

1 **Quantifying nitrous oxide emissions in the U.S.**
2 **Midwest - A top-down study using high resolution**
3 **airborne in situ observations**

4 **Maximilian Eckl¹, Anke Roiger¹, Julian Kostinek¹, Alina Fiehn¹, Heidi**
5 **Huntrieser¹, Christoph Knote², Zachary R. Barkley³, Stephen M. Ogle⁴,**
6 **Bianca C. Baier^{5,6}, Colm Sweeney⁶, and Kenneth J. Davis^{3,7}**

7 ¹Deutsches Zentrum für Luft- und Raumfahrt (DLR), Institut für Physik der Atmosphäre,
8 Oberpfaffenhofen, Germany

9 ²Ludwig-Maximilians-University (LMU), Meteorological Institute, Munich, Germany

10 ³Department of Meteorology and Atmospheric Science, Pennsylvania State University, University Park,
11 PA, USA

12 ⁴Natural Resource Ecology Laboratory, Colorado State University, Fort Collins, CO, USA

13 ⁵Cooperative Institute for Research in Environmental Sciences, University of Colorado-Boulder, Boulder,
14 CO, USA

15 ⁶NOAA Global Monitoring Laboratory, Boulder, CO, USA

16 ⁷Earth and Environmental Systems Institute, Pennsylvania State University, University Park, PA, USA

17 **Key Points:**

- 18 • Within the ACT-America project we gathered a unique airborne in situ N₂O data
19 set over the U.S. Midwest with enhancements up to 9 ppb
20 • N₂O emissions in the U.S. Midwest were on average $0.42 \pm 0.28 \text{ nmol m}^{-2} \text{ s}^{-1}$ in
21 Oct 2017 and $1.06 \pm 0.57 \text{ nmol m}^{-2} \text{ s}^{-1}$ in Jun-Jul 2019
22 • Bottom-up estimates from EDGAR and the often four times higher DayCent un-
23 derestimate U.S. Midwest N₂O emissions by factors up to 20

Corresponding author: Maximilian Eckl, Maximilian.Eckl@dlr.de

Abstract

The U.S. Midwest, with its intensive agriculture, is a prominent source of nitrous oxide (N_2O) but top-down and bottom-up N_2O emission estimates differ significantly. We quantify Midwest N_2O emissions by combining observations from the Atmospheric Carbon and Transport-America campaign with model simulations to scale the Emissions Database for Global Atmospheric Research (EDGAR). In October 2017 we increased agricultural EDGAR version 4.3.2/5.0 emissions by a factor of $6.3 \pm 4.6 / 3.5 \pm 2.7$, resulting in Midwest N_2O emissions of $0.42 \pm 0.28 \text{ nmol m}^{-2} \text{ s}^{-1}$. In June/July 2019, a period when extreme flooding was occurring in the Midwest, EDGAR was increased by a factor of $11.4 \pm 6.6 / 9.9 \pm 5.7$, resulting in N_2O emissions of $1.06 \pm 0.57 \text{ nmol m}^{-2} \text{ s}^{-1}$. Agricultural emissions estimated with the process-based model DayCent (Daily version of the CENTURY ecosystem model) were larger than in EDGAR but still substantially smaller than our estimates. Due to the complexity of N_2O emissions, further studies are necessary to fully characterize Midwest emissions.

Plain Language Summary

Nitrous oxide (N_2O) is the third most important anthropogenic greenhouse gas contributing to the warming of the planet and the dominant man-made ozone-depleting substance in the stratosphere. Its atmospheric concentrations have been rising since industrialization mainly due to an increase in anthropogenic sources, with agriculture being the dominant source. The densely farmed U.S. Midwest plays an important role in the global N_2O budget. However, previous studies that have collected observations of N_2O indicate that estimates of surface emissions in the Midwest are substantially underestimating the truth. In this study we combine unique aircraft-based N_2O measurements and model simulations to quantify Midwest emissions in October 2017 and June/July 2019. Agricultural inventory estimates had to be increased by factors up to 20 to match observations, revealing a large underestimation in current inventories. An extreme flooding event in 2019 when the summer observations occurred may be responsible for some of this discrepancy. Estimations of soil N_2O emissions calculated with a state-of-the-art biogeochemical model show less underestimation but are still too low compared to the fluxes derived from the aircraft observational data.

54 1 Introduction

55 Nitrous Oxide (N₂O) is the third most important anthropogenic greenhouse gas
56 (GHG) in terms of long-term radiative forcing (Myhre et al., 2013) and is the dominant
57 ozone depleting substance in the stratosphere (Ravishankara et al., 2009). Global N₂O
58 concentrations are 333 ppb as of April 2020, approximately a 20 % increase since prein-
59 dustrial times (MacFarling Meure et al., 2006; NOAA-ESRL, 2020). Anthropogenic sources
60 like agriculture and fossil fuel combustion contribute to this trend (Ciais et al., 2013).
61 In recent years, those N₂O emissions have increased at a higher rate than expected (Thompson
62 et al., 2019; Tian et al., 2020). Agricultural soil management associated with reactive
63 forms of nitrogen (N) (i.e. mineral fertilizer, livestock manure additions, and legumes)
64 accounts for half of global N₂O emissions (Paustian et al., 2016). Analyses of the iso-
65 topic composition of N₂O indicate that the observed rise in global atmospheric N₂O con-
66 centrations is mainly caused by the increased application of N-fertilizers (Park et al., 2012).

67 Bottom-up estimates, such as the Emissions Database for Global Atmospheric Re-
68 search (EDGAR, 2020), use emission factors and activity data to calculate emissions.
69 However, the nature of N₂O soil emissions complicates their quantification. Agricultural
70 practices (e.g. fertilizer application rate, crop type) as well as meteorological and soil con-
71 ditions (e.g. precipitation, soil moisture) directly influence emissions, resulting in large
72 temporal variability in N₂O surface fluxes (Stehfest & Bouwman, 2006). Process-based
73 biogeochemical models like DayCent (Daily version of the CENTURY ecosystem model)
74 provide a more sophisticated approach for estimation of N₂O emission by simulating soil
75 processes based on various environmental drivers. Nevertheless, fluxes at regional scale
76 are still highly uncertain due to insufficient direct observations (Reay et al., 2012).

