

The consolidated European synthesis of CH₄ and N₂O emissions for EU27 and UK: 1990-~~2017~~2019²⁰

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Abstract

Knowledge of the spatial distribution of the fluxes of greenhouse gases and their temporal variability as well as flux attribution to natural and anthropogenic processes is essential to monitoring the progress in mitigating anthropogenic emissions under the Paris Agreement and to inform its Global Stocktake. This study provides a consolidated synthesis of CH₄ and N₂O emissions using bottom-up (BU) and top-down (TD) approaches for the European Union and UK (EU27+UK) and updates earlier syntheses (Petrescu et al., 2020, 2021). The work integrates updated emission inventory data, process-based model results, data-driven sector model results, inverse modelling estimates, and extends the previous period 1990-2017 to 2020-2019. BU and TD products are compared with European National GHG Inventories (NGHGI) reported by Parties under the United Nations Framework Convention on Climate Change (UNFCCC) in 2021. Uncertainties in NGHGIs, as reported to the UNFCCC by the EU and its Member States are also included in the synthesis. Variation in estimates produced with other methods, such as atmospheric inversion models (TD) or spatially disaggregated inventory datasets (BU), arise from diverse sources including within-model uncertainty related to parameterization as well as structural differences between models. By comparing NGHGIs with other approaches, the activities included are a key source of bias between estimates e.g. anthropogenic and natural fluxes, which, in atmospheric inversions are sensitive to the prior geospatial distribution of emissions. **For CH₄ emissions**, over the updated 2015-2019 ~~2011-2015~~ period, which covers a sufficiently robust number of overlapping estimates, and most importantly the NGHGIs, the anthropogenic BU approaches are directly comparable, accounting for mean emissions of ~~20.8-5~~ Tg CH₄ yr⁻¹ (EDGAR ~~v5v6.0~~, last year 2018) and ~~49.0~~18.4 Tg CH₄ yr⁻¹ (GAINS, 2015), close to the NGHGI estimates of ~~48.9~~17.5 ± 2.1 Tg CH₄ yr⁻¹. TD inversions estimates give higher emission estimates, as they also detect natural emissions. Over the same period, high resolution regional TD inversions report a mean emission of ~~28.8~~34 Tg CH₄ yr⁻¹. Coarser-resolution global-scale TD inversions result in emission estimates of 23 Tg CH₄ yr⁻¹ and 24 Tg CH₄ yr⁻¹ inferred from GOSAT and surface (SURF) network atmospheric measurements, respectively. The magnitude of natural peatland and mineral soils emissions from the JSBACH-HIMMELI model, natural rivers, lakes and reservoirs emissions, geological sources and biomass burning together could account for the gap between NGHGI and inversions and account for ~~85.2~~ Tg CH₄ yr⁻¹. **For N₂O emissions**, over the ~~2011-2015~~2015-2019 period, both BU products (EDGAR ~~v5v6.0~~ and GAINS) report a mean value of anthropogenic emissions of 0.9 Tg N₂O yr⁻¹, close to the NGHGI data (0.8 ± 55 % Tg N₂O yr⁻¹). Over the same period, the mean of TD global and regional inversions was 1.4 Tg N₂O yr⁻¹ (excluding TOMCAT which reported no data). The TD and BU comparison method defined in this study can be ‘operationalized’ for future annual updates for the calculation of CH₄ and N₂O budgets at the national and EU27+UK scales. Future comparability will be enhanced with further steps involving analysis at finer temporal resolutions and estimation of emissions over intra-annual timescales, of great importance for CH₄ and N₂O, which may help identify sector contributions to divergence between prior and posterior estimates at the annual/inter-annual scale. Even if currently comparison between CH₄ and N₂O inversions estimates and NGHGIs is highly uncertain because of the large spread in the inversion results, TD inversions inferred from atmospheric observations represent the most independent data against which inventory totals can be compared. With anticipated improvements in atmospheric modelling and observations, as well as modelling of natural fluxes, TD inversions may arguably emerge as the most powerful tool for verifying emissions inventories for CH₄, N₂O and other GHGs. The

referenced datasets related to figures are visualized at <https://doi.org/10.5281/zenodo.6992472> (Petrescu et al., 2022) <https://doi.org/10.5281/zenodo.7553800> (Petrescu et al., 2023).

1. Introduction

Atmospheric concentrations of greenhouse gases (GHGs) reflect a balance between emissions from sources and removals by sinks, the former arising from both human activities and natural sources and the latter being found in the biosphere, oceans and atmospheric oxidation. Increasing levels of GHG in the atmosphere due to human activities have been the major driver of climate change since the pre-industrial period (pre-1750). In 2020⁴, GHG mole fractions were record highs, with globally averaged mole fractions reaching 1889 ± 2 parts per billion (ppb) for methane (CH_4) and 333.2 ± 0.1 ppb for nitrous oxide (N_2O)—333%, representing 262% and 123% of respective pre-industrial levels (WMO, 2021). Since 2004, when CH_4 registered a negative dip, the trend in the CH_4 concentration in the atmosphere continues to increase (NOAA, atmospheric data: https://www.esrl.noaa.gov/gmd/ccgg/trends_ch4/, last access May 2022). This increase was attributed to anthropogenic emissions from agriculture (livestock enteric fermentation and rice cultivation (12%) and fossil fuel related activities (17%)), combined with a contribution from natural tropical wetlands (Saunois et al., 2020, Thompson et al. 2018, Feng et al., 2022a,b). The increase in atmospheric N_2O also continues to rise with the highest annual increase ever recorded in 2020 (https://gml.noaa.gov/ccgg/trends_n2o/, last access: May 2022). The main sources remain linked to agriculture particularly the application of nitrogen fertilizers and livestock manure on agricultural land (FAO, 2020, 2015; IPCC, 2019b, Tian et al., 2020).

National GHG emission inventories (NGHGs) are prepared and reported on an annual basis by Annex I Parties¹ to the United Nations Framework Convention on Climate Change (UNFCCC). These inventories contain annual time series of each country's GHG emissions from the 1990 base year² until two years before the year of reporting and were originally set to track progress towards their reduction targets under the Kyoto Protocol (UNFCCC, 1997). Non-Annex I Parties³ to the UNFCCC also provide emissions estimates in Biennial Update Reports (BURs) as well as through National Communications (NCs); however, non-Annex I emissions are neither reported for annually nor use harmonized formats due to the comparatively less-stringent reporting requirements. Annex I NGHGs are reported according to the Decision 24/CP.19 of the UNFCCC Conference of the Parties (COP) which states that the national inventories shall be compiled using the methodologies provided in the IPCC Guidelines for National Greenhouse Gas Inventories (IPCC, 2006). The 2006 IPCC Guidelines provide methodological guidance for

¹ Annex I Parties include the industrialized countries that were members of the OECD (Organization for Economic Co-operation and Development) in 1992 plus countries with economies in transition (the EIT Parties), including the Russian Federation, the Baltic States, and several central and eastern European states (UNFCCC, <https://unfccc.int/parties-observers>, last access: February 2022). Under the Paris agreement all countries are requested to report their emissions.

² For most Annex I Parties, the historical base year is 1990. However, parties included in Annex I with an economy in transition during the early 1990s (EIT Parties) were allowed to choose one year up to a few years before 1990 as reference because of a non-representative collapse during the breakup of the Soviet Union (e.g., Bulgaria, 1988, Hungary, 1985–1987, Poland, 1988, Romania, 1989, and Slovenia, 1986).

³ Non-Annex I Parties are mostly developing countries. Certain groups of developing countries are recognized by the Convention as being especially vulnerable to the adverse impacts of climate change, including countries with low-lying coastal areas and those prone to desertification and drought. Others (such as countries that rely heavily on income from fossil fuel production and commerce) feel more vulnerable to the potential economic impacts of climate change response measures. The Convention emphasizes activities that promise to answer the special needs and concerns of these vulnerable countries, such as investment, insurance and technology transfer (UNFCCC, <https://unfccc.int/parties-observers>, last access: February 2022).

estimating emissions for well-defined sectors using national activity and available emission factors. Decision trees indicate the appropriate level of methodological sophistication (methodological *Tier*) based the absolute contribution of the sector to the national GHG balance (is the source or sink a *Key Category* or not) and the country's national circumstances (availability and resolution of national activity data and emission factors). Generally, Tier 1 methods are based on global or regional default emission factors that can be used with aggregated activity data, while Tier 2 methods rely on country-specific factors and/or activity data at a higher subsector resolution. Tier 3 methods are based on more detailed process-level modelling or even facility-level emission measurements. Annex I Parties are furthermore required to estimate and report uncertainties in emissions (95% confidence interval) following the 2006 IPCC guidelines using, as a minimum requirement, the Gaussian error propagation method (approach 1). Annex I Parties may use Monte-Carlo methods (approach 2) or a hybrid approach and are encouraged to do so.

Annex I NGHGs should follow principles of transparency, accuracy, consistency, completeness and comparability (TACCC) under the guidance of the UNFCCC (UNFCCC, 2014) and as mentioned above, shall be completed following the 2006 IPCC guidelines (IPCC, 2006). In addition, the IPCC 2019 Refinement (IPCC, 2019a), that may be used to complement the 2006 IPCC guidelines, has updated sectors with additional emission sources and provides guidance on the use of atmospheric data for independent verification of GHG inventories. Complementary to the NGHGs, research groups and international institutions produce estimates of national GHG emissions, with two kind of approaches: atmospheric inversions (top-down, TD) and GHG inventories based on the same principle as NGHGI but using activity and/or emissions factors from (partially) different sources (bottom-up, BU).

The two approaches (BU and TD) provide useful insights on emissions from two different point of view. First, TD approaches act as an additional quality control tool for BU and NGHGI approaches, and facilitates a deeper understanding of the processes driving changes in different elements of GHG budgets. Second, NGHGs cover regularly only a subset of countries (Annex I), and it is therefore necessary to construct BU estimates independently for all countries. Furthermore, while additional BU methods do not have prescribed standards like the IPCC Guidelines, independent BU methods can draw on different input data, or can provide estimates at higher-sectoral resolution, and therefore add complementary information to help quality control NGHGs and help inform climate mitigation policy processes. Additionally, BU estimates are needed as input for TD estimates. As there is no formal guideline to estimate uncertainties in TD or BU approaches, uncertainties are usually assessed from the spread of different estimates within the same approach, though some groups or institutions report uncertainties for their individual estimates using a variety of methods, for instance, by performing sensitivity tests (Monte Carlo approach) on input data parameters. However, this can be logistically and computationally difficult when dealing with complex process-based models.

Despite the important insights gained from complementary BU and TD emission estimates, it should be noted that comparisons with the official reported is not always straightforward. BU estimates often share common methodology and input data, and through harmonization, structural differences between BU estimates and NGHGs can be bridged. However, the use of common input data, albeit to varying extents, restricts the independence between the datasets and, from a verification perspective, may limit the conclusions drawn from the comparisons. On the other hand, TD estimates are constrained by independent atmospheric observations and can serve as an additional, almost-

160 fully independent quality control for NGHGs. Nonetheless, structural differences between NGHGs (what sources
and sinks are included, and where and when emissions/removals occur) and the actual fluxes of GHGs to the
atmosphere must be factored in to the comparison of estimates. While NGHGs go through a central QA/QC review
process, the IPCC procedures do not incorporate mandatory large-scale observation-derived verification.
165 Nevertheless, the individual countries may use atmospheric data and inverse modelling within their data quality
control, quality assurance and verification processes, with expanded and updated guidance provided in chapter 6 of
the 2019 Refinement of IPCC 2006 Guidelines (IPCC, 2019). So far, only a few countries (e.g. Switzerland, UK, New
Zealand and Australia) have used atmospheric observations to constrain national emissions and documented these
verification activities in their national inventory reports (Bergamaschi et al., 2018).

A key priority in the current policy process is to facilitate the 5 yearly Global Stocktakes (GSTs) of the Paris
170 agreement, the first of which is in 2023, and to assess collective progress towards achieving the near- and long-term
objectives, considering mitigation, adaptation and means of implementation. The GSTs are expected to create political
momentum for enhancing commitments in Nationally Determined Contributions (NDCs) under the Paris Agreement.
Though the modalities of the GSTs implementation are not clear, the key component of this process will be the NGHGI
reporting by countries under the Enhanced transparency framework of the Paris Agreement. Under the framework,
175 emissions reporting will move away from the differential Annex I and non-Annex I reporting requirements and
become more harmonized across Parties. Non-Annex I parties will be required to follow the 2006 IPCC guidelines
and provide regular (biennial) national GHG inventory reports to the UNFCCC, alongside developed countries, that
will continue to submit their inventories on an annual basis. Some developing countries will face challenges to
construct and subsequently update their NGHGs and meet the more-stringent reporting requirements.

180 The work presented in this paper covers dozens of distinct datasets and models, in addition to the individual
country submissions to the UNFCCC of the EU Member States and the UK. As Annex I Parties, the NGHGs of the
EU Member States and the UK are consistent with the general guidance laid out in IPCC (2006) yet still differ in
specific approaches, models, and parameters, in addition to differences underlying activity datasets. A comprehensive
investigation of detailed differences between all datasets is beyond the scope of this paper, though systematic analyses
185 have been previously made for specific sectors (e.g. agriculture Petrescu et al., 2020) and by the Global Carbon Project
CH₄ and N₂O syntheses (Saunio et al., 2020 and Tian et al., 2020). The focus of this paper is on updates of the
information from Petrescu et al., 2021a discussing whenever needed the changes in terms of emissions and trends.
The data from Petrescu et al., 2021a is labeled as v2019, while the latest results are labelled ~~v2020~~ and ~~v2021~~
respectively. Except for one on N₂O, the global inversions did not provide an update for v2021, and, therefore, the
190 earlier results are incorporated into this synthesis.

As Petrescu et al. (2021a) is the most comprehensive comparison of the NGHGI and research datasets
(including both TD and BU approaches) for the EU27+UK to date, the focus of the current paper is on improvement
of estimates in the most recent version in comparison with the previous one, including changes in the uncertainty
estimates and identification of the knowledge gaps and added value for policy making. Such exercises of yearly
195 updates are needed to improve the different respective approaches and furthermore can inform the development of
formal verification systems. Official NGHGI emissions are compared with research datasets, including necessary

200 harmonization of the latter on total emissions to ensure consistency. Differences and inconsistencies between emission
estimates were analyzed, and recommendations were made towards future evaluation of NGHGI data. It is important
to remember that uncertainties provided by the NGHGIs are intended to be used in prioritization and decision-making
(Volume 1, Chapter 3, IPCC, 2006) and not to enable comparisons between countries or other datasets. In addition,
individual spatially disaggregated research emission datasets often lack quantification of uncertainty. Here, the focus
is on the median and minimum/maximum (min/max) range of different research products of the same type to get a
first estimate of uncertainty (see Sect. 2). For those datasets providing uncertainties, new uncertainty reduction maps
are presented (see Section 3.1.5). For those models/inventories who did not provided an update for this study, the
205 previously published timeseries are shown. As this is the most comprehensive comparison of NGHGIs and research
datasets (including both TD and BU approaches) for the European continent to date, Official anthropogenic NGHGI
emissions were compared with research datasets, including necessary harmonization of the latter on total emissions to
ensure consistency. Differences and inconsistencies between emissions were analyzed, and recommendations were
made towards future evaluation of NGHGI data. While NGHGI include uncertainty estimates, individual spatially
210 disaggregated research datasets of emissions often lack quantification of uncertainty. Here, the median⁴ and
minimum/maximum (min/max) range of different research products of the same type were used in this work to get a
first estimate of uncertainty.

2. CH₄ and N₂O data sources and estimation approaches

215 The CH₄ and N₂O emissions in the EU27+UK from inversions and anthropogenic emissions inventories from
various BU approaches covering specific sectors were analyzed. The data (Table 2) span the period from 1990 to
2017-202019, with some of the data only available for shorter time periods or up until 2020. The estimates are available
both from peer-reviewed literature and from unpublished research results from the VERIFY project (Table 1 and
Appendix A) and in this work they are compared with NGHGIs reported in 2021 (time series for 1990-2017-2019).
220 Data sources are summarized in Table 2 with the detailed description of all products provided in Appendix A1-A3.

For both CH₄ and N₂O BU approaches, inventories of anthropogenic emissions covering all sectors (EDGAR
v5v6.0 and GAINS) and models and inventories limited to agriculture (CAPRI, FAOSTAT, DayCent, ECOSSE) were
used. For CH₄ biogeochemical models of natural peatland emissions (JSBACH-HIMMELI), and lakes and reservoirs
emissions (Lauerwald et al., 2019; Maisonnier et al., in prep.), as well as updated data for inland waters (rivers, lakes
225 and reservoirs; Lauerwald et al., in rev. GBC) and updated data for total geological emissions (Etiope et al., 2019)
were used. Emissions from gas hydrates and termites are not included as they are close to zero in the EU27+UK
(Saunois et al., 2020). Anthropogenic NGHGI CH₄ emissions from the Land Use, Land Use Change and Forestry
(LULUCF) sector are very small for EU27+UK (3 % in 2019 including biomass burning) (section 2.2).

TD approaches include both regional and global inversions, the latter having a coarser spatial resolution.
230 These estimates are described in section 2.3.

⁴-The reason for using median instead of mean for the ensembles is because there is a large spread between global inversions and we don't want to be biased by outliers/extremes.

For N₂O emissions, the same global BU inventories as for CH₄, natural emissions from inland waters (rivers, lakes and reservoirs) ([RECCAP2 Lauerwald et al., in rev. GBC](#)) were used, which did not change with respect to [Petrescu et al., 2021a](#). In this study, about 66 % of the N₂O emitted by Europe's natural rivers are considered anthropogenic indirect emissions, caused by leaching and run-off of N-fertilizers from the agriculture sector. [One important update is the inclusion of estimates of natural N₂O emissions from soils simulated with the O-CN model \(Zaehle et al., 2011\). These emissions are derived from model simulations in which land-use and atmospheric CO₂ remain constant, but climate varies through to 2020. These estimates are considered to be closer to what background natural N₂O emissions would be present day, so they were used for subtraction from outputs of inversions \(as it has a reasonable representation of the inter-annual variability \(IAV\)\).](#) The TD N₂O inversions include one regional inversion FLEXINVERT and three global inversions (Friedlingstein et al., 2019; Tian et al., 2020, [Patra et al., 2022](#)). Agricultural sector emissions of N₂O were presented in detail by Petrescu et al., 2020. In this current study, CAPRI and ECOSSE models and FAO provided updated emissions, with the latter additionally covering non-CO₂ emissions from biomass burning as a contribution to LULUCF. Fossil fuel related emissions and industrial emissions were obtained from GAINS (see Appendix A1). Table A2 in Appendix A presents the methodological differences of the current study with respect to Petrescu et al., 2020 and Petrescu et al., 2021a.

Table 1: Sectors included in this study and data sources providing estimates for these sectors.

Anthropogenic (BU) ⁵ CH ₄ and N ₂ O	Natural (BU) ⁶ CH ₄	Natural** (BU) N ₂ O	TD (CH ₄ and N ₂ O)
1. Energy (NGHGI, GAINS, EDGAR v65.0)	Peatlands, mineral soils (JSBACH-HIMMELI), inland waters fluxes (lakes, rivers and reservoirs and RECCAP2 estimate, Rosentreter et al. 2021), geological fluxes Etiope et al., 2019 with updated activity (this study) biomass burning (GFEDv4.1)	Inland water fluxes (lakes, rivers and reservoirs and RECCAP2 estimate, biomass burning (GFEDv4.1), pre-industrial natural soil emissions (O-CN)	No sectoral split – total emissions FLEXKF (CAMSV19r); TM5-4DVAR; FLEXINVERT; CTE-CH₄ InGOS inversions GCP-CH₄ 2019 anthropogenic partition from inversions GCP-CH₄ 2019 Natural partition from inversions GN2OB 2019 CHIMERE InTEM NAME (only for UK) VERIFY Community Inversion Framework (CIF): CHIMERE, FLEXPARTv10.4 (NILU)
2. Industrial Products and Products in Use (IPPU) (NGHGI, GAINS, EDGAR v65.0)			
3. Agriculture* (NGHGI, CAPRI, GAINS, EDGAR v5v6.0, FAOSTAT, ECOSSE and DayCent (only for N ₂ O))			
4. LULUCF total emissions (NGHGI Fig. 1,2,4,6, B1a for CH ₄ and Fig. 10, 11, 14 and B1b for N ₂ O)			

⁵ For consistency with the NGHGI, here we refer to the five reporting sectors as defined by the UNFCCC and the Paris Agreement decision (18/CMP.1), the IPCC Guidelines (IPCC, 2006), and their Refinement (IPCC, 2019a), with the only exception that the latest IPCC Refinement groups together Agriculture and LULUCF sectors in one sector (Agriculture, Forestry and Other land Use - AFOLU).

⁶ The term natural refers here to unmanaged natural CH₄ emissions ([peatlands, mineral soils, geological, inland waters and biomass burning](#)) not reported under the UNFCCC LULUCF sector.

5. Waste (NGHGI, GAINS, v5y6.0)	(NGHGI, EDGAR)			<u>and FELXPART (EMPA) (only CH₄)</u>
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* Anthropogenic (managed) agricultural soils can also have a level of natural emissions.

**Natural soils (unmanaged) can have both natural and anthropogenic emissions.

The units used in this paper are metric tonne (t) [1kt = 10⁹ g; 1Mt = 10¹²g] of CH₄ and N₂O. The referenced data used for the figures' replicability purposes are available for download at <https://doi.org/10.5281/zenodo.7553800> and [10.5281/zenodo.6992472](https://doi.org/10.5281/zenodo.6992472) (Petrescu et al., 2022, 2023). Upon request, the codes necessary to plot the figures in the same style and layout can be provided. The focus is on EU27+UK emissions. In the VERIFY project, an additional web tool was developed which allows for the selection and display of all plots shown in this paper (as well as the companion paper on CO₂), not only for the EU Member States and UK but for a total of 79 countries and groups of countries in Europe (Table A1, Appendix A). The data, located on the VERIFY project website: <http://webportals.ipsl.jussieu.fr/VERIFY/FactSheets/>, is free and can be accessed upon registration.

2.1. CH₄ and N₂O anthropogenic emissions from NGHGI

Anthropogenic CH₄ emissions from the four UNFCCC sectors (excluding LULUCF) were grouped together. Anthropogenic CH₄ emissions in 2019 account for 17.1 Tg CH₄ yr⁻¹ and represent 10.5 % of the total EU27+UK emissions (in CO₂e, GWP 100 years, IPCC AR4⁷). CH₄ emissions are predominantly related to agriculture (9.2 Tg CH₄ yr⁻¹ ± 0.8 Tg CH₄ yr⁻¹) or 53.8 % in 2019 (52.5 % in 2018) of the total EU27+UK CH₄ emissions. Anthropogenic NGHGI CH₄ emissions from the LULUCF sector are very small for EU27+UK e.g. 0.5 Tg CH₄ yr⁻¹ or 3 % in 2019, including emissions from biomass burning.

Regarding CH₄ emissions from wetlands, following the recommendations of the 2013 IPCC Wetlands supplement (IPCC, 2014) only emissions from managed wetlands are reported by Parties. According to NGHGI data between 2008 and 2018, managed wetlands in the EU27+UK for which emissions were reported under LULUCF (CRF table 4(II) accessible for each EU27+UK country⁸) represent one fourth of the total wetland area in EU27+UK (G. Grassi, EC-JRC, pers. comm.) and their emissions summed up in 2019 to 0.1 Tg CH₄ yr⁻¹.

Anthropogenic N₂O emissions (excluding LULUCF) in 2019 account for 0.8 Tg N₂O yr⁻¹ and represent 6.2 %- of the total EU27+UK emissions in CO₂eq. N₂O emissions are predominantly related to agriculture (0.6 Tg N₂O yr⁻¹ or 73.0 % in 2019 (73.5 % in 2018) of the total EU27+UK (including LULUCF+BB) N₂O emissions) but are also found in the other sectors (Tian et al. 2020). In addition, N₂O has natural sources, which are defined as the pre-industrial background emissions before the use of synthetic N-fertilizers and intensive agriculture, and derive from natural processes in soils but also in lakes, rivers and reservoirs (Lauerwald et al., in rev. GBC, Maavara et al., 2019; Lauerwald et al., 2019; Tian et al., 2020).

⁷ IPCC AR4 GWP 100 values are still used by the Member States in their NGHGI reporting to the UNFCCC.

⁸ <https://unfccc.int/process-and-meetings/transparency-and-reporting/reporting-and-review-under-the-convention/greenhouse-gas-inventories-annex-i-parties/national-inventory-submissions-2019>

2.2. CH₄ and N₂O anthropogenic and natural emissions from other bottom-up estimates

Data from five global data sets and models of CH₄ and N₂O anthropogenic emissions inventories were used, namely: CAPRI, DayCent, ECOSSE, FAOSTAT, GAINS and EDGAR v6.0 (Table 3). These estimates are not completely independent from NGHGIs (see Figure 4 in Petrescu et al., 2020) as they integrate their own sectorial modelling with the UNFCCC data (e.g., common activity data and IPCC emission factors) when no other source of information is available. The CH₄ biomass and biofuel burning emissions are included in NGHGI under the UNFCCC LULUCF sector, although they are identified as a separate category by the Global Carbon Project CH₄ budget synthesis (Saunois et al., 2020). For both CH₄ and N₂O, CAPRI (Britz and Witzke, 2014; Weiss and Leip, 2012) and FAOSTAT (FAO, 2021) report only agricultural emissions. DayCent and ECOSSE report only emissions for agriculture N₂O. Out of all BU inventories, only CAPRI reported new uncertainties for 2014, 2016 and 2018, while values for EDGARv6.0 were the same (Solazzo et al., 2021) as those reported in Petrescu et al., 2021a.

In this study, natural CH₄ emissions are included under the category “peatlands” and “other natural emissions”, the latter including geological emissions, biomass burning emissions and two estimates of inland waters (rivers, lakes and reservoirs). One inland water estimate comes from “process-based models” and is based on the Rosentreter et al., (2021) with ranges from Bastviken et al., (2011) and Stanley et al., (2016) and the second represent an upscaled estimate for inland waters from the RECCAP2 project (Lauerwald et al., in rev. GBC).

For peatlands and mineral soils, the JSBACH-HIMMELI framework was used. Additionally, the ensemble of thirteen monthly gridded estimates of peatland emissions based on different land surface models as calculated for Saunois et al. (2020) were used as described in Appendix B2. Geological emissions were initially based on the global gridded emissions from Etiope et al. 2019 and previously estimated to be 1.3 Tg CH₄ yr⁻¹ (Petrescu et al., 2021a). For this study these emissions were recalculated, using more detailed input data related to the activity, i.e., a more precise estimate of the continental oil-gas field area (which determines the potential area of microseepage) and offshore seepage area (Appendix A2) and now account for 3.3 Tg CH₄ yr⁻¹ (0.9 Tg CH₄ yr⁻¹ from offshore marine seepage and 2.4 Tg CH₄ yr⁻¹ onshore). This rescaled geological source represents the second largest natural component accounting for 42 % of the total EU27+UK natural CH₄ emissions. The upscaled inland waters, (rivers, lakes, and reservoirs, based on Lauerwald et al., in rev. GBC) are the largest component of natural emissions (3.3 Tg CH₄ yr⁻¹ and ranging from 2.7 Tg CH₄ yr⁻¹ to 4.3 Tg CH₄ yr⁻¹) and account for 44 %. The remaining 14 % emissions are attributed to peatlands, mineral soils and biomass burning. Overall, in EU27+UK the natural emissions thus accounted for 8 Tg CH₄ yr⁻¹. Finally, It should be noted that to a small extent the CH₄ natural emissions from waters are also due to an anthropogenic component, namely eutrophication following N-fertilizer leaching to inland waters. Globally, the contribution of eutrophication is estimated to lead to a further increase in lake and reservoir emission by 30 to 90% over the 21st century, which would be the result of a ~3 times higher nutrient loading to lakes and reservoirs (Beaulieu et al., 2019), similar to the review by Li et al. 2021 who gathered a lot of prove that eutrophication significantly increase CH₄ emissions. In temperate Europe, eutrophication contributes significantly to the overall increase natural emissions and Rinta et al., 2017 found that eutrophic, central European lakes show CH₄ emission rates which are about

one order of magnitude higher than those of oligotrophic boreal lakes and this study's model results are consistent with it.

The N₂O anthropogenic emissions from inventory datasets belong predominantly to agriculture and are associated to two main categories: 1) direct emissions from the agricultural sector where synthetic fertilizers and manure were applied, and from manure management, and 2) indirect emissions on non-agricultural land and water receiving anthropogenic N through atmospheric N deposition, leaching and run-off (also from agricultural land). Additional anthropogenic emissions result from industrial processes, in particular, adipic and nitric acid production, which are declining owing to the implementation of emission abatement technologies. Other N₂O emissions come from the wastewater treatment activities and fossil fuel combustion.

In this study, “natural” N₂O fluxes refer to emissions from inland waters (lakes, rivers and reservoirs, Maavara et al., 2019; Lauerwald et al., 2019, Lauerwald et al., in rev. GBC) which include also lakes with dams. The other component is the natural N₂O emissions from soils simulated with the O-CN model (Zaehle et al., 2011). Regarding the inland water emissions, more than half of the emissions (56 % globally, Tian et al., 2020, and 66 % for Europe this study) are due to enhanced N inputs from fertilizers, manure, sewage and, to a smaller extent, atmospheric N deposition. However, emissions from natural soils in this study are considered as “anthropogenic” because, according to the country specific National Inventory Reports (NIRs), all land in EU27+UK is considered to be managed

For both CH₄ and N₂O the natural biomass burning emissions from GFEDv4.1 (van der Werf et al., 2017) are included in Figs. 1, 4b, 5b, 9 and 13, while for CH₄ only, biomass burning emissions from the GCP 2020 (Saunois et al., 2020) are included in Fig 6.

2.3. CH₄ and N₂O emission data from inversions

Atmospheric inversions optimize prior estimates of emissions and sinks through modeling frameworks that utilizes atmospheric observations as a constraint on fluxes. Emission estimates from inversions depend on the data set of atmospheric measurements and the choice of the atmospheric model, as well as on other inputs (e.g., prior emissions and their uncertainties). Inversion results were taken from original publications without evaluation of their performance through specific metrics (e.g., fit to independent cross validation atmospheric measurements (Bergamaschi et al., 2013, 2018; Patra et al., 2016)). Some of the inversions allow for explicit attribution to different sectors, while others optimize all fluxes in each grid cell and then attribute emissions to sectors using prior grid-cell fractions (see details in Saunois et al. 2020 for global inversions).

For CH₄, the same set of nine regional inversions and 22 global inversions as listed in Table 3 and presented in Petrescu et al., 2021a was used. While many different inversions exist, it should be stressed that the variants are not completely independent of one another. Table B4, Appendix B in Petrescu et al., 2021a illustrates this by documenting to what extent the transport models, priors and atmospheric measurement data vary between the inversion datasets”. The subset of InGOS inversions (Bergamaschi et al., 2018a) belongs to a project where all models used the same atmospheric data over Europe covering the period 2006-2012. The global inversions from Saunois et al. 2020 were not updated for this work and cover a period until 2017.

The regional inversions generally use both higher-resolution prior data and higher-resolution transport models, and e.g. TM5-JRC runs simultaneously over the global domain at coarse resolution and over the European domain at higher resolution, with atmospheric CH₄ concentration boundary conditions taken from global fields. For CH₄, 11 global inversions use GOSAT for the period 2010-2017, eight global inversions use surface stations (SURF) since from 2000 to 2017, two global models use SURF since from 2010-2017 and one SURF from 2003-2017 (see “Appendix 4 Table” in Saunio et al. 2020). All regional inversions use observations from SURF stations as a base of their emission calculation.

