

# **CHAPTER 6**

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## **QUALITY ASSURANCE/QUALITY CONTROL AND VERIFICATION**

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## 6 QUALITY ASSURANCE/QUALITY CONTROL AND VERIFICATION

*Users are expected to go to Mapping Tables in Annex 1, before reading this chapter. This is required to correctly understand both the refinements made and how the elements in this chapter relate to the corresponding chapter in the 2006 IPCC Guidelines.*

### 6.1 INTRODUCTION

An important goal of IPCC inventory guidance is to support the development of national greenhouse gas inventories that can be readily assessed in terms of quality. It is *good practice* to implement quality assurance/quality control (QA/QC) and verification procedures in the development of national greenhouse gas inventories to accomplish this goal. The procedures as described in this chapter also serve to drive inventory improvement.

The guidance is designed to achieve practicality, acceptability, cost-effectiveness, incorporation of existing experience, and the potential for application on a worldwide basis. A QA/QC and verification system contributes to the objectives of *good practice* in inventory development, namely to improve transparency, consistency, comparability, completeness, and accuracy of national greenhouse gas inventories.

QA/QC and verification activities should be integral parts of the inventory process (see Section 1.6.3 of Chapter 1, Volume 1). The outcomes of QA/QC and verification may result in:

- improvements in the estimates of emissions and/or removals;
- reassessment of inventory compilation processes and category uncertainty estimates.

For example, the results of the QA/QC process may point to particular variables within the estimation methodology for a certain category that should be the focus of improvement efforts.

The terms ‘quality control’, ‘quality assurance’, and ‘verification’ are often used in different ways. The definitions of QC, QA, and verification in Box 6.1 will be used for the purposes of this guidance.

**BOX 6.1 (UPDATED)**  
**DEFINITIONS OF QA/QC AND VERIFICATION**

*Quality Control (QC)* is a system of routine technical activities to assess and maintain the quality of the inventory as it is being compiled. Personnel compiling the inventory perform it. The QC system is designed to:

- (i) Provide routine and consistent checks to ensure data integrity, correctness, and completeness;
- (ii) Identify and address errors and omissions;
- (iii) Document and archive inventory material and record all QC activities.

QC activities include general methods such as accuracy checks on data acquisition and calculations, and the use of approved standardised procedures for emission and removal calculations, measurements, estimating uncertainties, archiving information and reporting. QC activities also include technical reviews of categories, activity data, emission factors, other estimation parameters, and methods.

*Quality Assurance (QA)* is a planned system of review procedures conducted by personnel not directly involved in the inventory compilation/development process. Reviews, preferably by independent third parties, are performed upon a completed inventory following the implementation of QC procedures. Reviews verify that measurable objectives (data quality objectives, see Section 6.5, QA/QC Plan) were met, ensure that the inventory represents the best possible estimates of emissions and removals given the current state of scientific knowledge and data availability, and support the effectiveness of the QC programme.

*Verification* refers to the collection of activities and procedures conducted during the planning and development, or after completion of an inventory that can help to establish its reliability for the intended applications of the inventory. For the purposes of this guidance, verification refers specifically to those methods that are external to the inventory and apply independent data, including comparisons with inventory estimates made by other bodies or through alternative methods. Verification activities may be constituents of both QA and QC, depending on the methods used and the stage at which independent information is used. It is important to distinguish verification, as defined by the IPCC guidelines, from the term verification used in carbon markets, which is synonymous with an independent audit. Such an audit would fall under the scope of a QA procedure in the terminology of the IPCC Guidelines. For example, under the Kyoto Protocol Clean Development Mechanism (CDM) verification is defined as the periodic independent review and *ex post* determination by an auditing body of monitored reductions in anthropogenic emissions by sources of GHGs that have occurred as a result of a registered CDM project activity during the verification period. Verification has different meanings in different contexts, and in the case of carbon markets, results-based payments, etc. has a meaning more similar to the QA definition above.

Before implementing QA/QC and verification activities, it is necessary to determine which techniques should be used, and where and when they will be applied. QC procedures may be *general* with a possible extension to *category specific* procedures. There are technical and practical considerations in making these decisions. The technical considerations related to the various QA/QC and verification techniques are discussed in general in this chapter, and specific applications to categories are described in the category-specific guidance in Volumes 2 to 5. The practical considerations involve assessing national circumstances such as available resources and expertise, and the particular characteristics of the inventory (e.g., whether or not a category is *key*).

## 6.2 PRACTICAL CONSIDERATIONS IN DEVELOPING QA/QC AND VERIFICATION SYSTEMS

No refinement.

## 6.3 ELEMENTS OF A QA/QC AND VERIFICATION SYSTEM

No refinement.

## 6.4 ROLES AND RESPONSIBILITIES

No refinement.

## 6.5 QA/QC PLAN

No refinement.

## 6.6 GENERAL QC PROCEDURES

No refinement.

## 6.7 CATEGORY-SPECIFIC QC PROCEDURES

No refinement.

### 6.7.1 Emissions factor QC

No refinement.

#### 6.7.1.1 IPCC DEFAULT EMISSION FACTORS

No refinement.

#### 6.7.1.2 COUNTRY-SPECIFIC EMISSION FACTORS

No refinement.

#### 6.7.1.3 DIRECT EMISSION MEASUREMENTS

No refinement.

### 6.7.2 Activity data QC

No refinement.

#### 6.7.2.1 NATIONAL LEVEL ACTIVITY DATA

Following are fundamental QC checks that should be considered for assessing the quality of national level activity data. In all cases, it is important to have a well-defined and documented data set from which appropriate checks can be developed.

***QC checks of reference source for national activity data:*** When using national activity data from secondary data, it is *good practice* for the inventory compiler to evaluate and document the associated QA/QC activities. This is particularly important with regard to activity data, since most activity data are originally prepared for purposes other than as input to estimates of greenhouse gas emissions. Many statistical organisations, for example, have their own procedures for assessing the quality of the data independently of what the end use of the data may be.

The inventory compiler should determine if the level of QC associated with secondary activity data includes, at a minimum, those QC procedures listed in Table 6.1 of the *2006 IPCC Guidelines*. In addition, the inventory compiler may check for any peer review of the secondary data and document the scope of this review. If the QA/QC associated with the secondary data is adequate, then the inventory compiler can simply reference the data source and document the applicability of the data for use in its estimates (see Box 6.3 for an example of this procedure).

If the QC associated with the secondary data is inadequate or if the data have been collected using standards/definitions that deviate from this guidance, then the inventory compiler should establish QA/QC checks on the secondary data. The uncertainty of estimates should be reassessed in the light of the findings. The inventory compiler should also reconsider how the data are used and whether any alternative data and international data sets may provide a better estimate of emissions or removals. If no alternative data sources are available, the inventory compiler should document the inadequacies associated with the secondary data QC as part of its summary report on QA/QC.

#### Box 6.3

##### EVALUATION OF DATA QUALITY ON EXTERNAL DATA IN THE TRANSPORTATION SECTOR

Countries typically use either fuel usage or kilometre (km) statistics to develop emissions estimates. The national statistics on fuel usage and km travelled by vehicles are usually prepared by a specialised agency. However, it is the responsibility of the inventory compiler to determine which QA/QC activities were implemented by the agency that prepared the original fuel usage and km statistics for vehicles. Questions that may be asked in this context are:

- Does the statistical agency have a QA/QC plan that covers the collection and handling of the data?
- Was an adequate sampling protocol used to collect data on fuel usage or km travelled?
- How recently was the sampling protocol reviewed?
- Has any potential bias in the data been identified by the statistical agency?
- Has the statistical agency identified and documented uncertainties in the data?
- Has the statistical agency identified and documented errors in the data?

**Comparisons with independently compiled data sets:** Where possible, a comparison check of the national activity data with independently compiled activity data sources should be undertaken. For example, many of the agricultural source-categories rely on government statistics for activity data such as livestock populations and production by crop type. Comparisons can be made to similar national statistics disseminated via FAOSTAT<sup>1</sup> by the United Nations Food and Agriculture Organization (FAO). Similarly, the International Energy Agency (IEA) maintains a database on national energy production and usage that can be used for checks in the energy. Industry trade associations, university research, and scientific literature are also possible sources of independently derived activity data to use in comparison checks. Activity data may also derive from balancing approaches – see Section 6.7.2.2 of the *2006 IPCC Guidelines* for a description and an example. As part of the QC check, the inventory compiler should ascertain whether alternative activity data sets are really based on independent data. International information is often based on national reporting which is not independent from the data used in the inventory. Available scientific or technical literature may also be used for a national inventory. In some cases, the same data are treated differently by different agencies to meet varying needs. Comparisons may need to be made at a regional level or with a subset of the national data since many alternative references for such activity data have limited scope and do not cover the entire nation.

**Comparisons with samples:** The availability of partial data sets at sub-national levels may provide opportunities to check the reasonableness of national activity data. For example, if national production data are being used to calculate the inventory for an industrial category, it may also be possible to obtain plant-specific production or capacity data for a subset of the total population of plants. Extrapolation of the sample production data to a national level can then be done using a simple approximation method. The effectiveness of this check

<sup>1</sup> <http://www.fao.org/faostat>.

depends on how representative the sub-sample is of the national population, and how well the extrapolation technique captures the national population.

**Trend checks of activity data:** National activity data should be compared with previous year's data for the category being evaluated. *Activity* data for most categories tend to exhibit relatively consistent changes from year to year without sharp increases or decreases. If the national activity data for any year diverge greatly from the historical trend, they should be checked for errors. If a calculation error is not detected, the reason for the sharp change in activity should be confirmed and documented. A more thorough approach to take advantage of similarities between years has been described in Chapter 5, Time Series Consistency.

### 6.7.2.2 SITE-SPECIFIC ACTIVITY DATA

No refinement.

### 6.7.3 Calculation-related QC

No refinement.

## 6.8 QA PROCEDURES

No refinement.

## 6.9 QA/QC AND UNCERTAINTY ESTIMATES

No refinement.

## 6.10 VERIFICATION

No refinement.

### 6.10.1 Comparisons of national estimates

There are a number of practical verification techniques that do not require specialised modelling expertise or extended analyses. Most of these can be considered as method-based comparisons that consider the differences in national estimates based on using alternative estimation methodologies for the same category or set of categories. These comparisons look for major calculation errors and exclusion of major source categories or sub-source categories. Method-based comparisons can be designed around the multi-tier level of methods outlined for each category in the sector guidance, through comparisons to independent estimates developed by other institutions, and, to a limited extent, through cross-country comparisons. The choice of method will depend on the method used in the inventory, a clear definition and correlation of categories between methods, and the availability of alternative data.

These checks can be extremely useful in confirming the reasonableness of national inventory estimates and may help identify any gross calculation errors. Some of these techniques, such as the compilation of the reference approach for Energy Sector estimates, should be considered as part of the inventory development process.

Discrepancies between inventory data and data compiled using alternative methods do not necessarily imply that the inventory data are in error. When analysing discrepancies, it is important to consider that there may be large uncertainties associated with the alternative calculations themselves.

**Applying lower tier methods:** Lower tier IPCC methods typically are based on 'top-down' approaches that rely on highly aggregated data at a summary category level. Inventory compilers using higher tier, 'bottom-up' approaches may consider using comparisons to lower-tier methods as a simple verification tool. As an example, for carbon dioxide (CO<sub>2</sub>) from fossil fuel combustion, a reference calculation based on apparent fuel consumption per fuel type is specified as a verification check in the Energy Sector procedures (see Volume 2: Energy). As an additional example, since 2014 the EU performs annually a full QA of its EU-28 GHG Inventories for agriculture, using the FAOSTAT emissions estimates for verification<sup>2</sup>. This reference approach estimate can be compared to the sum of sectoral-based estimates from a Tier 1, 2, or 3 approach. While the quality of the reference approach is

<sup>2</sup> Data for agriculture and land use accessible at: <http://www.fao.org/faostat/en/?#data/GT>.

typically lower than that of the sectoral approach, it remains useful as a simple approximation method. It is less sensitive to errors due to its simplicity and can be used as a top-down completeness check. Another example, where emissions are calculated as the sum of sectoral activities based on the consumption of a specific commodity, e.g., fuels or products like hydrofluorocarbons (HFCs), perfluorocarbons (PFCs) or sulphur hexafluoride (SF<sub>6</sub>), the emissions could be estimated using apparent consumption figures, e.g., national total production + import – export ± stock changes, taking into consideration any possible time lags in actual emissions.

Similar checks can be performed for industrial type sources, e.g., nitrous oxide estimates for nitric acid production and adipic acid where inventory estimates were determined for each individual production plant based on plant-specific data. The check of emission estimates would consist of the comparison between the sum of the individual plant-level emission estimates and a top-down emission estimate based on national nitric acid production figures and IPCC default Tier 1 factors. Large differences do not necessarily indicate that there are problems with the inventory estimate. As lower tier methods typically rely on more highly aggregated data, there may be relatively large uncertainties with the Tier 1 approach compared to an inventory estimated using a bottom up approach based on *good practice*. If differences cannot easily be explained, the inventory compiler may consider the following questions in any further QA/QC checks:

- Are there inaccuracies associated with any of the individual plant estimates (e.g., an extreme outlier may be accounting for an unreasonable quantity of emissions)?
- Are the plant-specific emission factors significantly different from each other?
- Are the plant-specific production rates consistent with published national level production rates?
- Is there any other explanation for a significant difference, such as the effect of controls, the manner in which production is reported or possibly undocumented assumptions?

This is an example of how the results of a relatively simple emission check can lead to a more intensive investigation of the representativeness of the emissions data. Knowledge of the category is required to isolate the parameter that is causing the difference in estimates and to understand the reasons for the difference.

**Applying higher tier methods:** Higher tier IPCC methods typically are based on detailed ‘bottom-up’ approaches that rely on highly disaggregated data and a well-defined subcategorization of sources and sinks. Inventory compilers may find that they cannot fully implement a higher tier approach because they are lacking sufficient data or resources. However, the availability of even partial estimates for a subcategory of sources may provide a valuable verification tool for the inventory. An estimate based on higher tier data derived from a proportion of the total sources in a country can be extrapolated to the national level, provided that the sample is representative. Such an extrapolation can be used to corroborate the national estimate.

**Comparisons with independently compiled estimates:** Comparisons with other independently compiled inventory data on national level (if available) are useful options to evaluate completeness, assess approximate emission (removal) levels and correct category allocations. Although the inventory compiler is ultimately responsible for preparing the national greenhouse gas inventory, other independent publications on this subject may be available e.g., from scientific literature or publication by other institutes or agencies. For example, national level CO<sub>2</sub> emissions estimates associated with the combustion of fossil fuel are compiled by the IEA<sup>3</sup>, the Carbon Dioxide Information and Analysis Centre (CDIAC)<sup>4</sup>, the Emission Database for Global Atmospheric Research (EDGAR)<sup>5</sup> and by British Petroleum (BP)<sup>6</sup>. Likewise, FAO compiles and disseminates national-level CO<sub>2</sub> and non-CO<sub>2</sub> emissions and removals for AFOLU (<http://www.fao.org/faostat>), using underlying national statistics as activity data. Estimates of emissions of other gases are available from the EDGAR, Regional Emission inventory in Asia (REAS)<sup>7</sup>, and US Environment Protection Agency (EPA)<sup>8</sup>. The World Resources Institute (WRI)<sup>9</sup> combines data from several sources mentioned in this section to provide sector-specific emission estimates. Use of multiple data sources in the comparison is advantageous as the data show differences between datasets, even for relatively well-known emissions of carbon dioxide (Ciais et al. 2010). Additional, independently compiled data for use in national greenhouse gas inventory may sometimes be found in national accounts, specifically those

<sup>3</sup> <https://www.iea.org/statistics>.

<sup>4</sup> <https://cdiac.ess-dive.lbl.gov> (doi:10.3334/CDIAC/00001\_V2017).

<sup>5</sup> <http://edgar.jrc.ec.europa.eu>.

<sup>6</sup> <https://www.bp.com/en/global/corporate/energy-economics/statistical-review-of-world-energy/co2-emissions.html>.

<sup>7</sup> <https://www.nies.go.jp/REAS/>.

<sup>8</sup> <https://www.epa.gov/ghgemissions/global-greenhouse-gas-emissions-data>.

<sup>9</sup> <http://cait.wri.org>.

developed under the UN System of Environmental and Economic Accounts (SEEA). If independently compiled datasets use IPCC Tier 1 methodologies, the same considerations discussed above will apply.

While national data are normally considered more reliable as they are able to accommodate more detailed country-specific information, and international data are normally compiled at a lower tier, these international data sets provide a good basis for comparison, as they are consistent between countries. Additionally, databases from international agencies such as IEA and FAO, use as activity data the underlying national statistics, providing enhanced opportunities for QA analysis. Furthermore, the FAOSTAT Tier 1 AFOLU estimates are available together with the corresponding UNFCCC country data, and differences analysed in a dedicated "compare" section<sup>10</sup>. The comparisons can be made for different greenhouse gases at national, sectoral, category, and subcategory levels, as far as the differences in definitions enable them. Before conducting these types of comparisons, it is important to check the following items.