77 The U.S. Midwest is one of the most intensively cultivated agricultural regions world-
78 wide (FAO, 2020; USDA-NASS, 2020), thus contributing significantly to the global an-
79 thropogenic N₂O emissions (Miller et al., 2012). Previous top-down studies indicate that
80 emissions in the Midwest are underestimated by EDGAR, but are highly uncertain on
81 the magnitude of this underestimation (Kort et al., 2008; Miller et al., 2012; Griffis et
82 al., 2013; Chen et al., 2016; Fu et al., 2017). Kort et al. (2008) showed that EDGAR ver-
83 sion 32FT2000 underestimates emissions in May-June 2003 by a factor of 2.62 over the
84 central U.S. and southern Canada. Miller et al. (2012) derived scaling factors of 6.1 and
85 10.1 for EDGAR version 4 for June 2004 and June 2008, respectively. Fu et al. (2017)
86 concluded even higher scaling factors for agricultural EDGAR version 4.2 emissions in
87 the Corn Belt region of the Midwest, with scaling factors of 19.0-28.1 in June 2010. These
88 described top-down studies used tall tower measurements, characterized by long time se-
89 ries over several months but limited in their spatial coverage. Only Kort et al. (2008)
90 used aircraft-based flask measurements, which provide some spatial (central U.S. and south-
91 ern Canada) but limited temporal (May-June 2003) coverage. The large range in the quan-
92 titative results show that Midwest N₂O surface fluxes are underestimated by EDGAR
93 inventories, but their true values are highly uncertain.

94 In this study we quantify N₂O emissions for several flights conducted in parts of
95 the U.S. Midwest in October 2017 and June/July 2019 with a top-down approach. Un-
96 like previous studies which have relied on observations with limited spatial coverage, this
97 study uses continuous airborne in situ measurements of N₂O. By combining these ob-
98 servations with forward model simulations, we optimize agricultural fluxes from EDGAR
99 version 4.3.2 and version 5.0 to quantify Midwest N₂O emissions. The employed method
100 was already successfully applied in several methane top-down studies (Barkley et al., 2017;
101 Barkley, Davis, et al., 2019; Barkley, Lauvaux, et al., 2019). The derived emission rates
102 are finally compared to flux estimates of direct soil emissions produced with EDGAR
103 and the biogeochemical model DayCent (Parton et al., 1998; Del Grosso et al., 2001, 2011).

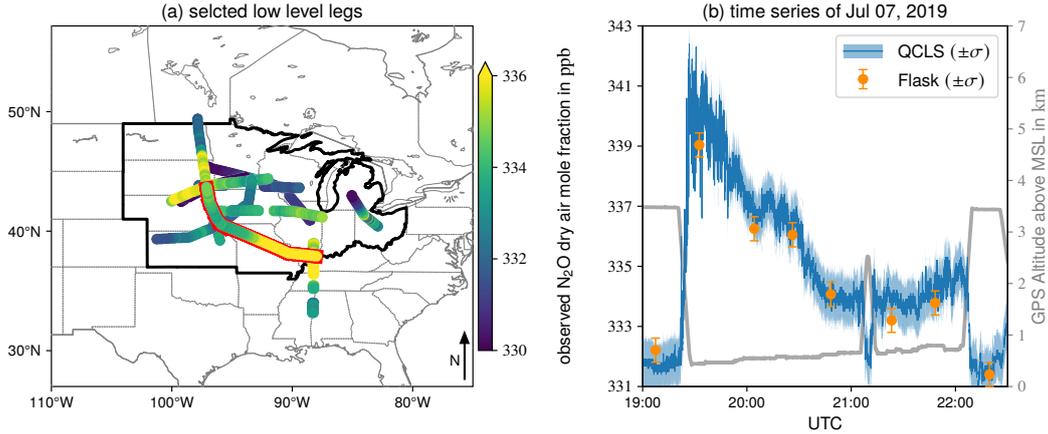


Figure 1. (a) Selected low level legs (at approx. 1000 ft AGL) of the ACT-America campaigns in 2017 and 2019, color-coded with observed N_2O dry air mole fractions. The study region (Midwest) is encircled by a thick black line. (b) Time series of N_2O dry air mole fraction of the flight on July 07, 2019 with error bars indicating ± 0.8 ppb and coincident NOAA/GML flask measurements of N_2O (± 0.4 ppb). The corresponding transect in (a) is encircled in red.

2 Data and Methods

2.1 Observational Data

We use measurements from the Atmospheric Carbon and Transport-America (ACT-America, <https://act-america.larc.nasa.gov/>) campaign. ACT-America includes five airborne campaigns from 2016 to 2019, providing a rich data set of in situ and remote greenhouse gas measurements in all four seasons. During the fall 2017 (10 Oct - 13 Nov) and summer 2019 (17 Jun - 27 Jul) field deployments, we collected approximately 60 h of in situ data onboard NASA's C-130 with an Aerodyne Quantum Cascade Laser Spectrometer (QCLS) measuring N_2O mole fractions (among others) at 2 Hz with an uncertainty of 0.8 ppb (Kostinek et al., 2019). Every 3-10 minutes in-flight calibrations were performed using standards that were cross-calibrated after the campaign against NOAA/GML standards traceable to the NOAA-2006A scale (Hall et al., 2007). Additionally, during each flight 6-12 whole-air flask samples were taken by NOAA/GML and measured for trace gases including N_2O with an uncertainty of 0.4 ppb (Sweeney et al., 2015, 2018; Baier et al., 2020). Those were merged into the QCLS time series to fill any data gaps.

For this study we selected four flights from 2017 (October) and six flights from 2019 (June/July). For each flight the C-130 flew low level legs well within the planetary boundary layer (PBL) (~ 1000 ft above ground level (AGL)) for at least 45 min during which Midwest air was sampled. Figure 1a shows the selected transects, color-coded with observed N_2O dry air mole fractions. These flights cover most parts of the Midwest. Mole fractions up to 341 ppb were observed (Figure 1b). We are not aware of comparable continuous N_2O measurements spanning most of Midwest across two seasons, highlighting the unique opportunity to quantify Midwest emissions with these data.