Table 2: Data sources for CH₄ and N₂O emissions used in this study:

<u>Name</u>	<u>CH₄</u>	<u>N₂O</u>	<u>Contact / lab</u>	<u>References</u>	<u>Status compared to Petrescu et al., 2021a</u>
<u>CH₄ and N₂O Bottom-up anthropogenic</u>					
<u>UNFCCC NGHGI (2021) CRFs</u>	<u>CH₄ emissions</u> <u>1990-2019</u>	<u>N₂O emissions</u> <u>1990-2019</u>	<u>MS inventory agencies</u> <u>Yearly uncertainties provided by the EU GHG inventory team</u>	<u>UNFCCC CRFs</u> <u>https://unfccc.int/process-and-meetings/transparency-and-reporting/reporting-and-review-under-the-convention/greenhouse-gas-inventories-annex-i-parties/national-inventory-submissions-2019</u>	<u>updated</u>
<u>EDGAR v6.0</u>	<u>CH₄ sectoral emissions</u> <u>1990-2018</u>	<u>N₂O sectoral emissions</u> <u>1990-2018</u>	<u>EC-JRC</u>	<u>Crippa et al., 2019a</u> <u>Crippa et al., 2019 EU REPORT</u> <u>Janssens-Maenhout et al., 2019</u> <u>Solazzo et al., 2020 (in review ACP)</u>	<u>Updated</u>
<u>CAPRI</u>	<u>CH₄ agricultural emissions</u> <u>1990-2014 and 2016, 2018</u>	<u>N₂O agricultural emissions</u> <u>1990-2014 and 2016, 2018</u>	<u>EC-JRC</u>	<u>Britz and Witzke, 2014</u> <u>Weiss and Leip, 2012</u>	<u>Updated</u>
<u>GAINS</u>	<u>CH₄ sectoral emissions</u> <u>1990-2015</u>	<u>N₂O sectoral emissions</u> <u>1990-2015</u> <u>(every five years)</u>	<u>IIASA</u>	<u>Höglund-Isaksson, L. 2017</u> <u>Höglund-Isaksson, L. et al., 2020</u> <u>Winiwarter et al., 2018</u>	<u>Not updated</u>
<u>FAOSTAT</u>	<u>CH₄ agriculture and land use emissions</u>	<u>N₂O agricultural emissions</u>	<u>FAO</u>	<u>Tubiello et al. 2013</u> <u>FAO, 2015, 2020</u>	<u>Updated</u>

	1990-2019	1990-2019		Tubiello, 2019	
ECOSSE		Direct N₂O emissions from agricultural soils 2000-2020	UNIABD N	Bradbury et al., 1993 Coleman., 1996 Jenkinson., 1977, 1987 Smith et al., 1996, 2010a,b	Updated
DayCent		N₂O emissions from direct agricultural soils avg. 2015-2019	EC-JRC	Orgiazzi et al., 2018 Lugato et al., 2018, 2017 Quemada et al., 2020	Updated
CH₄ and N₂O natural					
JSBACH-HIMMELI	CH₄ emissions from peatlands and mineral soils 2005-2020		FMI	Raivonen et al., 2017 Susiluoto et al., 2018	Updated
Non-wetland inland waters	One average value for CH₄ fluxes from rivers, lakes and reservoirs with uncertainty 2010-2019 One median upscaled value from RECCAP2 analysis 1990-2019	One median N₂O value for emissions from lakes, rivers, reservoirs from the RECCAP2 analysis 1990-2019	ULB	Maisonnier et al., in prep., after Maavara et al., 2017, 2019 and Lauerwald et al., 2019 Bastviken et al., 2011 Stanley et al. 2016 Rosentreter et al., 2021 Lauerwald et al., in rev. GBC	Updated
Geological emissions, (onshore and offshore)	Global grid geological CH₄ emission model (2019)		Istituto Nazionale di Geofisica e Vulcanologia (INGV)	Etiopie et al., 2019 and this work (updated activity data)	updated
GFED4.1	Biomass burning emissions	Biomass burning emissions	VU Amsterdam	van der Werf et al., 2017	new

	2000-2020	2000-2020			
O-CN		Background natural N₂O emissions from soils (model simulations in which land-use and atm. CO₂ remain constant, but climate varies through to 2020)	MPI-BGC	Zaehle et al., 2011 Zaehle & Friend, 2010	new
CH₄ and N₂O inversions					
Regional inversions over Europe (high transport model resolution)					
FLEXPART- FLEXKF-TM5- 4DVARCAMSv19r	Total CH₄ emissions from inversions with uncertainty 2005-2019		EMPA	Brunner et al., 2012 Brunner et al., 2017 Background concentrations from CAMSv19r (Arjo Segers)	Updated
TM5-4DVAR	CH₄ emissions from inversions, split into total, anthropogenic and natural 2005-2018		EC-JRC	Bergamaschi et al., 2018a	Not updated
FLEXINVERT	CH₄ total emissions from inversions 2005-2018	N₂O total emissions, 2005-2019	NILU	Thompson and Stohl, 2014	Updated for N₂O
CTE-CH₄	Total CH₄ emissions from inversions for Europe with uncertainty 2005-2018		FMI	Brühl et al., 2014 Houweling et al., 2014 Giglio et al., 2013 Ito et al., 2012 Janssens-Maenhout et al., 2013 Krol et al., 2005 Peters et al., 2005 Saunio et al., 2020 Stocker et al., 2014 Tsuruta et al., 2017	Not updated
InGOS inversions	Total CH₄ emissions from inversions		EC-JRC and InGOS	Bergamaschi et al., 2018a	Not updated

	2006-2012		project partners	TM5-4DVAR: Meirink et al., 2008; Bergamaschi et al. 2010; 2015 TM5-CTE: Tsuruta et al., 2017 LMDZ-4DVAR: Hourdin and Armengaud, 1999; Hourdin et al., 2006 TM3-STILT: Trusilova et al., 2010, Gerbig et al., 2003; Lin et al., 2003; Heimann and Koerner, 2003 NAME: Manning et al. 2011; Bergamaschi et al., 2015 CHIMERE: Berchet et al. 2015a; 2015b; Menut et al., 2013; Bousquet et al., 2011\ COMET: Eisma et al., 1995; Vermeulen et al., 1999; Vermeulen et al., 2006	
VERIFY Community Inversion Framework (CIF): CHIMERE, FLEXPARTv10.4 (NILU) and FELXPART (EMPA) (only CH₄)	Total CH₄ emissions from inversions 2006-2017	Total N₂O emissions from inversions 2005-2018	LSCE, NILU and EMAP	Berchet et al., 2021 Fortems-Cheiney et al., 2021	New datasets
Global inversions from the Global Carbon Project CH₄ and N₂O budgets (Saunois et al. 2020, Tian et al., 2020)					
GCP-CH₄ 2019 anthropogenic partition from inversions	22 models for CH₄ inversions, both <i>SURF</i> and <i>GOSAT</i> 2000-2017		LSCE and GCP-CH₄ contributors	Saunois et al., 2020 and model specific references in Appendix B, Table B4	Not updated
GCP-CH₄ 2019 Natural partition from inversions	22 models with optimized wetland CH₄ emissions 2000-2017		LSCE	Saunois et al., 2020 and model specific references in Appendix B, Table B3	Not updated
GN₂OB 2019		Inverse N₂O emissions - 3 Inversions PYVAR (CAMS-N₂O) TOMCAT MIROC4-ACTM 1998-2016	GN₂OB 2019 and contributors	Thompson et al., 2019 Tian et al., 2020	Not updated

For N₂O, one regional inversion (FLEXINVERT) for the 2005-2019 period and three global inversions for the period 1998-2016 from Tian et al., (2020) and Thompson et al. (2019) were used as listed in Table 3. These estimates were not updated for this paper. These inversions are not completely independent from each other since most of them use the same input information (Appendix B3). The regional inversion uses a higher resolution atmospheric transport model for Europe, with atmospheric N₂O concentration boundary conditions taken from global model fields. As all inversions produced total rather than anthropogenic emissions, emissions from soils (O-CN) and inland waters (lakes, rivers and reservoirs) estimated by Lauerwald et al., in rev. GBC were subtracted from the total emissions. Note that inland water emissions include anthropogenic emissions from N-fertilizer leaching accounting for 66 % of the inland water emissions in EU27+UK. In 2016-2019, emissions from inland waters represented 2-21.4 % of the total UNFCCC NGHGI (2021) N₂O emissions.

The largest share of N₂O emissions comes from agricultural soils (direct and indirect emissions from the applications of fertilizers, whether synthetic or manure) contributing in 2019 79 % of the total N₂O emissions (excluding LULUCF) in EU27+UK. In Petrescu et al., 2021a, “Table B1c, Appendix B1” presented the allocation of emissions by activity type covering all agricultural activities and natural emissions, following the IPCC (2006) sector classification scheme. Each data product has its own particular way of grouping emissions, and does not necessarily cover all emissions activities. The main inconsistencies between process-based models and inventories are observed regarding activity allocation in the two models, ECOSSE and DayCent. ECOSSE only estimates direct N₂O emissions, and does not estimate downstream emissions of N₂O, for example indirect emissions from nitrate leached into water courses, which also contributes to an underestimation of total N₂O emissions. Field burning emissions are also not included by most of the data sources.

3. Results and discussion

3.1. Comparing CH₄ emission estimates from different approaches

3.1.1. Estimates of European and regional total CH₄ fluxes

Total CH₄ fluxes from EU27+UK and five main regions in Europe: North, West, Central, East (non-EU) and South are presented in the paper. The countries included in these regions, which include countries outside the EU27+UK bloc, are all Annex I Parties to UNFCCC and are listed in Appendix A, Table A. Figure 1 shows the total CH₄ fluxes from the NGHGIS for base year 1990, as well as five-year mean values for the 2011-2015 and 2015-2019 periods. The five-year periods are informing on emission trends and what could be achieved by the GST process. Given that the GST is only repeated every five years, a five-year average is clearly of interest even if, in this current study 2021 estimates are not available. Figure 1 shows the total CH₄ fluxes from the NGHGIS for base year 1990, as well as five-year mean values for the 2011-2015 and 2015-2019 periods. We use the five-year periods as an exercise for what could be achieved in 2023, the year of the first GST, when for most parties to the Convention the reported inventories will include 2021. Given that the GST is only repeated every five years, a five-year average is clearly of interest.

The total NGHGI estimates include emissions from all sectors (excluding LULUCF) and are plotted and compared to fluxes from global datasets, BU models and inversions. There is a good agreement noted in absolute total values between inventories, as well as between regional and global inversion ensembles, but uncertainties (min/max ranges) are large. This match can be explained by interdependencies in input data (AD and EFs) for the BU estimates (Petrescu et al., 2020) and similar prior information used by inversions (Petrescu et al., 2021a). In Figure 1, hatched transparent bars represent the 2011-2015 mean while colour-filled bars represent the new updated 2015-2019 mean values. For GAINS and some inversions that do not have annual estimates for all five years, only the average of available years is calculated (e.g., 2015 for GAINS).

For all study regions, 2019 CH₄ emissions decreased by 24 % (Southern Europe) to 57 % (Eastern Europe), with respect to NGHGI 1990 values; and for EU27+UK emissions decreased by 39 %. The decrease in CH₄ emissions is mainly due to the EU legislation policies and strategies starting with the implementation in the early 90s of European and country-specific emission reduction policies on agriculture and the environment, as well as socioeconomic changes in the sector resulting in overall lower agricultural livestock and lower emissions from managed waste disposal on land and from agricultural soils. After 2005, these trends maintain their decreasing trajectory, even if, at a lower intensity. For the Central and Eastern Europe, reductions were abrupt and mainly due to the dissolution of the Soviet Union (1989–1991) and the consequent structural changes in the economy of the former eastern European communist centralized economy block (Petrescu et al., 2020). This is encouraging in the context of meeting EU total GHG commitments under the Paris Agreement (55% decrease in 2030 compared to 1990 levels and reaching carbon neutrality by 2050). This reduction will need to be achieved by strong reductions in top emitter sectors (e.g. Agriculture) and compensated by sinks in the LULUCF sector. It also shows that not only at EU27+UK level, but also at regional European level, the emissions from BU (anthropogenic and natural) and TD estimates agree in magnitude with reported NGHGI data despite the high uncertainty associated with the TD estimates. This uncertainty is represented here by the variability in the model ensembles and denotes the range (min and max) of estimates within each model ensemble. The comparison of TD to anthropogenic estimates (Fig. 1), suggests that the total CH₄ flux is dominated by natural emissions (i.e., Northern Europe) although comparison with EDGAR v6.0 would indicate that anthropogenic emissions are dominant (e.g. Northern, Central and Western Europe).

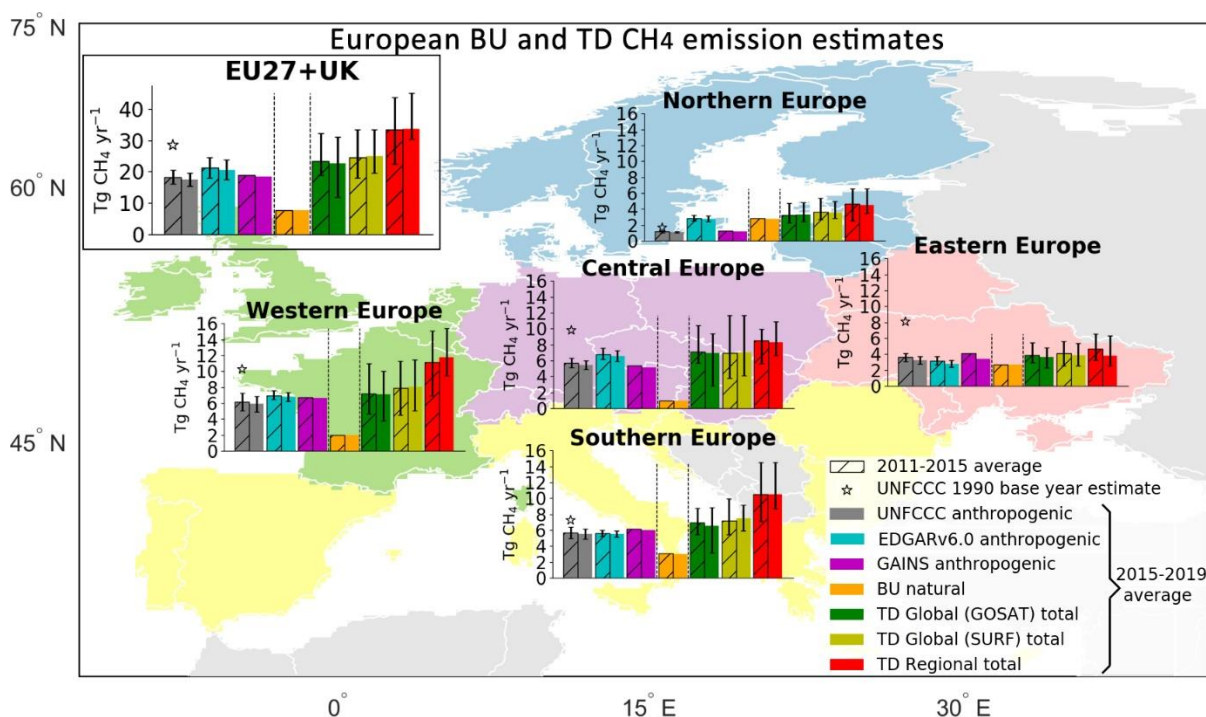


Figure 1: Five-year means (2011-2015 hashed bars and 2015-2019 full bars) in total CH₄ emission estimates (excluding LULUCF) for EU27+UK and five European regions (North, West, Central, South and East non-EU). Eastern European region does not include European Russia. Northern Europe includes Norway. Central Europe includes Switzerland. The data comes from UNFCCC NGHGI (2021) submissions (grey), which are plotted with respective base year 1990 (black star) estimates, two inventories (GAINS and EDGAR v6.0), natural unmanaged emissions (sum of peatland, geological, inland waters (RECCAP2) and GFEDv4.1 biomass burning emissions) and three inversion estimates: one regional European inversion (excluding InGOS unavailable for 2013-2015) and GOSAT and SURF ensemble estimates from global inverse models. The relative error on the UNFCCC value represents the NGHGI (2021) reported uncertainties computed with the error propagation method (95% confidence interval) and gap-filled to provide respective estimates for each year. Uncertainty for EDGAR v6.0 was calculated for 2015 based on the 95 % confidence interval of a lognormal distribution (Solazzo et al., 2021).

The EDGAR v6.0 updated estimates for Northern Europe remain two-times higher than NGHGI and GAINS ones. The EDGAR approach is to use a globally harmonised methods and sources of data, which means that country-specific detail is often replaced with global averages. In some countries and for some sectors or gases, these assumptions lead to huge differences. For example, fugitive emissions of methane in the oil and gas sector are estimated based on the level of production of oil and gas. In the case of Norway this ignores the substantial effects of regulation on reducing such fugitive emissions. Instead, EDGAR's methane emissions estimates for Norway follow the pattern of its total production of oil and gas (Olhoff et al., 2022). For Eastern Europe we note that all estimates decreased compared to the previous five year mean and the BU anthropogenic estimates remain similar in magnitude to the TD estimates of total CH₄ emissions. One possible explanation is that for TD estimates (i.e. using atmospheric

inversions) the fluxes are better constrained by a larger number of observations. Where there are fewer or no observations, like in Eastern Europe, the fluxes in the inversion will stay close to the prior estimates, since there is little or no information to adjust them.

In line with Bergamaschi et al., 2018a the potentially significant contribution from natural unmanaged sources (peatlands, mineral soils, geological and inland waters (RECCAP2)), which for EU27+UK accounted in 2019 for 5.24 8 Tg CH₄ yr⁻¹ (Figure 1) can be highlighted. Taking into account these natural unmanaged CH₄ emissions, and adding it to the range of the BU anthropogenic estimates (4722 Tg CH₄ yr⁻¹ (NGHGI) – 24–26 Tg CH₄ yr⁻¹ (EDGARv6.0)) improves agreement with the TD estimates. BU estimates become consistent with the lower range of the regional total TD estimates (23–32 Tg CH₄ yr⁻¹ (TM5_JRC)– 28–41 Tg CH₄ yr⁻¹ (FLEXINVERT)) and show even better agreement in absolute values with the global median SURF (24 Tg CH₄ yr⁻¹) and GOSAT (23 Tg CH₄ yr⁻¹) inversions. The broad consistency between the TD and BU estimates could be interpreted in two ways: 1) BU and TD regional estimates are similar given the large uncertainties and spread in TD results, or 2) regional TD higher estimates potentially indicate shortcomings of BU inventories, the latter interpretation being more consistent with the general atmospheric developments (WMO, 2021).

Is it notable to highlight that the regional TD total is considerably higher for all regions and EU27+UK total and by considering this estimate the best to date total estimate for the whole Europe, including all sources and sinks, this would infer a missing of 20 to 30 % of CH₄ emissions from the other BU approaches.

3.1.2. NGHGI sectoral emissions and decadal changes

According to the UNFCCC (20192021) NGHGI estimates, in 2017–2019 the EU27+UK emitted GHGs totaling 3.97 Gt CO₂e (including LULUCF), of this total, CH₄ emissions accounted for 11.8 % (0.4 Gt CO₂e or 1817.45 Tg CH₄ yr⁻¹ ± 2.2 Tg CH₄ yr⁻¹) (Appendix, B2, Figure B2a) with France, UK and Germany together contributing 3637 % of total CH₄ emissions.

The data in Figure 2 shows anthropogenic CH₄ emissions and their change from one decade to the next, from UNFCCC NGHGI (20192021), with the split between the different sectors. In 20172019, NGHGI report CH₄ from agricultural activities to be 52.4 % (± 10.8.7 %) of the total EU27+UK CH₄ emissions, followed by emissions from waste, 27.5 % (± 22.53 %). The large share of agriculture in total anthropogenic CH₄ emissions also holds at global level (IPCC Special Report on Climate Change and Land (SRCCCL), 2019). Between the 1990s and the 2000s, the net 17.7–6 % reduction originates largely from the energy and waste sectors, with only negligible contributions to emission trends and levels from IPPU (metal and chemical industry) and LULUCF. Between the 2000s and 2010–20172019, a further reduction by 16.5 % is observed with the waste sector as the largest contributor to this reduction. The two largest sectors contributing to total EU27+UK emission are agriculture and waste, but energy and waste are showing the higher reductions over the last decade.

The reduction observed in the waste sector coincide with the adoption of the first EU methane strategy published in 1996 (COM(96) 557, 1996). EU legislation addressing emissions in the waste sector may have been successful to trigger the largest reductions. Directive 1999/31/ EC on the landfill (also referred to as the Landfill Directive) required the Member States to separate waste, minimizing the amount of biodegradable waste disposed

untreated in landfills and to install landfill gas recovery at all new sites. Based on the 1999 Directive, the new 2018/1999 EU Regulation on the Governance of the Energy Union requires the European Commission to propose a strategic plan for methane, which will become an integral part of the EU's long-term strategy. In the waste sector, the key proposal included the adoption of EU legislation requiring the installation of methane recovery and use systems at new and existing landfills. Other suggested actions included measures aimed at the minimization, separate collection and material recovery of organic waste (Olczak and Piebalgs, 2019).

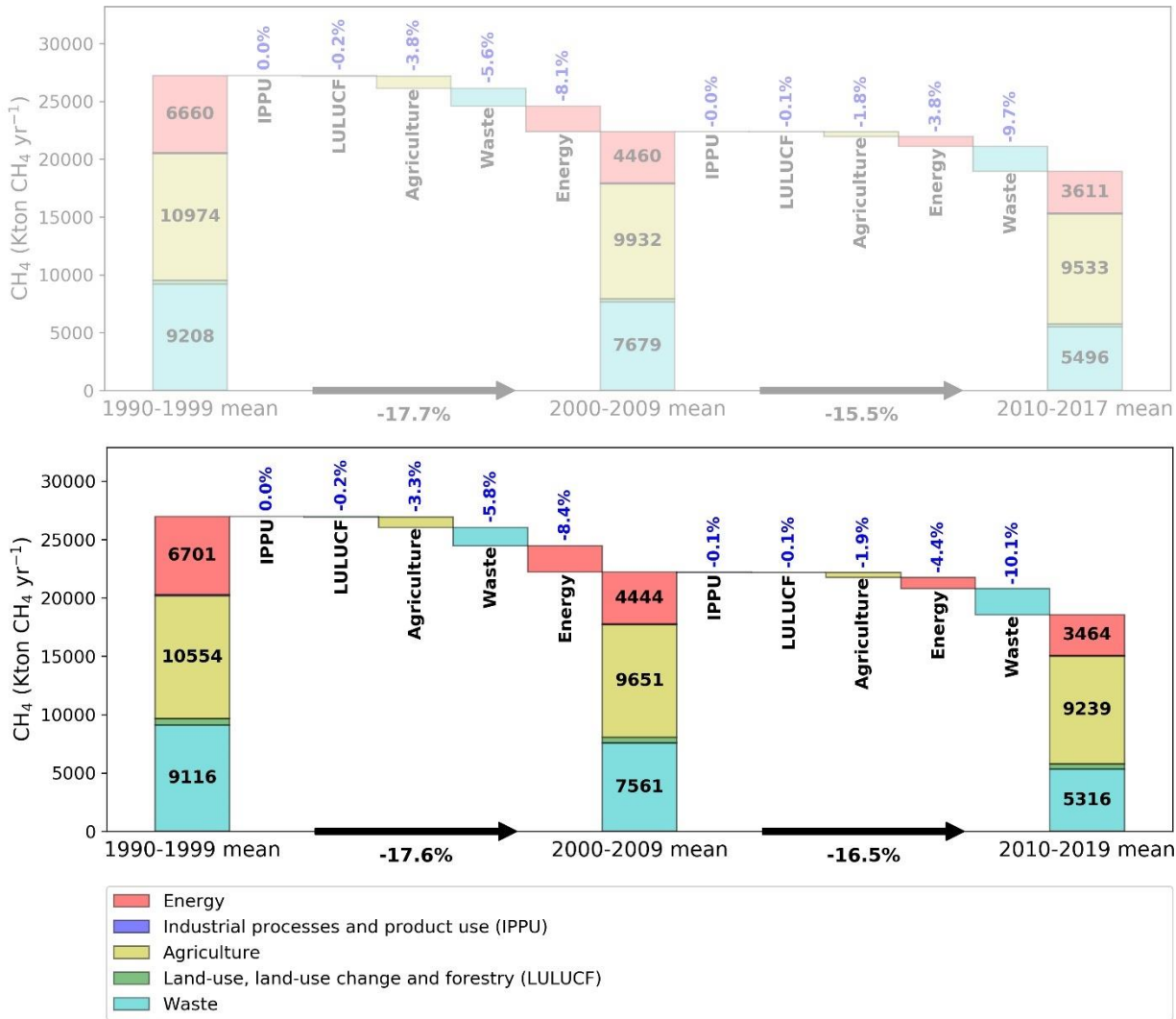


Figure 2: The contribution of changes (%) in CH₄ anthropogenic emissions in the five sectors to the overall change in decadal mean for the EU27+UK, as reported to UNFCCC. The top plot shows the previous NGHGI data from Petrescu et al., 2021a and bottom plot illustrates data from UNFCCC NGHGI (2021). The three stacked columns represent the average CH₄ emissions from each sector during three periods (1990-1999, 2000-2009 and 2010-2019) and percentages represent the contribution of each sector to the total reduction percentages (black arrows) between periods.

3.1.3. NGHGI estimates compared with bottom-up inventories

The data in Figure 3 presents the total anthropogenic CH₄ emissions from four BU inventories and UNFCCC NGHGI (2019/2021) submissions excluding emissions from LULUCF, which was identified to a non-significant contributor (Figure 2). According to NGHGI, in 2019 anthropogenic CH₄ emissions from the four sectors (Table 1, excluding LULUCF) amounted to 4817.1 Tg CH₄ yr⁻¹, representing 10.57 % of the total EU27+UK GHG emissions in CO₂eq.. Figure 3a shows EDGARv6.0 and GAINS trends being consistent with the ones of NGHGI (excluding LULUCF), although while GAINS and NGHGI agree in terms of emissions levels. EDGARv6.0 estimates are consistently higher estimates (8–19 %) than NGHGI. In contrast to the previous version, EDGAR v4.3.2, which was found by Petrescu et al. 2020 to be consistent with NGHGI (2018) data, EDGAR v6.0 reports higher estimates than EDGARv5.0 (~8% higher) and falls outside the 9.6 % UNFCCC uncertainty range. Over the 1990-2019 period, the trends in emissions agree well between the two BU data sets and NGHGI, showing linear trend reductions of 40 % for EDGAR v6.0 and 36 % for GAINS and NGHGIs. The average yearly reduction trend was 2 % yr⁻¹ for all three data sources .

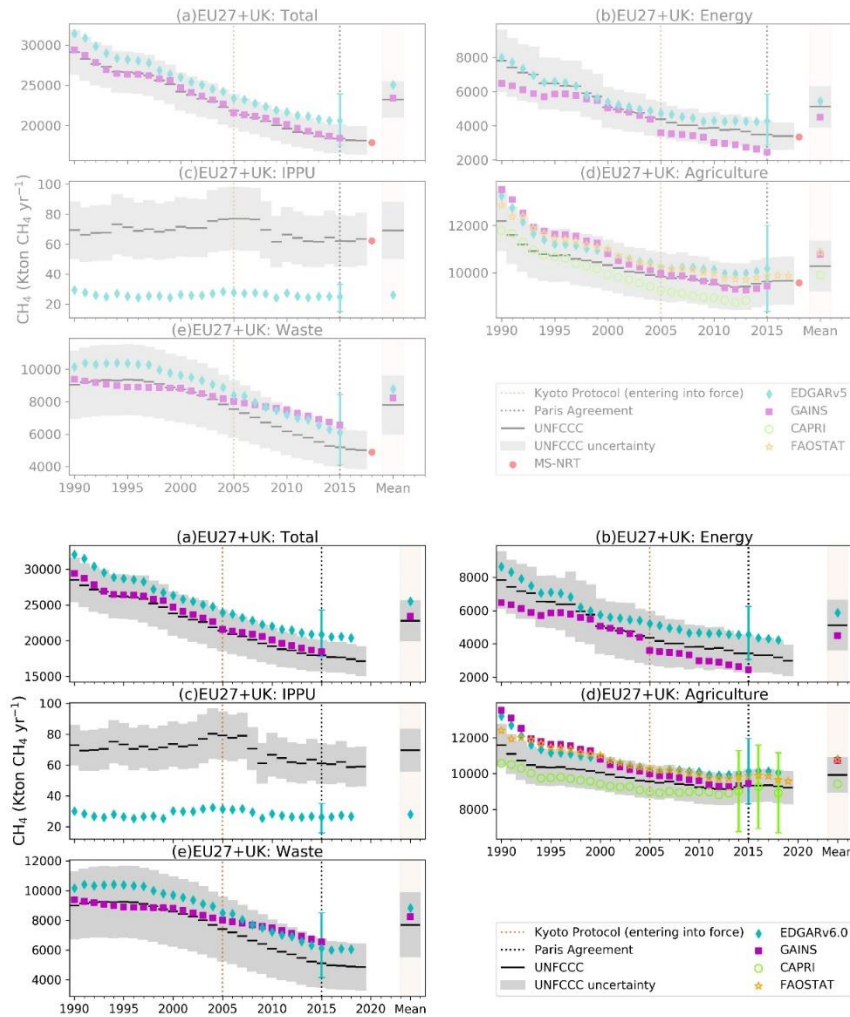


Figure 3: Total annual anthropogenic CH₄ emissions (excluding LULUCF) for the EU27+UK over time. The top plot presents previous data synthesized in Petrescu et al., 2021a while bottom plot data synthesized by the current study: a) EU27+UK and total sectoral emissions from: b) Energy, c) Industry and Products in Use (IPPU), d) Agriculture and e) Waste from UNFCCC NGHGI (2021) submissions compared to global bottom-up inventory models for agriculture (CAPRI, FAOSTAT) and all sectors excl. LULUCF (EDGAR v6.0, GAINS). CAPRI reports one estimate for Belgium and Luxembourg. The relative error on the UNFCCC value represents the UNFCCC NGHGI (2021) Member States reported uncertainties computed with the error propagation method (95% confidence interval) that were gap-filled and provided for every year. The uncertainty for EDGARv6.0 is the same as that of v5.0 and calculated for 2015 as min/max values for the total and each sector (Solazzo et al., 2021) and represents the 95 % confidence interval of a lognormal distribution. The mean column represents the common overlapping periods between datasets: 1990-2015 for Total EU27+UK, Energy, Agriculture and Waste, 1990-2018 for IPPU and 1990-2014 for CAPRI. Last years of the time series of the respective datasets are 2018 (EDGAR v6.0, CAPRI) 2019 (FAOSTAT, UNFCCC) and 2015 (GAINS). After 2014 CAPRI delivered estimates for two additional years, 2016 and 2018 as well as uncertainties for 2014 and 2018 (25.1 %) and 2016 (25.2 %).

Sectoral time series of anthropogenic CH₄ emissions (excluding LULUCF) and their means are shown in Figures 3b,c,d and e. For the energy sector (Figure 3b), both EDGAR v6.0 and GAINS agree in trends with the NGHGI thanks to updated methodology that derives emission factors and accounts for country-specific information about associated petroleum gas generation and recovery, venting and flaring (Höglund-Isaksson, 2017). After 2005, GAINS reports consistently lower emissions than UNFCCC due to a phase-down of hard coal production in Czech Republic, Germany, Poland and the UK, a decline in oil production in particular in the UK, and declining emission factors reflecting reduced leakage from gas distribution networks as old town gas networks are replaced. A difference in tiers is also one reason for the differences (Petrescu et al., 2020).

The consistently higher estimates (+6 % compared to the UNFCCC mean) of EDGAR ~~v5~~v6.0 might be due to the use of default emission factors for oil and gas production based on data from the US (Janssens-Maenhout et al. (2019)). There are several other reasons that could be the cause for the differences, including the use of Tier 1 emission factors for coal mines, assumptions for material in the pipelines (in the case of gas transport) and the activity data). Also EDGAR ~~v5~~v6.0, similar to the previous estimates from EDGARv5.0 uses the gas pipeline length as a proxy for the activity data however this may not be appropriate for the case of the official data, which could consider the total amount of gas being transported or both methods according to the countries. Using pipeline length may overestimate the emissions because the pipeline is not always at 100% capacity thus a larger amount of methane is assumed to be leaked (Rutherford et al., 2021). For coal mining, emissions are a function of the different types of processes being modelled.

The IPPU sector (Figure 3c), which has only a small share of the total emissions, is not included in GAINS, while EDGAR ~~v6~~v5.0 estimates are less than half of the emissions reported by NGHGI ~~2019-2021~~ in this sector. The discrepancy for this sector has negligible impact on discrepancy for the total CH₄ emission. However, we identified that the low bias of EDGAR ~~v5~~v6.0 could be explained by fewer activities included in EDGAR ~~v5~~v6.0 (e.g. missing solvent, electronics and other manufacturing goods) accounting for 5.5 % of the total IPPU emissions in 2015 reported

to UNFCCC. The reason for the remaining difference could be explained by the allocation of emissions from auto-producers⁹ in EDGAR v5.0 to the Energy sector (following the 1996 IPCC guidelines), while in NGHGI they are reported under the IPPU sector (following the 2006 IPCC guidelines).