- Confirm that the underlying data for the independent estimate are not the same as that used for the inventory; a comparison is only meaningful if the data being compared are different.
- Determine if the relationships between the sectors and categories in the different inventories can be defined and matched appropriately.
- Account for the data quality (e.g., QA/QC system or review) and for any known uncertainties in the estimate used for the comparison to help interpret results.

***Comparisons of intensity indicators between countries:*** Emission (removal) intensity indicators, e.g., those commonly referred to as "Implied emission (removal) factors", may be compared between countries (e.g., emissions per capita, industrial emissions per unit of value added, transport emissions per car, emissions from power generation per kWh of electricity produced, emissions from dairy ruminants per tonne of milk produced). These indicators provide a preliminary check and verification of the order of magnitude of the emissions or removals. Different practices and technological developments as well as the varying nature of the source categories will be reflected in the emission intensity indicators. Thus, differences between countries need to be expected. However, these checks may flag potential anomalies at the country or sector level.

## 6.10.2 Comparisons with atmospheric measurements

### 6.10.2.1 INTRODUCTION TO EMISSION ESTIMATES BASED ON ATMOSPHERIC CONCENTRATION MEASUREMENTS

This section addresses the state of science for emission estimates based on atmospheric measurements and their application to comparison with national emission inventories. Since the *2006 IPCC Guidelines* were published, the most notable advances have been achieved in the application of inverse models of atmospheric transport for estimating emissions at the national scale. An increasing number of countries are considering applying such models.

An ideal condition for verification is the use of fully independent data as a basis for comparison. Measurements of atmospheric concentrations provide such datasets, and recent scientific advances allow using such data as a basis for emission modelling. The approach is particularly valuable as it can be largely independent of standard estimation method drivers, such as sector activity data and implied emission factors. The scale of such models can be designed around local, regional, or global boundaries and can provide information on either level or trends in emissions. Some brief examples of these techniques are provided in this section; however, further discussion and elaboration can be found in more comprehensive summaries on the use of these methods for inventory verification (Rypdal et al. 2005; Benkovitz 2001; Benjey and Middleton 2002; NACP 2002; Jacob et al. 2016; DeCola et al. 2018; Miller & Michalak 2017; Bergamaschi et al. 2018).

Atmospheric measurements are being used to provide useful quality assurance of the national greenhouse gas emission estimates (Manning et al. 2011; Fraser et al. 2014; Henne et al. 2016). Under the right measurement and modelling conditions (discussed further in this section), they can provide a perspective on the trends and magnitude of greenhouse gas (GHG) emission estimates that is largely independent of inventories. It should be recognized that the technical complexity as well as the limited application potential of atmospheric models to inventory verification, particularly at a national level, can restrict their utility to many inventory compilers. In addition, many of the techniques will require specialised modelling skills, combined with a sufficient number and distribution of measurement locations, proxy data and adequate modelling and computing resources, in order to appropriately correlate the atmospheric data back to the inventory for comparison, and be cost- and labour intensive. Depending

<sup>10</sup> <http://www.fao.org/faostat/en/#compare>.

on specific conditions, results may be only applicable to parts of a country, to groups of countries, or to specific categories or gases. Currently, the required analysis time typically extends beyond an inventory cycle, thus making these types of comparisons more applicable for long-term verification programs. In many cases, the uncertainties associated with the atmospheric models themselves may not be sufficiently quantified or may be too large for the model to be used effectively as a verification tool.

In contrast to the other methods described in this chapter, comparisons with atmospheric measurements are not established as a standard tool for verification to be applied by an inventory compiler. Still, considerable scientific progress in this area needs to be noted and inventory compilers may wish to take advantage of the potential of this approach, as it gives independent data for verification. If applicable, national inventory compilers may also consider joining forces with neighbouring countries, in cases when emission modelling from atmospheric measurement is more reliable for larger entities than countries. There is a difficulty of separating emissions from neighboring countries in the inverse modelling estimates, especially in case of sparse observation network. Despite the limitations given, there are a number of evolving techniques that deserve to be mentioned here:

**Inverse Modelling:** The concentrations of greenhouse gases in air samples are measured at monitoring sites and can be used to provide emission estimates by a technique known as inverse modelling. Inverse models calculate emission fluxes from concentration measurements and atmospheric transport models. For local and regional estimation, complex mathematical and statistical models are required together with continuous, or quasi-continuous, measurements that capture all pollution incidents. The source discrimination of air sampling-derived emissions requires highly precise and labour-intensive analysis, which may prevent the applicability of inverse modelling approaches to source-specific emissions verification. In contrast to national inventories, flux assessments from inverse modelling have problems with separating anthropogenic emissions from natural sources/sinks as well as international transport (Desjardins et al. 2018). Considering the limited monitoring network currently available for many of the greenhouse gases and the resulting uncertainties in the model results, inverse modelling is not being widely applied as a verification tool of national inventories. However, there is increasing scientific recognition for the potential of these techniques for both level and trend verification of national inventories.

Inverse modelling techniques are undergoing rapid development and are being applied now in comparisons with national inventory estimates (O'Doherty et al. 2003; Manning et al. 2011; Fraser et al. 2014; Henne et al. 2016), European emission estimates (Manning et al. 2003) and to provide geographical distributions of emissions within the European Union (Ryall et al. 2001). A useful measure of the utility of these techniques can be provided by comparison of the uncertainties between the calculated inventory estimates and the inverse model-derived estimates (Bergamaschi et al. 2004; Rypdal et al. 2005). For example, Henne et al. 2016 estimated Switzerland's methane emissions with 9% uncertainty. Brunner et al. 2017 estimated emissions of HFC-125 for four large European countries with uncertainties ranging from 9% to 23%. Where the uncertainty of the model results is larger than calculated inventory uncertainty, the results of the comparison should be treated with caution. Also, where the model results are significantly different from the inventory, this can point to missing sources or possibly large calculation errors.

Fluorinated gases and methane (CH<sub>4</sub>) are considered the most suitable greenhouse gases for which inverse modelling could provide verification of emission estimates (Rypdal et al. 2005, Bergamaschi et al. 2004). The fluorinated compounds are considered good candidates for inverse modelling verification because: they have virtually no natural source interference in the atmospheric measurements, there can be considerable uncertainties in inventory methods, they are long-lived, and the loss mechanisms are well known. Despite of influences by natural sources, methane is considered a favourable candidate because of the generally high uncertainty in emission estimates resulting from inventory methodologies, and the strong atmospheric signal to noise ratio of measurements. Modelling of CO<sub>2</sub> emissions for national inventory verification is more difficult since the inventory methods already have low uncertainties, except where agriculture, forestry and other land-use is dominant. The impacts of large natural sources and sinks on atmospheric measurements make a correlation to strictly anthropogenic sources difficult. However, it may improve understanding of contributions from forests and natural sources and sinks. Due to the large uncertainties associated with some of the N<sub>2</sub>O inventory methodologies, verification through atmospheric measurements would be desirable. However, the influence of natural sources and sinks on measurements, as well as very long atmospheric lifetime lead to a poor signal to noise ratio in measured concentrations.

Inverse models calculate emissions by optimally combining concentration observations with an atmospheric transport model. In doing so, the inverse model must take into account estimates of uncertainty from both the observations and the atmospheric model. Flux assessments from inverse modelling necessarily include the contribution from all sectors (anthropogenic and natural sources/sinks) as well as international transport from country to country. As a result, it remains challenging to attribute estimated fluxes to specific source categories or regions using currently available sparse observation networks, which complicates the application of inverse modelling approaches for source-specific emissions verification (Miller & Michalak 2017). However, it is

expected to become less difficult in the future, with more dense observation networks (Pison et al. 2018), complemented by observations of radiocarbon, atmospheric potential oxygen (APO), and co-emitted tracers, such as carbon monoxide, that allows distinguishing the fossil CO<sub>2</sub> component (<sup>14</sup>C), carbon to hydrogen ratio of burned fuel (C/H ratio), and separating anthropogenically polluted air from natural background. Since <sup>14</sup>C is absent in fossil carbon, its measurements in CO<sub>2</sub> plumes give clear indication of fossil fuel originated fraction of the observed atmospheric CO<sub>2</sub> variation, thus observation of <sup>14</sup>C in CO<sub>2</sub> is considered as a powerful tool to distinguish between the fossil and biogenic CO<sub>2</sub> emission sources (Levin et al. 2003). Atmospheric potential oxygen is a proxy of the total oxygen ( $1.1\text{CO}_2 + \Delta(\text{O}_2/\text{N}_2)$ ) that is conserved during exchange with biosphere. As consumption of oxygen per unit of fuel carbon burned depends on fuel type, measurements of the atmospheric oxygen in the same time with CO<sub>2</sub> provide a tool for distinguishing burning hydrocarbons and natural gas from other CO<sub>2</sub> emission sources, such as coal combustion and biomass burning (Keeling 1988).

The quality of the derived emissions critically depends on the quality and quantity of measurements, and the quality of the gridded emission inventory and the atmospheric model, since inverse methods typically propagate estimated observation, inventory and model errors, the latter usually being the one of dominant components (Bergamaschi et al. 2018).

The most demanding, but proven, approach for verification through atmospheric measurements is establishment and operation of a national or regional/multi-national GHG observing network combined with inverse modelling and analysis (Andrews et al. 2014; Lopez-Coto et al. 2017; Bergamaschi et al. 2018). Despite the availability of inverse modelling tools, specialized training is required to apply them and obtain robust flux estimates that can be used to verify emission estimates from a greenhouse gas inventory. More details are presented in the Integrated Global Greenhouse Gas Information System (IG<sup>3</sup>IS) Science Implementation Plan prepared by the Global Atmosphere Watch (GAW) program of WMO, which documents *good practice* methodological guidelines for “how atmospheric measurements and analysis methods can deliver valuable information for inventory verification” (IG<sup>3</sup>IS Science Implementation Plan 2018). Operational verification systems already exist in the UK and Switzerland, where emission inventories for major non-CO<sub>2</sub> GHGs are verified annually and numbers are reported in the National Inventory Report to the UNFCCC. Another example of verification system based on inverse modelling is also in place in Australia.

At sub-national scales, such as city-, facility- and basin-scale, studies using regional atmospheric monitoring networks or targeted observation campaigns are being used for improving the knowledge about regional and facility level emissions and contributing to updating the emission factors for selected emission categories. These include the oil and gas sector, urban emissions, and emissions from agriculture, applying regional inverse modelling (Keller et al. 2011; Breon et al. 2015; McKain et al. 2015; Yver-Kwok et al. 2015; Lauvaux et al. 2016; Viatte et al. 2017), or mass-balance approaches (Zavala-Araiza et al. 2015; Conley et al. 2017).

**Continental Plumes:** A strong difference between source and non-source regions may generally be found between a continent and an ocean where routine measurements of the difference between background air concentrations and the offshore plume concentrations, coupled with wind vector analysis or trajectory analysis, may provide an indication of emissions on a broad scale (Cape et al. 2000; Derwent et al. 2001). For example, a number of greenhouse gases, including chlorofluorocarbons (CFCs), N<sub>2</sub>O and CH<sub>4</sub> from the European continental plume have been detected at Mace Head, Ireland. These results have then been used for subsequent quantification of the European emission source strength by inverse modelling (Derwent et al. 1998a, 1998b; Vermeulen et al. 1999).

**Use of Proxy Emission Databases:** In the cases where one of the components measured in the air samples has a well characterised emission inventory (a ‘marker’ or ‘tracer’ compound), the emissions of greenhouse gases may be estimated from atmospheric measurements of their concentration ratio to this marker compound. The technique is appropriate if sources of the compounds are co-located, and it has been used in the USA, for example with carbon monoxide (CO) as the marker (Barnes et al. 2003a, 2003b), and in the EU employing radon (<sup>222</sup>Rn; Biraud et al. 2000).

**Global Dynamic Approaches:** Trends over time in the atmospheric concentration of particular compounds may also indicate a change in the global balance between sources and sinks and give an estimate of the globally aggregated emissions, constraining the total of national emissions from an aggregate perspective and possibly indicating areas of weakness in the inventories. Such approaches have been taken for CH<sub>4</sub> (Dlugokencky et al. 1994; Saunio et al. 2016), SF<sub>6</sub> (Maiss and Brenninkmeijer 1998; Levin et al. 2010), PFC-14 and carbon tetrafluoride (CF<sub>4</sub>) (Harnisch and Eisenhauer 1998). These methods can be applicable to cover a large proportion of global emissions, and monitoring is possible on a routine basis. Although the global trend data can not be directly related to national inventory, the findings can be useful for identifying deficiencies in inventory methodology, such as a need for revising default emission factors. Year to year changes of global abundance of the long-lived atmospheric trace gases can be reliably measured at one or a few background monitoring stations (Prinn et al. 2018). Atmospheric measurements are useful for evaluating the global emissions of the new fluorinated compounds, even before reporting and inventory procedures are well established. For example,

emerging growth in the atmospheric content of HFC-365mfc, HFC-245fa, HFC-227ea, HFC-236fa, and NF<sub>3</sub> were quantified using background concentration monitoring (Stemmler et al. 2007; Vollmer et al. 2011; Arnold et al. 2012), and unexpected continuing emissions of CFC-11 were detected (Montzka et al. 2018). Measurements of the methane isotopic composition were used by Schwietzke et al. 2016, Rice et al. 2016 and others to propose corrections of the global emissions of methane, with implications for estimates of global methane emissions of both fossil (including oil and gas) and biogenic (wetlands and agriculture) origin. Continuous observations of multiple trace gases provide opportunity to use the strong correlations observed between short term variabilities of different tracers to deduce (approximately) the regional emission rate ratios (e.g. CH<sub>4</sub>/CO, CH<sub>4</sub>/CO<sub>2</sub>) and their trends over time, as shown by Fraser et al. 2014 for Australia, and Tohjima et al. 2014 for East Asia.

### 6.10.2.2 SUMMARY OF COMPONENTS NEEDED FOR GHG EMISSION INVENTORY VERIFICATION USING ATMOSPHERIC MEASUREMENTS

Establishing a verification system for national greenhouse gas inventories based on atmospheric observations and inverse modelling requires overcoming technical challenges and involves costs. Such verification of emission estimates needs to be undertaken by atmospheric observation scientists and modellers informed by GHG inventory priorities and needs. The following key elements needed are summarized below (see also report by Manning et al. 2017 and Section 6.10.2.6 for a description of the implementation steps):

- **Atmospheric observations.**
- **Surface-based and airborne observations** of atmospheric GHGs are made by, usually, networks of meteorological agencies, research institutes and site operators. The observations need to meet high standards in all procedures including air sample analysis, data processing, reference gas maintenance, calibration correction against international standards, accompanied by metadata on conditions of measurement. Data quality can be better monitored by data submission to global databases such as World Data Centre for Greenhouse Gases (WDCGG). Establishing a national GHG monitoring network involves optimal network design in order to set up the observation locations that maximize the effect of the observations on reducing the uncertainty of the emission estimates (Nickless et al. 2015; Lopez-Coto et al. 2017). The guidelines for observation techniques and reference gas maintenance are provided by the WMO Global Atmospheric Watch Program<sup>11</sup>, and Advanced Global Atmospheric Gases Experiment Network (AGAGE) (Prinn et al. 2018).
- **Satellite retrievals.** While in situ measurements have the advantage of directly measuring concentrations within the boundary layer, providing strong constraints on regional emissions, satellite retrievals are integrated over a larger portion of the atmospheric column and are subject to biases. To reduce biases, satellite observations are compared to observations of column abundance by a network of high-resolution ground-based spectrometers (Kulawik et al. 2016). Due to their greater spatial coverage, observations from satellites were used to improve the inverse model estimates for methane, by Ganesan et al. 2017 for India, and Turner et al. 2015 for USA. OCO-2 satellite observations have shown potential for quantifying carbon dioxide emissions from large power plants (Nassar et al. 2017).
- **Inverse modelling tools** backed by guidance from expert inverse model users and developers. A number of transport models - Flexpart (Stohl et al. 2005), NAME (Jones et al. 2007), STILT (Lin et al. 2003) - and inverse-modelling tools: Flexinvert (Thompson & Stohl 2014), NAME-InTEM (Manning et al. 2011), Carbontracker (van der Laan-Luijckx et al. 2017), GEOS-Chem (Henze et al. 2007), PYVAR (Chevallier et al. 2005) are available from the developer groups for use in emission estimates. It should be noted that, despite the availability of inverse modelling tools, experienced modellers are required to apply them. As a simple, but less accurate alternative to inverse modelling, tracer correlation methods are also being used, especially for selected halocarbons in Switzerland and Australia (Fraser et al. 2014).
- **Gridded prior inventory data as input for inverse modelling.** For use in inverse modelling the national GHG inventory should be spatially and temporally disaggregated and presented as a gridded emission dataset, typically at 1 km to 10 km spatial resolution for national scale estimates (Maasackers et al. 2016; Tsagatakis et al. 2017), while continental or global emission inventory dataset can be used for larger geographical domains, such as EDGAR (Janssens-Maenhout et al. 2017). While EDGAR is using Tier 1 emission factors and global activity maps for multiple sectors, national scale datasets have the advantage of applying country specific emission factors and have more detailed activity maps. In the absence of the current national gridded inventory data, global coverage is provided by the EDGAR database and is often used. Spatial distribution of the prior emission inventory influences the inverse model estimates, so it is of high value to develop realistic

<sup>11</sup> WMO reports (<https://www.wmo.int/pages/prog/arep/gaw/gaw-reports.html>).

gridded emissions based on the best available geospatial activity data. In several inverse modelling studies, the influence of the gridded inventory of estimated emissions is checked by applying sensitivity tests, where several versions of gridded inventories are used.