2.2 Model Setup

The Weather Research and Forecasting model with chemistry enabled version 4.0.2 (WRF-Chem; Grell et al. (2005)) is used to propagate N_2O enhancements emitted from emission inventories (Section 2.3) throughout the atmosphere. Initial N_2O concentra-

131 tions and the inflow at the boundaries of the model domain are set to zero. Thus, we
 132 simulate only enhancements caused by emissions within the model domain. We treat N₂O
 133 as a passive tracer due to its long atmospheric lifetime of ~ 116 years (Prather et al.,
 134 2015). The model domain consists of an outer and inner domain with a horizontal res-
 135 olution of $15\text{ km} \times 15\text{ km}$ and $3\text{ km} \times 3\text{ km}$, respectively. The outer domain, centered
 136 over the Midwest, covers nearly the whole continental U.S., northern Mexico, and south-
 137 ern Canada (Figure 2a), whereas the extension and position of the inner domain is sep-
 138 arately chosen for each flight so that the low level legs are spaciouly encapsulated. We
 139 perform each simulation with three different meteorological initial and boundary con-
 140 ditions: The 5th generation atmospheric reanalysis data (ERA5, 2017; Hersbach et al.,
 141 2020), the North American Regional Reanalysis (NARR, 2005), and the Global Data As-
 142 sssimilation System Final analysis (GDAS-FNL, 2015). As in Barkley, Davis, et al. (2019),
 143 we use these different simulations to estimate model transport errors (Díaz-Isaac et al.,
 144 2018). See the supporting information (SI) for additional information about the model
 145 setup.

146 2.3 Emission Inventories

147 The prior N₂O surface emission estimates for the optimization were obtained from
 148 EDGAR version 4.3.2 (EDGAR4.3.2, 2017; Janssens-Maenhout et al., 2019) and version
 149 5.0 (EDGAR5.0, 2019; Crippa et al., 2020). For this study the different sectors in the
 150 inventories were merged into three main sectors: agricultural E_{AGR} , anthropogenic non-
 151 agricultural E_{nonAGR} , and natural emissions E_N (see SI). We assume that these three
 152 sectors cover all N₂O emissions in the model domain. EDGAR4.3.2 and EDGAR5.0 pro-
 153 vide monthly resolved N₂O fluxes from anthropogenic source (E_{AGR} and E_{nonAGR}) on
 154 a $0.1^\circ \times 0.1^\circ$ grid for 2012 and 2015, respectively, but do not include fluxes from nat-
 155 ural sources. Hence, we supplemented both versions with yearly E_N on a $1^\circ \times 1^\circ$ grid
 156 from EDGAR version 2.0 (EDGAR2; Olivier et al. (1996, 1999)). All fluxes are assumed
 157 to originate from the surface.

158 With the process-based, biogeochemical model DayCent we estimated daily direct
 159 N₂O soil emissions from crop- and grassland on a $0.5^\circ \times 0.5^\circ$ grid in the Midwest from
 160 2011 to 2015, which were aggregated to a monthly time step. The model simulates fluxes
 161 of carbon and nitrogen between the atmosphere, vegetation, and soil thus deriving N₂O
 162 emissions. Incorporating several environmental drivers, including weather patterns, agri-
 163 cultural practices, soil characteristics, and crop features, this approach provides a more
 164 sophisticated estimate of soil emissions than the emission factor based EDGAR inven-
 165 tory. The GHG inventory of the United States Environmental Protection Agency (EPA,
 166 2020) uses DayCent estimates of direct soil emissions for emissions reporting of agricul-
 167 tural soil N₂O to the UN Framework Convention on Climate Change. DayCent does not
 168 calculate emissions from manure management, agricultural waste burning, indirect soil
 169 emissions, and those associated with minor crops such as vegetables. The EPA inven-
 170 tory quantifies these sources and subsources with an emission factor approach. We es-
 171 timate their contribution by employing the yearly estimates from EPA, calculating their
 172 relative fraction of the EPA direct soil emissions, and adding them to our monthly es-
 173 timates. As a result, our DayCent inventory properly accounts for the total agricultural
 174 emissions, but not the spatial distribution of agricultural sources which are not estimated
 175 by DayCent.

176 2.4 Optimization Technique

177 To solve for N₂O emissions, we use an approach similar to the optimization described
 178 in Barkley et al. (2017). First, we calculate the observed N₂O enhancements by subtract-
 179 ing a background from the measured absolute mole fraction. For each campaign we de-
 180 rive one background by taking the 2nd percentile of all low level legs of the entire cam-
 181 paign (see SI). The background is defined campaign-wise rather than transect-wise be-

182 cause during some transects we were not able to measure background mole fractions as
 183 we started a low level leg within a plume and did not exit the plume inside of the PBL
 184 (Figure 1b).

185 With observed N₂O enhancements calculated, we then compare modeled N₂O en-
 186 hancements emitted from our prior emission estimate ($E_{AGR} + E_{nonAGR} + E_N$) to the
 187 observed enhancements. Differences between model and observed enhancements are then
 188 minimized for each flight by scaling agricultural emissions E_{AGR} with a factor F_{AGR} thus
 189 quantifying emissions. This process is reliant on the assumption that the discrepancy
 190 between the observed and modeled N₂O is driven primarily by errors in the E_{AGR} . As
 191 agricultural emissions are the dominant N₂O source in our flights, we scale E_{AGR} , as-
 192 suming that errors in E_{nonAGR} and E_N are inconsequential to the overall solution. The
 193 complexity of N₂O soil emissions suggests that E_{AGR} exhibits a much higher uncertainty
 194 than other sources (Butterbach-Bahl et al., 2013), supporting the presented approach.

195 As an equation, this optimization technique is described by calculating F_{AGR} through
 196 the minimization of the following cost function:

$$197 \quad J(F_{AGR}) = |A_{obs} - \underbrace{(F_{AGR} \cdot A_{AGR} + A_{nonAGR} + A_N)}_{=A_{mod}(F_{AGR})}| \quad (1)$$

198 A_{obs} and A_{mod} are the time integral along a transect of observed and modeled enhance-
 199 ments, respectively (e.g., area below plume in Figure 3a). A_{mod} consists of an agricul-
 200 tural portion A_{AGR} scaleable with F_{AGR} , a non-agricultural anthropogenic portion A_{nonAGR} ,
 201 and a natural portion A_N . We compare integrals rather than enhancements themselves
 202 because we are interested in the amount of N₂O emitted in the atmosphere. Neither the
 203 model transport nor the inventory is perfect and even small uncertainties in just one of
 204 them could cause a shift or deformation in the alignment of the modeled plume relative
 205 to the observed plume. By minimizing the difference in the total N₂O enhancements rather
 206 than the point-by-point absolute error, we preserve the capability to solve for total N₂O
 207 emissions even when the modeled and observed plumes do not align. Due to the linear-
 208 ity between A_{AGR} and the area averaged E_{AGR} (see SI), a F_{AGR} derived with equation
 209 1 denotes a F_{AGR} -folded E_{AGR} .