As CAPRI and FAOSTAT report only emissions from agriculture, they are included only in Fig. 3d. The data (EDGAR v6.0, GAINS, CAPRI and FAOSTAT) shows good agreement, with CAPRI at the lower range of emissions (Petrescu et al., 2020) and on average 3% lower than that of NGHGI, and EDGAR v5.0 at the upper range. The reason for EDGAR v5.0 having the highest estimate (contrary to Petrescu et al., 2020 where NGHGI were the highest and EDGAR v4.3.2 was the second highest) is likely due to the activity data updates in EDGAR v6.0 based on FAOSTAT values, compared to EDGAR v4.3.2. When looking at the time series mean, EDGAR v6.0, GAINS and FAOSTAT show 5 % higher emissions than that of NGHGI. The three BU estimates and NGHGI estimates show similar mean values likely due to the use of similar activity data and emission factors (EFs) (i.e. Figure 4 in Petrescu et al., 2020). The updates submitted by CAPRI, for the years 2014, 2016 and 2018 match the NGHGI emission estimates and have uncertainties of 21 %. Compared to the previous version of CAPRI used in Petrescu et al., 2021a, the new runs report lower CH₄ emissions. Compared to previous results, in the last version some changes have been implemented in the last version (e.g. introduction of slope and altitude limits based on LUCAS¹⁰, improved distribution of grazing livestock etc.). The main activity triggering the differences was the emissions from enteric fermentation. Statistical information on most agricultural data required for the estimation of CH₄ and N₂O emissions are not available at high spatial (regional) and temporal (annual since 1990) resolution. Therefore, the CAPRI model features a module that provides generic data at regional level (CAPREG) and additionally a module that also estimates feed distribution and GHG emissions at the required resolution for VERIFY (CAPINV). As indicated in an internal VERIFY report (Leip et al., 2019), the results of the CAPINV module were scrutinized and shortcomings were identified. These concern mainly the distribution of feed, which is one of the most important parameter for CH₄ emissions from enteric fermentation, and manure excretion and subsequent GHG emissions. Other updates included addition of some regional input data (sources: FAOSTAT and EUROSTAT).

For the waste sector (Figure 3e) EDGAR v5.0 shows consistently higher estimates compared to the NGHGI data, while GAINS has higher emissions than the NGHGI after 2000 (mean 1990-2015 value 6% higher than NGHGI emissions). The two inventories, EDGAR v5.0 in its 2020 update for landfills, and GAINS used an approach based on the decomposition of waste into different biodegradable streams, with the aim of applying the methodology described in the 2019 Refinement of the 2006 IPCC guidelines and the IPCC waste model (IPCC, 2019) using the First-Order-Decay (FOD) method. The main differences between the two datasets come from i) sources for total waste generated per person, ii) assumption for the fraction composted and iii) the oxidation. The two inventories may have used different strategies to complete the waste database when inconsistencies were observed in the EUROSTAT database or in the waste emissions trends in NGHGI.

⁹ auto-producers of electricity and heat: cogeneration by industries and companies for housing management (central heating and other services) (Olivier et al., 2017 PBL report)

¹⁰ <https://ec.europa.eu/eurostat/web/lucas>

3.1.4. NGHGI estimates compared to atmospheric inversions

European estimates from regional inversions

Figure 4 compares TD regional estimates, NGHGI anthropogenic data for CH₄ emissions and natural BU emissions. Figure 4a presents TD estimates of total emissions (anthropogenic and natural) from Petrescu et al., 2021a while Fig. 4b shows the current study with updated total TD estimates. Figs. 4c and 4d show estimates of anthropogenic emissions (Petrescu et al., 2021a and current study) calculated by subtracting the total natural emissions from the total TD emissions.

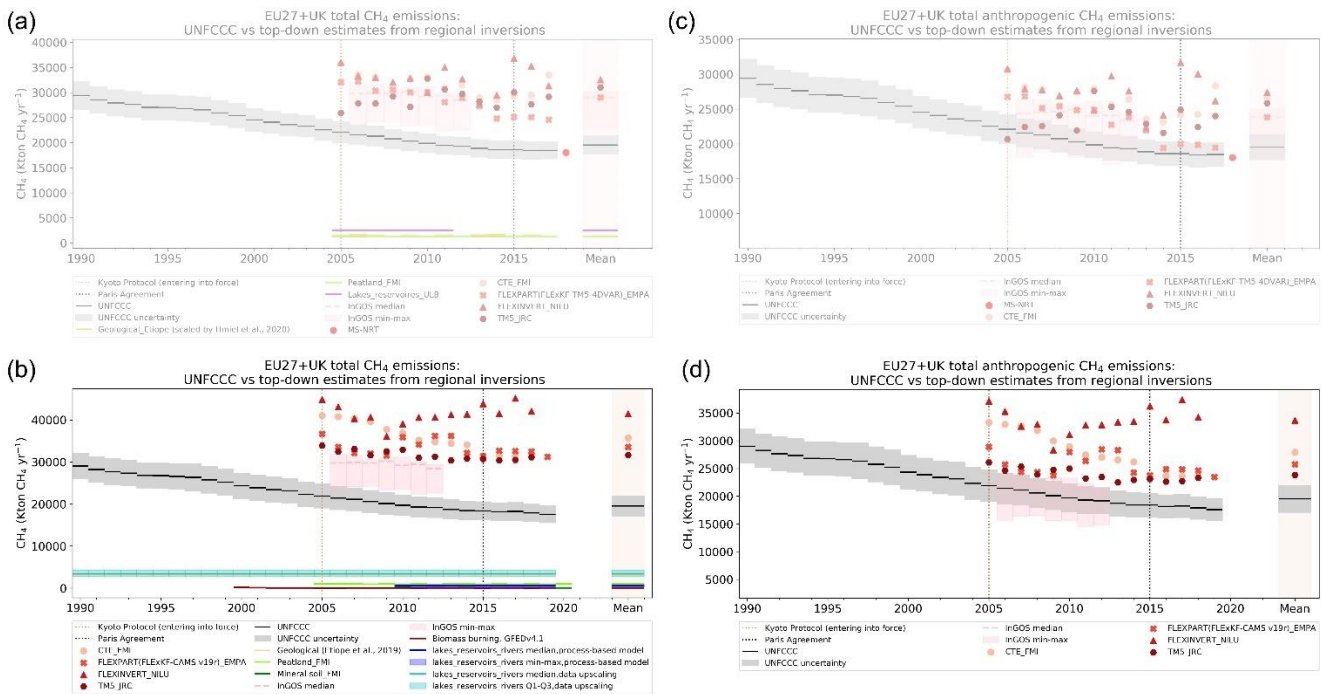


Figure 4: a) and b) Comparison of total CH₄ emissions for EU27+UK from four top-down regional inversions with UNFCCC NGHGI (grey) data and two estimates for inland waters (lakes_rivers_reservoirs—process-based models, blue and upscaled emissions, cyan), peatlands and mineral soils (from JSBACH-HIMMELI, light green and dark green), geological emissions (orange) and biomass burning, (from GFEDv4.1, brown) as following: a) shows previous data from Petrescu et al., 2021a and b) current study; c) and d) comparison of anthropogenic CH₄ emissions from four top-down regional inversions with UNFCCC NGHGI (grey) data as following: c) previous data from Petrescu et al., 2021a and d) current study. Anthropogenic emissions from these inversions are obtained by removing natural emissions and biomass burning from total TD CH₄ emissions shown in Figure 4a,b. UNFCCC NGHGI (2018-2021) reported uncertainties computed with the error propagation method (95% confidence interval) were calculated for each year of the time series and represents the gap-filled harmonized Member States reported uncertainty for all sectors (including LULUCF). The time series mean was computed for the common period 2005-2018 between datasets (excluding InGOS).

The TD estimates of European CH₄ emissions in Figure 4b use four European regional models for 2005-2018 period and an ensemble of five different inverse models (InGOS, Bergamaschi et al., 2015) for 2006-2012.

For the 2005-2018 period (excluding InGOS), the four regional inversions give a total CH₄ emissions mean of 25.836 (32-42) Tg CH₄ yr⁻¹ compared to anthropogenic total of 20 Tg CH₄ yr⁻¹ in NGHGI (Fig. 4b). The large positive difference between TD and NGHGI suggests a potentially significant contribution from BU natural sources, represented separate in Figure B1c, Appendix B (peatlands, geological sources, inland waters and biomass burning), which for the same period are estimated at 8 Tg CH₄ yr⁻¹. However, it needs to be emphasized that natural wetland emission estimates have large uncertainties and show large variability in the spatial (seasonal) distribution of CH₄ emissions but for Europe their inter-annual variability is not very strong (mean of 13-14 years from JSBACH-HIMMELI peatland emissions is 1.0 Tg CH₄ yr⁻¹). Overall, they do represent an important source, and could dominate the budget assessments in some regions such as Northern Europe (Figure 1). That TD and NGHGI diverged in terms of both emissions levels and trends is certainly significant and potentially has implications for bottom-up and NGHGI estimates of CH₄ emissions, if the discrepancies cannot be explained by natural fluxes alone.

The differences between inversion results in current study and Petrescu et al., 2021a can be summarized as following: for the version used in this study, FLExKF-CAMSv19r EMPA, the background mole fraction was taken from a global CAMS v19r assimilation run with assimilation of surface observations of CH₄ only (no satellite data) where the domain was cut out following the two-step approach of Rödenbeck et al. (2009). Background concentrations from CAMS v19r are on average about 5 ppb lower than those of the TM5-4DVAR system used previously, which results in somewhat higher emission estimates over Europe compared to Petrescu et al. (2021a). The major differences to previous CTE-FMI run are the prior fluxes, except for biomass-burning which remained GFED. The new VERIFY_S5 (core) run uses fluxes as described in Thompson et al., 2021. The new VERIFY_S5 (core) run uses fluxes as described in Thomson et al., 2021. Lake & geological emissions were not included in Petrescu et al., 2021a synthesis, but are included in the current CTE-FMI simulation, which probably also contributes to higher total emissions. On top of this, the assimilated data (i.e. observation network) contributes to the differences (enlarged observation network, more sites (five core sites for CH₄ located in Spain, France and UK were added). The FLEXINVERT version used in this study updated the atmospheric observation network (more sites were added) as well as the prior emissions. The background mole fraction was also coupled with that from CAMS v19r assimilation run, which, similar to FLExKF model, might imply higher emissions. Regarding decreasing trends seen for the current inversion CH₄ results, for FLExKF model the trend in CH₄ emission was slightly negative over 2005-2019, at -0.48% per year, which is lower than the decrease in the prior of -0.8% per year. For the other models, based on Thompson et al., 2021, the differences in trends might be due to regional vs. global inversion differences.

The geological emissions were recalculated based on the global grid model of Etiope et al (2019), using more precise “activity” data for EU27+UK (details in Appendix A2): the emission resulted to be 3.3 Tg CH₄ yr⁻¹, i.e. 42 % of the total natural CH₄ emissions in EU27+UK. Geological emissions are an important component of the EU27+UK emissions budget, but their temporal variability is unknown (Etiope and Schwietzke, 2019) and so their impact on climate warming cannot be predicted.

The other natural sources of CH₄ contribute as following: natural emissions from inland waters (based on Lauerwald et al., in [preprev. GBC](#), see Appendix A2) contribute 3.4 Tg CH₄ yr⁻¹, or 43 % of the total natural CH₄ emissions; peatlands and mineral soils (Raivonen et al. 2017 and Susiluoto et al. 2018) account for 1.0 Tg CH₄ yr⁻¹, i.e. 13.4 % of the total natural CH₄ emissions while biomass burning contributes only 0.6 % to the total CH₄ natural emissions.

Similar to peatlands, inland water emissions also remain highly uncertain. The compilation of emission estimates lead to a total flux that is 3.3 Tg CH₄ yr⁻¹ (min 2.7 Tg CH₄ yr⁻¹ and max 4.3 Tg CH₄ yr⁻¹) and about five times larger than the process-based model estimates for lakes+reservoirs and the spatially resolved flux for rivers (0.6 Tg CH₄ yr⁻¹ with min 0.2 and max 0.8 Tg CH₄ yr⁻¹) and about 25 % larger than the previous budget in Petrescu et al., 2021a (2.5 Tg CH₄ yr⁻¹), which ignored the contribution of rivers and relied on one observation-based estimate (extrapolation from late-summer data reported in Rinta et al. 2017) and four semi-empirical model assessments (Petrescu et al., 2021a). Interestingly, the new process-based estimate for natural lake+reservoirs CH₄ emissions matches well the data-driven assessment by Rinta et al. (2017) for the late summer season, with a relative difference smaller than 5 %. The first approach synthesizes 15 average annual CH₄ emissions fluxes for Europe that were rescaled to a consistent set of inland water surface area (Lauerwald et al., in [preprev. GBC](#)) and corrected for the effect of seasonal ice cover.

Model results however also reveal a strong seasonal variability in CH₄ emissions, with much lower fluxes during winter. This seasonality is driven by physical factors (changing ice cover and bottom-water temperature) and biogeochemical factors (autotrophic primary production) that are well established drivers of the temporal variability in lake CH₄ emissions (Del Sontro et al., 2018; Jansen et al., 2022). This finding provides a likely explanation as to why the spatio-temporally resolved model results lead to significantly lower estimates than observation-based methods that do not capture well the temporal variability in lake CH₄ emissions.

~~Model results however also reveal a strong seasonal variability in CH₄ emissions, with much lower fluxes during winter. This finding partly explains why the spatio-temporally resolved model for rivers results lead to significantly lower estimates than observation-based methods that do not capture well the temporal variability in lake CH₄ emissions.~~

According to the IPCC 2006 guidelines (IPCC, 2006) CH₄ emissions from wetlands are reported by the Member States to the NGHGI under the LULUCF sector and considered anthropogenic, if the wetlands in question are considered managed land. They are included in the total LULUCF values (Figure 1, 2, 4 and 6) and in 2019 reported CH₄ emissions from wetlands accounted for 0.1 Tg CH₄ yr⁻¹.

To quantify the anthropogenic CH₄ component in the European TD estimates, the BU peatland emissions from the regional JSBACH-HIMMELI model and those from geological, inland water sources and biomass burning were subtracted from the total TD emissions (Fig. 4d). It remains however uncertain to perform these corrections due to the prior inventory data allocation of emissions to different sectors (e.g. anthropogenic or natural) used in inversions, which can induce uncertainty of up to 100 % if for example an inventory allocates all emissions to natural emissions and the correction is made by subtracting the natural emissions. All regional inversion anthropogenic estimates are higher compared to the UNFCCC NGHGI (20192021), mean of 28 Tg CH₄ yr⁻¹ from inversions compared to 20 Tg

CH₄ yr⁻¹ from the NGHGs. Regarding trends, TD are stable except for CTE showing a linear decreasing trend up to 2015 followed by an increase over the next three years, while NGHGs and BU trends are declining. From this attempt we find that not many of the inversions showed the clear decline reported by the NGHGs. As NGHGI emissions are dominated by anthropogenic fluxes and decline by almost 30% compared to 1990, a similar decline was expected in the corrected anthropogenic inversions. Further investigation into how well the NGHGs reflect reality or how well the TD estimates capture the trends is clearly needed. Currently, in the UK NIR (<https://unfccc.int/documents/273439>) the national inversion system produced similar recent UK CH₄ emission levels, but did not validate the large declining trend since 1990 that is estimated by the UK inventory.

Spatial distribution of CH₄ emissions from regional inversions

A novelty in this study is represented by the new top-down estimates of CH₄ fluxes were also calculated in this reporting period using the Community Inversion Framework (CIF) (Berchet et al. 2021). For CH₄ (Figure 5), inversions using three atmospheric transport models (or model variants) were performed with the CIF, there were: i) the regional non-hydrostatic Eulerian model, CHIMERE (Fortems-Cheiney et al., 2021) used by LSCE, ii) the Lagrangian particle dispersion model, FLEXPART used by EMPA (from hereon, FLEXPART-EMPA), and iii) FLEXPART used by NILU (from hereon, FLEXPART-NILU).

The spatial distribution of CH₄ fluxes are similar for the three inversions with higher emissions in the Netherlands and Belgium, western France and southern UK. However, FLEXPART-NILU inversions show some spurious areas of very low fluxes in Italy, Switzerland and southern France, which are presumably owing to the positive bias in the prior modelling mixing ratios at mountain sites, which will be corrected in future simulations. The patterns of differences, however, are quite different between the CHIMERE and the two FLEXPART inversions. All inversions find positive increments (posterior high than prior) over northern Netherlands, but FLEXPART-EMPA finds negative increments over southern Netherlands and both FLEXPART inversions find negative increments over northern Italy, which is not the case in CHIMERE (Figure 5, top). The total mean emissions for EU27+UK over 2006-2017 (Figure 5) were 26, 22 and 24 Tg CH₄ yr⁻¹, for CHIMERE, FLEXPART-NILU and FLEXPART-EMPA, respectively. FLEXPART-EMPA is the same model as used in the comparison shown in Figure 4 (FLEXPART(FLExKF-CAMsv19r)) but in those inversions the total mean emissions for EU27+UK were higher at

33 Tg yr⁻¹. This difference is likely owing to the different dataset used for determining the background mixing ratios and farther analysis is ongoing.

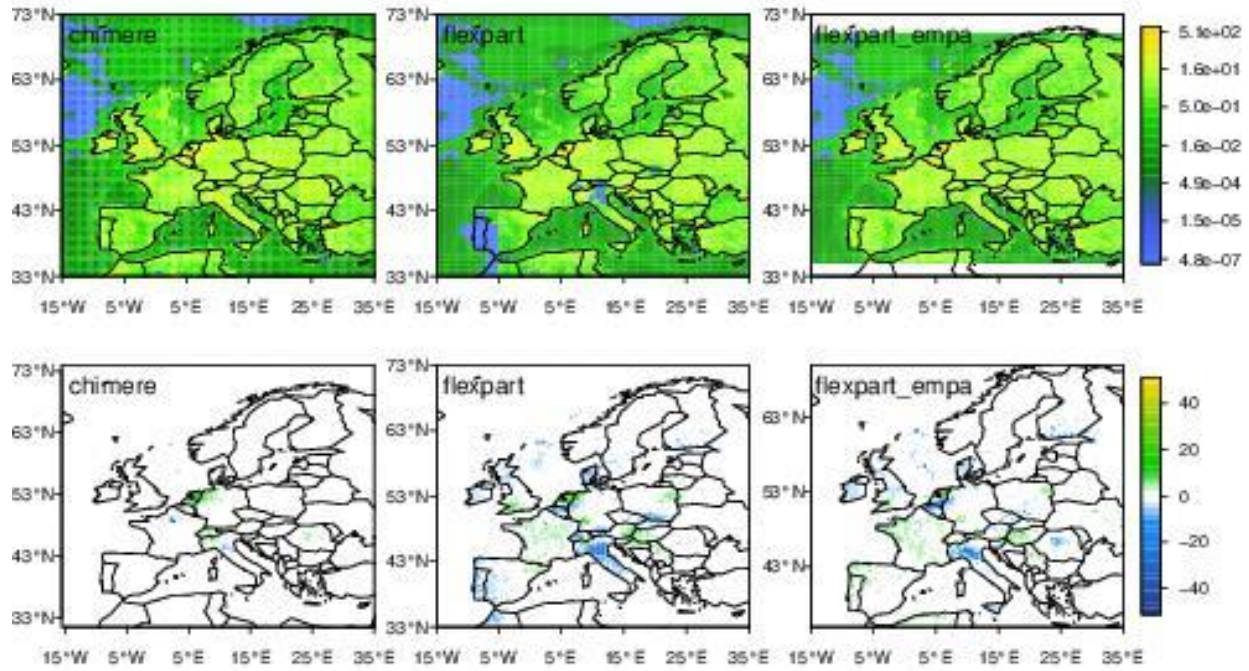


Figure 5: Posterior CH₄ fluxes averaged over 2006-2017 (g CH₄ m⁻² yr⁻¹) from three regional inversions, CHIMERE (LSCE), FELXPART (NILU) and FLEXPART (EMPA) shown with a log base 2 color scale (top) and the flux increments (posterior-prior)(g CH₄ m⁻² yr⁻¹) shown on a linear color scale (bottom).

European estimates from global inversions

Figures 6 compares TD global estimates, with NGHGI data and provides information about the wetland emissions from global wetland inversions (Saunois et al., 2020). Figure 6a presents TD estimates of total emissions (anthropogenic and natural) from Petrescu et al., 2021a while Fig. 6b shows the current study with updated total TD estimates. Figs. 6c and 6d show estimates of anthropogenic emissions (Petrescu et al., 2021a and current study) calculated by subtracting the total natural emissions from the global total TD emissions.

The global inversion models were split according to the type of observations used, 11 of them using satellites (GOSAT) and 11 using surface stations (SURF). Each of these 22 global inversions provided as well wetlands emissions used by the Global Methane Budget (Saunois et al., 2020) and are post-processed with prior ratios estimates for wetlands CH₄ emissions (Appendix B2, Table B2.4).

For the common period between datasets (2010-2016), the two ensembles of regional and global models give a total CH₄ emission mean (Figure 6a) of 23 Tg CH₄ yr⁻¹ (GOSAT) and 24 Tg CH₄ yr⁻¹ (SURF) for the EU27+UK compared to 19 Tg CH₄ yr⁻¹ ± 2.3 Tg CH₄ yr⁻¹ of NGHGI (Figure 6a). The mean of the natural wetland emissions from the global inversions is 1.3 Tg CH₄ yr⁻¹ and partly explains the positive difference between total emissions from inversions and NGHGI anthropogenic emissions.

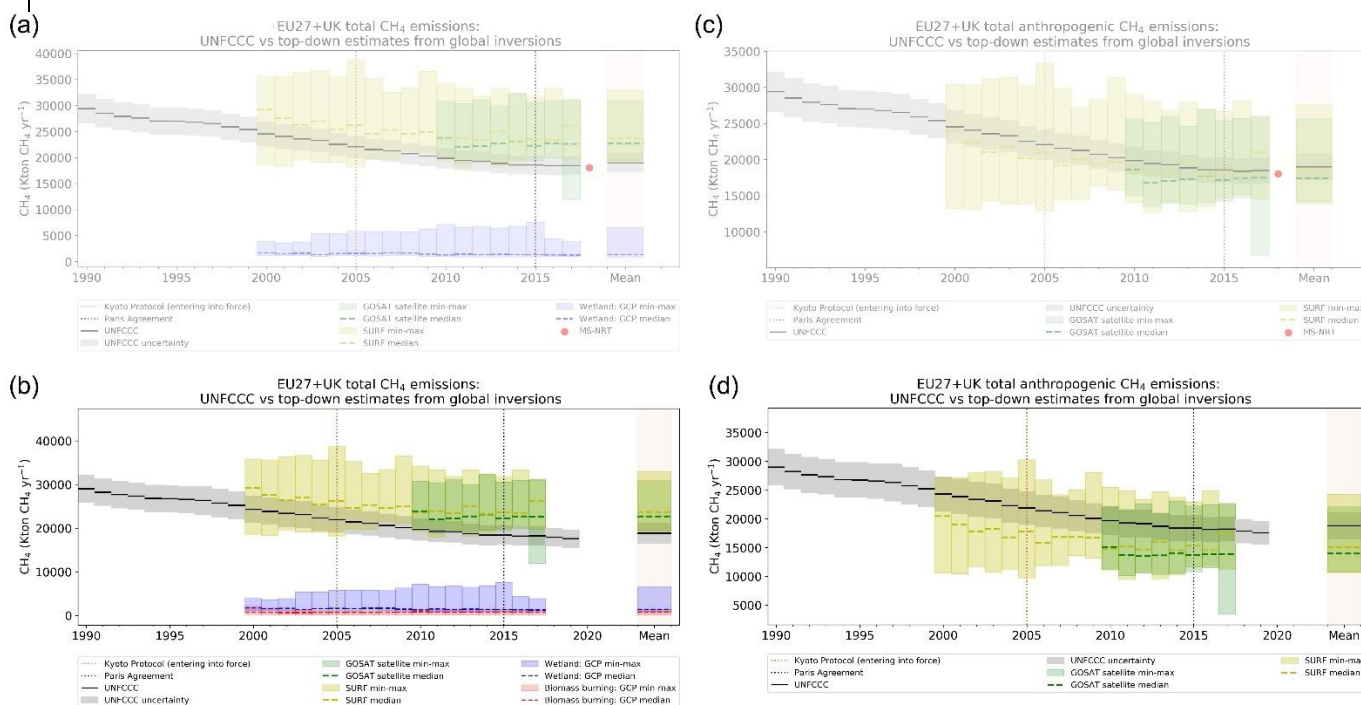


Figure 6: *a) and b)* Total CH_4 emissions from TD global ensembles based on surface stations data (SURF) (yellow) and satellite concentration observations (GOSAT) (green) from 22 global models compared with UNFCCC NGHGI (grey) data (including LULUCF) *as following: a) represents previous data from Petrescu et al., 2021a and b) the current study; c) and d)* Anthropogenic CH_4 emissions from top-down global inversions based on surface stations (SURF) (yellow) and on satellite concentration observations (GOSAT) (green) from different estimates *as following: c) previous data from Petrescu et al., 2021a and d) the current study*. Anthropogenic emissions from these inversions were obtained by removing the sum of the *natural emissions (global wetland GCP emissions (blue), the inland waters and geological fluxes as shown in figure 4a) from the total estimates. The biomass burning emissions included in each inversion results was removed as well.* UNFCCC NGHGI (2018/2021) Member States reported uncertainty computed with the error propagation method (95% confidence interval) *was gap-filled and provided for every year for all sectors (including LULUCF).* The time series mean was computed for the common period 2010-2016. Two out of 11 SURF products (GELCA-SURF_NIES, TOMCAT-SURF_UOL) were not available for 2016.

To quantify the European TD anthropogenic CH_4 component, the *GCP inversions wetlands emissions* and those from *geological, inland water sources and biomass burning emissions (reported by the global inversions) were subtracted from the total CH_4 emissions (Fig. 6d).*

For the 2010-2016 common period, the two ensembles of global models give an anthropogenic CH_4 emission ~~mean median~~ (Figure 6b) of *17.413 Tg $\text{CH}_4 \text{ yr}^{-1}$ with min and max values of 10 and 21 Tg $\text{CH}_4 \text{ yr}^{-1}$ (GOSAT) and 23.714 Tg $\text{CH}_4 \text{ yr}^{-1}$ with min and max values of 9 and 22 Tg $\text{CH}_4 \text{ yr}^{-1}$ (SURF) compared to $19 \pm 2.3 \text{ Tg } \text{CH}_4 \text{ yr}^{-1}$ for NGHGI. The TD ensemble that produced the closest anthropogenic estimate (Figure 6d) to the UNFCCC NGHGI (2021) is SURF, with the median of SURF inversions falling just below the uncertainty range of the NGHGI.*

Between 2010-2016, total TD CH₄ emissions (Figure 6b) from the SURF and GOSAT ensemble decreased by 0.5% and 4.6%, respectively. For anthropogenic CH₄ emissions (Figure 6d), the SURF and GOSAT ensembles show a decrease of 1.1 % and 6.3%, respectively, compared to the 7.7 % decrease for the NGHGI.

3.1.5 CH₄ uncertainty reduction maps

Bergamaschi et al (2010) used TM5 4DVAR to analyze the sensitivity of the modelling system to observations, for further interpretation of the derived emissions, in particular in the context of verification of BU inventories. For this purpose, Bergamaschi et al., 2010 calculated uncertainty reduction maps, as a measure of the sensitivity of the observational network used for the reference inversion. This reduction in uncertainty is calculated as the ratio between a posterior and a prior uncertainty with the formula $(1-\Delta_{\text{post}}/\Delta_{\text{prior}})$, where Δ_{post} represents the posterior uncertainties and Δ_{prior} the prior uncertainties of the inversion system. The same methodology was applied to two VERIFY regional inversions systems, CTE-CH₄ and FLE_xKF (Brunner et al., 2022).

The first inversion system, FLE_xKF, calculated the uncertainty reduction maps for CH₄ for the year 2018 with two different sets of observation stations (Figure 7). Maps of uncertainty reduction can be really informative and the results below (Figure 7) present the uncertainty reductions for two different sets of stations, which show the value of only considering ICOS sites (left figure) and when adding also other stations in the U.K. and Switzerland (right figure). However, the larger the prior uncertainties are, the stronger potential for uncertainty reduction is, therefore given that the prior uncertainty varies, the uncertainty reduction is not a direct indication of the information provided by observations.

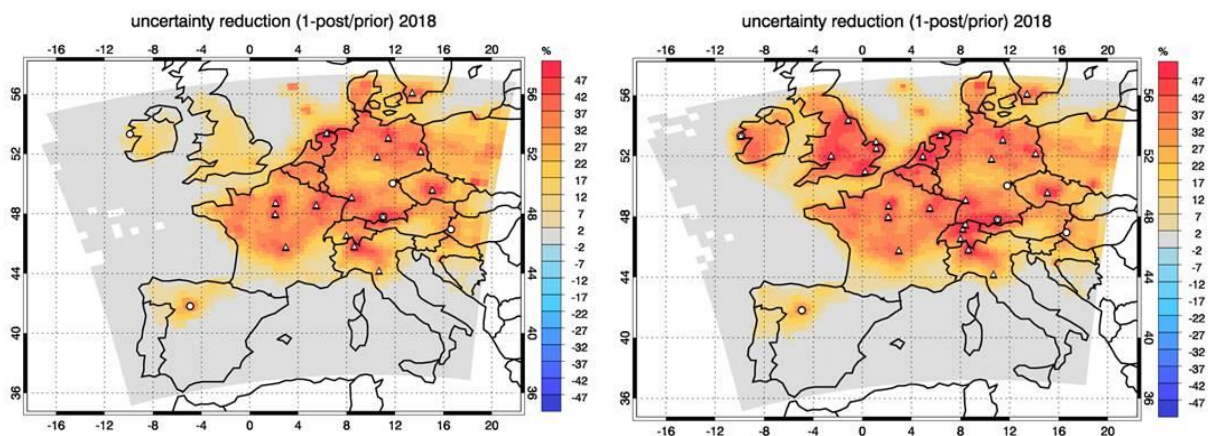


Figure 7: FLE_xKF uncertainty reduction maps computed as $(1-\Delta_{\text{post}}/\Delta_{\text{prior}})$ for the same year, 2018, but with two different sets of observation stations (white dots).

The second inversion system, CTE-CH₄ (Tsuruta et al., 2017) calculated the uncertainty reduction maps from surface inversions (SURF) for 2006 and 2018, as those used in Thompson et al., (2021), referred to here as VERIFY_S5 ("core" inversion) (Figure 8). The system included two sets of inversions with different observation sets assimilated. However, the degrees of freedom in the state of the system was low, and therefore, the uncertainty

estimates may not differ much between the two. The data from CTE-CH₄ includes uncertainties (standard deviations) and fluxes for 2006 and 2018. The differences in the simulations are observation sets and underlying prior covariance structure. "VERIFY S5" uses data from only those sites that have long-term measurements assimilated i.e. there is little differences in the assimilated sites between the years. From the two panels of the Fig.8, higher uncertainty reductions are seen in 2018 compared to 2006 in E Poland, N Italy and Spain.

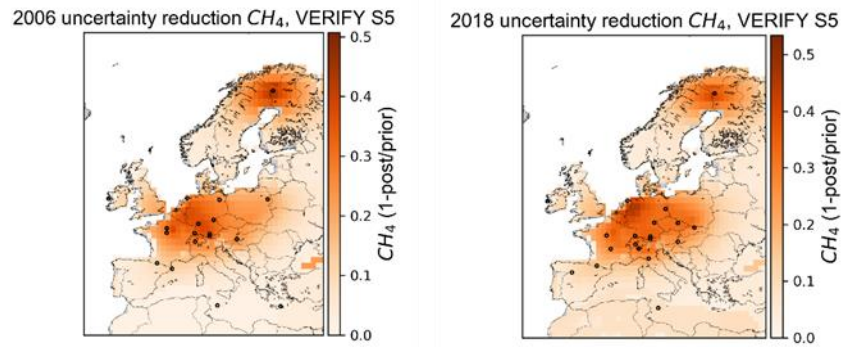


Figure 8: VERIFY S5 (core) inversion run, uncertainty reduction maps computed as $(1 - \Delta_{\text{post}}/\Delta_{\text{prior}})$ for 2006 (left) and 2018 (right) with different sets of observation stations.

The differences between the two years are mostly due to changes in the amount of observational data, although additional observation stations in certain locations may produce only a limited reduction in uncertainty. This can occur if: i) uncertainty assigned to the observations (i.e. how much weight/trust we put on it) are comparatively high, ii) prior emissions and/or their uncertainties around the sites are simply very small, and therefore the inversion does not change fluxes much; and/or iii) the location is not very sensitive to emissions in the surrounding area (e.g. mountain sites) due to the atmospheric transport to the observation site. Generally, sites that contribute to a larger uncertainty reduction should be included in the inversions and located closer to emission sources and/or sink areas.

CTE-CH₄ was also used to estimate fluxes utilizing prior information from GOSAT data, for 2010 and 2017. Figure 9 presents the associated uncertainty reduction maps. Because of the different inversion system set-up (e.g. resolution, spatial correlation) compared to previous results, where prior data was coming from observation networks, it is difficult to conclude on the effects satellites have on posterior emissions from the two years. However, it is interesting to note how satellite data assimilation infers changes on a regional scale. Unlike surface stations, satellite data have more power to constrain northern European emissions than central European emissions.