Spatial resolution of the gridded inventory should correspond to spatial coverage of observation network and resolution of the transport model, and its temporal resolution should reflect temporal changes in emissions modeled in the national inventory. The gridded inventory should to the extent possible match the methods, data, and results in the national inventory. It should rely on detailed activity data spatial information from the same activity data source as the inventory. In many cases, use of geospatial proxies is necessary (e.g., if data are unavailable on landfill locations, waste emissions could be mapped to population maps). Where emissions factors and activity are known to vary regionally (and if this is modeled in the national inventory), this should be reflected in the gridded inventory. Sources with seasonal changes in emissions rates should also be reflected in the gridded inventory. The information in each grid cell should allow tracing back the data sources to assess which emissions sources in a grid may be relevant to comparing the inventory to independent estimates. In addition, uncertainty information (both on emissions magnitude and spatial allocation) should be developed.

An example of a gridded inventory by Maasackers et al. 2016 is based on the same data as U.S. Inventory prepared by EPA. The authors (which include atmospheric scientists and the EPA Inventory compilers) disaggregate the 2012 national emissions reported by the 2016 version of the U.S. Inventory into a gridded  $0.1^\circ \times 0.1^\circ$  monthly inventory. The gridded inventory is consistent with the EPA national emission totals for each source type and distributes these emissions based on information at the state, county, sub-county, and point source levels. A key input to the U.S. Inventory for many source calculations is EPA's Greenhouse Gas Reporting Program (GHGRP), which collects methane emission and supporting data from large facilities with emissions greater than 25 Gg CO<sub>2</sub> equivalent per year. Where possible, facility-level emissions from the detailed GHGRP emission reports are used in both the gridded inventory and the U.S. Inventory. For facilities that do not report to the GHGRP, emissions are assigned based on the remainder of total national or regional emissions for the source type, and activity data. For source types outside of the scope of the GHGRP, emissions are apportioned by other methods, such as use of detailed state-level modelling for emissions from livestock. Spatial disaggregation is based on livestock population distribution, crop area maps, oil and gas well databases, maps of natural gas pipelines, compressor stations, landfills, vegetation fires and other available information. In the gridded inventory, emission estimates for each source in each month reflect temporal variation by incorporating factors such as changes in livestock populations, and certain temperature-dependent emissions sources such as manure management and landfills. This helps the gridded inventory to better align with studies taken over shorter time frames than the annual U.S. Inventory.

- **Collaboration** between inventory compilers and inverse modellers. As an example, UK inventory and inverse modelling teams have worked together historically to improve the value and credibility of the Inversion Technique for Emission Modelling (InTEM) model estimates to the inventory compilers. Brown et al. 2018 noted a case, when a comparison of inventory estimates of HFC-134a with those modelled through the InTEM system has suggested that the inventory may be over estimating its HFC-134a emissions. Further analysis of the mobile air conditioning sector of the inventory, the main UK source of HFC-134a, has suggested several parameters with high uncertainty that may be the source of the difference. Revisions to the refrigeration and air conditioning model (reviewing assumptions following the implementation of the EU F-gas regulations, and other corrections) have been made, resulting in better agreement between inverse modelling and inventory results.

### 6.10.2.3 OVERVIEW OF THE EMISSION ESTIMATES BY TARGET GAS

#### *Methane*

Methane (CH<sub>4</sub>) is considered a favorable candidate to which inverse modelling techniques can be applied because of the strong atmospheric signal to noise ratio of measurements (despite of the influence from seasonally varying natural sources) and the generally high uncertainty in emission estimates that arises from uncertainty of activity data and emission factors. Efforts to estimate national-scale methane emissions using atmospheric observations and inverse models of atmospheric transport have been made in Switzerland (see Table 6.4), the UK (see example in Box 6.6), the USA (Miller et al. 2013), the EU-28 countries, and other regions. Emission estimates for 28 EU countries (Bergamaschi et al. 2018) were made with a set of several inverse models over the period 2006-2012 using observations from a network with 18 stations. The advantage of applying several models is that the spread of individual inverse model results provides a measure of the errors and biases inherent to the transport and inverse modelling. As a summary of the study, it was mentioned that influence of natural wetland emissions over Northern Europe needs to be better quantified, transport models need to be improved, and a network with more monitoring stations is needed.

### ***Carbon dioxide***

Uncertainties of carbon dioxide anthropogenic emissions due to fuel combustion are usually lower than that of inverse model estimates. However, substantial effort is applied to quantify urban emissions (e.g. Lauvaux et al. 2016; Stauffer et al. 2016) that may lead to developing capability to track the emission reduction trends with an addition of dense urban monitoring networks, and supporting tracer measurements useful for discriminating between natural fluxes and fossil emissions, such as atmospheric potential oxygen (APO), as discussed by Minejima et al. 2012 and radiocarbon  $^{14}\text{C}$  in  $\text{CO}_2$  (Levin et al. 2003). High uncertainty makes carbon dioxide emissions and sinks by AFOLU one of the more challenging sectors to verify, particularly carbon stock changes and associated  $\text{CO}_2$  fluxes for land use and management. In this case, use of atmospheric observations is obstructed by strong interference from natural fluxes. In the studies by Ogle et al. 2015, and Steinkamp et al. 2017 the authors did find agreement between the results from the atmospheric  $\text{CO}_2$  concentration data and inverse modelling and an inventory of  $\text{CO}_2$  emissions based on data from US and New Zealand greenhouse gas inventories. The US study focused, in part, on a sub-region of the United States that is dominated by agricultural food production, and showed that in order to verify emissions from the AFOLU sector, compilers will need to address all sources of  $\text{CO}_2$  uptake and release, including lateral movement of carbon, such as transport of agricultural products.

### ***Nitrous oxide***

Nitrous oxide emissions by agricultural soils are known to have large uncertainty because of patchy heterogeneous emission patterns and significant temporal variability, leading to uncertainty in activity data, emission factors and emission rates, which makes it useful to test the estimated emissions with inverse modelling. Inverse model estimates of the nitrous oxide emissions based on atmospheric monitoring are made for many regions of the globe (Manning et al. 2011; Miller et al. 2012; Bergamaschi et al. 2015) and are also reported in UK National Inventory Report (see Box 6.6). In several studies, a reasonable match is found between inventory and inverse model estimates, for example  $\text{N}_2\text{O}$  inverse modelling results for Europe (Bergamaschi et al. 2015) confirm that the amount reported to UNFCCC by 15 EU countries are within the model uncertainty range.

### ***Fluorinated gases***

Fluorinated gases (HFCs, PFCs,  $\text{SF}_6$ ) are particularly suitable for inverse modelling as they are solely of anthropogenic origin and sufficiently long-lived. In addition, bottom up inventories for fluorinated gases are affected by considerable uncertainties. In the past decade, much progress has been made in the use of atmospheric measurements for estimating emissions of these powerful greenhouse gases. This has been made possible due to the increased capability of producing high-quality atmospheric datasets and to the rapid development of inverse modelling techniques that have been extensively applied from the global to the regional (national) scale (Stohl et al. 2009; Keller et al. 2011; Manning et al. 2011). Such studies are based on long-term and/or continuous observations of the atmospheric levels of fluorinated gases that are carried out within international and national programmes - AGAGE (Prinn et al. 2018). National Oceanic and Atmospheric Administration-Earth System Research Laboratory-Global Monitoring Division (NOAA-ESRL-GMD<sup>12</sup>) and others. Switzerland, United Kingdom and Australia (Fraser et al. 2014) included estimates of fluorinated gas emissions based on atmospheric measurements in their National Inventory Reports. Several regional and national scale estimates were made with available observations by Hu et al. 2017 for USA, Keller et al. 2011 and Graziosi et al. 2017 for European countries, Kim et al. 2010 and Fang et al. 2015 for East Asia (China). One of the most studied gases is HFC-134a, the most abundant HFC in the global atmosphere, mainly used as refrigerant in mobile air conditioners and stationary refrigeration. Differently to other HFCs, studies based on atmospheric measurements suggested that bottom-up HFC-134a inventories were likely to overestimate the emissions (Graziosi et al. 2017; Hu et al. 2017; and references therein).

As an example of using atmospheric observations for improving inventory procedures, Australia's annual  $\text{SF}_6$  loss rate from electricity supply and distribution has been calibrated to changes in atmospheric concentrations of  $\text{SF}_6$  measured at the Commonwealth Science and Industrial Research Organisation (CSIRO) Cape Grim monitoring station (Fraser et al. 2014; Australian Government 2018). Interspecies correlation and inverse modelling techniques are used to derive a national estimate of emissions of  $\text{SF}_6$  based on these atmospheric measurements at Cape Grim. Fluctuations in measured concentrations are reflected in changes to the loss rate for each inventory year from 2010 onwards. The strength of this approach is that it enables the inventory estimates to better reflect improvements in industry practice in terms of gas handling, equipment maintenance and decommissioning.  $\text{SF}_6$  is an ideal gas on which to use inverse modelling techniques to derive national estimates as the likely uncertainty of model results is less than the uncertainty of inventory estimates, especially as inventory leakage rates are based on limited measurements. There is also clarity over the interpretation of the observations of  $\text{SF}_6$  because this gas has no natural sources or sinks and the remote southerly location of the Cape Grim monitoring station reduces the likelihood of measurement error from trans-boundary sources.

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<sup>12</sup> <https://www.esrl.noaa.gov/gmd>.

### 6.10.2.4 STRENGTHS AND WEAKNESSES OF USING ATMOSPHERIC MEASUREMENTS FOR VERIFICATION OF GHG EMISSIONS

The current level of success with the use of atmospheric monitoring for testing anthropogenic GHG emission inventories varies by target gas and region. Usefulness of atmospheric observations depends on several factors, such as uncertainty of the emission inventory and of the models, number and location of available observations, contribution of the natural fluxes to the observed concentration variability (Bergamaschi et al. 2018). Table 6.2 provides an overview of the strengths and weaknesses of using atmospheric measurements for verification of anthropogenic emissions for particular greenhouse gases. More details on the uncertainty of inventory and feasibility of applying inverse modelling for comparison with emission inventories for particular target gases, regions and emission sectors are summarized by Rypdal et al. 2005, Rypdal & Winiwarter 2001 and Janssens-Maenhout et al. 2017. It is worth mentioning that the use of atmospheric measurement in countries like UK or Australia, mentioned in Table 6.2, is facilitated by absence of substantial emissions from surrounding oceans, while implementation of similar approaches in other countries may face different challenges, depending on geographical location.

Gas	Strengths/Successes <sup>13</sup>	Problems/Weaknesses	Future Development/Possibilities
CO <sub>2</sub>	Large number of observations, although historically focusing on natural fluxes.	With sparse observing networks, uncertainties of models may be significantly higher than those of national anthropogenic CO <sub>2</sub> emission inventories.	Need more CO <sub>2</sub> observations targeting anthropogenic emissions, complemented by APO and radiocarbon observations.
CO <sub>2</sub> city-scale	City-scale studies show some degree of success. Inventory uncertainties are relatively larger than at national scale.	Even with dense observation networks, errors in emission estimates are large, due to interference from strong vegetation fluxes. Not used in national reporting.	Large efforts are ongoing to develop observation networks, pilot projects for tracking urban emissions, trends. Radiocarbon, APO, satellite observations also expected to contribute.
CH <sub>4</sub>	Large anthropogenic emission fraction. National reporting <sup>14</sup> : UK, Switzerland. National-scale emission estimates <sup>15</sup> : EU-28, USA, India, China and others.	Few countries have observations, transport and inverse models have uncertainties, interference from natural emissions (wetlands) cited.	Regional observation networks and satellite observations are expanding.
N <sub>2</sub> O	National reporting: UK National-scale emission estimates: EU-28, US, and others.	Observation sites are few, gridded inventories are simplified, large contribution from natural sources.	Expansion of surface networks will contribute to better model estimates.
HFCs, SF <sub>6</sub>	Dominant anthropogenic emission fraction. National reporting: UK, Switzerland, Australia. National-scale emission estimates: China, US, EU. Revised EFs: Australia, UK.	Measurements are sophisticated and expensive. Observation sites are few, gridded inventories are simplified.	Expanding the monitoring network depends on funding.

<sup>13</sup> See references to country studies in Section 6.10.2.3.

<sup>14</sup> Here “National reporting” means the model estimates are included in National Inventory Report.

<sup>15</sup> “National-scale emission estimates” are made on research basis.

## 6.10.2.5 USE OF COMPLIMENTARY OBSERVATIONS AND GLOBAL MODELLING PRODUCTS

### Comparing national inventory to the global inverse model products

For many countries where the national observing networks or national scale inverse model estimates are not available, optionally, national scale emission estimates can still be derived from regional and global inverse modelling results. Regional methane emission assessments have been made by several groups for the EU, East Asia, and North America (Miller et al. 2013; Thompson et al. 2015; Bergamaschi et al. 2018). The data can be requested from the authors and national estimates can be extracted from those inverse modelling results. Regularly updated and publicly available inverse model estimates for CH<sub>4</sub> and N<sub>2</sub>O emissions are provided by operational global and regional inverse modelling products, such as Copernicus Atmosphere Monitoring Services (CAMS) for CH<sub>4</sub> (Segers & Houweling 2017) and N<sub>2</sub>O (Thompson 2017), NOAA Carbontracker-CH<sub>4</sub> (Bruhwiler et al. 2014). The work towards estimating anthropogenic CO<sub>2</sub> emissions is recognized as important (GEO carbon and GHG initiative, Ciais et al. 2014), and is being addressed by a number of national and international programmes, such as the Copernicus initiative for CO<sub>2</sub> observing systems (Pinty et al. 2017). The Global Carbon Project - Methane (GCP-methane) compares and makes available multiple global inverse model estimates (Saunois et al. 2016). Several institutions, such as LSCE, MPI BGC, and Wageningen University also make regular updates of their emission estimates at the global scale and make their gridded flux data available upon request. Step-by-step instructions for using global products for comparison to national inventory are provided in Table 6.5.

### Satellite observations

In regions with sparse ground-based observational coverage, emission estimates by global and regional inverse models have larger biases and uncertainties. This issue is being addressed by expansion of surface observing networks and satellite observations of atmospheric GHGs. Satellite observations by Greenhouse Gases Observing Satellite (GOSAT) were used for national scale methane emission estimates with regional inverse models by Ganesan et al. 2017 for India and Turner et al. 2015 for the USA. Currently several global inverse modelling products by the Copernicus atmospheric monitoring service (Segers & Houweling 2017), the GOSAT Level 4 product (Saito et al. 2016) and several others use satellite observations of methane in addition to the ground-based observations. Emission estimates with inverse models utilizing satellite data are included in the GCP-methane assessment. Use of satellite observations (GOSAT, SCIAMACHY, OCO-2) in inverse modelling for anthropogenic emission estimates is still in the experimental stage, due to multiple technical challenges of producing the high-quality concentration retrievals from the satellite-observed spectra. On the other hand, currently available products are checked for consistency by comparing with estimates made with the use of ground-based observations, and generally do not produce significantly different results (Bruhwiler et al. 2017).

In addition to emission estimates made using inverse methods, several studies have shown the sensitivity of satellite sensors to concentration enhancements around emission hot spots, as summarized in the review<sup>16</sup> (Matsunaga & Maksyutov 2018). A common technique applied in several estimates of anthropogenic CO<sub>2</sub> and CH<sub>4</sub> emissions with satellites, is to take the difference between satellite observations over an emission hot spot or a plume and background concentration defined as a mean of several observations away from polluted area. Local GHG concentration enhancements observed by the GOSAT satellite correlate well with transport model simulations (Janardanan et al. 2016; Janardanan et al. 2017), so that the anthropogenic emissions for large regions like the US or temperate Asia can be estimated by fitting model simulated enhancements to a large number of satellite observations. However, there was less success with country scale estimates due to a lack of observations. With the expected availability of GHG observations from new satellite sensors, such as TROPOMI (Hu et al. 2018), GOSAT-2, GeoCarb, TanSat and others, the limitations of observation numbers will be relaxed, and national scale emission estimates by hot-spot emission data analysis are expected to become possible (the assumption that localised emissions by megacities and other compact sources are representative of regional and national total is supported by large share of population and industrial production being concentrated in conglomerates). Multiple new satellite missions with enhanced capabilities for GHG observations are in preparation, such as listed in CEOS database<sup>17</sup>, so the emission estimates using satellite data will steadily improve.

