SOURCE-SPECIFIC RECALCULATIONS

In submission 2018 a major revision of the emission factors was made within the industrial sector. Briefly, the revision was to a large extent based on conclusions from the previous major emission factor revision that was made within the public electricity and heat production sector (NFR 1A1a) in submission 2017. This is due to the fact that many combustion processes in the industrial sector are similar to those in the heat and power sector. In conclusion, the revision includes several fuel types and recalculations were in general made for the whole time series for all the revised fuel types. The revision is described in more detail in Annex 2.

3.2.4.6  SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

3.2.5  Non-Ferrous Metals, NFR1A2b

3.2.5.1  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 4.4.

As for all subcategories to NFR1A2, for companies with less than 10 employees the Tier 2 method is used since country specific emission factors are used. Emissions from companies with less than 10 employees are allocated to NFR1A2g.

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. In recent years, the copper producers account for 40-50% of the fuel consumption in 1A2b and the aluminium producers account for 32-45%. The most common fuel is LPG (45-61% in recent years), followed by natural gas and heating oils.

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 key source analysis, NFR1A2b, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2015</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2b</td>
<td>Liquid –</td>
<td>Liquid –</td>
</tr>
<tr>
<td></td>
<td>Solid –</td>
<td>Solid –</td>
</tr>
</tbody>
</table>

3.2.5.2 METHODOLOGICAL ISSUES

3.2.5.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
As for NFR1A2a, time series consistency despite the changes in activity data source is discussed in Annex 2.

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.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Solid fuels for Zn
- Liquid fuels for all emissions
- All fuels for As, diox
- Other fuels for PM10, PM2.5, TSP

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.5.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
The same QA/QC procedures are used for NFR1A2b as for 1A2a described above. In addition to this, a detailed quality study of the non-ferrous metal industry was performed in 2010. The aim of this study was to investigate suspected errors in several CRF/NFR codes. Data for selected plants were verified with environmental reports per facility on an aggregate level, i.e. NFR1+2 together. This study showed no reasons for revisions in NFR1A2b. In submission 2018, a development project was carried out with the specific purpose to improve emission allocation between the energy sector and IPPU and to establish a procedure for annual cross-sectoral control of reported emissions. Within this project, a new quality control tool has been developed to ensure that comparisons of emissions in the energy and IPPU sector are done on a more regular basis, for the same range of facilities, and using a unified procedure. In the tool, for each of the relevant facilities, emissions reported in the energy sector (NFR 1A) are summed up with the emissions reported in the NFR 2, and the sum is compared to both EU ETS data (if available) and en-

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34 Skårman et.al, 2010.
environmental reporting provided by facilities. Quality control is therefore being conducted on a facility level. In case of discrepancies, they are easily identified and further investigated regarding potential gaps or double-counting. This work is ongoing, and feasible reallocations will be done in the submission 2019.

3.2.5.5 SOURCE-SPECIFIC RECALCULATIONS
In submission 2018 a major revision of the emission factors was made within the industrial sector. Briefly, the revision was to a large extent based on conclusions from the previous major emission factor revision that was made within the public electricity and heat production sector (NFR 1A1a) in submission 2017. This is due to the fact that many combustion processes in the industrial sector are similar to those in the heat and power sector. In conclusion, the revision includes several fuel types and recalculations were in general made for the whole time series for all the revised fuel types. The revision is described in more detail in Annex 2.

3.2.5.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.6 Chemicals, NFR1A2c
3.2.6.1 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 %.

As in other subcategories of NFR1A2, emissions from companies with less than 10 employees are allocated to NFR1A2g.

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 3-7.

### Table 3.7. Summary of key source analysis, NFR1A2c, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2c</td>
<td>Biomass –</td>
<td>Biomass –</td>
</tr>
<tr>
<td></td>
<td>Liquid – Ni, NO&lt;sub&gt;X&lt;/sub&gt;, SO&lt;sub&gt;2&lt;/sub&gt;</td>
<td>Liquid – Ni, NO&lt;sub&gt;X&lt;/sub&gt;</td>
</tr>
<tr>
<td></td>
<td>Other Fuels – SO&lt;sub&gt;2&lt;/sub&gt;</td>
<td>Other Fuels – SO&lt;sub&gt;2&lt;/sub&gt;</td>
</tr>
</tbody>
</table>

#### 3.2.6.2 METHODOLOGICAL ISSUES

The Tier 2 method is used. 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.3\textsuperscript{38} in the energy statistics are included in this sector, as recommended in IPCC 2006 Guidelines.

In submission 2009, after careful studies of different data sources regarding activity data of consumption of other petroleum fuels in this sector, it was found that the fuel used is a by-product of the process in one facility, a gas that consists mainly of methane. Since no specific emission factors for methane and methane based gas mixtures are available, emission factors for natural gas are used as these fuels are considered to have similar properties, but of course fuel consumption and emissions are still reported under liquid fuels. As natural gas contains around 90 molar % methane, the emission factors are considered to be accurate also for methane-rich gas mixtures of liquid origin.

In a development project in 2010\textsuperscript{39}, 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 NFR1 and 2 were analysed on facility level and verified against environmental reports, and when necessary the facilities were contacted for explanations or complementary data. Most of the data reported in submission 2010 was concluded to be correct, and only a few revisions had to be made in submission 2011. A few erroneous activity data records were detected and revised. The errors include double-counting, input data errors and miscoding, e.g. biogenic ethanol that had been coded as natural gas or hydrogen coded as other petroleum fuels.

The project also resulted in revisions of a couple of emission factors. Emission factors for hydrogen, which were previously set equal to those of “other petroleum fuels” for all substances containing nitrogen, i.e. including NH<sub>3</sub>, were corrected and set to zero for all substances except for NO<sub>X</sub>.

\textsuperscript{38} United Nations Statistics Division, 2010

\textsuperscript{39} Gustafsson, Nyström & Gerner, 2010
The revision that had the largest impact on the emissions is the conclusion drawn that some (not all) of the natural gas consumption previously reported in NFR1A2c 2004 and onwards is actually used as feedstock and not for energy production, and hence no emissions from this activity should be reported in NFR1A2c.

3.2.6.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The uncertainty in activity data is 1.5 % (2012) and the emission factor uncertainty is assumed to be 10 % based on the variation in plant specific values. The Activity data uncertainty for this fuel 2012 is as reported to the EU ETS. For the other fuels used and for all fuels for 1990, uncertainties are assigned by expert judgement by staff at the energy statistics department of Statistics Sweden.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As
- Liquid As, Cr, Cu, Hg, Ni, Se, Zn
- Other fuels for As, Cr, Cu, Hg, Ni Pb, PAH
- Gaseous fuels for PAH, PM10, PM2.5

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.6.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for NFR A2c as for 1A2a and 1A2b 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 2010 mentioned above, the activity data time series 1990-2008 for all fuel types and all facilities within the chemical industry were thoroughly reviewed. In submission 2018, a development project was carried out with the specific purpose to improve emission allocation between the energy sector and IPPU and to establish a procedure for annual cross-sectoral control of reported emissions. Within this project, a new quality control tool has been developed to ensure that comparisons of emissions in the energy and IPPU sector are done on a more regular basis, for the same range of facilities, and using a unified procedure. In the tool, for each of the relevant facilities, emissions reported in the energy sector (NFR 1A) are summed up with the emissions reported in the NFR 2, and the sum is compared to both EU ETS data (if available) and environmental reporting.
provided by facilities. Quality control is therefore being conducted on a facility level. In case of discrepancies, they are easily identified and further investigated regarding potential gaps or double-counting. This work is ongoing, and feasible reallocations will be done in the submission 2019.

### 3.2.6.5 SOURCE-SPECIFIC RECALCULATIONS

In submission 2018 a major revision of the emission factors was made within the industrial sector. Briefly, the revision was to a large extent based on conclusions from the previous major emission factor revision that was made within the public electricity and heat production sector (NFR 1A1a) in submission 2017\(^{41}\). This is due to the fact that many combustion processes in the industrial sector are similar to those in the heat and power sector\(^{42}\). In conclusion, the revision includes several fuel types and recalculations were in general made for the whole time series for all the revised fuel types. The revision is described in more detail in Annex 2.

### 3.2.6.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

### 3.2.7 Pulp, Paper and Print, NFR1A2d

#### 3.2.7.1 SOURCE CATEGORY DESCRIPTION

In 2016 there were 50 paper mill and pulp industry plants and 120 sawmills (production capacity >10 000 m\(^3\)/year) in Sweden. In total, they were producing 10.1 Mt of paper, 17.8 Mm\(^3\) of sawn timber and 11.6 Mt of pulp\(^{43}\). 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 83 % in 2016. As for NFR 1A2 in general, emissions from companies with less than 10 employees are allocated to NFR 1A2g.

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 3-8.

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\(^{41}\) Mawdsley, I., Wisell, T., Stripple, H., Ortiz, C. 2016.

\(^{42}\) Mawdsley, I., Stripple, H. 2017.

\(^{43}\) The Swedish Forest Industries Federation, 2017-11-10 http://www.skogsindustrierna.se/skogsindustrin/skogsindustrin-i-korthet/fakta--nycketal
Table 3. Summary of key source analysis, NFR1A2d, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2d</td>
<td>Biomass – As, Pb, Zn, TSP, Cr, DIOX, Cd, PM_{10}</td>
<td>Biomass – As, Pb, Zn, TSP, Cr, DIOX, Cd, PM_{10}, SO\textsubscript{2}, Ni, Hg, PM_{2.5}, NO\textsubscript{x}, NMVOC, Se, Cu</td>
</tr>
<tr>
<td></td>
<td>Liquid – SO\textsubscript{2}, Pb, Ni, NO\textsubscript{x}, Se, As</td>
<td>Liquid – SO\textsubscript{2}, Ni, NO\textsubscript{x}, PM_{10}, PM_{2.5}, Se</td>
</tr>
<tr>
<td>Solid Fuels –</td>
<td></td>
<td>Solid Fuels – SO\textsubscript{2}, Se, Hg, PM_{2.5}</td>
</tr>
<tr>
<td>Other Fuels –</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

3.2.7.2 METHODOLOGICAL ISSUES

The Tier 2 method is used. Emissions from processes in the Pulp, paper and print industry are reported under NFR2D1, see section 4.6 Other industrial processes and product use, NFR2H.

Activity data 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 and later. For more details on these surveys see Annex 2. There is no apparent trend in fuel consumption since 1990.

During 2009, an investigation of emissions of NO\textsubscript{x}, SO\textsubscript{2} and particulate matter from the pulp and paper industry was performed. A comparison between the total emissions from the facilities calculated with national emission factors and the corresponding emissions reported in the environmental reports of the corresponding facilities showed that the use of national emission factors leads to an overestimation of the emissions. In the environmental reports, however, emissions are not reported per fuel type, and hence it was not possible to develop revised emission factors per fuel. Instead, emissions of NO\textsubscript{x}, SO\textsubscript{2} and particulate matter from fuel combustion in pulp and paper production facilities are enumerated with the same mean factors for all fuels:

\[
\text{NO}_x: \quad \text{EM} = 0.736 \times \text{EF(NO}_x) \times \text{AD} \\
\text{SO}_2: \quad \text{EM} = 0.565 \times \text{EF(SO}_2) \times \text{AD} \\
\text{TSP/PM}_{10}/\text{PM}_{2.5}: \quad \text{EM} = 0.686 \times \text{EF(TSP/PM}_{10}/\text{PM}_{2.5}) \times \text{AD};
\]

\text{EM} = \text{emission} \\
\text{EF} = \text{national emission factor, specific for each substance} \\
\text{AD} = \text{activity data in TJ.}

The availability of environmental reports for the years before 2000 is very limited, why the correction factors quoted above are used only for the years 2000 and later. The investigation, and hence the correction factors, applies to the pulp and paper industry only, and not to the printing works. Emissions from combustion of sulphur lyes are presently not reported in 1A2d as this activity has been considered as an industrial process.
3.2.7.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As for NFR1A2 in general, time series consistency despite the changes in activity data source is discussed in Annex 2. 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.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As
- Gaseous fuels for As
- Liquid As, Cd, Cr, Cu, diox, Hg, Ni, Se, Zn
- Other fuels for As, Cr, Cu, Hg, Ni, Pb, Zn
- All fuels for PAH, PM10, PM2.5, TSP

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.7.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

In general, the same QA/QC procedures are used for NFR1A2d as for 1A1a and 1A2a–c described above.

3.2.7.5 SOURCE-SPECIFIC RECALCULATIONS

In submission 2018 a major revision of the emission factors was made within the industrial sector. Briefly, the revision was to a large extent based on conclusions from the previous major emission factor revision that was made within the public electricity and heat production sector (NFR 1A1a) in submission 2017. This is due to the fact that many combustion processes in the industrial sector are similar to those in the heat and power sector. In conclusion, the revision includes several fuel types and recalculation were in general made for the whole time series for all the revised fuel types. The revision is described in more detail in Annex 2.

3.2.7.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

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3.2.8 **Food Processing, Beverages and Tobacco, NFR1A2e**

3.2.8.1 **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 3000 companies, of which only around 650 have more than 10 employees\(^\text{46}\). 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 1A2e). 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. As for NFR 1A2 in general, emissions from companies with less than 10 employees are allocated to NFR 1A2g.

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 key source analysis, NFR1A2e, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2e</td>
<td>Biomass –</td>
<td>Biomass –</td>
</tr>
<tr>
<td></td>
<td>Liquid – Ni</td>
<td>Liquid – Ni, SO(_2), Se, NO(_x)</td>
</tr>
<tr>
<td></td>
<td>Other –</td>
<td>Other –</td>
</tr>
</tbody>
</table>

3.2.8.2 **METHODOLOGICAL ISSUES**

The Tier 2 method is used. 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.

The fuel consumption varies between years and decreased steadily during the years 1998-2008. Since 2008, the total annual fuel consumption in this sector is stable.

3.2.8.3 **UNCERTAINTIES AND TIME-SERIES CONSISTENCY**

As for NFR1A2 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.

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.

\(^\text{46}\) The Swedish Food Federation 2013-10-02
Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As, diox
- Gaseous fuels for As
- Liquid As, Cd, Cr, Cu, Hg, Ni, Pb, Se, Zn
- Other fuels for As, Cr, diox, Cu, Hg, Ni, Pb, Zn
- All fuels for PAH, PM10, PM2.5, TSP

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.8.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
Generally, the same QA/QC procedures are applied for 1A2e as for other 1A2 categories described above.

3.2.8.5 SOURCE-SPECIFIC RECALCULATIONS
In submission 2018 a major revision of the emission factors was made within the industrial sector. Briefly, the revision was to a large extent based on conclusions from the previous major emission factor revision that was made within the public electricity and heat production sector (NFR 1A1a) in submission 2017. This is due to the fact that many combustion processes in the industrial sector are similar to those in the heat and power sector. In conclusion, the revision includes several fuel types and recalculations were in general made for the whole time series for all the revised fuel types. The revision is described in more detail in Annex 2.

3.2.8.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.9 Non-Metallic Minerals, NFR1A2f
3.2.9.1 SOURCE CATEGORY DESCRIPTION
This source category includes stationary combustion of fuels in non-metallic mineral industries (ISIC 26). Cement production accounts for the major part of the emissions. 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.

---

Table 3-10. Summary of key source analysis, NFR1A2f, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2f</td>
<td>Biomass –</td>
<td>Biomass –</td>
</tr>
<tr>
<td></td>
<td>Liquid – Ni, Hg, SO₂, As, Se, NOₓ</td>
<td>Liquid – Hg, As, Ni, NOₓ</td>
</tr>
<tr>
<td></td>
<td>Solid – Pb, Hg, SO₂, As, Se, Cr, Ni, DIOX, Cd, NOₓ</td>
<td>Solid – Pb, Hg, As, Se, Cr, Ni, NOₓ, SO₂</td>
</tr>
<tr>
<td></td>
<td>Other –</td>
<td>Other – NOₓ</td>
</tr>
</tbody>
</table>

3.2.9.2 METHODOLOGICAL ISSUES

Tier 2 method is used for emissions from stationary combustion for NFR1A2f, because country-specific emission factors for the source category and fuel for each gas is used.

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 2008 and later, activity data for the three plants within the cement production industry is taken from the EU ETS system, as this data source provides more detailed information on fuel types. The total amount of fuels combusted is consistent with the quarterly fuel statistics.

National emission factors are used. For more details on these surveys and emission factors see Annex 2.

For practical reasons, SO₂ and NOₓ emission data available from environmental reports are reported in NFR2A7. All other energy related emissions for this facility are reported in NFR1A2f.

Due to confidentiality reasons liquid and solid fuels are reported as C for energy consumption.

3.2.9.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As for NFR1A2 in general, time series are considered consistent despite the changes in activity data source as discussed in Annex 2. The IEFs are slightly variable between years due to variations in the fuel mix.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As
- Gaseous fuels for As
• Liquid As, Cd, Cr, Cu, diox, Hg, Pb, Se, Zn
• Other fuels for As
• All fuels for diox, PM10, PM2.5, TSP

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.9.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
Generally, the same QA/QC procedures are applied for 1A2f as for other 1A2 categories described above. In some earlier submissions, extensive QA/QC and verification efforts have been made for the other sectors including the construction industry.

3.2.9.5 SOURCE-SPECIFIC RECALCULATIONS
In submission 2018 a major revision of the emission factors was made within the industrial sector. Briefly, the revision was to a large extent based on conclusions from the previous major emission factor revision that was made within the public electricity and heat production sector (NFR 1A1a) in submission 2017.\textsuperscript{49} This is due to the fact that many combustion processes in the industrial sector are similar to those in the heat and power sector.\textsuperscript{50} In conclusion, the revision includes several fuel types and recalculations were in general made for the whole time series for all the revised fuel types. The revision is described in more detail in Annex 2.

3.2.9.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.9.7 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been carried out in submission 2017.

\textsuperscript{49} Mawdsley, I., Wisell, T., Stripple, H., Ortiz, C. 2016.
\textsuperscript{50} Mawdsley, I., Stripple, H. 2017.
3.2.10 Other industries, NFR1A2g

3.2.10.1 SOURCE CATEGORY DESCRIPTION

This source category is by nature heterogeneous, as 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 1A2a-1A2f, and stationary combustion within all companies with less than 10 employees regardless of branch, and stationary combustion within the construction sector. The quarterly fuel statistics is a cut-off survey including enterprises with ten or more employees. The estimation of emissions from enterprises with less than ten employees is based on activity data from the annual energy balances, i.e. a model estimate of aggregate fuel consumption in all small enterprises within the entire manufacturing industry. These emissions are reported in 1A2gviii.

The mobile emission source included in this sector is combustion by off-road vehicles and other machinery (working machinery) used in the construction and manufacturing industry. The emissions of air pollution from this sector are reported in 1A2gvii.

In terms of stationary fuel combustion and emissions, two branches of industry are dominating; 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 1A2g in total, it has decreased slightly since 1990. Liquid and biomass fuels account for most of the decrease. For mobile combustion, i.e. working machinery, the fuel consumption for 2016 has nearly doubled since 1990. The emissions of CO, NOx, NMVOC and particles shows however a decreasing trend for the last 2-3 years.

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 3-11.

Table 3-11. Summary of key source analysis, NFR1A2g, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A2g vii</td>
<td>Biomass – Cu</td>
<td>Biomass – Cu</td>
</tr>
<tr>
<td></td>
<td>Liquid – Cu, NOₓ, PM₁₀, PM₂₅, TSP, Cd, CO, SO₂, NMVOC</td>
<td>Liquid – Cu, NOₓ, PM₁₀, PM₂₅, TSP, Cd, CO, SO₂, Ni</td>
</tr>
<tr>
<td></td>
<td>Other –</td>
<td>Other –</td>
</tr>
<tr>
<td>1A2g viii</td>
<td>Biomass – TSP, As, Pb, PM₁₀, PM₂₅, SO₂, Cd, NOₓ, Se, DIOX, Zn, Cr, Hg, Ni</td>
<td>Biomass – TSP, As, Pb, PM₁₀, PM₂₅, Cd, PAH 1-4, Se, DIOX, Zn, Cr, Hg, Ni, NMVOC, NOₓ</td>
</tr>
<tr>
<td></td>
<td>Liquid – As, Ni, SO₂, NOₓ</td>
<td>Liquid – Ni, SO₂, PM₁₀, PM₂₅, Se</td>
</tr>
<tr>
<td>Solid – As, Cr, DIOX, Hg, Ni, NOₓ, Pb, Se, SO₂</td>
<td>Solid – As, CO, DIOX, Hg, Ni, Pb, Se, SO₂</td>
<td></td>
</tr>
</tbody>
</table>
3.2.10.2 METHODOLOGICAL ISSUES

All consumption of motor gasoline and diesel oil, including low blended biofuel, in manufacturing industries and construction is allocated to mobile combustion while all other fuels (heating oils, natural gas etc.) are allocated to stationary combustion.

3.2.10.2.1 Stationary combustion

For emissions from stationary combustion, the Tier 2 method is used. 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 within the manufacturing industry are estimated and reported under NFR1A2g. Activity data are collected from the annual energy balances produced by Statistics Sweden\(^51\). The last emission year is estimated as a projection of the second last year by the trend from the quarterly energy balances, as the annual energy balances for the last emission year are not ready in time for the emission calculations. 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\(^52\) in the same way as for small companies described above.

The fuel consumption varies between years, but has totally decreased slightly since 1990, especially the consumption of liquid and biomass fuels.

Since 2002, for one glassworks plant, it is no longer possible to separate combustion emissions of \(\text{SO}_2\) from process emissions. The reason is that the facility has restructured its environmental report, and only reports emissions of \(\text{SO}_2\) on an aggregate level. The median value for the share of process-related \(\text{SO}_2\) missions of the total \(\text{SO}_2\) emissions is 2% for the years 1990 - 2001. The emission data reported in the plants environmental report are considered to be more accurate than emissions calculated from fuel combustion with standard emission factors, and thus for practical reasons, all data that is available from environmental reports from this plant, namely \(\text{SO}_2\) and \(\text{NO}_X\), are reported in NFR2A7 and all other emissions are reported in NFR1A2F.

For 2008 and later, activity data for the three plants within the cement production industry is taken from the EU ETS system because the reporting of waste-like fuels


\(^{52}\) Statistics Sweden, EN20SM 1990-2008. See also Annex 2.
to the energy statistics has shown to be partly incomplete for some years. In the CLRTAP stage 3 review of submission 2013 (and in earlier reviews) it was recommended that the emissions from the cement industry within NFR1A2f should be reported separately. This is however not possible, because data on emissions of NO\(_X\), SO\(_2\) and particulate matter from the cement industry are taken from environmental reports. In these reports, only the total emissions for each substance are reported, and it is not possible to isolate the combustion emissions. Because of this, all emissions of these substances from the cement industry are reported in NFR2A1. This means that the emissions reported under NFR1A2g in the NFR tables do not include combustion emissions from the cement industry. A table for the cement industry would hence show “IE” for the major pollutants.

Due to confidentiality reasons liquid, solid and biomass fuels are reported as C for energy consumption.

3.2.10.2.2 Mobile combustion

Emissions from mobile combustion in NFR1A2g ii refer to working machinery used in industry, including for example tractors, dumpers, cranes, excavators, generators and wheel loaders. A national model is used to estimate emissions from all working machinery used in Sweden and it is considered to correspond to Tier 3 for all emissions, except for SO\(_2\) which is estimated according to Tier 2. The model is further explained in Annex 2.

Emissions from working machinery are also reported in NFR1A3e ii, 1A4a ii, 1A4b ii and 1A4c ii. See Table 3-12.

Table 3-12. Distribution of emissions from off-road vehicles and other machinery.

<table>
<thead>
<tr>
<th>Category</th>
<th>NFR</th>
<th>Definition IPCC Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>1A2g vii</td>
<td>Mobile machineries in industry that run on petroleum fuels, as for example tractors, dumpers,</td>
</tr>
<tr>
<td></td>
<td></td>
<td>cranes, excavators, generators, wheel loaders, sorting works, pump units etc.</td>
</tr>
<tr>
<td>Other</td>
<td>1A3e ii</td>
<td>Combustion emissions from all remaining transport activities including ground activities</td>
</tr>
<tr>
<td></td>
<td></td>
<td>in airports and harbours, and off-road activities not otherwise reported under 1A4c ii or</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1A2g vii.</td>
</tr>
<tr>
<td>Commercial/</td>
<td>1A4a ii</td>
<td>Garden machinery, e.g. lawn mowers and clearing saws, not used by private users, Also tractors</td>
</tr>
<tr>
<td>Institutional</td>
<td></td>
<td>not used in industry ore forestry or agriculture.</td>
</tr>
<tr>
<td>Residential</td>
<td>1A4b ii</td>
<td>All emissions from mobile fuel combustion in households, as for example tractors, lawn</td>
</tr>
<tr>
<td></td>
<td></td>
<td>movers, snowmobiles, forklifts, trimmers, chainsaws and forklifts</td>
</tr>
<tr>
<td>Agriculture,</td>
<td>1A4c ii</td>
<td>Emissions from mobile fuel combustion in agriculture and forestry, as for example loader-</td>
</tr>
<tr>
<td>Forestry</td>
<td></td>
<td>excavator, tractor, harvester, clearing saw etc. High-way agricultural transportation is</td>
</tr>
<tr>
<td></td>
<td></td>
<td>excluded.</td>
</tr>
</tbody>
</table>
3.2.10.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

3.2.10.3.1 Stationary combustion
As for NFR1A2 in general, time series consistency despite the changes in activity data source is discussed in Annex 2. As for other categories in NFR1A2, the IEFs 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.

Uncertainties for activity data and emission factors are generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As
- Gaseous fuels for As
- Liquid As, Cd, Cr, Cu, diox, Hg, Pb, Se, Zn
- Other fuels for As
- All fuels for diox, PM10, PM2.5, TSP

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.10.3.2 Mobile combustion
No recalculations have been performed for emissions in the mobile sector for the eighties, and thus there are inconsistencies in time series between the eighties and later years. Time series from mobile combustion in NFR1A4b ii have been reviewed for later years and are considered to be consistent.

Uncertainties for activity data and emissions reported for working machinery in NFR1A4b ii can be seen in Annex 1 to the IIR.

3.2.10.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

3.2.10.5 SOURCE-SPECIFIC RECALCULATIONS
The model for working machinery has been updated with sales data for off-road vehicles for 2005-2016 in submission 2018, which was provided by the Swedish trade association for suppliers of mobile machines. The allocation key in the model was also updated, as new information regarding the sector distribution was received from both the trade association and the vehicle register. The age of some of
the oldest tractors in the model was adjusted as well, as they in previous submis-
sions mistakenly were grouped together in the oldest age category\textsuperscript{53}.

The updated model has resulted in a slightly decreased consumption of diesel (in-
cluding low blended biofuel) but an increased consumption of gasoline (including
low blended biofuel) for most years, in submission 2018 compared to submission
2017, for working machineries used in the construction and manufacturing indus-
try. The emissions from working machinery used in the industrial sector have in-
creased since the late 1990s, on a total level, in submission 2018 compared to sub-

In submission 2018 a major revision of the emission factors for stationary combus-
tion within this code was made within the industrial sector for stationary combus-
tion. Briefly, the revision was to a large extent based on conclusions from the pre-
vious major emission factor revision that was made within the public electricity
and heat production sector (NFR 1A1a) in submission 2017\textsuperscript{54}. This is due to the
fact that many combustion processes in the industrial sector are similar to those in
the heat and power sector\textsuperscript{55}. In conclusion, the revision includes several fuel types
and recalculations were in general made for the whole time series for all the revised
fuel types. The revision is described in more detail in Annex 2.

3.2.10.6 \hspace{1em} SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

3.2.11 \hspace{1em} Civil Aviation, NFR1A3a i-ii

3.2.11.1 \hspace{1em} SOURCE CATEGORY DESCRIPTION

Domestic Aviation is defined as emissions from flights that depart and arrive in the
same country. But for the reporting of air pollution according to the Long-Range
Transboundary Air Pollution Convention (LRTAP), emissions from national and
international aviation during the LTO cycle\textsuperscript{56} both belong to the national totals.
And emissions from both domestic and international aviation during the Cruise cy-
acle\textsuperscript{57} is reported separately as memo items and not included in national totals.

The national government administers 13 of 40 airports with regular and/or charted
air traffic in Sweden, for which activity data is provided at present. The remaining
27 airports are private and/or administered by local governments\textsuperscript{58}.

\textsuperscript{53}Eklund, V., Lidén, M., Jerksjö, M., 2017. Regelbunden indataförsörjning till beräkningsmodellen för
arbetsmaskiner. SMED PM 2017.

\textsuperscript{54}Mawdsley, I., Wisell, T., Stripple, H., Ortiz, C. 2016.

\textsuperscript{55}Mawdsley, I., Stripple, H. 2017.

\textsuperscript{56}Landing and take-off.

\textsuperscript{57}Cruise cycle: above 3000 feet.

\textsuperscript{58}Transportstyrelsen, 2013.
The energy consumption from the LTO cycle has increased by 3 % between 1990 and 2016 and by 13 % since 2015, whereas the energy consumption from the cruise cycle has increased by 62 % since 1990 and by 15 % since 2014.

The emissions of NO\textsubscript{X} from both the Cruise and LTO cycle have fluctuated over the years, but while the emissions NO\textsubscript{X} from the Cruise cycle have increased by 46% since 1990 have the emissions of NO\textsubscript{X} from LTO decreased by ~17% since 1990. The emissions of NO\textsubscript{X} increased from both Cruise and LTO cycle by 16 respectively 14 % between 2015 and 2016. The emissions of particles have also increased in the last year, but while the Cruise cycle have an increasing trend, the LTO cycle have a mildly decreasing trend.

The emissions of CO from Cruise reached their lowest in 2003 and then started to rise again. This also applies to CO from LTO, but these emissions fluctuate more. The emissions of CO from Cruise have increased by 5% since 1990 and by 11% in the last year, while the emissions of CO from the LTO phase have increased by 7% since 1990 and by 10% between 2015 and 2016.

The emissions of NMVOC from the Cruise phase have had a decreasing trend for the whole time series, with a couple of exceptions, and have dropped by 60% since 1990. The trend was however broken in the last year, when the emissions increased by 14%. The emissions of NMVOC from the LTO phase have fluctuated a lot for the whole time series and decreased by 17% since 1990 whereas they increased by 12% in the last year. The declining emissions of NMVOC in recent years is the result of phasing out a specific type of airplane (MD-80/82), which is a major contributor to these gases.
Table 3-13. Summary of key source analysis, NFR1A3a i-ii, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A3a</td>
<td>Aviation Gasoline - Pb</td>
<td>Aviation Gasoline - Pb</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Jet Kerosene – NOx, SO2</td>
<td>Jet Kerosene – NOx, SO2</td>
<td></td>
</tr>
</tbody>
</table>

3.2.11.2 METHODOLOGICAL ISSUES
Sweden uses Tier 1 to estimate emissions of SO2, Tier 2 to estimate the emissions of NMVOC and Tier 3a to estimate the emissions of all other gases. Emissions from aviation used by the agricultural and forestry sectors are reported as civil aviation. Emissions from military use of aviation fuels are reported under Other – mobile sources (NFR1A5b) or 1D2 Military operations abroad.

The Swedish Transport Agency (STAg) is responsible for reporting the emissions from aviation. But the fuel consumption and emissions published by STAg are calculated by the Swedish Defence Research Agency (FOI) by using an air emission model. STAg provides FOI statistics for the model regarding:

- Airport of departure and arrival
- Type of aircraft
- Number of flights
- Number of LTO cycles
- Number of passengers

A database with information regarding 200 different types of aircraft is also used. The emission data regarding different types of aircraft in the database originates from “ICAO Engine Exhaust Emission Data Bank”. All this data is used to calculate emissions and the amount of combusted fuel for the whole flights as well as for aircraft movements below 3000 feet at the airports, the so called LTO cycle. FOI has written a report which describes their method for estimating the emission from aviation59.

The model used to calculate the emissions from aviation underestimates the number of kilometres flown, as the model uses more direct flight routes in the calculations than the aircrafts do in reality. As a result, the consumption of fuel and emissions are underestimated, and need to be adjusted to be in line with data on national delivery of aviation fuel from the monthly survey on supply and delivery of petroleum products from Statistics Sweden (see Annex 2). The results from the emission calculations are aggregated into four groups: domestic landing and take-off (LTO), domestic cruise, international LTO and international cruise.

59 Calculation of exhaust emissions from air traffic. T. Mårtensson, A. Hasselrot. FOI R 3677 mSE
The methodology for calculating national emissions is the same for all years with a few exceptions for earlier years. All emissions for 1990-1994 were calculated by SMED in cooperation with the STAg 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 NOₓ were calculated in a similar way.

The mean value of the ratio between NOₓ 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, NOₓ, CO and HC are estimated by FOI as described above. The emissions of NMVOC and CH₄ are estimated based on the model estimated emissions of HC and emission factors from EMEP/EEA air pollutant emission inventory guidebook. N₂O emissions from LTO are estimated using information from STAg on the number of LTO cycles together with emission factors from EMEP/EEA guidebook. Emissions of N₂O from cruise are based on emission factors from EMEP/EEA guidebook together with the fuel consumption for cruise activities as well as emissions of particles.

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. For international flights, ICAO standard taxi time has been used for the part of the LTO cycle occurring on international airports.

The Swedish Transport Agency (STAg) 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.

In 2006, the STAg 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 STAg 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.

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60 Gustafsson, 2005.
61 Näs, 2005.
3.2.11.3  UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Data on domestic and international bunker fuel in the Monthly fuel, gas and inventory statistics has been found to be of good quality (See Annex 1 to IIR for more information). Regarding time-series consistency, see the Methodology section.

3.2.11.4  SOURCE-SPECIFIC QA/QC AND VERIFICATION
The activity data has been subject to QA/QC procedures.

3.2.11.5  SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculation was made in submission 2018.

3.2.11.6  SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.12  Road transport, NFR1A3b i-iv
3.2.12.1  SOURCE CATEGORY DESCRIPTION
Road traffic is the dominating mode for transport of both goods and people and the Swedish citizens travelled more than 67,000 millions of km by car in 2016. This is an increase by approximately 20 % since 1990. For all trucks (LCV & HGV) there has been an increase in number of travelled km by 83 % since 1990, while buses drive nearly the same number of km in 2016 as in 1990 (Table -3-14). The Swedish road network comprises of around 140,700 km of public roads, whereof 2,660 km freeways and motor-traffic roads.


<table>
<thead>
<tr>
<th>Year</th>
<th>Passenger cars</th>
<th>LCV</th>
<th>HGV</th>
<th>Motorcycles</th>
<th>Buses</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>55 696</td>
<td>3 709</td>
<td>3 653</td>
<td>407</td>
<td>964</td>
</tr>
<tr>
<td>2000</td>
<td>58 555</td>
<td>4 574</td>
<td>4 310</td>
<td>670</td>
<td>976</td>
</tr>
<tr>
<td>2016</td>
<td>67 071</td>
<td>8 781</td>
<td>4 716</td>
<td>798</td>
<td>979</td>
</tr>
</tbody>
</table>

Road transport includes five vehicle categories: passenger cars, light commercial vehicles (LCV), heavy goods vehicles (HGV), buses and mopeds & motorcycles. The total number of trucks and passenger cars (in traffic) has increased by 100 % respectively 32 % since 1990 in contrast to the emissions of NO\textsubscript{X} and NMVOC (from the same vehicles), where the emissions of NMVOC have decreased by 93 % by all vehicles while the emissions of NO\textsubscript{X} have decreased 64 % by trucks and 68 %

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62 Ministry of the Environment, 2001
64 Data from the road emission model HBEFA (Trafikverket).
% by passenger cars. This is mainly a result of the introduction of three-way catalytic converters on passenger cars and trucks. Gasoline evaporation, Automobile tyre and brake wear and Automobile road abrasion are, beside combustion of fuel, also sources of air pollution caused by road traffic.

The emissions of CO also show a downward trend for all years (1990-2016) as well as the emissions of SO₂, as the sulphur content in fuel has been heavily restricted over the years. The emissions of NH₃ showed an increasing trend until 2001 but have decreased steadily since then.

Gasoline has previously been the most common fuel used for road transportation, but as from 2011 the amount of diesel used by road traffic as well as the emissions of GHG from diesel surpassed gasoline. The increasing consumption of diesel by road traffic is primarily explained by a shift from gasoline cars to diesel cars, but also by an increased consumption of diesel by HGV and LCV. The total consumption of diesel by HGV and LCV correspond to 52 % of the total consumption of diesel by road traffic in 2016, while passenger cars consume around 42 % of the total diesel for road traffic.

The consumption of diesel by HGV increased by 59 % between 1990 and 2007, from an already high level, and then started to decrease (Figure 3-2). The consumption of diesel by HGV was 15 % higher in 2016 than in 1990. The consumption of diesel by LCV has increased by 719 % since 1990, levelled out in 2011 and started to decrease slightly in subsequent years. The diesel consumption by LCV is at a lower level than both passenger cars and HGV. The consumption of diesel by passenger cars has increased steadily since 1990 but showed a 3 % decrease between 2015 and 2016.
The total use of liquid biofuels (FAME and ethanol) has increased by more than a factor of eight since 2003, when large-scale blending of ethanol into petrol began. The increasing production and use of biofuels was initiated by advantageous policy regulations and tax reliefs for biofuels\textsuperscript{66}. The amount of biogas used by road traffic has also increased greatly since it was introduced on the market and has doubled every other to every third year between 1998 and 2008.

The main part of ethanol used by road transportation in Sweden is used as a blending component for gasoline. Large-scale blending of ethanol into petrol began in 2003 and the total amount of ethanol used for road traffic nearly tripled between 2003 and 2011. As from 2012, the amount of low-blended ethanol in gasoline started to decline as a result of the shift from gasoline cars to diesel cars. Today, just about all petrol sold in Sweden contains around 5% ethanol.

Ethanol is also used by ethanol buses and by E85 passenger cars (flexi fuel cars). The ethanol used by E85 cars and by buses, increased steadily until 2011 respectively 2012, when the trend turned downward again (Figure 3-3).

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\textsuperscript{66} Swedish Energy Agency, 2013.
Large-scale blending of FAME into diesel began in 2007/2008 and has increased steadily ever since (Figure 3-4). The total use of FAME by road traffic has increased by 33-43% each year starting 2011. Between 2015 and 2016, the use of FAME by road traffic showed a 49% increase. This is mainly a result of a growing trend for diesel cars and a rising fraction of FAME blended into diesel.
Table 3-15. Summary of key source analysis, NFR1A3b i-iv, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A3b i</td>
<td>Biomass – Cd, Zn, CO, NMVOC, NOx</td>
<td>Biomass – Cd, Zn, Hg, CO, NMVOC, NOx, SO2</td>
</tr>
<tr>
<td></td>
<td>Diesel oil – Hg, Cr, Cd, Zn, PM2.5, PM10</td>
<td>Diesel oil – Hg, Cr, Cd, Zn</td>
</tr>
<tr>
<td></td>
<td>Gasoline – Pb, NH3, Hg, DIOX, Cd, Zn, Cr, PM2.5</td>
<td>Gasoline – Pb, NH3, Hg, DIOX, Cd, Cr, PM10, PM2.5, TSP</td>
</tr>
<tr>
<td></td>
<td>Gasoline evaporation -</td>
<td>Gasoline evaporation – NMVOC</td>
</tr>
<tr>
<td>1A3b ii</td>
<td>Biomass –</td>
<td>Biomass –</td>
</tr>
<tr>
<td></td>
<td>Diesel oil – NOx, Cd, Zn, PM2.5, PM10</td>
<td>Diesel oil – NOx, Hg, Cd, Zn</td>
</tr>
<tr>
<td></td>
<td>Gasoline –</td>
<td>Gasoline – CO, NOx, Pb, NMVOC</td>
</tr>
<tr>
<td>1A3b iii</td>
<td>Biomass – Cd, Hg, NOx, Zn</td>
<td>Biomass – Zn, Cd, Cr, Hg, NOx, PM2.5, PM10, SO2, TSP</td>
</tr>
<tr>
<td></td>
<td>Diesel oil – NOx, Cr, Cd, PM2.5, Hg, PM10, TSP, CO, Zn</td>
<td>Diesel oil - NOx, Cr, Cd, PM2.5, Hg, PM10, TSP, SO2, Zn, NMVOC</td>
</tr>
<tr>
<td></td>
<td>Gaseous Fuels –</td>
<td>Gaseous Fuels –</td>
</tr>
<tr>
<td>1A3b iv</td>
<td>Gasoline – CO</td>
<td>Gasoline –</td>
</tr>
</tbody>
</table>

3.2.12.2 METHODOLOGICAL ISSUES

The road emission model HBEFA is used by the Swedish Transport Administration (STA) to estimate the fuel consumption and different emissions from road traffic. The fuel consumption is adjusted to be in line with national statistics on supply and delivery of petroleum products. The fuel consumption and emissions are allocated by fuel type and five vehicle categories: Passenger cars, Light commercial vehicles (LCV), Heavy goods vehicles (HGV), Buses and Mopeds & Motorcycles. The road traffic emission model HBEFA is updated yearly with new information regarding emission factors, vehicle fleet, composition of the fuel and the current traffic work. The HBEFA model is administrated by The Swedish National Road and Transport Research Institute (VTI) and the Swedish Transport Administration (STA).

Emissions of SO2 are based on the fuel consumption per vehicle type and country specific thermal values and emission factors. The emission factors for SO2 are based on the actual sulphur content for the different environmental classes of petrol and diesel fuel as from submission 2007. The thermal values and the country-specific emission factors for SO2 are provided by SPBI. 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.

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68 Swedish petroleum and biofuel institute
The emissions of CO, NO\textsubscript{X}, NMVOC, NH\textsubscript{3}, Pb and particles from road traffic are estimated by HBEFA. The emissions of Cd, Hg, As, Cr, Cu, Ni, Se and Zn from combustion were estimated for the first time in submission 2016, with default emission factors from EMEP/EEA Guidebook 2013.

Activity data for natural gas is available from 1990, while reliable activity data for biogas exists from 1996 and for ethanol and FAME from 1998. Thermal values for biogas have been collected from the Swedish Biogas Association, for ethanol from SPBI and for FAME from the “Eco fuel” web page\textsuperscript{69}.

The bottom-up estimations of the fuel consumption by the HBEFA model differ slightly from those reported to the UNFCCC (based on fuel delivery). According to IPCC Guidelines, the inventory should only account for emissions from fuel purchased in Sweden compared to the STA, who aims to describe what is emitted on Swedish roads, regardless of where the fuel was bought or the nationality of the vehicles. An overview of the two different objectives is presented in Table 3-16.

\textsuperscript{69} \url{http://www.ecobransle.se/}
Table 3.16. Emissions from road transport reported by the STA and in the NFR.

<table>
<thead>
<tr>
<th>Fuel bought in</th>
<th>Traffic on Swedish roads</th>
<th>Traffic in Sweden, not on roads</th>
<th>Traffic to/from other country</th>
<th>Traffic in other countries</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sweden</td>
<td>NFR1A3b i-iv STA</td>
<td>NFR1A3b i-iv</td>
<td>NFR1A3b i-iv* STA to the Swedish border</td>
<td>NFR1A3b i-iv*</td>
</tr>
<tr>
<td>Other country</td>
<td>STA</td>
<td>Not reported</td>
<td>STA to the Swedish border</td>
<td>Not reported</td>
</tr>
</tbody>
</table>

* 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 NFR1A3b.

Military transport emissions are reported under NFR1A5b. Military road transport is included in the road traffic emissions estimated by HBEFA. To subtract and separate emissions from military transport from emissions from civil road transport, emissions from HBEFA 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 the equation below:

\[ A = B - \sum \left( \frac{C-D}{C \cdot E_i} \right) \]

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

Data on particle emissions are lacking for the years 1981-1984 and 1986 and are therefore interpolated. It should also be noted that emissions of particles reported under NFR1A3b for the 1980s include military activities.

Emissions of dioxin from road transport were before submission 2008 not separated into different sub-sectors and were aggregated under Road Transportation (1A3b). In submission 2008 data from 1990 were updated since detailed background information (m3 gasoline and diesel) per vehicle category from the HBEFA model, has made it possible to report emitted dioxin separately for Passenger cars (1A3b i), Light duty vehicles (1A3b ii), Heavy duty vehicles (1A3b iii) and Mopeds & Motorcycles (1A3b iv). From submission 2012, data from the HBEFA 3.1 model were used instead of data from the ARTEMIS model. The emission factors used are from Finstad et al (2001)\(^7\).

Further, as from submission 2008 emissions of benzo(a)pyrene, benzo(b)fluoranthene, benzo(k)fluoranthene and indeno(1,2,3-cd)pyrene were reported for 1A3b.

\(^7\) Finstad et al, 2001
As for dioxin, detailed data from the ARTEMIS model (submission 2008-2011) and HBEFA 3.1 (from submission 2012) regarding yearly mileages (km x 10^6) per vehicle and fuel category, as well as emission factors, have been used. Emission factors presented by Westerholm et al. (2001) are used for the calculations of PAHs from Swedish environmental classified diesel (MK1) used in Heavy duty vehicles. Emission factors for MK1 diesel in Passenger cars and Light duty vehicles are calculated using the relationship Passenger car/Heavy duty vehicle and Light duty vehicle/Heavy duty vehicle in the EMEP-Corinair Guidebook and emission factors for Heavy duty vehicle according to Westerholm et al (2001). For MK1 diesel the emissions of benzo(k)fluoranthene are included in reported benzo(b)fluoranthene. All other emission estimates are based on emission factors in the EMEP-Corinair Guidebook. The emission factors used are shown in Table 3-17.

Table 3-17. Emission factors used for estimations of dioxin and PAH emissions from fuel combustion in NFR1 A 3 b i - iv.

<table>
<thead>
<tr>
<th></th>
<th>Gasoline</th>
<th>Diesel</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Leaded</td>
<td>Un-leaded</td>
</tr>
<tr>
<td></td>
<td>Passenger cars, light duty vehicles</td>
<td>Heavy duty vehicles</td>
</tr>
<tr>
<td></td>
<td>Not DI</td>
<td>Direct injection</td>
</tr>
<tr>
<td>Dioxin (µg/Mg)</td>
<td>2.00</td>
<td>0.10</td>
</tr>
<tr>
<td>Benzo(a)pyrene (µg/km)</td>
<td>0.48</td>
<td>0.32</td>
</tr>
<tr>
<td>Benzo(b)fluoranthene (µg/km)</td>
<td>0.88</td>
<td>0.36</td>
</tr>
<tr>
<td>Benzo(k)fluoranthene (µg/km)</td>
<td>0.30</td>
<td>0.26</td>
</tr>
<tr>
<td>Indeno(1,2,3-cd)pyrene (µg/km)</td>
<td>1.03</td>
<td>0.39</td>
</tr>
</tbody>
</table>

Time series per vehicle category are calculated for dioxin and PAH-4 from 1980, but data for 1980 - 1989 are not updated in the NFR-tables.

For the dioxin emission estimates the share of leaded gasoline of the total amount of gasoline must be taken into consideration (Table 3-18). For the PAH calculations the share of diesel Passenger cars and diesel Light duty vehicles with direct injection must be estimated, since these emission factors differ from the emission factors for diesel vehicles without direct injection (Table 3-18). All Heavy duty vehicles are assumed to have direct injection. Also the share of MK1 diesel of the total amount of diesel used has to be known (Table 3-18).

71 Westerholm et al., 2001. Comparison of Exhaust Emissions from Swedish Environmental Classified Diesel Fuel (MK1) and European Program on Emissions, Fuels and Engine Technologies (EPEFE) Reference Fuel: A Chemical and Biological Characterization, with Viewpoints on Cancer Risk
### Table 3-18. Distribution of vehicles with respect to fuel type and injection system.