210 **2.5 Uncertainty Assessment**

211 We adopted the method of Barkley, Davis, et al. (2019) to assess uncertainties in
 212 our solutions. F_{AGR} is affected by uncertainties in the following variables:

- 213 1. observed background mole fraction
- 214 2. A_{nonAGR}
- 215 3. A_N
- 216 4. model transport
- 217 5. model wind speed and PBL height
- 218 6. spatial distribution in EDGAR emissions

219 We quantify the influence of uncertainties 1 to 4 by using a Monte Carlo approach. For
 220 each flight we repeat the optimization 10 000 times with a perturbed background mole
 221 fraction, A_{nonAGR} , and A_N . For the background we take the value derived from the ob-
 222 servations and add a normal random number with $\mu = 0$ ppb and $\sigma = \pm 0.5$ ppb for
 223 2017 and $\sigma = \pm 0.9$ ppb for 2019. A_{nonAGR} and A_N are independently multiplied by
 224 a factor drawn from a normal distribution with $\mu = 1.0$ and $\sigma = \pm 0.21$ and $\sigma = \pm 0.42$,
 225 respectively. To account for the model transport error, we randomly select one of the three
 226 model runs with different meteorological initial and boundary conditions, creating vari-
 227 ability in the plume shape. The resulting spread in F_{AGR} is used as its uncertainty. Ex-
 228 planations of the values that represent the uncertainties are in the SI.

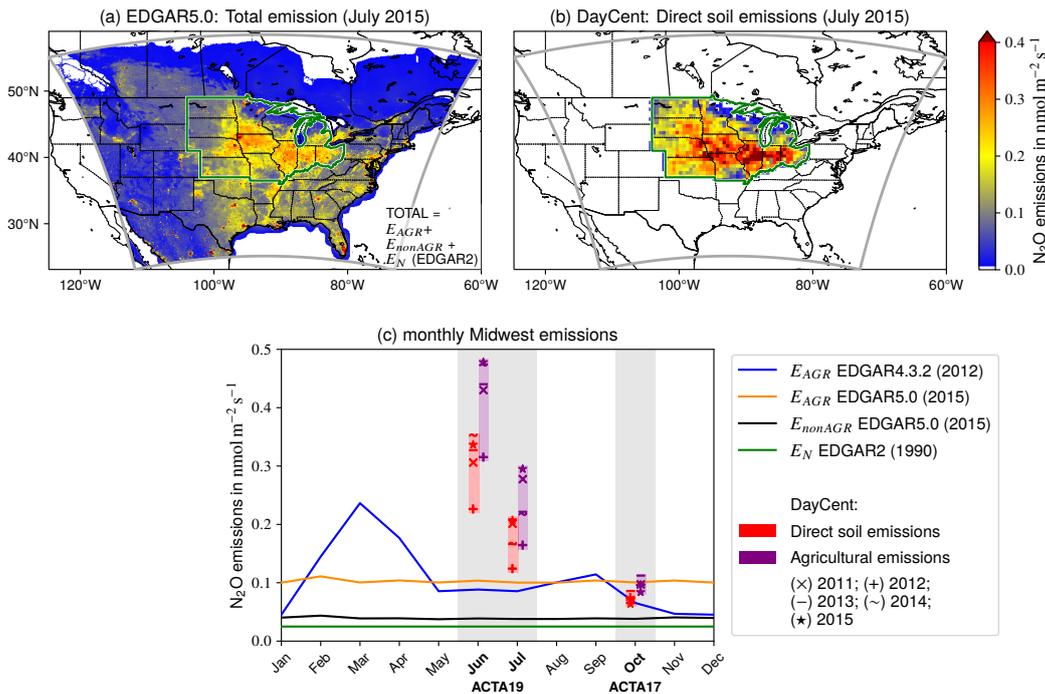


Figure 2. (a) EDGAR5.0 N₂O emissions (plus EDGAR2 E_N) within the model domain (gray box). The Midwest is encircled in green. (b) Direct soil emissions in July 2015 estimated with DayCent. (c) Monthly Midwest emissions. E_{nonAGR} in EDGAR4.3.2 is almost identical to EDGAR5.0. Total agricultural DayCent emissions are estimated utilizing the EPA GHG inventory (Section 2.3).

229 The modeled wind speed and PBL height uncertainty (source 5), cannot be covered
 230 by the Monte Carlo simulation. Errors in these variables cause lower or higher sim-
 231 ulated enhancements thus producing biases. Following Barkley et al. (2017) we correct
 232 for those biases by applying a correction factor based on the differences between the mod-
 233 eled and observed wind speed and PBL height. On average the modeled wind speed and
 234 PBL height is 8% and 3% higher than observations, respectively. The impact of this cor-
 235 rection on our results is insignificant. Results and further explanations can be found in
 236 the SI.

237 Our final source of uncertainty relates to uncertainties regarding errors in the spa-
 238 tial distribution of the fluxes in the prior inventory, and is difficult to quantify. However,
 239 the mapping of emissions in EDGAR is based on several high-resolution proxy data sets
 240 (Janssens-Maenhout et al., 2019). For this reason, we assume its spatial errors to be small.
 241 Given the insignificant difference between modeled and observed wind speeds and PBL
 242 heights, the good agreement between modeled and measured plume structures support
 243 this assumption (see SI). Furthermore, because we quantify large area sources and not
 244 point sources, slight misplacement in the inventory would only marginally affect our re-
 245 sults. At the same time, missing or strongly misplaced fluxes would produce errors that
 246 are not considered in this study.

3 Results and Discussion

3.1 Emission Inventory Comparison

Figure 2a shows prior July N_2O emissions in the outermost model domain from anthropogenic EDGAR5.0 and natural EDGAR2 sources. Compared to EDGAR4.3.2 no significant differences in the spatial distribution of emissions is seen, both versions just differ in the strength of the surface fluxes. The largest surface fluxes are concentrated in the Midwest, coinciding with the Corn Belt and its dominant agricultural emissions. Figure 2b shows DayCent direct soil emissions in July 2015. Similar to EDGAR emission maps, the Corn Belt within the Midwest is a prominent source of N_2O . We are not able to perform a detailed comparison of the spatial distributions in EDGAR and DayCent as both do not cover the same set of sources. However, in terms of the overall magnitude, DayCent estimates much higher surface fluxes compared to EDGAR, despite containing fewer sources (gridded total agricultural DayCent emissions are not available; Section 2.3).