<sup>16</sup> <http://www.nies.go.jp/soc/en/documents/guidebook>.

<sup>17</sup> <http://database.eohandbook.com>.

### 6.10.2.6 PROCEDURES FOR INVENTORY COMPARISON TO ESTIMATES BASED ON ATMOSPHERIC MEASUREMENTS

#### Key steps to follow in applying National scale observation program and inverse modelling for verification of a national GHG inventory

Several working examples (Manning et al. 2011; Fraser et al. 2014; Henne et al. 2016) of inverse modelling use for national reports are available, while implementing such a system requires advanced technological capability. Alternatively, the use of global data products (see Section 6.10.2.8) can be considered. For countries capable of developing their own observation program and inverse model, several key steps can be identified that are needed for the successful use of inverse modelling in verification of a national GHG inventory. These are summarized in Table 6.3.

Step	Work package	Responsible group
1	Acquisition of GHG observations from a surface network (and when available, from aircraft and satellites) that has sufficient coverage of the country's emissions. The observation data have to be linked to the same calibration scale and be processed by the compatible routines across the network.	Observation /atmospheric modelling
2	Preparing gridded (spatially and temporally disaggregated) prior emissions data.	Gridded inventory
3	Preparing and operating the inverse model, other observation-based emission estimation methods.	Atmospheric modelling
4	Quality Assurance / Quality Control to the inverse model output.	Atmospheric modelling
5	Comparison, verification, and reporting. Production of final outputs and update of the GHG inventory improvement plan.	Inventory/ Atmospheric modelling

In many cases, steps 1, 3, and 4 are conducted by research institutions/divisions not connected to the GHG Inventory compilation, and steps 2, and 5 are conducted in collaboration with the GHG Inventory compiler. To illustrate the content of the procedures made at each step, several examples of comparing the national inventory to the inverse model estimates are provided in the Table 6.4, while a UK example is presented in more detail in the Box 6.6.

**TABLE 6.4 (New)**  
**SUMMARY OF THE KEY STEPS IMPLEMENTED IN NATIONAL EXAMPLES**

<b>Examples</b>	<b>Example 1 Methane emissions in Switzerland<sup>18</sup></b>	<b>Example 2 Methane emissions in UK<sup>19</sup></b>	<b>Example 3 SF<sub>6</sub>, HFCs emissions in Australia<sup>20</sup></b>
<b>Comparison steps</b>			
Step 1: Acquisition of the concentration measurements on national GHGs network.	CarboCount-CH measurement network (4 sites).	Advanced Global Atmospheric Gases Experiment (AGAGE) / UK DECC network, four sites.	Background AGAGE site at Cape Grim (Tasmania), and urban site at Aspendale (Victoria).
Step 2: Preparation of the gridded prior emission data.	Swiss Greenhouse Gas Inventory (SGHGI).	Prior estimates not used.	Australian national inventory.
Step 3: Preparing and operating the inverse model.	Lagrangian particle dispersion model (LPDM) FLEXPART.	Numerical Atmospheric dispersion Modelling Environment (NAME), InTEM (Inversion Technique for Emission Modelling).	Interspecies correlation (ISC), forward CSIRO TAPM model, inverse model NAME-InTEM.
Step 4: Quality assurance/Quality Control to the inverse model.	Sensitivity analysis, Transport model validation.	Sensitivity analysis, Transport model validation.	Sensitivity analysis, Transport model validation.
Step 5: Comparison, verification, and reporting.	Estimated national CH <sub>4</sub> emissions of 196 ± 18 Gg yr <sup>-1</sup> , agrees with SGHGI estimation of 206 ± 33 Gg yr <sup>-1</sup> .	The InTEM methane emission estimates in 2013-2015 (with four DECC sites data) are consistent with UK GHG inventory.	Agreement found to within 2% for HFC-125, HFC- 134a, HFC-143a and HFC- 152a, within 15% for HFC- 23, HFC-365mfc and SF <sub>6</sub> , within 35% for HFC-32.

The examples above generally found good agreement between the national inventories and the observation-based emission estimates. In cases where there are discrepancies between the two estimates, the effort to reduce this discrepancy should be taken by both the inverse modelling and inventory compiling groups, as summarized in Box 6.5.

<sup>18</sup> Henne et al. 2016.

<sup>19</sup> Brown et al. 2018; Manning et al. 2017.

<sup>20</sup> Fraser et al. 2014.

**BOX 6.5 (NEW)****COMPARISON – VERIFICATION ACTIONS ON INVENTORY COMPILER SIDE**

In cases where there are discrepancies between the two estimates, the effort to reduce this discrepancy should be taken by both the inverse modelling and inventory compiling groups. On the inventory compiler side, following steps are suggested to take:

1. Confirm that the observation-based emission estimates and the inventories represent the same time period, areas.
2. Determine what emission dataset was used as a prior, and how it compares to the emission inventory.
3. Assess how the estimation procedure treats anthropogenic and natural emissions, to confirm that the estimates compare with anthropogenic and natural emissions included in the inventory.
4. Confirm that seasonal variability of the emissions and other effects have been considered in the comparison.
5. Assess the uncertainties of the estimated emissions, and note whether the discrepancy is statistically significant.
6. For sub-national scale regions with the larger discrepancies, determine which emissions activities are occurring there, based on the gridded or regional GHG inventory:
  - (i) Recheck inventory activity data in that region;
  - (ii) Assess factors that may make the regional emission rates different from the national inventory average (e.g. different regulations, different technologies), and assess the extent to which these have been taken into account in the national inventory and in its gridding/disaggregation.
7. In the national inventory improvement plan, prioritize emission sources/regions with larger discrepancies.

Example of national inventory comparison to inverse modelling estimates (UK CH<sub>4</sub> and N<sub>2</sub>O inverse modelling) is provided in Box 6.6 below.

**BOX 6.6 (NEW)****UK METHANE (CH<sub>4</sub>) AND NITROUS OXIDE (N<sub>2</sub>O) INVERSE MODELLING**

**Observation and modelling:** In order to provide verification of the UK Greenhouse Gas Inventory (GHGI), the UK government's Department for Business, Energy and Industrial Strategy (BEIS) maintains a high-quality remote observation station at Mace Head (MHD) (set up in 1987) on the west coast of Ireland. The station reports high-frequency concentrations of the key greenhouse gases under the supervision of the University of Bristol (O'Doherty et al. 2004). UK extended the measurement programme in 2012 with three new tall tower stations across the UK: Tacolneston (TAC) near Norwich; Ridge Hill (RGL) near Hereford; Tall Tower Angus (TTA) near Dundee, Scotland (replaced by Bilsdale (BSD) in North Yorkshire in Sept 2015). Methane, carbon dioxide, nitrous oxide and sulphur hexafluoride are measured across the UK network, whereas all of the other gases (e.g. HFCs and PFCs) are only measured at MHD and TAC. The UK Met Office, under contract, employs the Lagrangian dispersion model Numerical Atmospheric dispersion Modelling Environment (NAME) (Jones et al. 2007) driven by three-dimensional modelled meteorology to interpret the observations. By estimating the underlying baseline concentration trends (Northern Hemisphere mid-latitude atmospheric concentrations where the short-term impact of regional pollution has been removed from the data) and by modelling where the air has passed over en route to the observation stations on a regional scale, estimates of UK emissions are made. A methodology called Inversion Technique for Emission Modelling (InTEM) has been developed that uses a Bayesian minimization technique, to determine the emission map that most accurately reproduces the observations (Manning et al. 2003, 2011).

**Output, analysis and arising actions:**

In the UK National Inventory Report (Brown et al. 2018) emission estimates made for the UK using the InTEM methodology are compared to the GHGI emission estimates for the period 1990 onwards. It should be noted that findings, analysis and actions described in the UK National Inventory Report are presented here as examples, and will be different for another country or time period.

**Findings:**

- UK GHG inventory methane estimates have fallen steadily since 1990 largely due to estimated reductions in emission from the waste disposal and energy (fugitives) sectors.
- The InTEM methane emission estimates using all the available observations, including MHD, TAC, RGL, TTA/BSD, and Cabauw (CBW), in 2013-2015 are consistent with UK GHG inventory. Larger mismatch is found in earlier years when using the estimates based on data of two sites (MHD+CBW).
- The annual InTEM estimates for N<sub>2</sub>O are close to the GHG inventory estimates, with both showing declining UK totals. Unlike the GHG inventory however, the InTEM estimates are marginally higher than the GHGI post 2000 although well within the uncertainty. The GHG inventory estimates show a sharp decline (40 Gg) between 1998 and 1999 in line with the introduction of the clean technology at an adipic acid plant in Wilton, north east England.

**Actions:**

- The differences between the GHGI and the inverse modelling trends are a subject of active investigation by the modelling and GHG inventory teams.
- Inventory actions – assessment of missing / underrepresented methane sources:
  - (i) Consider how the yearly variability of emissions from enteric fermentation (specifically sheep) could impact emission estimates. A new agriculture model is being implemented but this is unlikely to have a significant impact.
  - (ii) Review fugitive emissions from offshore oil/gas and coal mines.
  - (iii) Review the extent to which conservative assumptions have been made regarding landfill emissions and methane capture in different parts of the time-series that may misrepresent the real time-series of emissions.

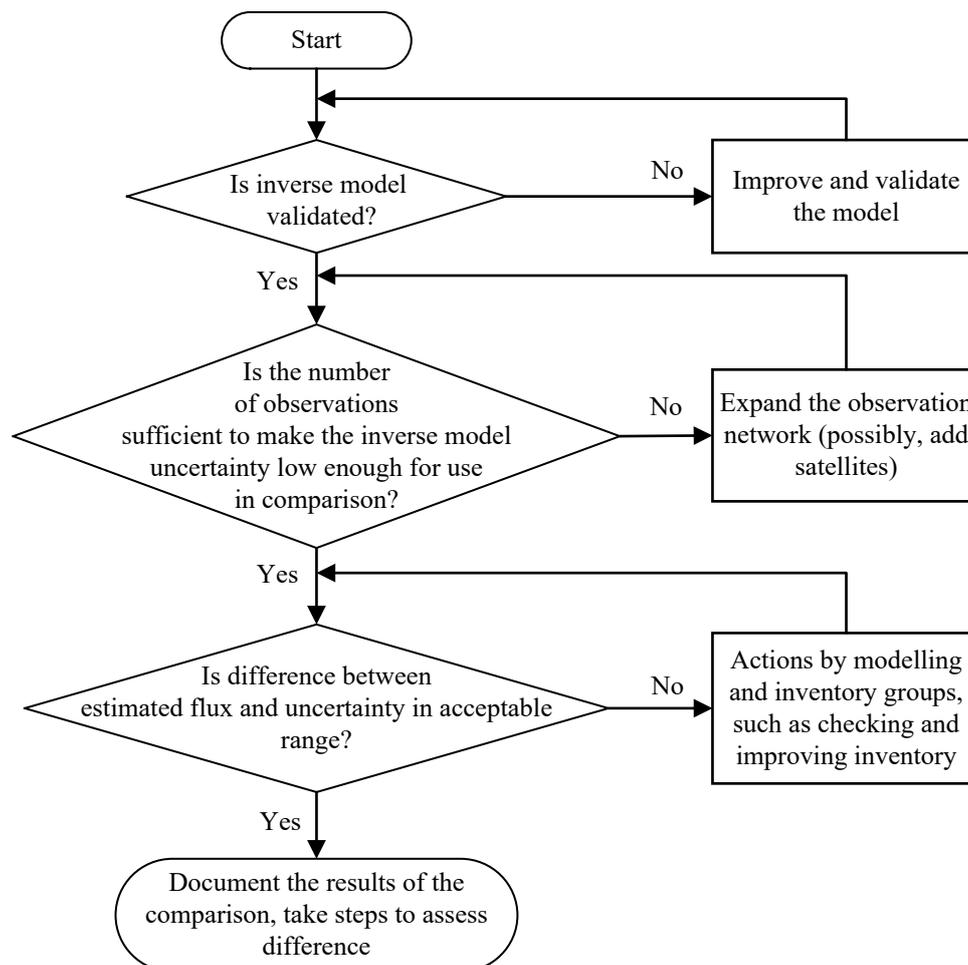
### 6.10.2.7 CHECK LIST FOR APPLYING INVERSE MODEL ESTIMATES FOR COMPARISON WITH NATIONAL INVENTORIES

Utility of inverse model estimates for quality checks and improving the inventory depends on the accuracy and precision of the emission estimates by inverse modelling. The inverse model estimates can be used for inventory verification with more confidence when several conditions are achieved:

- Inverse modelling system has been tested and validated by several methods, including transport model validation with well-known tracers, inverse model validation by model comparison and sensitivity studies.
- Sufficient number of observation sites, and measurement frequency are available. Three to four tall tower sites are used for CH<sub>4</sub> in the Swiss and UK cases, while useful estimates for HFCs were made with one/two sites for UK, Australia and Switzerland. General requirement is to establish large enough number of observation sites to ensure that inverse model estimates are guided more by the observations than by the prior emission inventory. Required number varies by specific gas and target region, and can be determined with the inverse model-guided observation network design, as implemented by (Nickless et al. 2015; Lopez-Coto et al. 2017) and other studies.
- Check if GHG inventory uncertainty is not too low already. This check is applied to avoid comparison of inventories with significantly lower uncertainty (such as carbon dioxide from fossil fuel) to the inverse model estimates. On the other hand, high emission inventory uncertainty is often linked to emissions of HFCs and other fugitive compounds.

Based on these three criteria listed above, a model decision tree for evaluating feasibility of using inverse modelling estimates for inventory verification is shown in Figure 6.1:

**Figure 6.1 (New) A decision tree for checking the conditions for using the inverse model estimates in the National Inventory verification**



### 6.10.2.8 NECESSARY STEPS FOR COMPARING NATIONAL INVENTORY TO THE GLOBAL/REGIONAL INVERSE MODELLING PRODUCTS

An outline of the necessary steps for comparing annual total emissions by national inventory to the emission estimates provided by global/regional inverse model products is given in Table 6.5.

Defining target gases and time periods	<ul style="list-style-type: none"> <li>Based on inverse modelling data available at the time of report preparation, select available gases (CO<sub>2</sub>, CH<sub>4</sub>, N<sub>2</sub>O, HFCs) and periods overlapping between inventory data and inverse model results. Use advice from the modellers on the degree of uncertainty the product is providing for a particular country's emissions.</li> </ul>
Data acquisition	<ul style="list-style-type: none"> <li>Download gridded emission data files (including prior emissions, inverse model estimated emissions and emission uncertainty data), file format descriptions and release notes. Check if the data can be read with available software.</li> </ul>
Remapping to make national total (if national estimate is not provided in the inverse model product)	<ul style="list-style-type: none"> <li>Prepare remapping table. Calculate area fraction of the national land in each grid cell of the emission data grid.</li> <li>Calculate national total emission for each time step, by summing grid emissions multiplied by fraction of national land. Make national total for each year.</li> <li>If data necessary for remapping emission uncertainties is available with inverse modelling results, remap emission uncertainty.</li> </ul>
Using multiple products	<ul style="list-style-type: none"> <li>When the number of available inverse modelling products is more than one, remapping to make national total can be made for all the available products. It is recommended to include in the report national total estimates for each inverse modelling product, along with average and standard deviation of the emissions across the set of inverse modelling products.</li> </ul>
Analysing differences between inverse model estimates and inventory	<ul style="list-style-type: none"> <li>When significant differences between inverse model estimates and inventory are found, check if activity data and emission factors used in inventory can be updated to more recent version, if available. Report differences to inverse modellers, request providing a feedback.</li> </ul>
Documenting the results of the comparison	<ul style="list-style-type: none"> <li>Outline the dataset (datasets) used in the report, cite the product release version, reference the release date, and version of the release note. Provide a description of the remapping procedure used in the remapping. Prepare comparison table showing the national emissions for all gases and years by inventory and emissions with emission uncertainties estimated with inverse models, average value and standard deviation across a set of inverse modelling products.</li> </ul>

## 6.11 DOCUMENTATION, ARCHIVING AND REPORTING

No refinement.

## 6.12 USE AND REPORTING OF MODELS

### 6.12.1 Use of models

The *2006 IPCC Guidelines* provide some guidance on how to ensure that data from models can comply with *good practice* when used in National Greenhouse Gas Inventories. For example, Table 6.6 indicates some of the specific reference in the *2006 IPCC Guidelines* related to the development and use of models. However, this guidance is not complete or systematic: this section addresses this gap.