<table>
<thead>
<tr>
<th>Year-1980</th>
<th>Gasoline</th>
<th>Without direct injection</th>
<th>With direct injection</th>
<th>Diesel type</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Leaded</td>
<td>Un-leaded</td>
<td>Light duty vehicles</td>
<td>Light duty vehicles</td>
</tr>
<tr>
<td>1980-1985</td>
<td>100%</td>
<td>0%</td>
<td>100%</td>
<td>0%</td>
</tr>
<tr>
<td>1990</td>
<td>50%</td>
<td>50%</td>
<td>100%</td>
<td>0%</td>
</tr>
<tr>
<td>1995</td>
<td>0%</td>
<td>100%</td>
<td>67%</td>
<td>100%</td>
</tr>
<tr>
<td>2000</td>
<td>0%</td>
<td>100%</td>
<td>30%</td>
<td>45%</td>
</tr>
<tr>
<td>2005</td>
<td>0%</td>
<td>100%</td>
<td>6%</td>
<td>20%</td>
</tr>
<tr>
<td>2010</td>
<td>0%</td>
<td>100%</td>
<td>0%</td>
<td>4%</td>
</tr>
<tr>
<td>2013</td>
<td>0%</td>
<td>100%</td>
<td>0%</td>
<td>0%</td>
</tr>
</tbody>
</table>

#### 3.2.12.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

For the energy sector, the largest uncertainties come from activity data for the 1980's and from emission factors. No recalculations have been performed for emissions in the mobile sector for the eighties, and thus there are inconsistencies in time series between the eighties and later years.

Activity data for gasoline, diesel and natural gas is available from 1990, while reliable activity data for biogas exists from 1996, for ethanol from 1998 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 2015 has driven the most part during 2014. 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. See Annex 1 for more details regarding uncertainties for activity data and emissions.

#### 3.2.12.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

All Tier 1 general inventory level QC procedures and all QC procedures applicable to this sector are used. The activity data has been subject to QA/QC procedures. 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 NFR-code and checked against the emission factors to make sure that no calculation errors have occurred when emissions were computed.
3.2.12.5 SOURCE-SPECIFIC RECALCULATIONS

The HBEFA model is updated annually regarding vehicle fleet composition, fuel composition, emission factors etc. The estimated consumption of gasoline and diesel by HBEFA was modified for all years by a residual of gasoline and diesel. The residual is the difference in the gasoline respectively the diesel consumption when comparing the national statistics on supply and delivery of petroleum products (top-down approach) to the bottom-up estimated fuel consumption. The residual is proportionally redistributed to the following sectors: to 1A3b (road transportation), 1A3d (domestic navigation), 1A2g vii, 1A3c, 1A4b and 1A4c (off-road vehicles and working machinery) and 1A4c (fishing). Beside these yearly updates, the following adjustments to the model has also taken place in submission 2018:

- The HBEFA model version 3.3 has implemented updated emission factors for NOx regarding Euro IV and Euro VI passenger cars. The transition to the new model with updated emission factors for Euro 4-6 passenger cars leads to higher emissions of nitrogen oxides and the difference is increasing with time. Emissions of NMVOC from gas buses is also included in the HBEFA 3.3 model.
- A reallocation of both biogas and natural gas from passenger cars to buses has been implemented in HBEFA in submission 2018. This applies to year 1995, 1999 and onwards and leads to decreased emissions for gas fueled passenger cars and increased emissions for gas fueled buses.
- Data on biogas and natural gas consumption has been revised for gas fueled passenger cars. There is in submission 2018 data on consumption for the years 1995, 1999 and onwards, whereas submission 2017 showed consumption for the years 1997-2015. Moreover, the delivered quantities of natural gas and biogas were also revised for the year 2015 in submission 2018.

3.2.12.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

3.2.13 Automobile tyre and brake wear, NFR1A3b vi, and automobile road abrasion, NFR1A3b vii

3.2.13.1 SOURCE CATEGORY DESCRIPTION

TSP emissions from tyre and brake wear have since 1990 increased by 28 %. Emissions from tyre and brake wear are heavily dependent on the total amount of vehicle driven kilometres which has also increased with 28 %.

Table 3-19. Summary of key source analysis, NFR1A3b vi-vii, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
</tr>
</thead>
</table>

72 (Statistic Sweden. Data from Monthly fuel, gas and inventory statistics) http://www.scb.se/. See Annex 2. chapter "1.4 Allocation of fuels for mobile combustion" for more information regarding the allocation of fuels for mobile combustion.
### 3.2.13.2 METHODOLOGICAL ISSUES

From submission 2017 and onwards, a national model is used to estimate emissions from tyre and brake wear and road abrasion. The national model takes into account several factors that have not been considered in previous estimates, such as meteorology, regional variations and measured PM$_{10}$ concentrations in urban street canyons for model verification. The new method has the advantage in that the emissions are calculated separately for eight regions in Sweden. Hence, the variability in both meteorological conditions and in the use of studded tyres is taken into account.

Activity data is obtained from the national model, divided by eight regions, and adjusted to fit activity data from the HBEFA 3.1 model.

#### 3.2.13.2.1 Particle emissions from Tyre and brake wear and road abrasion, NFR1A3b vi and 1A3bvii

Emission factors for PM$_{10}$ for the years 2008-2014 have been calculated using the national model, for previous years and following years, averages of 2008-2014 have been used. TSP and PM$_{2.5}$ emissions have been estimated based on the PM$_{10}$ emission factor according to the EMEP/EEA Guidebook 2013 and other literature. As the modelled PM$_{10}$ emissions include both tyre and brake wear and road abrasion, the emission sources have been separated by assuming a constant emission factor for tyre and brake wear of 10 mg vkm$^{-1}$, according to literature values.

TSP emissions are separated further into tyre wear and brake wear and passenger cars, light duty vehicles, mopeds and motorcycles, based on the time series from previous years (1990-2014).

#### 3.2.13.2.2 PAH emissions from Tyre and brake wear, NFR1A3b vi

The separation between particles from tyre wear and brake wear also makes it possible to calculate and report PAH-4 from tyre wear and brake wear in 1A3b vi. The emission factors used for the calculations of PAH emissions from tyre wear and brake wear are as presented in the EMEP/EEA air pollutant emission inventory guidebook (detailed methodology). The emission factors used for the PAH calculations are presented in Table 3-20.

Table 3-20 Emission factors used for PAH emission calculations in 1A3b vi.

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3.2.13.2.3 Metal emissions from Tyre and brake wear, NFR1A3b vi

In Hjortenkrans et al. (2006)\textsuperscript{74} mean metal concentrations in retread and non-re-tread tyre tread rubber are presented. As almost all tyres used on heavy duty vehicles are retread tyre tread rubber\textsuperscript{74} and emissions calculated for heavy duty vehicles are based on an emission factor representing retread tyre tread rubber. For all other vehicle categories the calculations are based on emission factors for non retread tyre tread rubber (Table 3-21).

Table 3-21. Emission factors used for metal emission calculations from tyre wear in 1A3b vi.

<table>
<thead>
<tr>
<th>Vehicle category</th>
<th>Emission factors, TYRE WEAR, (mg/kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pb</td>
</tr>
<tr>
<td>Passenger cars</td>
<td>9.4</td>
</tr>
<tr>
<td>Light duty vehicles</td>
<td>9.4</td>
</tr>
<tr>
<td>Heavy duty vehicles*</td>
<td>9.5</td>
</tr>
<tr>
<td>Mopeds &amp; Motorcycles</td>
<td>9.4</td>
</tr>
</tbody>
</table>

* retread tyres

Hjortenkrans et al. (2006)\textsuperscript{75} also studied the metal content in both branded brake linings and those from independent suppliers. A similar study was made in the late 1990s\textsuperscript{76} and the results show that there is a clear reduction of the Pb and Zn content in both branded linings and linings from independent suppliers. Also for Cu the metal content in linings from independent suppliers from 2005\textsuperscript{77} is much lower.

\textsuperscript{74} Hjortenkrans et al. 2006. Metallemission från trafiken i Stockholm – Däck. (in Swedish, results from a study on metal emissions from tyre wear)

\textsuperscript{75} Hjortenkrans et al. 2006. Metallemission från trafiken i Stockholm – Bromsbelägg. (in Swedish, results from a study on metal emissions from brake linings)


\textsuperscript{77} Hjortenkrans et al. 2006. Metallemission från trafiken i Stockholm – Bromsbelägg. (in Swedish, results from a study on metal emissions from brake linings)
than in 1998\textsuperscript{78}. For branded linings the results is the contrary, the Cu content in linings from 2005 is higher compared to linings from 1998. For the brake linings metal emission calculations the same assumption as both Hjortenkrans\textsuperscript{77} and Westerlund\textsuperscript{78} is made; 40\% of the traffic volume is related to new vehicles using branded linings and 60\% to older vehicles using linings from independent suppliers. For Pb, Cu and Zn the emission factors used are based on results presented by Westerlund\textsuperscript{78} for the years 1980 - 1998, and on results presented by Hjortenkrans\textsuperscript{77} for 2005 and onwards. The emission factors for 1999 - 2004 are interpolated. For Cd the same emission factor is set for the whole time series\textsuperscript{75}.

The emission factors used for calculating metal emissions from tyre wear and brake wear are presented in Table 3-22.

Table 3-1. Emission factors used for metal emission calculations from brake wear in 1A3b vi.

<table>
<thead>
<tr>
<th>Vehicle category</th>
<th>Emission factors, BRAKE WEAR, branded (ppm wt.)</th>
<th>Emission factors, BRAKE WEAR, independent (ppm wt.)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Pb</td>
<td>Cd</td>
</tr>
<tr>
<td>1980 - 1998</td>
<td>13854</td>
<td>2.6</td>
</tr>
<tr>
<td>1999</td>
<td>12090</td>
<td>2.6</td>
</tr>
<tr>
<td>2000</td>
<td>10327</td>
<td>2.6</td>
</tr>
<tr>
<td>2001</td>
<td>8563</td>
<td>2.6</td>
</tr>
<tr>
<td>2002</td>
<td>6800</td>
<td>2.6</td>
</tr>
<tr>
<td>2003</td>
<td>5037</td>
<td>2.6</td>
</tr>
<tr>
<td>2004</td>
<td>3273</td>
<td>2.6</td>
</tr>
<tr>
<td>2005 -</td>
<td>1510</td>
<td>2.6</td>
</tr>
</tbody>
</table>

3.2.13.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The same method is used throughout the time series, which is considered consistent. Uncertainties for activity data and emissions reported for automobile tyre and brake wear (NFR 1A3b vi) and automobile abrasion (NFR 1A3b vii) can be seen in Annex 1 to the IIR.

3.2.13.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

3.2.13.5 SOURCE-SPECIFIC RECALCULATIONS

Due to the implementation of a national model, all emissions are updated throughout the whole time series. The emissions of TSP have increased with about 95\%, PM\textsubscript{10} emissions with about 87\% and PM\textsubscript{2.5} emissions with about 94\%.

3.2.13.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No planned improvements for this category.

3.2.14 Railways, NFR1A3c

3.2.14.1 SOURCE CATEGORY DESCRIPTION

The majority of all railway traffic in Sweden runs on electricity. Only a small part runs on diesel fuel and the emissions related to the use of electricity for railway should not be included in this sector. Production of electricity is accounted for in NFR1A1A, regardless of where it’s consumed. The energy use by railways is very small compared to the total transport sector.

Table 3-23. Summary of key source analysis, NFR1A3c according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A3c</td>
<td>Liquid Fuels – NOₓ</td>
</tr>
</tbody>
</table>

3.2.14.2 METHODOLOGICAL ISSUES

Both Tier 1 and Tier 2 methods are used to estimate emissions from diesel. The Swedish Transport Administration (STA) estimates the emissions of SO₂, NOₓ, NMVOC and CO from railways, which are based on the consumption of diesel by railways and default as well as country specific emission factors which are described below.

The estimated diesel consumption is based on fees paid by the rail operators and is considered to be of very high quality. Emissions of SO₂ are based on country-specific thermal values and the actual sulphur content for diesel fuel.

The threshold limits for CO and NOₓ is used as emission factors for all emissions from engines that comply with the EU emission standards Stage IIIA and Stage IIIB. For engines introduced before the implementation of EU emissions standards, the emission factors from EMEP/EEA guidebook 2013 are used to estimate emissions of CO and NOₓ.

The conversion of g/kWh to g/litre is based on the fuel consumption factors in Table 3-5 in the EMEP/EEA Guidebook 2013 and a diesel density of 816 g / litre. The same density is used for all years.

The emissions of NMVOC and particles are estimated with emission factors from EMEP/EEA guidebook 2013.

3.2.14.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The emissions for NFR1A3c are 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.

Uncertainties for activity data and emissions reported for working machinery in NFR1A4b ii can be seen in Annex 1 to the IIR

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79 As of 2017, the Swedish Energy Agency took over the responsibility for data relating to energy consumption by railways; previously published by Traffic Analysis (TRAFA). SWEDISH OFFICIAL STATISTICS STATISTICAL NOTIFICATIONS EN0118 SM 1701.

80 http://www.dieselnet.com/standards/eu/nonroad.php#rail
3.2.14.4  SOURCE-SPECIFIC QA/QC AND VERIFICATION
All Tier 1 general inventory level QC procedures and all QC procedures applicable to this sector are used. The activity data has been subject to QA/QC procedures.

3.2.14.5  SOURCE-SPECIFIC RECALCULATIONS
The consumption of diesel was adjusted for 2012-2015 in submission 2018, due to updated information from the rail operators. It resulted in an increased consumption of diesel for 2012-2013 and a decreased consumption in 2014-2015. The same pattern applies to the emissions.

3.2.14.6  SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.15  Navigation, NFR1A3d ii
3.2.15.1  SOURCE CATEGORY DESCRIPTION
Domestic navigation is defined as emissions from diesel oil, domestic heating oil and residual fuel oil purchased and used in Sweden by large commercial ships and leisure boats. The energy consumption from all fuels by national navigation has decreased by 57 % since 1990. It’s foremost the consumption of residual fuel oils that have decreased 1990-2016, while the consumption of diesel had increased slightly. This is the result of the EU Sulphur Directive which entered into force the 1’st of January 2015, which requires a drastic cut in the sulphur emissions from all vessels operating in the Baltic sea (as well as in the North Sea and in the English channel).

Table 3-24. Summary of key source analysis, NFR1A3d ii, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A3d ii</td>
<td>Gas/diesel oil – CO, NMVOC, PM$_{1,5}$, NO$<em>x$, PM$</em>{10}$</td>
<td>Gas/diesel oil – SO$_x$, CO, NO$_x$, TSP</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Residual oil – Ni</td>
<td>Residual oil – Ni, SO$<em>x$, As, PM$</em>{1,5}$, NO$<em>x$, PM$</em>{10}$, Cr, TSP</td>
<td></td>
</tr>
</tbody>
</table>

3.2.15.2  METHODOLOGICAL ISSUES
This source category covers domestic navigation and leisure boats. Domestic navigation is defined as emissions from diesel oil, domestic heating oil and residual fuel oil purchased and used in Sweden. 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 3-25.

Table 3-2. Reporting of emissions from navigation, according to the Good Practice Guidance.

<table>
<thead>
<tr>
<th>Fuel bought in</th>
<th>Traffic between Swedish harbours</th>
<th>Traffic between Swedish and international harbours</th>
<th>Traffic between two international harbours</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sweden</td>
<td>Domestic, 1A3d ii</td>
<td>International bunkers, 1A3d i (i)</td>
<td>International bunkers, 1A3d i (i)</td>
</tr>
</tbody>
</table>
Emissions from national navigation are estimated using Tier 1. The activity data for domestic navigation is based on the amount of fuel purchased and consumed in Sweden by national navigation according to the monthly survey on supply and delivery of petroleum products from Statistics Sweden. The Swedish Maritime Administration (SMA) provides the emissions factors for NOX, CO, NMVOC and SO2 for 2005-2015 and the Swedish Transport Agency (STA) as from 2016 in accordance with the Swedish climate legislation. The emission factors for metals and PAH:s were updated in submission 2018 by the Swedish Environmental Research Institute (IVL). Emissions by national navigation in 1990-2004 are calculated using emission factors from a SMED study conducted in 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.

The gasoline and diesel consumption by leisure boats in Sweden 1990-2004 is based on a survey regarding leisure boat from 2004 and a study carried out by SMED in 2005. The gasoline consumption by leisure boats is estimated to 32,500 m³/year and the diesel consumption to 12,000 m³/year for the years 1990-2004 as a result of these studies.

The gasoline and diesel consumption by leisure boats in 2005-2013 is based on a leisure boat survey from 2010 and an analysis of the result in the survey carried out by SMED in 2014. The gasoline consumption is estimated to 37,768 m³/year and the diesel consumption to 18,172 m³/year for 2010 and onwards as a result of these studies. The consumption of both gasoline and diesel has been estimated by interpolation for the years 2005-2009.

Emissions of SO₂ from leisure boats are based on the fuel consumption and the same thermal values and emission factors as for civil road traffic regarding both transportation modes and sea transport.
gasoline and diesel. The emission of NO\textsubscript{X}, NMVOC and CO are based on the estimated fuel consumption and emission factors provided and updated by the Swedish Environmental Research Institute (IVL) in submissions 2018\textsuperscript{88}. The emission factors were in previous submissions taken from the EMEP/EEA guidebook 2009. The emissions of NO\textsubscript{X}, NMVOC and CO from leisure boats also depend on the ratio between 2-stroke and 4-stroke engines. The estimated ratio is based on a study by Statistics Sweden 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 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 the same as for 2004.

Emissions of particles from leisure boats have been estimated with the assumption that leisure boats generate the same amount of emissions per energy unit as for gasoline-run off road vehicles and other machinery for households\textsuperscript{89}.

\subsection*{3.2.15.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY}

The distribution of marine distillate fuels and residual fuel oils to domestic and international navigation (bunkers) entail uncertainties. The respondents of the survey on supply and delivery of petroleum products\textsuperscript{90} provide the current distribution, but they are suspected to lack full information on the end-use of all the fuels provided. The distribution between domestic and international use has varied considerably for some years and was therefore reviewed in a SMED study in 2011\textsuperscript{91}. The study showed that the respondents of the survey are capable to correctly separate these fuel amounts, and in line with IPCC Guidelines, since VAT is applied on national fuel consumption but not on international bunkers.

It should be noted that the consumption of fuel by national navigation is relatively small compared to the consumption by international navigation (bunkers). A minor shift in the fuel consumption between the two is clearly noticeable in the national consumption but not the other way around.

The Swedish Energy Agency (STEM) has for the last 2.5 years worked to improve the survey “Monthly fuel, gas and inventory statistics”\textsuperscript{92}, which is the source for activity data (fuel consumption) for both domestic and international navigation in the Swedish inventory. The survey collects fuel data from oil companies and other fuel providers who have stocks of petroleum products and coal as well as from

\textsuperscript{88} Fridell, E., Mawdsley, I., Wisell T. 2017.
\textsuperscript{89} Kindbom and Persson. 1999
\textsuperscript{90} Monthly fuel, gas and inventory statistics.
\textsuperscript{91} Eklund et al. 2011. Emissions from navigation and fishing including international bunkers
companies with a large consumption of oil in the manufacturing industries and energy industries. STEM implemented the new version of the survey in January 2018. The results from the new survey will be applied to the Swedish inventory in submission 2020.

Uncertainties for activity data and emissions reported for domestic navigation in NFR1A3d can be seen in Annex 1 to the IIR.

3.2.15.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
All Tier 1 general inventory level QC procedures and all QC procedures applicable to this sector are used. The activity data has been subject to QA/QC procedures.

In 2011 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 calculations 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 IDs 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 NFR1A4c) and military ships (1A5b). Further studies should also include fishing and military ships to get the whole picture.

3.2.15.5 SOURCE-SPECIFIC RECALCULATIONS
The Swedish Environmental Research Institute (IVL) updated several emission factors for both domestic and international navigation in submission 2018\(^\text{93}\) as well as for all fuel types. There was a greater adjustment of the emission factors for residual fuel oil, compared to the other fuels, and as a result the emissions. The following emission factors were updated:

\(^{93}\) Fridell, E., Mawdsley, I., Wisell T. 2017.
• NOx, NMVOC & CO for leisure boats
• TSP, PM10, PM2.5, BC, NH3, As, Cd, Cr, Cu, Hg, Ni, Pb, Se, Zn, benso(a)pyren, benso(b)fluoranten, benso(k)fluoranten, indeno(1,2,3-cd)pyren, PAH 4 total, Dioxin/Furan, PCB tot and HCB for ships.

On a total level, the updated emission factors caused increased emissions of PAH’s for the whole time-period. The emissions of most metals also increased on a total level for 1990-2005 and decreased for 2006-2016 comparing submission 2018 and 2017. The emissions of NH3 increased for the whole time-period, except for 1999-2000, while the emission of particles increased for 1990-2009 and decreased for 2010-2016 on a total level, due to the updated emission factors.

3.2.15.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.16 Other transportation, NFR1A3e ii
3.2.16.1 SOURCE CATEGORY DESCRIPTION
Emissions reported in NFR1A3e ii refer to off-road vehicles and other machinery including ground activities in airports and harbours, and off-road activities not otherwise reported under 1A2g, 1A4a, 1A4b or 1A4c. A list of NFR codes in which emissions from working machinery are reported is found in Table 3-26.

The consumption of diesel (including low blended biofuel) has had a decreasing trend in 2005-2014, but for the last two years the consumption has increased. The emissions of CO, NOx, particles and NMVOC keep however decreasing.
Table 3-26. Distribution of emissions from off-road vehicles and other machinery

<table>
<thead>
<tr>
<th>Category</th>
<th>NFR</th>
<th>Definition IPCC Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>1A2g vii</td>
<td>Mobile machineries in industry that run on petroleum fuels, as for example tractors, dumpers, cranes, excavators, generators, wheel loaders, sorting works, pump units etc.</td>
</tr>
<tr>
<td>Other</td>
<td>1A3e ii</td>
<td>Combustion emissions from all remaining transport activities including ground activities in airports and harbours, and off-road activities not otherwise reported under 1A4c ii or 1A2g vii.</td>
</tr>
<tr>
<td>Commercial/Institutional</td>
<td>1A4a ii</td>
<td>Garden machinery, e.g. lawn mowers and clearing saws, not used by private users. Also tractors not used in industry ore forestry or agriculture.</td>
</tr>
<tr>
<td>Residential</td>
<td>1A4b ii</td>
<td>All emissions from mobile fuel combustion in households, as for example tractors, lawn mowers, snow mobiles, forklifts, trimmers, chainsaws and forklifts.</td>
</tr>
<tr>
<td>Agriculture, Forestry</td>
<td>1A4c ii</td>
<td>Emissions from mobile fuel combustion in agriculture and forestry, as for example loader-excavator, tractor, harvester, clearing saw etc. High-way agricultural transportation is excluded.</td>
</tr>
</tbody>
</table>

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 key source analysis, NFR1A3e ii, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Diesel oil - NOₓ</td>
<td>Diesel oil - NOₓ, PM₁₀, TSP, PM₂.₅</td>
</tr>
</tbody>
</table>

3.2.16.2 METHODOLOGICAL ISSUES
A national model is used to estimate emissions from all working machinery used in Sweden, 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.

3.2.16.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
3.2.16.3.1 Mobile combustion
No recalculations have been performed for emissions in the mobile sector for the eighties, and thus there are inconsistencies in time series between the eighties and later years. Time series from mobile combustion in NFR1A4b ii have been reviewed for later years and are considered to be consistent.

Uncertainties for activity data and emissions reported for working machinery in NFR1A4b ii can be seen in Annex 1 to the IIR.

3.2.16.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
The model was implemented the first time in submission 2009. During 2010 the model underwent a second verification. Activity data and emissions factors were reviewed in 2012 and 2013. Time series are checked for consistency and recalculations are verified every year.
3.2.16.5 SOURCE-SPECIFIC RECALCULATIONS

The model for working machinery has been updated with sales data for off-road vehicles for 2005-2016 in submission 2018, which was provided by the Swedish trade association for suppliers of mobile machines. The allocation key in the model was also updated, as new information regarding the sector distribution was received from both the trade association and the vehicle register. The age of some of the oldest tractors in the model was adjusted as well, as they in previous submissions mistakenly were grouped together in the oldest age category\textsuperscript{94}.

The updated model has resulted in a decreased consumption of diesel (including low blended biofuel) for practically the entire time series, especially after 2005, as well as decreased emissions of NOx, CO, NH\textsubscript{3} and particles comparing submission 2018 to 2017. The emissions of NMVOC increased however for 1990-2006 and then decreased for the rest of the time series.

3.2.16.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

3.2.17 Commercial/institutional, NFR1A4a

3.2.17.1 SOURCE CATEGORY DESCRIPTION

This category includes stationary combustion for heating of premises used for commercial and institutional activities as well as emissions from working machinery used in these activities.

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 35000 TJ, around year 2000 it had decreased to about 20000 TJ, and in 2015 it was around 7000 TJ. Liquid fuels account for most of the decrease. The share of liquid fuels in 1990 was about 95\% and the corresponding share in 2016 was 32\%.

For premises, the energy consumption with district heating has increased with 14.7 TWh in 2002 to 16.8 TWh in 2016. The energy consumption with oil has decreased with from 3.3 TWh in 2002 to 0.4 TWh in 2016. For one- and two-dwellings, the energy consumption with district heating has increased with from 23.3 TWh in 2002 to 24 TWh in 2016. The energy consumption with oil has decreased with from 2.5 TWh in 2002 to 0.2 TWh in 2016. For one- and two-dwellings, the energy consumption with district heating has increased with 3 TWh in 2002 to 5.5 TWh in 2016. The energy consumption with oil has decreased from 9 TWh in 2002 to 0.4 TWh in 2016\textsuperscript{95}.


\textsuperscript{95}Swedish Energy Agency 2017.
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.

The mobile emission source included in this sector is combustion by off-road vehicles and other machinery (working machinery), for example gardening machines for professional use and tractors that are not used in industry, farming or forestry. The emissions of air pollution from this sector are reported in 1A4aii. The fuel consumption by mobile combustion has increased by 42% in 2016 compared to 1990, but while the consumption of diesel (including low blended biofuel) has shown an increasing trend, the consumption of gasoline (including low blended biofuel) has stagnated or even shown a slightly decreasing trend as from 2011. The emissions of CO, NOx, NH3 and NMVOC have decreased in the last years while the emissions of particles show a decreasing trend.

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 key source analysis, NFR1A4a, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A4a</td>
<td>Biomass – PAH 1-4, CO, Cd, PM2.5, PM10, TSP, Zn</td>
<td>Biomass – PAH 1-4, CO, Cd, PM2.5, PM10, TSP, Zn</td>
</tr>
<tr>
<td></td>
<td>Diesel oil – NOx, PM2.5</td>
<td>Diesel oil – NOx</td>
</tr>
<tr>
<td></td>
<td>Gasoline - CO, NMVOC, NOx</td>
<td>Gasoline - CO</td>
</tr>
<tr>
<td></td>
<td>Liquid fuels -</td>
<td>Liquid fuels - Ni, SO2, NOx, Se, PM2.5, TSP, PM10</td>
</tr>
</tbody>
</table>

3.2.17.2 METHODOLOGICAL ISSUES

3.2.17.2.1 Stationary combustion

For stationary combustion within NFR1A4a, all activity data and emission factors are on national level by fuel type and estimated emissions are therefore considered to correspond to Tier 2. 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 in Annex 2. Activity data for the latest emission year is preliminary as the annual energy balances are not published at the time when the emission calculations have to be finalized.

3.2.17.2.2 Mobile combustion

Emissions from mobile combustion in NFR1A4a refer mainly to gardening machines for professional use and tractors that are not used in industry, farming or forestry. A national model is used to estimate emissions from all working machinery used in Sweden and it is considered to correspond to Tier 3 for all emissions, except for SO2 which is estimated according to Tier 2. The model is further explained in Annex 2.
Emissions from working machinery are also reported in NFR1A2g vii, 1A3e ii, 1A4b ii and 1A4c ii. See Table 3-29.

**Table 3-29. Distribution of emissions from off-road vehicles and other machinery**

<table>
<thead>
<tr>
<th>Category</th>
<th>NFR</th>
<th>Definition IPCC Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>1A2g vii</td>
<td>Mobile machineries in industry that run on petroleum fuels, as for example tractors, dumpers,</td>
</tr>
<tr>
<td></td>
<td></td>
<td>cranes, excavators, generators, wheel loaders, sorting works, pump units etc.</td>
</tr>
<tr>
<td>Other</td>
<td>1A3e ii</td>
<td>Combustion emissions from all remaining transport activities including ground activities</td>
</tr>
<tr>
<td></td>
<td></td>
<td>in airports and harbours, and off-road activities not otherwise reported under 1A4c ii or</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1A2g vii.</td>
</tr>
<tr>
<td>Commercial/</td>
<td>1A4a ii</td>
<td>Garden machinery, e.g. lawn mowers and clearing saws, not used by private users, Also</td>
</tr>
<tr>
<td>Institutional</td>
<td></td>
<td>tractors not used in industry ore forestry or agriculture.</td>
</tr>
<tr>
<td>Residential</td>
<td>1A4b ii</td>
<td>All emissions from mobile fuel combustion in households, as for example tractors, lawn</td>
</tr>
<tr>
<td></td>
<td></td>
<td>movers, snow mobiles, forklifts, trimmers, chainsaws and forklifts</td>
</tr>
<tr>
<td>Agriculture,</td>
<td>1A4c ii</td>
<td>Emissions from mobile fuel combustion in agriculture and forestry, as for example loader-</td>
</tr>
<tr>
<td>Forestry</td>
<td></td>
<td>excavator, tractor, harvester, clearing saw etc. High-way agricultural transportation is</td>
</tr>
<tr>
<td></td>
<td></td>
<td>excluded.</td>
</tr>
</tbody>
</table>

3.2.17.2.3 **Verification of Estimation models and allocation methods for fuel in the other sectors**

In submission 2005 and earlier, there were large uncertainties in estimation models and allocation methods for fuel in the Other sectors and NFR1A2f, construction. In 2005, a study was performed by SMED, aiming at identifying and analysing the methods and models applied for each sub-sector and determine whether they were in line with the IPCC guideline recommendations. 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 recalculation of activity data and emissions in NFR1A4a and 1A4b. Due to these recalculations there are obvious inconsistencies between the national energy balances and the national emission inventory data.

Furthermore, all fuels proved to be correctly allocated on stationary and mobile combustion. In the Swedish air emission inventory, this means that all diesel oil and gasoline reported under Other sectors in the energy balances are used by mobile combustion, while all the other fuels are related to stationary combustion.

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3.2.17.2.4 Activity data for stationary combustion in other sectors

Activity data for the latest emission year is preliminary as the annual energy balances are not published at the time when the emission calculations have to be finalized.

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 study carried out in 2013 had shown that the fuel consumption estimate used in the national energy balance and the emission inventory is more complete than the data reported to Eurostat.

Every year, there are revisions in the annual energy balances for years t-2 and t-3, that is, data published in 2010 contain revisions in fuel consumption in 2007 and 2008. These sometimes large revisions in the annual energy balances lead to large revisions of GHG inventory data as well as for air pollutants. In submission 2016, activity data and hence also emissions have been revised for 2012 and 2013.

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 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 uncertainty is high in this sector and the time series 1990-2013 shows that inter-annual variations in total fuel consumption can be high.

3.2.17.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

3.2.17.3.1 Stationary combustion

The large activity data uncertainty in the stationary combustion is due to the use of data from the annual energy balances. Uncertainties for activity data and emission factors are in generally set by fuel type.

Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As, Cd, Cu, HCB, PAH, PCB
- All fuels for diox
- Liquid fuels for Cu, Hg, Cr
- Solid fuels for HCB, PCB

See Annex 1 for more details regarding uncertainties for activity data and emissions.
3.2.17.3.2 **Mobile combustion**

No recalculations have been performed for emissions in the mobile sector for the eighties, and thus there are inconsistencies in time series between the eighties and later years. Time series from mobile combustion in NFR1A4b ii have been re-viewed for later years and are considered to be consistent.

Uncertainties for activity data and emissions reported for working machinery in NFR1A4b ii can be seen in Annex 1 to the IIR.

3.2.17.4 **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. In 2005, a study was performed by SMED, aiming at identifying and analysing the methods and models applied for each sub-sector and determine whether they were in line with the IPCC guideline recommendations[98]. 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 NFR1A4a and 1A4b[99]. 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.

3.2.17.5 **SOURCE-SPECIFIC RECALCULATIONS**

In submission 2018 activity data for use of oils in forestry sector was revised for years 2005 to 2015. The revision resulted in very large relative changes within the sector.

In addition activity data for all categories in this sector has been revised between for year 2015 due to revision of the annual energy balances.

The model for working machinery has been updated with sales data for off-road vehicles for 2005-2016 in submission 2018, which was provided by the Swedish trade association for suppliers of mobile machines. The allocation key in the model

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was also updated, as new information regarding the sector distribution was received from both the trade association and the vehicle register. The age of some of the oldest tractors in the model was adjusted as well, as they in previous submissions mistakenly were grouped together in the oldest age category.  

The updated model resulted in a decreased consumption of diesel (including low blended biofuels) for most years 1991-2009, but an increased consumption for 2010-2016. The consumption of gasoline (including low blended biofuels) only increased slightly as from 2006. Most pollutants from working machinery used in this sector increased as from 2005/2006 and onwards in submissions 2018 compared to submission 2017, except for particles.

3.2.17.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.  

3.2.18 Residential, NFR1A4b

3.2.18.1 SOURCE CATEGORY DESCRIPTION

In this category both stationary and mobile combustion occur. Stationary combustion of fuels within residential decreased by 62% between 1990 and 2016, mainly due to a continuous increase in district heating use. In recent years, the use of heat pumps has also increased significantly. 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. Emissions of CO₂, CH₄ and N₂O from the use of charcoal are included in this source category.

Mobile combustion in NFR1A4bii refer to gardening machines used in households e.g. lawn mowers, hedged cutters, clearing saws and more. Also snow mobiles and four wheelers not used for professional purposes are allocated to NFR1A4bii. The emissions of CO, NOx, NMVOC, NH₃ and particles mainly arise from combustion of gasoline, but regarding NOx this applies after 2009 and for NH₃ after 1995. In 2016, the consumption of gasoline was 89% of the total fuel consumption.

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101 Swedish Energy Agency 2014a

102 Swedish Energy Agency 2014b
3.2.18.2 METHODOLOGICAL ISSUES

3.2.18.2.1 Stationary combustion

Emissions from dwellings within stationary combustion is reported in NFR1A4b i. For stationary combustion, all activity data is on national level by fuel type and estimated emissions are therefore considered to correspond to Tier 1.

The main data source is the annual energy balances. One- and two-dwellings statistics, Holiday cottages statistics and Multi-dwellings statistics are used as complementary data sources 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 well as the use of preliminary data for stationary combustion in the Other sectors as discussed in section 3.2.17 also applies to NFR1A4b.

Emissions arising from the use of charcoal are estimated using national statistics and default 2006 IPCC guidelines EFs.

3.2.18.2.2 Mobile combustion

Emissions from mobile machinery used in households are included in NFR1A4bii. Machines included here are mainly several types of gardening machines e.g. lawn movers, hedge cutters and chain saws. Also emissions from snow mobiles used in households are reported here. Emissions from gardening machines for professional use are reported in 1A4a ii. Also emissions from tractors that are not used in forestry, agriculture or the industry are reported here as are emissions from generator sets and mobile freezers and chillers. A national model is used to estimate most emissions from all working machinery used in Sweden, considered to correspond to Tier 3 for all emissions, except SO₂ which is estimated according to Tier 2. The model is further explained in Annex 2.

Emissions from working machinery are also reported in NFR1A2g vii, 1A3e ii, 1A4a ii and 1A4c ii. See Table 3-31.
Table 3-31. Distribution of emissions from off-road vehicles and other machinery

<table>
<thead>
<tr>
<th>Category</th>
<th>NFR</th>
<th>Definition IPCC Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>1A2g vii</td>
<td>Mobile machineries in industry that run on petroleum fuels, as for example tractors, dumpers, cranes, excavators, generators, wheel loaders, sorting works, pump units etc.</td>
</tr>
<tr>
<td>Other</td>
<td>1A3e ii</td>
<td>Combustion emissions from all remaining transport activities including ground activities in airports and harbours, and off-road activities not otherwise reported under 1A4c ii or 1A2g vii.</td>
</tr>
<tr>
<td>Commercial/Institutional</td>
<td>1A4a ii</td>
<td>Garden machinery, e.g. lawn mowers and clearing saws, not used by private users, Also tractors not used in industry ore forestry or agriculture.</td>
</tr>
<tr>
<td>Residential</td>
<td>1A4b ii</td>
<td>All emissions from mobile fuel combustion in households, as for example tractors, lawn movers, snow mobiles, forklifts, trimmers, chainsaws and forklifts</td>
</tr>
<tr>
<td>Agriculture, Forestry</td>
<td>1A4c ii</td>
<td>Emissions from mobile fuel combustion in agriculture and forestry, as for example loader-excavator, tractor, harvester, clearing saw etc. High-way agricultural transportation is excluded.</td>
</tr>
</tbody>
</table>

3.2.18.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

3.2.18.3.1 Stationary combustion

The activity data and emission factor uncertainties for stationary combustion are 20 % and 1 % respectively. The large activity data uncertainty is due to the use of input data from the annual energy balances.

The time series for NFR1A4b is considered to be consistent as there haven’t been any major changes in methodology or input data to the energy balances that affect this category.

Uncertainties for activity data and emission factors are in generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As, Cd, Cr, Cu, HCB, PAH, PCB, Zn
- Liquid fuels for Cd, Cr, Ni, PAH, Pb, Se, SO2 and Zn
- Other fuels for PM10, PM2.5

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.18.3.2 Mobile combustion

No recalculation has been performed for emissions in the mobile sector for the eighties, and thus there are inconsistencies in time series between the eighties and later years. Time series from mobile combustion in NFR1A4b ii have been reviewed for later years and are considered to be consistent.

Uncertainties for activity data and emissions reported for working machinery in NFR1A4b ii can be seen in Annex 1 to the IIR.
3.2.18.4 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.

3.2.18.5 SOURCE-SPECIFIC RECALCULATIONS
Following revisions of the energy balances, the activity data for stationary combustion within 1A4b was revised for all fuels for the year 2014.

The model for working machinery has been updated with sales data for off-road vehicles for 2005-2016 in submission 2018, which was provided by the Swedish trade association for suppliers of mobile machines. The allocation key in the model was also updated, as new information regarding the sector distribution was received from both the trade association and the vehicle register. The age of some of the oldest tractors in the model was adjusted as well, as they in previous submissions mistakenly were grouped together in the oldest age category\(^{103}\).

The updated model has resulted in an increased consumption of diesel (including low blended biofuels) for 1991-2009 as well as increased emissions of NOx, CO, NMVOC, NH3 and particles in submission 2018 compared to submission 2017. The consumption of gasoline (including low blended biofuels) has barely changed, except for a slight increase of the consumption in the end of the times series.

3.2.18.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.19 Agriculture/Forestry/Fisheries, NFR1A4c
3.2.19.1 SOURCE CATEGORY DESCRIPTION
This category includes emissions from stationary combustion for heating purposes and mobile combustion in working machinery within agriculture and forestry, and fishing vessels. The structure of the agricultural sector in Sweden is described in chapter 6. Changes in use of liquid and gaseous fuels in agriculture, fishing and forestry have been small since 1990. Due to availability of better data for the period 2003 and later years, there is a shift in the time series for biomass.

The consumption of both gasoline and diesel (including low blended biofuels) by working machinery used in agriculture and forestry has an increasing trend, but the consumption of diesel is dominant and correspond to 91% of the total fuel consumption in 2016.

The emissions of NOx from diesel (including biofuel) are on a much higher level than the same emissions from gasoline and have a strong decreasing trend. They have dropped by 70% since 1990. The same applies for emissions of particles. The greater part of the emissions of CO and NMVOC arise from combustion of gasoline and despite a decreasing trend for these emissions from diesel, they have increased by 100% respectively 16% on a total level since 1990.

Solid fuels consumption within this sector has decreased substantially since 1990.

### Table 3-32. Summary of key source analysis, NFR1A4c i, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A4c i</td>
<td>Biomass – PAH 1-4, CO, Cd, DIOX, PM2.5, PM10, TSP, NMVOC, Zn, Se, Pb, NOX, SO2</td>
<td>Biomass – PAH 1-4, CO, Cd, DIOX, PM2.5, Cr, PM10, TSP, NMVOC, Zn, Cu, Se, Hg, NOX, SO2, Ni</td>
</tr>
<tr>
<td></td>
<td>Liquid – Ni, SOX, NOX, PM2.5, PM10, TSP, Cu, CO, NMVOC, Cd</td>
<td>Liquid – Ni, SOX, PM2.5, PM10, TSP, Cu, CO, NMVOC, NOX</td>
</tr>
<tr>
<td></td>
<td>Solid –</td>
<td>Solid – SOX, Se, TSP</td>
</tr>
</tbody>
</table>

### 3.2.19.2 METHODOLOGICAL ISSUES

In this sector both stationary and mobile combustion occur.

#### 3.2.19.2.1 Stationary combustion

For stationary combustion, all activity data is on national level by fuel type and estimated emissions are therefore considered to correspond to Tier2. Activity data is based on models and results from a survey from 1985 repeated in 2007 (see Other statistics from Statistics Sweden in Annex 2).

Estimation models and allocation methods for fuel in the Other sectors, as well as the use of preliminary data for stationary combustion in the Other sectors as discussed in section 3.2.17 Commercial/institutional, NFR1A4a also applies to NFR1A4c. As a consequence, emissions from stationary combustion 2002-2006 were revised in submission 2009. Emissions for the most recent year will be revised in next submission when annual statistics are available. Note that as a consequence of this revision, emissions from biomass are inconsistent with a sharp increase to a higher level in 2003. There is no information available to improve data 2002 and earlier years. Emissions 1990 are considered to be of sufficient quality as the 1985 survey then was only five years old.

#### 3.2.19.2.2 Mobile combustion

Mobile combustion in 1A4c refers to working machinery used in agriculture (e.g. tractors and combine harvesters), forestry (e.g. forwarders and harvesters) and fisheries. A national model is used to estimate emissions from all land based working machinery used in Sweden, considered to correspond to Tier 3 for all emissions, except SO2 which is estimated according to Tier 2. The model is further explained in Annex 2.
Emissions from Fisheries, NFR1A4c, were first reported in submission 2006. The estimated fuel consumption is based on a survey on energy consumption within the fishing industry by Statistics Sweden\textsuperscript{104} together with data on the Swedish fishing fleet’s total installed effect in kW from the Swedish 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 fuel consumption by fisheries has decreased by 30% since 1990 and by 4% since last year.

The emissions factors used to estimate emissions from Fisheries are based on a SMED study from 2005\textsuperscript{105}, producing emission factors for $\text{SO}_2$, $\text{NO}_x$ and NMVOC, for 1990-2004. As from 2005, the estimates are based on the same consumption estimate and emission factors as for 2004. However, from 2007 and onwards the emission factors for $\text{SO}_2$ 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 $\text{NO}_x$ reduction) is assumed to be negligible.

Emissions from working machinery are also reported in NFR1A2g vii, 1A3e ii, 1A4a ii and 1A4b ii. See Table 3-33.

### Table 3-33. Distribution of emissions from off-road vehicles and other machinery

<table>
<thead>
<tr>
<th>Category</th>
<th>NFR</th>
<th>Definition IPCC Guidelines</th>
</tr>
</thead>
<tbody>
<tr>
<td>Industry</td>
<td>1A2g vii</td>
<td>Mobile machineries in industry that run on petroleum fuels, as for example tractors, dumpers, cranes, excavators, generators, wheel loaders, sorting works, pump units etc.</td>
</tr>
<tr>
<td>Other</td>
<td>1A3e ii</td>
<td>Combustion emissions from all remaining transport activities including ground activities in airports and harbours, and off-road activities not otherwise reported under 1A4c ii or 1A2g vii.</td>
</tr>
<tr>
<td>Commercial/Institutional</td>
<td>1A4a ii</td>
<td>Garden machinery, e.g. lawn mowers and clearing saws, not used by private users, Also tractors not used in industry ore forestry or agriculture.</td>
</tr>
<tr>
<td>Residential</td>
<td>1A4b ii</td>
<td>All emissions from mobile fuel combustion in households, as for example tractors, lawn movers, snow mobiles, forklifts, trimmers, chainsaws and forklifts</td>
</tr>
<tr>
<td>Agriculture, Forestry</td>
<td>1A4c ii</td>
<td>Emissions from mobile fuel combustion in agriculture and forestry, as for example loader-excavator, tractor, harvester, clearing saw etc. High-way agricultural transportation is excluded.</td>
</tr>
</tbody>
</table>

\textsuperscript{104} Statistics Sweden, 2006  ENFT0601.  
\textsuperscript{105} Cooper et al., 2005a.
3.2.19.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

3.2.19.3.1 Stationary combustion

The sharp increase in use of biomass in stationary combustion 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. Solid fuels have not been used in this sector since 2000.

Uncertainties for activity data and emission factors are in generally set by fuel type. Activity data uncertainty is relatively low for most fuel types. Emission factor uncertainty is for some fuels very high. The fuel groups and EF that have highest uncertainty are:

- Biomass for As, Cd, Cu, diox, HCB, PAH, PCB
- Liquid fuels for Cd, Cr, Cu, Ni, PAH, Pb, Se and Zn
- Solid fuels for As, diox, HCB, PAH, SO2, Zn

See Annex 1 for more details regarding uncertainties for activity data and emissions.

3.2.19.3.2 Mobile combustion

No recalculation have been performed for emissions in the mobile sector for the eighties, and thus there are inconsistencies in time series between the eighties and later years. The time series from mobile combustion in NFR1A4c ii have been reviewed for later years and are considered to be consistent.

Uncertainties for activity data and emissions reported for working machinery in NFR1A4c ii can be seen in Annex 1 to the IIR.

3.2.19.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Following revisions of the energy balances, the activity data for stationary combustion within 1A4c was revised for all fuels 2014. In addition revision of the energy consumption of biomass and natural gas within the agriculture and forestry sector was made for the years 2005-2013.

3.2.19.5 SOURCE-SPECIFIC RECALCULATIONS

The model for working machinery has been updated with sales data for off-road vehicles for 2005-2016 in submission 2018, which was provided by the Swedish trade association for suppliers of mobile machines. The allocation key in the model was also updated, as new information regarding the sector distribution was received from both the trade association and the vehicle register. The age of some of
the oldest tractors in the model was adjusted as well, as they in previous submissions mistakenly were grouped together in the oldest age category\textsuperscript{106}.

The updated model has resulted in a decreased consumption of diesel (including low blended biofuels) by working machineries used in forestry for the entire time series, while the opposite is true for agriculture. The emissions of NOx still have a decreasing trend in submission 2018, just as in submission 2017. But there is a shift regarding the emissions of NOx from diesel, between agriculture and forestry. The emissions of NOx increase from agriculture and decrease from forestry and the same applies for emissions of particles.

The consumption of gasoline (including low blended biofuels) is relatively small in both sectors and has only increased marginally for the last years when comparing submission 2018 and 2017. As most emissions of CO stem from combustion of gasoline, they have not changed significantly due to the updated model, but shows a more distinct upward trend for the last years in submission 2018.

3.2.19.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

3.2.20 Other stationary combustion, NFR1A5a
No emissions are reported in this category.

3.2.21 Military transport, NFR1A5b
3.2.21.1 SOURCE CATEGORY DESCRIPTION
NFR1A5b includes emissions from military transports. The fuel consumption and emissions have decreased over the years due to decreased activities.

\begin{table}
\centering
\begin{tabular}{|c|c|}
\hline
NFR & Key Source Assessment 2016 \\
Level & Trend \\
\hline
1A5b & Liquid Fuels - NOx & Liquid Fuels - NOx \\
\hline
\end{tabular}
\caption{Summary of key source analysis, NFR1A5b according to approach 1.}
\end{table}

3.2.21.2 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.

Emissions from military aviation are based on an average of LTO and cruise emission factors from civil aviation. 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, NFR1D2.

Emissions from military road transport are included in the emissions estimated by the road emission model HBEFA 3.3. The emissions by each vehicle type in HBEFA are consequently 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. These emissions are allocated to military road transport.