Figure 2c displays the monthly evolution of E_{AGR} , E_{nonAGR} , and E_N averaged over the Midwest. Both EDGAR versions have an annual average E_{AGR} of approximately $0.10 \text{ nmol m}^{-2} \text{ s}^{-1}$. However, unlike EDGAR5.0, EDGAR4.3.2 exhibits a strong seasonal cycle ranging from $0.05 \text{ nmol m}^{-2} \text{ s}^{-1}$ in winter up to $0.24 \text{ nmol m}^{-2} \text{ s}^{-1}$ in spring. In spring, when most N-fertilizer is applied, the amount peaks, followed by a plateau during summer at $0.09 \text{ nmol m}^{-2} \text{ s}^{-1}$. The harvest season in fall features a local peak at $0.11 \text{ nmol m}^{-2} \text{ s}^{-1}$. In a future EDGAR5.0 release a seasonal cycle for some crop related emissions will be implemented (Crippa et al., 2020). E_{nonAGR} shows no significant change over the year and is on average $0.04 \text{ nmol m}^{-2} \text{ s}^{-1}$ in both versions. Natural soil emissions account for $0.02 \text{ nmol m}^{-2} \text{ s}^{-1}$ per month.

From 2011 to 2015 DayCent emissions in the Midwest range between $0.23\text{--}0.35 \text{ nmol m}^{-2} \text{ s}^{-1}$, $0.12\text{--}0.21 \text{ nmol m}^{-2} \text{ s}^{-1}$, and $0.06\text{--}0.08 \text{ nmol m}^{-2} \text{ s}^{-1}$ in June, July, and October respectively. June and July DayCent emissions are significantly larger than in EDGAR, despite manure management, indirect soil, and agricultural waste burning emissions not being included. DayCent's October emissions are within the magnitude of agricultural EDGAR emissions. We estimate total agricultural Midwest emissions from 2011 to 2015 by combining DayCent direct soil emissions and the EPA GHG inventory (Section 2.3), resulting in $0.32\text{--}0.48 \text{ nmol m}^{-2} \text{ s}^{-1}$, $0.16\text{--}0.30 \text{ nmol m}^{-2} \text{ s}^{-1}$, and $0.08\text{--}0.11 \text{ nmol m}^{-2} \text{ s}^{-1}$ in June, July, and October, respectively. In June/July this is on average over four/two times higher than EDGAR E_{AGR} estimates. The 2012 emissions are significantly lower than in the other years causing the large range across years in the summer months. During this year, the most extensive drought since the 1930s occurred across a large swath of the U.S., including most of the Midwest, which lead to widespread harvest failure (NOAA-NCEI, 2020). This event might explain the low values and indicates that during an average climatological year DayCent emissions are at the upper end of the range. Furthermore, in contrast to EDGAR4.3.2 which states constant emissions in June and July, DayCent emissions are much higher in June than in July. This is consistent with the N_2O climatology in Sweeney et al. (2015).

3.2 Model Optimization

Here, we provide an example of the model optimization process for Oct 10, 2017 (Figure 3a). In the eastern part of the Midwest N_2O enhancements up to 7 ppb were observed within the PBL. The slightly negative values at the beginning of the time series occurred prior to the low level leg in the free troposphere. Our background is derived from air within the PBL and is representative for the time and location of the campaign. Free tropospheric air might have a different history and hence different background which can lead to negative values if we subtract our background. Model simulations with unmodified EDGAR emissions show only enhancements up to 1 ppb along the transect. How-