<b>TABLE 6.6 (NEW)</b> <b>GENERAL GUIDANCE RELATED TO MODELS IN VOLUMES 1 &amp; 4 OF THE 2006 IPCC GUIDELINES</b>	
Section in 2006 IPCC Guidelines	Guidance
<i>Chapter 3, Volume 1: Uncertainties</i>	
3.2.1 Sources of data and information (p 3.14).	Guidance on uncertainties associated with models.
<i>Chapter 5, Volume 1: Time Series Consistency</i>	
5.2.1 Recalculations due to methodological changes.	The calculation of emission factors and other parameters and refinements (Box 5.1, p 5.6) in AFOLU may require a combination of sampling and modelling work. Time series consistency must apply to the modelling work as well. Models can be viewed as a way of transforming input data to produce output results. In most cases where changes are made to the data inputs or mathematical relationships in a model, the entire time series of estimates should be recalculated. In circumstances where this is not feasible due to available data, variations of the overlap method could be applied.
<i>Chapter 6, Volume 1: Quality Assurance/Quality Control and Verification</i>	
6.7.1 Emissions factor QC (p 6.12)	Guidance on QC checks on models
<i>Chapter 2, Volume 4: Generic Methodologies Applicable to Multiple Land-Use Categories</i>	
2.5.2 Model-based Tier 3 inventories (p 2.52)	Guidance on developing model based Tier 3 inventories for AFOLU sector

## 6.12.2 Why use more complex methods?

Simple approaches to estimating greenhouse gas emissions and removals may be unsatisfactory for some specific categories in some countries because they fail to capture the complexity and diversity of systems and practices, in that sector. Therefore, some inventories rely on more sophisticated approaches, using models or direct measurements.

In general, models may be used to estimate those emissions or removals that cannot be easily otherwise obtained, to extend limited information to cover national emissions and removals, both spatially and temporally, or to improve the accuracy of the estimates. Model development relies on data from direct measurements and uses measured data for calibration and evaluation.

However, models should be used with care. Complex models are not necessarily improvements over simple ones (e.g. carbon dioxide emissions from road transport is best estimated from fuel sold and its carbon content: no transport model will provide a better estimate although they may allocate the emissions to specific vehicle types and estimate improved methane and nitrous oxide emissions). Models are limited by the underlying quality of the data. Use of models will require resources for additional QA/QC and documentation.

## 6.12.3 Models

Models aim to transform input data into outputs in a way that replicates the real world. For example, with inputs of the distance driven by road vehicles an appropriate model can estimate emissions of greenhouse gases. Thus, models add value to original data. Models are frequently used to assess complex systems and can be used to generate data; however, models are means of data transformation and do not remove the need for the original data to drive them.

Every emission or removal estimate has an underlying model and assumptions. Even a simple calculation assumes that units of activity, individually or on the average, carry the same emissions burden:

$$Emission = (Emission\ factor) \times (Activity\ data).$$

This assumption is the underlying model. More complex models are called for where this simple calculation seems inadequate e.g., the sigmoid growth of a stand of trees means that one cannot simply multiply the removal rate by the stand area to get a removal from the atmosphere: the age of the stand also matters. Linkages between processes

can be much more complicated than this. This situation can be captured by more complex models, but the greater complexity can lead to reduced transparency. This guidance aims to achieve greater transparency in these situations.

There are many benefits in using complex models in national greenhouse gas inventories. These may include:

- models may improve coverage and completeness as they can extend existing data to improve geographic coverage/distribution and coverage of source/sink categories by filling in gaps in data;
- models may increase spatial and temporal resolution of estimates;
- generally, models may increase the accuracy of results by an improved representation of the processes covered by the model and more systematic treatment of data;
- models can provide an opportunity to test our understanding of cause-and-effect relationships, hence to potentially assess the impacts of mitigation efforts;
- models may provide comparability with other countries and systems;
- models may improve transparency through stratification by making differences between strata (subcategories) explicit;
- models may improve time series consistency of inventory, for example, by providing annual estimates even where only occasional measurements exist;
- models may be a cost effective and, in many cases, the only possible option to estimate emissions and removals compared to extensive data collection;
- models can enable better projections by matching past estimates and future projections and treatment of nationally specific circumstances, technologies and practices and mitigation efforts;
- models can represent non-linear and dynamic systems better compared to the linear averaging done in most Tier 1 and 2 methods;
- models can be adapted to national circumstances;
- models can provide frameworks for uncertainty analyses and identification of research priorities to improve greenhouse gas inventories as far as is practicable.

However, using models may have some adverse effects in such cases where:

- the model is incorrectly used (e.g., applied outside the domain of application without appropriate adaptation);
- the key assumptions are not correct;
- there are errors in the model;
- inappropriate data are fed into the model;
- models lack transparency unless they are fully documented;
- model development may not be cost-effective;
- models are limited by the underlying data when such data is missing.

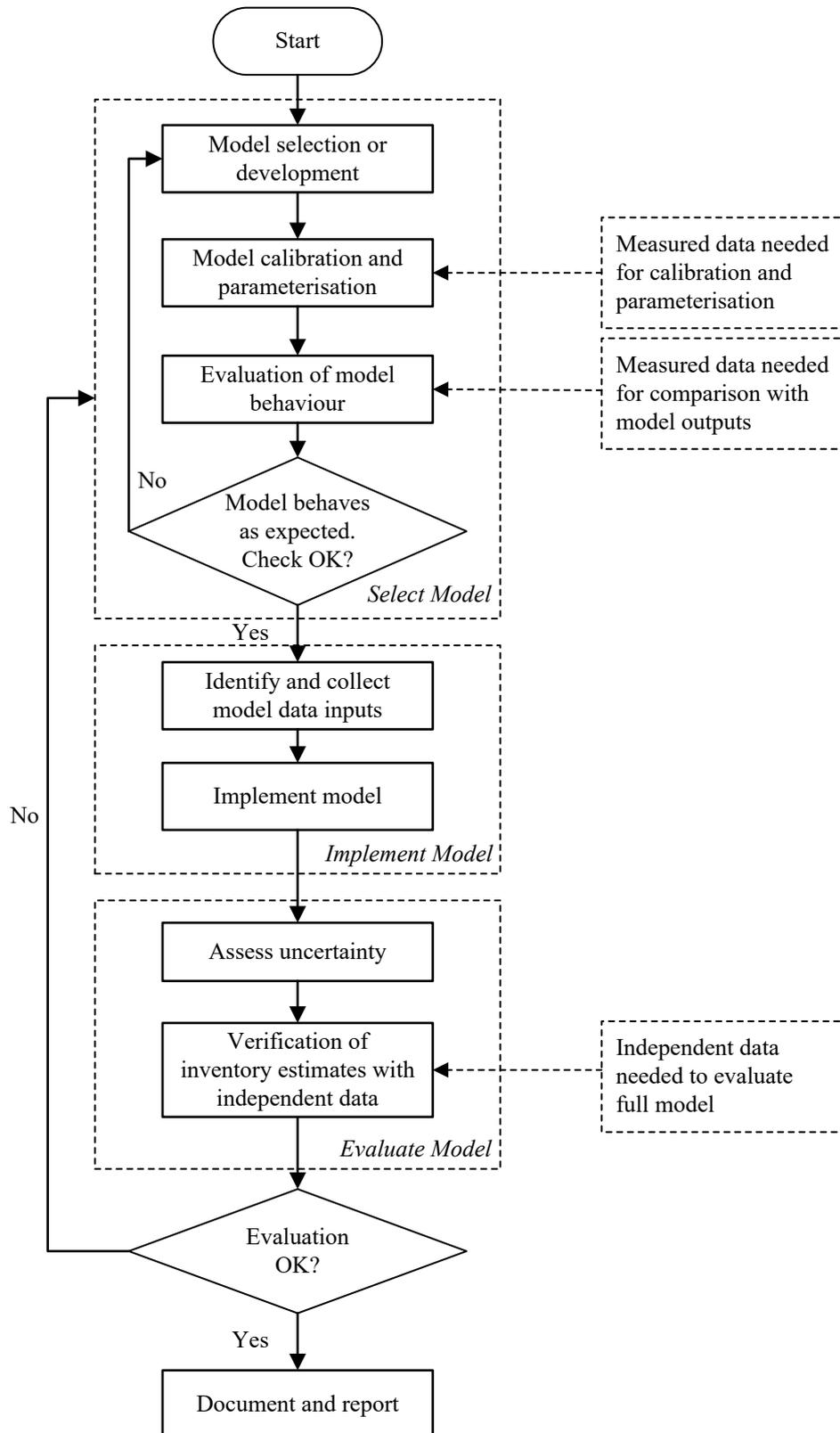
#### **6.12.4 Good Practice Use of Models in National Greenhouse Gas Inventories**

In the application of models in national greenhouse gas inventories, a critical issue is suitability. Suitability describes how well the model reflects the national circumstances: It may have been specifically developed or adapted from an existing model. A model should be correctly parameterized and calibrated, and this will be demonstrated through the model evaluation and the uncertainty assessment. Previously, lack of transparency and inconsistent documentation has been identified as a major concern (IPCC 2010). While these general guidelines will not specify how to choose, build, calibrate or evaluate a model it is crucial that models are reported and documented transparently in order for the model results to be understandable, assessable and credible and the guidelines concentrate on these issues. It is *good practice* to follow the approach given here.

Most complex models should be well-documented covering model description, suitability, calibration, model evaluation and uncertainty and where this exists, the documentation should be referenced: there is no need to reproduce it.

Established and well-known models (e.g. some transport models) are usually well documented, calibrated and validated already. For these inventory compilers can rely on published reports and peer-reviewed publications and simply reference this material. There is no need to duplicate the reports, calibration or validation work, or uncertainty analysis.

**Figure 6.2 (New) Schematic of typical model development/selection process**



### 6.12.4.1 IDENTIFY MODEL: SELECT OR ADAPT EXISTING MODEL OR DEVELOP NEW MODEL

A model must be suitable for its intended use. Suitability is the applicability of the model and any adaptation to the specific national situation in which the model is used for greenhouse gas inventory purposes. A model could be developed for the specific situation or could be a development or adaptation of an existing model. Where an existing model is selected, compilers need to consider and document the following questions:

- Is the model designed for, or portable to, the current national circumstances?
- Are the other conditions for which the model is applied different from those for which the model originally was developed (e.g. ecological or management)?

It is *good practice* to document the suitability of the model. The documentation (see Section 6.12.6) should include:

- The reason for choosing or designing the model (applicability).
- Are there differences in local conditions compared to those for which the model was constructed (or recently adapted to and used) were treated (e.g. ecological or management)?
- Is the model used outside the range of parameter for which the model was developed?

### DEVELOPING OR ADAPTING A MODEL: PARAMETERISATION, CALIBRATION AND EVALUATION OF MODEL BEHAVIOUR<sup>21</sup>

In order to set up, calibrate and parameterise the model real data (“calibration data”) is needed. The data used and outcome of this should be documented.

Following the establishment of the model and its calibration and parameterisation, it is *good practice* to compare model outputs with data independent of the calibration data (e.g. evaluation of model behaviour). This will check whether or not the model behaves as expected and indicates the extent to which the model reproduces the variation in the data that were used to establish its parameter values.

It is *good practice* to ensure that a model responds appropriately to variations in activity data and that the model is able to report results by the required categories. Re-calibration of the model or modifications to the structure (i.e., algorithms) may be necessary if the model does not capture general trends or there are large systematic biases. In some cases, a new model may be selected or developed based on this evaluation. Evaluation results are an important component of the reporting documentation, justifying the use of a particular model for quantifying emissions or removals in an inventory category.

The results of these checks should be documented and reported. It is *good practice* to document the input data needed, the model structure and material assumptions.

### 6.12.4.2 IMPLEMENTING AND EVALUATING THE MODEL

Following the selection of the model, it needs to be implemented. This involves the identification and collection of all the relevant input data and the refinement of the software implementation. Following this, the next step in model development is model results evaluation: comparing model results with independent measurements.

This is an important step in the use of models as it involves testing the fully implemented model, as it will be used in practice with independent data. Evaluation with independent data is done with a completely independent set of data from model calibration, providing a more rigorous assessment of model components and results. Optimally, independent evaluation should be based on measurements from a monitoring network or from research sites that were not used to calibrate model parameters. The sampling does not need to be as dense as needed for measurement-based estimates.

If this independent evaluation demonstrates that the model-based estimation system produces large differences between model results and the measurements this may not indicate the model is wrong. Problems may stem from two other possibilities: errors in the implementation step or poor input data. Implementation problems typically arise from computer programming errors, while model inputs may generate erroneous results if these data are not

<sup>21</sup> The term “*model evaluation*” is used instead of “*validation*” and “*verification*” in this section to describe activities used to determine the appropriateness of models for estimating GHG emissions and/or removals. The terms, “*validation*” and “*verification*” are often used differently in different contexts (e.g. in financial, engineering compared to GHG inventories) and therefore not used here to avoid confusion.

representative of the activity, management or environmental conditions. These possibilities need to be excluded before the model is revised or discarded.

It is *good practice* for the results of this evaluation to be documented and reported.

The evaluation should cover the following points:

- Testing should cover different conditions, circumstances and spatial scales.
- Partial or component tests for the measurable parts should be performed.
- Evaluation of the model output through model inter-comparison, if possible. This will show which models best represent local conditions.
- Evaluation of the model through comparison with Tier 1 or Tier 2 results. Differences between a complex model and lower tier approaches may reflect that the model is better representing the real world (e.g. temporal variability), by including effects not represented in the lower tier. Therefore, it is important to explain significant differences in terms of the physical processes represented in the model. Uncertainty assessment results from the lower tier approaches should be compared and findings documented.

In addition, it may be possible to produce some indicators that show the model is performing correctly. Reporting such indicators and showing they are correctly conserved will demonstrate model robustness. Examples include:

- AFOLU sector models should conserve mass and land area;
- Energy sector models should be consistent with the energy balance;
- In some industrial sectors, a mass balance is possible (e.g. carbon in refineries and iron and steel plant);
- Transport models should conserve vehicle number.

## UNCERTAINTY AND SENSITIVITY ANALYSIS

While an understanding of likely model uncertainty may be produced based on the model structure and algorithms, uncertainty and sensitivity analysis should also be performed as part of model evaluation. This is important so that a rigorous measure of model confidence, based on model inputs and structure, can be reported. When the model is created or materially modified, it is *good practice* to document (preferably in peer reviewed publications which can be referenced by an inventory report to avoid duplication):

- the error distribution of key parameters;
- the covariance matrix of the model parameters (if it is a parametric model);
- results of either error propagation or Monte-Carlo analysis;
- the results of an evaluation of uncertainties from the comparison of model outputs with the independent data;
- the results of a sensitivity analysis or identification of key parameters/inputs to which the model outputs are more sensitive.

## INTERPRETATION OF MODEL RESULTS

In order to assist the correct interpretation of the model results, experience suggests that it would be useful to also supply, as part of the model and inventory documentation:

- Either a comparison of implied emission factors with country-specific factors or, if not available, IPCC default values. This comparison should also provide an explanation for any significant differences.
- An explanation of any unusual input values and results (i.e. outliers with respect to some reference data).
- The distribution of input and output values.

### 6.12.5 QA/QC for selecting, adapting and using models

It is *good practice* for the selection, development and use of models to be part of the inventory QA/QC plan. The elements described in Section 6.3 of the *2006 IPCC Guidelines* are all relevant. There should be clear roles and responsibilities. The inventory QA/QC plan should include the checking and evaluation steps described and should check that documentation is available. References to appropriate documents and publications are acceptable. Do not replicate existing documents.

Regular use of the model should include checks on the input of data and the reasonableness of outputs.

When the model is created or materially modified, it is *good practice* to include external experts (those not involved in the model development) in the evaluation of the inventories. Publication of the model in peer-reviewed literature is desirable.

In planning the implementation of any model, allowance should be made for sufficient resources to allow adequate QA/QC.

### 6.12.6 Reporting on the use of models in emission inventories

To ensure transparency in the use of models it is *good practice* to document the following items (references should be made to existing model documentation and publications wherever possible):

- Basis and type of model (statistical, deterministic, process-based, empirical, etc.);
- Reasons for selecting the particular model;
- If an existing model is being used and adapted: Area of application of original model and adaptation of the model (description of why and how the model was adapted for conditions outside the originally intended domain of application);
- Main equations/processes;
- Material assumptions (important assumptions made in developing and applying the model);
- Domain of application (description of the range of conditions for which the model has been developed to apply)<sup>22</sup>;
- How the model parameters were estimated;
- Description of key inputs and outputs;
- Details of calibration and evaluation with calibration data and independent data (showing intermediate outputs at an adequately disaggregated level);
- Description of the approach taken to the uncertainty analysis and to the sensitivity analysis, and the results of these analyses;
- QA/QC procedures adopted;
- Findings of QA by experts not involved in the model development;
- Interpretation of model results;
- Comparison of model results with lower tier approaches<sup>23</sup>;
- References to peer-reviewed literature (where details of the research on the model can be found).

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<sup>22</sup> Model outputs should match the definitions and requirements of the IPCC Guidelines.

<sup>23</sup> It is not necessary to do this every year, but in establishing a model as part of a national inventory system, the impact of the model results compared with the lower tier approach should be considered. For example, a model may be able to better describe annual temporal changes and so better describe larger year-to-year variability: this would be averaged out in lower tiers.

## 6.12.7 Checklist for ensuring good practice in the use of complex, higher tier models in national greenhouse gas inventories

These can be reported in publications and available model documentation and referenced in inventory reports: duplication should be avoided.