To subtract and separate emissions from military road transport from civil road transport in HBEFA, the equation below is used:

\[ A = B - \sum \frac{(C-D)}{C}E_i \]

\[ A = \text{Military transport emissions} \]
\[ B = \text{Total HBEFA emissions} \]
\[ C = \text{Total fuel consumption National Statistics} \]
\[ D = \text{Military fuel consumption Swedish Armed Forces} \]
\[ E_i = \text{HBEFA emissions per vehicle type} \]

Please note that for 1980-1989, only emissions of particles from jet gasoline in military aviation are reported due to the lack of sufficient information for the other sub-sectors. Also, note that there was only a consumption of FAME for 1999-2002 and a consumption of Ethanol for 2007-2016 (with the exception of 2015).

3.2.21.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for activity data and emissions reported for military transport NFR1A5b can be seen in Annex 1 to the IIR.

3.2.21.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

3.2.21.5 SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculations have been made.

3.2.21.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.
3.2.22 Memo Items International bunkers, 1D, NFR1A3ai and 1A3dii

3.2.22.1 SOURCE CATEGORY DESCRIPTION

This sector includes emissions from bunker fuels, e.g. fuel bought in Sweden and used for international aviation and international navigation.

International Aviation is defined as emissions from flights that depart in one country and arrive in a different country. However, “Cruise emissions” from both domestic and international aviation should be reported separately as a memo items in NFR1A3ai and are not included in national totals. This applies according to the Long-Range Transboundary Air Pollution Convention (LRTAP). Emissions from both national and international aviation during the “LTO cycle” belong to the national totals. The emissions of NO\textsubscript{X} have increased since 2009, while the emissions of CO from cruise have had a decreasing trend before 2003 and again an increasing trend 2003 and onwards. The emissions of Pb have decreased noticeably for the whole period, as leaded gasoline has been phased out. The emissions of NMVOC fluctuate, but show a decreasing trend over time since 1990.

International navigation is defined as fuels bought in Sweden, by Swedish or foreign-registered ships, and used for transports to non-Swedish destinations, but excludes consumption by fishing vessels. Emissions from international navigation are not included in the national total, but are instead reported separately as a memo item in NFR1A3dii. The division on international and domestic fuels is based on information from the monthly survey on supply and delivery of petroleum products\textsuperscript{107}. The emissions from international navigation fluctuate over time in line with the fluctuating fuel consumption.

The distribution of marine distillate fuels and residual fuel oils over domestic and international navigation (bunkers) entail uncertainties. The current distribution is provided by the respondents of the survey on supply and delivery of petroleum products, but these are suspected to lack full information on the end-use of all the fuels provided. Hence, the distribution between domestic and international use might vary considerably for some years. The fuel data in the Monthly fuel, gas and inventory statistics was analysed in 2011 and found to be of relatively good quality\textsuperscript{108}. This is a consequence of that VAT is applied on national fuel consumption, but not on international bunkers, which makes all respondents to the survey separate the fuel used for national respectively international navigation with accuracy.

To evaluate Swedish emissions, international bunkers are of course important, especially as international bunkering of fuel is substantially greater than the fuel used for domestic navigation and aviation. Emissions have increased significantly since 1990 due to increased travelling and transportation of goods.


\textsuperscript{108} Eklund et al. 2011. Emissions from navigation and fishing including international bunkers
Table 3-35. Summary of key source analysis, NFR1D according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Level</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1D</td>
<td>International Aviation: Jet kerosene</td>
<td>$NO_x$</td>
<td>$NO_x$</td>
</tr>
</tbody>
</table>

3.2.22.2 METHODOLOGICAL ISSUES

International bunkers from aviation and navigation are defined as fuels bought in Sweden, by Swedish or foreign-registered airplanes or ships, and used for transport to non-Swedish destinations. Emissions from bunker fuels is reported separately as a memo items in NFR1A3ai respectively 1A3di.

The fuel consumption and emissions from both national and international aviation are calculated by the Swedish Defence Research Agency (FOI) by using an estimation model and data provided by Swedish Transport Agency (STAg) regarding:

- Airport of departure and arrival
- Type of aircraft
- Number of flights
- Number of passengers
- International or domestic flight

A database with information regarding 200 different types of aircraft is also used. The emission data regarding different types of aircrafts in the database originates from “ICAO Engine Exhaust Emission Data Bank”. All this data is used to calculate emissions and amounts of burnt fuel for total flight time as well as for aircraft movements below 3000 feet at the airports, the so called LTO cycle. The FOI has in a published report described their method for estimating the emission from aviation\(^\text{109}\).

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 compared to the International Civil Aviation Organization (ICAO) standards that the IPCC guidelines follow\(^\text{110}\). The traffic from Swedish airports consumes as a result less fuel and gives rise to less emission. The estimated fuel consumption and emissions are adjusted to match the statistics on delivered amount of aviation fuels from Statistics Sweden (see Annex 2).


\(^{110}\) Gustafsson, 2005.
The results from the emission calculations are aggregated into four groups; domestic landing and take-off (LTO), domestic cruise, international LTO and international cruise. The aggregation is based on estimated emissions from the LTO cycle & Cruise reported by STAg and the national/international (bunker) fuel consumption from the monthly survey on supply and delivery of petroleum products from Statistics Sweden.

3.2.22.2.1 INTERNATIONAL AVIATION, NFR1A3A I
International Aviation is defined as emissions from flights that depart in one country and arrive in a different country. However, “Cruise emissions” from both domestic and international aviation should be reported separately as a memo items in NFR1A3ai and not included in national totals. This applies according to the Long-Range Transboundary Air Pollution Convention (LRTAP). Emissions from both national and international aviation during the “LTO cycle” belong to the national totals.

3.2.22.2.2 INTERNATIONAL NAVIGATION, NFR1A3D I
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, but excludes consumption by fishing vessels. Emissions from international bunkers navigation are not included in the national total but instead reported separately as a memo item in NFR1A3di. The division on international and domestic fuels is based on information from the monthly survey on supply and delivery of petroleum products.

Emission from multilateral operations are not included in the national total but instead reported separately as a memo item in NFR1A3di. These emissions are derived from fuel purchased in Sweden and used abroad by Swedish forces participating in UN related operations. These emissions account for very small amounts.

3.2.22.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
The distribution of marine distillate fuels and residual fuel oils to domestic respectively international navigation (bunkers) entail uncertainties. The respondents of the survey on supply and delivery of petroleum products provide the current distribution, but the respondents are suspected to lack full information on the end-use of all the fuels provided. Hence, the distribution between domestic and international use might vary considerably for some years. The fuel data in the Monthly fuel, gas and inventory statistics was analysed in 2011\footnote{Eklund et al. 2011. Emissions from navigation and fishing including international bunkers} and found to be of reliable quality, as VAT is applied on national fuel consumption but not on international bunkers, which results in that the respondents to the survey are able to separate national from international fuel consumption correctly and in line with IPCC Guidelines.
However, the Swedish Energy Agency (STEM) has for the last 2.5 years worked to improve the quality of the survey “Monthly fuel, gas and inventory statistics”\(^{112}\), which is the source for activity data (fuel consumption) in the Swedish inventory. The survey collects fuel data from oil companies and other fuel providers who have stocks of petroleum products and coal as well as from companies with a large consumption of oil in the manufacturing industries and energy industries. STEM implemented the new version of the survey in January 2018. The results from the new survey will be applied to the Swedish inventory in submission 2020.

Uncertainties for activity data and emissions reported for domestic navigation in NFR1A3d can be seen in Annex 1 to the IIR.

3.2.22.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

3.2.22.5 SOURCE-SPECIFIC RECALCULATIONS
The Swedish Environmental Research Institute (IVL) updated several emission factors for both domestic and international navigation in submission 2018\(^{113}\) as well as for all fuel types. There was a greater adjustment of the emission factors for residual fuel oil, compared to the other fuels, and as a result the emissions. The following emission factors were updated:

- TSP, PM10, PM2.5, BC, NH3, As, Cd, Cr, Cu, Hg, Ni, Pb, Se, Zn, benso(a)pyren, benso(b)fluoranten, benso(k)fluoranten, indeno(1,2,3-cd)pyren, PAH 4 total, Dioxin/Furan, PCB tot and HCB for ships.

On a total level, the updated emission factors caused increased emissions of PAH’s for the whole time-period. The emissions of most metals also increased on a total level for 1990-2005 and decreased for 2006-2016 comparing submission 2018 and 2017. The emissions of NH\(_3\) increased for the whole time-period, except for 1999-2000, while the emission of particles increased for 1990-2009 and decreased for 2010-2016 on a total level, due to the updated emission factors.

3.2.22.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.


\(^{113}\)Fridell, E., Mawdsley, I., Wisell T. 2017.
3.3 Fugitive emissions from solid fuels and oil and natural gas, NFR1B

3.3.1 Coal mining and handling, NFR1B1a

3.3.1.1 SOURCE CATEGORY DESCRIPTION

There are no coal mines in Sweden and hence no fugitive emissions from coal mines occur (hence reported as NO).

3.3.2 Solid fuel transformation, NFR1B1b

3.3.2.1 SOURCE CATEGORY DESCRIPTION

NFR1B1b includes emissions of SO$_2$, HN$_3$, NMVOC, NO$_x$, Se and PAH from quenching and extinction at coke ovens. Particle emissions, also occurring during coke production, are allocated to NFR1A1c (industrial combustion). The summary of the latest key source assessment is presented in Table 3-36.

Table 3-36. Summary of key source analysis, NFR1B1b, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1B1b</td>
<td>PAH 1-4, Se</td>
<td>PAH 1-4</td>
</tr>
</tbody>
</table>

3.3.2.2 METHODOLOGICAL ISSUES

Information on SO$_2$ emissions are retrieved from the companies’ environmental reports and direct communication. PAH-4 emissions from quenching and extinction at coke ovens have been calculated for the time period 1987-2016. Estimated emissions of PAH-4 from coke production are based on activity data in combination with emission factors for extinction and quenching from US EPA (United States Environmental Protection Agency, EPA-454/R-98-014). The same emission factors have been used for the whole time series. For the time period before 1987 no reliable activity data are available.

NH$_3$ and Se emissions from coke production are reported from submission 2016, NMVOC and NO$_x$ emissions – from submission 2018 and onwards. Emission factors are obtained from EMEP/EEA Guidebook 2013 and applied to activity data. Activity data, produced amount of coke (Mton), has been acquired from official statistics (1987 - 2000) and from the annual environmental reports for the two facilities producing coke (2001 and onwards).

Fugitive emissions of particles from handling of coke have not been included since these emissions are included in the reporting of particle emissions from the industrial facilities that produce coke. These emissions are thus reported “IE”. Separate calculations based on statistics on coke and petroleum coke, using emission factors for handling of coal from CEPMEIP results in a rough estimate of 300 t TSP/year.
Emissions of As, Hg and dioxin occurring during coke production are reported in NFR2C1.

3.3.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for NFR1B1b are displayed in Annex 1. Uncertainties are expert estimates, except for HN$_3$, NMVOC, NO$_x$ and Se, for which the emission factor uncertainties are calculated based on the intervals in the EMEP/EEA Guidebook 2013.

3.3.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
The improvements in methodology and allocation of emissions from the integrated iron and steel industry in submission 2010 were made based on a study carried out in 2008 looking at emissions from several industrial plants, including the two largest iron and steel plants in Sweden, where inventory data from submission 2008 was compared with data from environmental reports.

3.3.2.5 SOURCE-SPECIFIC RECALCULATIONS
Emissions of NMVOC and NO$_x$ from coke quenching have been added in submission 2018, resulting in ~0.008-0.009 kt and ~0.0003-0.0004 kt increase, respectively, for this code. For As, Hg and dioxin notation keys are changed from NE to IE. Emissions occurring from coke production are reported in NFR2C1. For PCB and HCB notation keys are changed from NE to NA. According to both facilities, there are no indications that these emissions occur.

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.

3.3.3 Other, NFR1B1c
3.3.3.1 SOURCE CATEGORY DESCRIPTION
NFR1B1c includes emissions from flaring of coke oven gas at coke ovens handling (NO$_x$, CO, NMVOC, SO$_2$, TSP, PM$_{10}$ and PM$_{2.5}$) and particle emissions from solid fuels handling (TSP, PM$_{10}$ and PM$_{2.5}$). The summary of the latest key source assessment is presented in Table 3-37.

Table 3-37. Summary of key source analysis, NFR1B1c, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1B1c</td>
<td>Solid Fuels – TSP, PM$<em>{10}$, PM$</em>{2.5}$</td>
<td></td>
</tr>
</tbody>
</table>

3.3.3.2 METHODOLOGICAL ISSUES

3.3.3.2.1 Flaring of coke oven gas

Table 1B1 is not really designed to include flaring, but since NFR1B2 only refers to liquid and gaseous fuels, it is not possible to report flaring from coke oven gas, blast furnace gas and steel converter gas in NFR Table 1B2.

The emissions from flaring of coke oven gas (COG – by-product gas at the integrated iron and steel plants) are calculated with Tier 2, i.e. with activity data directly from the plants, in the same way as for emissions from stationary combustion. All emissions, with the exception of SO₂, are calculated with the same emission factors as for stationary combustion because no other information is available for COG flaring in particular (emission numbers in the environmental reports are given for the coke ovens in total, including both flaring and industrial combustion of COG). For SO₂, one facility provides the emissions from COG flaring, whereas for the other total emissions from COG are distributed between NFR codes using the fuel amounts allocated to each code.

Reported activity data is amounts of flared COG (Mton). The amounts vary considerably between years, and during some years (2009, 2015) they were unusually high, resulting in increasing emissions. According to environmental reports\(^\text{115}\), COG is flared when the production is temporarily stopped because of urgent needs of reparation of equipment or other maintenance measures.

3.3.3.2.2 Handling of solid fuels

Particulate emissions from handling of solid fuels have been calculated for all years since 1980. Emission factors used for handling of exported and imported fuels are those suggested in the CEPMEIP-project\(^\text{116}\). The TSP emission factor is 0.15 kg/tonne where PM\(_{10}\) constitutes 40 % and PM\(_{2.5}\) 4 % of the total particulate emissions. The same emission factors have been used for the entire time series.

No production of coal occurs in Sweden but peat production does occur and from submission 2011 particulate emissions from production of milled peat is included in the estimates of particle emissions. Activity data (as m\(^3\) produced peat) is available from official statistics from 1980 and onwards and is divided in peat used for energy purposes and peat used for agricultural purposes. Furthermore there are different methods for peat production. Most particle emissions arise from the production of milled peat.

Production data from official statistics divide peat used for energy purposes in milled peat and other types of peat. However, this split is not used when reporting


\(^{116}\) CEPMEIP, 2001. TNO. 
http://www.mep.tno.nl/wie_we_zijn_eng/organisatie/kenniscentra/centre_expertise_emissions_assessment.html
production data for peat used for agriculture purposes. Milled peat is mostly used for energy purposes, but some may also be used for agricultural purposes, hence the production data for milled peat may be underestimated.

The TSP emission factors used for milled peat production are from Nuutinen et al. (2007)\footnote{Nuutinen, J., Yli-Pirilä, P., Hytönen, K., Kärtevä, J., 2007, Turvetuotannon poly- ja melupäästöt sekä vaikutukset lähiöon ilmanlaatuun, Symo} and the share of PM\(_{10}\) and PM\(_{2.5}\) are from Tissari et al. (2006)\footnote{Tissari, J. M., Yli-Tuomi, T., Raunemaa, T. M., Tiitta, P. T., Nuutinen J. P., Willman, P. K., Lehtinen, K. E. J., Jokiniemi, J. K., 2006, Fine particle emissions from milled peat production, Boreal Environmental research 11:283-293, Helsinki 30 August 2006}. There are different methods that can be used when harvesting milled peat and the size of the particle emissions depends on which method is used. Since no information is available about the share between the different methods in Sweden an average emission factor is used, Table 3-38.

<table>
<thead>
<tr>
<th>Harvesting method</th>
<th>Particle emission factors (g/m(^3)) for peat production</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>TSP</td>
</tr>
<tr>
<td>HAKU method</td>
<td>0.09</td>
</tr>
<tr>
<td>Mechanical collector</td>
<td>0.14</td>
</tr>
<tr>
<td>Pneumatic collector</td>
<td>0.12</td>
</tr>
<tr>
<td><strong>Average</strong></td>
<td><strong>0.12</strong></td>
</tr>
</tbody>
</table>

Due to problems with data files on energy consumption in energy industries and manufacturing industries for 1980-1989, it has not been possible to recalculate emissions as has been done for different sectors for 1990 and onwards. Hence, time series for the eighties are not directly comparable with later years for NFR1B1c.

3.3.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Time series reported in NFR1B1c have been reviewed in later years and are considered to be consistent. Uncertainties for emissions are mostly expert estimates. For COG flaring, uncertainties in activity data are high since the amount of flared gas are not measured as carefully as combusted gas (this statement is true for any plant). Uncertainties for particulate emissions from handling of solid fuels are ±20% and based on expert judgement. More detailed information is to be found in Annex 1.

3.3.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

3.3.3.5 SOURCE-SPECIFIC RECALCULATIONS
Emissions of TSP, PM\(_{10}\) and PM\(_{2.5}\) from COG flaring have been corrected for 1990-2008 in submission 2018, resulting in ~0.9% – +2% change for this code.
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.

### 3.3.4 Hydrogen production plants at refineries, NFR1B2a i

#### 3.3.4.1 SOURCE CATEGORY DESCRIPTION

NFR1B2a i includes emissions from hydrogen production at refinery facilities – NOX, CO, NMVOC, SO2, NH3, TSP, PM10 and PM2.5 and BC. The summary of the latest key source assessment for aggregated emissions from NFR1B2a is presented in Table 3-39.

**Table 3-39. Summary of key source analysis, NFR1B2a, according to approach 1.**

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1B2a i,iv,v</td>
<td>Oil – NMVOC, SO2, DIOX</td>
<td>Oil – NMVOC, DIOX, SO2</td>
</tr>
</tbody>
</table>

#### 3.3.4.2 METHODOLOGICAL ISSUES

The first hydrogen production facility in Sweden was taken into operation in 2005, and in this facility, butane was used as a main raw material until 2011. Since 2011, several fuels are used as stock for hydrogen production, including liquefied natural gas (LNG) from 2014. The second hydrogen production facility is operative since 2006. At this facility, naphtha was used as raw material until 2011 and LNG from 2011 and onwards. Emissions from these facilities are reported in NFR1B2a i in accordance with 2006 IPCC Guidelines.

The Tier 2 method is used. Activity data is reported as NE because one of the plants has changed reporting method so that activity data no longer represents amounts of feedstock - a mixture of butane, off-gas from one of the refinery units, and LNG (from 2014). Instead, to calculate CO2 emissions reported to EU ETS, the facility from now one will use amounts of so called ‘PSA (pressure swing adsorption) gas’ - energy-poor off-gas from the hydrogen production unit. PSA gas is a good proxy for activity data for this particular plant with a complicated feedstock structure; however, it is not a feedstock and thus cannot be summed up with feedstock data (naphta for 2006 – 2011 and LNG from 2011 onwards) from the other plant.

The emissions are calculated using plant specific activity data and net calorific values. For natural gas, national emission factors are used, whereas national emission factors for “other petroleum fuels” were used for naphtha and PSA due to lack of specific emission factors for these fuels.

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119 Ortiz, C., et al.. Överlappande mellan CRF 1 och 2, SMED memorandum, 2017
The emissions of \( \text{SO}_2 \) from hydrogen production ended in 2011 when naphta was no longer used as a raw material in the production.

### 3.3.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for NFR1B2a i are displayed in Annex 1. Uncertainties are mostly expert estimates. According to data reported to the EU ETS, both hydrogen production plants use the level 2 method to measure activity data, which means that the activity data uncertainty is \( \pm 2.5 \% \) or less.

### 3.3.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

### 3.3.4.5 SOURCE-SPECIFIC RECALCULATIONS

Notation key for dioxin has been changed from NE to NA: there are no indications that these emissions occur at hydrogen production. Change of activity data for one of the plants resulted in emission changes within a range 55-96\% for the years 2012-2014, and -28\% for 2015 – for all substances. NOx emissions have been recalculated due to an emission factor correction, which resulted in 0.0002 – 0.06 kt (16-30\%) decrease for 2005-2015.

### 3.3.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

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

### 3.3.5 Refineries, NFR1B2a iv

#### 3.3.5.1 SOURCE CATEGORY DESCRIPTION

There are five refinery facilities in Sweden. Emissions from refineries reported in NFR1B2 i iv include:

- Fugitive emissions of \( \text{CH}_4 \) and NMVOC from the total refinery area;
- \( \text{SO}_2 \) emissions from desulphurisation (occur at four facilities);
- Emissions from make-up coke combustion (occur at three facilities) – NMVOC, \( \text{NH}_3 \), PAH and dioxin;
- Emissions from fluidized catalytic cracker (occur at one facility) – \( \text{SO}_2 \), TSP, PM\(_{10}\), PM\(_{2.5}\) and BC.

The summary of the latest key source assessment for aggregated emissions from NFR1B2a is presented in Table 3-40.
Table 3-40. Summary of key source analysis, NFR1B2a according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1B2a i,iv,v</td>
<td>Oil – NMVOC, SO$_2$, DIOX</td>
<td>Oil – NMVOC, DIOX, SO$_2$</td>
</tr>
</tbody>
</table>

3.3.5.2 METHODOLOGICAL ISSUES

Sweden estimates 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. Emissions are reported from catalytic cracking (TSP, PM$_{10}$, PM$_{2.5}$ and BC), desulphurisation (SO$_2$), and from the storage and handling of oil (NMVOC, CH$_4$). Catalytic cracking occurs at one plant in Sweden.

Emissions of SO$_2$ from desulphurisation decreased dramatically during the early 1990’s and have for most years after 1995 been between 0.4 and 1 Gg with the exception for 2006, when the emissions increased due to operational disturbances at one facility.

Particle emissions from catalytic cracking have only been estimated since 1990 due to lack of data for earlier years. The emissions from cracker have been obtained from the company. The particle size distributions have been estimated with expert judgement. The assumed size distribution is 95 % for PM$_{10}$ of TSP for 1990-2009 (from 2010 onwards PM$_{10}$ emissions are included in environmental reports) and 85 % for PM$_{2.5}$ of TSP for the whole time-series.

Fugitive emissions of NMVOC 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$^{120,121,122,123}$ and Statistics Sweden$^{124}$. The activity data, as throughput of crude oil, 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 3-41, the reported emissions as well as activity data can be seen.

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$^{120}$ Swedish EPA, 1990.
$^{121}$ Swedish EPA, 1994a.
$^{122}$ Swedish EPA, 1994b.
Table 3-41. Throughput of crude oil at oil refineries and estimated fugitive emissions of NMVOC (Mg) NFR1B2a iv.

<table>
<thead>
<tr>
<th>Year</th>
<th>Throughput of crude oil, Mg</th>
<th>Total emissions of NMVOC, Mg</th>
</tr>
</thead>
<tbody>
<tr>
<td>1988</td>
<td>15 600 000</td>
<td>16 088</td>
</tr>
<tr>
<td>1989</td>
<td>16 440 000</td>
<td>16 588</td>
</tr>
<tr>
<td>1990</td>
<td>17 330 000</td>
<td>14 160</td>
</tr>
<tr>
<td>1991</td>
<td>16 810 000</td>
<td>12 669</td>
</tr>
<tr>
<td>1992</td>
<td>17 870 000</td>
<td>10 745</td>
</tr>
<tr>
<td>1993</td>
<td>18 723 684</td>
<td>10 127</td>
</tr>
<tr>
<td>1994</td>
<td>18 192 000</td>
<td>8 765</td>
</tr>
<tr>
<td>1995</td>
<td>19 430 000</td>
<td>7 494</td>
</tr>
<tr>
<td>1996</td>
<td>20 305 000</td>
<td>9 443</td>
</tr>
<tr>
<td>1997</td>
<td>20 130 000</td>
<td>9 526</td>
</tr>
<tr>
<td>1998</td>
<td>20 254 000</td>
<td>9 294</td>
</tr>
<tr>
<td>1999</td>
<td>19 483 034</td>
<td>10 090</td>
</tr>
<tr>
<td>2000</td>
<td>20 253 120</td>
<td>11 225</td>
</tr>
<tr>
<td>2001</td>
<td>19 592 852</td>
<td>9 523</td>
</tr>
<tr>
<td>2002</td>
<td>19 681 182</td>
<td>9 923</td>
</tr>
<tr>
<td>2003</td>
<td>19 661 646</td>
<td>11 336</td>
</tr>
<tr>
<td>2004</td>
<td>20 611 941</td>
<td>8 707</td>
</tr>
<tr>
<td>2005</td>
<td>19 919 968</td>
<td>7 501</td>
</tr>
<tr>
<td>2006</td>
<td>20 050 576</td>
<td>8 068</td>
</tr>
<tr>
<td>2007</td>
<td>17 706 518</td>
<td>8 655</td>
</tr>
<tr>
<td>2008</td>
<td>20 420 061</td>
<td>8 359</td>
</tr>
<tr>
<td>2009</td>
<td>19 669 472</td>
<td>8 556</td>
</tr>
<tr>
<td>2010</td>
<td>20 278 888</td>
<td>8 693</td>
</tr>
<tr>
<td>2011</td>
<td>19 034 115</td>
<td>8 709</td>
</tr>
<tr>
<td>2012</td>
<td>21 021 566</td>
<td>8 381</td>
</tr>
<tr>
<td>2013</td>
<td>17 021 700</td>
<td>8 135</td>
</tr>
<tr>
<td>2014</td>
<td>19 320 478</td>
<td>7 710</td>
</tr>
<tr>
<td>2015</td>
<td>20 244 131</td>
<td>8 888</td>
</tr>
<tr>
<td>2016</td>
<td>19 976 308</td>
<td>7 441</td>
</tr>
</tbody>
</table>

Since submission 2009, emissions from combustion of cracker coke in refineries are reallocated from the energy sector to NFR1B2a iv (hence the combustion is not carried out for energy purposes). This was based on a study performed by SMED\textsuperscript{125}. 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. Combustion of cracker coke occurs at three facilities. Activity data as amount of cracker are taken from the company’s report to the EU ETS system. NO\textsubscript{X}, PAH, dioxins and

NMVOC are calculated with these plant specific activity data and national emission factors. SO\textsubscript{2} reported by one of the facilities originates from coke combustion as well – these emissions are obtained from the environmental reports.

### 3.3.5.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for NFR1B2a iv are displayed in Annex 1. Uncertainties are mostly expert estimates. Activity data uncertainties concern only cracker coke combustion (for other activities, emissions from environmental reports are used).

### 3.3.5.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

### 3.3.5.5 SOURCE-SPECIFIC RECALCULATIONS

Shares of NMVOC and CH\textsubscript{4} in the total reported VOC have been corrected, resulting in the decrease of NMVOC emissions by 0.15-0.34 kt for the whole time series. SO\textsubscript{2} emissions from one of the facilities have been corrected for 2015 resulting in 0.07 kt increase compared to submission 2017. NO\textsubscript{x} and CO previously reported as non-combustion emissions from catalytic cracker have been removed from NFR1B: an investigation and contacts with the facility have shown that these emissions are reported in the energy sector (combustion of refinery gas and other fuels) and shouldn’t be double-counted. NO\textsubscript{x} (comes from fuel combustion) is reported as NA. CO is reported as IE since part of it originates from raw material but all emitted CO is reported in the energy sector.

### 3.3.5.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

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

### 3.3.6 Gasoline handling and distribution, NFR1B2a v

#### 3.3.6.1 SOURCE CATEGORY DESCRIPTION

NFR1B2a v includes fugitive emissions of NMVOC from the storage of oil products (oil depots) and from gasoline handling at gasoline stations. The summary of the latest key source assessment for aggregated emissions from NFR1B2a is presented in Table 3-42.

#### Table 3-42. Summary of key source analysis, NFR1B2a, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1B2a i,iv,v</td>
<td>Oil – NMVOC, SO\textsubscript{2}, DIOX</td>
<td>Oil – NMVOC, DIOX, SO\textsubscript{2}</td>
</tr>
</tbody>
</table>
3.3.6.2 METHODOLOGICAL ISSUES

3.3.6.2.1 Gasoline stations

The calculation of the NMVOC time series for fugitive emissions from gasoline distribution, 1988-2016, is based on methods given by Concawe\textsuperscript{126}, 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 stations\textsuperscript{127}. The basic assumptions are presented in Table 3-43; ethanol both for use in blends with gasoline and for use unblended is included in the reported gasoline volumes.

Table 3-43. Assumptions for calculating fugitive emissions from the handling and distribution of gasoline.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Assumption</th>
</tr>
</thead>
<tbody>
<tr>
<td>Density of gasoline</td>
<td>730 kg/m(^3) 1988 - 1996</td>
</tr>
<tr>
<td></td>
<td>750 kg/m(^3) 1997 -</td>
</tr>
<tr>
<td>Distribution of gasoline to gas stations</td>
<td>0.16 % of distributed volume</td>
</tr>
<tr>
<td>Spill</td>
<td>0.01 % of distributed volume</td>
</tr>
<tr>
<td>Filling of car tanks</td>
<td>0.18 % of filled volume</td>
</tr>
<tr>
<td>Measures at distribution to gas station</td>
<td>90 % Efficiency of measures</td>
</tr>
<tr>
<td>Measures at filling cars</td>
<td>70 % Efficiency of measures</td>
</tr>
</tbody>
</table>

The measures at distribution and filling were introduced over a period of time from 1991-1994, to the extent presented in Table 3-44. The amount of gasoline sold at large and small gas stations, respectively, was assumed to be 50/50 for the years 1988-1994. Data on the distributed amounts of gasoline is taken from national statistics from Statistics Sweden for submission 2016 and onwards, Table 3-45. For earlier submissions data on the distributed amounts of gasoline was taken from the HBEFA model.

Table 3-44. Fraction of gasoline stations with technical measures installed.

<table>
<thead>
<tr>
<th>Year</th>
<th>Large gas stations &gt;2000 m(^3)</th>
<th>Small gas stations</th>
</tr>
</thead>
<tbody>
<tr>
<td>1988 - 1990</td>
<td>0%</td>
<td>0%</td>
</tr>
<tr>
<td>1991</td>
<td>50%</td>
<td>0%</td>
</tr>
<tr>
<td>1992</td>
<td>75%</td>
<td>25%</td>
</tr>
<tr>
<td>1993</td>
<td>100%</td>
<td>75%</td>
</tr>
<tr>
<td>1994 -</td>
<td>100%</td>
<td>100%</td>
</tr>
</tbody>
</table>

\textsuperscript{126} Concawe, 1986. Hydrocarbon emissions from gasoline storage and distribution systems, Report No 85/54.

\textsuperscript{127} Andersson, 2000.
3.3.6.2.2 Oil depots

Calculated fugitive emissions of NMVOC from the storage of oil products have been obtained from SPI\textsuperscript{128} or from the environmental reports of the oil depots. The calculations are based on the amount of gasoline handled in the depots. The calculations cover the years 1990 and onwards and are based on methods given by Concawe 85/54\textsuperscript{129} for the years 1990-2006 and on Concawe 03/07\textsuperscript{130} for the years 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 depots the reported NMVOC emissions are based on emission measurements in the depot areas and not on calculations based on the amount gasoline handled in the depots. For some years, for which no data was provided, emissions were by using interpolation. Handled amount of gasoline and fugitive emissions of NMVOC from depots and gasoline stations are presented in Table 3-45.

Table 3-45. Handled and distributed amount of gasoline and estimated fugitive emissions of NMVOC (Gg) from storage at depots and at gasoline stations.

<table>
<thead>
<tr>
<th>Year</th>
<th>Volume of gasoline and ethanol m\textsuperscript{3}</th>
<th>Fugitive emissions of NMVOC at depots Gg</th>
<th>Fugitive emissions of NMVOC at gasoline stations, Gg</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>5 629 582</td>
<td>2.48</td>
<td>14.38</td>
</tr>
<tr>
<td>1995</td>
<td>5 762 693</td>
<td>1.93</td>
<td>3.37</td>
</tr>
<tr>
<td>2000</td>
<td>5 372 801</td>
<td>2.07</td>
<td>3.22</td>
</tr>
<tr>
<td>2005</td>
<td>5 508 187</td>
<td>2.31</td>
<td>3.30</td>
</tr>
<tr>
<td>2006</td>
<td>5 363 256</td>
<td>2.47</td>
<td>3.22</td>
</tr>
<tr>
<td>2007</td>
<td>5 253 617</td>
<td>2.35</td>
<td>3.15</td>
</tr>
<tr>
<td>2008</td>
<td>4 928 238</td>
<td>2.53</td>
<td>2.96</td>
</tr>
<tr>
<td>2009</td>
<td>4 861 776</td>
<td>2.41</td>
<td>2.92</td>
</tr>
<tr>
<td>2010</td>
<td>4 550 207</td>
<td>2.21</td>
<td>2.73</td>
</tr>
<tr>
<td>2011</td>
<td>4 222 863</td>
<td>2.47</td>
<td>2.53</td>
</tr>
<tr>
<td>2012</td>
<td>3 880 860</td>
<td>2.45</td>
<td>2.33</td>
</tr>
<tr>
<td>2013</td>
<td>3 685 278</td>
<td>2.20</td>
<td>2.21</td>
</tr>
<tr>
<td>2014</td>
<td>3 545 193</td>
<td>1.93</td>
<td>2.13</td>
</tr>
<tr>
<td>2015</td>
<td>3 471 386</td>
<td>2.02</td>
<td>2.08</td>
</tr>
<tr>
<td>2016</td>
<td>3 338 553</td>
<td>1.93</td>
<td>2.00</td>
</tr>
</tbody>
</table>

\textsuperscript{128} Per Brännström, 2009-, personal communication; Leif Ljung -2009, personal communications

\textsuperscript{129} Concawe, 1986, Hydrocarbon emissions from gasoline storage and distribution systems, Report No 85/54.

\textsuperscript{130} Concawe Report No. 3/07, Air pollutant emission estimation methods for E-PRTR reporting by refineries
3.3.6.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Time series reported in NFR 1B2a v have been reviewed in later years and are considered to be consistent. Uncertainties for NMVOC emissions reported in NFR 1B2a v are ± 70% and based on expert judgement. More detailed information is to be found in IIR Annex 1.

3.3.6.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

3.3.6.5 SOURCE-SPECIFIC RECALCULATIONS

No recalculations have been performed in NFR1B2a v in submission 2018.

3.3.6.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

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

3.3.7 Transmission and distribution of natural gas and gasworks gas – fugitive emissions, 1B2b

3.3.7.1 SOURCE CATEGORY DESCRIPTION

NFR1B2b includes fugitive emissions of NMVOC from transmission and storage of natural gas and biogas (1B2b iv) and from distribution of natural gas, biogas and gasworks gas (1B2b v). The summary of the latest key source assessment is presented in Table 3-46.

Table 3-46. Summary of key source analysis, NFR1B2b, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>1B2b</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

3.3.7.2 METHODOLOGICAL ISSUES

In 2013, a national method for estimating fugitive emissions of natural gas and gasworks gas has been developed for the period 1990 to 2012\textsuperscript{131}. Emission of NMVOC from this subsector is for the first time reported in submission 2014.

3.3.7.2.1 Transmission and storage of natural gas 1B2b iv

Emission estimates for gas transmission and storage are based on information provided by Swedegas, the operator of the gas transmission pipeline and storage in Sweden.

\textsuperscript{131} Jerksjö M., Gerner A., Wängberg I. 2013
The Swedish network for gas storage and transmission includes several different types of facilities: metering and regulation stations (M/R stations), compressor stations, ramification stations, valve stations, pig launcher & receiver stations, and a storage facility. According to Swedegas, many of the facilities are combined, e.g., valves located close to M/R stations. To enable biogas transmission in the network, two compressor stations were put into operation in 2014 – one combined with M/R station and one stand-alone facility.

In 2016, the method for estimating the emissions from the gas transmission network was revised since new measurements of methane emissions became available. Methane leakage rates per hour have been measured at all major types of facilities. Estimated emission factors (see Table 3-47 below) have been applied to the number of facilities of each type.

### Table 3-47. Method for estimation of gas leakage from the national gas transmission network.

<table>
<thead>
<tr>
<th>Facility type</th>
<th>CH₄ EF g/hour</th>
<th>Number of facilities in 2016</th>
<th>Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>M/R station</td>
<td>91</td>
<td>42</td>
<td>Number of facilities is known for the whole time series</td>
</tr>
<tr>
<td>Storage</td>
<td>200</td>
<td>1</td>
<td>In operation since 2006</td>
</tr>
<tr>
<td>M/R + compressor station</td>
<td>222</td>
<td>1</td>
<td>In operation since 2014</td>
</tr>
<tr>
<td>Compressor station</td>
<td>100</td>
<td>1</td>
<td>In operation since 2014</td>
</tr>
<tr>
<td>Valve station</td>
<td>30</td>
<td>26</td>
<td>For the years 1990-2014, the number of facilities is assumed to be in direct proportion to the network’s length (320 km in 1990, 620 km in 2016)</td>
</tr>
<tr>
<td>Pig launcher &amp; receiver station</td>
<td>300</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Ramification station</td>
<td>30</td>
<td>39</td>
<td></td>
</tr>
</tbody>
</table>

Methane emissions have been further calculated to NMVOC emissions by using parameters shown in Table 3-48. Information on gas composition was obtained from Swedegas and constitutes average values from the period 2006 to 2012.

### Table 3-48. Composition and physical properties of natural gas.

<table>
<thead>
<tr>
<th>Property</th>
<th>Unit</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methane content in natural gas</td>
<td>% by weight</td>
<td>78.6</td>
</tr>
<tr>
<td>NMVOC content in natural gas</td>
<td>% by weight</td>
<td>19.0</td>
</tr>
<tr>
<td>Density of natural gas</td>
<td>kg/Nm³</td>
<td>0.817</td>
</tr>
<tr>
<td>Density of methane</td>
<td>kg/Nm³</td>
<td>0.716</td>
</tr>
</tbody>
</table>

132 Bjur & Lindsjö, 2016
133 Jerksjö, M., Salberg, H. 2016. Mätningar av metanläckage längs svenska naturgasnätets stamlä- ning, IVL report C202 (in cooperation with Fluxsense)
Emissions earlier reported as gas leakage have been re-allocated to the sector NFR1B2c Venting and flaring since these emissions are controlled and associated with regular network maintenance work rather than with uncontrolled gas leakage.

3.3.7.2.2 Natural gas distribution 1B2b v)

There are three types of gas networks for distribution of gas in Sweden.

1. The gas network for distribution of natural gas
2. Local biogas distribution network

The gas network for distribution of natural gas is connected to the national transmission pipeline via M/R stations as mentioned above and had a total length of 2620 km in year 2012. This network delivers natural gas to the end users, which are industries or municipalities which in turn use the gas for energy production, to feed their town gas networks, etc. There are about 40 small local distribution networks for biogas in Sweden\textsuperscript{134}. The total length was 146 km in 2012. The biogas is of similar quality as natural gas and is distributed in similar distribution pipes as natural gas.

Most of the gasworks gas networks use natural gas and their distribution system has been modernised and considered to be of the same standard as the distribution system for natural gas. However, the gasworks gas networks in Stockholm and Gothenburg (the two largest cities in Sweden) are different. These networks consist to a large part of old pipes with considerable high leaking rate. Between 1990 and 2011, a facility in Stockholm produced gasworks gas from cracking light petroleum. In 2011, they started to use a mixture of natural gas and air. The city of Gothenburg produced gasworks gas of a similar quality as that in Stockholm during the period 1990 – 1993. In 1993, the city of Gothenburg shifted to a mixture of natural gas and air and since the beginning of 2011, only pure natural gas is distributed in Gothenburg. Activity data in terms of leakage of gasworks gas has been obtained from the gasworks gas distributor in Stockholm for the years 2002-2012. For earlier years, only production data is available, and the average relation of leakage to production has been used to estimate leakage for the years 1990-2001. The emissions of NMVOC have been calculated with data on chemical composition of gas from cracking and natural gas/air mixture. The methodology is described in Jerksjö et al\textsuperscript{135}.


Since no measurement on fugitive methane emissions from distribution of gas has been made in Sweden, emission factors found in the literature were compared and examined. Information on the Swedish gas network was collected by contacting the operators. Based on this information an emission factor obtained from a Dutch investigation (Wikkerlink 2006\textsuperscript{136}) was chosen. The emission factor is the result of an evaluation of data from measurements of gas leaks at several places in the Netherlands and is equal to 120 Nm$^3$ methane per km distribution line. According to net operators of new or renewed Swedish networks for natural gas, the networks in Sweden are of similar standard and design as those in the Netherlands. The Dutch emission factor is considered to be valid for pipes made from PVC and polyethylene, etc., and can be used as an average value covering different pressure regimes. The emission factor from the Dutch study was adopted for estimating the methane emissions from Swedish gas networks 1. (Natural gas) and 2. (Biogas) and also gas networks in cities with new or renewed distribution systems.

Data on gas mixtures, sources of activity data and emission factors used for emission calculations in NFR1B2b v for each gas distribution network are summarized in Table 3-49.

Table 3-49. Summary of method for calculating emissions from Swedish gas distribution networks.

<table>
<thead>
<tr>
<th>Gas distribution networks</th>
<th>Natural gas*</th>
<th>Local biogas</th>
<th>Gasworks gas – Stockholm</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gas mixture used</td>
<td>Natural gas</td>
<td>Biogas of similar quality as natural gas</td>
<td>Mixture of natural gas and air. Until 2011 – gasworks gas and mixture of natural gas and air</td>
</tr>
<tr>
<td>Source of activity data</td>
<td>Gas distribution companies</td>
<td>Grönmij. 2009</td>
<td>Stockholm gas environmental reports</td>
</tr>
<tr>
<td>Type of activity data</td>
<td>km length</td>
<td>km length</td>
<td>Nm$^3$ gas leakage</td>
</tr>
<tr>
<td>Emission factor for NMVOC</td>
<td>No emission factors are used. Emissions are calculated based on estimated methane emissions and the content of CO$_2$ and NMVOC in the natural gas.</td>
<td>No emission factors are used. Emissions are calculated based on the content of NMVOC in the gas mixtures considered</td>
<td></td>
</tr>
</tbody>
</table>

* Including a number of city gas distribution networks, for instance Gothenburg gas distribution network since 2011.

The gas distribution networks in Stockholm and Gothenburg constitute of both old and new or re-lined pipes. The old pipes have a relatively high leaking rate. During 1990 to January 2011 gasworks gas (i.e. from cracking of light petroleum) was produced and distributed in the Stockholm gas network. In January 2011 one started to

\textsuperscript{136} Wikkerlink. 2006.
use a mixture of natural gas and air. The city of Gothenburg produced gasworks gas of a similar quality as that in Stockholm during the period 1990 – 1993. In 1993, the city of Gothenburg shifted to a mixture of natural gas and air and since the beginning of 2011, only pure natural gas is distributed in Gothenburg.

The fugitive emissions from distribution of gasworks gas in Stockholm and Gothenburg have been estimated from statistics on production of gasworks gas and natural gas air mixtures and leakage rates from Stockholm Gas. The content of NMVOC in gasworks gas and natural gas air mixture is shown in Table 3-50.

### Table 3-50. NMVOC content in gasworks gas.

<table>
<thead>
<tr>
<th>Property</th>
<th>Unit</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMVOC in gasworks gas</td>
<td>kg/Nm³</td>
<td>0.04</td>
</tr>
<tr>
<td>NMVOC in natural gas air mixture</td>
<td>kg/Nm³</td>
<td>0.08</td>
</tr>
</tbody>
</table>

3.3.7.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

*Gas transmission:* The associated emission uncertainty is ±50 % according to expert estimates. More detailed information is to be found in IIR Annex 1.

*Gas distribution:* Fugitive emissions from the distributing network in Stockholm constitute 80 – 90 % of the total emissions from gas distribution in Sweden. The emission data from the Stockholm distribution network is based on measurements provided by the operator and the associated uncertainty is estimated to ± 50 %. The total uncertainty concerning distribution of gas in Sweden is largely influenced by the contribution from the gas network in Stockholm, and is thus likewise estimated to ± 50 %. More detailed information is to be found in IIR Annex 1.

3.3.7.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

3.3.7.5 SOURCE-SPECIFIC RECALCULATIONS

No recalculations have been performed in NFR1B2b in submission 2018.

3.3.7.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

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

3.3.8 Venting and flaring, NFR1B2c

3.3.8.1 SOURCE CATEGORY DESCRIPTION

NFR1B2c includes emissions of NMVOC from gas venting as well as emissions from flaring of gas and/or oil products at refineries, three iron and steel plants, one pulp industrial plant, as well as during the national gas network maintenance (NOx, CO, NMVOC, SOx, NH3, TSP, PM10, PM2.5, and BC). The summary of the latest key source assessment is presented in Table 3-51.

**Table 3-51. Summary of key source analysis, NFR1B2c, according to approach 1.**
3.3.8.2 METHODOLOGICAL ISSUES
For flaring of oil products and refineries and other industrial sites, activity data has been collected directly from the plant operators. For 2007 and onwards, activity data is mainly taken from the EU ETS system. Plant specific net calorific values are used when available. The same emission factors are used as for stationary combustion.

The subcategory also includes occasional emissions from venting and flaring of natural gas during a network inspection at Swedegas. Network inspection conducted once in eight years (sometimes more often) requires so called piggling – emptying M/R stations, which means release of certain amounts of natural gas. A larger part of the released gas is flared but some is vented. For the years 2014-2016, estimated amounts of gas vented during the inspections have been obtained from the operator. For the years 2006, 1998 and 1990 estimates were made based on the relation of the amount of vented gas to the number of M/R stations in 2014-2015. Emissions of NMVOC from piggling are ~0.13-0.42 t.

Beside piggling operations, there is certain amount of gas annually vented to the atmosphere (earlier reported as diffuse emissions) from M/R stations – about 13 Nm³ gas per station, or about 1 tonne NMVOC in total per year. Much higher and much more varying venting emissions come from the storage facility (put into operation in 2006) – from 6 to 170 t NMVOC per year, depending on how well the compressor worked. In 2013 the compressor failure result in particularly high emissions.

3.3.8.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Estimates of emissions from natural gas venting are provided by the operator. The associated uncertainty is ± 50 % according to expert estimates. The activity data uncertainty for flaring of different fuels at refineries and other industrial facilities is as reported to EU ETS and is estimated to ±17.5 %. For gas and oil flaring, the total emission uncertainties are affected by uncertainties in the emission factors, which are the same as for industrial combustion. More detailed information is to be found in IIR Annex 1.

3.3.8.4 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

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137 Hellström 2013-2015
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. These facilities are regularly asked to verify that the default value is still valid.

3.3.8.5 SOURCE-SPECIFIC RECALCULATIONS
Corrections of types of gases flared at refineries resulted in minor increase of SO₂ (0.0004 – 0.0016 kt), NMVOC (0.0002 – 0.0008 kt), NH₃ (0.0002 – 0.0008 kt), TSP (0.0009 – 0.004 kt), PM₁₀ (0.0009 – 0.004 kt), PM₂.₅ (0.0009 – 0.004 kt), and BC (0.0002 – 0.001 kt) for 2007-2015. Emissions of NOₓ and CO have decreased by 0.004 – 0.016 kt and 0.001 – 0.004 kt, respectively, for the same time period, compared to Submission 2017.

3.3.8.6 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 Industrial processes and product use (NFR sector 2)

4.1 Overview

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 from industrial processes include the cement industry, primary aluminium production and some processes in the chemical industry.

Generally three sources of information concerning activity and emission data for the industrial process sector have been used:

- Emission data as reported annually by facilities in legally required environmental reports to the authorities.
- National production statistics or similar information at national level.
- Plant specific data collected by direct contacts with facilities

Under Swedish environmental laws, operators performing environmentally hazardous activities that require a permit by law are obliged to compile and submit an annual environmental report to their supervisory authority. The environmental report consists of three parts:

- Basic identification information about the facility.
- Text section (for example, a description of the facility and the processes, the use of energy, chemicals and raw materials, emissions and conditions in the permit).
- Emission declaration (for example, production data, fuel consumption data, emission data and, in some cases, information on how emission data has been determined).