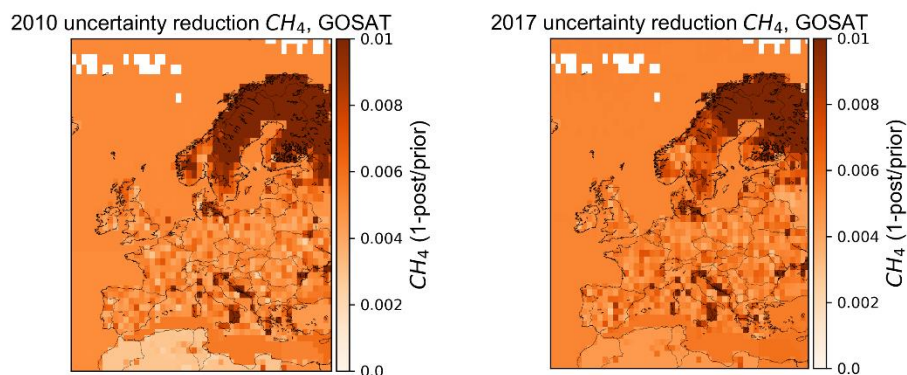


Figure 9: CTE-CH₄ GOSAT inversion run, uncertainty reduction maps computed as $(1-\Delta_{\text{post}}/\Delta_{\text{prior}})$ for 2010 (left) and 2017 (right).

3.2 Comparing N₂O emission estimates from different approaches

3.2.1. Estimates of European and regional total N₂O fluxes

Total N₂O fluxes from EU27+UK and five main regions in Europe are presented in a similar fashion as the CH₄. Figure 10 summarizes the total N₂O fluxes from NGHGI 2021 (excluding LULUCF) for the base year 1990 as well as mean annual emissions for the 2011-2015 and 2015-2019 five-year periods.

The total UNFCCC estimates that include emissions from all sectors are compared with the fluxes from global datasets, BU models and TD inversions. Relative to 1990, N₂O emissions in 2019 decreased by a minimum of 26 % (Eastern Europe) up to a maximum of 43–46 % (Western Europe) and by 39 % for EU27+UK. At European level, the emissions from BU estimates (anthropogenic NGHGI plus the sum of all natural, 991 kton N₂O) and TD total (including natural) regional estimate (1443 kton N₂O) averaged over 2015-2019, roughly agree within the uncertainty reported by UNFCCC ($\pm 59\text{--}80\%$). The TD uncertainty is represented as the variability in the model ensembles and denotes the range between the minimum and maximum estimates within each model ensemble. There is significant uncertainty in Northern Europe, where the TD average estimates indicate sources yet the ensemble ranges from a net sink to a net source (Figure 10). The current observation network is sparse, which currently limits the capability of inverse models to quantify N₂O emissions at country or regional scale.

For all other regions, the BU anthropogenic emissions agree in absolute values with the NGHGI given uncertainties, though consistently higher estimates are produced by TD regional and global models. The difference is still too high to be attributed to the sum of the natural emission, which ranges for all five regions in 2019 between a minimum of 13 kton N₂O yr⁻¹ (Northern Europe) to a maximum of 113 kton N₂O yr⁻¹ (Southern Europe), while the EU27+UK total natural emission is estimated at 178 kton N₂O yr⁻¹.

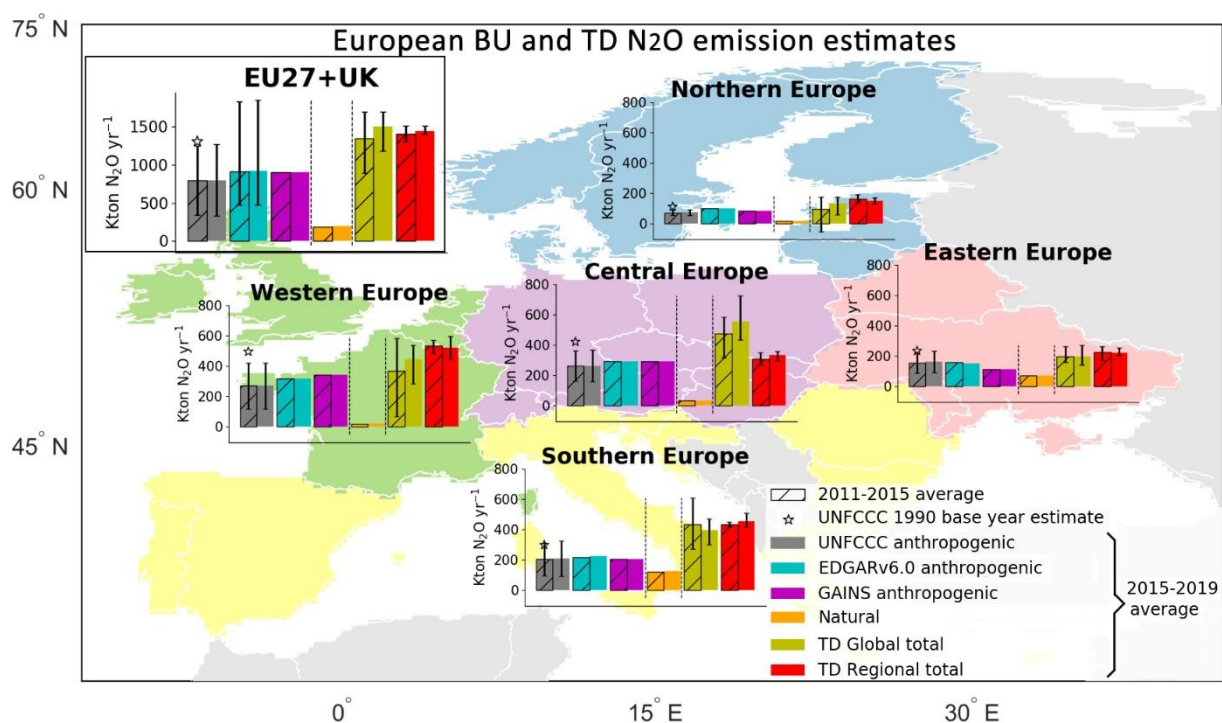


Figure 10: Five-yearly means (2011-2015 hashed bars and 2015-2019 full bars) in total N_2O emission estimates (excluding LULUCF) for EU27+UK and five European regions (Northern, Western Central, Southern and Eastern non-EU). The Eastern European region does not include European Russia, Northern Europe includes Norway and Central Europe includes Switzerland. The data are from the UNFCCC NGHGI (2021) submissions (grey), which are plotted with respective base year 1990 (black star) estimates, two inventories (GAINS and EDGAR v6.0), natural unmanaged emissions (lakes rivers reservoirs emissions from RECCAP2 and natural N_2O from O-CN) and two inversion total estimates (one regional European inversions (FLEXINVERT) and average of three global inverse models from GN₂OB, Tian et al. 2020). The relative error on the UNFCCC value represents the NGHGI (2021) reported uncertainties computed with the error propagation method (95% confidence interval) and gap-filled to provide respective estimates for each year. (see Appendix A); For Easter Europe non-EU the uncertainty value of 42.3 % was calculated from the NIRs. Northern Europe Tier 1 uncertainty for Norway was not available.

3.2.2. NGHGI sectoral emissions and decadal changes

According to the UNFCCC NGHGI (2019/2021) estimates for 2017-2019, the EU27+UK emitted GHGs totaling 3.9-7 Gt CO₂e (including LULUCF, using a GWP 100, IPCC AR4) (Appendix B1, Figure B1b), of which N_2O emissions accounted for ~76 % (254 Mt CO₂e or 854 kton N_2O yr⁻¹) (Figure 11). France, Germany and UK together contributed 4440 % of total N_2O emissions (338 kton N_2O yr⁻¹). For 2019, NGHGI reported anthropogenic emissions from the EU27+UK for the four activity sectors (excluding LULUCF) (Table 1), to be 793 kton N_2O yr⁻¹. Agricultural N_2O emissions accounted for 79 % (± 72.5 %) of total EU27+UK emissions in 2019, followed by emissions from the energy sector with 12 % (± 30 %).

Figure 11 shows anthropogenic N₂O emissions from [UNFCCC NGHGI \(2021\)](#) and their changes from one decade to the next, with the respective contributions from different sectors also illustrated.

Between the 1990s and the 2000s, the net reduction of ~~-17.7~~9 % originates mainly from IPPU ~~(-13.2 %)~~, ~~with a smaller contribution from~~ agriculture ~~(-4.2-4 %)~~. For the period between the 2000s and 2010-201~~9~~7, the net ~~reduction of~~ ~~-15.2-4~~ % was again mainly attributed to the IPPU sector (-14.1 %), despite very small increases from the LULUCF ~~(0.2 %)~~ and waste sectors (~~0.6-2~~ %).

By ~~2017~~2019, emissions from the IPPU sector ~~were only 36 kton N₂O yr⁻¹, a 98-91 % decrease~~ compared to 1990. Although the IPPU sector contributes in ~~2017-2019~~ only 4% to 2019 total N₂O emissions, it is the sector associated with the largest emissions reduction. IPPU sector emissions are mainly linked to the production of nitric acid (e.g. used in fertilizer production) and adipic acid (e.g. used in nylon production). In the late 1990's and early 2000's the five European adipic acid plants were equipped with efficient abatement technology, cutting emissions by 95-99 %, largely through voluntary agreements of the companies. Much of the remaining IPPU emissions, from nitric acid plants, were cut in a similar manner around 2010, a development that has been connected with the introduction of the European Emission Trading System that made it economically attractive for companies to apply emission abatement technologies (catalytic reduction of N₂O in the flue gas) to reduce their emissions.

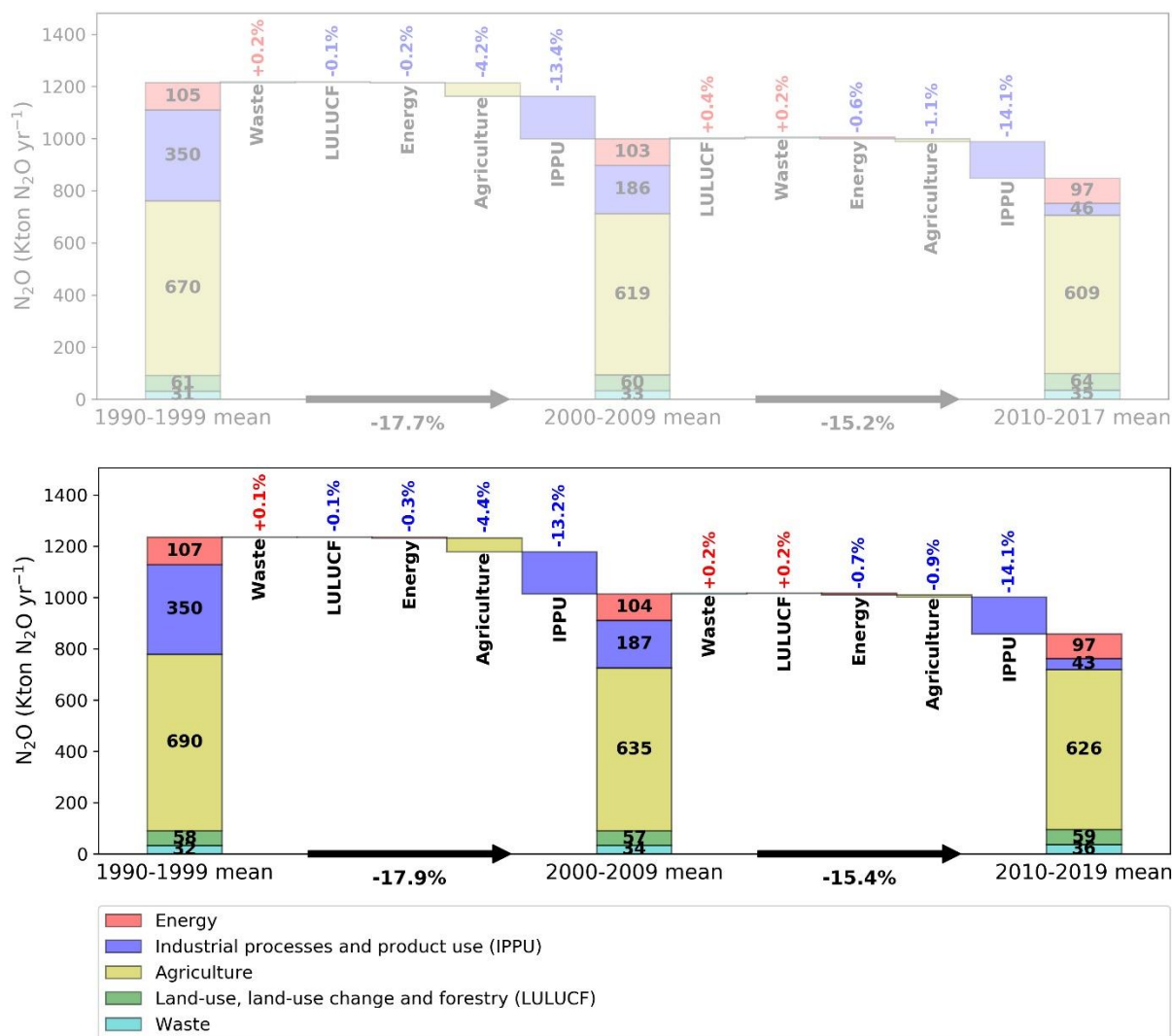


Figure 11: The contribution of changes (%) in N_2O anthropogenic emissions in the five sectors to the overall change in decadal mean for the EU27+UK as reported to UNFCCC. The top plot shows the previous NGHGI data in Petrescu et al., 2021a and bottom plot depicts data from UNFCCC NGHGI (2021). The three stacked columns represent the average N_2O emissions from each sector during three periods (1990-1999, 2000-2009 and 2010-2017/2019) and percentages represent the contribution of each sector to the total reduction percentages between periods.

3.2.3. NGHGI estimates compared with bottom-up inventories

Figure 12 compares the six bottom-up inventories with UNFCCC NGHGI (2019/2021) data, and shows that all of them are around the NGHGI estimates (Figure 12a), noting that GAINS only provides emissions every five years. The BU estimates show good agreement with one another and with the NGHGI estimates until 2005. After 2005 the slightly increasing trend is influenced by the IPPU (Figure 12c) and Waste (Figure 12e) sectors, with estimates of both EDGAR +5.6% and GAINS for total anthropogenic N_2O emissions in the year 2015-2018 being 9% and 13% higher than the respective UNFCCC NGHGI (2019/2021) estimates. Except for agriculture (Figure

12d), where four of the five models/inventories show good match in absolute mean values with the NGHGI and over 1990-~~2015~~2018 and have similar linear trends of -0.18, -0.17, -0.15 and -0.11 % yr⁻¹ in NGHGI, EDGAR v6.0, GAINS and FAOSTAT respectively, for the other sectors the trends differ. The match in agriculture trends reflects that the sources rely on the same basic activity data from FAOSTAT and follow the IPCC EF Tier 1 or 2 approach (Petrescu et al., 2020). However, the high reported uncertainty range from the NGHGIs contradicts the match of the BU estimate absolute values and represents an important research question to be further investigated. In contrast, ECOSSE shows lower estimates because it does not use the FAO fertilizer application rate data base, but instead calculates ideal fertilizer application rates from the nitrogen demand of the crops. ECOSSE uses fertilizer data derived by Mueller et al. (2012) and simulates only for winter wheat. It is very likely that the assumed fertilizer application rates are lower than those used in FAO for the country specific average, which could explain the lower estimates. This means that it may severely under-estimate the applied fertilizer amounts for some areas (e.g. Netherlands, Denmark or North-West Germany), and the results are more indicative of emissions under idealized fertilizer application rates. Additionally, as mentioned above, the model simulates only the direct emissions.

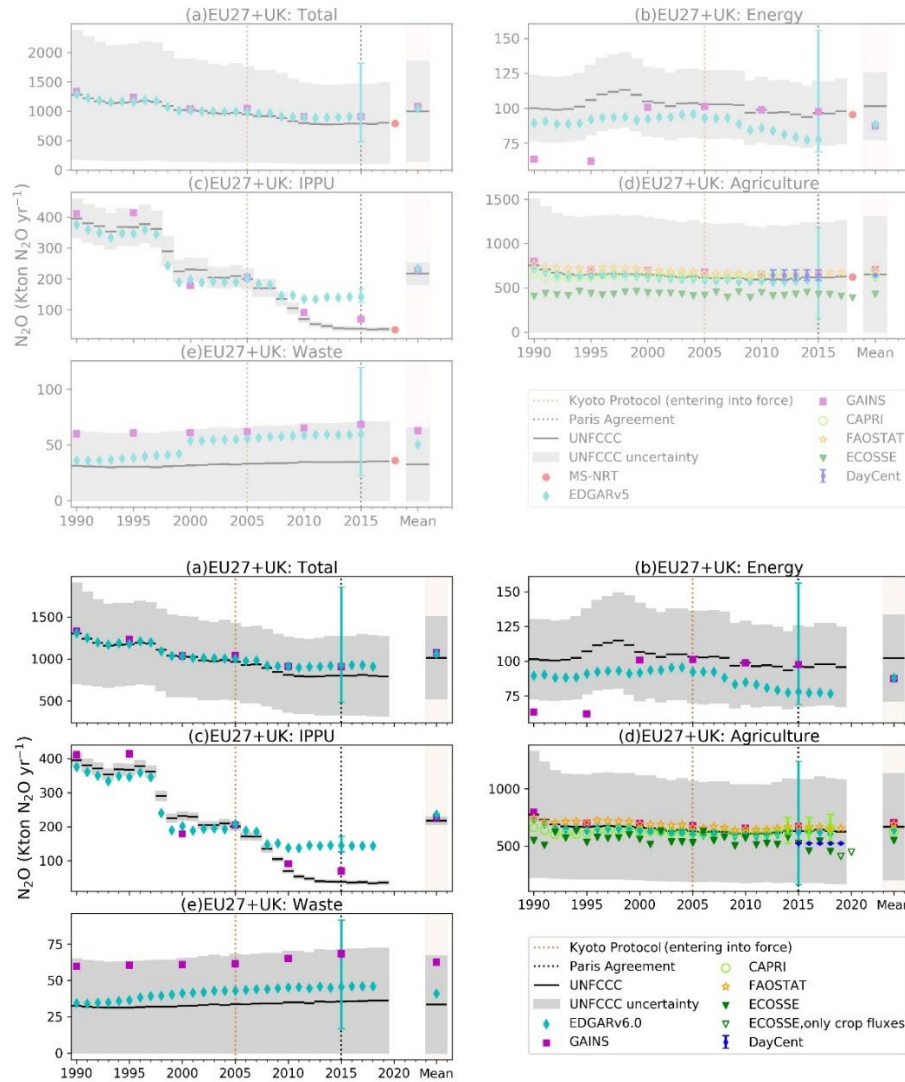


Figure 12: a) Total annual anthropogenic N₂O emissions (excluding LULUCF) for the EU27+UK over time. The top plot presents previous data synthesized in Petrescu et al., 2021a while the bottom plot depicts data synthesized by the current study: a) EU27+UK and total sectoral emissions from: b) Energy, c) IPPU, d) Agriculture, e) Waste from UNFCCC NGHGI (2021) submissions compared to global BU inventories for agriculture (CAPRI, ECOSSE, FAOSTAT, DayCent) and all sectors excl. LULUCF (EDGAR v6.0, GAINS). CAPRI reports one value for Belgium and Luxembourg. The relative error on the UNFCCC value represents the UNFCCC NGHGI (2021) Member States reported uncertainties computed with the error propagation method (95% confidence interval) that were gap-filled and provided for every year. The uncertainty for EDGARv6.0 is the same as that of v5.0 and calculated for 2015 as min/max values for the total and each sector (Solazzo et al., 2021) and represents the 95 % confidence interval of a lognormal distribution. CAPRI reports uncertainties for the last three years as following: 2014 and 2016 (17.6 %) and 2018 (17.8 %). The mean column represents the common overlapping period 1990-2018 between datasets Last years of the time series of the respective datasets are 2019 (UNFCCC and FAOSTAT), 2015-2018 (EDGAR v5.0, v6.0).

CAPRI), 2015 (GAINS every five years), 2015-2019 (DayCent) and 2020 ECOSSE, with last two years reporting only crop emissions.

In the NGHGI (~~2018~~2021) submissions, *for 2019*, the EU27+UK Tier 1 total uncertainty for the waste sector (based on the IPCC chapter 3 error propagation method described in detail by Petrescu et al., 2020) *and the gap-filling method described in Appendix A*, was ~~626-360~~ %. The sectoral activity responsible for this high uncertainty is the wastewater treatment and discharge (~~462913~~%) and this remains one of the most uncertain sources of N₂O having the highest emissions in the waste sector. Emissions are known to vary markedly in space and time even within a single wastewater treatment plant (Gruber et al., 2020), a fact that only recently has been properly accounted for in the inventory guidelines (IPCC, 2019a). However, the total emissions from the waste sector account for only 4.~~4~~₂ % of the total EU27+UK *2019* N₂O emissions (excl. LULUCF).

3.3.4. NGHGI estimates compared to atmospheric inversions

Figure 13 compares inversion estimates of total N₂O emissions, including natural, from regional (FLEXINVERT) and global (three models) N₂O inversions with the UNFCCC NGHGI (~~2019~~2021) estimates. The min-max range of all inversions is within the 2-sigma uncertainty of NGHGI, with the median of global inversions being on average 42 % or 0.4 Tg N₂O yr⁻¹ higher than that of NGHGI. Over the period 2005-~~2017~~2019, the regional FLEXINVERT is *almost double that of* UNFCCC NGHGI (~~2019~~2021). From the three global inversions, *MIROC4-ACTM shows* consistently higher estimates *until 2019, when it registers a drop in the estimated emission level (similar to FLEXINVERT)*. Similar reduction of emissions are seen for 2003 and 2005. In all these years, Europe registered record breaking heatwaves. One plausible explanation for the low N₂O is that high temperature accompanied with lesser soil moisture reduces N₂O emission, as seen in the tropics (Patra et al., 2022).

The other two global inverse models, TOMCAT and CAMS-N₂O register high estimates as well as *very* high variability. Regarding trends, FLEXINVERT shows a similar decreasing trend of *18* % over 2005-201~~9~~7, compared to *16 % for* UNFCCC NGHGI (~~2019~~2021). The global CAMS-N₂O inversion agrees *the* best in its *absolute* mean value (1.0 Tg N₂O yr⁻¹) with the *NGHGI* estimate (0.9 Tg N₂O yr⁻¹) but not in its *trend*. *In this updated synthesis*, natural *pre-industrial soil* emissions of N₂O *from the O-CN model, were included, but these* are not considered in NGHGI reporting, *and therefore cannot explain the gap between inventories and TD estimates*. In addition, the *emission factors* used in NGHGI reporting are *regarded to be* very uncertain (up to 300% for direct agricultural emissions) *which, based on the comparison with TD estimates, could imply that inventories underestimate N₂O emissions*. *The uncertainty reported by the NGHGIs in 2019 was 59% compared to 86% in 20187 (Petrescu et al., 2021a).*

Regarding the natural N₂O emissions, the median natural flux from inland waters, is very low (~~2.7~~12.7 kton N₂O yr⁻¹) and part of the inland water natural estimate is considered anthropogenic in Europe and is due to the leaching of N-fertilizers from agriculture. The anthropogenic share accounts for 66 % of the total inland waters emissions (Petrescu et al., 2021a). *In the current study more natural N₂O estimates have been added. The soil natural background emissions are estimated at 177 kton N₂O yr⁻¹ averaged over 2005-2014 (the common overlapping period of all data*

sources), while the biomass burning emissions account for only 1.6 kton N₂O yr⁻¹ for the same period. In the lower plot, the NGHGI uncertainty was recalculated to 56 % compared to 86% in Petrescu et al., (2021a) (upper plot).

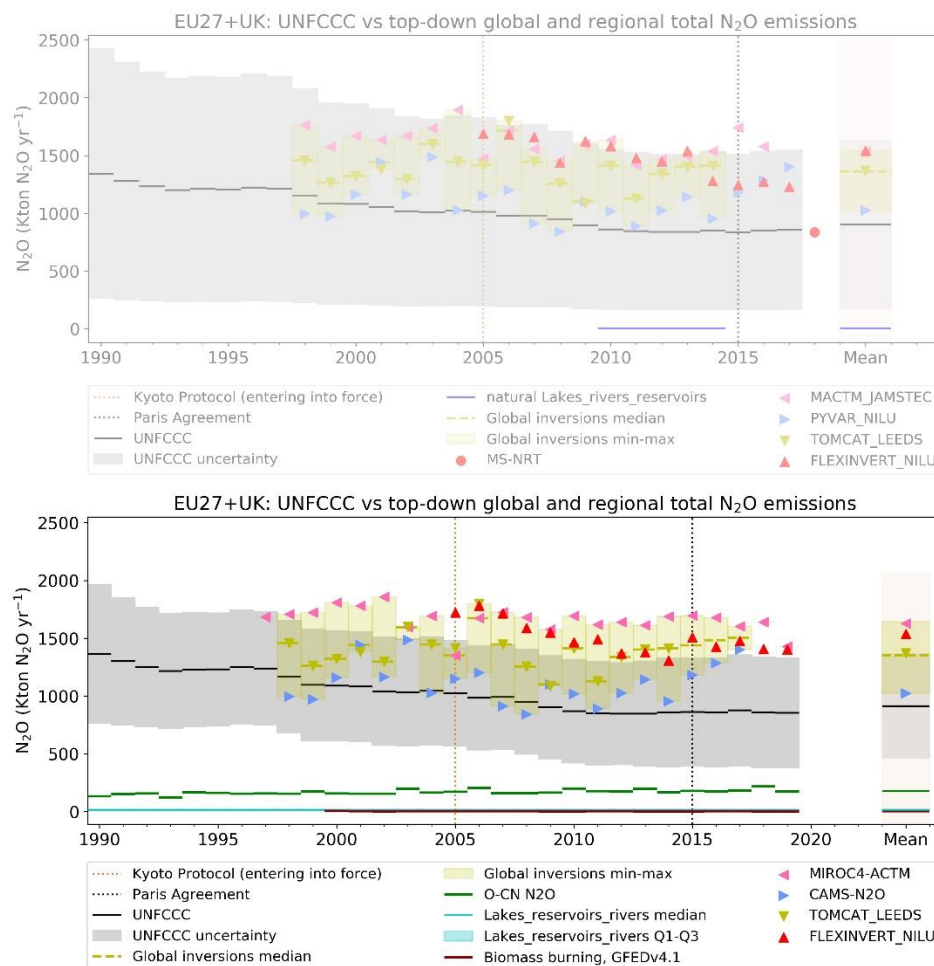


Figure 13: Comparison of total N₂O emissions for EU27+UK from one top-down regional inversion (FLEXINVERT) and three inversions (TOMCAT, CAMS-N₂O and MIROC4-ACTM) with UNFCCC NGHGI (grey) data and natural N₂O emissions (lakes_reservoirs_rivers from RECCAP2, natural pre-industrial soil emissions from the O-CN model and biomass burning from GFEDv4.1) as following: the top plot shows the previous synthesized data in Petrescu et al., 2021a and the bottom plot depicts data synthesized by the current study. The relative error on the UNFCCC value represents the UNFCCC NGHGI (2021) Member States reported uncertainties computed with the error propagation method (95% confidence interval) that were gap-filled and provided for every year (including LULUCF). Last years of the time series of the respective datasets are 2014 (TOMCAT), 2017 CAMS-N₂O, 2019 (UNFCCC, FLEXINVERT and MIROC4-ACTM). The mean column represents the common overlapping period (2005-2014) between datasets MACTM-JAMSTEC is the same with MIROC4-ACTM and PYVAR NILU with CAMS-N₂O.

For the N₂O results (FLEXINVERT in Figure 13), after updating the last runs, both the magnitude and trends of the N₂O emissions changed. This new decreasing trend is confirming the UNFCCC trend but shows a larger average

970 source after correcting for the estimate of natural emissions. Future work should strongly focus on establishing the
uncertainty and variability in the natural emissions of N₂O so that the results of inversion could be more directly
compared to emission inventories.

Spatial distribution of N₂O emissions from regional inversions

975 New top-down estimates of N₂O fluxes were produced using the CIF (Berchet et al. 2021). For N₂O,
inversions were performed by LSCE using three CHIMERE and by NILU using FLEXPART(v10.4). For the CIF
inversions, the prior fluxes and observations of N₂O were the same as those used in the Figure 13, and the background
mixing ratios were calculated using the CAMSv19r products for N₂O (Segers et al., 2020) (based on the global TM5-
980 4DVAR assimilation run (Bergamaschi et al. 2018a, Rödenbeck et al., 2009).

For N₂O, the FLEXPART inversion resulted in slightly larger fluxes over Europe compared to the CHIMERE
inversion, especially over the Netherlands, Belgium, northern France and England. In these regions, FLEXPART also
results in larger fluxes compared to the prior. FLEXPART estimated smaller fluxes compared to the prior and to
CHIMERE in northeastern Germany. CHIMERE, on the other hand remained close to the prior estimates. The total
985 mean emission for EU27+UK for the period 2005-2018 was 1538 kton N₂O and 1680 kton N₂O yr⁻¹ for CHIMERE
and FLEXPART, respectively, compared to the estimate of 1513 kton N₂O yr⁻¹ from FLEXPART (Figure 13). Both

inversions also found a decreasing trend over 2005-2018 with decreases of 157 kton N₂O yr⁻¹ and 298 kton N₂O yr⁻¹ per year for CHIMERE and FLEXPART, respectively, which was not seen in the prior estimates.

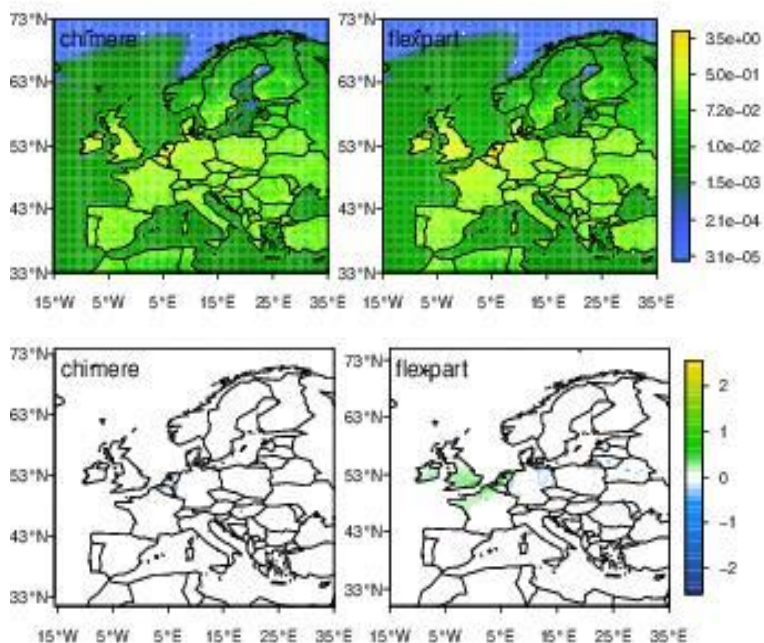


Figure 14. Posterior N₂O fluxes averaged over 2005-2018 (g N₂O m⁻² yr⁻¹) from two regional inversions CHIMERE (LSCE) and FLEXPART (NILU) shown with a log base 2 color scale (top) and the flux increments (posterior-prior) (g N₂O m⁻² yr⁻¹) shown on a linear color scale.

Similar to CH₄, the differences might be owing to the different dataset used for determining the background mixing ratios. Farther analysis is ongoing.

4. Data availability

Data files reported in this work which were used for calculations and figures are available for public download at <https://doi.org/10.5281/zenodo.7553800> ~~10.5281/zenodo.6992472~~ (Petrescu et al., 2022,2023). The data are reachable with one click (without the need for entering login and password), with a second click to download the data, consistent with the two click access principle for data published in ESSD (Carlson and Oda, 2018). The data and the DOI number are subject to future updates and only refers to this version of the paper. The raw gridded data, according to the VERIFY consortium governing document, will be made publicly available 12 months after its publication in ESSD.

5. Summary and concluding remarks

This study is an update of the first comprehensive synthesis of European CH₄ and N₂O emission estimates (Petrescu et al., (2021a), that compares total and sectoral European CH₄ and N₂O from BU (anthropogenic and natural) with TD estimates to assess their use for quality control and verification of UNFCCC NGHGI reporting. Using the most recent data, differences between TD and BU estimates were compared and comparisons were made with the

previous synthesis, Petrescu et al., (2021a). Identification of source specific uncertainty is key in understanding these differences and can lead to a reduction of the overall uncertainty in GHG inventories. Furthermore, the results have been synthesized in a way that would be compatible with the methodological framework of the first 2023 Global Stocktake of the Paris Agreement. Five-year means of CH₄ (Figure 1) and N₂O (Figure 10) emissions for the periods 2011-2015 and 2015-2019 from the different BU and TD datasets have been calculated for the EU27+UK bloc, as well as for five regions in Europe. These estimates are then compared with respective NGHGI emissions for the same periods and the 1990 base year.