Model Identification (covering selection, development or adaptation of existing models):

- Selection and applicability of model and adaptation to the situation in which the model is used for GHG inventory purposes:
  - (i) Document choice of model based on published studies using the model for the conditions in your country and/or how the model has been adapted to represent the conditions in your country;
  - (ii) Supplemental documentation may be needed to describe the adaptation of the model to the conditions in a country if publications are not available with this information.
- Basis and type of model (statistical, deterministic, process-based, empirical, top-down, bottom-up etc.):
  - (i) Document the conceptual approach (e.g. model represents statistical relationships or processes), and the mathematical formulation in general terms, such as the model is process-based with a bottom-up approach to estimate emissions.
- Identify main processes and equations:
  - (i) Document the main processes and describe the driving variables for those processes;
  - (ii) List the main equations if feasible (may not be feasible with highly complex models or not necessary with simple bookkeeping models);
  - (iii) Also, cite publications that describe the model in detail if they exist. It may be necessary to develop supplemental information documents if the model description has not been published or to provide regional parameter values that are too detailed to be publishable in a scientific journal.
- Material assumptions in model:
  - (i) Document material assumptions. For example, first order approximation was assumed to represent soil organic matter decomposition for three kinetically defined pools with a short, medium and long turnover time.
- Domain of application:
  - (i) Provide information about the extent of the model application to systems in the country, e.g., all agricultural lands with arable crops grown on upland soils.
- Model calibration and checks:
  - (i) Briefly describe the calibration of the model (i.e., parameterization) which may include tuning individual algorithms or the model in a single operation using informal (manual) adjustments to parameters or an automated optimization that attempts to derive a set of parameters based on minimizing the error in the predictions relative to a set of measurements.
- Document the model checks:
  - (i) Provide graphs or other summaries of the evaluation of calibrated model to measured emissions data. Evaluation data should be from sites that were not used in calibration or data from the calibration sites that were collected at different periods than the data used in the calibration step;
  - (ii) Other key predictions from the model may also be evaluated e.g. net primary production and respiration, litterfall, harvest transfers, or stock sizes that may be predicted in AFOLU sector models;
  - (iii) May also compare performance to other models if other models were evaluated;
  - (iv) Include references to published articles with more detail on the calibration and/or evaluation if available. Supplemental documentation may be needed if this information is not published.
- Model Implementation and Model Evaluation:
  - (i) Identify Model Inputs:

- i. Describe type of data inputs to the model. e.g., weather data were based on analysis of long-term precipitation and temperature data from the national weather service or transportation data were based a national scale monitoring of miles travelled by vehicle type, engine, condition and age;
  - ii. Include references to publications of the input data or online publication of the data;
  - iii. List any key assumptions that were necessary to use these data, such as representativeness of management data;
  - iv. Describe any special considerations about the domain of the inventory application using the model given input data. For example, were different input data sets used in different parts of the domain, or was the application of the model limited to specific parts of the country due to the domain of the input data.
- (ii) Implementation of Model:
  - i. Briefly describe computing framework including the hardware, databases and programs that were used to execute the inventory;
  - ii. Provide a description of output variables from the model and any conversions or modifications made to derive the final emissions and removal estimates;
  - iii. Summarise QA/QC procedures adopted to ensure the modelling systems performed appropriately, e.g. checking that of land area is conserved through the analysis; unit conversions are correct; and review of the procedures, inputs and/or outputs by experts not involved with the inventory. List any critical errors identified and corrective actions taken;
  - iv. Optionally provide examples of simple model calculations, such as emissions and removals by forest stands or landscapes in response to different forest management, natural disturbance, or mitigation scenarios. Examples of model performance may be easier to understand than lengthy and complex descriptions of intended model behaviour.
- (iii) Evaluation of inventory results:
  - i. Describe checks on emission results. This may include:
    - a) Estimating implied emissions factors and comparing to lower tier emission factors and/or expected ranges. Further explanation may be needed for differences;
    - b) Compare to lower tier methods if inventory also estimated with lower tiers;
    - c) Compare to independent measurements that were not used for calibration and evaluation of the model, such as data from a monitoring network in the country.
  - ii. Where conservation of mass is expected (e.g. carbon from fuel combustion, storage and leakage of fluorinated gases, carbon from land use and land use change, nitrogen in waste) check that the mass entering the system in combination with the existing storage, is accounted for through emissions and/or storage in the system. Note that losses of mass may not all be related to greenhouse gas emissions (e.g., nitrate leaching from soils which does not contribute to direct soil nitrous oxide emissions).
- (iv) Assess Uncertainties:
  - i. Provide a description of any sensitivity analysis conducted and a summary of findings in terms of key parameters influencing the model results;
  - ii. Describe the derivation of uncertainties in the model inputs and model structure, as well as any other key uncertainties;
  - iii. Provide references to articles that contain additional detail on sensitivity or uncertainty analysis from your application. Supplemental documentation may be needed if this information is not published.

## References

### References copied from the 2006 IPCC Guidelines

- Barnes, D.H., Wofsy, S.C., Fehlaw, B.P., Gottlieb, E.W., Elkins, J.W., Dutton, G.S. and Montzka S.A. (2003a) Urban/industrial pollution for the New York City-Washington, D. C., corridor, 1996-1998:1. Providing independent verification of CO and PCE emissions inventories, *Geophys J. Res.*, **108**(D6), 4185, 10.1029/2001JD001116, 2003a.
- Barnes, D.H., Wofsy, S.C., Fehlaw, B.P., Gottlieb, E.W., Elkins, J.W., Dutton, G.S., and Montzka, S.A. (2003b). Urban/industrial pollution for the New York City-Washington, D. C., corridor, 1996-1998: 2. A study of the efficacy of the Montreal Protocol and other regulatory measures, *Geophys J. Res.*, **108**(D6), 4186, 10.1029/2001JD001117, 2003b.
- Benjey, W. and Middleton, P. (2002). 'The Climate-Air Quality Scale Continuum and the Global Emission Inventory Activity.' Presented at the EPA Emissions Conference, April 15-18.
- Benkovitz C. (2001). 'Compilation of Regional to Global Inventories of Anthropogenic Emissions'. Submitted for publication in "Emissions of Chemical Species and Aerosols into the Atmosphere", Precursors of Ozone and their Effects in the Troposphere (POET), Kluwer Academic Publishers, Dordrecht, Netherlands.
- Bergamaschi, P., Behrend, H. and Andre, J., eds. (2004). Inverse Modeling of National and EU Greenhouse Gas Emission Inventories. Report of the October 23-24 workshop "Inverse Modeling for Potential Verification of National and EU Bottom-up GHG Inventories", held by the European Commission, Joint Research Centre. Report published.
- Biraud, S., Ciais, P., Ramonet, M., Simmonds, P., Kazan, V., Monfray, P., O'Doherty S., Spain T.G. and Jennings, S.G. (2000). European greenhouse gas emissions estimated from continuous atmospheric measurements and radon 222 at Mace Head, Ireland, *J. Geophys. Res.*, **105**(D1), 1351-1366.
- Cape, J.N., Methven, J. and Hudson L.E. (2000). The use of trajectory cluster analysis to interpret trace gas measurements at Mace Head, Ireland, *Atmospheric Environment*, **34**(22), 3651-3663.
- Derwent, R.G., Manning, A.J. and Ryall D.B. (2001). Interpretation of Long-Term Measurements of Ozone-Depleting Substances and Radiatively Active Trace Gases: Phase III, Final Report: DETR Contract No: EPG 1/1/103, Dec 2001.
- Derwent, R.G., Simmonds, P.G., O'Doherty, S., Ciais P., and Ryall, D.B. (1998b). European source strengths and northern hemisphere baseline concentrations of radiatively active trace gases at Mace Head Ireland, *Atmospheric Environment* **32**(21), 3703-3715.
- Derwent, R.G., Simmonds, P.G., O'Doherty, S. and Ryall, D.B. (1998a). The impact of the Montreal Protocol on halocarbon concentrations in northern hemisphere baseline and European air masses at Mace Head Ireland over a ten year period from 1987-1996, *Atmospheric Environment* **32**(21), 3689-3702.
- Dlugokencky, E.J., Steele, L.P., Lang, P.M. and Mesarie, K.A., (1994). The growth rate and distribution of atmospheric CH<sub>4</sub>. *J. Geophys. Res.* 99, 17021-17043.
- EDGAR. Emission Database for Global Atmospheric Research (EDGAR). RIVM-MNP, Bilthoven, TNO-MEP, Apeldoorn, JRC-IES, ISPRA and MPIC-AC, URL: <http://www.mnp.nl/edgar/>.
- Harnisch, J. and Eisenhauer, A. (1998). Natural CF<sub>4</sub> and SF<sub>6</sub> on Earth, *Geophys. Res. Lett.*, **25**(13), 2401-2404.
- Maiss, M. and Brenninkmeijer, C.A.M. (1998) Atmospheric SF<sub>6</sub>: trends, sources and prospects. *Environ. Sci. Techn.* 32, 3077-3086.
- Manning, A.J., Ryall, D.B., Derwent, R.G., Simmonds, P.G. and O'Doherty S. (2003). Estimating European emissions of ozone-depleting and greenhouse gases using observations and a modelling back-attribution technique, *J. Geophys. Res.* Vol. 108, No. D14, 4405, 10.1029/2002JD002312, 17 July 2003.
- NACP (2002). *The North American Carbon Programme*. NACP Committee of the U.S. Carbon Cycle Science Steering Group (Steven C. Wofsy, Robert C. Harris, co-chairs), Chapter 2, Major Elements of the North American Carbon Program Plan. U.S. Global Change Research Program, Washington, D.C., 2002. <http://www.esig.ucar.edu/nacp>.
- O'Doherty, S., McCulloch, A., O'Leary, E., Finn, J. and Cunningham, D. (2003). Climate Change: Emissions of Industrial Greenhouse Gases (HFCs, PFCs and Sulphur Hexafluoride), Final Report, Environmental Protection Agency ERDTI Report Series No. 10, EPA, Johnstown Castle, C. Wexford, Ireland, 2003.

- Ryall, D.B., Derwent, R.G., Manning, A.J., Simmonds, P.G. and O'Doherty S. (2001). Estimating source regions of European emissions of trace gases from observations at Mace Head, *Atmospheric Environment*, 35, 2507-2523.
- Rypdal, K., Stordal, F., Fuglestad, J.S. and Berntsen, T. (2005). Bottom-up vs. top-down methods in assessing compliance with the Kyoto Protocol, *Climate Policy* 5, 393-405.
- Vermeulen, A.T., Eisma, R., Hensen, A. and Slanina J. (1999). Transport model calculations of NW-European methane emissions, *Environmental Science & Policy*, 2, 315-324.

## References newly cited in the 2019 Refinement

- Arnold, T., Muhle, J., Salameh, P., Harth, C., Ivy, D. & Weiss, R. (2012) Automated Measurement of Nitrogen Trifluoride in Ambient Air. *Analytical Chemistry* **84**(11): 4798-4804.
- Bergamaschi P, Corazza M, Karstens U, Athanassiadou M, Thompson R, Pison I, Manning A, et al. (2015) Top-down estimates of European CH<sub>4</sub> and N<sub>2</sub>O emissions based on four different inverse models. *Atmospheric Chemistry and Physics* **15**: 715-736.
- Bergamaschi, P., Karstens, U., Manning, A., Saunio, M., Tsuruta, A., Berchet, A., Vermeulen, A., Arnold, T., Janssens-Maenhout, G., Hammer, S., Levin, I., Schmidt, M., Ramonet, M., Lopez, M., Lavric, J., Aalto, T., Chen, H., Feist, D., Gerbig, C., Haszpra, L., Hermansen, O., Manca, G., Moncrieff, J., Meinhardt, F., Necki, J., Galkowski, M., O'Doherty, S., Paramonova, N., Scheeren, H., Steinbacher, M. & Dlugokencky, E. (2018) Inverse modelling of European CH<sub>4</sub> emissions during 2006-2012 using different inverse models and reassessed atmospheric observations. *Atmospheric Chemistry and Physics* **18**(2): 901-920.
- Breon, F., Broquet, G., Puygrenier, V., Chevallier, F., Xueref-Remy, I., Ramonet, M., Dieudonne, E., Lopez, M., Schmidt, M., Perrussel, O. & Ciais, P. (2015) An attempt at estimating Paris area CO<sub>2</sub> emissions from atmospheric concentration measurements. *Atmospheric Chemistry and Physics* **15**(4): 1707-1724.
- Brown P., Broomfield M., Cardenas L., Choudrie S., Kilroy E., Jones L., MacCarthy J., et al. (2018) *UK Greenhouse Gas Inventory 1990 to 2016: Annual Report for submission under the Framework Convention on Climate Change*. London: Ricardo Energy & Environment.
- Bruhwyler, L., Dlugokencky, E., Masarie, K., Ishizawa, M., Andrews, A., Miller, J., Sweeney, C., Tans, P. & Worthy, D. (2014) CarbonTracker-CH<sub>4</sub>: an assimilation system for estimating emissions of atmospheric methane. *Atmospheric Chemistry and Physics* **14**(16): 8269-8293.
- Bruhwyler, L. M., Basu, S., Bergamaschi, P., Bousquet, P., Dlugokencky, E., Houweling, S., Ishizawa, M., Kim, H.-S., Locatelli, R., Maksyutov, S., Montzka, S., Pandey, S., Patra, P. K., Petron, G., Saunio, M., Sweeney, C., Schwietzke, S., Tans, P. & Weatherhead, E. C. (2017) US CH<sub>4</sub> emissions from oil and gas production: Have recent large increases been detected? *Journal of Geophysical Research-Atmospheres* **122**(7): 4070-4083.
- Chevallier, F., Fisher, M., Peylin, P., Serrar, S., Bousquet, P., Breon, F., Chedin, A. & Ciais, P. (2005) Inferring CO<sub>2</sub> sources and sinks from satellite observations: Method and application to TOVS data. *Journal of Geophysical Research-Atmospheres* **110**(D24).
- Ciais, P., Dolman, A., Bombelli, A., Duren, R., Peregón, A., Rayner, P., Miller, C., Gobron, N., Kinderman, G., Marland, G., Gruber, N., Chevallier, F., Andres, R., Balsamo, G., Bopp, L., Breon, F., Broquet, G., Dargaville, R., Battin, T., Borges, A., Bovensmann, H., Buchwitz, M., Butler, J., Canadell, J., Cook, R., DeFries, R., Engelen, R., Gurney, K., Heinze, C., Heimann, M., Held, A., Henry, M., Law, B., Luyssaert, S., Miller, J., Moriyama, T., Moulin, C., Myneni, R., Nussli, C., Obersteiner, M., Ojima, D., Pan, Y., Paris, J., Piao, S., Poulter, B., Plummer, S., Quegan, S., Raymond, P., Reichstein, M., Rivier, L., Sabine, C., Schimel, D., Tarasova, O., Valentini, R., Wang, R., van der Werf, G., Wickland, D., Williams, M. & Zehner, C. (2014) Current systematic carbon-cycle observations and the need for implementing a policy-relevant carbon observing system. *Biogeosciences* **11**(13): 3547-3602.
- Ciais, P., Paris, J. D., Marland, G., Peylin, P., Piao, S. L., Levin, I., Pregger, T., Scholz, Y., Friedrich, R., Rivier, L., Houweling, S., Schulze, E. D. & members of the, C. S. T. (2010) The European carbon balance. Part 1: fossil fuel emissions. *Global Change Biology* **16**(5): 1395-1408.
- Conley, S., Faloona, I., Mehrotra, S., Suard, M., Lenschow, D., Sweeney, C., Herndon, S., Schwietzke, S., Petron, G., Pifer, J., Kort, E. & Schnell, R. (2017) Application of Gauss's theorem to quantify localized surface emissions from airborne measurements of wind and trace gases. *Atmospheric Measurement Techniques* **10**(9): 3345-3358.