Data in the environmental reports often originate from measurements or mass balances. The use of default emission factors is limited. Only operators that exceed the thresholds for the substances, listed in the Swedish environmental law concerning environmental reports\(^{138}\), are obliged to compile the emission declaration.

The County Administrative Boards audit the data presented in the operators’ environmental reports. Since the beginning of year 2007 environmental reports can be submitted electronically via the Swedish Portal for Environmental Reporting (SMP)\(^{139}\). This database includes not only emissions, but also basic information

\(^{138}\) Swedish EPA, NFS 2000:13, Naturvårdsverkets föreskrifter om miljörapport för tillståndspikliga miljöfarliga verksamheter.

\(^{139}\) Swedish Portal for Environmental Reporting, Svenska Miljörapporteringssportalen. https://smp.lansstyrelsen.se/
about the facilities, such as their activity code (national code system, adjustment of NACE four digits), IPPC code and permit, location coordinates, etc. The procedure for updating the database is not regulated by legislation, which results in some incompleteness and inconsistencies in the database.

The use of emission factors in the Swedish inventory for industrial processes is limited and, when used, they are nationally derived or specific for a facility. Where there are a large number of companies within a specific sector, and when all environmental reports are not available, a combination of information from environmental reports and production statistics on the national level, are used to estimate the sector's emissions on a national scale.

Emissions of PM\textsubscript{10} and PM\textsubscript{2.5} are in many cases calculated as a fraction of TSP. An overview of the fractions that are used within the sector are included in Annex 3.2.

Emissions of metals have not been reported for the period 1980 - 1989 due to lack of reliable data for some important sources.

Sweden’s emission inventory is in accordance with EMEP/EEA Air Pollutant Emission Inventory Guidebook\textsuperscript{140}, 2006 IPCC Guidelines\textsuperscript{141}, and the Guidelines for Estimating and Reporting Emission Data under the Convention on Long-range Transboundary Air Pollution\textsuperscript{142}.

4.2 Mineral products, NFR2A

4.2.1 Cement production, NFR2A1

4.2.1.1 SOURCE CATEGORY DESCRIPTION

Cement production occurs at three facilities in Sweden, with one being dominant. Emission data is taken from environmental reports and by direct contacts with the facilities. Calculation methods have been discussed with the industry.

Table 4-1. Summary of key source analysis, NFR2A1, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2A1</td>
<td>PM\textsubscript{2.5}</td>
<td>SO\textsubscript{2}, PM\textsubscript{10}, TSP, PM\textsubscript{2.5}</td>
</tr>
</tbody>
</table>

\textsuperscript{140} The EMEP/EEA Guidebook: http://www.eea.europa.eu/themes/air/emep-eea-air-pollutant-emission-inventory-guidebook/emep


\textsuperscript{142} UNECE 2003, Guidelines for Estimating and Reporting Emission Data under the Convention on Long-range Transboundary Air Pollution. Air Pollution Studies No. 15.
4.2.1.2 METHODOLOGICAL ISSUES

All three cement-producing facilities (owned by one company) are covered in the reported estimates and the missions have been estimated based on direct information from the company or from environmental reports. Emissions originate mainly from fuel combustion and less from industrial processes. Hence IE is reported for NO\textsubscript{x} in NFR 2A1 and emissions are reported in NFR 1A2f.

SO\textsubscript{2} emissions are allocated to industrial processes. Reported SO\textsubscript{2} emissions for 2016 have slightly decreased compared to 2015, however emissions since 1990 have decreased substantially. Emissions of TSP, PM\textsubscript{10} and PM\textsubscript{2.5}, also allocated to industrial processes and product use, have been estimated for the time period 1980 – 2016.

Reported emissions of TSP, PM\textsubscript{10} and PM\textsubscript{2.5} for 2010 to 2016 are substantially lower than average emissions in the 2000’s. The decreased emissions are due to the installation of a new dust filter at the largest site in 2010. BC emissions are reported for 2000 onwards and are calculated as a fraction of PM\textsubscript{2.5} according to EMEP/EEA Guidebook 2013.

NH\textsubscript{3} emissions arise partly due to the selective non-catalytic reduction (SNCR) of NO\textsubscript{x} where NH\textsubscript{3} is injected and partly from the raw material. Emissions have been included for the years that such an SNCR has been installed at respective plant. To ensure double reporting does not occur, reported emissions in the energy sector are subtracted from total NH\textsubscript{3} emissions reported by the company. There is a substantial variation in NH\textsubscript{3} emissions over the time series, which is due to variations in ammonia slip and the fact that emissions in the energy sector are subtracted from the total as reported by the company.

Emissions of heavy metals, PAH-4 and dioxins from the fuels used are calculated based on energy statistics and reported in the energy sector (NFR1).

Table 4-2 gives an overview of the allocation of pollutants from cement production.
Table 4- 2. Allocation of pollutants from cement production.

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>NFR</th>
</tr>
</thead>
<tbody>
<tr>
<td>NO\textsubscript{X}</td>
<td>1A2f</td>
</tr>
<tr>
<td>NMVOC</td>
<td>1A2f</td>
</tr>
<tr>
<td>SO\textsubscript{2}</td>
<td>2A1</td>
</tr>
<tr>
<td>NH\textsubscript{3}</td>
<td>2A1 and 1A2f</td>
</tr>
<tr>
<td>PM\textsubscript{2.5}</td>
<td>2A1</td>
</tr>
<tr>
<td>PM\textsubscript{10}</td>
<td>2A1</td>
</tr>
<tr>
<td>TSP</td>
<td>2A1</td>
</tr>
<tr>
<td>BC</td>
<td>NE</td>
</tr>
<tr>
<td>CO</td>
<td>1A2f</td>
</tr>
<tr>
<td>Metals</td>
<td>1A2f</td>
</tr>
<tr>
<td>PAH-4</td>
<td>1A2</td>
</tr>
</tbody>
</table>

4.2.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions are based on expert judgement. Assessed uncertainties for SO\textsubscript{2}, NO\textsubscript{X} and NH\textsubscript{3} are ± 20%, ± 50%, and ± 400%, respectively. For TSP, PM\textsubscript{10} and PM\textsubscript{2.5}, uncertainties are estimated to ± 30% each. More information is given in IIR Annex 1.

Time series from cement production reported in NFR2A1 have been reviewed in later years and are considered to be consistent.

4.2.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

4.2.1.5 SOURCE-SPECIFIC RECALCULATIONS

No recalculations have been performed in NFR2A1 in submission 2018.

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, NFR2A2

4.2.2.1 SOURCE CATEGORY DESCRIPTION

Emissions of SO$_2$, particulate matter and BC from lime production are reported in NFR2A2. 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 since 1990 (about 440 kt) and peaked in 2005 (about 730 kt). In 2009 there was a large decrease in lime production due to the global economic recession. In 2016, there was a slight increase in production compared to 2015 with quantities of around 574 kt, marking however a considerably lower production compared to years 2010-2013.

Table 4-3. Summary of key source analysis, NFR2A2, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2A2</td>
<td>$PM_{10}$, $SO_2$</td>
<td>$PM_{10}$, $PM_{2.5}$, $SO_2$, TSP</td>
</tr>
</tbody>
</table>

4.2.2.2 METHODOLOGICAL ISSUES

Emissions of SO$_2$ from 1990 have been estimated for production of quick lime. The estimations from quick lime production are calculated using emission factors presented in environmental reports by one of the producers$^{143}$. The emission factor provided by the lime producer is substantially higher for 2008 than for earlier years. This resulted in an increase of reported SO$_2$ emissions for 2008 compared to earlier years. However in 2009 the reported SO$_2$ emissions were again on the same level as before 2008 due to less use of lime. For 2009-2016 the emission factor for 2008 has been used for the estimation of emissions of SO$_2$ due to lack of more recent information in the environmental reports.

Emissions of SO$_2$ from quick lime production intended for the pulp and paper industry are, as in earlier submissions, not included in the estimates reported in NFR2A2 but are reported in NFR2H1. SO$_2$ emissions from quick lime production within carbide production are from submission 2015 onwards included in NFR2A2.

Emissions of particles from lime production have also been estimated. Estimates were made for the period from 1980 onwards and emissions were calculated using emission factors presented in environmental reports by one of the producers and size fractions based on expert judgement.

$^{143}$ Nordkalk, http://www.nordkalk.com
For corresponding emissions from the production of other lime products, emission factors published by the CEPMEIP project\(^{144}\) were used. For particle emissions from quicklime production within carbide production, emissions are obtained from the company’s environmental report.

BC emissions are included as of submission 2015 and are calculated as a fraction of PM\(_{2.5}\) according to EMEP/EEA Guidebook 2013.

From 2005 onwards, facility-specific EU ETS data is used for activity data estimations, where the amount of burnt lime is calculated based on CO\(_2\) emissions. In previous submissions, statistics from the Swedish Lime Association have been used for activity data for the entire time series, however, the statistics have in recent years not been produced in time, resulting in the need of an alternative data source. In a study carried out in 2015\(^{145}\), different data sources were compared and it was concluded that EU ETS data is the more reliable one. For 1990-2004, statistics from the Swedish Lime Association is used for activity data. Although different data sources are used over the time series, it was concluded that both sources provide similar results and the time series can be considered consistent.

4.2.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions are based on expert judgement. Assessed uncertainty for SO\(_2\) is ± 20% and ± 50% for TSP, PM\(_{10}\) and PM\(_{2.5}\). More information is given in IIR Annex 1.

Time series from lime production reported in NFR code 2A2 have been reviewed in later years and are consistent.

4.2.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

4.2.2.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been performed in NFR2A2 in submission 2018.

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.


\(^{145}\) Mawdsley, I. 2015. Change of activity data for lime production
4.2.3 Glass production, NFR2A3

4.2.3.1 SOURCE CATEGORY DESCRIPTION

In Sweden there is one facility for container glass production and several small facilities for manual glass production. In addition, emissions from one glass wool producer are included. From glass wool production, emissions of NH$_3$, NMVOC, particles and BC are reported.

From the float glass production, the total emissions of SO$_2$ and NO$_X$ from the glass furnace are allocated to 2A3 since a separation in energy-related and process-related emissions is not possible. From the container glass production, SO$_2$ emissions originating from the raw material and small amounts of NMVOC are reported. Emissions of Cu, Se and Ni from the float glass production and emissions of Se and Ni from the container glass production are reported from Submission 2016 onwards.

The only producer of float glass shut down in 2013. The shut-down of the float glass production facility resulted in very low emissions for 2013 as this is the only source of NO$_X$ emissions in NFR2A3. Also total emissions of SO$_2$ and particles in 2A3 decreased to a small extent in 2013-2014 due to the ceased production of float glass. In 2014, NO$_X$ emissions in 2A3 are reported “NA”.

Emissions of particulate matter have been reported from the production of container and manual glass for the period 1990-2013, whereas particle emissions from float glass production are reported for the time period 1980–2012 until production shut down. BC emissions are reported for glass and glass wool production for year 2000 onwards and heavy metals from glass production are reported 1990-2016.

No data regarding CO, dioxin, PAH and HCB emissions are available, and the time series from 1990 and onwards are thus reported NE, in accordance with EMEP/EEA Guidebook 2013.

All other emissions from the glass production facilities originate from combustion for energy purposes, and are allocated to the Energy sector (NFR1).

Table 4-4. Summary of key source analysis, NFR2A3, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2A3</td>
<td>Se, Ni, SO$_2$,</td>
<td>Se, NH$<em>3$, PM$</em>{10}$, TSP, PM$_{2.5}$, As, NO$_X$</td>
</tr>
</tbody>
</table>

4.2.3.2 METHODOLOGICAL ISSUES

The emission data sources for glass production are mixed. Some data derives from reports from the Swedish EPA and some have been received from the companies’ environmental reports or from data bases containing data from the environmental
In informative Inventory Report Sweden 2018 reports. For earlier years in the time series emission data are from national reporting to HELCOM and from the Swedish EPA. Data for missing years and parameters have been estimated or interpolated.

Emission factors for Ni, Se and Cu are obtained from EMEP/EEA Guidebook 2013 and applied to activity data. Activity data, produced amount of float and container glass, has been acquired from the annual environmental reports for the two major producing facilities.

Manual glass production used to be an important source of lead emissions. In the early 1990's emissions of lead from the production of manual glass represented roughly 80% of the total reported lead emission from glass production. Ten years later, the manual glass production adds up to only around 10% of the lead emissions from glass production. This reduction is probably due to the reduced production of lead crystal glass. Today no lead is used in the production of manual glass.

4.2.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions are based on the guidebook 2008 or 2013 and expert judgement. More detailed information is to be found in IIR Annex 1.

Time series from glass production reported in NFR code 2A3 have been reviewed in later years and are consistent.

4.2.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

4.2.3.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been performed in NFR2A3 in submission 2018.

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 Quarrying and mining of minerals other than coal, NFR2A5a

4.2.4.1 SOURCE CATEGORY DESCRIPTION
Reported emissions include estimates for quarrying and mining of minerals other than coal.

The use of mining explosives causes emissions of mainly nitrogen oxides, NOx. The dominating source for emissions of particulate matter are limestone quarrying, crushing and grinding, re-allocated from NFR2A2 in Submission 2016. Ore dressing plants are the dominating source of heavy metals. Data on emissions of NOx,

particulate matter and heavy metals are mainly collected from the companies' environmental reports to the authorities.

Table 4-5. Summary of key source analysis, NFR2A5a, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2A5a</td>
<td></td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

4.2.4.2 METHODOLOGICAL ISSUES

NOX emissions originating from the use of mining explosives are only reported for the years 2002 - 2016 due to lack of data for earlier years. Emissions of particles are reported for the time period from 1980 and for metals from 1990. Emissions of particles from limestone quarrying, crushing and grinding are re-allocated from NFR2A2 in Submission 2016.

4.2.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions of NOx, TSP, PM10, PM2.5 are based on expert judgement. Assessed uncertainties for are ± 100%, ± 49%, ± 49% and ± 49%, respectively.

Uncertainties for emissions for metals are based on data in the Guidebook 2008. Uncertainty for Pb, Cd, Hg, As, Cr, Cu, Ni and Zn are ± 50%, ± 100%, ± 1000%, 100%, ± 100%, ± 50%, ± 50% and ± 50%, respectively.

More information is given in IIR Annex 1.

Time series from quarrying and mining of minerals other than coal reported in NFR code 2A5a have been reviewed in later years and are consistent.

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

For quarrying and mining of minerals other than coal, NFR2A5a, TSP emissions for 2015 were updated due to new information from the company resulting in an increase of 0.01%. No other recalculations have been performed in NFR2A5a in submission 2018.

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 Construction and demolition, NFR2A5b

4.2.5.1 SOURCE CATEGORY DESCRIPTION
Reported emissions include estimates for construction and demolition (2A5b). Emissions of particles from construction work are reported. The basis for the calculations is national data on construction activity. Emission factors from the CEPMEIP study are used\textsuperscript{148}.

No data concerning the NMVOC emissions are available, and the time series from 1990 and onwards is thus reported NE, in accordance with EMEP/EEA Guidebook 2013.

Table 4-6. Summary of key source analysis, NFR2A5b, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2A5b</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

4.2.5.2 METHODOLOGICAL ISSUES
The data chosen as a basis for the particle emission estimates from construction work are national statistics on building permits for housing and non-residential buildings (in m\textsuperscript{2})\textsuperscript{149}, 1996 - 2016, and economic statistics on annual investments in construction-related activities\textsuperscript{150}, 1980 - 2002. As only information on economic investments used for construction work are available for the years 1980 - 1995 this information had to be transformed into a unit where emission factors can be used. For the calculations of the time series of emissions, the economic information was normalised to the 1995 level, and the costs per square meter was assumed to be constant through-out the time series. The investments in construction work in 2002 expressed in SEK and normalised to the 1995 level was used as the base year for the transformation of investment information into constructed square meters 1980 - 1995. The data is divided into four sub-groups; dwellings, industries, other buildings and construction. The latter three are considered to be utilities and are treated as a sum. Emission factors used for calculations of particulate matter from construction activities are all found in the CEPMEIP database\textsuperscript{151}.

4.2.5.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions of TSP, PM\textsubscript{10}, PM\textsubscript{2.5} are based on expert judgement. Assessed uncertainties for are ± 100%, for each.

\textsuperscript{148} CEPMEIP, 2001. TNO. \url{http://www.mep.tno.nl/wie_we_zijn_eng/organisatie/kenniscentra/centre_expertise_emissions_assessment.html}
\textsuperscript{149} Statistics Sweden. \url{http://www.scb.se}. Housing and construction, Statistics on building permits for housing and non-residential buildings
\textsuperscript{150} The Swedish Construction Federation. \url{http://www.bygg.org}. Personal communication
\textsuperscript{151} CEPMEIP, 2001. TNO. \url{http://www.mep.tno.nl/wie_we_zijn_eng/organisatie/kenniscentra/centre_expertise_emissions_assessment.html}
More information is given in IIR Annex 1.

Time series from construction work reported in NFR code 2A5b have been reviewed in later years and 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
Particle emissions for 1999 and 2004-2015 for NFR2A5b Construction and demolition have been corrected due to updated data in national statistics, resulting in small changes, between -0.2 to 1.2 % change from the previous submission.

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 Storage, handling and transport of mineral products, NFR2A5c
Emissions from the source category Storage, handling and transport of mineral products, NFR2A5c, have not been separated from the relevant mineral chapter and are included in respective source category, 2A1, 2A2, 2A3 or 2A5.

4.2.7 Other mineral products, NFR2A6
4.2.7.1 SOURCE CATEGORY DESCRIPTION
Reported emissions include estimates for other mineral products (2A6). In the source category other mineral products, emissions from battery manufacturing and mineral wool production are reported.

Under NFR2A6 “Other mineral products” emissions from battery manufacturing and mineral wool production are reported.

4.2.7.1.1 Mineral wool production
Mineral wool has been produced at approximately 2-5 different facilities during the time period 1980-2013. Presently, glass and mineral wool production occurs at two facilities run by one company.

4.2.7.1.2 Batteries manufacturing
There is currently one battery producing facility in Sweden. This battery producer of NiCd-batteries previously used iso-propanol in their processes, which gave rise to emissions of NMVOC. The process was changed in 1998 and, since then, no NMVOC emissions occur from this source. Before 2000 another two battery producing facilities were included of which one was emitting NMVOC until 1991.
Emissions of lead, cadmium and nickel are reported for the time period from 1990 and onwards.

**Table 4-7. Summary of key source analysis, NFR2A6, according to approach 1.**

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2A6</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

4.2.7.2 METHODOLOGICAL ISSUES

4.2.7.2.1 Mineral wool production

For mineral wool production, the reported emission data on NMVOC consists of the sum of formaldehyde and phenol.

The data on particulate emissions from mineral wool production provided for the 1990’s and 2000 - 2016 are primarily based on measurements whereas for earlier years, estimates made by the companies are based on known circumstances influencing emissions. Concerning the particle emissions, only the TSP emissions were provided by industry, and the fractions of TSP as PM$_{10}$ and PM$_{2.5}$ were calculated from emission factors for production of glass fibres provided in the CEPMEIP study.$^{152}$

4.2.7.2.2 Batteries manufacturing

The time series from 1990 - 2016 is based on emission data representing three individual facilities. From 2000 there is only one active facility. Between 1988 and 1991 reported NMVOC represents emissions from two facilities. From 1992 to 1998 only NMVOC from one facility is included. This battery manufacturer of NiCd-batteries used isopropanol in their processes, which gave rise to emissions of NMVOC. The process was changed in 1998 and, since then, no NMVOC emissions occur from this source. The heavy metal emissions from the battery manufacture nowadays originate from one facility producing nickel-cadmium batteries. For some years information on emissions is not available, and data has been interpolated.

4.2.7.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions of NMVOC, NH$_3$, TSP, PM$_{10}$, PM$_{2.5}$ are based on expert judgement. Assessed uncertainties for are ± 400%, ± 400%, ± 100%, ± 100% and ± 100%, respectively.

Uncertainties for emissions for metals are based on data from the Guidebook 2008. Uncertainties for Pb, Cd, Hg, As, Cr, Cu, Ni and Zn are ± 50%, ± 100%, ± 1000%, 100%, ± 100%, ± 50%, ± 50% and ± 50%, respectively.

More information is given in IIR Annex 1.

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Time series from construction work reported in NFR code 2A6 have been reviewed in later years and are consistent.

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 were performed during submission 2018.

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, NFR2B

4.3.1 Ammonia production, NFR2B1

4.3.1.1 SOURCE CATEGORY DESCRIPTION
There is an annual production of about 5 Gg of ammonia in Sweden, according to UN statistics\(^\text{153}\). This 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 NTA\(^\text{154}\). Emissions from this industry are included in NFR code 2B5. Ammonia production, 2B1, is thus reported as NO in the NFR-tables.

4.3.1.2 METHODOLOGICAL ISSUES
Emission data on NO\(_X\) and NH\(_3\) originating from the nitric acid production has been obtained directly from the facilities and from official statistics. Emissions for all years, except 1991-1993, are as reported from the facilities. The reduction of the reported NO\(_X\) emissions in 2001 and 2002, compared to earlier years, is a result of one facility being shut down in late 2000 and a second one during 2001. 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 year 2007 catalytic abatement was installed at one of the production units at the active facility and as a result the emissions of NO\(_X\) and NH\(_3\) were reduced compared to previous years. According to the company the increased NH\(_3\) emissions in 2010-2014 is a result of prioritizing low NO\(_X\) emissions. NH\(_3\) is used as a reducing agent in the de-NO\(_X\) catalyst and hence lower NO\(_X\) implies more injected NH\(_3\). NH\(_3\) that do not react in the catalyst is emitted to the air. From 2007 emissions are continuously measured in one of the two production lines, from 2011 emissions are continuously measured in both production lines.

\(^{153}\) UN. Commodity Production Statistical Database. Department of Economic and Social Affairs, Statistics Division, as referred in FCCC Synthesis and Assessment report 2002 Part I.

\(^{154}\) Kindbom, 2004.
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 and measurements. However, this information is confidential.

4.3.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Time series from industrial processes reported in NFR codes 2A-2H have been reviewed in later years and are consistent.

4.3.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
Experts at the Swedish EPA conduct a review of the inventory, estimates, methodology and emissions factors used. The experts also identify areas of improvement, which constitute part of the basis for improvements in coming submissions.

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.

All Tier 1 general inventory level QC procedures and some specific Tier 2 QC procedures, listed in Good Practice Guidance section 8, have been performed and are documented in check-lists.

The time series for all revised data have been studied carefully in search for outliers and to make sure that levels are reasonable. Data have, when possible, been compared with information in environmental reports and/or other independent sources. Remarks in reports from the UNFCCC and CLRTAP/NEC reviews have been carefully read and taken into account.

According to the Good Practice Guidance, the method of calculating emissions at facilities should be documented. This is currently not done in most cases and will be improved in the future.

4.3.1.5 SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculations have been made.

4.3.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are currently planned.

4.3.2 Nitric acid production, NFR2B2
4.3.2.1 SOURCE CATEGORY DESCRIPTION
Production of nitric acid has taken place at three facilities in Sweden. One of these was shut down late 2000, and a second one was shut down during 2001. Therefore, there is currently only one facility producing nitric acid in Sweden. Data on emissions have been obtained directly from the facilities and from official statistics.
4.3.2.2 METHODOLOGICAL ISSUES

Emission data on NO$_X$ and NH$_3$ originating from the nitric acid production has been obtained directly from the facilities and from official statistics. Emissions for all years, except 1991-1993, are as reported from the facilities. The reduction of the reported NO$_X$ emissions in 2001 and 2002, compared to earlier years, is a result of one facility being shut down in late 2000 and a second one during 2001. 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 year 2007 catalytic abatement was installed at one of the production units at the active facility and as a result the emissions of NO$_X$ and NH$_3$ were reduced compared to previous years. According to the company the increased NH$_3$ emissions in 2010-2014 is a result of prioritizing low NO$_X$ emissions. NH$_3$ is used as a reducing agent in the de-NO$_X$ catalyst and hence lower NO$_X$ implies more injected NH$_3$. NH$_3$ that do not react in the catalyst is emitted to the air. From 2007 emissions are continuously measured in one of the two production lines, from 2011 emissions are continuously measured in both production lines.

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 and measurements. However, this information is confidential.

4.3.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Time series from industrial processes reported in NFR codes 2A-2H have been reviewed in later years and are consistent. Uncertainties for NFR2B2 are displayed in Table 4-9.

Uncertainties for emissions are based on information from the company. Assessed uncertainties for NO$_X$ and for NH$_3$ are ± 5%. More information is given in IIR Annex 1.

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 recalculations have been performed in submission 2018.
4.3.2.6  SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are currently planned.

4.3.3  Adipic acid production, NFR2B3
No production of adipic acid occurs in Sweden, and thus NO is reported for NFR2B3.

4.3.4  Caprolactam, glyoxal and glyoxylic acid production (NFR2B4)
4.3.4.1  SOURCE CATEGORY DESCRIPTION
No production of caprolactam, glyoxal or glyoxylic acid occurs in Sweden, and thus NO is reported for NFR2B3.

4.3.5  Carbide production, NFR2B5
4.3.5.1  SOURCE CATEGORY DESCRIPTION
Carbide production occurs at only one facility in Sweden. Emissions of TSP, PM$_{10}$, PM$_{2.5}$ and BC are reported from carbide production and estimates are based on information from the company. The distribution of particulates between TSP, PM$_{10}$ and PM$_{2.5}$ has been determined by expert judgement.

Silicium carbide production does not occur in Sweden.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2B5</td>
<td>PM$_{2.5}$</td>
<td></td>
</tr>
</tbody>
</table>

4.3.5.2  METHODOLOGICAL ISSUES
The time series of particle emissions from carbide production are considered complete and consistent in methodology. TSP, PM$_{10}$, PM$_{2.5}$ and BC emissions from carbide production are included and the dominating part of reported emissions arises from flaring of carbide oven gas. The partitioning of particles between TSP, PM$_{10}$ and PM$_{2.5}$ has been done by expert judgement after discussions with the carbide producing company and the emitted mass of PM$_{10}$ and PM$_{2.5}$ are estimated to constitute 90% and 80% of the total TSP emission, respectively. Particle emissions from quicklime production in the carbide industry are included in NFR2A2.

4.3.5.3  UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Time series from industrial processes reported in the NFR code 2B5 have been reviewed in later years and are consistent. Uncertainties for NFR2B5 are displayed in Annex I.

4.3.5.4  SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.
4.3.5.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been performed in submission 2018.

4.3.5.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are currently planned.

4.3.6 Titanium dioxide production, NFR2B6
No production of titanium dioxide occurs in Sweden, and thus NO is reported for NFR2B6.

4.3.7 Soda ash production, NFR2B7
In 2004\textsuperscript{155} a study was carried out to collect data on soda ash use and calculate CO\textsubscript{2} emissions. From this study it became clear that no production of soda ash occurs in Sweden, and is hence reported as NO in NFR2B7.

4.3.8 Other chemical industry, NFR2B10a
4.3.8.1 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 is not covered elsewhere. Approximately 70 larger industrial facilities are included in the emission estimates. Emissions of NO\textsubscript{X}, CO, NMVOC, SO\textsubscript{2}, NH\textsubscript{3} and TSP are reported. From submission 2016 onwards, emissions of PM\textsubscript{2.5}, PM\textsubscript{10} and BC from three inorganic chemical processes are included: production of carbon black, production of phosphate fertilizers and NPK, and production of PVC. It is possible that some emissions of NMVOC reported in NFR2B10 should be reported in NFR2D3g (e.g. pharmaceutical industries), but since it has been difficult to make the distinction clear between process emissions and solvent use, all NMVOC emissions from these facilities are included in NFR2B10.

The mercury emissions reported originate from the chloralkali and the sulphuric acid industries. The dioxin emissions reported in 2001 originate from three facilities, in 2002 from four, and for 2003 from six facilities. Due to lack of information about emissions in earlier years, dioxin emissions are reported NE (Not Estimated) for 1980 – 2000.

From some chemical processes, emissions of Se, As and PAH may occur, according to EMEP/EEA Guidebook 2013 (e.g. arsenic from production of PVC and phosphate fertilizers). Data on these emissions are not available, and the time series from 1990 and onwards are thus reported NE.

Time series for HCB emissions has been included to the inventory as a result of a development project\textsuperscript{156}. These emissions occur at one facility within inorganic chemical production, and have significantly (by more than 90\%) decreased between 1990 and 2016.

Table 4-11. Summary of key source analysis, NFR2B10a, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2B10</td>
<td>Hg, SO(_2), NMVOC</td>
<td>Hg, SO(_2), NH(_3), NMVOC</td>
</tr>
</tbody>
</table>

4.3.8.2 METHODOLOGICAL ISSUES

The primary information on emissions of NO\(_x\), CO, NMVOC, SO\(_2\), NH\(_3\) and TSP are as reported by the companies in their environmental reports. A total of approximately 70 facilities are included, but not all of them report on all emissions. The time series have been reviewed and are considered to be consistent.

From submission 2016 onwards, emissions of PM\(_{2.5}\), PM\(_{10}\) and BC from three inorganic chemical processes are included. Emission factors are based on particulate fractions as in EMEP/EEA Guidebook 2013, and applied to TSP emissions reported by facilities. Activity data on production of PVC and carbon black has been acquired from the facilities’ environmental reports. Activity data on production of phosphate fertilizers was obtained from FAOSTAT\textsuperscript{157} for 1990-1995 and from Statistics Sweden database for 1996 onwards.

Mercury emissions reported in 2B10a originate from processes in the chloralkali industry and from sulphuric acid production. Reported emissions of mercury were derived from information in the SMP database, from the industries’ environmental reports or unpublished earlier estimates\textsuperscript{158}.

Hardly any information on dioxin emissions from the chemical industry has been available, and it has not been possible to compile any time series of dioxin emissions from these sources. There is only information on dioxin emissions available from a few facilities from 2001.

The SO\(_2\) emissions reported in 2B10a decreased dramatically in 2004 in comparison to earlier years. This is due to that in December 2003 one facility for production of viscose staple fibre was shut down. The yearly SO\(_2\) emissions from this facility represented between 8 and 20 \% of the totally reported SO\(_2\) emission in NFR2 – Industrial Processes, 1990 - 2003. In 2007 the CO-emissions were very

\textsuperscript{156} Hagström, P. 2017, SMED memorandum on HCB emissions from Swedish industrial processes

\textsuperscript{157} FAOSTAT \texttt{http://faostat3.fao.org}

\textsuperscript{158} Levander, 1989.
low from one facility producing PVC. NH$_3$ emissions decreased since 2007 due to that one facility are working on replacing NH$_3$ in the production process.

In 2010, emissions in this sub-category were reviewed as part of a quality control project carried out by SMED on behalf of the Swedish EPA, aiming at increasing the quality and reducing the uncertainties of the emissions of the most important substances from chemicals industries in Sweden\textsuperscript{159}. Emissions reported in the environmental reports were compared to plant-specific data. Significant discrepancies were investigated, and recommendations were provided on feasible improvements for submission 2011 as well as recommendations on further investigations\textsuperscript{160}.

Overall, the QC-project showed that total reported emissions from the chemical industries in the Swedish inventory are in coherence with the emission data reported by the plants.

4.3.8.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for NFR2B10a are displayed in Annex I.

Time series for various chemical industries reported in NFR code 2B10a have been reviewed in later years and are considered to be consistent.

4.3.8.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Emissions reported in the plant-specific environmental reports are carefully studied annually to retrieve the most appropriate data.

4.3.8.5 SOURCE-SPECIFIC RECALCULATIONS

Inorganic chemical production (relative changes in % below are given for this particular sub-sector):

Minor changes have been implemented for the year 2015 with regard to NH$_3$, and particle fractions from one plant, due to one year delay in in-data and other corrections – they resulted in a decrease by 0.4 % and an increase by 0.03-0.2%, respectively, compared to submission 2017.

Hg emissions from one facility (1 kg) are subtracted from the emissions 2015, since the facility was closed in 2014 and emissions no longer occur. This resulted in the emission decrease by 6%

\textsuperscript{159} Swedish EPA. 2010.

\textsuperscript{160} Most recommendations on further investigations refer to the energy sector
NMVOC emissions for 2007-2015 from one facility have been recalculated due to updates in the method used by the facility. NMVOC emissions reported in the sub-sector decrease by 0.5-15% (up to 26 tonnes) compared to Submission 2017.

In submission 2018, a time series for HCB was added in the inventory. The trend is descending, with annual emissions varying from 15 kg in the nineties to around 1-3 kg in the recent years.

Organic chemical production (relative changes in % below are given for this particular sub-sector):

Minor corrections have been implemented for the year 2015 with regard to CO and NH3 emissions from one plant – they resulted in a decrease by 0.4 tonnes (1%) and 2 tonnes (17%), respectively, compared to submission 2017.

For another facility, NOx emissions from boilers have been recalculated taking into account the share of internal gas combustion. The recalculation resulted in the significant emission decrease – by up to 71%, or 15 tonnes – for the whole time series.

NMVOC emissions from one plant have been corrected by subtracting methane, which resulted in the emission decrease by 0.1-0.8% (up to 6 t) for 2004-2015.

4.3.8.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are currently planned.

4.3.9 Storage, handling and transport of chemical products, NFR2B10b
Emissions from the source category Storage, handling and transport of chemical products, NFR2B10b, have not been separated from the relevant chemical production chapter and are included in respective source category, 2B1, 2B2, 2B5 or 2B10a.

4.4 Metal production, NFR2C
4.4.1 Iron and steel production NFR2C1
4.4.1.1 SOURCE CATEGORY DESCRIPTION
Processes that are included in this category are primary and secondary iron and steel production, direct reduced iron production, iron ore mining, dressing, sintering and iron ore pellets production. The summary of the latest key source assessment is presented in Table 4-12.

In Sweden, there are three primary iron and steel facilities equipped with blast furnaces, producing iron and steel products from virgin materials, and about ten secondary steel plants equipped with electric arc furnaces, producing iron and steel products from scrap and direct reduced iron. One of the facilities is using a shaft
furnace process to produce stainless steel from recovered flue gas dust and other waste products. 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 refinement processes. From submission 2009 and onwards, emissions from two major iron ore mines and three facilities producing pellets in Sweden are reported in 2C1. Emissions from one sinter producing facility are reported in 2C1 until 1995, when the production closed down.

Table 4-12. Summary of key source analysis, NFR2C1, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2C1</td>
<td>Se, Cr, Ni, DIOX, Zn, SO(<em>2), PM(</em>{10}), TSP, Cd, PM(_{10}), Hg, Pb, Cu, NO(_x), PAH 1-4, As</td>
<td>Se, Cr, Ni, DIOX, Zn, SO(<em>2), PM(</em>{10}), TSP, Cd, PM(_{10}), Hg, Pb, Cu, NO(_x), As</td>
</tr>
</tbody>
</table>

4.4.1.1 Primary iron and steel production

There are two plants in Sweden that produce pig iron and steel as part of their integrated coke ovens, blast furnaces and steel converters. The basis of the production is iron ore pellets.

4.4.1.2 Secondary iron and steel production

There are about ten secondary steel plants equipped with electric arc furnaces, producing iron and steel products from scrap and direct reduced iron. One of the facilities is using a shaft furnace process to produce stainless steel from recovered flue gas dust and other waste products.

4.4.1.3 Direct reduced iron

There is one plant in Sweden which produces iron sponge and iron powder using direct reduction of iron ore pellets.

4.4.1.4 Iron ore mining, dressing, sintering and iron ore pellets production

In Sweden, there are three primary iron and steel facilities equipped with blast furnaces, producing iron and steel products from virgin materials. Emissions from three major iron ore mines and three facilities producing pellets in Sweden are reported in 2C1.

Emissions from a sinter producing facility are also included until 1995, when the production closed down.

4.4.1.2 METHODOLOGICAL ISSUES

Process emissions arising from reducing agents in the primary steel works and secondary iron and steel works are reported in NFR2C1. As the plants also generate emissions from fuel combustion (NFR1A1c and NFR1A2a) and fugitive emissions
(NFR1B1c) the text in this section is closely connected to the text in the energy section.

In the Swedish inventory, emissions from primary iron and steel production and secondary steel production are estimated separately but reported together under 2C1 iron and steel production. In 2C1 also emissions from three major iron ore mines and three facilities producing pellets in Sweden are included.

4.4.1.2.1 Primary iron and steel production

Two plants reported in this sector are primary iron and steel producing plants as part of 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 to act as reducing agents, and the associated emissions are reported as industrial processes from iron and steel production in NFR2C1, according to the 2006 IPCC Guidelines.

Figure 4-1 gives an overview of the input and output materials, the carbon flows between the different processes (plant stations), and the CO₂-emitting sources. Note that for non-CO₂ emissions, the different emission sources may vary considerably. The flow chart is however giving a general introduction to the two integrated iron and steel production 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 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 metallurgical 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.
Figure 4-1. Carbon flow chart of integrated primary iron and steel plants in Sweden. CRF = NFR.
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 NFR1A1a 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 in line with the 2006 IPCC Guidelines\textsuperscript{161} where allocation of emissions from delivered gases is described. During the whole process from raw material to final product, emissions are released.

From 2003 onwards, the plant specific annual environmental reports consist of plant station data on consumed amounts of energy gases (coke oven gas, blast furnace gas and LD-gas) and other fuels, emissions of NO\textsubscript{X}, SO\textsubscript{2}, several heavy metals, TSP and dioxin (one plant only), but lack information on emissions of NMVOC, CO and some heavy metals. In previous submissions, time series for several pollutants (NO\textsubscript{X}, SO\textsubscript{2}, NMVOC and CO) were based on information from various sources (e.g. Statistics Sweden and environmental reports). As of submission 2010, the inventory reporting of all emissions is based on information from the environmental reports and some additional information from direct contact with the plants. In order to achieve consistent time series and to estimate emissions of missing pollutants, different IPCC splicing techniques were applied.

Emissions of NO\textsubscript{X}, SO\textsubscript{2} and TSP are derived from the environmental reports and direct contact with the plants for the entire time series. The allocation of both plants total emissions of NO\textsubscript{X}, SO\textsubscript{2} and TSP on plant stations and consequently NFR category is presented in Table 4-13 for the year 2016.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Plant station</th>
<th>NO\textsubscript{X} emissions (kt)</th>
<th>SO\textsubscript{2} emissions (kt)</th>
<th>TSP emissions (kt)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A1c</td>
<td>Coke Oven</td>
<td>0.35</td>
<td>0.13</td>
<td>71</td>
</tr>
<tr>
<td>1A2a</td>
<td>Combustion in Rolling Mills + Power and Heat Production</td>
<td>0.37</td>
<td>0.32</td>
<td>15</td>
</tr>
<tr>
<td>1B1b</td>
<td>SO\textsubscript{2} from quenching and extinction at coke ovens</td>
<td>0.03</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1B1c</td>
<td>Flare in Coke Oven (COG)</td>
<td>0.01</td>
<td>0.01</td>
<td>0.22</td>
</tr>
<tr>
<td>2B5</td>
<td>Sulphuric acid production</td>
<td></td>
<td>0.01</td>
<td></td>
</tr>
<tr>
<td>2C1</td>
<td>Blast Furnace + Steelworks (including Flaring of BFG and LD-gas)</td>
<td>0.17</td>
<td>0.57</td>
<td>221</td>
</tr>
<tr>
<td></td>
<td><strong>Total</strong></td>
<td><strong>0.90</strong></td>
<td><strong>1.08</strong></td>
<td><strong>307</strong></td>
</tr>
</tbody>
</table>

NMVOC and CO emissions are estimated based on consumed amounts (including flared amounts) of energy gases multiplied by country specific emission factors (see Annex 2). Emissions of NMVOC and CO from coke oven gas, blast furnace

\textsuperscript{161} See 2006 IPCC Guidelines: Volume 3: Industrial Processes and Product Use, Box 1.1 (page 1.8)
gas and LD-gas in the blast furnace and steel converter are allocated to NFR2C1. 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 CO₂ emissions as the surrogate parameter.

Inventory emissions of heavy metals, TSP and dioxin are mostly obtained from the environmental reports, except for selenium, which is included for the first time in submission 2016, calculated using default emission factor from EMEP/EEA Guidebook 2013. In some cases, especially for the early 1990’s and for one of the plants, information on heavy metal emissions are lacking and thus estimated by extrapolation using IEF and TSP as a surrogate parameter. Emissions of heavy metals and dioxin are all reported under NFR2C1. The PM size fractioning has been made according to reported emissions of PM₁₀ and PM₂.₅ from one of the plants. From submission 2015, BC emissions are also reported from year 2000, calculated as a fraction of PM₂.₅, according to the EMEP/EEA Guidebook 2013.

Reported activity data are amounts of pig iron produced (Mton).

4.4.1.2.2 Secondary iron and steel production
For reported emissions from secondary iron and steel production, the companies’ environmental reports are the main source of information. NOₓ, NMVOC and SO₂ emissions emitted from electric arc furnaces are reported in 2C1. NOₓ emissions may also arise from pickling and acid regeneration and NMVOC emissions may arise from rolling mills. These sources are also included in the estimates.

The estimated TSP emissions are based on information from the trade association¹⁶² for almost all years 1980-1999 but for 2000 and onwards the information was derived from the companies’ environmental reports. TSP data for missing years have been interpolated. The PM size fractioning has been made according to expert judgement, and are based on knowledge about changes in production methods and abatement technology. From submission 2015 onwards, BC emissions are reported from secondary iron and steel production from 2000 onwards. Emissions are calculated as a fraction of PM₂.₅ according to EMEP/EEA Guidebook 2013.

The estimated metal emissions from secondary iron and steel processes are based on produced amount of steel, published by the trade association¹⁶², and emission factors, for the years 1990 - 2000. The emission factors used are based on compiled information from older trade specific reports made by the Swedish EPA for some years during the 1990’s. Emission factors have been calculated for Cd, Cr; Cu, Hg, Ni, Pb and Zn. For years where the Swedish EPA did not provide trade specific reports, or when the trade was not fully covered in the reports, data has been interpolated. Data on As emissions from iron and steel production is somewhat uncertain since reported data are scarce. From 2001 and onwards the emissions are mainly

derived from the companies’ environmental reports. For years when information is missing in the environmental reports, emissions are estimated using IEF for earlier years and production volumes or amounts of particles emitted.

Dioxin emissions have been compiled for the whole time series. According to the US-EPA\textsuperscript{163}, dioxin emissions from steel production are strongly dependent on a number of parameters, likely to vary between steel plants. Whether steel is produced from primary metals or from scrap metal is one very important factor, with the latter giving much higher dioxin emissions. Since the emission factors vary widely depending on several process factors, no straightforward calculations using an emission factor were made when compiling a time series of national dioxin emissions from the iron and steel industry. Instead, the estimates for the time period 1980 - 2000 are based on a combination of information concerning production data for scrap-based steel, results from dioxin measurements, earlier estimates and expert judgement in co-operation with the trade association\textsuperscript{164}. From 2001 the information concerning the dioxin emissions were derived from the companies’ environmental reports.

In submission 2016 emissions of PAH1-4 and PCB from secondary iron and steel processes were reported for the first time, based on national figures on produced amount of steel and emission factors from EMEP/EEA Guidebook 2013.

\subsection*{4.4.1.2.3 Direct reduced iron}

For emissions estimates for the producer of iron ore based iron powder, almost all reported emissions are obtained from the plant’s environmental reports and are verified by collecting and comparing the carbon contents in the amounts of coke, anthracite and output material. 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 NFR2C1. Reported activity data is produced amount of direct-reduced iron (iron sponge). For estimation of PM\textsubscript{10} and PM\textsubscript{2.5} emissions, the same fractions of TSP as in the CEPMEIP project\textsuperscript{165} are used.

Starting from submission 2016, emissions of Se, HCB and PCB are reported, calculated with default emission factors from the EMEP/EEA Guidebook 2013.

\subsection*{4.4.1.2.4 Iron ore mining, dressing, sintering and iron ore pellets production}

There are currently two major iron ore mines and three facilities producing pellets in Sweden. Until 1995, emissions from a now closed sinter producing facility are

\begin{itemize}
\item\textsuperscript{163} U.S. Environmental Protection Agency, 1997.
\item\textsuperscript{164} The Swedish Steel Producers Association, \url{http://www.jernkontoret.se}.
\item\textsuperscript{165} CEPMEIP, 2001. TNO. \url{http://www.mep.tno.nl/wie_we_zijn_eng/organisatie/kenniscentra/centre_expertise_emissions_assessment.html}
\end{itemize}
also included. Emissions considered are SO₂ from the sulphur content in the ore and NOₓ emitted as a result of the use of explosives. The use of mining explosives also causes emissions of carbon monoxide, CO. No data concerning the CO emissions are available and the time series from 1990 is thus reported NE. Metals are reported for the time period from 1990, whereas emissions of dioxins and particles are reported for the period 1980 onwards.

The figures are based on data reported by the companies in their environmental reports. For years with missing data figures have been interpolated or estimated, using expert judgement in cooperation with industry. For distributing the emission of particulates between TSP, PM₁₀ and PM₂.₅ the same ratio has been used as the one used in the CEPMEIP study. Content of heavy metals in particulate matter has been calculated using an analysis provided by the leading company. The analysed samples were from the pellets production, but in the emission estimates the factors have been used on the sum of produced sinter and pellets. BC emissions are reported from submission 2015 and are calculated as a fraction of PM₂.₅ according to EMEP/EEA Guidebook 2013.

In submission 2016 emissions of PAH1-4, PCB, HCB from sintering and iron ore pellets production are reported for the first time. For iron ore pellets production also emissions of selenium are reported for the first time. The estimates are based on produced amount and emission factors from EMEP/EEA Guidebook 2013.

The reported emissions of dioxins are based on an emission factor developed using measurements in 2001 and known production data for the same year. Since the production methodology and other circumstances have not changed significantly during the time period, this developed emission factor has been used to calculate the time series of dioxin emissions for the period 1980 - 2001. For 2002 onwards, data on dioxin emissions provided by the companies are reported.

4.4.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Time series from industrial processes reported in NFR codes 2C1 has been reviewed in later years and are consistent. The emission of dioxin was 2.3 times higher during submission 2017 in comparison to the year before. However, the difference only amounted 0.4 g and was an effect of that two smelters reported higher emissions than earlier.

Uncertainties for NFR2C1 are displayed in Annex 1.

Primary iron and steel plants (including direct reduced iron production): Uncertainties are mostly expert estimates, except for Se, PCB and HCB, for which the

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166 Wieland, 2004
emission factor uncertainties are calculated based on the intervals in the EMEP/EEA Guidebook 2013.

4.4.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

4.4.1.5 SOURCE-SPECIFIC RECALCULATIONS

4.4.1.5.1 Primary iron and steel plants:
Emissions of TSP, PM$_{10}$, PM$_{2.5}$ and BC have been corrected for the entire time series in submission 2018, resulting in $\sim$3% – +2%, 15–64%, 15–43% and 11–43% change, respectively, for this code.
Minor corrections have been made in the data from environmental reports for 1990-2002, resulting in 0.00002-0.00003% change.

4.4.1.5.2 Secondary iron production:
No recalculations have been made in submission 2017.

4.4.1.5.3 Iron ore mining, dressing, sintering and iron ore pellets production:
New stack gas cleaning technique was installed in the Svappavara pellet plant during December 2015. After installation the cleaning efficiency in respect to SO$_2$ was increased from 10% to 97% as proved by continuous measurements of the SO$_2$ emission during 2016. The new SO$_2$ cleaning factor was implemented in submission 2018.