Inconsistencies between CH₄ BU estimates and NGHGI data at EU27+UK level (Figure 3), are mainly caused by different methodologies in calculating emissions as highlighted in Petrescu et al. (2020, 2021a). Both BU inventories and the NGHGI use similar activity data and, to varying extents, the default EFs reported in the IPCC 2006 guidelines meaning that the estimates are predestinated to agree rather well. Thus, the spread in all BU estimates may not be indicative of the uncertainty. For global consistency purposes, EDGAR v5.0 and FAOSTAT mostly uses Tier 1 approaches in calculating emissions (and uncertainties), a fact which triggers differences with other data sources using Tier 2 or 3 methods (GAINS for all sectors and CAPRI for agriculture). Within the UNFCCC reporting process, the agriculture sector was the highest contributor to the CH₄ emissions, followed by energy and waste (Figure 1).

A reason for the small inconsistencies between datasets is the allocation of emissions to different sectors and that some data sources use updated methods and emission factors from different versions of the IPCC guidelines (e.g. 1996 versus 2006 or 2019 Refinements to the 2006 (i.e. EDGAR v6.0)).

For N₂O anthropogenic emissions, all BU data sources show good agreement with the UNFCCC NGHGI (2021) data in both trends and means (Figure 12), agriculture remaining the largest emitter (e.g. soil emissions due to fertilizer additions), within the reported uncertainties. As with CH₄, the different BU estimates share some common elements such as activity data and emissions factors, and the agreement between estimates may not be relevant to the underlying uncertainties.

An important improvement compared to Petrescu et al., 2021a, was the harmonization of UNFCCC Member States uncertainty estimates, which were gap-filled and calculated for each year of the time series. VERIFY interaction with the EU inventory team has helped improve the uncertainty estimations on the EU GHG emissions reported under UNFCCC. For both CH₄ and N₂O the uncertainties reported by NGHGI are large and underline the need for further improvement in the inventories of these two GHGs.

Regarding the TD estimates, this analysis shows that comparison between CH₄ inversions estimates and NGHGIs is highly uncertain because of the large spread in the inversion results. Nevertheless, in contrast to BU methods, TD inversions inferred from atmospheric observations represent the most independent data against which CH₄ inventory totals can be compared. With anticipated improvements in atmospheric modelling and observations, as well as modelling of natural fluxes, TD inversions may arguably emerge as the most powerful tool for verifying emissions inventories for CH₄ and other GHGs.

As TD inversions do not fully distinguish between all emission sectors used by NGHGI and report either total emissions or a coarse sectorial partitioning, their comparison to NGHGI is only possible for total emissions. It is also necessary to make an adjustment for natural emissions, which are included in TD inversions but not reported by

the NGHGs. A future improvement for the natural CH₄ emissions are the consistent time series of measurements to make clear statements about how wetland and lake fluxes change over time. For lakes there is virtually no long-term monitoring, while for wetlands variability (e.g. area) is a key uncertainty but Fronzek et al., (2018) have shown that model ensembles work well in simulating highly uncertain variables. In general regional inversions show less spread than the global inversions as they used recent updates of transport models and simulate atmospheric transport at higher resolutions.

The global models use fewer observations for Europe compared to the included regional inversions, and thus are expected to have larger uncertainties for the European fluxes. In addition, the global models are at coarser resolution, and thus likely have larger model representation errors compared to the regional ones, which may contribute to further systematic uncertainty for the European fluxes. Currently, for Europe the regional TD total is considerably higher than the global estimates (Figure 1). If the regional TD estimate for whole EU27+UK including all sources and sinks is considered to be the best total estimate in place, and if the natural fluxes are assumed to have been accurately subtracted from the optimized net flux, then NGHGI and BU approaches may be underestimating total EU27+UK CH₄ emissions by approximately 20-30%.

For N₂O, the TD estimates fall within the large range of the NGHGI uncertainties, and, in fact, the spread in the regional ensembles is much smaller than the inventory uncertainty range (Figure 13). Compared to Petrescu et al., 2021a, the natural emissions consisting of pre-industrial natural soil emissions and biomass burning emissions were included, however, for N₂O natural emissions do not explain the 415 kton N₂O difference between NGHGI (and BU) estimates and the average TD estimate. More research is thus needed to identify the source of discrepancies.

A key challenge for the inversion CH₄ community remains the separation of emissions in specific source sectors, as derived total emissions may also include natural emissions (or removals). In the case of N₂O this won't be possible since the anthropogenic emissions due to agriculture are caused by a perturbation to microbial processes (i.e., nitrification and denitrification) and cannot be cleanly separated from "natural" emissions (defined as the level of emission in the pre-industrial period, i.e., before perturbation by anthropogenic N-inputs). However, some TD quantification of industrial emissions should be possible in high-resolution inversions. In any case, TD inversions for estimating N₂O emissions should mainly focus on the trends. Furthermore, the accuracy of derived emissions and the spatial scales at which emissions can be estimated depend on the quality and density of measurements and the quality of the atmospheric models (Bergamaschi et al., 2018b). Significant further developments of the global observations system and the top-down methods would be required to support the implementation of the Paris Agreement.

The exercise of presenting uncertainty reduction maps illustrated the effect on uncertainty reductions with removing/adding ground-based observation stations. This is one of the elements informing policy makers on the need for further investing into a denser and efficient surface observation network used by inverse systems to calibrate their estimates and better inform climate policy with respect to emissions verification. This might serve to monitor and build more accurate budget of country estimates as well as provide data for inferring subnational (e.g. city-scale) emissions.

This synthesis makes use and brings together state-of-the-art BU and TD estimates from different sources and compares these data with the official NGHGI estimates reported to UNFCCC. The exercise underlines the

1095 uncertainties in the emissions of these important GHGs and illustrates the importance of regional consistent analyses
and synthesis of available estimates for informing climate policy. Specifically, the approach demonstrated here could
form the basis of the Multilateral Facilitative Consideration of Progress under the enhanced transparency framework
of the Paris Agreement. The implementation of the Paris Agreement requires accurate quantification of GHG
emissions in order to track the progress of all parties with their "Nationally Determined Contributions" and to assess
1100 collective progress towards achieving the purpose of this Agreement and its long-term goals (GST). As this will be
mainly achieved and built upon BU methodologies developed by the IPCC, we need to take into consideration the
potential to quantify GHG emissions by using "top-down" methods ("inverse modelling") (Bergamaschi et al., 2018b).
One advantage of the inverse estimate is that it provides total emission estimates inferred from atmospheric GHG
measurements. Therefore, the capability to quantify anthropogenic emissions depends on the magnitude of natural
1105 sources and sinks and the capability to quantify them and subtract them from the TD estimates.

As stated in the introduction, the main aim was to explore and discuss the issues causing differences between
NGHGI, BU and TD approaches. Such an exercise can help to improve the different respective approaches and
furthermore can inform the development of formal verification systems. Some differences in BU and NGHGI
estimates were observed and were traced back to factors such as the variations activity data, emission factors and
1110 sectoral allocation of emissions (CH₄). Nevertheless, BU and NGHGI estimates generally converged at the total
emission level for the EU27+UK bloc and the five European regions. The overall agreement is generally due to similar
sources of input activity data and emission factors (albeit with some aforementioned variations) and is not indicative
of the true uncertainties in the respective CH₄ and N₂O inventories. Indeed, NGHGI report CH₄ and N₂O emissions
with large uncertainties and, furthermore, NGHGI estimates generally diverged from the respective TD fluxes. Despite
1115 the significant spread in the inversion estimates (due to e.g. use of different transport models and/or observation
datasets, while priors might be the same (Table B2.4)), TD estimates were generally higher than NGHGI, even when
accounting for the (albeit uncertain) natural fluxes.

The analysis done here generally compared estimates in terms of long-term trends and averages over five or
more years, and thus provides a working example of how such syntheses could inform future Global Stocktakes under
1120 the Paris Agreement. A further step could involve analysis at finer temporal resolutions. While NGHGI are reported
at annual scales, analyzing emissions over intra-annual timescales, of great importance for CH₄ (wetland emission
estimates have large uncertainties and show large variability in space and between seasons) and N₂O (agricultural
fertilizer application), may help to identify sector contributions to divergence between prior and posterior estimates at
the annual/inter-annual scale. To do this, however requires expanded in-situ monitoring so that such dynamics can be
1125 better represented in the temporally-resolved prior estimates that feed into the top-down inversions.

6. Appendices

1130 **Appendix A: Data sources, methodology and uncertainty descriptions**

The country specific plots are found at: <http://webportals.ipsl.jussieu.fr/VERIFY/FactSheets/> v1.2428

VERIFY project

VERIFY's primary aim was to develop scientifically robust methods to assess the accuracy and potential biases in national inventories reported by the parties through an independent pre-operational framework. The main concept is to provide observation-based estimates of anthropogenic and natural GHG emissions and sinks as well as associated uncertainties. The proposed approach is based on the integration of atmospheric measurements, improved emission inventories, ecosystem data, and satellite observations, and on an understanding of processes controlling GHG fluxes (ecosystem models, GHG emission models).

Two complementary approaches relying on observational data-streams were combined in VERIFY to quantify GHG fluxes:

1) atmospheric GHG concentrations from satellites and ground-based networks (top-down atmospheric inversion models) and

2) bottom-up activity data (e.g. fuel use and emission factors) and ecosystem measurements (bottom-up models).

For CH₄ and N₂O, agricultural emissions were separated from fossil fuel and industrial emissions. Finally, trends in the budget of the three GHGs were analyzed in the context of NDC targets.

The objectives of VERIFY were:

Objective 1. Integrate the efforts between the research community, national inventory compilers, operational centres in Europe, and international organizations towards the definition of future international standards for the verification of GHG emissions and sinks based on independent observation.

Objective 2. Enhance the current observation and modelling ability to accurately and transparently quantify the sinks and sources of GHGs in the land-use sector for the tracking of land-based mitigation activities.

Objective 3. Develop new research approaches to monitor anthropogenic GHG emissions in support of the EU commitment to reduce its GHG emissions by 40% by 2030 compared to the year 1990.

Objective 4. Produce periodic scientific syntheses of observation-based GHG balance of EU countries and practical policy-oriented assessments of GHG emission trends, and apply these methodologies to other countries.

For more information on project team and products/results check <https://verify.lsce.ipsl.fr/>.

Table A1: *Country grouping used for reconciliation purposes between BU and TD estimates. The countries and groups of countries in italic are not directly used by this study but their figures and data is available on the VERIFY project web portal at: <http://webportals.ipsl.jussieu.fr/VERIFY/FactSheets/>.*

Country name – geographical Europe	BU-ISO3	Aggregation from TD-ISO3
Luxembourg	LUX	
Belgium	BEL	BENELUX
Netherlands	NLD	BNL
Bulgaria	BGR	BGR
Switzerland	CHE	

<i>Lichtenstein</i>	<i>LIE</i>	<i>CHL</i>
Czech Republic	CZE	Former Czechoslovakia
Slovakia	SVK	CSK
Austria	AUT	AUT
Slovenia	SVN	North Adriatic countries
Croatia	HRV	NAC
Romania	ROU	ROU
Hungary	HUN	HUN
Estonia	EST	
Lithuania	LTU	Baltic countries
Latvia	LVA	BLT
Norway	NOR	NOR
Denmark	DNK	
Sweden	SWE	
Finland	FIN	DSF
Iceland	ISL	ISL
Malta	MLT	MLT
Cyprus	CYP	CYP
France (Corsica incl.)	FRA	FRA
<i>Monaco</i>	<i>MCO</i>	
<i>Andorra</i>	<i>AND</i>	
Italy (Sardinia, Vatican incl.)	ITA	ITA
<i>San Marino</i>	<i>SMR</i>	
United Kingdom (Great Britain + N Ireland)	GBR	UK
<i>Isle of Man</i>	<i>IMN</i>	
Iceland		
Ireland	IRL	IRL
Germany	DEU	DEU
Spain	ESP	IBERIA
Portugal	PRT	IBE
Greece	GRC	GRC
<i>Russia (European part)</i>	<i>RUS European</i>	
<i>Georgia</i>	<i>GEO</i>	<i>RUS European+GEO</i>

<i>Russian Federation</i>	<i>RUS</i>	<i>RUS</i>
Poland	POL	POL
<i>Turkey</i>	<i>TUR</i>	<i>TUR</i>
EU27+UK (Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Germany, Denmark, Spain, Estonia, Finland, France, Greece, Croatia, Hungary, Ireland, Italy, Lithuania, Latvia, Luxembourg, Malta, Netherlands, Poland, Portugal, Romania, Slovakia, Slovenia, Sweden, United Kingdom)	AUT, BEL, BGR, CYP, CZE, DEU, DNK, ESP, EST, FIN, FRA, GRC, HRV, HUN, IRL, ITA, LTU, LVA, LUX, MLT, NDL, POL, PRT, ROU, SVN, SVK, SWE, GBR	E28
Western Europe (Belgium, France, United Kingdom, Ireland, Luxembourg, Netherlands)	BEL, FRA, UK, IRL, LUX, NDL	WEE
Central Europe (Austria, Switzerland, Czech Republic, Germany, Hungary, Poland, Slovakia)	AUT, CHE, CZE, DEU, HUN, POL, SVK	CEE
Northern Europe (Denmark, Estonia, Finland, Lithuania, Latvia, Norway, Sweden)	DNK, EST, FIN, LTU, LVA, NOR, SWE	NOE
<i>South-Western Europe (Spain, Italy, Malta, Portugal)</i>	<i>ESP, ITA, MLT, PRT</i>	<i>SWN</i>
<i>South-Eastern Europe (all) (Albania, Bulgaria, Bosnia and Herzegovina, Cyprus, Georgia, Greece, Croatia, Macedonia, the former Yugoslav, Montenegro, Romania, Serbia, Slovenia, Turkey)</i>	<i>ALB, BGR, BIH, CYP, GEO, GRC, HRV, MKD, MNE, ROU, SRB, SVN, TUR</i>	<i>SEE</i>
<i>South-Eastern Europe (non-EU) (Albania, Bosnia and Herzegovina, Macedonia, the former Yugoslav, Georgia, Turkey, Montenegro, Serbia)</i>	<i>ALB, BIH, MKD, MNE, SRB, GEO, TUR</i>	<i>SEA</i>
<i>South-Eastern Europe (EU) (Bulgaria, Cyprus, Greece, Croatia, Romania, Slovenia)</i>	<i>BGR, CYP, GRC, HRV, ROU, SVN</i>	<i>SEZ</i>
<i>Southern Europe (all) (SOE) (Albania, Bulgaria, Bosnia and Herzegovina, Cyprus, Georgia, Greece, Croatia, Macedonia, the former Yugoslav, Montenegro, Romania, Serbia, Slovenia, Turkey, Italy, Malta, Portugal, Spain)</i>	<i>ALB, BGR, BIH, CYP, GEO, GRC, HRV, MKD, MNE, ROU, SRB, SVN, TUR, ITA, MLT, PRT, ESP</i>	<i>SOE</i>
<i>Southern Europe (non-EU) (SOY) Albania, Bosnia and Herzegovina, Georgia,</i>	<i>ALB, BIH, GEO, MKD, MNE, SRB, TUR,</i>	<i>SOY</i>

<i>Macedonia, the former Yugoslav, Montenegro, Serbia, Turkey)</i>		
Southern Europe (EU) (SOZ) (Bulgaria, Cyprus, Greece, Croatia, Romania, Slovenia, Italy, Malta, Portugal, Spain)	BGR, CYP, GRC, HRV, ROU, SVN, ITA, MLT, PRT, ESP	SOZ
Eastern Europe (non-EU) (Belarus, Moldova, Republic of, Russian Federation, Ukraine)	BLR, MDA, RUS, UKR	EAE
<i>EU-15 (Austria, Belgium, Germany, Denmark, Spain, Finland, France, United Kingdom, Greece, Ireland, Italy, Luxembourg, Netherlands, Portugal, Sweden)</i>	<i>AUT, BEL, DEU, DNK, ESP, FIN, FRA, GBR, GRC, IRL, ITA, LUX, NDL, PRT, SWE</i>	<i>E15</i>
<i>EU-27 (Austria, Belgium, Bulgaria, Cyprus, Czech Republic, Germany, Denmark, Spain, Estonia, Finland, France, Greece, Croatia, Hungary, Ireland, Italy, Lithuania, Latvia, Luxembourg, Malta, Netherlands, Poland, Portugal, Romania, Slovakia, Slovenia, Sweden)</i>	<i>AUT, BEL, BGR, CYP, CZE, DEU, DNK, ESP, EST, FIN, FRA, GRC, HRV, HUN, IRL, ITA, LTU, LVA, LUX, MLT, NDL, POL, PRT, ROU, SVN, SVK, SWE</i>	<i>E27</i>
<i>All Europe (Aaland Islands, Albania, Andorra, Austria, Belgium, Bulgaria, Bosnia and Herzegovina, Belarus, Switzerland, Cyprus, Czech Republic, Germany, Denmark, Spain, Estonia, Finland, France, Faroe Islands, United Kingdom, Guernsey, Greece, Croatia, Hungary, Isle of Man, Ireland, Iceland, Italy, Jersey, Liechtenstein, Lithuania, Luxembourg, Latvia, Moldova, Republic of, Macedonia, the former Yugoslav, Malta, Montenegro, Netherlands, Norway, Poland, Portugal, Romania, Russian Federation, Svalbard and Jan Mayen, San Marino, Serbia, Slovakia, Slovenia, Sweden, Turkey, Ukraine)</i>	<i>ALA, ALB, AND, AUT, BEL, BGR, BIH, BLR, CHE, CYP, CZE, DEU, DNK, ESP, EST, FIN, FRA, FRO, GBR, GGY, GRC, HRV, HUN, IMN, IRL, ISL, ITA, JEY, LIE, LTU, LUX, LVA, MDA, MKD, MLT, MNE, NDL, NOR, POL, PRT, ROU, RUS, SJM, SMR, SRB, SVK, SVN, SWE, TUR, UKR</i>	<i>EUR</i>

*countries highlighted in *italic* are not discussed in the current 2019 synthesis mostly because unavailability of NGHGI data (non-Annex I countries¹¹) but are present on the web-portal: <http://webportals.ipsl.jussieu.fr/VERIFY/FactSheets/>. Results of Annex I countries (NOR, CHE, ISL) and non-EU EAE countries/groups are represented in Figures 1 and 9.

¹¹Non-Annex I countries are mostly developing countries. The reporting to UNFCCC is implemented through national communications (NCs) and biennial update reports (BURs): <https://unfccc.int/national-reports-from-non-annex-i-parties>.

1165 **Table A2: Main methodological changes (in bold) of current study with respect to Petrescu et al., 2020 and 2021a;**
n/a cells mean that there is no data available.

<u>Publication year</u>	<u>Gas</u>	<u>Bottom-up anthropogenic CH₄ / N₂O emissions</u>			<u>Bottom-up natural CH₄ / N₂O emissions</u>	<u>Top-down CH₄ / N₂O emissions</u>		<u>Uncertainty and other changes</u>
		<u>Inventories</u>	<u>Global databases</u>	<u>Emission models</u>	<u>Emission models</u>	<u>Regional models</u>	<u>Global models</u>	
<u>Petrescu et al., 2020</u>	<u>CH₄</u>	<u>National emissions from UNFCCC (2018) 1990-2016</u> <u>AFOLU sector (Agriculture and LULUCF) EU28 data for four years (1990, 2005, 2010 and 2016)</u>	<u>EGDAR v4.3.2 1990-2012</u> <u>EDGAR FT2017 1990-2016</u> <u>FAOSTAT 1990-2016</u> <u>Agriculture sector EU28 data for four years (1990, 2005, 2010 and last reported year)</u>	<u>CAPRI 1990-2013</u> <u>GAINS 1990-2015</u> <u>Agriculture sector EU28 data for four years (1990, 2005, 2010 and last reported year)</u>	<u>Natural (wetlands) CH₄ emissions model ensemble GCP (2018) Poulter et al. (2017)</u> <u>Time series 1990-2017</u>	<u>n/a</u>	<u>n/a</u>	<u>UNFCCC (2018) uncertainty estimates for 2016 (error propagation 95% interval method)</u> <u>EDGAR v.4.3.2. reports only for 2012</u>
	<u>N₂O</u>	<u>National emissions from UNFCCC (2018) 1990-2016</u> <u>Agriculture sector EU28 data for four years (1990, 2005, 2010 and 2016)</u>	<u>EGDAR v4.3.2 1990-2012</u> <u>EDGAR FT2017 1990-2016</u> <u>FAOSTAT 1990-2016</u> <u>Agriculture sector EU28 data for four years (1990, 2005, 2010 and last reported year)</u>	<u>CAPRI 1990-2013</u> <u>GAINS 1990-2015</u> <u>Agriculture sector EU28 data for four years (1990, 2005, 2010 and last reported year)</u>	<u>n/a</u>	<u>n/a</u>	<u>n/a</u>	<u>UNFCCC (2018) uncertainty estimates for 2016</u> <u>EDGAR v.4.3.2. reports only for 2012</u>
<u>Petrescu et al., 2021a</u>	<u>CH₄</u>	<u>National emissions from UNFCCC (2019)</u>	<u>EGDAR v 5.0 1990-2015</u>	<u>CAPRI 1990-2013</u>	<u>Non-wetland inland waters</u>	<u>Total CH₄ column Time series 2005-2017:</u>	<u>Anthropogenic and natural partitions</u>	<u>UNFCCC (2018) uncertainty estimates for 2016 (error</u>

			1990-2017 All UNFCCC sectors EU27+UK time series and 2018 MS-NRT estimate (EEA, 2019) Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR)	FAOSTAT (only agriculture) 1990-2017 Anthropogenic EU27+UK time series (excl. LULUCF) Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR) Excl. LULUCF	GAINS 1990-2015 Agriculture sector EU27+UK Times series	Average 2005-2011 Geological fluxes Total pre-industrial era JSBACH-HIMMELI 2005-2017	FLEXPART - FLExKF TM5-4DVAR FLEXINVERT NILU CTE-FMI InTEM-NAME Only for UK InGOS inversions 2006-2012	GCP-GCB 2019 2000-2017 EDGAR v.4.3.2. reports only for 2015 For model ensembles reported as variability in extremes (min/max)
	N₂O	National emissions from UNFCCC (2019) 1990-2017 All UNFCCC sectors EU27+UK time series and 2018 MS-NRT estimate (EEA, 2019) Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR)	EGDAR v 5.0 1990-2015 (excl. LULUCF) FAOSTAT (only agriculture) 1990-2017 Anthropogenic EU27+UK time series Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR) Excl. LULUCF	Agriculture CAPRI 1990-2013 ECOSSE 1990-2018	N₂O missions from lakes, rivers, reservoirs Average 2010-2014	Total N₂O column Time series FLEXINVERT NILU 2005-2017	Total N₂O column Time series GCP - GN₂OB 2019 CAMS-N₂O TOMCAT MIROC4-ACTM 1998-2016	UNFCCC (2018) uncertainty estimates for 2016 (error propagation 95% interval method) EDGAR v.4.3.2. reports only for 2015 For model ensembles reported as variability in extremes (min/max)
Current study	CH₄	National emissions from UNFCCC (2021) 1990-2019 All UNFCCC sectors	EGDAR v6.0 1990-2018 FAOSTAT (only agriculture) 1990-2020	CAPRI 1990-2014 and 2016 and 2018 GAINS	One median value from process-based models for non-wetland inland waters (lakes, rivers, reservoirs) 2010-2019	Total CH₄ column Time series 2005-2018: FLEXPART - FLExKF	Anthropogenic and natural partitions GCP-GCB 2000-2017	UNFCCC (2021) uncertainty estimates for 2019 (error propagation 95% interval method) and yearly uncertainties provided by the EU GHG inventory team EDGAR reports only for 2015 values from v6.0

			EU27+UK time series Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR) excl. LULUCF Two means: 2011-2015 2015-2019	Anthropogenic EU27+UK time series (excl. LULUCF) Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR) excl. LULUCF	1990-2015 Agriculture sector EU27+UK Times series	One median value from upscaled results in RECCAP2 1990-2019 Geological fluxes Total pre-industrial era updates for EU27+UK JSBACH-HIMMELI peatlands and mineral soils 2005-2020 Biomass burning GFEDv4.1 emissions	TM5-4DVAR CTE-FMI FLEXINVERT NILU 1990-2019 InGOS inversions 2006-2012 not included anymore in the mean InTEM-NAME (only for UK plots on the VERIFY website) VERIFY Community Inversion Framework (CIF): CHIMERE, FLEXPARTv10.4 (NILU) and FELXPART (EMPA) 2006-2017	For model ensembles reported as variability in extremes (min/max) CAPRI uncertainties for 2014, 2016 and 2018
	N₂O	National emissions from UNFCCC (2021) 1990-2019 All UNFCCC sectors EU27+UK time series Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR) excl. LULUCF Two means: 2011-2015 2015-2019	EGDAR v 6.0 1990-2018 FAOSTAT (v2021) (only agriculture) 1990-2020 Anthropogenic EU27+UK time series (excl. LULUCF) Regional EU27+UK totals (incl. NOR, CHE, UKR, MLD and BLR) excl. LULUCF	CAPRI 1990-2014 and 2016 and 2018 GAINS 1990-2015 Agriculture sector EU27+UK Times series ECOSSE 1990-2020	One N₂O median value for emissions from lakes, rivers, reservoirs (RECCAP2) Average 1990-2019 Natural N₂O pre-industrial emissions from O-CN model 1990-2020 Biomass burning GFEDv4.1 emissions 2000-2019	Total N₂O column Time series FLEXINVERT NILU 2005-2019 VERIFY Community Inversion Framework (CIF): CHIMERE, FLEXPARTv10.4 (NILU) 2005-2018	Total N₂O column Time series CAMS-N₂O 1998-2017 TOMCAT 1998-2014 MIROC4-ACTM 1997-2019 FELXINVERT 2005-2019	UNFCCC (2021) uncertainty estimates for 2019 (error propagation 95% interval method) and yearly uncertainties provided by the EU GHG inventory team EDGAR reports only for 2015 values from v6.0 For model ensembles reported as variability in extremes (min/max) CAPRI uncertainties for 2014, 2016 and 2018

A1: Anthropogenic CH₄ emissions (sectors Energy, IPPU, Agriculture, LULUCF and Waste)

Bottom-up CH₄ emissions estimates

UNFCCC NGHGI (2019/2021)

Under the UNFCCC and its Kyoto Protocol national greenhouse gas (GHG) inventories are the most important source of information to track progress and assess climate protection measures by countries. In order to build mutual trust in the reliability of GHG emission information provided, national GHG inventories are subject to standardized reporting requirements, which have been continuously developed by the Conference of the Parties (COP)¹². The calculation methods for the estimation of greenhouse gases in the respective sectors is determined by the methods provided by the 2006 IPCC Guidelines for National Greenhouse Gas Inventories (IPCC, 2006). These Guidelines provide detailed methodological descriptions to estimate emissions and removals, as well as provide recommendations to collect the activity data needed. As a general overall requirement, the UNFCCC reporting guidelines stipulate that reporting under the Convention and the Kyoto Protocol follow the five key principles of transparency, accuracy, completeness, consistency and comparability (TACCC).

The reporting under UNFCCC should meet the TACCC principles. The three main GHGs are reported in time series from 1990 up to two years before the due date of the reporting. The reporting is strictly source category based and is done under the Common Reporting Format tables (CRF), downloadable from the UNFCCC official submission portal: <https://unfccc.int/ghg-inventories-annex-i-parties/2021>.

The UNFCCC NGHGI anthropogenic CH₄ and N₂O emissions analyzed in this study include estimates from 4 key sectors for the EU27+UK: 1 Energy, 2 Industrial processes and product use (IPPU), 3 Agriculture and 5 Waste. The methodological tiers a country applies depends on the source contribution to the national total (*Key Category* or not), on the national circumstances and the individual conditions of the land, which explains the variability of uncertainties among the sector itself as well as among EU countries. The LULUCF CH₄ and N₂O emissions are very small but are included in some figures (see Table 1).

Gap-filling harmonization procedure for NGHGI uncertainties

The presented uncertainties in the emission levels of the individual countries and the EU27+UK bloc were calculated by using the methods and data used to compile the official GHG emission uncertainties that are reported by the EU under the UNFCCC (NIRs, 2022). The EU uncertainty analysis reported in the bloc's National Inventory Report (NIR) is based on country-level, Approach 1 uncertainty estimates (IPPC, 2006, Vol. 1, Chap. 3) that are reported by EU Member States, Iceland and United Kingdom under Article 7(1)(p) of Regulation (EU) 525/2013. These country-level uncertainty estimates are typically reported at beginning of a submission cycle and are not always revised with updated CRF submissions later in the submission cycle. Furthermore, the compiled uncertainties of some countries are incomplete (e.g. uncertainties not estimated for LULUCF and/or indirect CO₂ emissions, certain

¹² The last revision has been made by COP 19 in 2013 (UNFCCC, 2013)

subsector emissions are confidential) and the sector and gas resolution at which uncertainties are provided varies between the countries. The EU inventory team therefore implements a procedure to harmonise and gap-fill these uncertainty estimates. A processing routine reads the individual country uncertainty files that are pre-formatted manually to assign consistent sector and gas labels to the respective estimates of emissions/removals and uncertainties. The uncertainty values are then aggregated to a common sector resolution, at which the emissions and removals reported in the uncertainty tables of the countries are then replaced with the respective values from the final CRF tables of the countries. Due to the issue of incompleteness mentioned above, the country-level data are then screened to identify residual GHG emissions and removals for which no uncertainty estimates have been provided. Where sectors are partially complete, the residual net emission is quantified in CO₂ equivalents and incorporated. An uncertainty is then estimated, by calculating the overall sector uncertainty of the sources and sinks that were included in that country's reported uncertainties estimates and assigning this percentage average to the residual net emission. In cases where for certain sectors no uncertainties have been provided at all (e.g. indirect CO₂ emissions, LULUCF), an average (median) sector uncertainty in percent is calculated from all the countries for which complete sectoral emissions and uncertainties were reported, and this average uncertainty is assigned to the country's sector GHG total reported in its final CRF tables.

The country-level uncertainties presented in this paper were provided by the EU GHG inventory team and have been compiled using this same processing routine and using the uncertainties and CRF data reported by the countries in the 2021 submission. However, for the purpose of this synthesis work, the method has been expanded to gap-fill at the individual greenhouse gas level (CH₄ and N₂O emissions only) rather than at the aggregate GHG level. Furthermore, the expanded method here assigns the sub-sectoral uncertainties to the emissions and removals of the entire time series (1990-2019), rather than just the base year and latest year of the respective time series. This allows uncertainties to be sensitive to the sub-sectoral contributions to sectoral and national total emissions, which of course change over time. For each year of the time series, uncertainties in the total and sectoral CH₄ and N₂O emissions are calculated using Gaussian error propagation, by summing the respective sub-sectoral uncertainties (expressed in kton CH₄ and N₂O) in quadrature and assuming no error correlation. In contrast, for the EU27+UK bloc, uncertainties in the total and sectoral CH₄ and N₂O emissions were calculated to take into account error correlations between the respective country estimates at the subsector level. This was done by applying the same methods and assumptions described in the 2022 EU NIR (UNFCCC NIR, 2022). The subsector resolution applied for gap-filling allows the routine to access respective data on emission factors from CRF Table *Summary 3* and apply correlation coefficients (r) when aggregating the uncertainties. For a given subsector, it is assumed that the errors of countries using default factors are completely correlated ($r = 1$), while errors of countries using country-specific factors are assumed uncorrelated ($r = 0$). For countries using a mix of default and country-specific factors at the given subsector level, it is assumed that these errors are partially correlated ($r = 0.5$) with one another and with the errors of countries using the default factors only.

Based on these correlation assumptions, the routine then aggregates CH₄ and N₂O emissions and uncertainties for the specified subsector resolution at the EU27+UK level. Uncertainties at sector total level are then aggregated from the subsector estimates assuming no correlation between subsectors. However, for countries reporting very

coarse resolution estimates (e.g. total sector CH₄ and N₂O emissions) or where the sector has been partially or completely gap-filled, it is assumed that these uncertainties are partially correlated ($r = 0.5$) with one another and with the other reported subsector level estimates. Level uncertainties on the total EU27+UK CH₄ and N₂O emissions (with and without LULUCF) are then aggregated from the sector estimates assuming no error correlation between sectors.