- DeCola P., Tarasova O., Brunner D., Maksyutov S., Manning A., Vogel F., Gurney K., et al. (2018) *An Integrated Global Greenhouse Gas Information System (IG<sup>3</sup>IS) Science Implementation Plan*. Geneva: WMO.
- Desjardins, R., Worth, D., Pattey, E., VanderZaag, A., Srinivasan, R., Mauder, M., Worthy, D., Sweeney, C. & Metzger, S. (2018) The challenge of reconciling bottom-up agricultural methane emissions inventories with top-down measurements. *Agricultural and Forest Meteorology* **248**: 48-59.
- Fang, X., Stohl, A., Yokouchi, Y., Kim, J., Li, S., Saito, T., Park, S. & Hu, J. (2015) Multiannual Top-Down Estimate of HFC-23 Emissions in East Asia. *Environmental Science & Technology* **49**(7): 4345-4353.
- Fraser P., Dunse B., Krummel P., Steele P., Derek N. (2014) *Australian HFC, PFC, Sulfur Hexafluoride and Sulfuryl Fluoride emissions*. Aspendale, Australia: Centre for Australian Weather and Climate Research, CSIRO Oceans and Atmosphere Flagship.
- Ganesan, A. L., Rigby, M., Lunt, M. F., Parker, R. J., Boesch, H., Goulding, N., Umezawa, T., Zahn, A., Chatterjee, A., Prinn, R. G., Tiwari, Y. K., van der Schoot, M. & Krummel, P. B. (2017) Atmospheric observations show accurate reporting and little growth in India's methane emissions. *Nature Communications* **8**(1): 836.
- Government A. (2018) *National Inventory Report 2016 Volume 1, Commonwealth of Australia*. Department of the Environment and Energy.
- Graziosi, F., Arduini, J., Furlani, F., Giostra, U., Cristofanelli, P., Fang, X., Hermanssen, O., Lunder, C., Maenhout, G., O'Doherty, S., Reimann, S., Schmidbauer, N., Vollmer, M., Young, D. & Maione, M. (2017) European emissions of the powerful greenhouse gases hydrofluorocarbons inferred from atmospheric measurements and their comparison with annual national reports to UNFCCC. *Atmospheric Environment* **158**: 85-97.
- Henne, S., Brunner, D., Oney, B., Leuenberger, M., Eugster, W., Bamberger, I., Meinhardt, F., Steinbacher, M. & Emmenegger, L. (2016) Validation of the Swiss methane emission inventory by atmospheric observations and inverse modelling. *Atmospheric Chemistry and Physics* **16**(6): 3683-3710.
- Henze, D., Hakami, A. & Seinfeld, J. (2007) Development of the adjoint of GEOS-Chem. *Atmospheric Chemistry and Physics* **7**(9): 2413-2433.
- Hu, L., Montzka, S. A., Lehman, S. J., Godwin, D. S., Miller, B. R., Andrews, A. E., Thoning, K., Miller, J. B., Sweeney, C., Siso, C., Elkins, J. W., Hall, B. D., Mondeel, D. J., Nance, D., Nehrkorn, T., Mountain, M., Fischer, M. L., Biraud, S. C., Chen, H. & Tans, P. P. (2017) Considerable contribution of the Montreal Protocol to declining greenhouse gas emissions from the United States. *Geophysical Research Letters* **44**(15): 8075-8083.
- Hu H., Landgraf J., Detmers R., Borsdorff T., de Brugh J., Aben I., Butz A., et al. (2018) Toward Global Mapping of Methane With TROPOMI: First Results and Intersatellite Comparison to GOSAT. *Geophysical Research Letters* **45**: 3682-3689.
- Jacob, D. J., Turner, A. J., Maasakkers, J. D., Sheng, J., Sun, K., Liu, X., Chance, K., Aben, I., McKeever, J. & Frankenberg, C. (2016) Satellite observations of atmospheric methane and their value for quantifying methane emissions. *Atmos. Chem. Phys.* **16**(22): 14371-14396.
- Janardanan, R., Maksyutov, S., Ito, A., Yukio, Y. & Matsunaga, T. (2017) Assessment of Anthropogenic Methane Emissions over Large Regions Based on GOSAT Observations and High Resolution Transport Modeling. *Remote Sensing* **9**(9).
- Janardanan, R., Maksyutov, S., Oda, T., Saito, M., Kaiser, J., Ganshin, A., Stohl, A., Matsunaga, T., Yoshida, Y. & Yokota, T. (2016) Comparing GOSAT observations of localized CO<sub>2</sub> enhancements by large emitters with inventory-based estimates. *Geophysical Research Letters* **43**(7): 3486-3493.
- Janssens-Maenhout, G., Crippa, M., Guizzardi, D., Muntean, M., Schaaf, E., Dentener, F., Bergamaschi, P., Pagliari, V., Olivier, J. G. J., Peters, J. A. H. W., van Aardenne, J. A., Monni, S., Doering, U. & Petrescu, A. M. R. (2017) EDGAR v4.3.2 Global Atlas of the three major Greenhouse Gas Emissions for the period 1970-2012. *Earth Syst. Sci. Data Discuss.* **2017**: 1-55.
- Jones, A., Thomson, D., Hort, M. & Devenish, B. (2007) The U.K. Met Office's Next-Generation Atmospheric Dispersion Model, NAME III. *Air Pollution Modeling and Its Application XVII*, eds. C. Borrego & A.-L. Norman, pp. 580-589. Boston, MA: Springer US.
- Keeling, R. F. (1988) Measuring correlations between atmospheric oxygen and carbon dioxide mole fractions: A preliminary study in urban air. *Journal of Atmospheric Chemistry* **7**(2): 153-176.
- Keller, C. A., Brunner, D., Henne, S., Vollmer, M. K., O'Doherty, S. & Reimann, S. (2011) Evidence for under-reported western European emissions of the potent greenhouse gas HFC-23. *Geophysical Research Letters* **38**(15).

- Kim, J., Li, S., Kim, K., Stohl, A., Muhle, J., Kim, S., Park, M., Kang, D., Lee, G., Harth, C., Salameh, P. & Weiss, R. (2010) Regional atmospheric emissions determined from measurements at Jeju Island, Korea: Halogenated compounds from China. *Geophysical Research Letters* **37**.
- Kulawik, S., et al. (2016). "Consistent evaluation of ACOS-GOSAT, BESD-SCIAMACHY, CarbonTracker, and MACC through comparisons to TCCON." *Atmospheric Measurement Techniques* **9**(2): 683-709.
- Lauvaux, T., Miles, N., Deng, A., Richardson, S., Cambaliza, M., Davis, K., Gaudet, B., Gurney, K., Huang, J., O'Keefe, D., Song, Y., Karion, A., Oda, T., Patarasuk, R., Razlivanov, I., Sarmiento, D., Shepson, P., Sweeney, C., Turnbull, J. & Wu, K. (2016) High-resolution atmospheric inversion of urban CO<sub>2</sub> emissions during the dormant season of the Indianapolis Flux Experiment (INFLUX). *Journal of Geophysical Research-Atmospheres* **121**(10): 5213-5236.
- Levin, I., Kromer, B., Schmidt, M. & Sartorius, H. (2003) A novel approach for independent budgeting of fossil fuel CO<sub>2</sub> over Europe by (CO<sub>2</sub>)-C-14 observations. *Geophysical Research Letters* **30**(23).
- Levin, I., Naegler, T., Heinz, R., Osusko, D., Cuevas, E., Engel, A., Ilmberger, J., Langenfelds, R., Neininger, B., Von Rohden, C., Steele, L., Weller, R., Worthy, D. & Zimov, S. (2010) The global SF<sub>6</sub> source inferred from long-term high precision atmospheric measurements and its comparison with emission inventories. *Atmospheric Chemistry and Physics* **10**(6): 2655-2662.
- Lin, J., Gerbig, C., Wofsy, S., Andrews, A., Daube, B., Davis, K. & Grainger, C. (2003) A near-field tool for simulating the upstream influence of atmospheric observations: The Stochastic Time-Inverted Lagrangian Transport (STILT) model. *Journal of Geophysical Research-Atmospheres* **108**(D16).
- Lopez-Coto, I., Ghosh, S., Prasad, K. & Whetstone, J. (2017) Tower-based greenhouse gas measurement network design - The National Institute of Standards and Technology North East Corridor Testbed. *Advances in Atmospheric Sciences* **34**(9): 1095-1105.
- Maasackers, J., Jacob, D., Sulprizio, M., Turner, A., Weitz, M., Wirth, T., Hight, C., DeFigueiredo, M., Desai, M., Schmeltz, R., Hockstad, L., Bloom, A., Bowman, K., Jeong, S. & Fischer, M. (2016) Gridded National Inventory of US Methane Emissions. *Environmental Science & Technology* **50**(23): 13123-13133.
- Manning, A.J., O'Doherty, S., Jones, A. R., Simmonds, P. G. & Derwent, R. G. (2011) Estimating UK methane and nitrous oxide emissions from 1990 to 2007 using an inversion modeling approach. *Journal of Geophysical Research* **116**(D2).
- Manning A.J., Stanley K., Redington A., O'Doherty S., Young D. (2017) *Methodology Report Verification of Emissions using Atmospheric Observations*. BEIS.
- Matsunaga T., Maksyutov S. (2018) *A Guidebook on the Use of Satellite Greenhouse Gases Observation Data to Evaluate and Improve Greenhouse Gas Emission Inventories*. Japan: Satellite Observation Center, National Institute for Environmental Studies.
- McKain, K., Down, A., Raciti, S., Budney, J., Hutyrá, L., Floerchinger, C., Herndon, S., Nehr Korn, T., Zahniser, M., Jackson, R., Phillips, N. & Wofsy, S. (2015) Methane emissions from natural gas infrastructure and use in the urban region of Boston, Massachusetts. *Proceedings of the National Academy of Sciences of the United States of America* **112**(7): 1941-1946.
- Miller, S., Kort, E., Hirsch, A., Dlugokencky, E., Andrews, A., Xu, X., Tian, H., Nehr Korn, T., Eluszkiewicz, J., Michalak, A. & Wofsy, S. (2012) Regional sources of nitrous oxide over the United States: Seasonal variation and spatial distribution. *Journal of Geophysical Research-Atmospheres* **117**.
- Miller, S. & Michalak, A. (2017) Constraining sector-specific CO<sub>2</sub> and CH<sub>4</sub> emissions in the US. *Atmospheric Chemistry and Physics* **17**(6): 3963-3985.
- Miller, S., Wofsy, S., Michalak, A., Kort, E., Andrews, A., Biraud, S., Dlugokencky, E., Eluszkiewicz, J., Fischer, M., Janssens-Maenhout, G., Miller, B., Miller, J., Montzka, S., Nehr Korn, T. & Sweeney, C. (2013) Anthropogenic emissions of methane in the United States. *Proceedings of the National Academy of Sciences of the United States of America* **110**(50): 20018-20022.
- Minejima, C., Kubo, M., Tohjima, Y., Yamagishi, H., Koyama, Y., Maksyutov, S., Kita, K. & Mukai, H. (2012) Analysis of Delta O-2/Delta CO<sub>2</sub> ratios for the pollution events observed at Hateruma Island, Japan. *Atmospheric Chemistry and Physics* **12**(5): 2713-2723.
- Montzka, S., Dutton, G., Yu, P., Ray, E., Portmann, R., Daniel, J., Kuijpers, L., Hall, B., Mondeel, D., Siso, C., Nance, D., Rigby, M., Manning, A., Hu, L., Moore, F., Miller, B. & Elkins, J. (2018) An unexpected and persistent increase in global emissions of ozone-depleting CFC-11. *Nature* **557**(7705): 413-417.

- Nassar, R., Hill, T., McLinden, C., Wunch, D., Jones, D. & Crisp, D. (2017) Quantifying CO<sub>2</sub> Emissions From Individual Power Plants From Space. *Geophysical Research Letters* **44**(19): 10045-10053.
- Nickless, A., Ziehn, T., Rayner, P., Scholes, R. & Engelbrecht, F. (2015) Greenhouse gas network design using backward Lagrangian particle dispersion modelling - Part 2: Sensitivity analyses and South African test case. *Atmospheric Chemistry and Physics* **15**(4): 2051-2069.
- O'Doherty, S., Cunnold, D., Manning, A., Miller, B., Wang, R., Krummel, P., Fraser, P., Simmonds, P., McCulloch, A., Weiss, R., Salameh, P., Porter, L., Prinn, R., Huang, J., Sturrock, G., Ryall, D., Derwent, R. & Montzka, S. (2004) Rapid growth of hydrofluorocarbon 134a and hydrochlorofluorocarbons 141b, 142b, and 22 from Advanced Global Atmospheric Gases Experiment (AGAGE) observations at Cape Grim, Tasmania, and Mace Head, Ireland. *Journal of Geophysical Research-Atmospheres* **109**(D6).
- Ogle, S., Davis, K., Lauvaux, T., Schuh, A., Cooley, D., West, T., Heath, L., Miles, N., Richardson, S., Breidt, F., Smith, J., McCarty, J., Gurney, K., Tans, P. & Denning, A. (2015) An approach for verifying biogenic greenhouse gas emissions inventories with atmospheric CO<sub>2</sub> concentration data. *Environmental Research Letters* **10**(3).
- Pinty B., Janssens-Maenhout G., Dowell M., Zunker H., Brunhes T., Ciais P., Dee D., et al. (2017) An Operational Anthropogenic CO<sub>2</sub> Emissions Monitoring & Verification Support capacity - Baseline Requirements, Model Components and Functional Architecture. European Commission Joint Research Centre.
- Pison, I., Berchet, A., Saunio, M., Bousquet, P., Broquet, G., Conil, S., Delmotte, M., Ganesan, A., Laurent, O., Martin, D., O'Doherty, S., Ramonet, M., Spain, T., Vermeulen, A. & Kwok, C. (2018) How a European network may help with estimating methane emissions on the French national scale. *Atmospheric Chemistry and Physics* **18**(5): 3779-3798.
- Prinn, R. G., Weiss, R. F., Arduini, J., Arnold, T., DeWitt, H. L., Fraser, P. J., Ganesan, A. L., Gasore, J., Harth, C. M., Hermansen, O., Kim, J., Krummel, P. B., Li, S., Loh, Z. M., Lunder, C. R., Maione, M., Manning, A. J., Miller, B. R., Mitrevski, B., Mühle, J., O'Doherty, S., Park, S., Reimann, S., Rigby, M., Saito, T., Salameh, P. K., Schmidt, R., Simmonds, P. G., Steele, L. P., Vollmer, M. K., Wang, R. H., Yao, B., Yokouchi, Y., Young, D. & Zhou, L. (2018) History of chemically and radiatively important atmospheric gases from the Advanced Global Atmospheric Gases Experiment (AGAGE). *Earth Syst. Sci. Data* **10**(2): 985-1018.
- Rice, A., Butenhoff, C., Teama, D., Roger, F., Khalil, M. & Rasmussen, R. (2016) Atmospheric methane isotopic record favors fossil sources flat in 1980s and 1990s with recent increase. *Proceedings of the National Academy of Sciences of the United States of America* **113**(39): 10791-10796.
- Rypdal, K. & Winiwarter, W. (2001) Uncertainties in greenhouse gas emission inventories - evaluation, comparability and implications. *Environmental Science & Policy* **4**(2): 107-116.
- Saito, M., Kim, H., Ito, A., Yokota, T. & Maksyutov, S. (2016) Enhanced Methane Emissions during Amazonian Drought by Biomass Burning. *Plos One* **11**(11).
- Saunio, M., Bousquet, P., Poulter, B., Peregon, A., Ciais, P., Canadell, J., Dlugokencky, E., Etiope, G., Bastviken, D., Houweling, S., Janssens-Maenhout, G., Tubiello, F., Castaldi, S., Jackson, R., Alexe, M., Arora, V., Beerling, D., Bergamaschi, P., Blake, D., Brailsford, G., Brovkin, V., Bruhwiler, L., Crevoisier, C., Crill, P., Covey, K., Curry, C., Frankenberg, C., Gedney, N., Hoglund-Isaksson, L., Ishizawa, M., Ito, A., Joos, F., Kim, H., Kleinen, T., Krummel, P., Lamarque, J., Langenfelds, R., Locatelli, R., Machida, T., Maksyutov, S., McDonald, K., Marshall, J., Melton, J., Morino, I., Naik, V., O'Doherty, S., Parmentier, F., Patra, P., Peng, C., Peng, S., Peters, G., Pison, I., Prigent, C., Prinn, R., Ramonet, M., Riley, W., Saito, M., Santini, M., Schroeder, R., Simpson, I., Spahni, R., Steele, P., Takizawa, A., Thornton, B., Tian, H., Tohjima, Y., Viovy, N., Voulgarakis, A., van Weele, M., van der Werf, G., Weiss, R., Wiedinmyer, C., Wilton, D., Wiltshire, A., Worthy, D., Wunch, D., Xu, X., Yoshida, Y., Zhang, B., Zhang, Z. & Zhu, Q. (2016) The global methane budget 2000-2012. *Earth System Science Data* **8**(2): 697-751.
- Schwietzke, S., Sherwood, O., Ruhwiler, L., Miller, J., Etiope, G., Dlugokencky, E., Michel, S., Arling, V., Vaughn, B., White, J. & Tans, P. (2016) Upward revision of global fossil fuel methane emissions based on isotope database. *Nature* **538**(7623): 88-91.
- Segers A., Houweling S. (2017) *Description of the CH<sub>4</sub> Inversion Production Chain*. Copernicus Atmosphere Monitoring Service.
- Stauffer, J., Broquet, G., Breon, F., Puygrenier, V., Chevallier, F., Xueref-Remy, I., Dieudonne, E., Lopez, M., Schmidt, M., Ramonet, M., Perrussel, O., Lac, C., Wu, L. & Ciais, P. (2016) The first 1-year-long estimate of the Paris region fossil fuel CO<sub>2</sub> emissions based on atmospheric inversion. *Atmospheric Chemistry and Physics* **16**(22): 14703-14726.