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 NFR2C2

4.4.2.1 SOURCE CATEGORY DESCRIPTION
Ferroalloy production is reported for only one facility in Sweden. There is also ferroalloy production at another plant, but since the main production at this facility is of iron and steel, these emissions are reported in NFR2C1, Iron and steel production.
Table 4-14. Summary of key source analysis, NFR2C2, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2C2</td>
<td>Cr</td>
<td>Cr</td>
</tr>
</tbody>
</table>

4.4.2.2 METHODOLOGICAL ISSUES

Emission data for SO$_2$ and NO$_X$ has been obtained directly from the company for all years. Production of ferrosilicon leads to larger emissions of SO$_2$ compared to production of ferrochromium. From 2005 the production of ferrosilicon has been much reduced and during 2008 – 2011 and 2014 no ferrosilicon was produced. This led to a distinct decrease in SO$_2$ emissions during these years. In 2012 the production of ferrosilicon was relatively large compared to adjacent years, which resulted in high emissions of SO$_2$. Also emissions of NO$_X$ were higher in 2012 compared to years with no ferrosilicon production. In 2013 only small amounts of ferrosilicon was produced, resulting in lower emissions of SO$_2$ and NO$_X$ compared to 2012.

TSP emissions for 1980-1999 have been calculated based on activity data provided by the company and emission factors derived from reported emissions of TSP in the company’s environmental reports in later years. The calculated average emission factor has been used for all years during the 1990’s and was doubled for the period 1980-1989, as suggested by the company experts. From 2000, data on TSP emissions from the company’s environmental report were used. An expert has performed measurements at the facility since the beginning of the 1980’s, and has estimated the size distribution of particulates between PM$_{10}$ and PM$_{2.5}$\cite{peterson2002}. The same particle size distribution was assumed for the time series 1980 - 2001. From 2002 and onwards the reported PM$_{10}$ emissions are as presented by the company in their environmental reports. For the years after 2001 the reported PM$_{2.5}$ emissions are estimated as for earlier years due to lack of specific information on PM$_{2.5}$ emissions from the company.

Metals emitted to air from ferroalloy production are primarily Cr, Pb, Ni and Zn. Chromium emission have been reported in the environmental reports to the emission database from 1992. The chromium (Cr) emissions in the database and the activity data obtained from the company have been used to derive emission factors. The average emission factor for 1992-1994 was used for 1990-1991. Zinc and lead emissions have only been sporadically reported to the database during the 1990’s. In order to estimate emissions of Zn and Pb, information from older Swedish EPA reports were combined with the reported data on emissions to calculate emission factors for the 1990’s. Emissions of Ni from ferroalloys production has been derived from the company’s environmental reports or by information from the producer for the years 2003 – 2014. For earlier years no data is available and Ni emissions are hence reported NE (Not Estimated) for the time period 1980 – 2002.

\cite{peterson2002} Peterson, 2002.
4.4.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Time series from industrial processes reported in NFR codes 2C2 have been reviewed in later years and are consistent.

Uncertainties for NFR2C2 are displayed in Annex 1.

4.4.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

4.4.2.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been performed in submission 2018

4.4.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are currently planned.

4.4.3 Aluminium production, NFR2C3

4.4.3.1 SOURCE CATEGORY DESCRIPTION
The process included in this category is primary aluminium production (2C3).

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. 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 big 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.

PAHs emissions occur in Söderberg plants due to the self-baking anode. Emissions of PAHs during the electrolysis process are small for Prebake plants but to submission 2018 these emissions are calculated from 2009 and onwards using emission factors emissions presented in EMEP/EEA Guidebook 2016.

Table 4-15. Summary of key source analysis, NFR2C3, according to approach 1.
4.4.3.2 METHODOLOGICAL ISSUES

Primary aluminium production takes place in one facility, where historically both the Prebake and the Söderberg technologies have been used. All pot-lines operating the Söderberg technology were shut-down by December 2008.

The time series of emissions compiled for primary aluminium production include emissions of NO\(_X\), CO, NMVOC and SO\(_2\), particles and PAH. Reported production statistics and emissions data are based on information in the environmental reports or received directly from the company.

In submission 2016 emissions of dioxin from aluminium production are reported for the first time. The estimates are based on produced amount of aluminium and emission factors from EMEP/EEA Guidebook 2013.

Emissions of NO\(_X\) have been calculated from production statistics using emission factors defined by Swedish EPA\(^{169}\). NMVOC emissions have been calculated from reported emissions of tar, assuming that 70% of the tar is emitted as NMVOC. Closing down the Söderberg ovens also ended the need for anode production in late 2008. The shutdown of the anode production ended the tar emissions which meant that also the NMVOC emissions fell sharply. From 2009 and onwards, emissions of NMVOC are reported NE since no emission factor is specified in the EMEP/EEA Guidebook. CO emissions were reported for the first time in submission 2008 and are for 2002-2015 as reported in the company's environmental reports. For the period 1990-2001 the CO emissions are calculated using production statistics and emission factor provided by the company as also the SO\(_2\) emissions, 1990-2005. For 2006-2015 SO\(_2\) is as reported by the company in their environmental reports.

The elevated SO\(_2\) emission in 2012 is primarily due to high sulphur content in delivered anodes. The desulfurization of flue gases in the flue gas treatment facilities was not sufficiently efficient. In 2014 the SO\(_2\) emissions were lower than previous year due to improved abatement technology. The improved abatement technology is also shown in low SO\(_2\) emissions in 2015 and 2016. Also the CO emissions were higher for 2012 compared to previous years. The reason for this is, according to the company, that a new calculation method has been used from 2012 and onwards.

Information concerning production statistics and emissions of TSP and benzo(a)pyrene (BaP) were provided by industry, and only a few missing years have to be interpolated. The reported emissions also include particles from the foundry located at the site of the primary production plant. The particle size fractions of PM\(_{10}\)

\(^{169}\) Ahmadzai, H. Swedish EPA. Personal communication. 2000.
and PM$_{2.5}$ have been assumed for the whole time period, as given in the CEPMEIP project\textsuperscript{170} for primary aluminium production. For particles from the foundry the same particle size fractions of PM$_{10}$ and PM$_{2.5}$ have been used. The assumption is thus that PM$_{10}$ constitutes 95 % and PM$_{2.5}$ 43 % of the reported TSP emissions.

Emissions of benzo(a)pyrene and “PAH” have been reported from the facility as far back as 1984. It is not known which compounds are included in the term “PAH”. In 1984 and 1986, benzo(a)pyrene emissions occurred from plant 1 and 2. From 1987 until 2008, emissions occurred only from plant 2, which represents the production of Söderberg anodes and anode baking in the so-called Söderberg ovens. Emissions of PAHs during the electrolysis process are small for Prebake plants but to submission 2018 these emissions are calculated from 2009 and onwards using emission factors emissions presented in EMEP/EEA Guidebook 2016. For 1990 – 2008 emissions of benzo(a)pyrene are as reported by the facility and emissions of benzo(b)fluoranthene, benzo(k)fluoranthene and indeno(1,2,3-de)pyrene are calculated based on information in EMEP/EEA Guidebook 2016.

According to UNEP\textsuperscript{171} primary production of aluminium has no significant emissions of dioxins to air. This was confirmed by measurements made at the facility in the late 1970’s and early 1980’s. The measurements in the early 1980’s showed no detectable amounts.

4.4.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions calculated with emission factors from EMEP/EEA Guidebook are based on data in the Guidebook (2013 for PCDD/F and 2016 for PAHs). Uncertainty for SO$_2$, NO$_X$, NMVOC and particles are ± 30%, ± 50%, ± 75% and ± 40%, respectively.

More information is given in IIR Annex 1.

Time series from aluminium production reported in NFR code 2C3 have been reviewed in later years and are 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

In earlier submissions emissions of PAH 1-4 from 2009 have been reported NA. In submission 2018 emission factors from EMEP/EEA have been used to calculate these emissions. For the years 1990-2008 emissions of benzo(b)fluoranthene, benzo(k)fluoranthene and indeno(1,2,3-cd)pyrene have been recalculated. In earlier

\textsuperscript{170} CEPMEIP, 2001. TNO. \url{http://www.mep.tno.nl/wie_wij_zaai_kennis/centre_expertise_emissions_assessment.html}

submissions emissions of benzo(b)fluoranthene and indeno(1,2,3-cd)pyrene were based on PAH profile from UNEP (benzo(k)fluoranthene were reported NE). In submission 2018 emissions of benzo(b)fluoranthene, benzo(k)fluoranthene and indeno(1,2,3-cd)pyrene are based on the relationship between the emission factors for these gases and the EF for benzo(a)pyrene (Söderberg technology) in the EMEP/EEA Guidebook 2016.

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 Magnesium production, NFR2C4
No production of magnesium occurs in Sweden, and thus NO is reported for NFR2C4.

4.4.5 Lead, Zinc, Copper and Nickel production, NFR2C5, 2C6, 2C7a and 2C7b
Production of lead, zinc, copper and nickel does occur in Sweden. However, since Swedish non-ferrous metal smelters produce several metals in the same process, emissions cannot be separated and are all included in NFR2C7c Other metal production. Thus IE is reported in NFR2C5, 2C6, 2C7a and 2C7b.

4.4.6 Other metal production, NFR2C7c
4.4.6.1 SOURCE CATEGORY DESCRIPTION
This sub-category includes emission estimates from one large smelter producing different non-ferrous metals such as copper, lead, zinc etc., one metal recycling company mainly producing lead and seven smaller smelters of various kinds. Emissions of particles have been obtained from the large smelter from 1980, for one facility from 1985 and for most of the smaller smelters from 1990. Time series of metal emissions are reported from 1990, and includes also the smaller facilities. In the dioxin time series reported emissions from the large smelter, from the metal recycling company and from two smaller smelters are included.
Table 4-16. Summary of key source analysis, NFR2C7c, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2C7c</td>
<td>Cu, Cd, Pb, As, SO₂, Cr, Zn, Hg, DIOX, Ni</td>
<td>Cu, Cd, Pb, As, SO₂, Cr, Zn, Hg, DIOX, Ni, PM₁₀, TSP, PM₂.₅</td>
</tr>
</tbody>
</table>

4.4.6.2 METHODOLOGICAL ISSUES

The reported emissions of SO₂ mainly originate from the sulphur content in the raw materials used in the large non-ferrous smelter, but also represent emissions from the metal recycling company and from one of the smaller smelters. Reported NOₓ in 2C7c represents the same facilities. The SO₂ and NOₓ time series are considered complete and consistent.

At the large smelter, a variety of processes occur, including both primary and secondary processes, and a number of products are produced. This facility has a long history of submitting environmental reports to the authorities, why emission estimates for all substances were readily available, except for the size fractions of emitted particles. Emission factors for PM₁₀ and PM₂.₅, as fractions of emitted TSP, have for the period before 1995 been assigned by expert judgement, in cooperation with company experts. Fractions range from 60 to 95 % for PM₁₀ from 1980 until 2003 and from 30 to 80 % for PM₂.₅ during the same period of time. The suggested emission factors according to CEPMEIP¹⁷², valid for 1995, correspond to a value of 90 % for PM₁₀ and 80 % for PM₂.₅. For the years after 2003 the emission factors for PM₁₀ and PM₂.₅ are the same as for 2003.

Emissions of particles and metals from nine secondary non-ferrous metal smelters have been compiled. Emissions are for TSP and the metals as reported by the companies in environmental reports, and further into an emission database. The data in the database are for early years not complete and consistent, and several instances of missing values have had to be interpolated in order to complete the time series. Estimates of the emissions of PM₁₀ and PM₂.₅ were made using the same assumptions concerning particle size fractions as was applied for the large smelter.

Primary non-ferrous metal production is not associated with major dioxin emissions to air. From secondary processes, however, dioxin emissions are known to occur. Dioxin emissions from the large smelter, from the metal recycling company and from two smaller smelters are included for the whole time series, 1990 – 2014.

Chrome and nickel emissions from copper production are reported from submission 2016 onwards. For two smaller smelters, almost all emission data are obtained from the environmental reports and interpolated/extrapolated to cover the period 1990-2010. For the other two facilities, emission factors are obtained from

EMEP/EEA Guidebook 2013 with assumptions on applied abatement (see Table 4-17). Activity data is acquired from the facilities’ environmental reports.

Table 4-17. Cr and Ni emissions from copper production (NFR2C7c) at four major facilities in Sweden – sources and emission factors used.

<table>
<thead>
<tr>
<th>Facility</th>
<th>Type of production</th>
<th>Emissions of Ni</th>
<th>Emissions of Cr</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Source</td>
<td>EF, g/Mg copper</td>
<td>Source</td>
</tr>
<tr>
<td>1</td>
<td>Primary Facility production data and default EF from Guidebook 2013</td>
<td>0.57 (abatement with 97% efficiency assumed)</td>
<td>Facility production data and default EF from Guidebook 2013</td>
</tr>
<tr>
<td>2</td>
<td>Secondary Facility production data and default EF from Guidebook 2013</td>
<td>0.0039 (abatement with 97% efficiency assumed)</td>
<td>Not estimated</td>
</tr>
<tr>
<td>3</td>
<td>Secondary Emissions from environmental reports available for 2014+extrapolation</td>
<td>Emissions from environmental reports available for 2008-2014+extrapolation</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>Secondary Facility production data and default EF from Guidebook 2013</td>
<td>0.0039 (abatement with 97% efficiency assumed)</td>
<td>Emissions from environmental reports available for 2006 and 2008 + interpolation/extrapolation</td>
</tr>
</tbody>
</table>

4.4.6.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Time series from industrial processes reported in NFR codes 2C7c have been reviewed in later years and are consistent.

Uncertainties for NFR2C7c are displayed in Annex 1.

4.4.6.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No SOURCE-specific QA/QC or verification is performed.

4.4.6.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been performed in submission 2018.

4.4.6.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are currently planned.

4.4.7 Storage, handling and transport of metal products, NFR2C7d
Emissions from the source category Storage, handling and transport of metal products, NFR2C7d, have not been separated from the relevant metal production chapter and are included in respective source category, 2C1, 2C2, 2C3 or 2C7.
4.5 Other solvent and product use, NFR2D3

4.5.1 Road paving with asphalt, NFR2D3b

4.5.1.1 SOURCE CATEGORY DESCRIPTION

In this source category, emissions from road paving with asphalt are included. Due to confidentiality reasons emissions from asphalt roofing (NFR2D3c) are reported together with emissions from road paving.

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 varied within the interval 17-50% during 2002-2014. In this inventory, NMVOC and particles emitted in the process of asphalt paving of roads are included. CO emissions have not been estimated due to lack of information and reported NE in accordance with EMEP/EEA Guidebook 2013.

Table 4-18. Summary of key source analysis, NFR2D3, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2D3b</td>
<td>NMVOC</td>
<td>NMVOC</td>
</tr>
</tbody>
</table>

4.5.1.2 METHODOLOGICAL ISSUES

NMVOC emission estimates for the late 1980s and early 1990s are taken from investigations and inventories made in the early 1990s. Data from 2002 onwards 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 emitted when used. In the calculations emissions from imported solvent-based bitumen are not included. The amount of imported solvent-based bitumen is most likely very small. Emissions of NMVOC reported for the years in mid- and late 1990s were interpolated. For the years 2014, 2015 and 2016, data on amount of road paving mixtures delivered in Sweden was unavailable, and estimates are made based on the asphalt statistics\textsuperscript{173} and an implied emission factor for the latest year when this data is available (2013). Since production data regarding the total amount of road paving mixtures delivered in Sweden is only available with a lag of one year, production data and emissions for 2015 are updated. As for 2016, activity data has been set equal to 2015 and will be updated in Submission 2019.

\textsuperscript{173} Asphalt in figures http://www.eapa.org/asphalt.php available 2016-10-15
The emissions of NMVOC in 2005 were much higher than adjacent years. This is due to that the production of solvent-based bitumen in 2005 was extremely high, since it was used to repair roads that had been damaged by a severe storm ("Gud-run") that hit large areas of southern Sweden in early 2005.

Particle emissions from road paving with asphalt are reported from submission 2016 onwards. Activity data is obtained from asphalt statistics\textsuperscript{174}. Emissions factors are obtained from EMEP/EEA Guidebook 2013. According to the Guidebook, there are two main types of asphalt production technologies – batch mix and drum mix technologies – with different emission factors for particle fractions. Both technologies are applied in Sweden, but the exact proportion is not known. In particle emissions calculations, it is assumed that 50 % of all the asphalt is produced with batch mix technology and another 50 % with drum mix technology. Based on personal communication with branch representatives\textsuperscript{175}, it is also assumed that all the facilities use fabric filters to catch major part of the particles.

4.5.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions are based on expert judgement. Assessed uncertainties for emissions, from Road paving with asphalt and Asphalt roofing, of CO, NMVOC, TSP, PM\textsubscript{10} and PM\textsubscript{2.5} are ± 216\%, ± 191\%, ± 109\%, ± 97\%, and ± 100\%, respectively. More information is given in IIR Annex 1.

Time series from road paving with asphalt reported in NFR code 2D3b have been reviewed in later years and are considered to be consistent.

4.5.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC has been performed.

4.5.1.5 SOURCE-SPECIFIC RECALCULATIONS

Since production data regarding the total amount of road paving mixtures delivered in Sweden is only available with a lag of one year, production data and emissions for 2015 are updated. This leads to a decrease of calculated emissions by 4 \% for NMVOC and particles compared to the provided data in Submission 2017.

4.5.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are currently planned.

\textsuperscript{174} Asphalt in figures http://www.eapa.org/asphalt.php available 2016-10-15

\textsuperscript{175} Jan Wikström and Lorentz Lundqvist, NCC Roads AB
4.5.2 Asphalt roofing, NFR2D3c

4.5.2.1 SOURCE CATEGORY DESCRIPTION

In this source category, emissions from asphalt roofing are included. Due to confidentiality reasons emissions from asphalt roofing are reported together with emissions from road paving (NFR2D3b).

Emissions to air linked to the asphalt roofing industry consist mainly of particles, CO and non-methane volatile organic compounds (NMVOC), which are emitted from asphalt storing tanks and blowing stills, as well as from coater-mixer tanks and coaters. 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 and particle emissions. CO emissions are estimated with the default emission factors from EMEP/EEA Guidebook 2013. No measurements or estimations on Ni, Pb, Cd or Cr emissions have been performed by the industry and are consequently reported NE for the whole time-series, in accordance with EMEP/EEA Guidebook 2013.

4.5.2.2 METHODOLOGICAL ISSUES

Data on the total Swedish production of asphalt-saturated felt was provided by the producing companies. Emission factors for NMVOC and particles are based on measurements and calculations made by the manufacturers. The NMVOC emissions from the production of asphalt-saturated felt originate from the felt saturation and coating processes. In submission 2018 new information and measurements from both companies were presented leading to new calculations of NMVOC emissions for the whole time series. The new information also showed that no NVMOC emissions came from leakage from the asphalt storage tanks. The NMVOC and TSP emissions, 1990-2008, for one company are calculated by emission factors based on measurements in 2009. The NMVOC and TSP emissions for 2009-2016 are based on measurements. For the other company the NMVOC emissions, 1990-2015, are calculated by an emission factor based on the measurements in 2016. The factor used for estimating the TSP emission for the other company includes particles emitted from the mineral surfacing process as well as from storage and handling of the mineral products (0.005 kg/Mg), and are based on data from 1997. Emission factors for CO are obtained from EMEP/EEA Guidebook 2013. CO emissions from Swedish production of asphalt-saturated felt are reported from submission 2016 onwards. The notation key for activity data is C (classified). However, emissions from asphalt roofing are reported in 2D3 Road paving with asphalt due to confidentiality reasons.

4.5.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions are based on expert judgement. Assessed uncertainties for emissions, from Road paving with asphalt and Asphalt roofing, of CO, NMVOC, TSP, PM_{10} and PM_{2.5} are ± 216%, ± 191%, ± 109%, ± 97%, and ± 100%, respectively. More information is given in IIR Annex 1.

Time series from asphalt roofing have been reviewed in later years and are considered to be consistent.

4.5.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No SOURCE-specific QA/QC has been performed.

4.5.2.5 SOURCE-SPECIFIC RECALCULATIONS

Due to new information from the companies the time series for NMVOC and TSP have been recalculated and updated leading to a NMVOC emission decrease by on average 0.1 kt per year for the time period 1990-2015 and a decrease in TSP by on average 0.0006 kt per year for the time period 2009-2015. Emissions of BC have also been recalculated due to the new information for the time period 2009-2015.

4.5.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are currently planned.

4.5.3 Solvent use, NFR2D3a, NFR2D3d, NFR2D3e, NFR2D3f, NFR2D3g, NFR2D3h and NFR2D3i

4.5.3.1 SOURCE CATEGORY DESCRIPTION

Use of solvents and products containing solvents result in emissions of non-methylene volatile organic compounds (NMVOC). The model used for estimating the NMVOC emissions reported in the various solvent use categories is described in more detail in Annex 3.1 and is fully described in Skårman et al., 2016.

Emission estimates reported for solvent use in NFR2D3 include emissions from the source groups NFR2D3a “Domestic solvent use” (all domestic use except use of coatings), NFR2D3d “Coating applications” (industrial coating, domestic coating, non-industrial coating), NFR2D3e “Degreasing” (use of degreasing in industry), NFR2D3f “Dry cleaning” (non-domestic dry cleaning), NFR2D3g “Chemical product use” (vehicle industry, rubber industry, paint industry, textile industry, leather industry), NFR2D3h “Printing” (printing industry) and NFR2D3i “Other solvent and product use” (all other use of solvents).

Emissions of NMVOC from solvents and products containing solvents have decreased by 32 % from 84 kt NMVOC in 1990 to 57 kt NMVOC in 2015. This can

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largely be explained by the reduced use of solvents in coating application due to a shift to water-based paints.

NMVOC emissions from “Coating applications” (NFR2D3d) have decreased by almost 70% from 35 kt NMVOC in 1990 to 11 kt NMVOC in 2015. The largest source of NMVOC from solvents reported in NFR2D3 is, in later years, NFR2D3i “Other product and solvent use”. In this sub-sector an increase of emitted NMVOC from 1990 (13 kt) to 2015 (31 kt) can be observed.

4.5.3.1.1 NFR2D3a “Domestic solvent use”
Domestic solvent use is a moderate source of NMVOC but increases over time. This increase, starting in 2002, is due to an increased use of the product groups washer fluid, degreasing agents, and ignition fluids. However, a decrease in emissions from the use of ignition fluids can be seen for later years.
Two different emission factors are used for NFR2D3a “Domestic solvent use” for the whole time series:
- Diluted 0.275 (product groups that are used diluted in water)
- Not diluted 0.95 (product groups that are not used diluted in water)
The separation between diluted and not diluted products is a new approach compared to the old calculation model.

4.5.3.1.2 NFR2D3d “Coating applications”
Coating applications is a moderate source of NMVOC and has decreased over time. Coating in industry is the dominating source, followed by domestic coating, and that non-industry coating is of less importance. Emissions of NMVOC from coating application have decreased for the whole time series from 1990. The decrease is both due to reduced use of paints containing solvents and more efficient abatement technologies as indicated in available environmental reports.

4.5.3.1.3 NFR2D3e “Degreasing”
Degreasing within the industry is a minor source of NMVOC and has decreased over time. The estimates are based on abatement efficiency factors given in EMEP/EEA guidebook and the distribution between different abatement technologies has been based on information available in the GAINS-model (scenario: EGEO_Baseline_CLE) for 1995, 2000, 2005 and 2010. Emissions of both NMVOC have decreased from 1990, mainly due to a decreased use of degreasing products, but also a shift in technology, i.e. lower emission factors for the later years.

4.5.3.1.4 NFR2D3f “Dry cleaning”
Dry cleaning is a minor source of NMVOC. The time series for emissions of NMVOC from dry cleaning has decreased from 1990 mainly due to less use of dilution and thinner products.
4.5.3.1.5  **NFR2D3g “Chemical product use”**

Chemical product use is a minor source of NMVOC. The vehicle industry is the predominant source of emissions for chemical product use. The emissions are decreasing over time. The decrease during the 90’s is both due to reduced solvent content in used products, as well as more efficient abatement technologies according to information available in environmental reports for the rubber and vehicle industry. The sources in Chemical product use are:

- Vehicle industry
- Rubber industry
- Paint industry
- Textile and leather industry

4.5.3.1.6  **NFR2D3h “Printing industry”**

Printing industry is a minor source of NMVOC. A steady decrease in the emissions of NMVOC from 1990 depends on a reduced use of solvent products within the industry as well as a technology shift.

4.5.3.1.7  **NFR2D3i “Other solvent and product use”**

Other solvent and product use is a major source of NMVOC and has increased over time. The increased emissions for the activity are mainly due to a greater use of the product groups preservatives, refrigerants, metal mordants/etchants and coolant agents. These products account for about 70% of the increase.

4.5.3.2  **METHODOLOGICAL ISSUES**

Activity data regarding all solvent use sub-categories for year 1995 and onwards has been obtained from the Product register at the Swedish Chemicals Agency.

The Products Register does not provide reliable data for the period 1990-1994 for most industry categories. Data from reported time series compiled in a dedicated study on NMVOC emissions carried out by SMED in 2002 (Kindbom et. al, 2004) has been used for the estimations of emissions for 1990 for most sources. Exceptions are the emissions for 1990 for NFR2D3e “Degreasing”, “Vehicle industry” (included in NFR2D3g) and NFR2D3i “Other solvent and product use”. The 1990 emissions for “Degreasing” have been calculated with activity data from the GAINS-model and emission factors from EMEP/EEA. The 1990 emissions for the “Vehicle industry” are based on the information that the number of produced vehicles was around 22% lower in 1990 than in 1995, and this information has been used to calculate the NMVOC emissions for 1990. The 1990 emissions for “Other product and solvent use” are based on the correlation between GDP (gross domestic product) (Ekonomifakta, 2016) and emissions from 1995 to 2013. From known GDP for 1990 and the mathematical function for the correlation between emissions and GDP, emissions of NMVOC have been calculated.
The emissions for 1991-1994 have been interpolated based on the available information for 1990 and the known data for 1995.

Emission factors given in the literature, for example the EMEP/EEA Guidebook 2013, EU legislations, and other countries IIR’s, have been compiled and included in the model. The used emission factors are presented in Annex 3.1. The model has been developed in order to enable to test different datasets of emission factors. Two emission factors have been developed for each activity; one for solvents used as raw material and one for the remaining quantities. The emission factors for raw material have been set to 0.001 for all SNAP codes, since most of the solvents will end up in the product and will not be emitted during production.

A new emission factor for products used diluted in water has been introduced in the new model. The new emission factor is set to 0.275 and it has been calculated based on available information given in the EMEP/EEA Guidebook (Domestic solvent use). In the previous estimates these products were not treated separately and consequently the emission factor of 0.95 was used also for water diluted products.

The country specific emission factors have been developed in order to adjust to the old time series 1990-2001, developed by SMED in 2002 (Kindbom et. al., 2004). However, for some activities errors have been identified in previously reported data for 1990, and consequently those emissions have been corrected. Furthermore, application techniques, available information in the environmental reports for specific industries, as well as other pathways of release (e.g. water), have been considered when developing the country specific emission factors.

4.5.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Reported time series are considered to be consistent, except for the last year (2016) where data for previous year (2015) has been reported. This practice has been questioned by the ERT several times. The reason for Sweden to report activity data and emissions from solvent use 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.

Uncertainties for NMVOC emissions for NFR2D3a, NFR2D3d, NFR2D3e, NFR2D3f, NFR2D3e, NFR2D3f, NFR2D3h and NFR2D3i are ± 25% for years 1990-1994 and ± 15% for 1995 and onwards. More detailed information is to be found in IIR Annex 1.

The uncertainties have been discussed and assigned in co-operation with the Swedish Chemicals Agency. Uncertainty estimates for the emission factors were estimated by expert judgement. Information available in environmental reports, in the GAINS model and in the EMEP/EEA guidebook has been taken into account when developing the emission factors.
4.5.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

4.5.3.5 SOURCE-SPECIFIC RECALCULATIONS
In Table 4-19 it can be seen that the estimates according to the new model results in a significant reduction in the emissions of NMVOC from solvent use in comparison with the old estimates (-18% for 1990, -10% for 2005 and -28% for 2010). The decrease is mainly due to following changes within the model:

- Reallocation between sub-categories. Mainly from NFR2D3i “Other solvent and product use” to NFR2D3a “Domestic solvent use” but also to other sub-categories. In previous estimates an emission factor of 0.95 was used for all activities reported in NFR2D3i “Other solvent and product use”. In the new estimates the emission factor depends on which source group the activity is allocated to.
- Introduction of the new emission factor for products that are used diluted in water reported in “Domestic solvent use” and in “Other solvent use”. In previous estimates these products were not treated separately and consequently the emission factor of 0.95 was used. In the new model an emission factor of 0.275 is used.
Table 4-19. Recalculations of NMVOC.

<table>
<thead>
<tr>
<th>Year</th>
<th>NMVOC, old model</th>
<th>NMVOC, new model</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>102</td>
<td>84</td>
</tr>
<tr>
<td>1991</td>
<td>97</td>
<td>78</td>
</tr>
<tr>
<td>1992</td>
<td>92</td>
<td>72</td>
</tr>
<tr>
<td>1993</td>
<td>88</td>
<td>66</td>
</tr>
<tr>
<td>1994</td>
<td>83</td>
<td>60</td>
</tr>
<tr>
<td>1995</td>
<td>78</td>
<td>51</td>
</tr>
<tr>
<td>1996</td>
<td>75</td>
<td>51</td>
</tr>
<tr>
<td>1997</td>
<td>78</td>
<td>52</td>
</tr>
<tr>
<td>1998</td>
<td>76</td>
<td>52</td>
</tr>
<tr>
<td>1999</td>
<td>72</td>
<td>52</td>
</tr>
<tr>
<td>2000</td>
<td>69</td>
<td>53</td>
</tr>
<tr>
<td>2001</td>
<td>68</td>
<td>56</td>
</tr>
<tr>
<td>2002</td>
<td>68</td>
<td>61</td>
</tr>
<tr>
<td>2003</td>
<td>72</td>
<td>66</td>
</tr>
<tr>
<td>2004</td>
<td>76</td>
<td>68</td>
</tr>
<tr>
<td>2005</td>
<td>79</td>
<td>71</td>
</tr>
<tr>
<td>2006</td>
<td>80</td>
<td>72</td>
</tr>
<tr>
<td>2007</td>
<td>80</td>
<td>68</td>
</tr>
<tr>
<td>2008</td>
<td>77</td>
<td>62</td>
</tr>
<tr>
<td>2009</td>
<td>82</td>
<td>60</td>
</tr>
<tr>
<td>2010</td>
<td>86</td>
<td>62</td>
</tr>
<tr>
<td>2011</td>
<td>87</td>
<td>61</td>
</tr>
<tr>
<td>2012</td>
<td>83</td>
<td>59</td>
</tr>
<tr>
<td>2013</td>
<td>80</td>
<td>57</td>
</tr>
<tr>
<td>2014</td>
<td>80</td>
<td>57</td>
</tr>
</tbody>
</table>

The differences between the old and the new model are described in detail in Skårman et al., (2016)\textsuperscript{178}

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

\textsuperscript{178} Skårman et al., 2016. Swedish method for estimating emissions from Solvent Use. Further development of the calculation model. SMED report 192.
4.6 Other product use, NFR2G4

4.6.1 Source category description

NFR2G4 includes emissions from tobacco smoking and use of fireworks.

4.6.1.1 TOBACCO SMOKING

Emissions of NH₃, Ni, Zn and individual PAHs from tobacco smoking are reported for the first time in submission 2016, using emission factors from EMEP/EEA Guidebook 2013. The emission factors for NOₓ, CO, NMVOC, particles, BC and dioxin have been updated to EMEP/EEA 2013 and emission factors for As, Cd, Cr, Cu, Hg and Pb have been updated to those presented in the Norwegian IIR submission 2015.

4.6.1.2 FIREWORKS

Emissions of NOₓ, SO₂, CO, As, Cd, Cr, Cu, Hg, Ni and Zn are reported for the first time in submission 2016, using emission factors from EMEP/EEA 2013. Emission factors for particles have been updated to EMEP/EEA 2013. In addition, activity data has been updated for the entire time series, to consider export as well as import of fireworks. As a result, all emissions have been updated.

These changes result in large increases of some substances relative to previously reported emissions from the source, however these two emission sources are overall very small compared to the national total emissions.

Table 4-20. Summary of key source analysis, NFR2G, according to approach 1.

<table>
<thead>
<tr>
<th>NFR Level</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2G</td>
<td>Cu, Ni, PM₂.₅, PM₁₀, TSP, Cd</td>
<td>Cu, Ni</td>
</tr>
</tbody>
</table>

4.6.2 Methodological issues

Emissions of NOₓ, SO₂, NMVOC, NH₃, CO, particles, BC, most heavy metals, dioxins and PAH from tobacco smoking and use of fireworks are included in NFR2G. Emissions from tobacco smoking are based on activity data from official statistics on sold amounts of tobacco for the whole time series from 1980. Activity data include only “legal” purchases of tobacco products in Sweden; products that are purchased through tax-free and cross-border trading are not included. For fireworks, activity data is based on national statistics on imported and exported amounts of fireworks. No significant production of fireworks occurs in Sweden. The activity data for 1980-1987 has been assumed based on available data after 1987, which shows an increasing trend in the use of fireworks. An increasing trend has been applied also to the years where activity data is lacking. Emission factors from EMEP/EEA 2013¹⁷⁹ are used for estimates of emissions from fireworks, and

for tobacco smoking, the emission factors are taken from the Norwegian IIR submission 2015, and EMEP/EEA 2013. Emission factors for the two sources are listed in Table 4-21 and in Table 4-22, corresponding activity data is shown.

Table 4-21. Emission factors for tobacco smoking and use of fireworks.

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Tobacco smoking</th>
<th>Emission factor</th>
<th>Fireworks</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>NO&lt;sub&gt;x&lt;/sub&gt;</td>
<td>1.8 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>0.26 kg/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>SOx</td>
<td>NE</td>
<td>-</td>
<td>3.02 kg/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>CO</td>
<td>55.1 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>7.15 kg/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>NMVOC</td>
<td>4.84 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>NE</td>
<td>-</td>
</tr>
<tr>
<td>NH&lt;sub&gt;3&lt;/sub&gt;</td>
<td>4.15 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
<tr>
<td>TSP</td>
<td>27 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>109.83 kg/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>PM&lt;sub&gt;10&lt;/sub&gt;</td>
<td>27 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>99.92 kg/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>PM&lt;sub&gt;2.5&lt;/sub&gt;</td>
<td>27 kg/ton</td>
<td>EMEP/EEA 2013</td>
<td>51.94 kg/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>BC</td>
<td>0.45 % of PM&lt;sub&gt;2.5&lt;/sub&gt;</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
<tr>
<td>Pb</td>
<td>0.05 g/ton</td>
<td>Norwegian IIR, 2015</td>
<td>NE</td>
<td>-</td>
</tr>
<tr>
<td>Cd</td>
<td>0.1 g/ton</td>
<td>Norwegian IIR, 2015</td>
<td>1.48 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>Hg</td>
<td>0.1 g/ton</td>
<td>Norwegian IIR, 2015</td>
<td>0.057 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>As</td>
<td>0.159 g/ton</td>
<td>Norwegian IIR, 2015</td>
<td>1.33 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>Cr</td>
<td>0.354 g/ton</td>
<td>Norwegian IIR, 2015</td>
<td>15.6 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>Cu</td>
<td>0.152 g/ton</td>
<td>Norwegian IIR, 2015</td>
<td>444 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>Ni</td>
<td>2.7 g/ton</td>
<td>EMEP/EEA 2013</td>
<td>30 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>Zn</td>
<td>2.7 g/ton</td>
<td>EMEP/EEA 2013</td>
<td>260 g/ton</td>
<td>EMEP/EEA 2013</td>
</tr>
<tr>
<td>Dioxin</td>
<td>0.1 ug/ton</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
<tr>
<td>B(a)P</td>
<td>0.111 g/ton</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
<tr>
<td>B(b)F</td>
<td>0.045 g/ton</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
<tr>
<td>B(k)F</td>
<td>0.045 g/ton</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
<tr>
<td>I(1,2,3-cd)P</td>
<td>0.045 g/ton</td>
<td>EMEP/EEA 2013</td>
<td>NA</td>
<td>-</td>
</tr>
</tbody>
</table>
Table 4-22. Activity data for tobacco smoking and use of fireworks.

<table>
<thead>
<tr>
<th>Year</th>
<th>Amount consumed tobacco</th>
<th>Amount used fireworks</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>8475</td>
<td>987</td>
</tr>
<tr>
<td>1995</td>
<td>6688</td>
<td>1696</td>
</tr>
<tr>
<td>2000</td>
<td>5599</td>
<td>2556</td>
</tr>
<tr>
<td>2005</td>
<td>5719</td>
<td>2346</td>
</tr>
<tr>
<td>2006</td>
<td>5397</td>
<td>2425</td>
</tr>
<tr>
<td>2007</td>
<td>4882</td>
<td>3115</td>
</tr>
<tr>
<td>2008</td>
<td>4162</td>
<td>1920</td>
</tr>
<tr>
<td>2009</td>
<td>4326</td>
<td>1600</td>
</tr>
<tr>
<td>2010</td>
<td>4324</td>
<td>1749</td>
</tr>
<tr>
<td>2011</td>
<td>4417</td>
<td>1627</td>
</tr>
<tr>
<td>2012</td>
<td>4154</td>
<td>1381</td>
</tr>
<tr>
<td>2013</td>
<td>3993</td>
<td>1537</td>
</tr>
<tr>
<td>2014</td>
<td>4068</td>
<td>1543</td>
</tr>
<tr>
<td>2015</td>
<td>3863</td>
<td>1307</td>
</tr>
<tr>
<td>2016</td>
<td>3826</td>
<td>1328</td>
</tr>
</tbody>
</table>

Generally, emissions from tobacco smoking have decreased during the years. Emissions from the use of fireworks show an increasing trend during the years 1990 to 2007. The reported emissions for 2008 – 2015 have decreased compared to 2007 since fewer fireworks were imported.

4.6.3 Uncertainties and time-series consistency

Time series from industrial processes reported in NFR codes 2A-2H have been reviewed in later years and are consistent.

Uncertainties for NFR2G4 are displayed in Annex 1.

4.6.4 Source-specific QA/QC and verification

No SOURCE-specific QA/QC or verification is performed.

4.6.5 Source-specific recalculation

No recalculation have been performed in submission 2018

4.6.6 Source-specific planned improvements

No major improvements are currently planned.
4.7 Pulp and paper industry, NFR2H1

4.7.1 Source category description

NFR2H1 includes emissions from pulp and paper.

The pulp and paper industry in Sweden is an important source of industrial process emissions. Emissions from approximately 45 individual pulp and paper facilities were reported before 2002. After 2002 some plants were closed down and for 2016 emissions from 37 individual pulp and paper facilities are included in reported emissions. The Kraft process (sulphate) dominates in Sweden but there are also emissions from four sulphite facilities and 10 facilities that are mainly CTMP (Chemo Thermo Mechanical Pulp) or TMP (Thermo Mechanical Pulp) facilities reported in NFR2H1.

Reported emissions from the pulp and paper industry are for SO$_2$, NO$_X$ and TSP based on information in the companies’ environmental reports, while other air pollutants are calculated using nationally derived emission factors.

Table 4-23. Summary of key source analysis, NFR2H, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2H1</td>
<td>TSP, PM$<em>{10}$, PM$</em>{2.5}$, NMVOC, SO$_2$, As, CO, NH$_3$, NO$_X$, Pb, Hg, Cd, DIOX, Ni, Cr</td>
<td>TSP, PM$<em>{10}$, PM$</em>{2.5}$, NMVOC, SO$_2$, As, CO, NH$_3$, NO$_X$, Pb, Hg, DIOX, Cu, Zn, Ni</td>
</tr>
</tbody>
</table>

4.7.2 Methodological issues

Reported SO$_2$, NO$_X$ and TSP emissions from the pulp and paper industry are primarily based on information on production (Figure 4-2) and emissions in the companies’ environmental reports. The industrial organisation within this sector has for several years co-operated closely with its members in developing sector-specific methods of measuring and calculating emissions, which have resulted in high-quality emissions data. The emission factors that are used for the other pollutants are derived from national measurements and from international literature. The reported emissions of NMVOC do not include terpenes.
4.7.3 Uncertainties and time-series consistency

Uncertainties for emissions are based on the guidebook 2008 and expert judgement. For TSP, PM, NH$_3$ as well as heavy metals information from a national good practice guidance project is taken into account. More detailed information is to be found in IIR Annex 1.

Time series from industrial processes for pulp and paper industries reported in NFR codes 2H1 have been reviewed in later years and are consistent.

4.7.4 Source-specific QA/QC and verification

No source-specific QA/QC or verification is performed.

4.7.5 Source-specific recalculations

Emissions of NMVOC for 2015 for one plant have been corrected. Total NMVOC emissions in 2H1 for 2015 decrease therefore by 0.1 kt or 2%.

4.7.6 Source-specific planned improvements

Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.
4.8 Food and beverages industry, NFR2H2

4.8.1 Source category description

NFR2H2 includes emissions from the food and beverages industry. 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.

4.8.2 Methodological issues

Estimations of NMVOC emissions are based on activity data from different official statistics. For wine the estimation of NMVOC emissions are based on data on sold amount\(^1\) togethe\r
r with figures on import and export\(^2\). NMVOC emissions from beer production are based on the Swedish annual total production of beer\(^2\). 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\(^1\). For the NMVOC emission estimations emission factors presented in Table 4- 24, were used.

Table 4- 24. NMVOC emission factors for the reported production activities in NFR2H2 - Food and drink.

<table>
<thead>
<tr>
<th>Production activity</th>
<th>Emission factor</th>
<th>Unit</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wine</td>
<td>0.8</td>
<td>kg/1000 litres</td>
<td>180</td>
</tr>
<tr>
<td>Beer</td>
<td>0.35</td>
<td>kg/1000 litres</td>
<td>182</td>
</tr>
<tr>
<td>Liquors</td>
<td>0.6</td>
<td>kg/1000 litres</td>
<td>EF based on emission and activity data from one producer, 2001</td>
</tr>
<tr>
<td>Bread (sponge dough)</td>
<td>8</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Bread (white)</td>
<td>4.5</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Bread (wholemeal and light rye)</td>
<td>3</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Bread (dark rye)</td>
<td>0</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Cakes</td>
<td>0.1</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Biscuits</td>
<td>0.1</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Breakfast cereals</td>
<td>0.1</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Sugar</td>
<td>10</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Yeast</td>
<td>18</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Margarine and solid cooking fats</td>
<td>10</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Coffee roasting</td>
<td>0.55</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
<tr>
<td>Animal feed</td>
<td>0.1</td>
<td>kg/Mg</td>
<td>181</td>
</tr>
</tbody>
</table>

182 Bryggeriföreningen. http://sverigesbryggerier.se
4.8.3 Uncertainties and time-series consistency

Uncertainties for emissions are based on expert judgement. Assessed uncertainties for NMVOC are ± 200%. More information is given in IIR Annex 1.

Time series from industrial processes for food and beverages industries reported in NFR codes 2H2 have been reviewed in later years and are consistent.

4.8.4 Source-specific QA/QC and verification

No SOURCE-specific QA/QC or verification is performed.

4.8.5 Source-specific recalculations

Activity data, thus affecting reported NMVOC emissions, have been updated, due to new information for statistics available at Statistics Sweden’s website, for:

- Bread (sponge dough): Produced amount 2015
- Bread (white): Produced amount 2008-2015
- Bread (dark rye): Produced amount 2015
- Cakes: Produced amounts 2015
- Biscuits: Produced amounts 2012-2015
- Breakfast cereals: Produced amounts 2015
- Sugar: Produced amounts 2008-2015
- Margarine and solid cooking fats: Produced amounts 2015
- Animal feed: Produced amounts for 2015
- Coffee roasting: Produced amounts 2015
- Yeast: Produced amounts 2015
- Wine spirits, beer and cider: 2013-2015

The recalculations resulted in decreased NMVOC emissions of about 0.076 kt 2015.

4.8.6 Source-specific planned improvements

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

4.9 Wood processing, NFR2I

4.9.1 Source category description

TSP emissions from wood processing are included for the first time in submission 2016. Emissions from production of plywood, MDF boards and fibre boards are included in NFR2I Wood processing. The number of companies, as well as the amount of produced wood products have decreased over the time series, with 18 companies in 1990 producing around 670 kt to three companies in 2016, producing around 380 kt.
Table 4-25. Summary of key source analysis, NFR2I, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>2I</td>
<td>TSP</td>
<td></td>
</tr>
</tbody>
</table>

4.9.2 **Methodological issues**
Activity data is retrieved from Trä- och Möbelföretagen\(^183\), a Swedish trade organisation for wood and furniture products. For the years 1991-1999, production quantities are interpolated as there is no available data for those years.

TSP emissions are calculated using activity data and the emission factor from EMEP/EEA air pollutant emission inventory guidebook 2013.

4.9.3 **Uncertainties and time-series consistency**
The time series reported in the NFR code 2I has been reviewed and is consistent. Uncertainties for emissions of TSP are ± 20%. More detailed information is to be found in IIR Annex 1.

4.9.4 **Source-specific QA/QC and verification**
No source-specific QA/QC or verification is performed.

4.9.5 **Source-specific recalculations**
As data was not available in time for submission, emissions for 2015 have been updated. Emissions of TSP increase therefore by about 0.03 kt.

4.9.6 **Source-specific planned improvements**
No major improvements are currently planned.

\(^{183}\) [http://www.tmf.se/english_1/about_tmf_1](http://www.tmf.se/english_1/about_tmf_1)
4.10 Production of POPs, NFR2J

To the knowledge of the Swedish inventory compilers, there is no production of POPs in Sweden. Thus no emissions are reported from the source category NFR2J.

4.11 Consumption of POPs and heavy metals, NFR2K

Emissions from consumption of POPs and heavy metals are not included in the Swedish emission inventory. Calculations have been made using default emission factors from the EEA/EMEP Guidebook 2013, however resulting emissions were judged to be unreasonably high for Swedish conditions.\textsuperscript{184}

4.12 Other production, consumption, storage, transportation or handling of bulk products, NFR2L

No other production, consumption, storage, transportation or handling of bulk products occur in Sweden, thus no emissions are reported in NFR2L.

\textsuperscript{184} Yaramenka, K., Mawdsley, I., Gustafsson, T. 2014. Utveckling av rapportering till CLRTAP NFR 1B, 2 och 5 map EMEP Guidebook, steg 1. SMED rapport nr 161 2014 (available in Swedish)
5 Agriculture (NFR sector 3)

5.1 Overview

In this sector emissions of ammonia, NMVOC, NO\textsubscript{X} and particulate matter (TSP, PM\textsubscript{10} and PM\textsubscript{2.5}) are reported. The general trend for all emissions is a continuous decline. One of the driving forces to this is a decreasing agricultural sector in Sweden which has resulted in a decrease in agriculture land and decreasing livestock populations. Over the past 50 years the agriculture in Swedish has undergone radical structural changes and rationalisations. 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. 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. From 1990 there has been a steady decrease in the number of dairy cows. However, milk yield per head has increased.