EDGAR v5.0

The Emissions Database for Global Atmospheric Research (EDGAR) is an independent global emission inventory of greenhouse gases (GHG) and air pollutants developed by the Joint Research Centre of the European Commission (<https://edgar.jrc.ec.europa.eu/index.php>). The first edition of the Emissions Database for Global Atmospheric Research was published in 1995. The dataset now includes almost all sources of fossil CO₂ emissions, is updated annually, and reports data for 1970 to n-1. Estimates are provided by sector. Emissions are estimated fully based on statistical data from 1970 till 2018 <https://data.jrc.ec.europa.eu/dataset/97a67d67-c62e-4826-b873-9d972c4f670b>. For complete description see Andrew (2020).

Uncertainties: EDGAR uses emission factors (EFs) and activity data (AD) to estimate emissions. Both EFs and AD are uncertain to some degree, and when combined, their uncertainties need to be combined too. To estimate EDGAR's uncertainties (stemming from lack of knowledge of the true value of the EF and AD), the methodology devised by IPCC (2006, Chapter 3) is adopted, that is the sum of squares of the uncertainty of the EF and AD (uncertainty of the product of two variables). A log-normal probability distribution function is assumed to avoid negative values, and uncertainties are reported as 95 % confidence interval according to IPCC (2006, chapter 3, equation 3.7). For emission uncertainty in the range 50 % to 230% a correction factor is adopted as suggested by Frey et al (2003) and IPCC (2006, chapter 3, equation 3.4). Uncertainties are published in Solazzo et al., 2021.

CAPRI

CAPRI is an economic, partial equilibrium model for the agricultural sector, focused on the EU (Britz and Witzke, 2014¹³; Weiss and Leip, 2012¹⁴). CAPRI stands for 'Common Agricultural Policy Regionalised Impact analysis', and the name hints at the main objective of the system: assessing the effect of CAP policy instruments not only at the EU or Member State level but at sub-national level too. The model is calibrated for the base year (currently 2012) and then baseline projections are built, allowing the ex-ante evaluation of agricultural policies and trade policies on production, income, markets, trade and the environment.

Among other environmental indicators, CAPRI simulates CH₄ emissions from agricultural production activities (enteric fermentation, manure management, rice cultivation, agricultural soils). Activity data is mainly based on FAOSTAT and EUROSTAT statistics and estimation of emissions follows IPCC 2006 methodologies, with a higher or lower level of detail depending on the importance of the emission source. Details on CAPRI methodology

¹³ https://www.capri-model.org/docs/CAPRI_documentation.pdf

¹⁴ <https://www.sciencedirect.com/science/article/pii/S0167880911004415>

for emissions calculations is referenced in the Annex Table A1. For this study CAPRI updated emissions for three years: 2014, 2016 and 2018.

Uncertainties were calculated for the updated years, 2014, 2016 and 2018. The uncertainty of the spatial allocation of emissions for CH₄ and N₂O has been calculated by taking into account the uncertainty of the spatial disaggregation and the uncertainty of the emission sources, assuming:

- the disaggregation having an uncertainty of 50% for N₂O and 20% for CH₄.

- the emission processes have uncertainty of:

○ 50%: N₂O soil processes;

○ 30%: N₂O manure processes;

○ 30%: CH₄ manure and enteric, and

○ 10%: rice.

Then, for each cell, the uncertainty of the disaggregation and that of the process are combined as they are independent (sum of squares, see Solazzo et al., 2021) and then the total uncertainty for the grid cell is aggregated using emission weighted sum of squares.

GAINS

Specific sectors and abatement technologies in GAINS vary by the specific emitted compound, with source sector definition and emission factors largely following the IPCC methodology at the Tier 1 or Tier 2 level. GAINS includes in general all anthropogenic emissions to air, but does not cover emissions from forest fires, savannah burning and land use / land use change. Emissions are estimated for 174 countries/regions, with the possibility to aggregate to a global emission estimate, and spanning a timeframe from 1990 to 2050 in five-year intervals. Activity drivers for macroeconomic development, energy supply and demand, and agricultural activities are entered externally, GAINS extends with knowledge required to estimate “default” emissions (emissions occurring due to an economic activity without emission abatement) and emissions and costs of situations under emission control (see Amann et al., 2001).

The GAINS model covers all source sectors of anthropogenic methane (CH₄) emissions; agricultural sector emissions from livestock, rice cultivation and agricultural waste burning, energy sector emissions from upstream and downstream sources in fossil fuel extraction and use, and emissions from handling and treatment of solid waste and wastewater source sectors. A description of the modelling of CH₄ emissions in GAINS is presented in Höglund-Isaksson (2020). Generation of solid waste and the carbon content of wastewater are derived within the model in consistency with the relevant macroeconomic scenario. The starting point for estimations of anthropogenic CH₄ is the methodology recommended in the IPCC (2006 and revision in 2019) guidelines, for most source sectors using country-specific information to allow for deriving country- and sector/technology- specific emission factors at a Tier 2 level. Consistent methodologies were further developed to estimate emissions from oil and gas systems (Höglund-Isaksson, 2017) and solid waste (Höglund-Isaksson, 2018; Gómez-Sanabria et al., 2018). Emission factors are specified in a consistent manner across countries for given sets of technology and with past implementation of emission abatement measures reflected as changes in technology structures. The resulting emission estimates are well comparable across geographic and temporal scales. The GAINS approach to calculate waste emissions is developed in consistency with

the First-Order-Decay method recommended by IPCC (2006 and 2019 revision), applying different decay periods when estimating emissions from flows of different types of organic waste, i.e., food & garden, paper, wood, textile and other. Data on waste generation, composition and treatment are taken from EUROSTAT (2019) and complemented with national information from the UNFCCC (2019) Common Reporting Format tables on the amounts of waste diverted to landfills of various management levels and to treatment e.g., recycling, composting, biodigestion and incineration.

Uncertainties: Uncertainty is prevalent among many different dimensions both in the estimations of emissions, abatement potentials and costs. When constructing global bottom-up emission inventories at a detailed country and source level, it is inevitable that some information gaps will be bridged using default assumptions. As it is difficult to speculate about how such sources of uncertainty affect resulting historical and future emission estimates, we instead address uncertainty in historical emissions by making comparisons to estimates by other publicly available and independently developed bottom-up inventories and various top-down estimates consistent with atmospheric measurements and inverse model results. Although existing publicly available global bottom-up inventories adhere to the recommended guidelines of the IPCC (2006), the flexibility in these is large and results will depend on the availability and quality of gathered source information. There is accordingly a wide range of possible sources of uncertainty built into estimations in such comprehensive efforts. Having a pool of independently developed inventories, each with its own strengths and weaknesses, can improve the understanding of the scope for uncertainty, in particular when compared against top-down atmospheric measurements.

FAOSTAT

FAOSTAT: The Food and Agricultural Organization of the United Nations (FAO), provides CH₄ -emissions totals or per gas/activity from agriculture and LULUCF available at: <https://www.fao.org/faostat/en/#data/GT>. The FAOSTAT emissions database is computed following Tier 1 IPCC 2006 Guidelines for National GHG Inventories (<http://www.ipcc-nggip.iges.or.jp/public/2006gl/index.html>). Country reports to FAO on crops, livestock and agriculture use of fertilizers are the source of activity data. Geospatial data are the source of AD for the estimates from cultivation of organic soils, biomass and peat fires. GHG emissions are provided by country, regions and special groups, with global coverage, relative to the period 1961-present (with annual updates, currently to 20172019) and with projections for 2030 and 2050, expressed as CO₂e for CH₄, by underlying agricultural emission sub-domain and by aggregate (agriculture total, agriculture total plus energy, agricultural soils). LULUCF emissions consist of CH₄ (methane) associated with biomass and peat fires. Comparison to the UNFCCC submissions is also provided. **Uncertainties** were computed by Tubiello et al., 2013 but are not available in the FAOSTAT database.

Top-down anthropogenic CH₄ emission estimates

FLEXPART – FLExKF

FLExKF applies an Extended Kalman Filter (Brunner et al. 2012) in combination with backward Lagrangian transport simulations using the model FLEXPART (Stohl et al. 2005; Pisso et al. 2019). It optimizes surface-atmosphere fluxes by assimilating atmospheric observations in a sequential manner, which allows for an analytical solution for relatively large inversion problems (long time periods, number stations O(100)). Since model-observation

residuals typically follow a log-normal distribution, the method optimizes log-transformed emissions, which also guarantees a positive solution. Source-Receptor Matrices (Seibert and Frank, 2004) were computed at $0.25^\circ \times 0.25^\circ$ resolution with FLEXPART driven by ECMWF Era Interim meteorological fields in the same way as for FlexInvert. Backward simulations were limited to 10 days prior to each observation and to the domain $15^\circ\text{W} - 35^\circ\text{E}$, $30^\circ\text{N} - 75^\circ\text{N}$. Fluxes were estimated for this domain on a monthly basis at $0.5^\circ \times 0.5^\circ$ resolution. For the version used in this study, FLEXPART-CAMSv19r-EMPA, the background mole fraction was taken from CAMSv19r which is based on the global TM5-4DVAR assimilation run (Bergamaschi et al. 2018a) where the above domain was cut out following the two-step approach of Rödenbeck et al. (2009).

Uncertainties: The uncertainty in the posterior fluxes is composed of random and systematic errors. The random uncertainties are represented by the posterior error covariance matrix provided by the Kalman Filter, which combines errors in the prior fluxes with errors in the observations and model representation. Systematic uncertainties primarily arise from systematic errors in modelled atmospheric transport and in background mole fractions, but also include aggregation errors, i.e. errors arising from the way the flux variables are discretized in space and time.

FLEXINVERT

The FlexInvert framework is based on Bayesian statistics and optimizes surface-atmosphere fluxes using the maximum probability solution (Rodgers, 2000). Atmospheric transport is modelled using the Lagrangian model FLEXPART (Stohl et al., 2005; Pisso et al., 2019) run in the backwards time mode to generate a so-called Source-Receptor Matrix (SRM). The SRM describes the relationship between the change in mole fraction and the fluxes discretized in space and time (Seibert and Frank, 2004) and was calculated for 8 days prior to each observation. For use in the inversions, FLEXPART was driven using ECMWF operational analysis wind fields. The state vector consisted of prior fluxes discretized on an irregular grid based on the SRMs (Thompson et al. 2014). This grid has finer resolution (in this case the finest was $0.25^\circ \times 0.25^\circ$) where the fluxes have a strong influence on the observations and coarser resolution where the influence is only weak (the coarsest was $2^\circ \times 2^\circ$). The fluxes were solved at 10-days temporal resolution. The state vector also included scalars for the background contribution. The background mixing ratio, i.e., the contribution to the mixing ratio that is not accounted for in the 8-day SRMs, was estimated by coupling the termination points of backwards trajectories (modelled using virtual particles) to initial fields of methane simulated with the Lagrangian FLEXPART-CTM model, which was developed at Empa based on FLEXPART (Stohl et al., 2005; Pisso et al., 2019). ~~In these simulations, we applied the d~~Data assimilation method described by Groot Zwaafink et al. (2018) that constrains modelled fields with surface observations through nudging was applied for current simulations.

Uncertainties: The posterior fluxes are subject to systematic errors primarily from: 1) errors in the modelled atmospheric transport; 2) aggregation errors, i.e. errors arising from the way the flux variables are discretized in space and time; 3) errors in the background methane fields; and 4) the incomplete information from the observations and hence the dependence on the prior fluxes. In addition, there is, to a smaller extent, some error due to calibration offsets between observing instruments. Uncertainties in the observation space were inflated to take into account the model representation errors

InGOS and TM5-4DVAR

The atmospheric models used within the European FP7 project InGOS (Integrated non-CO₂ Greenhouse gas Observing System) are described by Bergamaschi et al., 2018a and Supplement (<https://www.atmos-chem-phys.net/18/901/2018/acp-18-901-2018-supplement.pdf>). The models include global Eulerian models with a zoom over Europe (TM5-4DVAR, TM5-CTE, LMDZ), regional Eulerian models (CHIMERE), and Lagrangian dispersion models (STILT, NAME, COMET). The horizontal resolutions over Europe are ~ 1.0 – 1.2° (longitude) \times ~ 0.8 – 1.0° (latitude) for the global models (zoom) and ~ 0.17 – 0.56° (longitude) \times ~ 0.17 – 0.5° (latitude) for the regional models. Most models are driven by meteorological fields from the European Centre for Medium-Range Weather Forecasts (ECMWF) ERA-Interim reanalysis (Dee et al., 2011). In the case of STILT, the operational ECMWF analyses were used, while for NAME meteorological analyses of the Met Office Unified Model (UM) were employed. The regional models use boundary conditions (background CH₄ mole fractions) from inversions of the global models (STILT from TM3, COMET from TM5-4DVAR, CHIMERE from LMDZ) or estimate the boundary conditions in the inversions (NAME) using baseline observations at MaceHead as prior estimates. In the case of NAME and CHIMERE, the boundary conditions are further optimised in the inversion. The inverse modelling systems applied in this study use different inversion techniques. TM5-4DVAR, LMDZ, and TM3-STILT use 4DVAR variational techniques, which allow optimisation of emissions of individual grid cells. These 4DVAR techniques employ an adjoint model in order to iteratively minimise the cost function using a quasi-Newton (Gilbert and Lemaréchal, 1989) or conjugate gradient (Rödenbeck, 2005) algorithm. The NAME model applies a simulated annealing technique, a probabilistic technique for approximating the global minimum of the cost function. In CHIMERE and COMET, the inversions are performed analytically after reducing the number of parameters to be optimised by aggregating individual grid cells before the inversion. TM5-CTE applies an ensemble Kalman filter (EnKF) (Evensen, 2003), with a fixed-lag smoother (Peters et al., 2005).

Uncertainty: In general, the estimated model uncertainties depend on the type of station and for some models (TM5-4DVAR and NAME) also on the specific synoptic situation. In InGOS the uncertainty of the ensemble was calculated as 1σ estimate. Bergamaschi et al. (2015) showed that the range of the derived total CH₄ emissions from north-western and eastern Europe using four different inverse modelling systems was considerably larger than the uncertainty estimates of the individual models because the latter typically use Bayes' theorem to calculate the reduction of assumed a prior emission uncertainties by assimilating measurements (propagating estimated observation and model errors to the estimated emissions). An ensemble of inverse models may provide more realistic overall uncertainty estimates, since estimates of model errors are often based on strongly simplified assumptions and do not represent the total uncertainty.

InTEM – NAME

The Inverse Technique for Emission Modelling (InTEM) (Arnold et al., 2018) uses the NAME (Numerical Atmospheric dispersion Modelling Environment) (Jones et al, 2007) atmospheric Lagrangian transport model. NAME is driven by analysis 3-D meteorology from the UK Met Office Unified Model (Cullen, 1993). The horizontal and vertical resolution of the meteorology has improved over the modelled period from 40 km to 12 km (1.5 km over the UK). InTEM is a Bayesian system that minimises the mismatch between the model and the atmospheric observations

given the constraints imposed by the observation and model uncertainties and prior information with its associated uncertainties. The direction (latitude and longitude) and altitude varying background concentration and observation station bias are solved for within the inverse system along with the spatial distribution and magnitude of the emissions. The time-varying prior background concentration for the DECC network stations is derived from the MHD observations when they are very largely sensitive only to Northern Canada (Arnold et al., 2018). The prior bias (that can be positive or negative) for each station is set to zero with an uncertainty of 1 ppb. The observations from each station are assumed to have an exponentially decreasing 12-hr time correlation coefficient and, between stations, a 200 km spatial correlation coefficient. The observations are averaged into 2-hr periods. The uncertainty of the observations is derived from the variability of the observations within each 2-hr period. The modelling uncertainty for each 2-hr period at each station varies and is defined as the larger of; the median pollution events in that year at that station, or 16.5% of the magnitude of the pollution event. These values have been derived from analysis of the observations of methane at multiple heights at each station across the DECC network. Each inversion is repeated 24 times, each time 10% of the observations per year per station are randomly removed in 5-day intervals and the results and uncertainty averaged.

Uncertainty: This random removal of observations allows a greater exploration of the uncertainty, given the potential for some of the emission sources to be intermittent within the time-period of the inversion.

CTE-CH₄ Europe, CTE-SURF and CTE-GOSAT

CarbonTracker Europe CH₄ (CTE-CH₄) (Tsuruta et al., 2017) applies an ensemble Kalman filter (Peters et al. 2005) in combination with the Eulerian transport model TM5 (Krol et al. 2005). It optimizes surface fluxes weekly, and assimilates atmospheric CH₄ observations. TM5 was run at 1° x 1° resolution over Europe and 6° x 4° resolution globally, constrained by 3-hourly ECMWF ERA-Interim meteorological data. The photochemical sink of CH₄ due to tropospheric and stratospheric OH, and stratospheric Cl and O(¹D) was pre-calculated based on Houweling et al. (2014) and Brühl and Crutzen (1993) and not adjusted in the optimization scheme.

Three experiments were conducted, which differ in (1) sets of prior fluxes, (2) sets of assimilated observations, and (3) optimization resolution over the Northern Hemisphere.

In Petrescu et al., 2021a study, CTE-FMI uses-used sets of prior fluxes from LPX-Bern DYPTOP (Stocker et al., 2014) for biospheric, EDGAR v4.2 FT2010 (Janssens-Maenhout et al., 2013) for anthropogenic, GFED v4 (Giglio et al., 2013) for biomass-burning, Ito and Inatomi (2012) for termites and Tsuruta et al. (2017) for ocean sources. CTE-SURF and CTE-GOSAT use sets of prior fluxes from Global Carbon Project (Saunois et al., 2020). CTE-FMI and CTE-SURF assimilated ground-based surface CH₄ observations, while CTE-GOSAT assimilated GOSAT XCH₄ retrievals from NIES v2.72. CTE-FMI optimized fluxes at 1° x 1° resolution over Northern Europe, northeast Russia and southeast Canada, 6° x 4° resolution over other parts of the Northern Hemisphere land, and region-wise (combined TransCom regions and soil-type) over the Southern Hemisphere and ocean. CTE-SURF and CTE-GOSAT fluxes were optimized at 1° x 1° resolution over Europe and region-wise elsewhere globally.

For the current study, CTE-FMI S5 simulations updated priors as described in Table B2.4.

Uncertainty: The prior uncertainty is assumed to be a Gaussian probability distribution function, where the error covariance matrix includes errors in prior fluxes, observations and transport model representations. The uncertainty

for the prior fluxes were assumed to be 80 % of the fluxes over land and 20 % over ocean, with correlation between grid cells or regions to be 100-500 km over land and 900 km over ocean. The uncertainty for observations and transport model representations vary between observations, with min. aggregated uncertainty to be 7.5 ppb for surface observations and 15 ppb for GOSAT data. The posterior uncertainty is calculated as standard deviation of the ensemble members, where the posterior error covariance matrix is driven by the ensemble Kalman filter.

VERIFY Community Inversion Framework (CIF)

In the current study, new CIF results are presented (Berchet et al., 2021). For CH₄ inversions were run with CHIMERE, FLEXPART (NILU) and FLEXPART-EMPA. The results from CHIMERE differed somewhat from the two FLEXPART models in where the posterior fluxes departed from the prior estimates, but were reasonably comparable in terms of the annual emissions at regional scales. The models were run for EU27 plus Norway, Switzerland and UK, with annual mean emissions for 2005-2018.

CHIMERE (see <https://www.lmd.polytechnique.fr/chimere/>) is a non-hydrostatic Eulerian chemistry-transport model. Its area-limited domains can be designed to cover the hemispheric to the urban scales, with horizontal resolutions from several degrees to one kilometer. The time-steps usually cover a few minutes, depending on the CFL and choices made by the user for minimizing computation costs. For the purpose of flux inversions, the tangent-linear and the adjoint codes have been developed and parallelized, only for trace-gases (see (Fortems-Cheiney et al., 2021), their Section 3.2 for more details). The required input data are meteorological 3D and 2D fields (e.g. temperature, wind speed), boundary conditions for concentrations at the four sides and at the top of the domain and emission fluxes. The comparison to surface measurements is done by extracting from the model the simulated concentrations in the grid cell matching the stations' locations for the time-steps matching the measurement date and time. If the measurement covers a longer time (e.g. hourly means), the simulated concentrations for the matching time-steps are averaged. For this deliverable, CHIMERE was run using meteorological data from the ECMWF's IFS operational forecast (every three hours) retrieved at $0.25^\circ \times 0.25^\circ$ and interpolated onto the model's grid ($0.5^\circ \times 0.5^\circ$). CHIMERE extends from the surface to 200 hPa with 17 sigma-pressure levels.

For the inversions provided by NILU, FLEXPART-v10.4 was run using ECMWF's operational IFS forecast with backward trajectory lengths of 10 days for CH₄. The footprints were saved at $0.25^\circ \times 0.25^\circ$ for CH₄ inversions. For the inversions provided by EMPA, FLEXPART-v9.2 was run using ECMWF's ERA-Interim (up to 2018) and ERA5 for 2019 with backward trajectory lengths of 10 days and footprints were saved at $0.25^\circ \times 0.25^\circ$. For CH₄, prior anthropogenic emissions come from (EDGAR-v6) and for fluxes from natural soils (JSBACH-HIMMELI) and inland waters (CSLM-CH₄ model), and were supplemented by estimates from previous studies for geological, ocean and termite emissions (Etiope et al., 2019, Weber et al., 2019 and Saunio et al., 2020 (GCP-CH₄)). The EDGAR estimates were updated to include new activity data for 2016 to 2018, whereas in the previous version emissions for these years were based on a so-called fast-track estimation using proxies. The new estimate for lakes combines an empirical estimate, which provides only an annual climatology, with a monthly-resolved estimate from a mechanistic-stochastic modelling approach (MSM) accounting for nutrient (N and P) loads delivered to each lake and control of CH₄ emission by the lake trophic status.

Uncertainties: In the CIF, the uncertainties are described by matrices, with variances on the diagonal and covariances on the off-diagonal. They are called respectively the prior error (for the error statistics on the prior controlled variables) and the observation error (for the error statistics on the difference between the observed and modelled mixing ratios). Observations errors were assumed to be independent from each other in all inversions, i.e., the observation error covariance matrix is diagonal. For CHIMERE, measurement errors were combined with the transport, representation, and aggregation errors calculated for each station following (Szénási et al., 2021) using simulations based on the TNO emission inventory. For FLEXPART (NILU), the observations errors were taken from the observation files, but set to a minimum of 9 ppb. In addition, 50% of the contribution to from the fluxes outside the inversion domain (used in the calculation of the background mixing ratio) was added to the observation error. For FLEXPART-EMPA measurement errors were composed of an observation error and a model error, which are summed quadratically to obtain the total error. Observation errors were taken from the observation files and an average model error was determined for each station separately by an iterative procedure, following Koohkan and Bocquet (2012): An initial value was computed from the root-mean-square-differences between a priori simulated values and the observations. With this initial value, an inversion was performed. The model error was then scaled using the Eq. (20) of Koohkan and Bocquet (2012) until convergence.

MIROC4-ACTM:

The MIROC4-ACTM time dependent inversions solve for emissions from 53 regions for CH₄ and 84 regions for N₂O. The inversion framework is based on Bayesian statistics and optimizes surface-atmosphere fluxes using the maximum probability solution. Atmospheric transport is modelled using the JAMSTEC's Model for Interdisciplinary Research on climate, version 4 based atmospheric chemistry-transport model (MIROC4-ACTM) (Watanabe et al. 2008; Patra et al. 2018). The Source-Receptor Matrix (SRM) is calculated by simulating unitary emissions from 53 or 84 basis regions, for which the fluxes are optimised. The SRM describes the relationship between the change in mole fraction at the measurement locations for the unitary basis region fluxes. The MIROC4-ACTM meteorology was nudged to the JMA 55-year reanalysis (JRA55) horizontal wind fields and temperature. The calculation of photo-chemical losses is performed online. The hydroxyl (OH) radical concentration for reaction with CH₄ vary monthly but without any interannual variations. The simulated mole fractions for the total a priori fluxes are subtracted from the observed concentrations before running the inversion calculation (as in Patra et al., 2016 for CH₄ inversion). Both the inversion results are contributed to the GCP-CH₄ and GCP-N₂O activities (Saunois et al., 2020; Thompson et al., 2019; Tian et al., 2020).

Uncertainties: The posterior fluxes are subject to systematic errors primarily from: 1) errors in the modelled atmospheric transport; 2) aggregation errors, i.e. errors arising from the way the flux variables are discretized in space (84 regions) and time (monthly-means); 3) errors in the background mole fractions (assumed to be a minor factor); and 4) the incomplete information from the sparse observational network and hence the dependence on the prior fluxes. In addition, there is, to a much smaller extent, some error due to calibration offsets between observing instruments, which is more pertinent for N₂O than for other GHGs. We have validated model transport in troposphere using SF₆ for the inter-hemispheric exchange time, and the using SF₆ and CO₂ for the age of air in the stratosphere. The simulated

N₂O concentrations are also compared with aircraft measurements in the upper troposphere and lower stratosphere for evaluating the stratosphere-troposphere exchange rates. Comparisons with ACE-FTS vertical profiles in the stratosphere and mesosphere indicate good parameterisation of N₂O loss by photolysis and chemical reactions, and thus the lifetime, which affect the global total N₂O budgets. Random uncertainties are calculated by the inverse model depending on the prior flux uncertainties and the observational data density and data uncertainty. Only 37 sites are used in the inversion and thus the reduction in priori flux uncertainties have been minimal. The net fluxes from the inversion from individual basis regions are less reliable compared to the anomalies in the estimated fluxes over a period of time.

Global Carbon Project – Global Methane Budget (GMB)

GMB uses an ensemble of 22 top-down global inversions for anthropogenic CH₄ emissions presented in Saunio et al. (2020) for the Global Methane Budget. These inversions were simulated by nine atmospheric inversion systems based on various chemistry transport models, differing in vertical and horizontal resolutions, meteorological forcing, advection and convection schemes, and boundary layer mixing. Surface-based inversions were performed over the period 2000-2017 while satellite-based inversions cover the GOSAT data availability 2010-2017. The protocol established for these simulations was not stringent as the prior emission flux data set was not mandatory, and each group selected their constraining observations. More information can be found in Saunio et al. (2020) in particular in their Table 6 and S6.

Uncertainties: currently there are no uncertainties reported for the GMB models. This study uses the median and the min/max as uncertainty range estimation from the 22 models ensemble. In general uncertainties might be due to factors like: different transport models, physical parametrizations, prior fluxes, observation data sets etc.

A2: Natural CH₄ emissions

Bottom-up natural CH₄ emissions estimates

CH₄ emissions from inland waters

The CH₄ estimate from inland waters represents a climatology of diffusive and ebullitive CH₄ emissions from rivers, lakes and reservoirs. It is based on two approaches. The first approach synthesizes average annual CH₄ emissions fluxes for Europe that were rescaled to a consistent set of inland water surface area (Lauerwald et al., in rev. GBC) and corrected for the effect of seasonal ice cover. To obtain fluxes for the EU27+UK domain, the median and first interquartile range values are scaled down by a factor 0.75 based on the surface area of the two domains. The second approach provides a spatially-resolved climatology of inland water fluxes at the resolution of 0.1°. The river estimate relies on the 0.1°x0.1° river water surface area of Lauerwald et al. (2015) and 3 observation-based assessments (Bastviken et al., 2011; Stanley et al., 2016; Rosentreter et al., 2021) of mean CH₄ flux densities for European rivers. Note the very large range encompassed by the 3 studies (0.07-0.66 Tg CH₄ yr⁻¹ for EU27+UK), reflecting high uncertainty in the assessment of the river CH₄ flux. The lake estimate not only resolves the spatial variability, but also the temporal variability in CH₄ emissions. To do so, the mechanistic-stochastic-modeling (MSM)

approach of Maavara et al. (2017, 2019) and Lauerwald et al. (2019) was expanded to resolve the lake seasonal dynamics and the biogeochemical processes of the CH₄ and O₂ cycles occurring in the water column and sediments. To constrain the lake physics, the Canadian Small Lake Model (CSLM) was used (MacKay, 2012). CSLM represents lake stratification and mixing events by simulating vertical temperature profiles, thermocline and light penetration depths, and lake ice dynamics. For carbon, the MSM simulates a lake-mean trophic state from the balance between Net Primary Production (limited by light and nutrient inputs from the watershed) and heterotrophic decomposition of organic matter. It was upgraded to simulate vertical profiles of O₂ and CH₄ by accounting for eddy-diffusive transport and the set of consumption/production processes of the O₂-CH₄ cycles at the (sub)-daily resolution with a vertical resolution of less than one meter (Maisonnier et al., in prep.). In the sediment, net CH₄ production was split into diffusive and ebullitive pathways using an approach modified from Langenegger et al. (2019). The new process-based model was then applied to the European domain, using a lake clustering approach whereby within each grid of the simulation domain (2.5°x2.5°), lakes are binned into different classes as a function of the key drivers of CH₄ fluxes that are lake-size and depth (Messenger et al., 2016), and lake trophic status. Then, for each grid and each class, a representative simulation forced by high-resolution local climate forcings extracted from the lake sector of ISIMIP was performed. To carry out the spatial upscaling, the resulting diffusive and ebullitive areal CH₄ fluxes through the water-air interface were multiplied by the surface area of each lake class in each grid of the domain, extracted from the HydroLAKES database (Messenger et al., 2016).

JSBACH-HIMMELI

The model framework, JSBACH-HIMMELI (Raivonen et al., 2017; Susiluoto et al., 2018) is used to estimate wetland and mineral soil emissions, and an empirical model is used to estimate the emissions from inland water bodies.

JSBACH-HIMMELI is a combination of two models, JSBACH, that is the land-surface model of MPI-ESM (Reick et al., 2013), and HIMMELI, that is a specific model for northern peatland emissions of CH₄ (Raivonen et al., 2017). HIMMELI (Helsinki Model of Methane build-up and emission for peatlands) has been developed especially for estimating CH₄ production and transport in northern peatlands. It simulates both CH₄ and CO₂ fluxes and can be used as a module within different modelling environments (Raivonen et al., 2017; Susiluoto et al., 2018). HIMMELI is driven with soil temperature, water table depth, the leaf area index and anoxic respiration. These parameters are provided to HIMMELI from JSBACH, which models hydrology, vegetation and soil carbon input from litter and root exudates. CH₄ emission and uptake of mineral soils are calculated applying the method by Spahni et al. (2011) based on soil moisture estimated by JSBACH.

The distribution of terrestrial vegetation types in JSBACH-HIMMELI is adopted from CORINE land cover data and from native JSBACH land cover for the areas that CORINE does not cover. The HIMMELI methane model is applied for peatlands and the mineral soil approach for the rest. The map of inland water CH₄ emissions has been combined with JSBACH-HIMMELI land use map so that the map of inland waters is preserved and JSBACH grid-based fractions of different land use categories adjusted accordingly. In order to avoid double-counting the terrestrial CH₄ flux estimates have been normalized by the ratio of the two inland water body distributions.

Uncertainties: As in any process modeling the uncertainties of the bottom up modeling of CH₄ arise from three primary sources: parameters, forcing data (including spatial and temporal resolution), and model structure. An important source of uncertainty in the case of terrestrial CH₄ flux modeling is the spatial distribution of peatlands.

The uncertainties of JSBACH-HIMMELI peatland emissions were estimated by comparing the annual totals of measured and simulated methane fluxes at five European observation sites. Two of the sites are located in Finnish Lapland, one in middle Sweden, one in southern Finland and one in Poland.

For the sensitivity of mineral soil fluxes Spahni et al. (2011) tested two soil moisture thresholds, 85% or 95% of water holding capacity, below which mineral soils were assumed to be only CH₄ sinks, above which sources. We used the higher value, 95% of water holding capacity. The uncertainty was estimated using CH₄ flux simulations of one year (2005). We did two new model runs, using moisture thresholds 95±15%, and derived the uncertainty from the resulting range in the annual emission sum.

Geological fluxes

Framework and previous works

Geological methane emissions to the atmosphere, including natural gas seepage in petroliferous sedimentary basins and geothermal exhalations, were estimated at the global scale by multiple authors, based on bottom-up and top-down procedures (see review, including discussions on conflicting estimates, in Etiope and Schwietzke, 2019; Thornton et al. 2021 and references therein), and accounting for about 40-50 Tg CH₄ yr⁻¹.

For the European continent, a first geo-methane emission estimate was proposed by Simpson et al. (1999) with “a best guess” of 0.01 Tg yr⁻¹ and a speculative upper limit of 3 Tg yr⁻¹, only on the basis of an extrapolation of a few submarine emission data. At the time of the Simpson’s work, very few data on geological methane fluxes on land were available, and emission factors were basically unknown. Bottom-up emission estimates at European level including onshore seepage and geothermal exhalations were proposed ten years later, by Etiope (2009), on the basis of published regional emission estimates, suggesting around 3 Tg yr⁻¹ for geographic, onshore and offshore, Europe (including Azerbaijan; practically corresponding to present EU49).