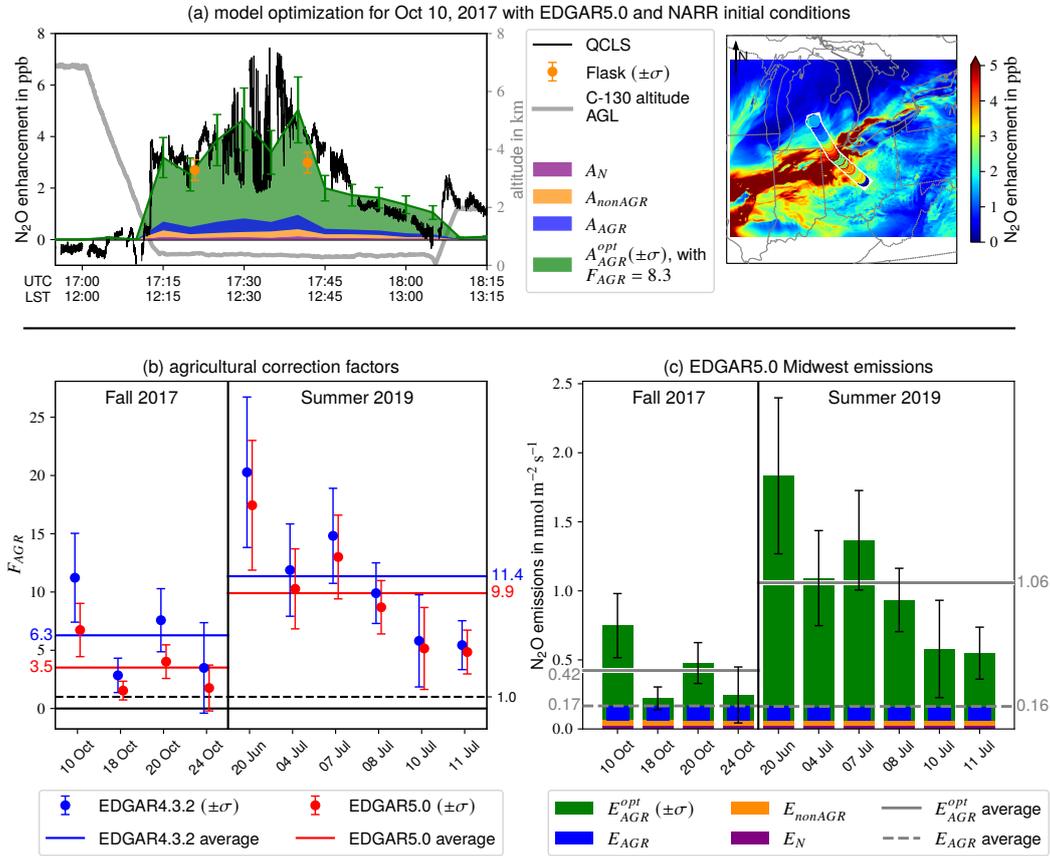


Figure 3. (a) Sample model optimization for Oct 10, 2017 with EDGAR5.0 (plus EDGAR2 E_N) and NARR initial conditions. The left panel shows the prior and optimized modeled N_2O enhancements along the flight track together with observed enhancements. The right panel shows a map of optimized modeled N_2O enhancements (from $E_{AGR}^{opt} + E_{nonAGR} + E_N$) at 300 m AGL at 17:30 UTC and the flight track color-coded with the observed enhancements. (b) Mean and standard deviation of agricultural correction factors F_{AGR} for the investigated research flights resulting from Monte Carlo simulations. (c) EDGAR5.0 Midwest N_2O emissions with optimized and prior E_{AGR} .

297 ever, by applying an agricultural correction factor F_{AGR} of 8.3 the model is able to
 298 reproduce our measurements. Optimizations of the remaining days can be found in the SI.

299 Figure 3b shows the mean and standard deviation for F_{AGR} of the Monte Carlo
 300 simulations of the ten research flights for the two EDGAR versions. As both invento-
 301 ries have a comparable spatial distribution, factors vary due to differences in total emis-
 302 sions. EDGAR4.3.2 correction factors are considerably higher for October 2017 and slightly
 303 higher for June/July 2019 than EDGAR5.0. For EDGAR4.3.2, F_{AGR} ranges from $2.9 \pm$
 304 1.5 to 11.3 ± 3.8 in 2017, with an average factor of 6.3 ± 4.6 . EDGAR5.0 F_{AGR} is cal-
 305 culated to be lower but still ranges from 1.6 ± 0.8 to 6.8 ± 2.3 , with an average factor
 306 of 3.5 ± 2.7 . For 2019 we modified EDGAR4.3.2 with a F_{AGR} between 5.5 ± 2.1 and
 307 20.2 ± 6.3 and EDGAR5.0 between 4.9 ± 1.9 and 17.4 ± 5.5 . On average this denotes an
 308 agricultural correction factor of 11.4 ± 6.6 and 9.9 ± 5.7 for EDGAR4.3.2 and EDGAR5.0,
 309 respectively. Altogether, both EDGAR versions exhibit a significant underestimation of
 310 agricultural emissions. Seasonal differences are likely one cause for the large difference
 311 in correction factors between 2017 and 2019. Additionally, during the 2019 aircraft cam-
 312 paign, an extreme flooding event occurred that likely influenced our results (discussed
 313 below). Although EDGAR4.3.2 exhibits a seasonal cycle, its agricultural correction fac-
 314 tor also varies considerably between 2017 and 2019. Hence, the seasonality is not cap-
 315 tured in the EDGAR inventory for the Midwest, which appears to be caused by the flood-
 316 ing. Figure 3c displays the EDGAR5.0 average Midwest emissions for each flight day with
 317 non-optimized and optimized agricultural emissions. For EDGAR4.3.2 the optimized re-
 318 sult is (nearly) the same as both versions differ (nearly) only in their strength of E_{AGR}
 319 which is adjusted in the course of the optimization. On average, optimized total N_2O
 320 emissions are $0.42 \pm 0.28 \text{ nmol m}^{-2} \text{ s}^{-1}$ in 2017 and $1.06 \pm 0.57 \text{ nmol m}^{-2} \text{ s}^{-1}$ in 2019.

321 Optimized emissions for June/July 2019 are 2–3 times higher compared to Day-
 322 Cent emissions. Despite this, DayCent emissions are closer to our optimized emissions
 323 compared to EDGAR during the same period. In contrast, DayCent and EDGAR emis-
 324 sions are both too low by a similar magnitude in October compared to our optimized
 325 results. Hence, as DayCent considers regional characteristics, it performs much better
 326 on the regional scale in the summer than the emission factor approach that is used in
 327 the EDGAR inventory. A more quantitative evaluation of DayCent would require sur-
 328 face flux calculations for 2017 and 2019 incorporating the corresponding regional con-
 329 ditions like weather, soil conditions, and N-fertilizer application rate and time. DayCent
 330 has not been applied to estimate emissions specific to 2017 and 2019 so it is not clear
 331 if the model would underestimate the values for these years although this may be the
 332 case given the historical data from 2011–2015.

333 Fu et al. (2017) reported emissions of $3.00\text{--}4.38 \text{ nmol m}^{-2} \text{ s}^{-1}$ during June 1–20, 2010
 334 for the Corn Belt, which is significantly higher than our estimates for June/July 2019.
 335 Griffis et al. (2013) estimated the Corn Belt emissions to be around $2 \text{ nmol m}^{-2} \text{ s}^{-1}$ and
 336 $1 \text{ nmol m}^{-2} \text{ s}^{-1}$ in June/July 2010 and 2011, respectively, which is consistent with our
 337 findings. Kort et al. (2008) and Miller et al. (2012) derived scaling factors for the cen-
 338 tral U.S. To be able to compare their results to ours, we estimated the corresponding
 339 flux densities for the Midwest region using their scaling factors for the respective EDGAR
 340 versions. Kort et al. (2008) derived $0.54 \text{ nmol m}^{-2} \text{ s}^{-1}$ for May/June 2003 and Miller et
 341 al. (2012) $0.57/0.25 \text{ nmol m}^{-2} \text{ s}^{-1}$ and $0.94/0.53 \text{ nmol m}^{-2} \text{ s}^{-1}$ for June/July 2004 and
 342 2008, respectively. Both studies show lower values than our estimate. Miller et al. (2012)
 343 stated that maximum emissions occurred in June. Our DayCent calculations are also high-
 344 est in June. This could partly explain our lower estimates compared to Fu et al. (2017)
 345 as we report for the end of June/beginning of July after the expected emission peak. More-
 346 over, Fu et al. (2017) only scaled Corn Belt emissions and kept other regions unmodi-
 347 fied which could lead to higher estimates, if they sampled other regions with lower emis-
 348 sion rates than the Corn Belt. Overall, our estimates are in the range of previous top-
 349 down studies. However, the spread among the studies is large.