5.2 Manure management, NFR3B

5.2.1.1 SOURCE CATEGORY DESCRIPTION

Manure management and housing of livestock give rise to emissions of ammonia, NMVOC, NO\textsubscript{X} and particulate matters. Emissions of ammonia mainly derive from manure management. That is storage, handling and application of stable manure. Emissions from application of manure are calculated here but reported under sector 3D. NMVOC emissions from animal husbandry originate from feed, especially silage, degradation and decomposition of feed in the rumen and in manure. In the Swedish inventory emissions of NMVOC from manure management from dairy cattle is the main contributor. Nitric oxide is formed through biological oxidation (i.e. nitrification) of ammonia or ammonium by aerobic bacteria. Nitric oxide is emitted from the surface layers of stored manure, during application of manure to soil and from deposition of excreta during grazing. Simultaneously emissions of nitrous oxide (N\textsubscript{2}O) occur, but are instead accounted for in the reporting to UNFCCC, being a greenhouse gas. Housing of livestock causes emissions of particulate matter. The emissions originate mainly from feed, but bedding materials such as straw or wood shavings can also give rise to airborne particulates.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>3B1 Dairy cattle</td>
<td>NH\textsubscript{3}, NMVOC, PM\textsubscript{2.5}, TSP, PM\textsubscript{10}</td>
<td>NH\textsubscript{3}, NMVOC, PM\textsubscript{2.5}</td>
</tr>
<tr>
<td>3B1 Non-dairy cattle</td>
<td>NH\textsubscript{3}, NMVOC, TSP, PM\textsubscript{2.5}, PM\textsubscript{10}</td>
<td>NH\textsubscript{3}, NMVOC, PM\textsubscript{2.5}</td>
</tr>
<tr>
<td>3B3 Swine</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3B4 Fur-bearing a.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3B4 Horses</td>
<td>NMVOC, NH\textsubscript{3}, TSP</td>
<td>NMVOC, NH\textsubscript{3}</td>
</tr>
<tr>
<td>3B4 Poultry</td>
<td>PM\textsubscript{10}, TSP, PM\textsubscript{2.5}, NMVOC, NH\textsubscript{3}</td>
<td>PM\textsubscript{10}, TSP, PM\textsubscript{2.5}, NMVOC, NH\textsubscript{3}</td>
</tr>
<tr>
<td>3B4 Sheep</td>
<td>NH\textsubscript{3}, PM\textsubscript{2.5}, PM\textsubscript{10}, TSP</td>
<td>NH\textsubscript{3}, TSP</td>
</tr>
</tbody>
</table>
5.2.1.2 METHODOLOGICAL ISSUES

5.2.1.2.1 Activity data

One of the main sources of activity data to calculate the emission from the agricultural sector is the survey “Use of fertilisers and animal manure in agriculture”. The survey is performed by Statistics Sweden and was carried out every second year until 2013, but after that it is schemed for every third year instead. The latest was for 2015/2016. The field investigation collects data from 5150 agricultural holdings which is an increase of about 1500 agricultural holdings since 2013. Detailed information about the design and implementation of the field investigation can be found in the report MI 30 SM 1702. From this survey data on, distribution of different manure management systems (solid manure, liquid manure and deep litter), design of manure containers (e.g. with or without a roof), time and method of manure application, timespan before mulching manure and data on stable periods is obtained. Since dairy cows often are stabled at night and also spend time in the stables during milking, the data on stable periods is combined with the assumption that 38% of the dairy cows manure is produced in the stable during the grazing period.

Another important source of activity data is the farm register that the Swedish Board of Agriculture is responsible for. From this register data on livestock population is obtained for most animal categories (table 5-2 and 5-3). Although, some other sources are used for slaughter chicken, horses and furred animals. Concerning horses the Farm Register underestimates the number of horses because only horses on farms are included (i.e. not horses for leisure activities). Three separate surveys estimated total number of horses in Sweden in 2004, 2010 and 2016. These estimates are used in the calculations instead. To estimate the number of slaughter chickens we use the Swedish official slaughter statistics together with timespan between production rounds to estimate the average yearly population. The number of minks is provided by the Swedish Furred Animals Association.

Data on manure and nitrogen excretion for different animals are compiled by the Swedish board of agriculture and based on nutrient balance calculations. The underlying data are based on a variety of sources. The data for the most significant animal groups (i.e. cattle and swine) are from public reports produced by the Swedish Board of Agriculture. Some of the data for the less significant animal groups are based on expert opinions.

Figures about milk yield for dairy cows, which affects the amount of nitrogen and manure excreted, are obtained from the Swedish Dairy Association. For the calculation of particle emissions from laying hens, information on the proportion of hens

\[185\] Statistics Sweden, report series MI 30 SM. http://www.scb.se/mi1001

\[186\] Swedish Board of Agriculture, report series JO 25 SM http://www.scb.se/jo0107
held in cages and in percheries is needed since different emission factors are used for these two production systems. This information is obtained from The Swedish Board of Agriculture.\(^{187}\)

### Table 5.2. Population size for cattle and swine (1000s heads).

<table>
<thead>
<tr>
<th>Year</th>
<th>Dairy cows</th>
<th>Suckler cows</th>
<th>Heifers</th>
<th>Bulls and steers</th>
<th>Calves</th>
<th>Sows (*)</th>
<th>Pig for meat production</th>
<th>Piglets</th>
<th>Boars</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>576</td>
<td>75</td>
<td>337</td>
<td>206</td>
<td>524</td>
<td>221</td>
<td>1 276</td>
<td>758</td>
<td>8.6</td>
</tr>
<tr>
<td>1995</td>
<td>482</td>
<td>157</td>
<td>370</td>
<td>226</td>
<td>542</td>
<td>237</td>
<td>1 300</td>
<td>768</td>
<td>7.6</td>
</tr>
<tr>
<td>2000</td>
<td>428</td>
<td>167</td>
<td>365</td>
<td>224</td>
<td>500</td>
<td>202</td>
<td>1 146</td>
<td>566</td>
<td>4.2</td>
</tr>
<tr>
<td>2005</td>
<td>393</td>
<td>177</td>
<td>327</td>
<td>200</td>
<td>508</td>
<td>185</td>
<td>1 085</td>
<td>538</td>
<td>2.7</td>
</tr>
<tr>
<td>2010</td>
<td>348</td>
<td>197</td>
<td>322</td>
<td>191</td>
<td>479</td>
<td>154</td>
<td>937</td>
<td>427</td>
<td>2.3</td>
</tr>
<tr>
<td>2014</td>
<td>344</td>
<td>186</td>
<td>312</td>
<td>179</td>
<td>472</td>
<td>143</td>
<td>857</td>
<td>376</td>
<td>1.7</td>
</tr>
<tr>
<td>2015</td>
<td>340</td>
<td>184</td>
<td>311</td>
<td>178</td>
<td>467</td>
<td>140</td>
<td>830</td>
<td>384</td>
<td>1.5</td>
</tr>
<tr>
<td>2016</td>
<td>331</td>
<td>194</td>
<td>305</td>
<td>184</td>
<td>475</td>
<td>139</td>
<td>835</td>
<td>378</td>
<td>1.5</td>
</tr>
</tbody>
</table>

(*) 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.

### Table 5.3. Population size for the other animal groups (1000s heads).

<table>
<thead>
<tr>
<th>Year</th>
<th>Sheep</th>
<th>Lamb</th>
<th>Horses (*)</th>
<th>Goats (***</th>
<th>Kids (***</th>
<th>Reindeer</th>
<th>Fur-bearing animals</th>
<th>Laying hens</th>
<th>Turkeys</th>
<th>Chickens</th>
<th>Slaughter Chickens (**)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>162</td>
<td>244</td>
<td>316</td>
<td>2.9</td>
<td>1.4</td>
<td>271</td>
<td>297</td>
<td>6 400</td>
<td>122</td>
<td>2 200</td>
<td>4 476</td>
</tr>
<tr>
<td>1995</td>
<td>195</td>
<td>266</td>
<td>316</td>
<td>3.5</td>
<td>1.8</td>
<td>253</td>
<td>254</td>
<td>6 100</td>
<td>122</td>
<td>1 800</td>
<td>7 055</td>
</tr>
<tr>
<td>2000</td>
<td>198</td>
<td>234</td>
<td>316</td>
<td>3.5</td>
<td>1.8</td>
<td>221</td>
<td>276</td>
<td>5 700</td>
<td>122</td>
<td>1 700</td>
<td>7 896</td>
</tr>
<tr>
<td>2005</td>
<td>222</td>
<td>249</td>
<td>323</td>
<td>3.7</td>
<td>1.8</td>
<td>251</td>
<td>290</td>
<td>5 100</td>
<td>122</td>
<td>1 700</td>
<td>8 453</td>
</tr>
<tr>
<td>2010</td>
<td>273</td>
<td>292</td>
<td>363</td>
<td>3.7</td>
<td>1.8</td>
<td>250</td>
<td>180</td>
<td>6 061</td>
<td>130</td>
<td>1 647</td>
<td>9 159</td>
</tr>
<tr>
<td>2014</td>
<td>287</td>
<td>301</td>
<td>363</td>
<td>3.7</td>
<td>1.8</td>
<td>254</td>
<td>210</td>
<td>6 549</td>
<td>156</td>
<td>1 713</td>
<td>10 319</td>
</tr>
<tr>
<td>2015</td>
<td>289</td>
<td>306</td>
<td>363</td>
<td>3.7</td>
<td>1.8</td>
<td>250</td>
<td>210</td>
<td>7 571</td>
<td>156</td>
<td>1 842</td>
<td>11 044</td>
</tr>
<tr>
<td>2016</td>
<td>281</td>
<td>297</td>
<td>356</td>
<td>3.7</td>
<td>1.8</td>
<td>247</td>
<td>200</td>
<td>8 174</td>
<td>130</td>
<td>1 575</td>
<td>11 659</td>
</tr>
</tbody>
</table>

Data are from the farm register, except: (*) estimated number of horses 2004, 2010 and 2016, by the Swedish Board of Agriculture. (**) Slaughter statistics, Swedish board of agriculture. (*** Data on goats were available until 1992, this data have been extrapolated.

**Table 5.4. Livestock stable periods (months).**

<table>
<thead>
<tr>
<th>Year</th>
<th>Dairy cows</th>
<th>Suckler cows</th>
<th>Heifers</th>
<th>Bulls and steers</th>
<th>Calves</th>
<th>Sheep, horses, goats</th>
<th>Reindeer</th>
<th>Poultry, Swine and Fur-bearing animals</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>7.2</td>
<td>6.2</td>
<td>6.5</td>
<td>7.6</td>
<td>7.8</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>1995</td>
<td>7.2</td>
<td>6.2</td>
<td>6.5</td>
<td>7.6</td>
<td>7.8</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>2000</td>
<td>7.2</td>
<td>5.8</td>
<td>6.1</td>
<td>7.6</td>
<td>7.6</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>2005</td>
<td>6.9</td>
<td>4.9</td>
<td>5.6</td>
<td>8.4</td>
<td>7.6</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>2010</td>
<td>7.2</td>
<td>5.5</td>
<td>5.8</td>
<td>8.8</td>
<td>8.2</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>2014</td>
<td>7.3</td>
<td>5.8</td>
<td>6.0</td>
<td>9.8</td>
<td>8.2</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>2015</td>
<td>7.3</td>
<td>5.5</td>
<td>5.9</td>
<td>9.2</td>
<td>8.1</td>
<td>6</td>
<td>0</td>
<td>12</td>
</tr>
<tr>
<td>2016</td>
<td>7.3</td>
<td>5.5</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(*) Data from Statistics Sweden’s survey “Use of fertilisers and animal manure in agriculture”.

**Table 5.5. Liquid waste management systems (fractions).**

<table>
<thead>
<tr>
<th>Year</th>
<th>Dairy cattle (*)</th>
<th>Suckler cows (*)</th>
<th>Heifers (*)</th>
<th>Bulls and steers (*)</th>
<th>Calves (*)</th>
<th>Pigs for meat production (*)</th>
<th>Other swine (*)</th>
<th>Sheep, goats</th>
<th>Reindeer</th>
<th>Poultry, Laying Hens, Chickens</th>
<th>Swine, Slaughter Chickens, Turkeys</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>0.23</td>
<td>0.16</td>
<td>0.19</td>
<td>0.20</td>
<td>0.44</td>
<td>0.44</td>
<td>0</td>
<td>0.25</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1995</td>
<td>0.31</td>
<td>0.21</td>
<td>0.26</td>
<td>0.26</td>
<td>0.63</td>
<td>0.63</td>
<td>0</td>
<td>0.25</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2000</td>
<td>0.39</td>
<td>0.13</td>
<td>0.16</td>
<td>0.16</td>
<td>0.81</td>
<td>0.81</td>
<td>0</td>
<td>0.25</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2005</td>
<td>0.51</td>
<td>0.06</td>
<td>0.23</td>
<td>0.19</td>
<td>0.93</td>
<td>0.93</td>
<td>0</td>
<td>0.21</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2010</td>
<td>0.58</td>
<td>0.11</td>
<td>0.23</td>
<td>0.17</td>
<td>0.94</td>
<td>0.94</td>
<td>0</td>
<td>0.12</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2014</td>
<td>0.64</td>
<td>0.11</td>
<td>0.26</td>
<td>0.17</td>
<td>0.95</td>
<td>0.95</td>
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<td>0.09</td>
<td>0.00</td>
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</tr>
<tr>
<td>2015</td>
<td>0.64</td>
<td>0.11</td>
<td>0.26</td>
<td>0.17</td>
<td>0.95</td>
<td>0.95</td>
<td>0</td>
<td>0.09</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2016</td>
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<td>0.27</td>
<td>0.22</td>
<td>0.97</td>
<td>0.97</td>
<td>0</td>
<td>0.09</td>
<td>0.00</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

(*) Data from Statistics Sweden’s survey “Use of fertilisers and animal manure in agriculture”.

**Table 5.6. Solid waste management systems (fractions).**

<table>
<thead>
<tr>
<th>Year</th>
<th>Dairy cattle (*)</th>
<th>Suckler cows (*)</th>
<th>Heifers (*)</th>
<th>Bulls and steers (*)</th>
<th>Calves (*)</th>
<th>Pigs for meat production (*)</th>
<th>Other swine (*)</th>
<th>Sheep, goats</th>
<th>Reindeer</th>
<th>Poultry, Laying Hens, Chickens</th>
<th>Swine, Slaughter Chickens, Turkeys</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>0.52</td>
<td>0.29</td>
<td>0.30</td>
<td>0.35</td>
<td>0.36</td>
<td>0.54</td>
<td>0.45</td>
<td>0.5</td>
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<td>0.48</td>
<td>0.55</td>
</tr>
<tr>
<td>1995</td>
<td>0.44</td>
<td>0.23</td>
<td>0.25</td>
<td>0.29</td>
<td>0.29</td>
<td>0.35</td>
<td>0.26</td>
<td>0.5</td>
<td>1</td>
<td>0.48</td>
<td>0.55</td>
</tr>
<tr>
<td>2000</td>
<td>0.35</td>
<td>0.28</td>
<td>0.29</td>
<td>0.37</td>
<td>0.37</td>
<td>0.37</td>
<td>0.67</td>
<td>0.5</td>
<td>1</td>
<td>0.48</td>
<td>0.55</td>
</tr>
<tr>
<td>2005</td>
<td>0.22</td>
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<td>0.17</td>
<td>0.27</td>
<td>0.23</td>
<td>0.06</td>
<td>0.46</td>
<td>0.5</td>
<td>1</td>
<td>0.48</td>
<td>0.73</td>
</tr>
<tr>
<td>2010</td>
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<td>0.20</td>
<td>0.15</td>
<td>0.26</td>
<td>0.22</td>
<td>0.05</td>
<td>0.30</td>
<td>0.5</td>
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<td>0.48</td>
<td>0.88</td>
</tr>
<tr>
<td>2014</td>
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<td>0.20</td>
<td>0.12</td>
<td>0.25</td>
<td>0.16</td>
<td>0.03</td>
<td>0.36</td>
<td>0.5</td>
<td>1</td>
<td>0.48</td>
<td>0.78</td>
</tr>
<tr>
<td>2015</td>
<td>0.11</td>
<td>0.20</td>
<td>0.12</td>
<td>0.25</td>
<td>0.16</td>
<td>0.03</td>
<td>0.36</td>
<td>0.5</td>
<td>1</td>
<td>0.48</td>
<td>0.78</td>
</tr>
<tr>
<td>2016</td>
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<td>0.15</td>
<td>0.09</td>
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<td>1</td>
<td>0.48</td>
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</tbody>
</table>

(*) Data from Statistics Sweden’s survey “Use of fertilisers and animal manure in agriculture”.
Table 5-7. Deep litter waste management systems (fractions).

<table>
<thead>
<tr>
<th>Year</th>
<th>Dairy cattle (*)</th>
<th>Suckler cows (*)</th>
<th>Heifers (*)</th>
<th>Bulls and steers (*)</th>
<th>Calves (*)</th>
<th>Pigs for meat production (*)</th>
<th>Other swine (*)</th>
<th>Sheep, goats, reindeer, Fur-bearing animals</th>
<th>Horses</th>
<th>Laying Hens, Chickens</th>
<th>Slaughter Chickens, Turkeys</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>0.01</td>
<td>0.07</td>
<td>0.08</td>
<td>0.09</td>
<td>0.02</td>
<td>0.11</td>
<td>0</td>
<td>0.02</td>
<td>0.20</td>
<td>1.00</td>
<td></td>
</tr>
<tr>
<td>1995</td>
<td>0.01</td>
<td>0.07</td>
<td>0.08</td>
<td>0.09</td>
<td>0.02</td>
<td>0.11</td>
<td>0</td>
<td>0.02</td>
<td>0.20</td>
<td>1.00</td>
<td></td>
</tr>
<tr>
<td>2000</td>
<td>0.00</td>
<td>0.16</td>
<td>0.10</td>
<td>0.20</td>
<td>0.22</td>
<td>0.01</td>
<td>0.07</td>
<td>0.02</td>
<td>0.02</td>
<td>0.07</td>
<td>1.00</td>
</tr>
<tr>
<td>2010</td>
<td>0.01</td>
<td>0.15</td>
<td>0.11</td>
<td>0.19</td>
<td>0.30</td>
<td>0.01</td>
<td>0.07</td>
<td>0.02</td>
<td>0.02</td>
<td>0.00</td>
<td>1.00</td>
</tr>
<tr>
<td>2014</td>
<td>0.01</td>
<td>0.17</td>
<td>0.12</td>
<td>0.23</td>
<td>0.35</td>
<td>0.02</td>
<td>0.06</td>
<td>0.02</td>
<td>0.13</td>
<td>1.00</td>
<td></td>
</tr>
<tr>
<td>2015</td>
<td>0.01</td>
<td>0.17</td>
<td>0.12</td>
<td>0.23</td>
<td>0.35</td>
<td>0.02</td>
<td>0.06</td>
<td>0.02</td>
<td>0.13</td>
<td>1.00</td>
<td></td>
</tr>
<tr>
<td>2016</td>
<td>0.01</td>
<td>0.20</td>
<td>0.12</td>
<td>0.23</td>
<td>0.31</td>
<td>0.01</td>
<td>0.17</td>
<td>0.02</td>
<td>0.01</td>
<td>1.00</td>
<td></td>
</tr>
</tbody>
</table>

(*) Data from Statistics Sweden’s survey “Use of fertilisers and animal manure in agriculture”.

5.2.1.2.2 Emissions of ammonia (Tier 2) and emissions of nitric oxide

To estimate the emission of ammonia until 2004 and the emission of NOX for the complete time series, Sweden use a slightly modified version of the default Tier 2 model described in the EMEP/EEA guidebook 2016. The ammonia emissions from 2005 and onwards are instead estimated with a country specific model (described in detail below together with an explanation of how we use this approach). The main modification of the Tier 2 model is that we use total N instead of TAN to estimate the emissions from stable ventilation and manure storage because the country specific emissions factors we use are developed in that way. The emission factors we use are the same as in the country specific model but here aggregated to fit the calculation level used in the default model (e.g. for different livestock categories instead of different spreading methods as we use in the CS model). The rationale for not using the default emission factors is that the climate in Sweden is considerably cooler than the European average, and this has a profound effect on the average annual ammonia emission factors.\(^{188}\)

The default Tier 2 methodology for both ammonia and NOX follows the same stepwise procedure in the guidebook. We have followed this methodology for all the livestock categories in table 5-2 and 5-3. To ensure consistency with the greenhouse gas inventory the same parameters have been used in both inventories for nitrogen excretion rates and distribution of different manure management systems for all livestock categories.

\(^{188}\) Swedish Institute of Agricultural and Environmental Engineering (JTI) 2002
Step 1
By using the annual nitrogen excretion rates (table 5-10) combined with the distribution of different manure management systems (table 5-5 to 5-7) and stable periods (table 5-4) we calculate the amount of total-N that is excreted in the stables and on the grazing grounds.

\[ m_{\text{build\_Solid\_N}}, m_{\text{build\_Liquid\_N}}, m_{\text{build\_Deep\_N}}, m_{\text{graz\_N}} \]

Step 2
Here we calculate the NH$_3$-N losses from the livestock buildings by multiplying the amount of excreted nitrogen with the emission factor for solid manure, liquid manure and deep litter, respectively (table 5-8).

\[
E_{\text{build\_Solid\_NH}_3N} = m_{\text{build\_Solid\_N}} \times EF_{\text{build\_Solid}} \\
E_{\text{build\_Liquid\_NH}_3N} = m_{\text{build\_Liquid\_N}} \times EF_{\text{build\_Liquid}} \\
E_{\text{build\_Deep\_NH}_3N} = m_{\text{build\_Deep\_N}} \times EF_{\text{build\_Deep}}
\]

Step 3
In this step we estimate the amount of total-N in the manure that remains after the ventilation losses. This is the amount that is stored in the different manure management systems.

\[
m_{\text{storage\_Solid\_N}} = m_{\text{build\_Solid\_N}} - E_{\text{build\_Solid\_NH}_3N} \\
m_{\text{storage\_Liquid\_N}} = m_{\text{build\_Liquid\_N}} - E_{\text{build\_Liquid\_NH}_3N} \\
m_{\text{storage\_Deep\_N}} = m_{\text{build\_Deep\_N}} - E_{\text{build\_Deep\_NH}_3N}
\]

Step 4
Here we estimate the nitrogen losses that occur from emissions of NH$_3$ as well as NO$_X$ during storage. In Sweden no manure is spread directly without being stored before applied so this simplifies the calculations compared to the guidelines. The emission factors used can be seen in table 5-9. To estimate the emissions of NO$_X$ we have used the default emission factors from the guidebook together with the country specific values of the proportion of total N that is ammonical nitrogen (table 5-10).

\[
E_{\text{storage\_Solid\_NH}_3N} = m_{\text{storage\_Solid\_N}} \times EF_{\text{Solid\_Storage}} \\
E_{\text{storage\_Liquid\_NH}_3N} = m_{\text{storage\_Liquid\_N}} \times EF_{\text{Liquid\_Storage}} \\
E_{\text{storage\_Deep\_NH}_3N} = m_{\text{storage\_Deep\_N}} \times EF_{\text{Deep\_Storage}}
\]

\[
E_{\text{storage\_Solid\_NO}_X} = (m_{\text{storage\_Solid\_N}} \times TAN_{\text{Solid}} + m_{\text{storage\_Deep\_N}} \times TAN_{\text{Deep}}) \times 0.01 \\
E_{\text{storage\_Liquid\_NO}_X} = m_{\text{storage\_Liquid\_N}} \times TAN_{\text{Liquid}} \times 0.0001
\]
Step 5
In this step we calculate the amount of N that is available for application to agricultural soil. That is, subtracting the amount of nitrogen lost during storage.

\[
\begin{align*}
    m_{\text{applic Solid N}} &= m_{\text{storage Solid N}} - E_{\text{storage Solid NH}_3N} \\
    m_{\text{applic Liquid N}} &= m_{\text{storage Liquid N}} - E_{\text{storage Liquid NH}_3N} \\
    m_{\text{applic Deep N}} &= m_{\text{storage Deep N}} - E_{\text{storage Deep NH}_3N}
\end{align*}
\]

Step 6
Emissions of ammonia from application of manure are estimated. First the available total N is translated to amount of TAN, because the country specific emission factors we use are expressed as fraction of TAN lost as ammonia (table 5-10).

\[
\begin{align*}
    E_{\text{applic Solid NH}_3N} &= m_{\text{applic Solid N}} \times \text{TAN Solid} \times E_{\text{Solid Appl}} \\
    E_{\text{applic Liquid NH}_3N} &= m_{\text{applic Liquid N}} \times \text{TAN Liquid} \times E_{\text{Liquid Appl}} \\
    E_{\text{applic Deep NH}_3N} &= m_{\text{applic Deep N}} \times \text{TAN Deep} \times E_{\text{Deep Appl}}
\end{align*}
\]

Step 7
The emissions of NO_x are estimated based on total amount of TAN applied to agricultural soils. The emission factor used is the default one from the guidebook.

\[
m_{\text{applic Tot TAN}} = ((m_{\text{applic Solid N}} \times \text{TAN solid}) + (m_{\text{applic Shurry N}} \times \text{TAN Shurry}) + (m_{\text{applic Deep N}} \times \text{TAN Deep})) \times 0.026
\]

Step 8
Here we calculate the emissions of ammonia from grazing (although reported under NFR3D). Emission factors in table 5-13.

\[
E_{\text{graz NH}_3N} = m_{\text{graz N}} \times E_{\text{grazing}}
\]

Step 9
In this final step the different emissions are aggregated to the relevant NFR categories and converted to the mass of the specific compounds. The reason we use 46/14 for NO_x instead of the value stated in the guidebook (30/14) is that the value for nitric oxide should be reported as NO_2 (i.e. not NO) in the NFR-tables.

Reported per animal (m) in 3B, manure management

\[
\begin{align*}
    EM_{3.B.m.}\text{NH}_3 &= (E_{\text{build Solid NH}_3N} + E_{\text{build Liquid NH}_3N} + E_{\text{build Deep NH}_3N} + E_{\text{storage Solid NH}_3N} + E_{\text{storage Liquid NH}_3N} + E_{\text{storage Deep NH}_3N}) \times 17/14 \\
    EM_{3.B.m.}\text{NO}_2 &= (E_{\text{storage Solid NO}_2N} + E_{\text{storage Liquid NO}_2N}) \times 46/14
\end{align*}
\]

Reported as a sum for all animals in 3Da2a, animal manure applied to soils

\[
\begin{align*}
    EM_{3.D.a.2.a.}\text{NH}_3 &= (E_{\text{applic Solid NH}_3N} + E_{\text{applic Liquid NH}_3N} + E_{\text{applic Deep NH}_3N}) \times 17/14
\end{align*}
\]

Reported as a sum for all animals in 3Da3, urine and dung deposited by grazing animals

\[
EM_{3.D.a.3.}\text{NH}_3 = E_{\text{graz NH}_3N} \times 17/14
\]
5.2.1.2.3  Emissions of ammonia (country specific model)

Sweden has also developed a country specific methodology to estimate emissions of ammonia from agriculture. The methodology\textsuperscript{189,190} is developed in collaboration between the Swedish EPA, Statistics Sweden, the Swedish Board of Agriculture and the Swedish Institute of Agricultural and Environmental Engineering. Several of the questions to the farmers in the Statistics Sweden’s field investigation among farmers are also designed to provide the model with correct and accurate activity data. This model is only possible to use from 2005 and forward. The reason is that prior to this year it is not possible to acquire the detailed micro data from the Statistics Sweden’s field investigation among farmers that is needed as activity data in the model. However, when 2005 is Sweden’s base year for ammonia reduction commitments in the EU national emission ceilings directive (NEC) and the Gothenburg protocol this is the most important part of the time series. Hence, the only available alternative strategy, to use the default Tier 2 model for the complete time series, would result in a less accurate monitoring of these commitments.

The Swedish method estimates the emissions separately from all four stages of the manure handling; stable ventilation, storage, manure application and grazing. The emission factors that describe the share of ammonia lost during the different stages of the manure handling are developed by the Swedish University of Agricultural Sciences and the Swedish Institute of Agricultural and Environmental Engineering (JTI)\textsuperscript{191}. The main difference between the country specific model and the Tier 2 model is that the former takes more variables into account when estimating the emissions from storage and application of manure. For example, if the manure is stored with or without a roof, and also type of roof, or application method as well as timespan between spreading and mulching.

The calculations are carried out as:
\[
\begin{align*}
\text{NH}_3N_{\text{ventilation}} &= D \times N \times P \times EF_1 \\
\text{NH}_3N_{\text{storage}} &= D \times N \times P \times (1 - EF_1) \times EF_2 \\
\text{NH}_3N_{\text{application}} &= D \times N \times (1 - EF_1) \times (1 - EF_2) \times EF_3 \\
\text{NH}_3N_{\text{grazing}} &= D \times N \times (1 - P) \times EF_4
\end{align*}
\]

Where, \(D\) = number of animal, \(N\) = yearly production of nitrogen per type of animal, \(P\) = stable period, \(EF_1\) = nitrogen losses through ventilation (fraction of total nitrogen content), \(EF_2\) = nitrogen losses during storage (fraction of total nitrogen content), \(EF_3\) = nitrogen losses during application of animal manure (fraction of ammonium nitrogen content) and \(EF_4\) = nitrogen losses during grazing (fraction of

\textsuperscript{189} Swedish Environmental Protection Agency 1997
\textsuperscript{190} Swedish Institute of Agricultural and Environmental Engineering (JTI) 2002
\textsuperscript{191} Swedish Institute of Agricultural and Environmental Engineering (JTI) 2002
total nitrogen content). In table 5-8 and 5-11 to 5-13 the emission factors used in the calculations are presented.

The ammonia emissions per animal from 3B manure management are then calculated as:

$$NH_3 = (NH_3N_{\text{Ventilation}} + NH_3N_{\text{Storage}}) \times \frac{17}{14}$$

Ammonia emissions from application of manure and grazing are calculated in this model but instead reported under 3D, crop production and agricultural soils. Concerning grazing, the length of the grazing periods for cattle are from the field investigation among farmers, while for horses, sheep and goats the grazing period is fixed to 6 months and to 12 months for reindeers.

Table 5-8. Emission factors (EF1) used to calculate emissions of ammonia from stable ventilation (% of total N) in both the Tier 2 and the country specific model.

<table>
<thead>
<tr>
<th>Animal category</th>
<th>EF solid manure</th>
<th>EF liquid manure</th>
<th>EF deep litter</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cattle</td>
<td>4%</td>
<td>4%</td>
<td>20%</td>
</tr>
<tr>
<td>Swine</td>
<td>10%</td>
<td>14%</td>
<td>25%</td>
</tr>
<tr>
<td>Sheep</td>
<td>4%</td>
<td>-</td>
<td>15%</td>
</tr>
<tr>
<td>Goats</td>
<td>4%</td>
<td>-</td>
<td>15%</td>
</tr>
<tr>
<td>Horses</td>
<td>4%</td>
<td>-</td>
<td>15%</td>
</tr>
<tr>
<td>Laying hens</td>
<td>10%</td>
<td>10%</td>
<td>35%</td>
</tr>
<tr>
<td>Chickens</td>
<td>10%</td>
<td>10%</td>
<td>20%</td>
</tr>
<tr>
<td>Slaughter Chickens</td>
<td>-</td>
<td>-</td>
<td>10%</td>
</tr>
<tr>
<td>Turkeys</td>
<td>10%</td>
<td>10%</td>
<td>35%</td>
</tr>
<tr>
<td>Fur-bearing animals</td>
<td>10%</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

Table 5-9. Emission factors used to calculate emissions of ammonia from manure storage (% of total N) in the Tier 2 model.

<table>
<thead>
<tr>
<th>Animal category</th>
<th>EF solid manure</th>
<th>EF liquid manure</th>
<th>EF deep litter</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dairy Cows</td>
<td>18%</td>
<td>3%</td>
<td>30%</td>
</tr>
<tr>
<td>Suckler cows</td>
<td>17%</td>
<td>3%</td>
<td>30%</td>
</tr>
<tr>
<td>Heifers</td>
<td>18%</td>
<td>3%</td>
<td>30%</td>
</tr>
<tr>
<td>Bulls and steers</td>
<td>17%</td>
<td>3%</td>
<td>30%</td>
</tr>
<tr>
<td>Calves</td>
<td>18%</td>
<td>3%</td>
<td>30%</td>
</tr>
<tr>
<td>Sows</td>
<td>18%</td>
<td>4%</td>
<td>30%</td>
</tr>
<tr>
<td>Boars</td>
<td>18%</td>
<td>4%</td>
<td>30%</td>
</tr>
<tr>
<td>Pigs for meat production</td>
<td>18%</td>
<td>4%</td>
<td>30%</td>
</tr>
<tr>
<td>Piglets</td>
<td>18%</td>
<td>4%</td>
<td>30%</td>
</tr>
<tr>
<td>Sheep</td>
<td>25%</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Goats</td>
<td>25%</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Horses</td>
<td>25%</td>
<td>-</td>
<td>33%</td>
</tr>
<tr>
<td>Laying hens</td>
<td>12%</td>
<td>4%</td>
<td>20%</td>
</tr>
<tr>
<td>Turkeys</td>
<td>20%</td>
<td>-</td>
<td>20%</td>
</tr>
<tr>
<td>Chickens</td>
<td>12%</td>
<td>4%</td>
<td>20%</td>
</tr>
<tr>
<td>Slaughter Chickens</td>
<td>5%</td>
<td>-</td>
<td>5%</td>
</tr>
<tr>
<td>Fur-bearing animals</td>
<td>30%</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Reindeer*</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>
Table 5-10. Nitrogen excretion, proportion of TAN and emission factors (% of TAN) used for ammonia emissions from application of manure in the Tier 2 model.

<table>
<thead>
<tr>
<th>Animal groups</th>
<th>Nitrogen kg/year/head</th>
<th>TAN in liquid manure</th>
<th>TAN in solid manure</th>
<th>TAN in deep litter</th>
<th>EF for application of liquid manure</th>
<th>EF for application of solid manure</th>
<th>EF for application of deep litter</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dairy Cows* (Milk production 6,000 kg/yr)</td>
<td>97</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Dairy Cows* (Milk production 8,000 kg/yr)</td>
<td>117</td>
<td>60%</td>
<td>51%</td>
<td>10%</td>
<td>33%</td>
<td>38%</td>
<td>41%</td>
</tr>
<tr>
<td>Dairy Cows* (Milk production 10,000 kg/yr)</td>
<td>137</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Suckler cows</td>
<td>63</td>
<td>60%</td>
<td>52%</td>
<td>10%</td>
<td>32%</td>
<td>39%</td>
<td>46%</td>
</tr>
<tr>
<td>Heifers</td>
<td>47</td>
<td>60%</td>
<td>51%</td>
<td>10%</td>
<td>32%</td>
<td>38%</td>
<td>47%</td>
</tr>
<tr>
<td>Bulls and steers</td>
<td>47</td>
<td>60%</td>
<td>52%</td>
<td>10%</td>
<td>32%</td>
<td>39%</td>
<td>46%</td>
</tr>
<tr>
<td>Calves</td>
<td>28</td>
<td>60%</td>
<td>51%</td>
<td>10%</td>
<td>33%</td>
<td>39%</td>
<td>47%</td>
</tr>
<tr>
<td>Sows</td>
<td>18.5</td>
<td>70%</td>
<td>51%</td>
<td>10%</td>
<td>30%</td>
<td>35%</td>
<td>48%</td>
</tr>
<tr>
<td>Boars</td>
<td>13</td>
<td>70%</td>
<td>51%</td>
<td>10%</td>
<td>29%</td>
<td>34%</td>
<td>46%</td>
</tr>
<tr>
<td>Pigs for meat production</td>
<td>9.5</td>
<td>70%</td>
<td>51%</td>
<td>10%</td>
<td>29%</td>
<td>35%</td>
<td>44%</td>
</tr>
<tr>
<td>Piglets</td>
<td>1.2</td>
<td>70%</td>
<td>51%</td>
<td>10%</td>
<td>29%</td>
<td>35%</td>
<td>48%</td>
</tr>
<tr>
<td>Sheep</td>
<td>14</td>
<td>25%</td>
<td>25%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Goats</td>
<td>13</td>
<td>25%</td>
<td>25%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Horses</td>
<td>48</td>
<td>24%</td>
<td>25%</td>
<td>10%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Laying hens</td>
<td>0.60</td>
<td>75%</td>
<td>60%</td>
<td>40%</td>
<td>29%</td>
<td>40%</td>
<td>56%</td>
</tr>
<tr>
<td>Turkeys</td>
<td>0.69</td>
<td>40%</td>
<td>40%</td>
<td>40%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chickens</td>
<td>0.22</td>
<td>75%</td>
<td>60%</td>
<td>40%</td>
<td>30%</td>
<td>41%</td>
<td>56%</td>
</tr>
<tr>
<td>Slaughter Chickens</td>
<td>0.29</td>
<td>40%</td>
<td>40%</td>
<td>40%</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fur-bearing animals</td>
<td>4.59</td>
<td>100%</td>
<td>100%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Reindeer*</td>
<td>10</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*For dairy cows the nitrogen excretion is estimated from milk production. **Due to a more intense swine production the nitrogen production for sows and pigs for meat production was updated in 2002. For the years prior to 2002 the values 18.5 and 9.5 kg were used.
Table 5-11. Emission factors (EF\textsubscript{2}) used to calculate emissions of ammonia from storage (% of total N) in the country specific model.

<table>
<thead>
<tr>
<th>Type of manure, handling</th>
<th>Cattle</th>
<th>Swine</th>
<th>Sheep/goats/horses</th>
<th>Laying hens/chicken</th>
<th>Slaughter chicken</th>
<th>Turkeys</th>
<th>Fur-bearing animals</th>
</tr>
</thead>
<tbody>
<tr>
<td>Liquid manure, uncovered</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from underneath</td>
<td>6%</td>
<td>8%</td>
<td>8%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from above</td>
<td>7%</td>
<td>9%</td>
<td>9%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Liquid manure, covered</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from underneath:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Roof</td>
<td>1%</td>
<td>1%</td>
<td>1%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>floating crust</td>
<td>3%</td>
<td>4%</td>
<td>4%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>2%</td>
<td>2%</td>
<td>2%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from above:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Roof</td>
<td>1%</td>
<td>1%</td>
<td>1%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>floating crust</td>
<td>4%</td>
<td>5%</td>
<td>5%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>3%</td>
<td>3%</td>
<td>3%</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Urine, uncovered</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from underneath</td>
<td>37%</td>
<td>37%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from above</td>
<td>40%</td>
<td>40%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Urine, with cover</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from underneath:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Roof</td>
<td>5%</td>
<td>5%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>floating crust</td>
<td>17%</td>
<td>17%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>10%</td>
<td>10%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Filled from above:</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Roof</td>
<td>5%</td>
<td>5%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>floating crust</td>
<td>20%</td>
<td>20%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>12%</td>
<td>12%</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Solid manure</td>
<td>20%</td>
<td>20%</td>
<td>25%</td>
<td>12%</td>
<td></td>
<td></td>
<td>30%</td>
</tr>
<tr>
<td>Deep litter manure</td>
<td>30%</td>
<td>30%</td>
<td>33%</td>
<td>20%</td>
<td>5%</td>
<td>20%</td>
<td></td>
</tr>
</tbody>
</table>
Table 5-12. Emission factors (EFs) used to calculate emissions of ammonia from spreading (% of TAN) in the country specific model.

<table>
<thead>
<tr>
<th>Tillage timing</th>
<th>Spreading strategy</th>
<th>Season</th>
<th>EF solid manure or deep litter</th>
<th>EF urine</th>
<th>EF liquid manure or slurry</th>
</tr>
</thead>
<tbody>
<tr>
<td>Immediately</td>
<td>Broadcast</td>
<td>Early autumn</td>
<td>20%</td>
<td>15%</td>
<td>5%</td>
</tr>
<tr>
<td>Immediately</td>
<td>Trailing hoses</td>
<td>Early autumn</td>
<td>10%</td>
<td>3%</td>
<td></td>
</tr>
<tr>
<td>Mulching within 4 h</td>
<td>Broadcast</td>
<td>Early autumn</td>
<td>35%</td>
<td>23%</td>
<td>18%</td>
</tr>
<tr>
<td>Mulching within 4 h</td>
<td>Trailing hoses</td>
<td>Early autumn</td>
<td>18%</td>
<td>9%</td>
<td></td>
</tr>
<tr>
<td>Mulching within 5-24 h</td>
<td>Broadcast</td>
<td>Early autumn</td>
<td>50%</td>
<td>30%</td>
<td>30%</td>
</tr>
<tr>
<td>Mulching within 5-24 h</td>
<td>Trailing hoses</td>
<td>Early autumn</td>
<td>25%</td>
<td>15%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on unseeded ground</td>
<td>Broadcast</td>
<td>Early autumn</td>
<td>70%</td>
<td>45%</td>
<td>70%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on unseeded ground</td>
<td>Trailing hoses</td>
<td>Early autumn</td>
<td>30%</td>
<td>40%</td>
<td></td>
</tr>
<tr>
<td>Immediately</td>
<td>Broadcast</td>
<td>Late autumn</td>
<td>10%</td>
<td>10%</td>
<td>5%</td>
</tr>
<tr>
<td>Immediately</td>
<td>Trailing hoses</td>
<td>Late autumn</td>
<td>4%</td>
<td>3%</td>
<td></td>
</tr>
<tr>
<td>Mulching within 4 h</td>
<td>Broadcast</td>
<td>Late autumn</td>
<td>15%</td>
<td>15%</td>
<td>8%</td>
</tr>
<tr>
<td>Mulching within 4 h</td>
<td>Trailing hoses</td>
<td>Late autumn</td>
<td>11%</td>
<td>4%</td>
<td></td>
</tr>
<tr>
<td>Mulching within 5-24 h</td>
<td>Broadcast</td>
<td>Late autumn</td>
<td>20%</td>
<td>20%</td>
<td>10%</td>
</tr>
<tr>
<td>Mulching within 5-24 h</td>
<td>Trailing hoses</td>
<td>Late autumn</td>
<td>18%</td>
<td>5%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on unseeded ground</td>
<td>Broadcast</td>
<td>Late autumn</td>
<td>30%</td>
<td>25%</td>
<td>30%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on unseeded ground</td>
<td>Trailing hoses</td>
<td>Late autumn</td>
<td>25%</td>
<td>15%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on unseeded ground</td>
<td>Broadcast</td>
<td>Winter</td>
<td>20%</td>
<td>40%</td>
<td>30%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on unseeded ground</td>
<td>Trailing hoses</td>
<td>Winter</td>
<td>30%</td>
<td>20%</td>
<td></td>
</tr>
<tr>
<td>Immediately</td>
<td>Broadcast</td>
<td>Spring</td>
<td>15%</td>
<td>8%</td>
<td>10%</td>
</tr>
<tr>
<td>Immediately</td>
<td>Trailing hoses</td>
<td>Spring</td>
<td>7%</td>
<td>5%</td>
<td></td>
</tr>
<tr>
<td>Mulching within 4 h</td>
<td>Broadcast</td>
<td>Spring</td>
<td>33%</td>
<td>14%</td>
<td>15%</td>
</tr>
<tr>
<td>Mulching within 4 h</td>
<td>Trailing hoses</td>
<td>Spring</td>
<td>14%</td>
<td>8%</td>
<td></td>
</tr>
<tr>
<td>Mulching within 5-24 h</td>
<td>Broadcast</td>
<td>Spring</td>
<td>50%</td>
<td>20%</td>
<td>20%</td>
</tr>
<tr>
<td>Mulching within 5-24 h</td>
<td>Trailing hoses</td>
<td>Spring</td>
<td>20%</td>
<td>10%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on pasture</td>
<td>Broadcast</td>
<td>Spring</td>
<td>70%</td>
<td>35%</td>
<td>40%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on pasture</td>
<td>Trailing hoses</td>
<td>Spring</td>
<td>25%</td>
<td>30%</td>
<td></td>
</tr>
<tr>
<td>Immediately</td>
<td>Shallow injection</td>
<td>Spring</td>
<td>8%</td>
<td>15%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on grain</td>
<td>Broadcast</td>
<td>Spring</td>
<td>70%</td>
<td>11%</td>
<td>20%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on grain</td>
<td>Trailing hoses</td>
<td>Spring</td>
<td>10%</td>
<td>15%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on pasture</td>
<td>Broadcast</td>
<td>Summer</td>
<td>90%</td>
<td>60%</td>
<td>70%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on pasture</td>
<td>Trailing hoses</td>
<td>Summer</td>
<td>40%</td>
<td>50%</td>
<td></td>
</tr>
<tr>
<td>Immediately</td>
<td>Shallow injection</td>
<td>Summer</td>
<td>15%</td>
<td>30%</td>
<td></td>
</tr>
<tr>
<td>After 24 hours or no mulching on grain</td>
<td>Broadcast</td>
<td>Summer</td>
<td>90%</td>
<td>10%</td>
<td>20%</td>
</tr>
<tr>
<td>After 24 hours or no mulching on grain</td>
<td>Trailing hoses</td>
<td>Summer</td>
<td>10%</td>
<td>7%</td>
<td></td>
</tr>
</tbody>
</table>
Table 5-13. Emission factors (EFs) used to calculate emissions of ammonia from grazing animals (% of total N) in both the Tier 2 and the country specific model.

<table>
<thead>
<tr>
<th>Animal category</th>
<th>EF Grazing</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cattle</td>
<td>8%</td>
</tr>
<tr>
<td>Horses</td>
<td>8%</td>
</tr>
<tr>
<td>Sheep</td>
<td>4%</td>
</tr>
<tr>
<td>Goats</td>
<td>4%</td>
</tr>
<tr>
<td>Reindeer</td>
<td>4%</td>
</tr>
</tbody>
</table>

5.2.1.2.4 Non-methane volatile organic compounds (NMVOCs)

The emissions of NMVOC from manure management are estimated with the Tier 2 methods described in the EMEP/EEA guidebook. The emissions are calculated as the sum of six different sources:

- from feeding of silage
- from silage stores
- from housing (feeding beside silage)
- from outdoor manure stores
- from manure application (reported in 3D)
- from grazing animals (reported in 3D)

The calculation methods differ slightly between cattle and other animals. For cattle the methodology is based on feed intake, but on excreted volatile substance for the other animals. That is, the factor $MJ_i$ is replaced with $kg VS_i$ (kg volatile solids excreted).

The emissions from cattle for the different subcategories are calculated as:

Feeding of silage:
\[
E_{NMVOC,\text{silage_feeding}_i} = AAP_i \times MJ_i \times x_{\text{house}_i} \times EF_{NMVOC,\text{silage_feeding}_i} \times \text{Frac}_{\text{silage}_i}
\]

Silage stores:
\[
E_{NMVOC,\text{silage_store}_i} = E_{NMVOC,\text{silage_feeding}_i} \times \text{Frac}_{\text{silage_store}}
\]

Housing (feeding beside silage):
\[
E_{NMVOC,\text{house}_i} = AAP_i \times MJ_i \times x_{\text{house}_i} \times EF_{NMVOC,\text{house}_i}
\]

Outdoor manure stores:
\[
E_{NMVOC,\text{manure_store}_i} = E_{NMVOC,\text{house}_i} \times (\text{ENH}_{3,\text{storage}_i} / \text{ENH}_{3,\text{house}_i})
\]

And the total emission of NMVOC reported in 3B, manure management is:
\[
\sum_i [E_{NMVOC,\text{silage_feeding}_i} + E_{NMVOC,\text{silage_store}_i} + E_{NMVOC,\text{house}_i} + E_{NMVOC,\text{manure_store}_i}]
\]
where;
AAP_{i} is the annual average population of animal $i$, MJ_{i} is the annual gross feed intake for animal $i$. The estimated gross feed is country specific and the same ones as used in the reporting of greenhouse gases to the UNFCCC. $X_{\text{house},i}$ is country specific data on the share of time an animal $i$ spends in the animal house in a year. The value is corrected for that part of the manure is deposited in the stables also during the grazing period when dairy cows return to the stables for milking. Frac-silage_store is the share of the emission from the silage store compared to the emission from the feeding table in the barn. The default tentative value of 0.25 from the guidebook is used. The emission factors used are from the EMEP/EEA guidebook 2013. ENH_{i,\text{storage}}, ENH_{i,\text{house}} and ENH_{i,\text{appl}} are the emissions of ammonia from storage of manure, stables and application of manure for animal category $i$, respectively. The values are taken from the Swedish ammonia inventory.

When no country specific emission factors on emissions of NMVOC exists in Sweden the default emission factors from the EMEP/EEA guidebook 2013 are used. See table 5-14 for a list of the factors used in the calculations of NMVOC from manure management.

Table 5-14. Parameters and emission factors used for estimating emissions of NMVOC from manure management.