Again ten years later, thanks to a wider data-set of CH₄ seepage flux from different geological environments in different countries and global inventories of geo-CH₄ emission sites, a process-based model using statistically derived emission factors and activity (areas) was developed to derive a global grid map of geo-CH₄ emissions (Etiope et al. 2019). The global grid model, developed by ArcGIS at 1°x1° resolution, can be, in theory, used to derive (scale-up) geo-CH₄ emissions at continental or regional scale.

This exercise was done, for Europe (EU27+UK) by Petrescu et al. (2021a), obtaining a value of 8.8 Tg yr⁻¹. The authors wished, however, to scale-down this value taking into account a global top-down estimate based on radiocarbon-free CH₄ in ice cores by Hmiel et al. (2020), who suggested, for the entire planet, 1.6 Tg yr⁻¹. This global estimate has more recently been contextualized and questioned by Thornton et al. (2021); in fact, the Hmiel et al. (2020) estimate has the same order of magnitude of emissions estimated by multiple authors for single local and regional seepage areas, so that the global emission must be considerable higher.

Petrescu et al. (2021) used however the upper limit of Hmiel et al. (2020), 5.4 Tg yr⁻¹, to scale-down the global gridded emission of Etiope et al. (2019), i.e. 37.4 Tg yr⁻¹, which is not the estimated global emission (43-50 Tg yr⁻¹, which included some factors that could not be considered in the grid model). From the 8.8 Tg yr⁻¹ (EU27 + UK), Petrescu et al. (2021) obtained then 1.3 Tg yr⁻¹ (for marine and land geological emissions) as follows:

$$8.8 \times 5.4/37.4 = 1.3 \text{ Tg CH}_4 \text{ yr}^{-1} \quad (1)$$

Besides the subjective use of the upper limit of Hmiel et al. (2020) and of the gridded (not the estimated global) value of Etiope et al. (2019), it is important to note that the initial gridded value derived for EU27+UK, 8.8 yr⁻¹, is affected by the relatively low precision, at the European scale, of the input data used for the global grid model. The global model was, in fact, developed on the basis of a global, large scale distribution of geological factors (for example the area of petroleum fields which determines the microseepage area), which lack the necessary precision for lower (continental and country) scale application. The main purpose of the global gridding was to offer a global spatial distribution of geo-CH₄ sources, with emission potential and methane isotopic values; it could not provide locally precise geo-CH₄ emissions because the datasets, developed for gridding purposes, was not complete and did not contain all the information necessary for improving previous estimates. A refinement of bottom-up estimates was possible only for mud volcanoes and microseepage because their gridding implied a more careful assessment of the spatial distribution and emission factors.

Re-assessment of geo-CH₄ emissions in Europe

For the current study, the global grid model of Etiope et al. (2019) using more detailed input data for Europe, with reference to the potential area of microseepage (activity) derivable by a more precise estimate of the continental oil-gas field area. The same microseepage emission factors statistically derived on global scale. For the other categories of geo-CH₄ sources (mud volcanoes, onshore seeps, submarine seepage and geothermal manifestations) European country masks from VERIFY (relative to EU27+UK and EU49) for the calculation of the onshore emission, and related EEZ (Exclusive Economic Zone) for the calculation of the sub-marine seepage were applied, to derive the global emission grid.

In the global microseepage emission model, the area where microseepage can potentially exist (named EMA - effective microseepage area) was estimated taking into account the distribution of microseepage observed by direct measurements in several oil-gas fields: statistical analysis of available data (summarized in Etiope et al. 2019) suggested that microseepage fluxes occur in about 57% of the petroleum field area (PFA). In theory, microseepage is expected also in the regions where seepage phenomena, manifested by macro-seeps (oil and gas seeps), exist. In Etiope et al. (2019) the global PFA was derived from Petrodata (Lujala et al. 2007), based on an oil-gas field digital map of USGS (Pawlewicz et al. 1997), which considered for each oil-gas field center point a buffer area of 30 km radius. The reason for this radius was not explained in Lujala et al. (2007).

In this work, the PFA of Europe (EU49) was derived by digitising all petroleum field center points from the USGS map (Pawlewicz et al. 1997). The microseepage CH₄ emission related to EMA is then derived by using the

same emission factors (four levels of microseepage; Table 2) established on global scale in Etiope et al. (2019). The results of the gridding, integrating activity (EMA) and emission factors, for EU49 are reported in Table A2.1.

Table A2.1: Results of microseepage gridding (0.05°x0.05°) for EU49

	<u>N. cell</u>	<u>Area (km²)</u>	<u>MS (t km⁻² yr⁻¹)</u>	<u>Total output (t yr⁻¹)</u>
<u>Gridded EMA</u>	<u>9457</u>	<u>199,703</u>		<u>2,985,570</u>
<u>Gridded Level 1</u>	<u>6843</u>	<u>143,307</u>	<u>0.474</u>	<u>67,927</u>
<u>Gridded Level 2</u>	<u>1899</u>	<u>40,094</u>	<u>11.366</u>	<u>455,708</u>
<u>Gridded Level 3</u>	<u>134</u>	<u>2959</u>	<u>40.15</u>	<u>118,804</u>
<u>Gridded Level 4</u>	<u>581</u>	<u>13,008</u>	<u>180.13</u>	<u>2,343,131</u>

MS: Microseepage emission factor statistically derived as described in Etiope et al. (2019).

The EMA and related microseepage emission for EU27+UK are derived by applying the related mask on EU49, resulting in:

$$\text{EU27+UK EMA} = 177,439 \text{ km}^2 \text{ and microseepage EU27+UK} = 2,161,060 \text{ t yr}^{-1}.$$

The total microseepage emission is quite sensitive to the activity (area); This estimate can be improved by increasing the number of measurements worldwide and related spatial (activity) and emission factor statistics.

For the other geological sources categories, i.e., onshore seeps (OS; including mud volcanoes), geothermal manifestations (GM) and submarine seepage (SS), the masks of the European territories (for EU27+UK and EU49, and the EEZ for the marine areas) on the global 1°x1° emission grid of Etiope et al. (2019) were applied. The result is summarised in Table A2.2.

Table A2.2: Gridded European geo-CH₄ emissions (t yr⁻¹).

	<u>Microseepage</u> <u>(MS)</u>	<u>Onshore seeps</u> <u>(OS)</u>	<u>Geothermal</u> <u>(GM)</u>	<u>Submarine</u> <u>Seeps (SS)</u>	<u>TOTAL</u>
<u>EU49</u>	<u>2,985,570</u>	<u>2,162,539</u>	<u>404,205</u>	<u>1,653,049</u>	<u>7,205,363</u>
<u>EU27+UK</u>	<u>2,183,733</u>	<u>69,618</u>	<u>206,705</u>	<u>863,368</u>	<u>3,323,424</u>

Table A2.3 compares the new results with previous European estimates (Etiope, 2009; Petrescu et al. 2021a).

Table A2.3: European geo-CH₄ emission estimates (Tg yr⁻¹)

	<u>Etiope</u> <u>(2009)</u>	<u>Petrescu et al. (2021a)</u>	<u>This work</u>

<u>Geographic Europe</u> <u>(onshore+ offshore, including Azerbaijan)</u>	<u>3</u>		
<u>EU27+UK onshore</u>		<u>8.8 (from global grid model)</u> <u>1.3 (scaled-down as Hmiel, 2020)</u>	<u>2.4</u>
<u>EU27+UK onshore + offshore</u>			<u>3.3</u>
<u>EU49 onshore</u>			<u>5.5</u>
<u>EU49 onshore + offshore</u>			<u>7.2</u>

The overall uncertainties of the spatial distribution of the geo-CH₄ sources and CH₄ emissions depend on individual uncertainties of the four categories of seepage, which are discussed in Etiope et al. (2019). Compared to the global grid model, the uncertainty of the microseepage at European scale by refining the activity (microseepage area) has been reduced.

The new EU27+UK geo-CH₄ emission estimate is lower than the one derived by Petrescu et al. (2021a) using the global gridding, but higher than the scaled-down value (eq. 1). The EU49 (onshore+offshore) emission is higher than, but of the same order of magnitude of, the preliminary, rough estimate of geographic Europe, which included Azerbaijan, by Etiope (2009). Onshore geo-CH₄ emissions occur mostly in Azerbaijan, Italy, Romania, which are actually the EU49 countries with major onshore oil-gas reserves and production (BP, 2020), and thus with greater natural seepage potential. Offshore emissions are dominated by the large estimates published for the UK shelf (Judd et al, 1997, revised by Tizzard, 2008). These estimates need to be verified and improved. Beyond the emission values, our gridding provides, however, the first detailed map of natural geological methane emission in Europe, which can be used for continental scale methane budget modelling.

Top-down natural CH₄ emissions estimates

Global Carbon Project - Global Methane Budget (Saunois et al., 2020)

For this study, none of the global inversions were updated.

GMB uses an ensemble of thirteen monthly gridded estimates of wetland emissions based on different land surface models as calculated for Saunois et al. (2020). Each model conducted a 30-year spin-up and then simulated net methane emissions from wetland ecosystems over 2000-2017. The models were forced by CRU-JRA reconstructed climate fields (Harris, 2019), and by the remote sensing-based wetland dynamical area dataset WAD2M (Wetland Area Dynamics for Methane Modeling). This data set provides monthly global areas over 2000-2017 based on a combination of microwave remote sensing data from Schroeder et al. (2015) and various regional inventory data sets.

More information is available in Saunio et al. (2020) and more details will be presented in a future publication led by Poulter et al., 2017 and colleagues.

1720 **Uncertainty:** As described by Saunio et al. (2020) uncertainties are reported as minimum and maximum values of the available studies, in brackets. They do not take into account the uncertainty of the individual estimates, but rather express the uncertainty as the range of available mean estimates, i.e., the standard error across measurements/methodologies considered.

1725

A3: Anthropogenic and natural N₂O emissions

Bottom-up N₂O emission estimates

1730 *UNFCCC NGHGI (2019), EDGAR v5.0 and CAPRI:* descriptions are found in Appendix A1.

ECOSSE

ECOSSE is a biogeochemical model that is based on the carbon model ROTH-C (Jenkinson and Rayner, 1977; Jenkinson et al. 1987; Coleman and Jenkinson, 1996) and the nitrogen-model SUNDIAL (Bradbury et al. 1993; Smith et al. 1996). All processes of the carbon and nitrogen dynamics are considered (Smith et al., 2010a,b). Additionally, in ECOSSE processes of minor relevance for mineral arable soils are implemented as well (e.g. methane emissions) to have a better representation of processes that are relevant for other soils (e.g. organic soils). ECOSSE can run in different modes and for different time steps. The two main modes are site specific and limited data. In the later version, basis assumptions/estimates for parameters can be provided by the model. This increases the uncertainty but makes ECOSSE a universal tool that can be applied for large scale simulations even if the data availability is limited. To increase the accuracy in the site-specific version of the model, detailed information about soil properties, plant input, nutrient application and management can be added as available.

During the decomposition process, material is exchanged between the SOM pools according to first order rate equations, characterised by a specific rate constant for each pool, and modified according to rate modifiers dependent on the temperature, moisture, crop cover and pH of the soil. The N content of the soil follows the decomposition of the SOM, with a stable C:N ratio defined for each pool at a given pH, and N being either mineralised or immobilised to maintain that ratio. Nitrogen released from decomposing SOM as ammonium (NH₄⁺) or added to the soil may be nitrified to nitrate (NO₃⁻).

For spatial simulations the model is implemented in a spatial model platform. This allows us to aggregate the input parameter for the needed resolution. ECOSSE is a one-dimensional model and the model platform provides the input data in a spatial distribution and aggregates the model outputs for further analysis. While climate data are interpolated, soil data are represented by the dominant soil type or by the proportional representation of the different soil types in the spatial simulation unit (this is in VERIFY a grid cell).

Uncertainties in ECOSSE arise from three primary sources: parameters, forcing data (including spatial and temporal resolution), and model structure.

DayCent

DayCent was designed to simulate soil C dynamics, nutrient flows (N, P, S) and trace gas fluxes (CO₂, CH₄, N₂O, NO_x, N₂) between soil, plants and the atmosphere at daily time-step. Submodels include soil water content and temperature by layer, plant production and allocation of net primary production (NPP), decomposition of litter and soil organic matter, mineralization of nutrients, N gas emissions from nitrification and denitrification, and CH₄ oxidation in non-saturated soils.

The DayCent modelling application at the EU level is a consolidated model framework running on LUCAS point (Orgiazzi, 2018) which was extensively explained in previous works (Lugato et al., 2017, 2018; Quemada et al., 2020) where a detailed description of numerical and geographical datasets and uncertainty estimations is reported.

Information directly derived from LUCAS (2009-2015) included the soil organic carbon content (SOC), particle size distribution and pH. Hydraulic properties and bulk density was also calculated with an empirically-derived pedotransfer. Management information was derived from official statistics (Eurostat, 2019) and included crop shares at NUTS2 level. The amount of mineral N was partitioned according to the regional crop rotations and agronomic crop requirements. Organic fertilization and irrigated areas were derived from the ‘Gridded Livestock of the World’ FAO dataset and the FAO-AQUASTAT product.

Meteorological data were downloaded from the E-OBS gridded dataset (<http://www.ecad.eu>) at 0.1° resolution. For the climatic projection, the gridded data from CORDEX database (<https://esgf-node.ipsl.upmc.fr/search/cordex-ipsl/>) were used. The average annual (2006-2010) atmospheric N deposition from the EMEP model (v4.5) were also implemented into the simulations. For current study, the results were updated and use the 2015-2019 mean.

Uncertainty: The starting year of the simulation was set in 2009 and projected in the future. The uncertainty analysis, based on the Montecarlo approach, was done running the model 52 times in each point and, contemporary, randomly sampling model inputs from probability density functions for: SOC pool partition, irrigation and both mineral and organic fertilization rates. The model outputs (including uncertainties) at point level were up-scaled regionally at 1 km resolution by a machine learning approach based on Random Forest regression.

N₂O emissions from inland waters

The N₂O estimate represents a climatology of average annual N₂O emissions from rivers, lakes, reservoirs and estuaries at the spatial resolution of 0.1°. Based on a spatially explicit representation of water bodies and point and non-point sources of N and P, this model quantifies the global scale spatial patterns in inland water N₂O emissions in a consistent manner at 0.5° resolution, which were then downscaled to 0.1° using the spatial distribution of European inland water bodies. The procedure to calculate the cascading loads of N and P delivered to each water body along the river–reservoir–estuary continuum and to topologically connect 1.4 million lakes (extracted from the HYDROLAKES database) is described in Maavara et al., 2019 and Lauerwald et al., 2019. The methodology to quantify N₂O emissions

is based on the application of a mechanistic stochastic model (MSM) to estimate inland water C-N-P cycling as well as N₂O production and emission generated by nitrification and denitrification. Using a Monte Carlo analysis, the MSM allows to generate relationships relating N processes and N₂O emissions to N and P loads and water residence time from the mechanistic model outputs, which are subsequently applied for the spatially resolved upscaling. For the estimation of N₂O emission, we ran two distinct model configurations relying on EFs scaling to denitrification and nitrification rates: one assuming that N₂O production equals N₂O emissions, the other taking into account the kinetic limitation on N₂O gas transfer and progressive N₂O reduction to N₂ during denitrification in water bodies with increasing residence time (Maavara et al., 2019). The model outputs from the two scenarios are used to constrain uncertainties in N₂O emission estimates. [For this study, the upscaled RECCAP2 estimates were used \(Lauerwald et](#)

[al., in rev. GBC\)](#)

GAINS

Specific sectors and abatement technologies in GAINS vary by the specific emitted compound, with source sector definition and emission factors largely following the IPCC methodology at the Tier 1 or Tier 2 level. GAINS includes in general all anthropogenic emissions to air, but does not cover emissions from forest fires, savannah burning and land use / land use change. Emissions are estimated for 174 countries/regions, with the possibility to aggregate to a global emission estimate, and spanning a timeframe from 1990 to 2050 in five-year intervals. Activity drivers for macroeconomic development, energy supply and demand, and agricultural activities are entered externally, GAINS extends with knowledge required to estimate “default” emissions (emissions occurring due to an economic activity without emission abatement) and emissions and costs of situations under emission control (Amann et al., 2001).

Emissions of nitrous oxide derive from energy, industry, agriculture, and waste. Land use change emissions are not included. In the energy sector, certain technologies implemented to improve air quality affect N₂O emission factors (like catalytic converters in vehicles), sometimes also negatively. That is also the case for non-selective catalytic reduction devices for NO_x abatement in power plants, or for fluidized bed combustion. Relevant industrial processes cover nitric acid and adipic acid, with other processes (glyoxal, if relevant, or caprolactam) included. Both processes allow for two different levels of abatement technologies, which both are relatively easily accessible and low cost. The use of N₂O in gaseous form, often as an anesthetic for medical purposes, is associated with population numbers and scaled by availability of hospital beds. Marked emission reductions (at low costs) as well as complete phase out of emissions (high costs) are implemented as technologies. Agricultural emissions in part derive from manure handling, where different management strategies have repercussions on emissions. The larger fraction of emission is from application of nitrogen compounds in different forms to grassland, crops and rice, with rice using a different emission factor. Application of manure and of mineral fertilizer in GAINS can be reduced by advanced computer technology such as automatic steering and variable rate application, or by agrochemistry (nitrification inhibitors). Costs of implementation are considered to depend on the size of a farm, hence farm size is an important parameter. In the waste sector, composting and wastewater treatment are considered relevant sources. For wastewater treatment, GAINS also considers a specific emission reduction option when optimizing processes towards N₂O

reduction (e.g. via favoring the anammox process). All details have been reported by Winiwarter et al. (2018) in their supplementary material.

Uncertainties: The same paper provides full information on the uncertainty of N₂O emissions in the GAINS model, which is a consequence of uncertainty provided in the activity data, in the emission factors, and in the actual structure of the respective management strategies that also include the share of abatement technology already implemented. Further parameters also described (on uncertainty of future projections and on costs) are not relevant here.

FAOSTAT

FAOSTAT: Statistics Division of the Food and Agricultural Organization of the United Nations, provides N₂O emissions from agriculture: <https://www.fao.org/faostat/en/#data/GT> and its sub-domains, as well as N₂O emissions from land use linked to biomass burning. The FAOSTAT emissions database is computed following Tier 1 IPCC 2006 Guidelines for National GHG Inventories (<http://www.ipcc-nggip.iges.or.jp/public/2006gl/index.html>). Country reports to FAO on crops, livestock and agriculture use of fertilizers are the source of activity data. Geospatial data are the source of AD for the estimates from cultivation of organic soils, biomass and peat fires. N₂O emissions are provided by country, regions and special groups, with global coverage, relative to the period 1961-present (with annual updates, currently ~~2017~~2019) and with projections for 2030 and 2050 for agriculture only, expressed in both CO₂e and N₂O by underlying agricultural and land use emission sub-domain and by aggregate (agriculture total, agriculture total plus energy, agricultural soils). The main N₂O emissions are reported for the following agricultural activities: manure management, synthetic fertilizers, manure applied to the soils, manure left in pasture, crop residues, cultivation of organic soils and burning crop residues. LULUCF emissions consist of N₂O associated with burning biomass and peat fires, as well as from the drainage of organic soils. [Comparison to the UNFCCC submissions is also provided.](#)

Uncertainties were computed by Tubiello et al., 2013 but are not available in the FAOSTAT database.

Top-down N₂O emission estimates

FLEXINVERT

The FlexInvert framework is based on Bayesian statistics and optimizes surface-atmosphere fluxes using the maximum probability solution (Rodgers 2000). Atmospheric transport is modelled using the Lagrangian model FLEXPART (Stohl et al. 2005; Pisso et al. 2019) run in the backwards time mode to generate a so-called Source-Receptor Matrix (SRM). The SRM describes the relationship between the change in mole fraction and the fluxes discretized in space and time (Seibert and Frank, 2004) and was calculated for 7 days prior to each observation. For use in the inversions, FLEXPART was driven using ECMWF Era Interim wind fields.

The state vector consisted of flux increments (i.e. offsets to the prior fluxes) discretized on an irregular grid based on the SRMs (Thompson et al. 2014). This grid has finer resolution (in this case the finest was 0.5°×0.5°) where the fluxes have a strong influence on the observations and coarser resolution where the influence is only weak (the coarsest was 2°×2°). The flux increments were solved at 2-weekly temporal resolution. The state vector also included scalars for the background mole fractions. The optimal (posterior) fluxes were found using the Conjugate Gradient method (e.g. Paige and Saunders, 1975).

The background mole fractions, i.e., the contribution to the modelled mole fractions that is not accounted for in the 7-day SRMs, was estimated by coupling the termination points of backwards trajectories (modelled using virtual particles) to initial fields of mole fractions from the optimized Eulerian model LMDz (i.e. the CAMS N₂O mole fraction product v18r1) following the method of Thompson et al. 2014.

Uncertainties: The posterior fluxes are subject to systematic errors primarily from: 1) errors in the modelled atmospheric transport; 2) aggregation errors, i.e. errors arising from the way the flux variables are discretized in space and time; 3) errors in the background mole fractions; and 4) the incomplete information from the observations and hence the dependence on the prior fluxes. In addition, there is, to a smaller extent, some error due to calibration offsets between observing instruments, which is more pertinent for N₂O than for other GHGs. Random uncertainties are calculated from a Monte Carlo ensemble of inversions following Chevallier et al. (2007) and uncertainties in the observation space were inflated to take into account the model representation errors.

VERIFY Community Inversion Framework (CIF)

In the current study, new CIF results are presented (Berchet et al., 2021). For N₂O, inversions were run with CHIMERE and FLEXPART (NILU). The results using the two models were very comparable at the level of EU27 plus Norway, Switzerland and UK, with annual mean emissions for 2005-2018.

CHIMERE (see <https://www.lmd.polytechnique.fr/chimere/>) is a non-hydrostatic Eulerian chemistry-transport model. Its area-limited domains can be designed to cover the hemispheric to the urban scales, with horizontal resolutions from several degrees to one kilometer. The time-steps usually cover a few minutes, depending on the CFL and choices made by the user for minimizing computation costs. For the purpose of flux inversions, the tangent-linear and the adjoint codes have been developed and parallelized, only for trace-gases (see (Fortems-Cheiney et al., 2021), their Section 3.2 for more details). The required input data are meteorological 3D and 2D fields (e.g. temperature, wind speed), boundary conditions for concentrations at the four sides and at the top of the domain and emission fluxes. The comparison to surface measurements is done by extracting from the model the simulated concentrations in the grid cell matching the stations' locations for the time-steps matching the measurement date and time. If the measurement covers a longer time (e.g. hourly means), the simulated concentrations for the matching time-steps are averaged. For this deliverable, CHIMERE was run using meteorological data from the ECMWF's IFS operational forecast (every three hours) retrieved at $0.25^\circ \times 0.25^\circ$ and interpolated onto the model's grid ($0.5^\circ \times 0.5^\circ$). CHIMERE extends from the surface to 200 hPa with 17 sigma-pressure levels.

For the inversions provided by NILU, FLEXPART-v10.4 was run using ECMWF's operational IFS forecast with backward trajectory lengths of 7 days for N₂O. The footprints were saved at $0.5^\circ \times 0.5^\circ$ for N₂O inversions. For N₂O anthropogenic emissions were taken from EDGAR-v6. An estimate for the natural N₂O flux was provided by the O-CN land surface model – a monthly climatology of the pre-industrial (or baseline) fluxes (Zaehle et al., 2011) and an estimate for the ocean N₂O flux was provided by the coupled ocean-biogeochemistry model, PlankTOM-v10 – also a monthly climatology (Buitenhuis et al., 2018).

Uncertainties: In the CIF, the uncertainties are described by matrices, with variances on the diagonal and covariances on the off-diagonal. They are called respectively the prior error (for the error statistics on the prior controlled variables)

and the observation error (for the error statistics on the difference between the observed and modelled mixing ratios). Observations errors were assumed to be independent from each other in all inversions, i.e., the observation error covariance matrix is diagonal. For CHIMERE, measurement errors were provided from observation files. Transport, representation and aggregation errors were deduced from the CH₄-based errors. For FLEXPART (NILU), the observations errors were calculated in the same way as for CH₄ but a minimum error of 0.3 ppb was used.

Global N₂O Budget – GCP (Tian et al., 2020)

CAMS-N₂O

Within the GCP 2019 results, N₂O fluxes are estimated using the atmospheric inversion framework, CAMS-N₂O. Atmospheric inversions use observations of atmospheric mixing ratios, in this case, of N₂O, and provide the fluxes that best explain the observations while at the same time being guided by a prior estimate of the fluxes. In other words, the fluxes are optimized to fit the observations within the limits of the prior and observation uncertainties. To produce the optimized (*a posteriori*) fluxes a number of steps are involved: first, the observations are pre-processed, second, a prior flux estimate is prepared, third mixing ratios are simulated using the prior fluxes and are used to estimate the model representation error, and fourth, the inversion is performed. In total 140 ground-based sites, ship and aircraft transects are included in the inversion. The term “site” refers to locations where there is a long-term record of observations and includes ground-based measurements, both from discrete samples (or “flasks”) and quasi-continuous sampling by in-situ instruments, as well as aircraft measurements. A prior estimate of the total N₂O flux with monthly resolution and inter-annually varying fluxes is prepared from a number of models and inventories. For the soil fluxes (including anthropogenic and natural) an estimate from the land surface model OCN-v1.2 is used, which is driven by observation-based climate data, N-fertilizer statistics and modelled N-deposition (Zaehle et al. 2011). For the ocean fluxes, an estimate from the ocean biogeochemistry model PlankTOM-v10.2 is used, which is a prognostic model (Buitenhuis et al. 2018). Atmospheric transport is modelled using an offline version of the Laboratoire de Meteorologie Dynamique model, LMDz5, which computes the evolution of atmospheric compounds using archived fields of winds, convection mass fluxes and planetary boundary layer (PBL) exchange coefficients that have been calculated using the online version nudged to ECMWF ERA interim winds.

CAMS-N₂O uses the Bayesian inversion method to find the optimal fluxes of N₂O given prior information about the fluxes and their uncertainty, and observations of atmospheric N₂O mole fractions. The method is the same as that used in Thompson et al. (2014). For this study, inversions were not updated.

Uncertainty: Uncertainties in CAMS-N₂O simulations pertain to observation space and to state space. Uncertainty in the observation space is calculated as the quadratic sum of the measurement and transport uncertainties. The measurement uncertainty is assumed to be 0.3 ppb (approximately 0.1%) based on the recommendations of data providers. The transport uncertainty includes estimates of uncertainties in advective transport (based on the method of Rödenbeck et al. (2003)) and from a lack of subgrid-scale variability (based on the method of Bergamaschi et al. (2010)). For the error in each land grid cell, the maximum magnitude of the flux in the cell of interest and its 8 neighbours is used, while for ocean grid cells the magnitude of the cell of interest only is used. Posterior flux uncertainties are calculated from a Monte Carlo ensemble of inversions, based on the method of Chevallier et al. (2005).

1940 ***TOMCAT-INVICAT***

TOMCAT-INVICAT (Wilson et al., 2014) is a variational inverse transport model, which is based on the global chemical transport model TOMCAT, and its adjoint. It uses a 4-D variational (4D-VAR) optimization framework based on Bayesian theory which seeks to minimize model-observation differences by altering surface fluxes, while allowing for prior knowledge of these fluxes to be retained. TOMCAT (Monks et al., 2017) is an offline chemical transport model, in which meteorological data is taken from ECMWF ERA-Interim reanalyses (Dee et al., (2011)). The model grid resolution, and therefore the optimised surface flux estimates, have a horizontal resolution of 5.6 x 5.6 degrees. The model has 60 vertical levels running from the surface to 0.1 hPa. For each individual year's fluxes, which are optimised on a monthly basis, 30 minimisation iterations are carried out. For this study, inversions were not updated.

1950 **Uncertainty:** Uncertainties in TOMCAT-INVICAT N₂O inversions are described as follows and further in Thompson et al., (2019). Uncertainty in the observations is calculated as the quadratic sum of the measurement and transport uncertainties. The measurement uncertainty for each observation is assumed to be 0.4 ppb. For the transport error for each observation is assumed to be the mean difference between the observation grid cell and its 8 neighbours. Prior flux errors are assumed to be 100% or the prior estimate, and are uncorrelated in space and time. Posterior flux

1955 uncertainties are not currently able to be calculated.

MIROC4-ACTM

The MIROC4-ACTM time dependent inversion for 84 regions (TDI84) framework is based on Bayesian statistics and optimizes surface-atmosphere fluxes using the maximum probability solution (Rodgers 2000). Atmospheric transport is modelled using the JAMSTEC's Model for Interdisciplinary Research on climate, version 4 based atmospheric chemistry-transport model (MIROC4-ACTM) (Watanabe et al. 2008; Patra et al. 2018, 2022). The Source-Receptor Matrix (SRM) is calculated by simulating unitary emissions from 84 basis regions, for which the fluxes are optimised. The SRM describes the relationship between the change in mole fraction at the measurement locations for the unitary basis region fluxes (similar to Rayner et al., 1997). The MIROC4-ACTM meteorology was nudged to the JMA 55-year reanalysis (JRA55) horizontal wind fields and temperature. The simulated mole fractions for the total a priori fluxes are subtracted from the observed concentrations before running the inversion calculation (as in Patra et al., 2016 for CH₄ inversion).

In this study, the simulation have been updated to 2019 (Patra et al., 2022).

Uncertainties: The posterior fluxes are subject to systematic errors primarily from: 1) errors in the modelled atmospheric transport; 2) aggregation errors, i.e. errors arising from the way the flux variables are discretized in space (84 regions) and time (monthly-means); 3) errors in the background mole fractions (assumed to be a minor factor); and 4) the incomplete information from the sparse observational network and hence the dependence on the prior fluxes. In addition, there is, to a much smaller extent, some error due to calibration offsets between observing instruments, which is more pertinent for N₂O than for other GHGs. We have validated model transport in the troposphere using SF₆ for the inter-hemispheric exchange time, and the using SF₆ and CO₂ for the age of air in the stratosphere. The

simulated N₂O concentrations are also compared with aircraft measurements in the upper troposphere and lower stratosphere for evaluating the stratosphere-troposphere exchange rates. Comparisons with ACE-FTS vertical profiles in the stratosphere and mesosphere indicate good parameterisation of N₂O loss by photolysis and chemical reactions, and thus the lifetime, which affect the global total N₂O budgets.

1980 Random uncertainties are calculated by the inverse model depending on the prior flux uncertainties and the observational data density and data uncertainty. Only 37 sites are used in the inversion and thus the reduction in priori flux uncertainties have been minimal. The net fluxes from the inversion from individual basis regions are less reliable compared to the anomalies in the estimated fluxes over a period of time.

1985 **B1: Overview figures**

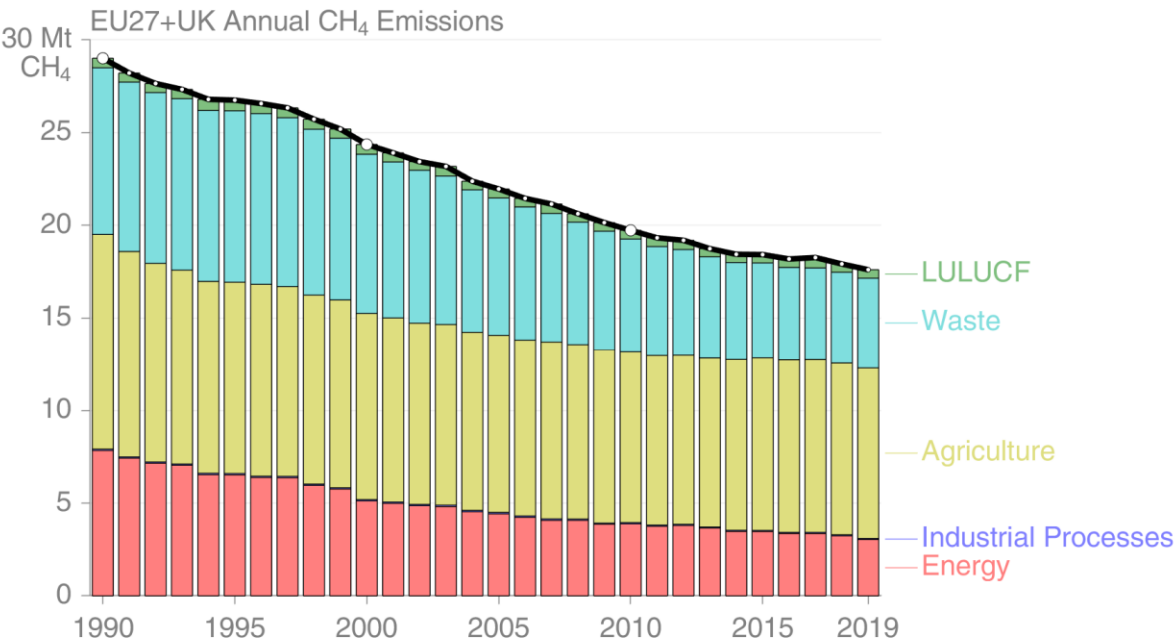


Figure [B1a](#): EU27+UK total CH₄ emissions time series per sectors as reported by UNFCCC NGHGI ([2021](#)~~2019~~).