- Steinkamp, K., Fletcher, S., Brailsford, G., Smale, D., Moore, S., Keller, E., Baisden, W., Mukai, H. & Stephens, B. (2017) Atmospheric CO<sub>2</sub> observations and models suggest strong carbon uptake by forests in New Zealand. *Atmospheric Chemistry and Physics* **17**(1): 47-76.
- Stemmler, K., Folini, D., Uhl, S., Vollmer, M., Reimann, S., Doherty, S., Grealley, B., Simmonds, P. & Manning, A. (2007) European emissions of HFC-365mfc, a chlorine-free substitute for the foam blowing agents HCFC-141b and CFC-11. *Environmental Science & Technology* **41**(4): 1145-1151.
- Stohl, A., Forster, C., Frank, A., Seibert, P. & Wotawa, G. (2005) Technical note: The Lagrangian particle dispersion model FLEXPART version 6.2. *Atmospheric Chemistry and Physics* **5**(9): 2461-2474.
- Stohl, A., Seibert, P., Arduini, J., Eckhardt, S., Fraser, P., Grealley, B. R., Lunder, C., Maione, M., Mühle, J., O'Doherty, S., Prinn, R. G., Reimann, S., Saito, T., Schmidbauer, N., Simmonds, P. G., Vollmer, M. K., Weiss, R. F. & Yokouchi, Y. (2009) An analytical inversion method for determining regional and global emissions of greenhouse gases: Sensitivity studies and application to halocarbons. *Atmospheric Chemistry and Physics* **9**(5): 1597-1620.
- Thompson R. (2017) *Description of the N<sub>2</sub>O inversion production chain*. Reading: Copernicus Atmosphere Monitoring Service.
- Thompson, R. & Stohl, A. (2014) FLEXINVERT: an atmospheric Bayesian inversion framework for determining surface fluxes of trace species using an optimized grid. *Geoscientific Model Development* **7**(5): 2223-2242.
- Thompson, R., Stohl, A., Zhou, L., Dlugokencky, E., Fukuyama, Y., Tohjima, Y., Kim, S., Lee, H., Nisbet, E., Fisher, R., Lowry, D., Weiss, R., Prinn, R., O'Doherty, S., Young, D. & White, J. (2015) Methane emissions in East Asia for 2000-2011 estimated using an atmospheric Bayesian inversion. *Journal of Geophysical Research-Atmospheres* **120**(9): 4352-4369.
- Tohjima, Y., Kubo, M., Minejima, C., Mukai, H., Tanimoto, H., Ganshin, A., Maksyutov, S., Katsumata, K., Machida, T. & Kita, K. (2014) Temporal changes in the emissions of CH<sub>4</sub> and CO from China estimated from CH<sub>4</sub>/CO<sub>2</sub> and CO/CO<sub>2</sub> correlations observed at Hateruma Island. *Atmospheric Chemistry and Physics* **14**(3): 1663-1677.
- Tsagatakis I., Brace S., Passant N., Pearson B., Kiff B., Richardson J., Ruddy M. (2017) *UK Emission Mapping Methodology - A report of the National Atmospheric Emission Inventory 2015*. London: Ricardo Energy & Environment.
- Turner, A., Jacob, D., Wecht, K., Maasakkers, J., Lundgren, E., Andrews, A., Biraud, S., Boesch, H., Bowman, K., Deutscher, N., Dubey, M., Griffith, D., Hase, F., Kuze, A., Notholt, J., Ohyama, H., Parker, R., Payne, V., Sussmann, R., Sweeney, C., Velazco, V., Warneke, T., Wennberg, P. & Wunch, D. (2015) Estimating global and North American methane emissions with high spatial resolution using GOSAT satellite data. *Atmospheric Chemistry and Physics* **15**(12): 7049-7069.
- Van der Laan-Luijkx, I. T., van der Velde, I. R., van der Veen, E., Tsuruta, A., Stanislawski, K., Babenhausenheide, A., Zhang, H. F., Liu, Y., He, W., Chen, H., Masarie, K. A., Krol, M. C. & Peters, W. (2017) The CarbonTracker Data Assimilation Shell (CTDAS) v1.0: implementation and global carbon balance 2001–2015. *Geosci. Model Dev.* **10**(7): 2785-2800.
- Viatte, C., Lauvaux, T., Hedelius, J. K., Parker, H., Chen, J., Jones, T., Franklin, J. E., Deng, A. J., Gaudet, B., Verhulst, K., Duren, R., Wunch, D., Roehl, C., Dubey, M. K., Wofsy, S. & Wennberg, P. O. (2017) Methane emissions from dairies in the Los Angeles Basin. *Atmos. Chem. Phys.* **17**(12): 7509-7528.
- Vollmer, M., Miller, B., Rigby, M., Reimann, S., Mühle, J., Krummel, P., O'Doherty, S., Kim, J., Rhee, T., Weiss, R., Fraser, P., Simmonds, P., Salameh, P., Harth, C., Wang, R., Steele, L., Young, D., Lunder, C., Hermansen, O., Ivy, D., Arnold, T., Schmidbauer, N., Kim, K., Grealley, B., Hill, M., Leist, M., Wenger, A. & Prinn, R. (2011) Atmospheric histories and global emissions of the anthropogenic hydrofluorocarbons HFC-365mfc, HFC-245fa, HFC-227ea, and HFC-236fa. *Journal of Geophysical Research-Atmospheres* **116**.
- Yver-Kwok, C., Muller, D., Caldow, C., Lebeque, B., Monster, J., Rella, C., Scheutz, C., Schmidt, M., Ramonet, M., Warneke, T., Broquet, G. & Ciais, P. (2015) Methane emission estimates using chamber and tracer release experiments for a municipal waste water treatment plant. *Atmospheric Measurement Techniques* **8**(7): 2853-2867.
- Zavala-Araiza, D., Lyon, D., Alvarez, R., Davis, K., Harriss, R., Herndon, S., Karion, A., Kort, E., Lamb, B., Lan, X., Marchese, A., Pacala, S., Robinson, A., Shepson, P., Sweeney, C., Talbot, R., Townsend-Small, A., Yacovitch, T., Zimmerle, D. & Hamburg, S. (2015) Reconciling divergent estimates of oil and gas methane emissions. *Proceedings of the National Academy of Sciences of the United States of America* **112**(51): 15597-15602.

## **Annex 6A.1 QC checklists**

### **FORMS AND CHECKLISTS FOR QUALITY CONTROL FOR SPECIFIC SOURCE CATEGORIES**

This annex contains a number of example forms that provide means to record both general and category-specific QC activities. These forms are only examples, and inventory compilers may find other means to effectively record their QA/QC activities (to be defined in the QA/QC plan). Refer to the *2006 IPCC Guidelines* chapters on QA/QC and Verification, Data Collection, and for each category as described in Volumes 2-5 for more detailed guidance on developing QC checks.

#### **A1. GENERAL QC CHECKLIST**

**(to be completed for each category and for each inventory)**

#### **A2. CATEGORY-SPECIFIC QC CHECKLIST (CHECKS TO BE DESIGNED FOR EACH CATEGORY)**

*Part A: Data Gathering and Selection*

*Part B: Secondary Data and Direct Emission Measurement*

## A1. GENERAL QC CHECKLIST

Inventory Report: \_\_\_\_\_ Source/Sink Category<sup>24</sup>: \_\_\_\_\_

Title(s) and Date(s) of Inventory Spreadsheet(s): \_\_\_\_\_

Source (sink) category estimates prepared by (name/affiliation): \_\_\_\_\_

### **INSTRUCTIONS FOR COMPLETING THIS FORM:**

This form is to be completed for each source/sink category, and provides a record of the checks performed and any corrective actions taken. The form may be completed by hand or electronically. The form should be distributed and filed according as specified in the QA/QC plan. If appropriate actions to correct any errors that are found are not immediately apparent, the QC staff performing the check should discuss the results according to the procedures predefined in the QA/QC plan.

The first page of this form summarises the results of the checks (once completed) and highlights any significant findings or actions. The remaining pages in this form list categories of checks to be performed. The analyst has discretion over how the checks are implemented. Not all checks will be applicable to every category. Checks/rows that are not relevant or not available should indicate 'n/r' (not relevant) or 'n/a' (not available) so that no check and no row is left blank or deleted. Rows for additional checks that are relevant to the source/sink category should be added to the form.

The column for supporting documentation should be used to reference any relevant Supplemental Reports or Contact Reports providing additional information.

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### **Summary of general QC checks and corrective action**

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Summary of results of checks and corrective actions taken:

Example -  
design  
your own

Suggested checks to be performed in the future:	Any residual problems after corrective actions have been taken:
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<sup>24</sup> Use IPCC recognized source/sink category names. See Table 8.2 of Chapter 8.

**Checklist for general QC checks (complete table for each category):**

Item	Check completed			Corrective action		Supporting documents (provide reference)
	Date	Individual (first initial, last name)	Errors (Y/N)	Date	Individual (first initial, last name)	
<b>DATA GATHERING, INPUT, AND HANDLING ACTIVITIES: QUALITY CHECKS</b>						
1.	Check a sample of input data for transcription errors					
2.	Review spreadsheets with computerised checks and/or quality check reports					
3.	Identify spreadsheet modifications that could provide additional controls or checks on quality					
4.	Other (specify):					
<b>DATA DOCUMENTATION: QUALITY CHECKS</b>						
5.	Check project file for completeness					
6.	Confirm that bibliographical data references are included (in spreadsheet) for every primary data element					
7.	Check that all appropriate citations from the spreadsheets appear in the inventory document					
8.	Check that all citations in spreadsheets and inventory are complete (i.e., include all relevant information)					
9.	Randomly check bibliographical citations for transcription errors					
10.	Check that originals of new citations are in current docket submittal					
11.	Randomly check that the originals of citations (including Contact Reports) contain the material & content referenced					
12.	Check that assumptions and criteria for selection of activity data, emission factors and other estimation parameters are documented					
13.	Check that changes in data or methodology are documented					
14.	Check that citations in spreadsheets and inventory document conform to acceptable style guidelines					
15.	Other (specify):					

**Checklist for general QC checks (complete table for each category) (Continued):**

Item	Check completed			Corrective action		Supporting documents (provide reference)
	Date	Individual (first initial, last name)	Errors (Y/N)	Date	Individual (first initial, last name)	
<b>CALCULATING EMISSIONS AND CHECKING CALCULATIONS</b>						
16.	Check that all calculations are included (instead of presenting results only)					
17.	Check whether units, parameters, and conversion factors are presented appropriately					
18.	Check if units are properly labelled and correctly carried through from beginning to end of calculation					
19.	Check that conversion factors are correct					
20.	Check that temporal and spatial adjustment factors are used correctly					
21.	Check the data relationships (comparability) and data processing steps (e.g., equations) in the spreadsheets					
22.	Check that spreadsheet input data and calculated data are clearly differentiated					
23.	Check a representative sample of calculations, by hand or electronically					
24.	Check some calculations with abbreviated calculations					
25.	Check the aggregation of data within a category					
26.	When methods or data have changed, check consistency of time series inputs and calculations					
27.	Check current year estimates against previous years (if available) and investigate unexplained departures from trend					
28.	Check value of implied emission/removal factors across time series and investigate unexplained outliers					
29.	Check for any unexplained or unusual trends for activity data or other calculation parameters in time series					
30.	Check for consistency with IPCC inventory guidelines and <i>good practices</i> , particularly if changes occur					
31.	Other (specify):					

## A2. CATEGORY-SPECIFIC QC CHECKLIST

Inventory Report: \_\_\_\_\_ Source/sink Category<sup>25</sup>: \_\_\_\_\_

Key category (or includes a key subcategory): (Y/N): \_\_\_\_\_

Title(s) and Date(s) of Inventory Spreadsheet(s): \_\_\_\_\_

Category estimates prepared by (name/affiliation): \_\_\_\_\_

### **GENERAL INSTRUCTIONS FOR COMPLETING THIS FORM:**

Category-specific checks focus on the particular data and methodology used for an individual source or sink category. The specificity and frequency of these checks will vary across source categories. The form may be completed by hand or electronically. Once completed, the form should be saved and included as part of the inventory archive, as defined in the QA/QC plan.

The first table on this form summarises generally the results of the category-specific checks and highlights any significant findings or corrective actions. The remaining pages in this form list categories of checks to be performed or types of questions to be asked. Part A checks are designed to identify potential problems in the estimates, factors, and activity data. Part B checks focus on the quality of secondary data and direct emission measurement. The analyst has discretion over how the checks are implemented. Checks/rows that are not relevant or not available should indicate 'n/r' (not relevant) or 'n/a' (not available) so that no check and no row is left blank or deleted. Rows for additional checks that are relevant to the category should be added to the form.

The column for supporting documentation should be used to reference any relevant Supplemental Reports or Contact Reports that provide additional information. Other sources may be included here, if they can be clearly referenced. Any documents associated with the category specific plan should be clearly referenced in the column for supporting documentation.

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### **Summary of category-specific QC activities**

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Summary of results of checks and corrective actions taken:

Suggested checks to be performed in the future:

Any residual problems after corrective actions have been taken:

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<sup>25</sup> Use IPCC recognized source/sink category names.

**ADDITIONAL INSTRUCTIONS FOR PART A:**

The checklist below indicates the types of checks and comparisons that can be performed and is not intended to be exhaustive. Supplemental Reports, Contact Reports, or other documents may be used to report detailed information on the checks conducted. For example, a Supplemental Report could provide information on the variables or sub-variables checked, comparisons made, conclusions that were drawn and rationale for conclusions, sources of information (published, unpublished, meetings, etc.) consulted, and corrective actions required.

**Category-specific checklist - Part A: Data gathering and selection**

Item	Check completed			Corrective action		Supporting documents (provide reference)
	Date	Individual (first initial, last name)	Errors (Y/N)	Date	Individual (first initial, last name)	
<b>EMISSION DATA QUALITY CHECKS</b>						
1.	Emission comparisons: historical data for source, significant sub-source categories					
2.	Checks against independent estimates or estimates based on alternative methods					
3.	Reference calculations					
4.	Completeness					
5.	Other (detailed checks)					
<b>EMISSION FACTOR QUALITY CHECK</b>						
6.	Assess representativeness of emission factors, given national circumstances and analogous emissions data					
7.	Compare to alternative factors (e.g., IPCC default, cross-country, literature)					
8.	Search for options for more representative data					
9.	Other (detailed checks)					
<b>ACTIVITY DATA QUALITY CHECK: NATIONAL LEVEL ACTIVITY DATA</b>						
10.	Check historical trends					
11.	Compare multiple reference sources					
12.	Check applicability of data					
13.	Check methodology for filling in time series for data that are not available annually					
14.	Other (detailed checks)					
<b>ACTIVITY DATA QUALITY CHECK: SITE-SPECIFIC ACTIVITY DATA</b>						
15.	Check for inconsistencies across sites					
16.	Compare aggregated and national data					
17.	Other (detailed checks)					

**ADDITIONAL INSTRUCTIONS FOR PART B:**

Completing the QC checks on secondary data and direct emission measurement may require consulting the primary data sources or authors. The checklist below is intended to be indicative, not exhaustive. Additional information on appropriate checks can be found in the QA/QC, Data Collection, and sectoral chapters of the *2006 IPCC Guidelines*.

Additional documentation is likely to be necessary to record the specific actions taken to check the data underlying the category estimates. For example, Supplemental Reports may be needed to record the data or variables that were checked, and the published references and individuals or organisations consulted as part of the investigation. Contact Reports should be used to report the details of personal communications. Supplemental Reports may also be used to explain the rationale for a finding reported in the summary, the results of research into the QC procedures associated with a survey, or checks of site measurement procedures. Be sure to provide references to all supporting documentation.

**Category-specific checklist - Part B: Secondary data and direct emission measurement**

Item	Check completed			Corrective action		Supporting documents (provide reference)
	Date	Individual (first initial, last name)	Errors (Y/N)	Date	Individual (first initial, last name)	
<b>SECONDARY DATA: SAMPLE QUESTIONS REGARDING THE QUALITY OF INPUT DATA</b>						
1. Are QC activities conducted during the original preparation of the data (either as reported in published literature or as indicated by personal communications) consistent with and adequate when compared against (as a minimum), general QC activities?						
2. Does the statistical agency have a QA/QC plan that covers the preparation of the data?						
3. For surveys, what sampling protocols were used and how recently were they reviewed?						
4. For site-specific activity data, are any national or international standards applicable to the measurement of the data? If so, have they been employed?						
5. Have uncertainties in the data been estimated and documented?						
6. Have any limitations of the secondary data been identified and documented, such as biases or incomplete estimates? Have errors been found?						
7. Have the secondary data undergone peer review and, if so, of what nature?						
8. Other (detailed checks)						
<b>DIRECT EMISSION MEASUREMENT: CHECKS ON PROCEDURES TO MEASURE EMISSIONS</b>						
9. Identify which variables rely on direct emission measurement						
10. Check procedures used to measure emissions, including sampling procedures, equipment calibration and maintenance						
11. Identify whether standard procedures have been used, where they exist (such as IPCC methods or ISO standards)						
12. Other (detailed checks)						