350 The nature of soil N₂O emissions leads to significant temporal variability in the emis-
 351 sions that is not represented in the EDGAR inventory. Unlike EDGAR, DayCent is ca-
 352 pable of representing those variations to a certain extent. In our 2011–2015 calculations
 353 the monthly standard deviations range from 10 % in October to 21 % in July, demon-
 354 strating the strong interannual variability. Furthermore, weather conditions in the study
 355 domain in 2019 were unusually extreme. During the campaign, the U.S. was experienc-
 356 ing its wettest period in 125 years, with severe flooding in the Midwest (NOAA, 2020)
 357 forcing the farmers to significantly delay planting in the affected regions (USDA, 2020)
 358 and postponing the peak emission period. Depending on whether the zenith is shifted
 359 closer to or further away from our investigated period in June/July this event may have
 360 either amplified or lowered our emission estimates. Additionally, the above-average hu-
 361 midity might have enhanced soil N₂O emissions leading to higher estimates (Butterbach-
 362 Bahl et al., 2013). The influence of this flooding event cannot be quantified within this
 363 study, as this would require more data over longer periods spanning the whole event. How-
 364 ever, in a follow-up study we plan to use DayCent simulations driven with those flood-
 365 ing conditions to gain insights on how soil N₂O emissions were affected.

366 4 Conclusion

367 Unique continuous in situ airborne N₂O measurements of ten research flights were
 368 used to quantify N₂O emissions in the U.S. Midwest using a top-down approach. In Oc-
 369 tober 2017 and June/July 2019 agricultural Midwest emission were on average $6.3 \pm 4.6 / 3.5 \pm$
 370 2.7 and $11.4 \pm 6.6 / 9.9 \pm 5.7$ times higher than EDGAR4.3.2/EDGAR5.0 estimates re-
 371 sulting in $0.42 \pm 0.28 \text{ nmol m}^{-2} \text{ s}^{-1}$ and $1.06 \pm 0.57 \text{ nmol m}^{-2} \text{ s}^{-1}$ Midwest emissions,
 372 respectively. Our 2019 estimates were most likely influenced by an extreme flooding event,
 373 which is difficult to capture in EDGAR as the inventory uses a more climatological av-
 374 erage emissions dataset. Agricultural soil emissions estimated with DayCent in 2011–
 375 2015 were 0.32–0.48, 0.16–0.30, and 0.08–0.11 $\text{nmol m}^{-2} \text{ s}^{-1}$ in June, July, and October,
 376 respectively. Based on these historical emission estimates, this is higher than non-optimized
 377 EDGAR emissions, but still significantly lower than our optimized fluxes. Our findings
 378 are in the range of previous top-down estimates for the Corn Belt and central U.S. How-
 379 ever, a quantitative comparison of those studies show that the range of derived N₂O sur-
 380 face fluxes is large, likely due to the temporal complexity of N₂O soil emissions.

381 More N₂O focused studies are necessary to fully understand the drivers of Midwest
 382 N₂O emissions and the most appropriate modeling methods to estimate emission pat-
 383 terns. To cover the high temporal variability on various scales, long term projects with
 384 regular airborne measurements spanning wide areas of the Midwest are necessary. Com-
 385 bining a process-based model like DayCent capable of simulating the temporal and spa-
 386 tial variability of N₂O emissions, with extensive airborne and tall tower top-down stud-
 387 ies at selected spots and times, could be a cost effective approach that would limit the
 388 number of flights needed to produce accurate estimates for the region and improve na-
 389 tional reporting of emissions (Ogle et al., 2020). As interest grows in expanding efforts
 390 to reduce N₂O emissions (Kanter et al., 2020), improved quantification of N₂O surface
 391 fluxes is mandatory for policy makers to be able to develop effective mitigation strate-
 392 gies.

393 Acknowledgments

394 The ACT-America project is a NASA Earth Venture Suborbital-2 project funded by NASA’s
 395 Earth Science Division (grant NNX15AG76G to the Pennsylvania State University). All
 396 ACT-America flask and in situ data used in this manuscript can be found online (at [https://
 397 daac.ornl.gov/cgi-bin/dataset_lister.pl?p=37](https://daac.ornl.gov/cgi-bin/dataset_lister.pl?p=37)). We thank DLR VO-R for fund-
 398 ing the young investigator research group “Greenhouse Gases”.