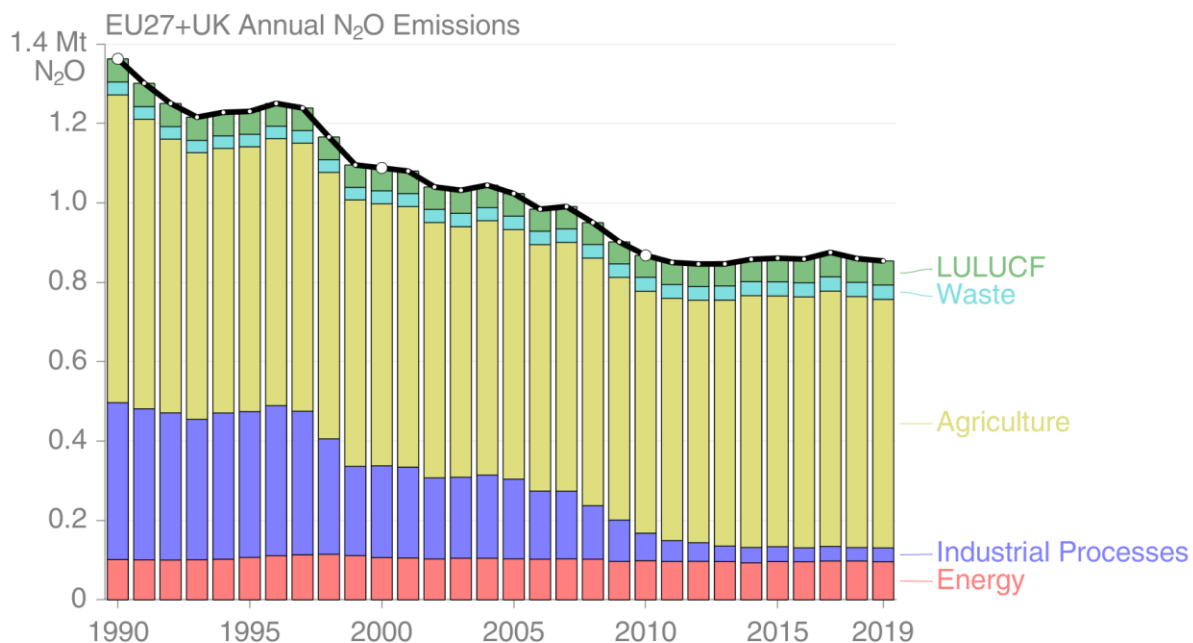


Figure B1b: EU27+UK total N_2O emissions time series per sectors as reported by UNFCCC NGHGI (2019-2021).

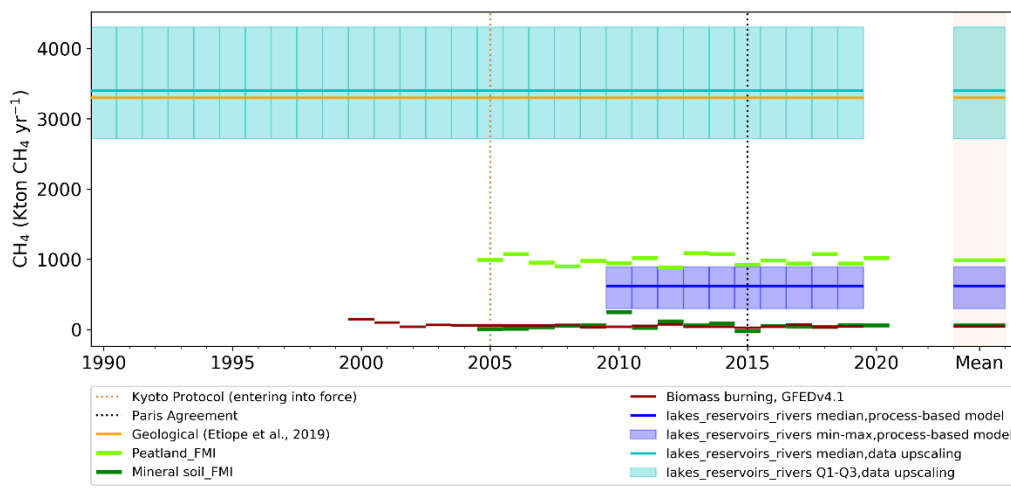


Figure B1c: EU27+UK natural CH_4 emissions from Figure 4b, as following: two estimates for inland waters (lakes rivers reservoirs process-based models, blue and upscaled emissions, cyan), peatlands and mineral soils (from JSBACH-HIMMELI, light green and dark green), geological emissions (orange) and biomass burning, (from GFEDv4.1, brown).

B2: Source specific methodology: AD, EF and uncertainties

Table B2.1: Source specific activity data (AD), emission factors (EF) and uncertainty methodology for all current VERIFY and non-VERIFY 2019-2021 data product collection.

CH ₄ bottom-up anthropogenic emissions				
Data source	AD/Tier	EFs/Tier	Uncertainty assessment method	Emission data availability
UNFCCC NGHGI (20192021)	Country-specific information consistent with the IPCC GLs.	IPCC GLs/country-specific information for higher tiers.	IPCC GLs (https://www.ipcc-nggip.iges.or.jp/public/2006gl/ , last access: December 2019) for calculating the uncertainty of emissions based on the uncertainty of AD and EF, two different approaches: (1) error propagation and (2) Monte Carlo simulation. <u>The EU GHG inventory team provided yearly harmonized and gap-filled uncertainties</u>	<u>NGHGI official data (CRFs) are found at https://unfccc.int/ghg-inventories-annex-i-parties/2021</u>
EDGAR v5v6.0	International Energy Agency (IEA) for fuel combustion Food and Agricultural Organisation (FAO) for agriculture US Geological Survey (USGS) for industrial processes (e.g. cement, lime, ammonia and ferroalloys) GGFR/NOAA for gas flaring World Steel Association for iron and steel production International Fertilisers Association (IFA) for urea consumption and production Complete description of the data sources can be found in	IPCC 2006, Tier 1 or Tier 2 depending on the sector	Tier 1 with error propagation by sectors for CH ₄	<u>https://edgar.jrc.ec.europa.eu/dataset_ghg60</u>

	Janssens-Maenhout et al. 2019 and in Crippa et al. (2019b).			
CAPRI	Farm and market balances, economic parameters, crop areas, livestock population and yields from EUROSTAT, parameters for input-demand functions at regional level from FADN (EC), data on trade between world regions from FAOSTAT, policy variables from OECD.	IPCC 2006: Tier 2 for emissions from enteric fermentation of cattle and from manure management of cattle. Tier 1 for all other livestock types and emission categories. N-flows through agricultural systems (including N excretion) calculated endogenously.	<u>Spatial uncertainties computed for 2014, 2016 and 2018</u>	Detailed gridded data CH ₄ and N ₂ O emissions can be obtained by contacting the data provider: Adrian.Leip@ec.europa.eu
GAINS	Livestock numbers by animal type (FAOSTAT, 2010; EUROSTAT, 2009; UNFCCC, 2010) Growth in livestock numbers from FAOSTAT (2003), CAPRI model (2009) Rice cultivation Land area for rice cultivation (FAOSTAT, 2010) Projections for EU are taken from the CAPRI Model	Country-specific information and: Livestock - Implied EFs reported to UNFCCC and IPCC Tier 1 (2006, Vol.4, Ch. 10) default factors Rice cultivation - IPCC Tier 1–2 (2006, Vol. 4, p. 5.49) Agricultural waste burning - IPCC Tier 1 (2006, Vol. 5, p. 520)	IPCC (2006, Vol.4, p.10.33) uncertainty range	Detailed gridded data CH ₄ and N ₂ O emissions can be obtained by contacting the data providers: for CH ₄ , contact Lena Höglund Isaksson (hoglund@iiasa.ac.at); for N ₂ O, contact Wilfried Winiwarter (winiwart@iiasa.ac.at).
FAOSTAT	FAOSTAT Crop and Livestock Production domains from country reporting; FAOSTAT Land Use Domain;	IPCC guidelines Tier 1	IPCC (2006, Vol.4, p.10.33) Uncertainties in estimates of GHG emissions are due to uncertainties in emission factors and activity data. They may be related to, inter	Agriculture total and subdomain specific GHG emissions are found for download at http://www.fao.org/faostat/en/#data/GT (last access: <u>April 2022</u>).

	Harmonized world soil; ESA CCI and Copernicus Global Land Cover Service (C3S) maps; MODIS MCD12Q1 v6; FAO Gridded Livestock of the World; MODIS MCD64A1.006 burned area products		alia, natural variability, partitioning fractions, lack of spatial or temporal coverage, or spatial aggregation.	
CH₄ bottom-up natural emissions				
Data source	AD/Tier	EFs/Tier	Uncertainty assessment method	Emission data availability
Mechanistic Stochastic Model CH₄ emissions from inland waters	Hydrosheds 15s (Lehner et al., 2008) and Hydro1K (USGS, 2000) for river network, HYDROLAKE S for lakes and reservoirs network and surface area (Messenger et al., 2016); Worldwide Typology of estuaries by Dürr et al. (2011)	N/A	Four model configurations for CH ₄	Detailed gridded data can be obtained by contacting the data providers: Ronny Lauerwald ronny.lauerwald@inrae.fr Pierre Regnier Pierre.Regnier@ulb.ac.be
JSBACH-HIMMELI	JSBACH vegetation and soil carbon and physical parameters provided to HIMMELI to simulate wetland methane fluxes HydroLAKES database (Messenger et al., 2016). CORINE land cover data VERIFY climate drivers $0.1^\circ \times 0.1^\circ$	CH ₄ fluxes from peatlands <u>and mineral soils</u>	the standard deviation and the resulting range in the annual emission sum represents a measure of uncertainty.	Detailed gridded data CH ₄ emissions can be obtained by contacting the data providers: Tuula.Aalto@fmi.fi tiina.markkanen@fmi.fi

Geological emissions, including marine and land geological)	Areal distribution activity: $1^\circ \times 1^\circ$ maps include the four main categories of natural geo-CH ₄ emission: (a) onshore hydrocarbon macro-seeps, including mud volcanoes, (b) submarine (offshore) seeps, (c) diffuse microseepage and (d) geothermal manifestations.	CH ₄ fluxes, measurements and estimates based on size and activity	95% confidence interval of the median emission-weighted mean sum of individual regional values	Etiope et al, 2019 with updated activity for current study) Detailed gridded data on geological CH₄ emissions can be obtained by contacting the data providers: Giuseppe Etiope: giuseppe.etiope@ingv.it Giancarlo Ciotoli: giancarlo.ciotoli@gmail.com
CH₄ Top-down inversions				
Regional inversions over Europe (high transport model resolution)				
Data source	AD/Tier	EFs/Tier	Uncertainty assessment method	Emission data availability
FLEXPART - FLExKF	Extended Kalman Filter in combination with backward Lagrangian transport simulations using the model FLEXPART Atmospheric observations ECMWF Era Interim meteorological fields	FLExKF- CAMSv19r _EMPA specific background	The random uncertainties are represented by the posterior error covariance matrix provided by the Kalman Filter, which combines errors in the prior fluxes with errors in the observations and model representation (see description in Appendix A1)	Detailed gridded data can be obtained by contacting the data provider: Dominik.Brunner@empa.ch
TM5-4DVAR	Global Eulerian models with a zoom over Europe, ERA-Interim reanalysis	4DVAR variational techniques	Uncertainty was calculated as 1σ estimate. See descriptions in Appendix A1	Detailed gridded data can be obtained by contacting the data provider: Peter.BERGAMASCHI@ext.ec.europa.eu
FLEXINVERT	Bayesian statistics Atmospheric transport is modelled using the Lagrangian model FLEXPART	prior fluxes from LPX-Bern DYPTOP, EDGAR v4.2 FT2010 GFED v4 Termites and ocean fluxes ground-based surface CH ₄ observations. Background fields based on nudged FLEXPART-CTM simulations (Groot Zwaafink et al., 2018)		Detailed gridded data CH ₄ emissions can be obtained by contacting the data provider: Christine Groot Zwaafink cgz@nilu.no

InTEM-NAME	Atmospheric Lagrangian transport model analysis 3-D meteorology from the UK Met Office Unified Model.	(a) the UK National Atmospheric Emissions Inventory (NAEI) 2015 within the UK. (b) Outside the UK – EDGAR 2010 emissions distributed uniformly over land (excluding the UK).	Derived from the variability of the observations within each 2 h period: a) 40 %; b) 50 %.	Detailed gridded data can be obtained by contacting the data provider: Alistair Manning (alistair.manning@metoffice.gov.uk).
CTE-FMI	Ensemble Kalman filter Eulerian transport model TM5 ECMWF ERA-Interim meteorological data	prior fluxes from LPX-Bern DYPTOP, EDGAR v4.2 FT2010 GFED v4 Termites and ocean fluxes ground-based surface CH ₄ observations GOSAT XCH ₄ retrievals from NIES v2.72	The prior uncertainty is assumed to be a Gaussian probability distribution function The posterior uncertainty is calculated as standard deviation of the ensemble members, where the posterior error covariance matrix are driven by the ensemble Kalman filter.	Detailed gridded data can be obtained by contacting the data provider: aki.tsuruta@fmi.fi
InGOS	18 European monitoring stations EDGARv4.2FT-InGOS wetland inventory of J.Kaplan and LPX-Bern v1.0 ERA-Interim reanalysis Met Office Unified Model	For Priors please see Table B2.4	The uncertainty of the model ensemble was calculated as 1σ estimate. Individual models use Bayes' theorem to calculate the reduction of assumed a priori emission uncertainties by assimilating measurements.	Detailed gridded data can be obtained by contacting the data provider: Peter.BERGAMASCHI@ext.ec.europa.eu
<u>VERIFY Community Inversion Framework (CIF): CHIMERE, FLEXPARTv10.4 (NILU) and FELXPART (EMPA) (only CH₄)</u>	<u>Extended Kalman Filter in combination with backward Lagrangian transport simulations using the model FLEXPART Atmospheric observations ECMWF Era Interim meteorological fields CHIMERE is a non-hydrostatic Eulerian chemistry-transport model</u>	<u>For Priors please see Table B2.4</u>	<u>The uncertainty in each grid cell (0.25°x0.25° for CH₄ and 0.5°x0.5° for N₂O) includes that due to the spatial disaggregation plus that due to emission-weighted uncertainty of a specific process.</u>	<u>Detailed gridded data can be obtained by contacting the data providers: Antoine Berchet antoine.berchet@lsce.ipsl.fr Brunner, Dominik Dominik.Brunner@empa.ch Rona Thompson rlt@nilu.no Gregoire Broquet gregoire.broquet@lsce.ipsl.fr</u>

Global inversions from the Global Carbon Project CH ₄ budget (Saunois et al. 2020)				
GCP-CH₄ 2019 anthropogenic and natural partitions from inversions	ensemble of inversions gathering various chemistry transport models surface or satellite data	For Priors please see Table B4	Uncertainties are reported as minimum and maximum values of the available studies, as the range of available mean estimates, i.e., the standard error across measurements/met hodologies considered. Posterior uncertainty mostly use Monte Carlo methods	Detailed gridded data can be obtained by contacting the data provider: Marielle Saunois marielle.saunois@lsce.ipsl.fr
N ₂ O bottom-up anthropogenic emissions				
Data source	AD/Tier	EFs/Tier	Uncertainty assessment method	Emission data availability
UNFCCC NGHGI (2019 2021), EDGAR v6.0, CAPRI, GAINS and FAOSTAT see above				
ECOSSE	The model is a point model, which provides spatial results by using spatial distributed input data (lateral fluxes are not considered). The model is a TIER 3 approach that is applied on grid map data, polygon organized input data or study sites.	IPCC 2006: Tier 3 The simulation results will be allocated due to the available information (size of spatial unit, representation of considered land use, etc.).	N/A	Detailed gridded data can be obtained by contacting the data provider: Kuhnert, Matthias matthias.kuhnert@abdn.ac.uk
DayCent	Spatial explicit simulations at point level, up-scaled at 1km for agricultural areas.	Tier 3; Land management and input factors for the cropland remaining cropland category based on datasets covering the 2005-2015 period.	Monte Carlo	Detailed gridded data can be obtained by contacting the data provider: Emanuele.LUGATO@ec.europa.eu
N ₂ O bottom-up natural emissions				
Mechanistic Stochastic Model for N₂O emissions from inland waters	Hydrosheds 15s (Lehner et al., 2008) and Hydro1K (USGS, 2000) for river network, HYDROLAKE S for lakes and reservoirs network and surface area	EFs applied to denitrification and nitrification rates for N ₂ O emissions. Values constrained from the range reported in Beaulieu et al., 2011.	<u>Upscaled emission estimates from RECCAP2</u>	Detailed gridded data can be obtained by contacting the data providers: Ronny Lauerwald ronny.lauerwald@inrae.fr Pierre Regnier Pierre.Regnier@ulb.ac.be

	(Messenger et al., 2016); Worldwide Typology of estuaries by Dürr et al. (2011); terrestrial N and P loads by Global-NEWS (Van Drecht et al., 2009; Bouwman et al., 2009), redistributed at 0.5° resolution by Maavara et al., 2019.			
Regional N₂O inversions over Europe (high transport model resolution)				
FLEXINVERT	Bayesian statistics Atmospheric transport is modelled using the Lagrangian model FLEXPART	background mole fractions	Random uncertainties are calculated from a Monte Carlo ensemble of inversions	Detailed gridded N ₂ O data can be obtained by contacting the data provider: Rona Thompson rlt@nilu.no
Global N₂O inversions over Europe from GN₂OB (Tian et al., 2020)				
CAMS-N₂O	Bayesian inversion method observations of atmospheric mixing ratios fluxes from ground-based sites, ship and aircraft transects soil fluxes OCN-v1.2 ocean biogeochemistry model PlankTOM-v10.2 GFED-v4.1s EDGAR-4.32 ECMWF ERA interim	Fires emission factors from Akagi et al., 2011	Uncertainty in the observation space is calculated as the quadratic sum of the measurement and transport uncertainties. For the error in each land grid cell, the maximum magnitude of the flux in the cell of interest and its 8 neighbours is used; for ocean grid cells the magnitude of the cell of interest only is used.	Detailed gridded N ₂ O data can be obtained by contacting the data provider: Rona Thompson rlt@nilu.no
TOMCAT-INVICAT	Variational Bayesian inverse model assimilating surface flask observations of atmospheric mixing ratios. ECMWF ERA-Interim	Prior emissions estimates are from OCN-v1.1 model (soils), EDGARv4.2FT2010 (anthro. non-soil), PlankTOM5 (oceans) and GFEDv4.1s (biomass burning).	Uncertainty in the observation space is calculated as the quadratic sum of the measurement and transport uncertainties. For the error in each land grid cell, the maximum	Detailed gridded N ₂ O data can be obtained by contacting the data provider: Christopher Wilson [GEO] C.Wilson@leeds.ac.uk

	meteorological driving data.		magnitude of the flux in the cell of interest and its 8 neighbours is used. Prior emission uncertainties are 100% and uncorrelated.	
MIROC4-ACTM	Matrix inversion for calculation of fluxes from 53 and 84 partitions of the globe for CH ₄ and N ₂ O, respectively. Forward model transport is nudged to JRA-55 horizontal winds and temperature.	Fire emissions for CH ₄ are taken from GFEDv4s	A posteriori uncertainties are obtained from the Bayesian statistics model. A priori emissions uncertainties are uncorrelated.	Detailed gridded data can be obtained by contacting the data provider: Prabir Patra prabir@jamstec.go.jp

2005 Table B2.2: Biogeochemical models that computed wetland emissions used in this study. Runs were performed for the whole period 2000-2017. Models run with prognostic (using their own calculation of wetland areas) and/or diagnostic (using WAD2M) wetland surface areas (see Sect 3.2.1) From Saunio et al., 2020.

Model	Institution	Prognostic	Diagnostic	References
CLASS-CTEM	Environment and Climate Change Canada	y	y	Arora, Melton and Plummer (2018) Melton and Arora (2016)
DLEM	Auburn University	n	y	Tian et al., (2010;2015)
ELM	Lawrence Berkeley National Laboratory	y	y	Riley et al. (2011)
JSBACH	MPI	n	y	Kleinen et al. (2019)
JULES	UKMO	y	y	Hayman et al. (2014)
LPJ GUESS	Lund University	n	y	McGuire et al. (2012)
LPJ MPI	MPI	n	y	Kleinen et al. (2012)
LPJ-WSL	NASA GSFC	y	y	Zhang et al. (2016b)
LPX-Bern	University of Bern	y	y	Spahni et al. (2011)

ORCHIDEE	LSCE	y	y	Ringeval et al. (2011)
TEM-MDM	Purdue University	n	y	Zhuang et al. (2004)
TRIPLEX_GHG	UQAM	n	y	Zhu et al., (2014;2015)
VISIT	NIES	y	y	Ito and Inatomi (2012)

2010 *Table B2.3: Top-down studies used in our new analysis, with their contribution to the decadal and yearly estimates noted. For decadal means, top down studies have to provide at least 8 years of data over the decade to contribute to the estimate, from Saunio et al., 2020*

Model	Institution	Observation used	Time period	Number of inversions	References
Carbon Tracker-Europe CH ₄	FMI	Surface stations	2000-2017	1	Tsuruta et al. (2017)
Carbon Tracker-Europe CH ₄	FMI	GOSAT NIES L2 v2.72	2010-2017	1	Tsuruta et al. (2017)
GELCA	NIES	Surface stations	2000-2015	1	Ishizawa et al. (2016)
LMDz-PYVAR	LSCE/CEA	Surface stations	2010-2016	2	Yin et al. (2019)
LMDz-PYVAR	LSCE/CEA	GOSAT Leicester v7.2	2010-2016	4	Yin et al. (2019)
LMDz-PYVAR	LSCE/CEA	GOSAT Leicester v7.2	2010-2017	2	Zheng et al. (2018a, 2018b)
MIROC4-ACTM	JAMSTEC	Surface stations	2000-2016	1	Patra et al. (2016; 2018)
NICAM-TM	NIES	Surface stations	2000-2017	1	Niwa et al. (2017a; 2017b)
NIES-TM-FLEXPART-VAR (NTFVAR)	NIES	Surface stations	2000-2017	1	Maksyutov et al. (2020); Wang et al. (2019b)
NIES-TM-FLEXPART-VAR (NTFVAR)	NIES	GOSAT NIES L2 v2.72	2010-2017	1	Maksyutov et al. (2020); Wang et al., (2019b)
TM5-CAMS	TNO/VU	Surface stations	2000-2017	1	Segers and Houweling (2018); Bergamaschi et al. (2010; 2013), Pandey et al. (2016)
TM5-CAMS	TNO/VU	GOSAT ESA/CCI	2010-2017	1	Segers and Houweling (2018,report); Bergamaschi

		v2.3.8 (combined with surface observations)			et al. (2010; 2013), Pandey et al. (2016)
TM5-4DVAR	EC-JRC	Surface stations	2000-2017	2	Bergamaschi et al. (2013, 2018)
TM5-4DVAR	EC-JRC	GOSAT OCPR v7.2 (combined with surface observations)	2010-2017	2	Bergamaschi et al. (2013, 2018)
TOMCAT	Uni. of Leeds	Surface stations	2003-2015	1	McNorton et al. (2018)

Table B2.4: List of prior datasets for natural CH₄ and N₂O emissions used by all inverse models

Project	Model	Prior						
		Wetlands	Geological	Fire	Termites	Soil sink	Ocean/Lakes	Wild animals
VERIFY	CTE_FMI <u>S5</u>	<u>JSBACH-</u> <u>HIMMELI LPX-</u> <u>Bern — DYPTOP</u> (Stocker et al., 2014)	<u>GCP_CH4</u> <u>Etiopie et al.,</u> <u>2019</u>	<u>RCO</u> plus <u>GFED4s.1 GFE</u> <u>D4s</u>	<u>Castaldi</u> as <u>GCP_CH4 Ito</u> <u>and Inatomi</u> <u>2012</u>	LPX-Bern DYPTOP (Stocker et al., 2014)	<u>Weber (in</u> <u>prep) for</u> <u>oceans and</u> <u>ULB for</u> <u>lakes in</u> <u>Europe, 0</u> <u>for the rest</u> <u>of the</u> <u>world Tsur</u> <u>uta et al.,</u> <u>2017</u>	
VERIFY	FLEXPART(F LExKF- <u>CAMSv19r)_E</u> MPA	JSBACH- HIMMELI		<u>GFED4s</u>	<u>GCP_2020</u>	Ridgwell /GCP	GCP/ULB	
VERIFY	FLEXINVERT	LPX-Bern DYPTOP (Stocker et al., 2014)		GFED4s	Ito and Inatomi, 2012	LPX-Bern DYPTOP (Stocker et al., 2014)	Tsuruta et al., 2017	
VERIFY	TM5_4DVAR JRC	GCP_CH4_2019	GCP_CH4 2019 (global total: 15 Tg CH ₄ yr ⁻¹)		GCP_CH4_2019	GCP_CH4_ 2019	GCP_CH4_ 2019	
<u>VERIFY</u> <u>-CIF</u>	<u>FLEXPART-</u> <u>NILU</u> <u>FLEXPART-</u> <u>EMPA</u>	<u>JSBACH-</u> <u>HIMMELI</u>	<u>Etiopie, 2019</u>	<u>EDGAR-v6</u> <u>(biofuel) and</u>	<u>Saunois et al.,</u> <u>2020 (GCP-</u> <u>CH4)</u>		<u>Weber et</u> <u>al. 2019</u>	

	<u>CHIMERE</u>			<u>GFED-v41s</u> <u>(biomass)</u>				
InGOS	INGOS-CTE-S4_EC	LPX-Bern v1.0 (Spahni et al., 2013)		GFED	Ito and Inatomi 2012	LPX-Bern v1.0 (Spahni et al., 2013)	Tsuruta et al., 2015	
InGOS	INGOS-LMDZEU-S4_EC	wetland inventory of J. Kaplan (Bergamaschi et al., 2007)						
InGOS	INGOS-TM3STILT-S4_EC	wetland inventory of J. Kaplan (Bergamaschi et al., 2007)						
InGOS	INGOS-TM5VAR-S4_EC	wetland inventory of J. Kaplan (Bergamaschi et al., 2007)			Sanderson /GCP	Ridgwell /GCP	Lambert /GCP	Osison climatology
InGOS	INGOS-NAME-S4_EC	wetland inventory of J. Kaplan (Bergamaschi et al., 2007)						
GCP	GELCA-SURF_NIES	VISIT (Ito and Inatomi, 2012)	n/a	GFEDv3.1 then GFAS v1.2 after 2011	Sanderson (TransCom-CH ₄ / GCP)	VISIT (Ito and Inatomi, 2012)	n/a	
GCP	MIROCv4-SURF_JAMA STEC	VISIT (Ito and Inatomi, 2012) (global total range : 173-197 Tg CH ₄ yr ⁻¹)	Etiopie and Milkov, 2004 (global total: 7.5 Tg CH ₄ yr ⁻¹)	GFEDv4s (global total range : 14-35 Tg CH ₄ yr ⁻¹)	Sanderson (TransCom-CH ₄) (global total: 20.5 Tg CH ₄ yr ⁻¹)	VISIT (Ito and Inatomi, 2012)	Lambert/Houweling (TransCom-CH ₄) (global total: 18.5 Tg CH ₄ yr ⁻¹)	
GCP	NICAM-SURF_NIES	VISIT (Ito and Inatomi, 2012)	GCP based on Etiopie 2015	GFEDv4s / GCP	Sanderson (TransCom-CH ₄ / GCP)	VISIT (Ito and Inatomi, 2012)	Lambert/Houweling (TransCom-CH ₄ / GCP)	

GCP	TOMCAT-SURF_ECMWF	JULES emissions from Mc Norton 2016a	Tomcat 2006	GFED V4	Matthews and Fung 2006	Patra et al. 2011	Tomcat 2006 Matthews and Fung 1987 - all emissions total rescaled to Schwietzke et al. 2016	
GCP	NTFVAR-GOSAT_NIES	VISIT (Ito and Inatomi, 2012)	Etiopie and Milkov, 2004	GFAS v1.2	Ito and Inatomi 2012	VISIT (Ito and Inatomi, 2012)	TransCom-CH4	
GCP	NTFVAR-SURF_NIES	VISIT (Ito and Inatomi, 2012)	Etiopie and Milkov, 2004	GFAS v1.2	Ito and Inatomi 2012	VISIT (Ito and Inatomi, 2012)	TransCom-CH4	
GCP	LMDZ-GOSAT1_LSC E	Bloom 2017	n/a	GFED V41s	Sanderson /GCP	Ridgwell /GCP	Lambert /GCP	
GCP	LMDZ-GOSAT2_LSC E	GCP - ensemble mean ESSD Saunois et al . 2016	GCP based on Etiopie 2015	GFED V41s	Sanderson /GCP	Ridgwell /GCP	Lambert /GCP	
GCP	LMDZ-GOSAT1 LMDZ-GOSAT2 LMDZ-GOSAT3 LMDZ-GOSAT4 LMDZ-SURF1 LMDZ-SURF2	Kaplan 2002 rescaled by Bergamaschi 2007	n/a	GFED V41	Sanderson 1996 /GCP	Ridgwell /GCP	Lambert and Schmidt 1993	
GCP	TM5-CAMS-GOSAT_TNO	Kaplan climatology	n/a	GFED V31 climatology after 2011	Sanderson /GCP	Ridgwell /GCP	Lambert /GCP	Osmon climatolgy
GCP	TM5-GOSAT1_EC	WETCHIMP ensemble mean;	GCP_CH4 2019 (global total: 15 Tg CH4 yr ⁻¹)		Sanderson /GCP	Ridgwell /GCP	Lambert /GCP	Osmon climatolgy
GCP	TM5-GOSAT2_EC	GCP_CH4_2019	GCP_CH4 2019 (global	GCP_CH4_2019	GCP_CH4_2019	GCP_CH4_2019	GCP_CH4_2019	

			total: 15 Tg CH ₄ yr ⁻¹)					
GCP	TM5-SURF1_EC	WETCHIMP ensemble mean;	GCP_CH4 2019 (global total: 15 Tg CH ₄ yr ⁻¹)		Sanderson /GCP	Ridgwell /GCP	Lambert /GCP	Osloon climatology
GCP	TM5-SURF2_EC	GCP_CH4_2019	GCP_CH4 2019 (global total: 15 Tg CH ₄ yr ⁻¹)	GCP_CH4_2019	GCP_CH4_2019	GCP_CH4_2019	GCP_CH4_2019	
GCP	CTE-GOSAT_FMI	GCP_CH4_2019	Etiopie 2015	GCP_CH4_2019 (=GFED4s)	GCP_CH4_2019	GCP_CH4_2019	GCP_CH4_2019	
GCP	CTE-SURF_FMI	GCP_CH4_2019	Etiopie 2015	GCP_CH4_2019 (=GFED4s)	GCP_CH4_2019	GCP_CH4_2019	GCP_CH4_2019	
	NAME-SURF_MetOff ice							
N₂O VERIFY CIF	FLEXPART-NILU CHIMERE			GFED-v41s		O-CN	PlankTom-v10	

2015

Author contributions:

AMRP designed this research and led the discussions with all co-authors, AMRP wrote the initial draft of the paper and edited all the following versions; CQ made the figures ~~and helped with the review process~~, PPe, MJM, V B processed the original data submitted to the VERIFY portal, MJM, PPe and PB designed and are managing the VERIFY web portal, BM provided the new gap-filled yearly UNFCCC NGHGI Member States uncertainty calculations and checked language and content of the final [reviewed](#) version, GPP provided the figures [B1a and B1b](#) in Appendix B1 and checked language and content of the final version, [AB provided the figures 5 and 14](#), [DBr provided the figure 7](#), AT provided the data for the ~~_gridded~~ figures [8 and 9](#) in section 3.1.5. and AMRP made the figures [8 and 9](#) ~~from section 3.1.5.~~, [AT and CQ provided input to the review answers](#), PPa, OT, RLT, CQ, GE, MK, LH-J, PR, RL, DBa, WW, provided in depth advice and commented/edited the initial versions of the manuscript, MB, RLT, AT, DBr, MK, PGE, GC, PR, RL, LH-J, WW, TA, AB, GC, MC, FD, CDGZ, AL, EL, AJM, MMa, TM, GDO, PKP, MS, AJJ, ES, HT, FNT, SZ. are data providers.

Competing interests

The authors declare that they have no conflict of interest.

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