<table>
<thead>
<tr>
<th>Source</th>
<th>Fraction silage feeding</th>
<th>Silage feeding (kg NMVOC/MJ feed intake)</th>
<th>Housing (kg NMVOC/MJ feed intake)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dairy cows</td>
<td>0.76</td>
<td>0.0002002</td>
<td>0.0000353</td>
</tr>
<tr>
<td>Suckler cows</td>
<td>0.59</td>
<td>0.0002002</td>
<td>0.0000353</td>
</tr>
<tr>
<td>Bulls and steers</td>
<td>0.63</td>
<td>0.0002002</td>
<td>0.0000353</td>
</tr>
<tr>
<td>Heifers</td>
<td>0.81</td>
<td>0.0002002</td>
<td>0.0000353</td>
</tr>
<tr>
<td>Calves</td>
<td>-</td>
<td>0.0002002</td>
<td>0.0000353</td>
</tr>
<tr>
<td>Sows and boars</td>
<td>-</td>
<td>-</td>
<td>0.007042</td>
</tr>
<tr>
<td>Pigs for meat production and piglets</td>
<td>-</td>
<td>-</td>
<td>0.001703</td>
</tr>
<tr>
<td>Sheep</td>
<td>0.68</td>
<td>0.01076</td>
<td>0.001614</td>
</tr>
<tr>
<td>Horses</td>
<td>1</td>
<td>0.01076</td>
<td>0.001614</td>
</tr>
<tr>
<td>Goats</td>
<td>0.67</td>
<td>0.01076</td>
<td>0.001614</td>
</tr>
<tr>
<td>Laying hens and chickens</td>
<td>-</td>
<td>-</td>
<td>0.005684</td>
</tr>
<tr>
<td>Slaughter chickens</td>
<td>-</td>
<td>-</td>
<td>0.009147</td>
</tr>
<tr>
<td>Turkeys</td>
<td>-</td>
<td>-</td>
<td>0.005684</td>
</tr>
<tr>
<td>Reindeers</td>
<td>-</td>
<td>-</td>
<td>0.001614</td>
</tr>
<tr>
<td>Fur-bearing animals</td>
<td>-</td>
<td>-</td>
<td>0.005684</td>
</tr>
</tbody>
</table>
5.2.1.2.5 Particulate matter

In those cases a Tier 2 methodology is described in the Guidebook it is used to estimate the emissions from animals as well as from crops. The only cases where Tier 1 methodologies are used are for the animal categories where no Tier 2 methodology exists. That is, emissions from sheep, horses, goats, chickens, turkeys and fur-bearing animals. The emission factors used are the default ones from the EMEP/EEA guidebook 2013 (see table 5-15 to 5-17).

Table 5-15. Tier 2 emission factors used to estimate the emissions from cattle and swine (kg/AAP/year).

<table>
<thead>
<tr>
<th>Substance</th>
<th>Manure</th>
<th>Dairy cows</th>
<th>Beef cows</th>
<th>Steers and bulls</th>
<th>Heifers</th>
<th>Calves</th>
<th>Pigs for meat production</th>
<th>Piglets</th>
<th>Sows and boars</th>
</tr>
</thead>
<tbody>
<tr>
<td>TSP</td>
<td>Slurry</td>
<td>1.81</td>
<td>0.69</td>
<td>0.69</td>
<td>0.34</td>
<td>0.70</td>
<td>0.36</td>
<td>1.36</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Solid</td>
<td>0.94</td>
<td>0.52</td>
<td>0.52</td>
<td>0.35</td>
<td>0.83</td>
<td>0.00</td>
<td>1.77</td>
<td></td>
</tr>
<tr>
<td>PM_{10}</td>
<td>Slurry</td>
<td>0.83</td>
<td>0.32</td>
<td>0.32</td>
<td>0.15</td>
<td>0.31</td>
<td>0.16</td>
<td>0.61</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Solid</td>
<td>0.43</td>
<td>0.24</td>
<td>0.24</td>
<td>0.16</td>
<td>0.37</td>
<td>0.00</td>
<td>0.80</td>
<td></td>
</tr>
<tr>
<td>PM_{2.5}</td>
<td>Slurry</td>
<td>0.54</td>
<td>0.21</td>
<td>0.21</td>
<td>0.10</td>
<td>0.06</td>
<td>0.03</td>
<td>0.11</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Solid</td>
<td>0.28</td>
<td>0.16</td>
<td>0.16</td>
<td>0.10</td>
<td>0.07</td>
<td>0.00</td>
<td>0.14</td>
<td></td>
</tr>
</tbody>
</table>

Table 5-16. Tier 2 emission factors used to estimate emissions from laying hens (kg/AAP/year).

<table>
<thead>
<tr>
<th>Substance</th>
<th>Housing</th>
<th>Laying hens</th>
</tr>
</thead>
<tbody>
<tr>
<td>TSP</td>
<td>Cages</td>
<td>0.025</td>
</tr>
<tr>
<td></td>
<td>Percheries</td>
<td>0.119</td>
</tr>
<tr>
<td>PM_{10}</td>
<td>Cages</td>
<td>0.025</td>
</tr>
<tr>
<td></td>
<td>Percheries</td>
<td>0.119</td>
</tr>
<tr>
<td>PM_{2.5}</td>
<td>Cages</td>
<td>0.003</td>
</tr>
<tr>
<td></td>
<td>Percheries</td>
<td>0.023</td>
</tr>
</tbody>
</table>

Table 5-17. Tier 1 emission factors used to estimate emissions from other animals (kg/AAP/year).

<table>
<thead>
<tr>
<th>Substance</th>
<th>Sheep</th>
<th>Horses</th>
<th>Goats</th>
<th>Chickens</th>
<th>Turkeys</th>
<th>Fur-bearing animals</th>
</tr>
</thead>
<tbody>
<tr>
<td>TSP</td>
<td>0.139</td>
<td>0.48</td>
<td>0.139</td>
<td>0.069</td>
<td>0.52</td>
<td>0.018</td>
</tr>
<tr>
<td>PM_{10}</td>
<td>0.056</td>
<td>0.22</td>
<td>0.056</td>
<td>0.069</td>
<td>0.52</td>
<td>0.0081</td>
</tr>
<tr>
<td>PM_{2.5}</td>
<td>0.017</td>
<td>0.14</td>
<td>0.017</td>
<td>0.009</td>
<td>0.07</td>
<td>0.0042</td>
</tr>
</tbody>
</table>

5.2.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

As described above it was not possible to apply the Swedish country specific ammonia model for the years prior to 2005. To minimize the time series break between the default Tier 2 model (1990-2004) and the country specific model (2005 and onwards) we have used country specific emission factors in the Tier 2 model. These emission factors are developed according to the country specific model by weighing various detailed variables to the aggregates that are used by the Tier 2 model. For example the Swedish model use much more disaggregated emissions
factors for application of animal manure (i.e. table 5-11). From these we have calculated weighted country specific factors that fit the default Tier 2 model (i.e. table 5-9).

Between 1995 and 1996 there is 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. Due to more intense swine production, the nitrogen production for sows and pigs for meat production were updated in 2002. Since no estimate on the number of horses exists before 2004, the value for 2004 is used for all preceding years.

The calculations are to a large degree based on information from farmers given in Statistics Sweden’s field investigation and these results are afflicted with standard errors. Hence all results must be considered with caution regarding the uncertainty in the input data. The emission factors are of course also encumbered with significant uncertainties. The emission factor uncertainties are likely more substantial than the activity data uncertainties. We estimate the uncertainty interval for the activity data for a specific emission category in 3B is some 20%. The uncertainty intervals for the different emission factor are estimated to: NH$_3$ 50%, NMVOC 200%, NO$_x$ 80%, PM 150%-200%.

5.2.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
Annual increase or decrease is verified for the whole time series for all sub sources to decide that all annual changes are reasonable. We compare the times series for the emission with the time series for the activity data to confirm that are in agreement. We conduct regular meetings with the different authorities that provide activity data to the inventory to ensure that the quality of the data is of satisfactory quality and that they in turn use appropriate QC methods.

5.2.1.5 SOURCE-SPECIFIC RECALCULATIONS
Only minor changes have with small effects on emissions have been done. i) The number of decimals in the length of stable periods in the calculations are changed for cattle from 14 to three decimals, a more realistic length. ii) The number of turkeys have been updated for 2012, 2013 and 2014 which have a slight but observed effect on all emissions in 3.B Poltry. iii) Updated amount of VS for sow which effect the NMVOC. iv) Minor changes in calculation method effecting NO$_x$ for swine.

5.2.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
Category-specific improvements will be decided after the finalization of the submission as part of the national QA/QC plan.
5.3 Crop production and agricultural soils, NFR3D

5.3.1.1 SOURCE CATEGORY DESCRIPTION

From the subsector, crop production and agricultural soils, Sweden report emissions of ammonia, NMVOC, NO\textsubscript{X} and particulate matters. The most significant source of ammonia is emissions from application of animal manure. Other sources of ammonia are the use of inorganic fertilisers, grazing animals, applications of sewage sludge and use of other organic fertilisers. The same sources also give rise to emissions of nitric oxide. Emissions of NMVOC from crop can arise to attract pollinating insects, eliminate waste products or as a means of losing surplus energy. Factors that can influence the emissions of NMVOC include temperature and light intensity, plant growth stage, water stress, air pollution and senescence. The main source from crop production in Sweden is emissions from ley. Particulate matters are emitted during production of crops, and the main sources are soil cultivation and crop harvesting. These emissions originate from operation of tractors and other machinery and are thought to consist of a mixture of organic fragments from the crop and soil mineral and organic matter.

Table 5-18. Summary of key source analysis, NFR3D, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016 Level</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>3Da1 Inorganic N fertilizers</td>
<td>NO\textsubscript{X}</td>
<td>NO\textsubscript{X}</td>
</tr>
<tr>
<td></td>
<td>NH\textsubscript{3}</td>
<td>NH\textsubscript{3}</td>
</tr>
<tr>
<td>3Da2a Animal manure applied to soils</td>
<td>NO\textsubscript{X}</td>
<td>NO\textsubscript{X}</td>
</tr>
<tr>
<td></td>
<td>NMVOC</td>
<td>NMVOC</td>
</tr>
<tr>
<td></td>
<td>NH\textsubscript{3}</td>
<td>NH\textsubscript{3}</td>
</tr>
<tr>
<td>3Da2b Sewage sludge applied to soils</td>
<td>NH\textsubscript{3}</td>
<td>NH\textsubscript{3}</td>
</tr>
<tr>
<td>3Da2c Other organic fertilizers applied to soils</td>
<td>NH\textsubscript{3}</td>
<td>NH\textsubscript{3}</td>
</tr>
<tr>
<td>3Da3 Urine and dung deposited by grazing animals</td>
<td>NO\textsubscript{X}</td>
<td>NO\textsubscript{X}</td>
</tr>
<tr>
<td></td>
<td>NH\textsubscript{3}</td>
<td>NH\textsubscript{3}</td>
</tr>
<tr>
<td>3Dc Farm-level agricultural operations</td>
<td>PM\textsubscript{2.5}</td>
<td>PM\textsubscript{2.5}</td>
</tr>
<tr>
<td></td>
<td>PM\textsubscript{10}</td>
<td>PM\textsubscript{10}</td>
</tr>
<tr>
<td>3De Cultivated crops</td>
<td>NMVOC</td>
<td>NMVOC</td>
</tr>
</tbody>
</table>
5.3.1.2  METHODOLOGICAL ISSUES

5.3.1.2.1  Activity data

To estimate the emissions of ammonia and nitric oxide, data on applied nitrogen is needed for a number of different sources. To ensure consistency with the greenhouse gases inventory the same data is used in both inventories. The data on total nitrogen content in different types of mineral fertilisers is from the Swedish Board of Agriculture’s sales statistics in Sweden (table 5-19). Application of sludge and nitrogen content in sludge is collected intermittently by Statistics Sweden and the Swedish EPA from sewage treatment plants. The nitrogen content in other organic fertilisers applied to soils is estimated from Statistics Sweden’s survey on “Use of fertilisers and animal manure in agriculture”. Nitrogen content in animal manure is estimated as the amount remaining after ventilation and storage losses calculated above in category 3B.

The formula used to calculate N content in animal manure applied to soils ($F_{AM}$) is:

$$F_{AM} = \sum T \left( \frac{N_T \times \text{Nex}_T \times (1 - \text{Frac}_{\text{LossMS}}) \times (1 - \text{Frac}_{\text{PRP}})}{T} \right)$$

Where $N_T$ is the number of heads of livestock in category $T$ in the country, $\text{Nex}_T$ is the annual average excretion of N per head of category $T$ in the country, $\text{Frac}_{\text{LossMS}}$ is the amount of N lost before application, $\text{Frac}_{\text{PRP}}$ is the fraction of the nitrogen in pasture, range and paddock manure. The amount of nitrogen in grazing manure is also calculated above under 3B.

To estimate emissions of particulate matter and NMVOC from crop production, statistics on crop areas is needed. This is produced by the Swedish Board of Agriculture192.

5.3.1.2.2  Emissions of Ammonia

For a methodological description of the emissions from application of manure and grazing animals, see paragraph 5.2.3.1 and 5.2.3.2. To estimate the ammonia emissions from mineral fertilisers we have used the default Tier 2 methodology from the EMEP/EEA guidebook 2016. To estimate the emissions from the relatively small sources, sewage sludge and other organic fertilisers, we have used the average nitrogen loss from application of animal manure as an approximation.

5.3.1.2.3  Emissions of Non-methane volatile organic compounds (NMVOCs)

The estimated value for the emissions of NMVOC from housing is used as the basis for estimating the emissions from manure application. See above in paragraph 5.2.3.3, for a description of how $E_{\text{NMVOC,house}}$ is calculated.

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192 Swedish Board of Agriculture, JO 10-series
Emissions from animal manure applied to soils (3Da2a):

\[ E_{\text{NMVOC, appl}_i} = E_{\text{NMVOC, house}_i} \times \left( \frac{\text{ENH}_3,\text{appl}_i}{\text{ENH}_3,\text{house}_i} \right) \]

For application of sewage sludge no methodology is described in the guidebook. As an approximation of the NMVOC emissions we have used the same emission factor as for manure from pasture, range and paddock and assumed the same volatile solid content as swine.

Emissions from sewage sludge applied to soils (3Da2b):

\[ E_{\text{NMVOC, Sludge}} = N_{\text{sludge}} \times \text{Frac}_{\text{VS, swine}} \times EF_{\text{NMVOC, graz}} \]

Emissions from urine and dung deposited by grazing animals (3Da3):

\[ E_{\text{NMVOC, graz}_i} = A A P_i \times M J_i \times (\text{or kg VS}_j) \times (1 - x_{\text{house}_i}) \times EF_{\text{NMVOC, graz}_i} \]

To estimate the emissions from crop production we have used the Tier 1 method, because no Tier 2 method is currently described in the guidebook.

The emissions are calculated as:

\[ E_{\text{NMVOC, crops}} = A R_{\text{area}} \times EF_{\text{NMVOC, crop}} \]

where;

- \( A R_{\text{area}} \) is the area covered with crops, and the emissions factor is amount of NMVOC per hectare (0.86 kg/ha/year).
Table 5-19. Parameters and emission factors used to estimate emissions of NMVOC from manure management.

<table>
<thead>
<tr>
<th>Source</th>
<th>Grazing (kg NMVOC/ MJ feed intake)</th>
<th>Grazing (kg NMVOC/kg VS excreted)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dairy cows</td>
<td>0.0000069</td>
<td></td>
</tr>
<tr>
<td>Beef cows</td>
<td>0.0000069</td>
<td></td>
</tr>
<tr>
<td>Steers and bulls</td>
<td>0.0000069</td>
<td></td>
</tr>
<tr>
<td>Heifers</td>
<td>0.0000069</td>
<td></td>
</tr>
<tr>
<td>Calves</td>
<td>0.0000069</td>
<td></td>
</tr>
<tr>
<td>Sows and boars</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Pigs for meat production and piglets</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Sheep</td>
<td>0.00002349</td>
<td></td>
</tr>
<tr>
<td>Horses</td>
<td>0.00002349</td>
<td></td>
</tr>
<tr>
<td>Goats</td>
<td>0.00002349</td>
<td></td>
</tr>
<tr>
<td>Laying hens and chickens</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Slaughter chickens</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Turkeys</td>
<td>-</td>
<td></td>
</tr>
<tr>
<td>Reindeers</td>
<td>0.00002349</td>
<td></td>
</tr>
<tr>
<td>Fur-bearing animals</td>
<td>-</td>
<td></td>
</tr>
</tbody>
</table>

5.3.1.2.4 Nitrogen oxides

The estimate of NOX emissions from crop production are based on the default Tier 1 methodology when no Tier 2 methodology yet exists in the guidebook.

To estimate these emissions the annual sum of all nitrogen applied to soil is required. That is, the sum of all applied nitrogen in: inorganic fertilisers, animal manure, sewage sludge, other organic fertilisers and excreta from grazing animals (table 5-20). This value is multiplied with the default Tier 1 emission factor of 0.026 kg of NO per kg of fertiliser-N applied. Finally the amount of NO is translated to amount of NO2 to be reported in the NFR-tables.

Table 5-20. Amount of nitrogen applied from different sources (t/year).

<table>
<thead>
<tr>
<th>Year</th>
<th>N in inorganic fertilisers (t)</th>
<th>N in applied organic fertilisers (t)</th>
<th>N in sewage sludge (t)</th>
<th>N in other organic fertilisers (t)</th>
<th>N deposited by grazing animals</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>224 500</td>
<td>76 631</td>
<td>1 180</td>
<td>1 700</td>
<td>44 245</td>
</tr>
<tr>
<td>1995</td>
<td>198 300</td>
<td>79 794</td>
<td>2 304</td>
<td>1 700</td>
<td>46 663</td>
</tr>
<tr>
<td>2000</td>
<td>189 400</td>
<td>75 587</td>
<td>1 758</td>
<td>1 800</td>
<td>46 724</td>
</tr>
<tr>
<td>2005</td>
<td>161 568</td>
<td>74 029</td>
<td>1 053</td>
<td>1 743</td>
<td>47 475</td>
</tr>
<tr>
<td>2010</td>
<td>168 000</td>
<td>72 703</td>
<td>2 224</td>
<td>2 712</td>
<td>45 243</td>
</tr>
<tr>
<td>2014</td>
<td>181 090</td>
<td>73 264</td>
<td>2 374</td>
<td>4 386</td>
<td>43 384</td>
</tr>
<tr>
<td>2015</td>
<td>190 200</td>
<td>73 381</td>
<td>2 802</td>
<td>4 386</td>
<td>43 187</td>
</tr>
<tr>
<td>2016</td>
<td>186 000</td>
<td>73 413</td>
<td>2 802</td>
<td>5 358</td>
<td>43 959</td>
</tr>
</tbody>
</table>
5.3.1.3 PARTICULATE MATTER

Emissions from agricultural crop operations are estimated based on cultivated area for the different crops. Statistics on crop areas and data on agricultural crop operations for the different crops are used according to the default Tier 2 model. The frequency of soil cultivation, harvesting, cleaning and drying has been set to one time per year for all crops except for grass for hay making. For this category, soil cultivation is assumed to take place every third year and harvest on average 2.4 times per year. Average number of harvests is estimated based on data from the Swedish survey on nitrogen and phosphorus balances for agricultural land.

Table 5-21. Tier 2 emission factors used to estimate emissions from crop production (kg/ha/year).

<table>
<thead>
<tr>
<th>Substance</th>
<th>Crop operation</th>
<th>Winter wheat</th>
<th>Spring wheat</th>
<th>Winter rye</th>
<th>Triticale</th>
<th>Mixed grain</th>
<th>Winter barley</th>
<th>Spring barley</th>
<th>Oats</th>
<th>Pasture ground</th>
<th>Other crops</th>
<th>Ley</th>
</tr>
</thead>
<tbody>
<tr>
<td>PM$_{10}$</td>
<td>Soil cultivation</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td>0.25</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Harvesting</td>
<td>0.49</td>
<td>0.49</td>
<td>0.37</td>
<td>0.37</td>
<td>0.41</td>
<td>0.41</td>
<td>0.62</td>
<td>NA</td>
<td>NA</td>
<td>0.25</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Cleaning</td>
<td>0.19</td>
<td>0.19</td>
<td>0.16</td>
<td>0.16</td>
<td>0.16</td>
<td>0.16</td>
<td>0.25</td>
<td>NA</td>
<td>NA</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Drying</td>
<td>0.56</td>
<td>0.56</td>
<td>0.37</td>
<td>0.37</td>
<td>0.43</td>
<td>0.43</td>
<td>0.66</td>
<td>NA</td>
<td>NA</td>
<td>0</td>
<td></td>
</tr>
</tbody>
</table>

| PM$_{2.5}$ | Soil cultivation | 0.015        | 0.015        | 0.015      | 0.015     | 0.015       | 0.015         | 0.015         | 0.015| 0.015        | 0.015      |     |
|            | Harvesting      | 0.02         | 0.02         | 0.015      | 0.015     | 0.016       | 0.016         | 0.025         | NA  | NA            | 0.01       |     |
|            | Cleaning        | 0.009        | 0.009        | 0.008      | 0.008     | 0.008       | 0.008         | 0.0125        | NA  | NA            | 0          |     |
|            | Drying          | 0.168        | 0.168        | 0.111      | 0.111     | 0.129       | 0.129         | 0.198         | NA  | NA            | 0          |     |

5.3.1.4 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

There is a decrease in the amount of sold fertilisers in 2009. This is due to an over-consumption in 2008 due to a dropped tax on fertilisers. Statistics on the use of sewage sludge have been published irregularly and in different reports, and the time series for the earlier years in the time series has been created through interpolation/extrapolation and certain assumptions. Gradually the quality of the data has increased and is for the latest years of adequate quality.

We estimate the uncertainty interval for the activity data for a specific emission category in 3D is 20%-35%. Uncertainty intervals for the different emission factor are estimated to: NH$_3$ 50%, NMVOC 200%, NO$_x$ 80%-400%. PM 150%-200%.

5.3.1.5 SOURCE-SPECIFIC QA/QC AND VERIFICATION

Annual increase or decrease is verified for the whole time series for all sub sources to decide that all annual changes are reasonable. We compare the times series for the emission with the time series for the activity data to confirm that are in agreement. Every year we ask experts from the Swedish board of agriculture to conduct
expert peer reviews of the methods used. Regular meetings are held with the authorities that provide activity data to the inventory to ensure that the quality of the data is of satisfactory quality and that they in turn use appropriate QC methods.

5.3.1.6 SOURCE-SPECIFIC RECALCULATIONS
Most recalculations have smaller effect but i) the usage of EMEP/EEA air pollutant emission inventory guidebook 2016 in the emission of ammonia changes the ammonia emissions from mineral fertilizers, sewage sludg and other organic fertilizers applied to soil effected the ammonia emissions in a significant way. ii) The changes in ammonia emissions also effect the NMVOC levels as ammonia is used as a proxy in the calculations. iii) Classification of N-fertilizers is updated in the later years of the timeseries. iv) Changes in handling of slurry and an increased proportion of slurry lead to a decrease in NH3-emissions. v) N-losses as nitrogen gas (N2) during storage is included. vi) The time series of other organic fertilizers applied to soils is revised from Statistics Sweden. vii) The number of decimals in the length of stable periods in the calculations are changed for cattle from 14 to three decimals, a more realistic length witch effect the emissions from urine and dung deposited by grazing animals slightly. viii) Revised population of animals. ix) Minor change in area of oat for 2015 effect the PM2.5 from farm-level agricultural operations.

5.3.1.7 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 Waste (NFR sector 5)

6.1 Overview

Emission estimates from the waste sector include emissions from solid waste disposal on land, biological treatment of waste, waste-water handling, incineration of hazardous waste (including cremation) and various types of fires such as landfill fires, house and car fires, bonfires and open burning of garden waste. Combustion of municipal waste is accounted for in the energy sector, since it is used as fuel for energy production. Emission estimates also includes emissions from sludge spreading (mechanical dewatering of digested sludge) and pets.

6.2 Solid waste disposal on land, NFR 5A

This category includes Solid waste disposal on land (NFR 5A).

6.2.1.1 Source category description

Sweden is reporting of emissions of NMVOC, TSP, PM\textsubscript{10} and PM\textsubscript{2.5}. Emissions of NH\textsubscript{3}, Hg and CO are reported as not estimated (NE). Other emissions are reported as not applicable (NA).

6.2.1.2 Methodological issues

6.2.1.2.1 Emission factors

Tier 1 default emission factors from the EMEP/EEA Emission Inventory Guidebook 2013/2016 are used for NMVOC, TSP, PM\textsubscript{10} and PM\textsubscript{2.5}. See further information in the table 6-1 below.

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Lower</td>
<td>Upper</td>
</tr>
<tr>
<td>NMVOC</td>
<td>1.56</td>
<td>kg/Mg</td>
<td>0.5</td>
<td>3.0</td>
</tr>
<tr>
<td>TSP</td>
<td>0.463</td>
<td>g/Mg</td>
<td>0.006</td>
<td>2.21</td>
</tr>
<tr>
<td>PM\textsubscript{10}</td>
<td>0.219</td>
<td>g/Mg</td>
<td>0.003</td>
<td>1.05</td>
</tr>
<tr>
<td>PM\textsubscript{2.5}</td>
<td>0.033</td>
<td>g/Mg</td>
<td>0.0004</td>
<td>1.16</td>
</tr>
</tbody>
</table>

6.2.1.2.2 Activity data

Activity data on emissions of landfill gas is used for emissions of NMVOC. The data is calculated from emission data on methane from solid waste disposal on land, reported to UNFCCC.

For TSP, PM\textsubscript{10} and PM\textsubscript{2.5}, activity data on landfilled waste at landfills for municipal solid waste is used. The data has been compiled and published by the association Swedish Waste Management (RVF/Avfall Sverige).
Table 6-2 Activity data used for NFR 5A Solid waste disposal on land.

<table>
<thead>
<tr>
<th>Year</th>
<th>Emission of landfill gas* (m³) (AD used for NMVOC)</th>
<th>Landfilled waste at landfills for municipal solid waste* (t, wet weight) (AD used for TSP, PM₁₀ and PM₂.₅)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>381 513 038¹</td>
<td>7 000 000²</td>
</tr>
<tr>
<td>1991</td>
<td>387 325 314²</td>
<td>6 770 000</td>
</tr>
<tr>
<td>1992</td>
<td>387 440 535³</td>
<td>6 540 000</td>
</tr>
<tr>
<td>1993</td>
<td>374 396 920⁴</td>
<td>6 310 000</td>
</tr>
<tr>
<td>1994</td>
<td>358 913 227⁴</td>
<td>6 080 000</td>
</tr>
<tr>
<td>1995</td>
<td>357 976 710⁴</td>
<td>5 340 000</td>
</tr>
<tr>
<td>1996</td>
<td>354 903 980⁴</td>
<td>5 050 000</td>
</tr>
<tr>
<td>1997</td>
<td>351 430 403⁴</td>
<td>4 750 000</td>
</tr>
<tr>
<td>1998</td>
<td>345 702 109⁴</td>
<td>4 800 000</td>
</tr>
<tr>
<td>1999</td>
<td>330 900 990⁵</td>
<td>4 900 000</td>
</tr>
<tr>
<td>2000</td>
<td>320 485 283⁵</td>
<td>4 450 000</td>
</tr>
<tr>
<td>2001</td>
<td>314 665 020⁵</td>
<td>4 240 000</td>
</tr>
<tr>
<td>2002</td>
<td>293 879 439⁵</td>
<td>3 770 000</td>
</tr>
<tr>
<td>2003</td>
<td>277 121 981⁵</td>
<td>2 940 000</td>
</tr>
<tr>
<td>2004</td>
<td>274 360 927⁵</td>
<td>2 480 000</td>
</tr>
<tr>
<td>2005</td>
<td>255 200 701⁵</td>
<td>1 940 000</td>
</tr>
<tr>
<td>2006</td>
<td>245 123 805⁵</td>
<td>1 830 000</td>
</tr>
<tr>
<td>2007</td>
<td>223 208 499⁵</td>
<td>1 994 000</td>
</tr>
<tr>
<td>2008</td>
<td>192 264 956⁵</td>
<td>1 670 000</td>
</tr>
<tr>
<td>2009</td>
<td>180 759 206⁵</td>
<td>1 618 500</td>
</tr>
<tr>
<td>2010</td>
<td>168 656 949⁵</td>
<td>1 567 000</td>
</tr>
<tr>
<td>2011</td>
<td>155 481 492⁵</td>
<td>1 515 500</td>
</tr>
<tr>
<td>2012</td>
<td>142 439 972⁵</td>
<td>1 555 300</td>
</tr>
<tr>
<td>2013</td>
<td>133 179 780⁵</td>
<td>1 391 900</td>
</tr>
<tr>
<td>2014</td>
<td>120 816 120⁵</td>
<td>1 432 200</td>
</tr>
<tr>
<td>2015</td>
<td>110 489 001⁵</td>
<td>1 662 200</td>
</tr>
<tr>
<td>2016</td>
<td>101 276 190⁵</td>
<td>1 983 400</td>
</tr>
</tbody>
</table>


*Data in bold are compiled data, other data is extrapolated or interpolated.

6.2.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The used uncertainties are presented below.

NMVOC

Emission factor:
0.5-3.0 kg/Mg
( Default).

Activity data “Emission of landfill gas”:
± 64 % (1990),
± 55 % (2017),
(Expert judgement).

TSP, PM₁₀ and PM₂.₅

Emission factor TSP:
0.006-2.21 g/Mg
( Default).
Emission factor $\text{PM}_{10}$:
0.003-1.05 g/Mg  
(Default).

Emission factor $\text{PM}_{2.5}$:
0.0004-0.16 g/Mg  
(Default).

Activity data “Landfilled waste at landfills for municipal solid waste”:
± 15% (1990),
± 10% (2017),  
(Expert judgement).

The time series in the waste sector are calculated consistently. When statistics are not produced annually, interpolation and extrapolation have been necessary tools for imputation.

6.2.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

6.2.1.5 SOURCE-SPECIFIC REcalculations
Recalculations have been made in submission 2018 for NMVOC for the years 2008-2015, because of adjustments of activity data (emissions of methane). New data on landfill gas recovery has been implemented as a result of a study on landfill gas recovery at closed landfills. The emissions have changed by between -1.97% and +0.32%.

6.2.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.3 Biological treatment of waste, NFR 5B
This category includes Composting (NFR 5B1) and Anaerobic digestion at biogas facilities (NFR 5B2).

Emissions from compost production from waste are calculated.

Because of lack of methods in the EMEP/EEA Guidebook 2013, no emissions from anaerobic digestion at biogas facilities are calculated. Therefore emissions from anaerobic digestion are reported as NA (Not applicable). The provided methods and EFs for NH$_3$ in the EMEP/EEA Guidebook 2016 has not been implemented in the inventory for submission 2018.
6.3.1 Composting, NFR 5B1

6.3.1.1 SOURCE CATEGORY DESCRIPTION
Sweden is reporting of emissions of NH$_3$ (from covered composting and windrow composting) and CO (from windrow composting). Emissions of NO$_X$, NMVOC, TSP, PM$_{10}$, PM$_{2.5}$ and BC are reported as not estimated (NE). Other emissions are reported as not applicable (NA).

Table 6-3. Summary of key source analysis, NFR5B1, according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>5B1</td>
<td>NH$_3$</td>
<td></td>
</tr>
</tbody>
</table>

6.3.1.2 METHODOLOGICAL ISSUES

6.3.1.2.1 Emission and abatement factors used
Tier 2 default emission factors from the EMEP/EEA Emission Inventory Guidebook 2013/2016 are used for NH$_3$ from compost production (covered composting). When composting food waste and household waste in Sweden, the composting process is normally covered.

The abatement factor is used in from year 2005. From year 1994 to 2005, this factor is estimated to be gradually increasing from zero to the default factor due to reflect an increasing degree of practicing abatement techniques. See further in the tables below.

Table 6-4. Emission factor used for NFR 5B1 Composting (covered composting).

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Lower</td>
<td>Upper</td>
</tr>
<tr>
<td>NH$_3$</td>
<td>0.24</td>
<td>kg/Mg organic waste</td>
<td>0.1</td>
<td>0.7</td>
</tr>
</tbody>
</table>

Table 6-5. Abatement factor used for NFR 5B1 Composting (covered composting).

<table>
<thead>
<tr>
<th>Abatement technology</th>
<th>Pollutant</th>
<th>Efficiency</th>
<th>Default value</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Biofilter</td>
<td>NH$_3$</td>
<td>90 %</td>
<td></td>
<td>70 %</td>
<td>97 %</td>
</tr>
</tbody>
</table>

Tier 2 default emission factors from the EMEP/EEA Emission Inventory Guidebook 2013/2016 are used for NH$_3$ and CO (from windrow composting of garden and park waste). See further in the table below.

Table 6-6. Emission factors used for NFR 5B1 Composting (windrow composting)

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>Lower</td>
<td>Upper</td>
</tr>
<tr>
<td>CO</td>
<td>0.56</td>
<td>kg/Mg waste</td>
<td>0.05</td>
<td>1</td>
</tr>
<tr>
<td>NH$_3$</td>
<td>0.66</td>
<td>kg/Mg waste</td>
<td>0.05</td>
<td>1</td>
</tr>
</tbody>
</table>
6.3.1.2.2 Activity data used

Activity data on composted waste (covered composting and windrow composting) is used for emissions of CO and NH₃. The data has been compiled and published by the association Swedish Waste Management (RVF/Avfall Sverige).

Table 6-7. Activity data used for NFR5B1, Composting.

<table>
<thead>
<tr>
<th>Year</th>
<th>Composted waste* (t, wet weight) (AD used for NH₃ from covered composting)</th>
<th>Composted waste* (t, wet weight) (AD used for CO and NH₃ from windrow composting of garden and park waste)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>50 000¹</td>
<td>20 000¹</td>
</tr>
<tr>
<td>1991</td>
<td>44 940</td>
<td>60 460</td>
</tr>
<tr>
<td>1992</td>
<td>39 880</td>
<td>100 920</td>
</tr>
<tr>
<td>1993</td>
<td>34 820</td>
<td>141 380</td>
</tr>
<tr>
<td>1994</td>
<td>29 760</td>
<td>181 840</td>
</tr>
<tr>
<td>1995</td>
<td>24 700²</td>
<td>222 300²</td>
</tr>
<tr>
<td>1996</td>
<td>43 350</td>
<td>197 650</td>
</tr>
<tr>
<td>1997</td>
<td>62 000³</td>
<td>173 000³</td>
</tr>
<tr>
<td>1998</td>
<td>72 333</td>
<td>185 167</td>
</tr>
<tr>
<td>1999</td>
<td>82 667</td>
<td>197 333</td>
</tr>
<tr>
<td>2000</td>
<td>93 000⁴</td>
<td>197 000⁴</td>
</tr>
<tr>
<td>2001</td>
<td>102 492</td>
<td>193 271</td>
</tr>
<tr>
<td>2002</td>
<td>111 984⁵</td>
<td>189 546⁵</td>
</tr>
<tr>
<td>2003</td>
<td>108 745⁵</td>
<td>273 215⁵</td>
</tr>
<tr>
<td>2004</td>
<td>99 950⁵</td>
<td>289 430⁵</td>
</tr>
<tr>
<td>2005</td>
<td>234 640⁶</td>
<td>225 190⁶</td>
</tr>
<tr>
<td>2006</td>
<td>248 230⁷</td>
<td>204 160⁷</td>
</tr>
<tr>
<td>2007</td>
<td>261 450⁸</td>
<td>253 840⁸</td>
</tr>
<tr>
<td>2008</td>
<td>278 000⁹</td>
<td>290 700⁹</td>
</tr>
<tr>
<td>2009</td>
<td>284 940ⁱ⁰</td>
<td>345 560¹⁰</td>
</tr>
<tr>
<td>2010</td>
<td>297 180¹¹</td>
<td>269 030¹¹</td>
</tr>
<tr>
<td>2011</td>
<td>257 110¹²</td>
<td>432 990¹²</td>
</tr>
<tr>
<td>2012</td>
<td>246 680¹³</td>
<td>312 150¹³</td>
</tr>
<tr>
<td>2013</td>
<td>211 260¹⁴</td>
<td>317 210¹⁴</td>
</tr>
<tr>
<td>2014</td>
<td>197 140¹⁵</td>
<td>270 780¹⁵</td>
</tr>
<tr>
<td>2015</td>
<td>208 430¹⁶</td>
<td>209 910¹⁶</td>
</tr>
<tr>
<td>2016</td>
<td>181 047¹⁷</td>
<td>295 093¹⁷</td>
</tr>
</tbody>
</table>


*Data in bold are compiled, other data is extrapolated or interpolated.

6.3.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The used uncertainties are presented below.

**NH₃ (from covered composting)**

Emission factor:
0.1-0.7 kg/Mg organic waste
(Default).

Abatement factor:
70%-97%
(Default).
Activity data “Composted waste”:
± 15 % (1990),
± 10 % (2017),
(Expert judgement).

NH₃ and CO (from windrow composting)
Emission factor (NH₃):
0.05-1 kg/Mg waste
(Default).

Emission factor (CO):
0.05-1 kg/Mg waste
(Default).

Activity data “Composted waste”:
± 20 % (1990),
± 15 % (2017),
(Expert judgement).

The time series in the waste sector are calculated consistently. When statistics are not produced annually, interpolation and extrapolation have been necessary tools for imputation.

6.3.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

6.3.1.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been made in submission 2018 for CO and NH₃ from composting.

6.3.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.4 Waste incineration, NFR 5C

6.4.1 Emissions from incineration of municipal waste, industrial waste, clinical waste and sewage sludge, NFR 5C1a, 5C1bi, 5C1bii and 5C1biv

Emissions from these sources reported for one plant in Sweden, and included in NFR 5C1bii. Since 2003, also MSW incineration occurs at the plant.

Table 6-8. Summary of key source analysis, NFR5C1 according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>5C1</td>
<td>DIOX, Hg</td>
<td>DIOX, Hg</td>
</tr>
</tbody>
</table>
6.4.2 Emissions from incineration of hazardous waste, NFR 5C1bii

6.4.2.1 SOURCE CATEGORY DESCRIPTION
Emissions from incineration of hazardous and industrial waste and since 2003 also MSW, from one large plant are reported in NFR 5C1bii. In NFR 5C1bv emissions of mercury, dioxin, benzo(a)pyrene and PAH-4 from cremation are reported, and from submission 2016 onwards also NO\textsubscript{X}, SO\textsubscript{2}, NMVOC, CO, HCB, PCB, heavy metals other than mercury, and PAH other than benzo(a)pyrene are included in the reporting. Particulate matter from cow and sheep burn using air curtain incinerator is also reported in NFR 5C1bv starting from submission 2016. Particulate matter and PAH-4 from domestic open waste burning, such as garden fires, are reported in 5C2. Other possible emissions from garden fires and bonfires are currently not reported due to lack of suitable emission factors.

Regarding incineration of medical waste, no national activity and emission data for this source category is available.

6.4.2.2 METHODOLOGICAL ISSUES
Incineration of hazardous waste, other than cremation, occurs at nine plants in Sweden. There is one major plant for handling and destruction of hazardous waste, which is the only one for which emission data is available. For 2004 around 88\% of the total amount of incinerated hazardous waste was incinerated at this plant. The emissions from the plant are reported in 5C1bii. Emissions from incineration of hazardous waste not reported in 5C1bii are included in 1A1a and in 1A2c, d, e and f.

The facility included in 5C1bii was operated with an electrostatic precipitator (ESP) from the start in 1983 until 1990, when a textile filter with coal injection replaced the ESP. During 2000, wet flue gas cleaning was installed after the textile filter.

Reported emissions are for the whole time series obtained from the facility’s environmental report or directly from the facility on request. Reported emissions are NO\textsubscript{X}, SO\textsubscript{2}, NMVOC, CO, NH\textsubscript{3}, particulate matter, Pb, Cd, Hg, As, Cr, Cu, Ni, Se, Zn, PAH1-4, B(a)P, B(b)F, B(k)F, I(cd)P, HCB, PCB and dioxin. SO\textsubscript{2}, NO\textsubscript{X}, CO, particulate matter and Hg are continuously measured in the flue gases. Dioxins in flue gases have been measured by spot tests, but are continuously collected and analysed once a week since June 2001. Emissions of NH\textsubscript{3} have been obtained from the facility from 2008 and onwards. For 2003 – 2007 reported NH\textsubscript{3} emissions are calculated based on implied emission factor for 2008. The estimates for Se, PAH1-4, B(a)P, B(b)F, B(k)F, I(cd)P, HCB, PCB and Zn are based on the amount of incinerated waste and emission factors from EMEP/EEA Guidebook 2013.
The time series for Pb, Cd, As, Cr, Cu and Ni are not consistently reported due to lack of data in the environmental reports for 2006 - 2009. For the years with reported notation key NE, the reason is that the amounts of metals emitted in the flue gas are below the detection limit.

The activity has increased over time. In 1995 the plant combusted about 22 000 t and in 2002 the corresponding value was about 33 000 t. In 2003 the capacity of the plant was increased substantially by taking a new incinerator into operation. In this new incinerator, the facility incinerates a mixture of MSW, industrial waste and hazardous waste. As a consequence of increased capacity, emissions from 2003 are increased compared to earlier years.

6.4.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for emissions continuously measured in the flue gases are expected to be quite low, ± 30%. Uncertainties for emissions calculated with emission factors from EMEP/EEA Guidebook 2013 are based on data in the Guidebook. For all other emissions, measured by spot tests, the uncertainties are set to ± 50%.

More information is given in IIR Annex 1.

Time series for incineration of hazardous waste reported in NFR code 5C1bii have been reviewed in later years and considered to be consistent.

6.4.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No SOURCE-SPECIFIC QA/QC has been performed.

6.4.2.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations are performed.

6.4.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

6.4.3 Emissions from cremation, NFR 5C1bv

6.4.3.1 SOURCE CATEGORY DESCRIPTION

In NFR 5C1bv emissions of mercury, dioxin, benzo(a)pyrene and PAH-4 from cremation are reported, and from submission 2016 onwards also NOx, SO2, NMVOC, CO, HCB, PCB, heavy metals other than mercury, and PAH other than benzo(a)pyrene are included in the reporting.

Particulate matter from cow and sheep burn using air curtain incinerator is reported in NFR 5C1bv from submission 2016 onwards.
6.4.3.2 METHODOLOGICAL ISSUES

Estimated emissions of PAH-4, benzo(a)pyrene and dioxin from cremation have been calculated based on national emission factors and statistics on the number of annual cremations. In submission 2016 emissions of NO\textsubscript{X}, TSP, PM\textsubscript{10}, PM\textsubscript{2.5}, SO\textsubscript{2}, NMVOC, CO, As, Cr, Cu, Ni, Pb, Se, Zn, PCB, HCB, B(b)F, B(k)F, and I(cd)P from cremation are reported for the first time. The estimates are made with emission factors from EMEP/EEA Guidebook 2013. Emissions from PAH-4 have been adjusted with respect to available estimates for B(b)F, B(k)F and I(cd)P. BC emissions have not been estimated due to lack of information and reported NE in accordance with EMEP/EEA Guidebook 2013.

The emissions of mercury are estimated using a methodology presented by Wängberg (2013)\textsuperscript{193}. From the late 1990’s, abatement techniques have been considered in the estimations. The method is based on statistics on the annual amount of cremations at each of the Swedish crematories in combination with information on installation of emission control, i.e. filter with activated carbon. The implied emission factors for 1980 - 2015 are given in Table 6-9.

<table>
<thead>
<tr>
<th>Year</th>
<th>IEF Submission 2018</th>
</tr>
</thead>
<tbody>
<tr>
<td>1980</td>
<td>0.00300</td>
</tr>
<tr>
<td>1985</td>
<td>0.00300</td>
</tr>
<tr>
<td>1990</td>
<td>0.00296</td>
</tr>
<tr>
<td>1995</td>
<td>0.00287</td>
</tr>
<tr>
<td>2000</td>
<td>0.00167</td>
</tr>
<tr>
<td>2001</td>
<td>0.00103</td>
</tr>
<tr>
<td>2002</td>
<td>0.00100</td>
</tr>
<tr>
<td>2003</td>
<td>0.00099</td>
</tr>
<tr>
<td>2004</td>
<td>0.00093</td>
</tr>
<tr>
<td>2005</td>
<td>0.00087</td>
</tr>
<tr>
<td>2006</td>
<td>0.00077</td>
</tr>
<tr>
<td>2007</td>
<td>0.00073</td>
</tr>
<tr>
<td>2008</td>
<td>0.00071</td>
</tr>
<tr>
<td>2009</td>
<td>0.00072</td>
</tr>
<tr>
<td>2010</td>
<td>0.00071</td>
</tr>
<tr>
<td>2011</td>
<td>0.00064</td>
</tr>
<tr>
<td>2012</td>
<td>0.00059</td>
</tr>
<tr>
<td>2013</td>
<td>0.00051</td>
</tr>
<tr>
<td>2014</td>
<td>0.00038</td>
</tr>
<tr>
<td>2015</td>
<td>0.00034</td>
</tr>
<tr>
<td>2016</td>
<td>0.00034</td>
</tr>
</tbody>
</table>

Emission factors used to calculate PAH-4 emissions from cremation are from USEPA\textsuperscript{194} and for dioxin a suggested emission factor from the European Dioxin

\textsuperscript{193} Wängberg, I. 2013. PM Utredning nr 7 Hg från krematorier
\textsuperscript{194} USEPA. 1998. Locating and Estimating Air Emissions from Sources of Polycyclic organic matter. EPA-454/R-98-014. Office of Air Quality Planning and Standards, USA
Inventory\textsuperscript{195} was used. UNEP\textsuperscript{196} presents emission factors for dioxins in the range 0.4 – 90 µg TEQ/cremation, while an earlier Swedish Inventory\textsuperscript{197} suggested 6-12 µg TEQ/cremation, referred to in the European Dioxin Inventory. An average of 9 µg TEQ/cremation has been used in the present emission estimates. This agrees with a recent experimental study that recommends 6-13 µg TEQ/cremation\textsuperscript{198}. The number of annual cremations has increased from 47000 in 1980 to more than 71000 in 2014, and associated dioxin emissions have thus increased from 0.42 g TEQ to approximately 0.64 g TEQ during the same period.

Emissions of particulate matter (TSP, PM\textsubscript{10} and PM\textsubscript{2.5}) from cow and sheep burn using air curtain incinerator are for the first time reported in submission 2016. The estimates are made with emission factors from EMEP/EEA Guidebook 2013. Cow and sheep burn using air curtain incinerator occurs rather seldom in Sweden; according to the Swedish Farming Services – the company responsible for collection and handling of cadavers – this only happens when there is a significant risk of infection, otherwise alternative utilization methods are used. The company estimates that only a few animals are burned with air curtain incinerator annually\textsuperscript{199}.

6.4.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

Uncertainties for NFR 5C1bv are displayed in Annex 1.

Time series for emissions from cremations reported in NFR code 5C1bv have been reviewed in later years and considered to be consistent.

6.4.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No SOURCE-SPECIFIC QA/QC has been performed.

6.4.3.5 SOURCE-SPECIFIC RECALCULATIONS

No source-specific recalculations are performed in submission 2018.

6.4.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.

\textsuperscript{195} Quass et al., 2001. \url{http://europa.eu.int/comm/environment/dioxin/pdf/stage1/cremation.pdf}

\textsuperscript{196} UNEP, 2001. \url{www.chem.unep.ch/pops/pdf/toolkit/toolkit.pdf}

\textsuperscript{197} deWit. 1993, unpublished

\textsuperscript{198} Wang, 2003.

\textsuperscript{199} Mikael Lidholm, Svensk Lantbrukstjänst AB
6.4.4 Particles and PAH from garden burning and bonfires, NFR 5C2

6.4.4.1 SOURCE CATEGORY DESCRIPTION
Particulate matter and PAH-4 from domestic open waste burning, such as garden fires, are reported in 5C2. Other possible emissions from garden fires and bonfires are currently not reported due to lack of suitable emission factors.

Table 6-10. Summary of key source analysis, NFR5C2 according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
<th>Trend</th>
</tr>
</thead>
<tbody>
<tr>
<td>5C2</td>
<td>$PM_{2.5}$, As, TSP, Cd, $PM_{10}$, DIOX, Zn</td>
<td>$PM_{2.5}$, As, TSP, Cd, $PM_{10}$, DIOX</td>
</tr>
</tbody>
</table>

6.4.4.2 METHODOLOGICAL ISSUES
In order to estimate emission of PAH from burning of garden waste, emission factors from USEPA were used, while emission factors for open burning of waste suggested by CEPMEIP\textsuperscript{200} were used to estimate emissions of TSP, $PM_{10}$ and $PM_{2.5}$. Emission factors presented in EMEP/EEA Guidebook 2009\textsuperscript{201}, chapter “Small-scale waste burning” represents emissions from open burning of agricultural waste. A study in 2004\textsuperscript{202} reveals that it is very rare that the farmers practice field burning in Sweden. Thus, only PAH and particle emissions are reported in NFR 5C2.

In submission 2016 emissions of Pb, Cd, As, Cr, Cu, Se, Zn and dioxines from burning of garden waste are reported for the first time for 1996 and onwards. The estimates are calculated with emission factors from EMEP/EEA Guidebook 2013.

As there are no national statistics regarding the extent of garden burning and bonfires, instead statistics on number of small houses have been used. The data should be considered as indicative levels of emissions from these sources.

6.4.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions of TSP, $PM_{10}$, $PM_{2.5}$ are based on expert judgement. Assessed uncertainties for are ± 100%, ± 100% and ± 100%, respectively. Uncertainties for emissions for dioxin and metals are based on expert judgement. Uncertainty for DIOX, Pb, Cd, As, Se, Zn, Cr and Cu are ± 200%, ± 200%, ± 200%, 200%, ± 200%, ± 20%, ± 250% and ± 195%, respectively.

More information is given in IIR Annex 1.

Time series for particles and PAH from garden burning and bonfires reported in NFR code 5C2 have been reviewed in later years and considered to be consistent.

\textsuperscript{200} CEPMEIP, 2001. TNO.


\textsuperscript{202} Wikström, H. and Adolfsson, R. 2004. Field Burning of Crop Residues.
6.4.4.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No SOURCE-SPECIFIC QA/QC has been performed.

6.4.4.5 SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculation are performed in submission 2018.

6.4.4.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.5 Waste-water handling, NFR 5D

6.5.1 Domestic wastewater handling, NFR 5D1

6.5.1.1 SOURCE CATEGORY DESCRIPTION
Sweden is reporting emissions of NMVOC (from municipal wastewater treatment plants) and NH$_3$ (from latrines).

Emissions of TSP, PM$_{10}$, PM$_{2.5}$, Pb, Cd, Hg, As, Cr, Cu, Ni, Se and Zn are reported as not estimated (NE). Other emissions are reported as not applicable (NA).

6.5.1.2 METHODOLOGICAL ISSUES

6.5.1.2.1 Emission factors used
Tier 1 default emission factor from the EMEP/EEA Emission Inventory Guidebook 2013/2016 are used for NMVOC (from wastewater handling at municipal wastewater treatment plants). See further in the table below.

Table 6-11. Emission factor used for NFR 5D1 Domestic wastewater handling (at municipal wastewater treatment plants).

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>NMVOC</td>
<td>15</td>
<td>mg/m$^3$ waste water</td>
<td>5 - 50</td>
<td>Atasoy et al. (2004)</td>
</tr>
</tbody>
</table>

Tier 2 default emission factor from the EMEP/EEA Emission Inventory Guidebook 2013 are used for NH$_3$ (from latrines). See further in the table below.

Table 6-12. Emission factor used for NFR 5D1 Domestic wastewater handling (latrines).

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>NH$_3$</td>
<td>1.6</td>
<td>kg/person/year</td>
<td>0.8 - 3.2</td>
<td>Guidebook</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>(2006)</td>
</tr>
</tbody>
</table>
6.5.1.2.2  Activity data used

Activity data on discharged volumes of treated wastewater is used for NMVOC emissions, compiled and published by Statistics Sweden for Swedish EPA.

In the previous Swedish inventory, activity data on population not connected to municipal wastewater treatment plants has been used as a substitute for data on population with latrines. The data has been compiled by Statistics Sweden for Swedish EPA. The average population for the years 1990-2015 is 1 300 000 people.

The most common sanitary solution for the population not connected to municipal wastewater treatment plants is septic tanks. The population with latrines is very small, but has not been properly investigated. The technical correction from the NECD (National Emission Ceilings Directive) Review 2017 estimated, based on the situation in five comparable countries, the population with latrines to be 1% of the population not connected to municipal wastewater treatment plants. This equals to a population of 13 000 people with latrines in Sweden.

Table 6-13. Activity data used for NFR 5D1 Domestic wastewater handling.

<table>
<thead>
<tr>
<th>Year</th>
<th>Discharged volumes of treated wastewater* (1000 m³)</th>
<th>Population with latrines* (persons)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>(AD used for NMVOC from wastewater handling at municipal wastewater treatment plants)</td>
<td>(AD used for NH₃ from latrines)</td>
</tr>
<tr>
<td>1990</td>
<td>1 305 000²</td>
<td>13 000</td>
</tr>
<tr>
<td>1991</td>
<td>1 276 050</td>
<td>13 000</td>
</tr>
<tr>
<td>1992</td>
<td>1 247 100²</td>
<td>13 000</td>
</tr>
<tr>
<td>1993</td>
<td>1 263 600</td>
<td>13 000</td>
</tr>
<tr>
<td>1994</td>
<td>1 280 100</td>
<td>13 000</td>
</tr>
<tr>
<td>1995</td>
<td>1 296 600¹</td>
<td>13 000</td>
</tr>
<tr>
<td>1996</td>
<td>1 315 067</td>
<td>13 000</td>
</tr>
<tr>
<td>1997</td>
<td>1 333 533</td>
<td>13 000</td>
</tr>
<tr>
<td>1998</td>
<td>1 352 000¹</td>
<td>13 000</td>
</tr>
<tr>
<td>1999</td>
<td>1 357 459</td>
<td>13 000</td>
</tr>
<tr>
<td>2000</td>
<td>1 362 917⁵</td>
<td>13 000</td>
</tr>
<tr>
<td>2001</td>
<td>1 295 459</td>
<td>13 000</td>
</tr>
<tr>
<td>2002</td>
<td>1 228 000⁶</td>
<td>13 000</td>
</tr>
<tr>
<td>2003</td>
<td>1 206 612</td>
<td>13 000</td>
</tr>
<tr>
<td>2004</td>
<td>1 185 223⁷</td>
<td>13 000</td>
</tr>
<tr>
<td>2005</td>
<td>1 212 514</td>
<td>13 000</td>
</tr>
<tr>
<td>2006</td>
<td>1 239 805⁸</td>
<td>13 000</td>
</tr>
<tr>
<td>2007</td>
<td>1 249 172</td>
<td>13 000</td>
</tr>
<tr>
<td>2008</td>
<td>1 258 539⁹</td>
<td>13 000</td>
</tr>
<tr>
<td>2009</td>
<td>1 222 653</td>
<td>13 000</td>
</tr>
<tr>
<td>2010</td>
<td>1 186 767¹⁰</td>
<td>13 000</td>
</tr>
<tr>
<td>2011</td>
<td>1 227 949</td>
<td>13 000</td>
</tr>
<tr>
<td>2012</td>
<td>1 269 131¹¹</td>
<td>13 000</td>
</tr>
<tr>
<td>2013</td>
<td>1 243 112</td>
<td>13 000</td>
</tr>
<tr>
<td>2014</td>
<td>1 217 093¹²</td>
<td>13 000</td>
</tr>
<tr>
<td>2015</td>
<td>1 217 093</td>
<td>13 000</td>
</tr>
<tr>
<td>2016</td>
<td>1 217 093</td>
<td>13 000</td>
</tr>
</tbody>
</table>


*Data in bold are compiled, other data is extrapolated or interpolated.
6.5.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The used uncertainties are presented below.

**NMVOC (from municipal wastewater treatment plants)**

Emission factor:
5-50 mg/m$^3$ wastewater
(Default).

Activity data “Discharged volumes of treated wastewater”:
± 10 % (1990),
± 10 % (2017),
(Expert judgement).

**NH$_3$ (from latrines)**

Emission factor:
0.8-3.2 kg/person/year
(Default).

Activity data “Population with latrines”:
± 10 % (1990),
± 5 % (2017),
(Expert judgement).

The time series in the waste sector are calculated consistently. When statistics are not produced annually, interpolation and extrapolation have been necessary tools for imputation.

6.5.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION

No source-specific QA/QC or verification is performed.

6.5.1.5 SOURCE-SPECIFIC RECALCULATIONS

No recalculations have been made in submission 2018 for NMVOC from municipal wastewater treatment plants.

For NH$_3$ from latrines, new estimates have been calculated as a result of implementation of a technical correction from the NECD (National Emission Ceilings Directive) Review 2017. The recalculation led to a 99% decrease of NH$_3$ emissions from latrines for the years 1990-2015.

6.5.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS

No major improvements are planned for the next submission.
6.5.2 Industrial wastewater handling, NFR 5D2

6.5.2.1 SOURCE CATEGORY DESCRIPTION
Sweden is reporting of emissions of NMVOC. Emissions of NH₃, TSP, PM₁₀, PM₂.₅, Pb, Cd, Hg, As, Cr, Cu, Ni, Se and Zn are reported as not estimated (NE). Other emissions are reported as not applicable (NA).

6.5.2.2 METHODOLOGICAL ISSUES

6.5.2.2.1 Emission factors used
Tier 1/Tier 2 default emission factor from the EMEP/EEA Guidebook 2013/2016 is used for NMVOC (from wastewater treatment in industry).

Table 6-14. Emission factor used for NFR 5D2 Industrial wastewater handling.

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95 % confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NMVOC</td>
<td>15</td>
<td>mg/m³ wastewater</td>
<td>5</td>
<td>50</td>
</tr>
</tbody>
</table>

6.5.2.2.2 Activity data used
Activity data on discharged volumes of treated wastewater is used for emissions of NMVOC. The data has been compiled and published by Statistics Sweden.

Table 6-15. Activity data used for NFR 5D2 Industrial wastewater handling (NMVOC).

<table>
<thead>
<tr>
<th>Year</th>
<th>Discharged volumes of treated wastewater* (1000 m³)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>933 056</td>
</tr>
<tr>
<td>1991</td>
<td>933 056</td>
</tr>
<tr>
<td>1992</td>
<td>933 056</td>
</tr>
<tr>
<td>1993</td>
<td>933 056</td>
</tr>
<tr>
<td>1994</td>
<td>933 056</td>
</tr>
<tr>
<td>1995</td>
<td>933 056</td>
</tr>
<tr>
<td>1996</td>
<td>930 496</td>
</tr>
<tr>
<td>1997</td>
<td>927 936</td>
</tr>
<tr>
<td>1998</td>
<td>925 375</td>
</tr>
<tr>
<td>1999</td>
<td>922 815</td>
</tr>
<tr>
<td>2000</td>
<td>920 255</td>
</tr>
<tr>
<td>2001</td>
<td>931 282</td>
</tr>
<tr>
<td>2002</td>
<td>942 310</td>
</tr>
<tr>
<td>2003</td>
<td>953 337</td>
</tr>
<tr>
<td>2004</td>
<td>964 365</td>
</tr>
<tr>
<td>2005</td>
<td>975 392</td>
</tr>
<tr>
<td>2006</td>
<td>988 232</td>
</tr>
<tr>
<td>2007</td>
<td>1 001 072</td>
</tr>
<tr>
<td>2008</td>
<td>1 013 911</td>
</tr>
<tr>
<td>2009</td>
<td>1 026 751</td>
</tr>
<tr>
<td>2010</td>
<td>1 039 591</td>
</tr>
<tr>
<td>2011</td>
<td>1 010 917</td>
</tr>
<tr>
<td>2012</td>
<td>982 242</td>
</tr>
<tr>
<td>2013</td>
<td>953 568</td>
</tr>
<tr>
<td>2014</td>
<td>924 893</td>
</tr>
<tr>
<td>2015</td>
<td>896 219</td>
</tr>
<tr>
<td>2016</td>
<td>896 219</td>
</tr>
</tbody>
</table>


*Data in bold are compiled, other data is extrapolated or interpolated.
6.5.2.3  UNCERTAINTIES AND TIME-SERIES CONSISTENCY
The used uncertainties are presented below.

NMVOC (from industrial wastewater handling)
Emission factor:
5-50 mg/m³ wastewater
(Default).

Activity data “Discharged volumes of treated wastewater”:
± 50 % (1990),
± 50 % (2017),
(Expert judgement).

The time series in the waste sector are calculated consistently. When statistics are
not produced annually, interpolation and extrapolation have been necessary tools
for imputation.

6.5.2.4  SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

6.5.2.5  SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been made in submission 2018 for NMVOC from
wastewater treatment in industry.

6.5.2.6  SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.6  Other waste, NFR 5E

The NFR category 5E Other waste, includes emissions from:
- Sludge spreading (mechanical dewatering of digested sludge)
- Landfill fires
- House and Car fires
- Pets

Table 6-16. Summary of key source analysis, NFR5E according to approach 1.

<table>
<thead>
<tr>
<th>NFR</th>
<th>Key Source Assessment 2016</th>
</tr>
</thead>
<tbody>
<tr>
<td>5E</td>
<td>DIOX, PM2.5, NH3, PM10, TSP, As, Cd</td>
</tr>
<tr>
<td></td>
<td>DIOX, PM2.5, NH3, PM10, Hg, TSP</td>
</tr>
</tbody>
</table>

6.6.1  Other waste, sludge spreading, NFR 5E

6.6.1.1  SOURCE CATEGORY DESCRIPTION
Sweden is reporting of emissions of NH3 from sludge spreading (mechanical de-
watering of digested sludge at wastewater treatment plants).
6.6.1.2 METHODOLOGICAL ISSUES

6.6.1.2.1 Emission factors used

Tier 2 default emission factor from the EMEP/EEA Emission Inventory Guidebook 2013/2016 are used for NH$_3$ (from sludge spreading). See further in the table below.

Table 6-17. Emission factor used for NFR 5E Other waste, Sludge spreading.

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Value</th>
<th>Unit</th>
<th>95% confidence interval</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>NH$_3$</td>
<td>50</td>
<td>g/kg NH$_3$ in the sludge</td>
<td>10 - 150</td>
<td>Guidebook (2006)</td>
</tr>
</tbody>
</table>

Table 6-18. Activity data used for NFR 5E Other waste, Sludge spreading.

<table>
<thead>
<tr>
<th>Year</th>
<th>Quantity of N in anaerobically digested sludge* (t) (AD used for NH$_3$ from sludge spreading)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1990</td>
<td>8 073</td>
</tr>
<tr>
<td>1991</td>
<td>8 073</td>
</tr>
<tr>
<td>1992</td>
<td>8 073</td>
</tr>
<tr>
<td>1993</td>
<td>8 073</td>
</tr>
<tr>
<td>1994</td>
<td>8 073</td>
</tr>
<tr>
<td>1995</td>
<td>8 073$^1$</td>
</tr>
<tr>
<td>1996</td>
<td>8 296</td>
</tr>
<tr>
<td>1997</td>
<td>8 518</td>
</tr>
<tr>
<td>1998</td>
<td>8 741$^2$</td>
</tr>
<tr>
<td>1999</td>
<td>8 656</td>
</tr>
<tr>
<td>2000</td>
<td>8 571$^3$</td>
</tr>
<tr>
<td>2001</td>
<td>8 878</td>
</tr>
<tr>
<td>2002</td>
<td>9 185$^4$</td>
</tr>
<tr>
<td>2003</td>
<td>8 802</td>
</tr>
<tr>
<td>2004</td>
<td>8 419$^5$</td>
</tr>
<tr>
<td>2005</td>
<td>8 565</td>
</tr>
<tr>
<td>2006</td>
<td>8 710$^6$</td>
</tr>
<tr>
<td>2007</td>
<td>8 908</td>
</tr>
<tr>
<td>2008</td>
<td>9 105$^7$</td>
</tr>
<tr>
<td>2009</td>
<td>9 038</td>
</tr>
<tr>
<td>2010</td>
<td>8 971$^8$</td>
</tr>
<tr>
<td>2011</td>
<td>8 948</td>
</tr>
<tr>
<td>2012</td>
<td>8 925$^9$</td>
</tr>
<tr>
<td>2013</td>
<td>8 989</td>
</tr>
<tr>
<td>2014</td>
<td>9 053$^{10}$</td>
</tr>
<tr>
<td>2015</td>
<td>8 919</td>
</tr>
<tr>
<td>2016</td>
<td>8 919</td>
</tr>
</tbody>
</table>

*Data in bold are compiled, other data is extrapolated or interpolated.

6.6.1.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY

The used uncertainties are presented below.

NH$_3$ (from mechanical dewatering of digested sludge)

Emission factor:
10-150 g/kg N in the sludge
(Default).
Activity data “Quantity of nitrogen in anaerobically digested sludge”:
± 10 % (1990),
± 2 % (2017),
(Expert judgement).

The time series in the waste sector are calculated consistently. When statistics are
not produced annually, interpolation and extrapolation have been necessary tools
for imputation.

6.6.1.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC or verification is performed.

6.6.1.5 SOURCE-SPECIFIC RECALCULATIONS
No recalculations have been made in submission 2018 for NH₃ from mechanical
dewatering of digested sludge at wastewater treatment plants.

6.6.1.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.6.2 Other waste, landfill fires, NFR 5E
6.6.2.1 SOURCE CATEGORY DESCRIPTION
In the sector 5E Other waste, emissions of dioxin, PAH-4 and Hg from landfill
fires (1996 and onwards) are included.

6.6.2.2 METHODOLOGICAL ISSUES
All emissions from landfill fires are in this submission based on the frequency and
duration of fires in Sweden²⁰³ and emission factors derived from measurements
performed during landfill fires²⁰⁴. The fires were classified as “underground fires”
(> 48 hours) or “surface fires” (< 48 hours) depending on the duration of the fire.
In Table 6-25, the emission factors used are presented, and in Table 6-26, the re-
ported emissions of Hg, dioxin and PAH-4 from landfill fires are presented.

<table>
<thead>
<tr>
<th>Fire category</th>
<th>Hg g/hour</th>
<th>Dioxin g/hour</th>
<th>benzo(a)pyrene g/hour</th>
<th>benzo(b)fluoranthene g/hour</th>
<th>benzo(k)fluoranthene g/hour</th>
<th>Indeno(1,2,3-cd)-pyrene g/hour</th>
</tr>
</thead>
<tbody>
<tr>
<td>Surface</td>
<td>1.30</td>
<td>200 × 10⁻⁶</td>
<td>0.15</td>
<td>0.24</td>
<td>NE</td>
<td>0.09</td>
</tr>
<tr>
<td>Underground</td>
<td>0.031</td>
<td>12.6 × 10⁻⁶</td>
<td>------</td>
<td>------</td>
<td>------</td>
<td>------</td>
</tr>
</tbody>
</table>

²⁰³ The Swedish Civil Contingencies Agency, personal communication
²⁰⁴ Pettersson et al., 1996.
Table 6-20. Number of hours of landfill fires and estimated Hg, dioxin and PAH emissions in Sweden 1996 – 2016.

<table>
<thead>
<tr>
<th>Year</th>
<th>Surface fire, no. of hours</th>
<th>Underground fire, no. of hours</th>
<th>Hg emissions, kg</th>
<th>Dioxin emissions, g</th>
<th>Total PAH emissions, kg</th>
</tr>
</thead>
<tbody>
<tr>
<td>1996</td>
<td>1284</td>
<td>966</td>
<td>1.7</td>
<td>0.27</td>
<td>0.62</td>
</tr>
<tr>
<td>1997</td>
<td>1108</td>
<td>1772</td>
<td>1.5</td>
<td>0.24</td>
<td>0.53</td>
</tr>
<tr>
<td>1998</td>
<td>654</td>
<td>1174</td>
<td>0.9</td>
<td>0.15</td>
<td>0.31</td>
</tr>
<tr>
<td>1999</td>
<td>733</td>
<td>2563</td>
<td>1.0</td>
<td>0.18</td>
<td>0.35</td>
</tr>
<tr>
<td>2000</td>
<td>969</td>
<td>717</td>
<td>1.3</td>
<td>0.20</td>
<td>0.46</td>
</tr>
<tr>
<td>2005</td>
<td>1799</td>
<td>1506</td>
<td>2.4</td>
<td>0.38</td>
<td>0.86</td>
</tr>
<tr>
<td>2006</td>
<td>1983</td>
<td>2783</td>
<td>2.7</td>
<td>0.43</td>
<td>0.95</td>
</tr>
<tr>
<td>2007</td>
<td>1683</td>
<td>2695</td>
<td>2.3</td>
<td>0.37</td>
<td>0.81</td>
</tr>
<tr>
<td>2008</td>
<td>1680</td>
<td>3110</td>
<td>2.3</td>
<td>0.38</td>
<td>0.81</td>
</tr>
<tr>
<td>2009</td>
<td>1540</td>
<td>1143</td>
<td>2.0</td>
<td>0.32</td>
<td>0.74</td>
</tr>
<tr>
<td>2010</td>
<td>1032</td>
<td>1604</td>
<td>1.4</td>
<td>0.23</td>
<td>0.50</td>
</tr>
<tr>
<td>2011</td>
<td>1574</td>
<td>922</td>
<td>2.1</td>
<td>0.33</td>
<td>0.76</td>
</tr>
<tr>
<td>2012</td>
<td>1043</td>
<td>1218</td>
<td>1.4</td>
<td>0.22</td>
<td>0.50</td>
</tr>
<tr>
<td>2013</td>
<td>1542</td>
<td>2782</td>
<td>2.1</td>
<td>0.34</td>
<td>0.74</td>
</tr>
<tr>
<td>2014</td>
<td>494</td>
<td>414</td>
<td>0.7</td>
<td>0.10</td>
<td>0.24</td>
</tr>
<tr>
<td>2015</td>
<td>460</td>
<td>541</td>
<td>0.6</td>
<td>0.10</td>
<td>0.22</td>
</tr>
<tr>
<td>2016</td>
<td>507</td>
<td>930</td>
<td>0.7</td>
<td>0.11</td>
<td>0.24</td>
</tr>
</tbody>
</table>

6.6.2.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions of DIOX, Hg are based on expert judgement and they are ± 1000%, and ± 100%, respectively.

More information is given in IIR Annex 1.

Time series for landfill fires reported in NFR code 5E have been reviewed in later years and considered to be consistent.

6.6.2.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC has been performed.

6.6.2.5 SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculations have been performed.

6.6.2.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.6.3 Other waste, house and car fires, NFR 5E
6.6.3.1 SOURCE CATEGORY DESCRIPTION
Since submission 2016 also emissions of PM$_{2.5}$, PM$_{10}$, TSP, Pb, Cd, Hg, As, Cr, Cu and dioxin from House/car fires are included in 5E (1990 and onwards).
6.6.3.2 METHODOLOGICAL ISSUES

Emissions of PM$_{2.5}$, PM$_{10}$, TSP, Pb, Cd, Hg, As, Cr, Cu and dioxin from house and car fires are reported for the first time in submission 2016. The emissions are based on the frequency and duration of fires in Sweden$^{205}$ and emission factors (Tier 2) from EMEP/EEA Guidebook 2013. Used activity data are: car fires, fires in detached and undetached house fires, apartment building fires and industrial building fires. Statistics for 1990-1997 are missing, and therefore the average amount of fires for 1998-2003 is used instead. In Table 6-29, the emission factors used are presented and in Table 6-18, the reported emissions of TSP, PM, metals and dioxin emissions from house and car fires are presented. In submission 2018 data for 2016 could not be available in time an average for 2011-2015 have been used. This will be adjusted in submission 2019.

Table 6-21. Emission factors used for estimation of TSP, PM, metals and dioxin emissions from house and car fires.

<table>
<thead>
<tr>
<th>Fire category</th>
<th>TSP kg/fire</th>
<th>PM$_{10}$ kg/fire</th>
<th>PM$_{2.5}$ kg/fire</th>
<th>Pb g/fire</th>
<th>Cd g/fire</th>
<th>Hg g/fire</th>
<th>As g/fire</th>
<th>Cr g/fire</th>
<th>Cu g/fire</th>
<th>Dioxin mg/fire</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cars</td>
<td>2.3</td>
<td>2.3</td>
<td>2.3</td>
<td>NE</td>
<td>NE</td>
<td>NE</td>
<td>NE</td>
<td>NE</td>
<td>NE</td>
<td>0.048</td>
</tr>
<tr>
<td>Detached house</td>
<td>143.82</td>
<td>143.82</td>
<td>143.82</td>
<td>0.42</td>
<td>0.85</td>
<td>0.85</td>
<td>1.35</td>
<td>1.29</td>
<td>2.99</td>
<td>1.44</td>
</tr>
<tr>
<td>Undetached house</td>
<td>61.62</td>
<td>61.62</td>
<td>61.62</td>
<td>0.18</td>
<td>0.36</td>
<td>0.36</td>
<td>0.58</td>
<td>0.55</td>
<td>1.28</td>
<td>0.62</td>
</tr>
<tr>
<td>Apartment building</td>
<td>43.78</td>
<td>43.78</td>
<td>43.78</td>
<td>0.13</td>
<td>0.26</td>
<td>0.26</td>
<td>0.41</td>
<td>0.39</td>
<td>0.91</td>
<td>0.44</td>
</tr>
<tr>
<td>Industrial building</td>
<td>27.23</td>
<td>27.23</td>
<td>27.23</td>
<td>0.08</td>
<td>0.16</td>
<td>0.16</td>
<td>0.25</td>
<td>0.24</td>
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<th>TSP kt</th>
<th>PM$_{10}$ kt</th>
<th>PM$_{2.5}$ t</th>
<th>Pb t</th>
<th>Cd t</th>
<th>Hg t</th>
<th>As t</th>
<th>Cr t</th>
<th>Cu t</th>
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<td>2009</td>
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<td>2015</td>
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6.6.3.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions from house and care fires reported in NFR 5E are based on expert judgement. Uncertainties for emissions for TSP, PM$_{10}$ and PM$_{2.5}$ are based on data from the Guidebook 2013 and they are ± 100% each. Uncertainties for emissions of NH$_3$, DIOX and metals are based on expert judgement. Assessed uncertainties for NH$_3$, DIOX, Cd, Hg, As, Cr, Cu and Pb are ± 200%, ± 100%, 100%, ± 100%, ± 100%, ± 100% and ± 110%, respectively. More information is given in IIR Annex 1.

Time series for house and car fires reported in NFR code 5E have been reviewed in later years and considered to be consistent.

6.6.3.4 SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC has been performed.

6.6.3.5 SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculations have been performed. In submission 2018 data for 2016 could not be available in time an average for 2011-2015 have been used. This will be adjusted in submission 2019.

6.6.3.6 SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.

6.6.4 Other waste, pets, NFR 5E
6.6.4.1 SOURCE CATEGORY DESCRIPTION
Emissions of NH$_3$ from cats and dogs are also included in NFR 5E Other waste (1990 and onwards).

6.6.4.2 METHODOLOGICAL ISSUES
The estimates of emissions of ammonia from cats and dogs are based on a calculation made in the beginning of the nineties and the same value (0.5 kt NH$_3$) has been used for the whole time period from 1990$^{206}$. The calculation is based on data on the number of cats and dogs and an estimated value on the amount of emissions from cats and dog relative to emissions from humans.

6.6.4.3 UNCERTAINTIES AND TIME-SERIES CONSISTENCY
Uncertainties for emissions of NH$_3$ from cats and dogs are based on expert judgement. Assessed uncertainties for NH$_3$ are ± 200%. More information is given in IIR Annex 1.

Time series for pets reported in NFR code 5E have been reviewed in later years and considered to be consistent.

6.6.4.4  SOURCE-SPECIFIC QA/QC AND VERIFICATION
No source-specific QA/QC has been performed.

6.6.4.5  SOURCE-SPECIFIC RECALCULATIONS
No source-specific recalculations are performed.

6.6.4.6  SOURCE-SPECIFIC PLANNED IMPROVEMENTS
No major improvements are planned for the next submission.
7 Other (NFR sector 6)

No other sectors are included in the Swedish emission inventory, the sector is reported “Not occurring”.

8 Recalculations and Improvements

In this submission, recalculations are explained under each sector and NFR-code.

As requested by the EEA, a copy of table 3 of the 2017 Review Report with recommendations from TERT and comments on the status of implementation is attached.

Table 8-1. Recommendations from TERT, considering revised estimates (RE) and technical corrections (TC).

<table>
<thead>
<tr>
<th>Observation</th>
<th>Key Category</th>
<th>NFR, Pollutant(s), Year(s)</th>
<th>Recommendation</th>
<th>RE or TC</th>
<th>Comment on implementation and reference to IIR section/chapter (if relevant)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SE-1A3b-2017-0002</td>
<td>Yes</td>
<td>1A3b Road transport, SO$_2$, NO$_x$, NH$<em>3$, NMVOC, PM$</em>{2.5}$, 1990-2015</td>
<td>For 1A3b Road Transport, the TERT noted that the use of alternative fuels has been described in the IIR, but activity data were not reported in the NFR tables. In response to a question raised during the review, Sweden clarified that emissions from the use of gaseous fuels and biomass were included in the NFR tables but notation keys were incorrectly used to report activity data for these fuels. The TERT recommends that Sweden reports these activity data in the next submission and Sweden has also indicated such plan.</td>
<td>no</td>
<td>This has been fixed. The activity data for alternative fuels are included in the NFR tables in submission 2018.</td>
</tr>
<tr>
<td>SE-1A3b-2017-0003</td>
<td>Yes</td>
<td>1A3b Road transport, SO$_2$, NO$_x$, NH$<em>3$, NMVOC, PM$</em>{1.5}$, 1990-2015</td>
<td>For 1A3b Road Transport, the TERT noted that there is a lack of transparency in the IIR regarding whether cold start emissions and the impact of emission degradation have been taken into account when estimating 1A3b emissions. In response to a question raised during the review, Sweden clarified that the HBEFA model has been used to estimate emissions from road traffic and both cold start emissions and the impact of emission degradation is included in the model. For comparability and transparency purposes, the TERT recommends that Sweden includes descriptions of methods as well as the relevant input assumptions used to take account of cold start emissions and the impact of emission degradation in future IIRs.</td>
<td>no</td>
<td>The method used in HBEFA to estimate emissions from cold starts is based on ARTEMIS 4. The next version of HBEFA, which will be available in the fall of 2018, will be based on ARTEMIS 5. Sweden cannot adjust the emission</td>
</tr>
<tr>
<td>Observation</td>
<td>Key Category</td>
<td>NFR, Pollutant(s), Year(s)</td>
<td>Recommendation</td>
<td>RE or TC</td>
<td>Comment on implementation and reference to IIR section/chapter (if relevant)</td>
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<tr>
<td>SE-1A3b-2017-0009</td>
<td>Yes</td>
<td>1A3b Road Transport, NH₃, 1990-2015</td>
<td>For category 1A3biii Road Transport: Heavy Duty Vehicles and Buses and pollutant NH₃ for years 2010 onwards, the TERT noted that in response to a question raised during the review Sweden did not provide a clear response to the trend of NH₃ implied emission factors (IEFs) for 1A3biii. The trend is not in line with expectation when compared with other Member States, whereby an upward trend is expected for recent years due to the penetration of Euro V and VI vehicles in the fleet with higher NH₃ emission factors. The TERT noted that the issue is below the threshold of significance for a technical correction. The TERT recommends that Sweden reviews its NH₃ emission factor assumptions for Euro V and VI heavy duty vehicles against the 2016 EMEP/EEA Guidebook and revises the estimates if necessary in the next submission.</td>
<td>no</td>
<td>The emission factor for NH₃ and 1A3biii is 0.003 mg/vehicle km for all euro classes and years. The emission factors will be adjusted in the next version of HBEFA, which will be available in the fall of 2018. Sweden cannot adjust the emission factors themselves. This is done on a central level by INFRAS and TUG.</td>
</tr>
<tr>
<td>SE-1A3bv-2017-0001</td>
<td>No</td>
<td>1A3bv Road transport: Gasoline evaporation, NMVOC, 1990-2015</td>
<td>For 1A3bv Road Transport: Gasoline Evaporation, the TERT notes that there is a lack of transparency in the IIR regarding the methodology used to estimate NMVOC emission factors themselves. This is done on a central level by INFRAS and TUG. There is some information about the method to estimate the emissions from cold starts under section “2.5.2.1 Vehicle mileages, loads, trip lengths and fuels” in IIR Annex.</td>
<td>no</td>
<td>The method used in HBEFA to estimate NMVOC from gasoline evaporation is</td>
</tr>
<tr>
<td>Observation</td>
<td>Key Category</td>
<td>NFR, Pollutant(s), Year(s)</td>
<td>Recommendation</td>
<td>RE or TC</td>
<td>Comment on implementation and reference to IIR section/chapter (if relevant)</td>
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<tr>
<td>SE-1A3ei-2017-0001</td>
<td>No</td>
<td>1A3ei Pipeline Transport, SO$_2$, NO$_x$, 2000-2015</td>
<td>For 1A3ei Pipeline Transport, the pollutants NO$_x$, SO$_2$, NMVOCs, NH$<em>3$ and PM$</em>{2.5}$ and the years 2000 - 2015, the TERT noted that in response to a question raised during the review, Sweden explained that it is a small source, but that they will look at estimating emissions for the next submission. The TERT noted that the issue is below the threshold of significance for a technical correction. The TERT recommends that emission estimates are included in the next submission.</td>
<td>no</td>
<td>This will be looked into in 2018 and if emission factors are available, it will implemented in submission 2019.</td>
</tr>
</tbody>
</table>

In response to a question raised during the review, Sweden explained the parameters used to estimate emissions from cold starts and evaporation for light duty vehicles using the HBEFA methodology. However, the TERT understands that the HBEFA methodology for estimating evaporative emissions is not as up to date as the 2013 or 2016 EMEP/EEA Guidebook. The TERT believes that Sweden is under estimating NMVOC emissions for 1A3bv and that a revision may have an impact on the total emissions that is above the threshold of significance. To illustrate the impact, the TERT made an attempt to model 1A3bv emissions for Sweden by running the COPERT 5 model with its own activity data for Sweden, and shared the results (SE_Copert5_1A3bv_v2.xlsx) with Sweden. The total petrol vehicles as assumed in the COPERT 5 model are in close agreement (<2% difference) with the total petrol vehicles provided by Sweden during the review (3 Antal SE_TrafficScenario_90_15_35_5_JUNI.XLSX). The emissions comparison shows that Sweden is potentially under estimating NMVOC emissions and the impact of a revision is close to the threshold of significance (1.5% to 1.8% of National Total). The TERT recommends that Sweden reviews and updates its methodology for estimating 1A3bv emission to be in line with the 2016 EMEP/EEA Guidebook in the next submission. Based on ARTEMIS 4. The next version of HBEFA, which will be available in the fall of 2018, will be based on ARTEMIS 5. Sweden cannot adjust the emission factors themselves. This is done on a central level by INFRAS and TUG.
<table>
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<th>Observation</th>
<th>Key Category</th>
<th>NFR, Pollutant(s), Year(s)</th>
<th>Recommendation</th>
<th>RE or TC</th>
<th>Comment on implementation and reference to IIR section/chapter (if relevant)</th>
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<tr>
<td>SE-1B1a-2017-0001</td>
<td>No</td>
<td>1B1a Fugitive Emissions from Solid Fuels: Coal Mining and Handling, PM$_{2.5}$, 2005-2015</td>
<td>For category 1B1a Fugitive Emissions from Solid Fuels: Coal Mining and Handling and pollutant PM$_{2.5}$ for all years the TERT noted that PM emissions from coal handling are not considered in this source category (notation key ‘NO’ was used). In response to a question raised during the review, Sweden explained that these emissions are reported in 1A1c Manufacture of Solid Fuel and Other Energy Industries together with emissions from peat production. The TERT agreed with the explanation. The TERT recommends that Sweden reports these emissions in category 1B1a in the next submission. If this is not possible, the TERT recommends using the notation key ‘IE’ (Included Elsewhere) and describe in the IIR where emissions from coal handling have been allocated.</td>
<td>no</td>
<td>Notation key NO will be replaced by IE in next submission and a description regarding the allocation of emissions will be included.</td>
</tr>
<tr>
<td>SE-1B1b-2017-0001</td>
<td>No</td>
<td>1B1b Fugitive Emissions from Solid Fuels: Solid Fuel Transformation, SO$_2$, NO$_x$, NMVOC, 2005-2015</td>
<td>For category 1B1b Fugitive Emissions from Solid Fuels: Solid Fuel Transformation and pollutants NO$_x$ and NMVOC and for all years the TERT noted that the emissions were reported as ‘NA’ (not applicable), while the 2016 EMEP/EEA Guidebook provides default emission factors for this source. This issue was also flagged during the CLRTAP Stage 3 review in 2013. In response to a question raised during the review, Sweden explained that in the Swedish emission inventory, emissions from coke oven gas combustion is the main source of NMVOC and NO$_x$ and that these emissions are reported in 1A1c Manufacture of Solid Fuel and Other Energy Industries. The TERT partly agreed to this explanation. Fugitive emissions are likely to occur as well, and therefore the TERT recommends that Sweden either quantifies these emissions, or in case that is not possible, replaces the notation key ‘NA’ with the notation key ‘IE’ and describe in the IIR how emissions have been allocated.</td>
<td>no</td>
<td>Fugitive emissions of NO$_x$ and NMVOC are calculated with default emission factors and included in Submission 2018. See section 3.3.2.2</td>
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<tr>
<td>SE-1B2c-2017-0001</td>
<td>No</td>
<td>1B2c Venting and Flaring (Oil, Gas, Combined Oil and Gas), SO$_2$, NO$<em>x$, NMVOC, PM$</em>{2.5}$, 2005-2015</td>
<td>For category 1B2c Venting and Flaring (Oil, Gas, Combined Oil and Gas) and pollutants SO$_2$, NO$<em>x$, NMVOC and PM$</em>{2.5}$, the TERT noted that there is a potential underestimate which is likely below the threshold of significance. The reason is that emissions from stationary combustion have been used to estimate emissions from flaring, as</td>
<td>no</td>
<td>Revision of emission factors for flaring will be further investigated within a new ongoing development</td>
</tr>
<tr>
<td>Observation</td>
<td>Key Category</td>
<td>NFR, Pollutant(s), Year(s)</td>
<td>Recommendation</td>
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<tr>
<td><strong>SE-2B5-2017-0001</strong></td>
<td>Yes</td>
<td>2B5 Carbide production, PM$_{2.5}$, 1990-2015</td>
<td>For category 2B5 Carbide Production the TERT noted that in response to a question raised during the review, Sweden explained that the high particle emissions reported for carbide production are due to flaring at the single Swedish production site for carbide. The TERT noted that flaring emissions are misallocated in this category. The TERT recommends that Sweden allocates emissions from flaring to category 2B10a Chemical Industry: Other to improve the comparability of category 2B5 to that of other Member States.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

In the last Stage 3 review under UNECE (2013), it was recommended by the ERT to review and update these emission factors. In response to a question raised during the review, Sweden explained that the use of the 2016 EMEP/EEA Guidebook Tier 1 approach would overestimate the emissions substantially, while the 2016 EMEP/EEA Guidebook Tier 2 approach is missing EFs for some pollutants. The TERT agreed with the explanation provided by Sweden. The TERT recommends that Sweden use the 2016 EMEP/EEA Guidebook Tier 2 method for pollutants where EFs are included, and estimate emissions of other relevant pollutants with county specific data where this is available. |

<table>
<thead>
<tr>
<th>Project</th>
<th>Flaring at the carbide production facility will be further investigated within a new ongoing development project. The efforts will be made to implement the results in Submission 2019.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>SE-2D3i-2017-0001</strong></td>
<td>Yes</td>
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</table>

EFs for 060412i (Not diluted) and 060412ii (Not diluted) are given in Table A3-5 in IIR Annex. EFs for raw materials are set to 0.001 for all SNAP codes as now mentioned in section 4.5.3.2. A table with data
<table>
<thead>
<tr>
<th>Observation</th>
<th>Key Category</th>
<th>NFR, Pollutant(s), Year(s)</th>
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<th>RE or TC</th>
<th>Comment on implementation and reference to IIR section/chapter (if relevant)</th>
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<tr>
<td>SE-3B4h-2017-0001</td>
<td>No</td>
<td>3B4h Manure Management - Other Animals, PM(_{2.5}), 2005, 2010, 2015</td>
<td>For category 3B4h Manure Management - Other Animals and PM(<em>{2.5}) emissions for years 2005, 2010 and 2015 the TERT noted that there may be an under estimate of emissions. The TERT noted that this under estimate may be because Sweden reports ‘NE’ for 3B4h Manure Management - Other animals although animal numbers are reported in the NFR. The EMEP/EEA Guidebook 2016 provides default EFs for Other animals (Fur animals). The impact of the potential under estimate is probably below the threshold of significance. In response to a question raised during the review, Sweden explained that it estimated particulate matter (PM(</em>{2.5}), PM(_{10}) and TSP) from fur-bearing animals (minks) but due to a technical error during the transfer of data to the NFR-tables, these numbers were not included. The method used is described in the IIR. Sweden attached the complete time series and explained that it will ensure that these emissions are included in the NFR-tables for submission 2018. The TERT notes that these estimates indicate that any correction is below the threshold of significance and therefore does not need to be included as an official revised estimate for this review. The TERT recommends that Sweden improves its QA/QC procedures connected with the data transfer to NFR and that it includes estimates in its next submission as announced in its answer.</td>
<td>Resolved. Emissions of all PM fractions for minks are now reported in the NFR-tables.</td>
<td></td>
</tr>
<tr>
<td>SE-5C1bi-2017-0001</td>
<td>No</td>
<td>5C1bi Hazardous Waste Incineration, SO(_2), NO(<em>x), NMVOC, PM(</em>{2.5}), 2005;2010;2015</td>
<td>For NO(_x), SO(<em>2), and PM(</em>{2.5}) emissions from 5C1bi Hazardous Waste Incineration the TERT noted that in response to a question raised during the review, Sweden explained that it has not contacted the waste incineration plant with regard to the matter of adjusting reported data according to the IED directive. The TERT noted that the issue is far below the threshold of significance for a technical correction. The TERT recommends that Sweden engages with operators to make sure that there is no underestimation.</td>
<td>The company has not yet been contacted in response to the question from TERT.</td>
<td></td>
</tr>
<tr>
<td>Observation</td>
<td>Key Category</td>
<td>NFR, Pollutant(s), Year(s)</td>
<td>Recommendation</td>
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</tr>
<tr>
<td>SE-SD-2017-0001</td>
<td>No</td>
<td>SD Wastewater Handling, NH₃, 2005;2010;2015</td>
<td>For 5D1 Domestic wastewater handling the TERT noted that the default NH₃ emission factor proposed in the 2016 EMEP/EEA Guidebook, which relates to latrines and not to septic tanks, is applied to both septic tanks and latrines in the Swedish inventory. In response to a question raised during the review, Sweden explained that it will not be able to provide a reliable revised estimate for this source within the time frame of the review. The TERT decided to calculate a technical correction for the years 2005, 2010 and 2015 which was accepted by the Sweden. The estimates demonstrate that the issue is above the threshold of significance. The TERT recommends that Sweden includes a revised estimate in its next submission.</td>
<td></td>
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</tbody>
</table>

In emissions values estimated and reported by the operators on the basis of measurements, especially dealing with the subtraction of "the value of the confidence interval" (at a maxima, 20% of SO₂, 20% of NOₓ, 30% of dust) stated in the IED to determine the validated hourly average values of concentration.

TC TERT’s technical correction is used as an estimate for the source’s emissions of NH₃ until reliable results on AD from latrines are available.
No projections are reported in this Submission.
10 Reporting of gridded emissions and LPS

Gridded emission data gives information of the geographical distribution of emissions and is used for control purpose and serves as an important input data for atmospheric dispersion modelling.

10.1 Scope

Swedish gridded emissions were last reported to UNECE in 2017. The years 1990, 2000, 2005, 2010 and 2015 were included in the submission for the following components: NO$_x$, NMVOC, SO$_x$, NH$_3$, PM$_{2.5}$, PM$_{10}$, BC, CO, Pb, Cd, Hg, PCDD/PCDF (dioxins/furans), PAHs, HCB and PCB.

The gridded emissions were aggregated into GNFR-sectors: A_PublicPower, B_Industry, C_OtherStationaryComb, E_Solvents, F_RoadTransport, G_Shipping, H_Aviation, I_Offroad, J_Waste, K_AgriLivestock and L_AgriOther. Due to data confidentiality it has not been possible to report the GNFR-sector D_Fugitive as a separate sector; these emissions have been included in the sector B_Industry.

The new EMEP grid resolution of 0.1° × 0.1°, introduced in the 2014 Reporting Guidelines, was implemented in the last reported gridded emissions in 2017. The previous gridded emissions reported in 2012 followed the old EMEP grid resolution of 50 km × 50 km.

10.2 Methodology

Both top-down and bottom-up approaches are used in order to calculate gridded emissions in Sweden. The methodology to allocate emissions into grid cells is schematically described below:

- For some emissions sources, the emissions are allocated with a bottom-up approach:
  - Emissions from large point sources are allocated directly into the appropriate grid cells.
  - Emissions from road transport are allocated to grid cells using a national road database (NVDB) containing the Swedish road network, vehicle kilometers and vehicle compositions.
  - Emissions from national shipping are allocated to grid cells with a system that uses AIS-data.

- For other emission sources where the emissions are not known for every grid cell, a top-down approach is used:
For many emission sources, activity data (or surrogate data) on an aggregated geographical level (for example municipality level) is used together with geographical data, in order to create a proxy. For example, emissions from civil aviation (LTO) are allocated with a proxy containing the numbers of LTOs per Swedish airport as activity data and the locations of the airports as the geographical data. Another example is emissions from manure management that are allocated with a proxy containing the numbers of animals per municipality as activity data and agriculture areas as the geographical data.

For some emission sources, activity data is only available on national level. For such emissions, only geographical data is used as a proxy. One example is emissions of NH$_3$ from pets; these emissions are allocated with population density as a proxy/surrogate data.

A detailed description of the methodology and the quality of the gridded emissions for each sector is given in Andersson et al. (2017).

10.3 Recent improvements

A summary of the major improvements of the gridded emissions reported in 2017, in comparison with gridded emissions reported in 2012, is given below:

- The new EMEP grid resolution of $0.1^\circ \times 0.1^\circ$ has been implemented.
- Three new components have been included; BC, HCB and PCB.
- The methodology used for gridding emissions from public power and stationary combustion in manufacturing industries has been improved. Information from registers has been included in the geographical data. This enables improved distribution of emissions for companies with facilities in two or more Swedish municipalities.
- The database used for gridding of emissions from road transport has been updated with a new year (2015).
- The methodology to allocate emissions from small-scale stationary residential combustion (aggregated in the GNFR-sector C_OtherStationaryComb) has been improved. For the reported emissions in 2012, emissions were calculated in four temperature zones in Sweden; within each temperature zone, emissions were distributed over the total small house areas [m$^2$] per grid cell. The improved gridded emissions reported in 2017 have been calculated with activity data on municipality level (number of boilers/stoves) with energy need calculated on country level. Within each municipality, the emissions are distributed over the total small house areas [m$^2$] per grid cell, also taking into account the availability of district heating in every grid cell.
• The proxy used for gridding emissions from mobile combustion in manufacturing industries and construction has been improved. Emissions from working machineries at airports are gridded using the total number of LTO for each Swedish airport as surrogate data. Emissions from working machineries on railways are distributed equally over the railway network in Sweden.

• Facility specific data is now used for allocating emissions from lime production to grid cells (these emissions are aggregated in the GNFR sector B_Industry).

• In the GNFR sector J_Waste, two new emission sources have been included; house and car fires and carcasses cremation. Emissions from house and car fires are gridded using a proxy with the number of house/car fires per Swedish municipality. For carcasses cremation, activity data is only available on national level, thus the emissions are equally distributed over agriculture areas.

• Consistent activity/surrogate data is now used in the whole time series for the GNFR sectors K_AgriLivestock and L_AgriOther.

10.4 Planned improvements

Potential major improvements for the next reported gridded Swedish emissions (in 2021) are listed below:

• The methodology for gridding emissions from national and international aviation (LTO) will most likely be improved by using DS-B, MLAT and radar data.

• Emissions from national shipping might be updated with an improved system for AIS-data.

• The database used for gridding emissions from road transport will be updated with a new year.
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