References

399

- 400 Baier, B. C., Sweeney, C., Choi, Y., Davis, K. J., DiGangi, J. P., Feng, S., . . . Weibring, P. (2020). Multispecies Assessment of Factors Influencing Regional CO₂ and CH₄ Enhancements During the Winter 2017 ACT-America Campaign. *Journal of Geophysical Research: Atmospheres*, *125*, e2019JD031339. doi: 10.1029/2019JD031339
- 405 Barkley, Z. R., Davis, K. J., Feng, S., Balashov, N., Fried, A., DiGangi, J., . . . Halliday, H. S. (2019). Forward Modeling and Optimization of Methane Emissions in the South Central United States Using Aircraft Transects Across Frontal Boundaries. *Geophysical Research Letters*, *46*(22), 13564–13573. doi: 10.1029/2019gl084495
- 410 Barkley, Z. R., Lauvaux, T., Davis, K. J., Deng, A., Fried, A., Weibring, P., . . . Dickerson, R. R. (2019). Estimating Methane Emissions From Underground Coal and Natural Gas Production in Southwestern Pennsylvania. *Geophysical Research Letters*, *46*(8), 4531–4540. doi: 10.1029/2019GL082131
- 414 Barkley, Z. R., Lauvaux, T., Davis, K. J., Deng, A., Miles, N. L., Richardson, S. J., . . . Maasackers, J. D. (2017). Quantifying methane emissions from natural gas production in north-eastern Pennsylvania. *Atmospheric Chemistry and Physics*, *17*(22), 13941–13966. doi: 10.5194/acp-17-13941-2017
- 418 Butterbach-Bahl, K., Baggs, E. M., Dannenmann, M., Kiese, R., & Zechmeister-Boltenstern, S. (2013). Nitrous oxide emissions from soils: how well do we understand the processes and their controls? *Philosophical Transactions of the Royal Society B: Biological Sciences*, *368*, 20130122. doi: 10.1098/rstb.2013.0122
- 423 Chen, Z., Griffis, T. J., Millet, D. B., Wood, J. D., Lee, X., Baker, J. M., . . . Wells, K. C. (2016). Partitioning N₂O emissions within the U.S. Corn Belt using an inverse modeling approach. *Global Biogeochemical Cycles*, *30*(8), 1192–1205. doi: 10.1002/2015gb005313
- 427 Ciais, P., Sabine, C., Bala, G., Bopp, L., Brovkin, V., Canadell, J., . . . Thornton, P. (2013). Carbon and Other Biogeochemical Cycles. In T. F. Stocker et al. (Eds.), *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change* (pp. 465–570). Cambridge, United Kingdom and New York, NY, USA: Cambridge University Press.
- 433 Crippa, M., Solazzo, E., Huang, G., Guizzardi, D., Koffi, E., Muntean, M., . . . Janssens-Maenhout, G. (2020). High resolution temporal profiles in the Emissions Database for Global Atmospheric Research. *Scientific Data*, *7*(121). doi: 10.1038/s41597-020-0462-2
- 437 Del Grosso, S. J., Parton, W. J., Keough, C. A., & Reyes-Fox, M. (2011). Special features of the DayCent modeling package and additional procedures for parameterization, calibration, validation, and applications. In L. R. Ahuja & L. Ma (Eds.), *Methods of Introducing System Models into Agricultural Research* (pp. 155–176). Madison, WI, USA: American Society of Agronomy, Crop Science Society of America, Soil Science Society of America. doi: 10.2134/advagricsystmodel2.c5
- 444 Del Grosso, S. J., Parton, W. J., Mosier, A. R., Hartman, M. D., Brenner, J., Ojima, D. S., & Schimel, D. S. (2001). Simulated Interaction of Carbon Dynamics and Nitrogen Trace Gas Fluxes Using the DAYCENT Model. In M. Schaffer, L. Ma, & S. Hansen (Eds.), *Modeling Carbon and Nitrogen Dynamics for Soil Management* (pp. 303–332). Boca Raton, Florida, USA: CRC Press.
- 449 Díaz-Isaac, L. I., Lauvaux, T., & Davis, K. J. (2018). Impact of physical parameterizations and initial conditions on simulated atmospheric transport and CO₂ mole fractions in the US Midwest. *Atmospheric Chemistry and Physics*, *18*(20), 14813–14835. doi: 10.5194/acp-18-14813-2018
- 453 EDGAR. (2020). *Emission Database for Global Atmospheric Research*. Retrieved

- 454 from <https://edgar.jrc.ec.europa.eu/> (last accessed: 20 Jul 2020)
- 455 EDGAR4.3.2. (2017). *Emissions Database for Global Atmospheric Research, version*
- 456 *4.3.2*. European Commission. Retrieved from [https://edgar.jrc.ec.europa](https://edgar.jrc.ec.europa.eu/overview.php?v=432_GHG)
- 457 [.eu/overview.php?v=432_GHG](https://edgar.jrc.ec.europa.eu/overview.php?v=432_GHG) doi: [https://data.europa.eu/doi/10.2904/JRC](https://data.europa.eu/doi/10.2904/JRC-DATASET-EDGAR)
- 458 [-DATASET-EDGAR](https://data.europa.eu/doi/10.2904/JRC-DATASET-EDGAR)
- 459 EDGAR5.0. (2019). *Emissions Database for Global Atmospheric Research, version*
- 460 *5.0*. European Commission. Retrieved from [https://edgar.jrc.ec.europa](https://edgar.jrc.ec.europa.eu/overview.php?v=50_GHG)
- 461 [.eu/overview.php?v=50_GHG](https://edgar.jrc.ec.europa.eu/overview.php?v=50_GHG) doi: [https://data.europa.eu/doi/10.2904/JRC](https://data.europa.eu/doi/10.2904/JRC-DATASET-EDGAR)
- 462 [-DATASET-EDGAR](https://data.europa.eu/doi/10.2904/JRC-DATASET-EDGAR)
- 463 EPA. (2020). *Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990–2018*.
- 464 United States Environmental Protection Agency. EPA 430-R-20-002. Retrieved
- 465 from [https://www.epa.gov/ghgemissions/inventory-us-greenhouse-gas](https://www.epa.gov/ghgemissions/inventory-us-greenhouse-gas-emissions-and-sinks-1990-2018)
- 466 [-emissions-and-sinks-1990-2018](https://www.epa.gov/ghgemissions/inventory-us-greenhouse-gas-emissions-and-sinks-1990-2018)
- 467 ERA5. (2017). *Copernicus Climate Change Service (C3S) (2017): ERA5: Fifth gen-*
- 468 *eration of ECMWF atmospheric reanalyses of the global climate*. Copernicus
- 469 Climate Change Service Climate Data Store (CDS). Retrieved from [https://](https://cds.climate.copernicus.eu/cdsapp#!/home)
- 470 cds.climate.copernicus.eu/cdsapp#!/home (last accessed: 02 Mar 2020)
- 471 FAO. (2020). *Food and Agriculture Organization of the United Nations - FAOSTAT*.
- 472 Retrieved from <http://www.fao.org/faostat/en/#compare> (last accessed:
- 473 20 Jul 2020)
- 474 Fu, C., Lee, X., Griffis, T. J., Dlugokencky, E. J., & Andrews, A. E. (2017). Inves-
- 475 tigation of the N₂O emission strength in the U. S. Corn Belt. *Atmospheric Re-*
- 476 *search*, 194, 66–77. doi: 10.1016/j.atmosres.2017.04.027
- 477 GDAS-FNL. (2015). *National Centers for Environmental Prediction, National*
- 478 *Weather Service, NOAA, U.S. Department of Commerce: NCEP GDAS/FNL*
- 479 *0.25 Degree Global Tropospheric Analyses and Forecast Grids, updated daily*.
- 480 Research Data Archive at the National Center for Atmospheric Research,
- 481 Computational and Information Systems Laboratory. (last accessed: 28 May
- 482 2020) doi: 10.5065/D65Q4T4Z
- 483 Grell, G. A., Peckham, S. E., Schmitz, R., McKeen, S. A., Frost, G., Skamarock,
- 484 W. C., & Eder, B. (2005). Fully coupled “online” chemistry within
- 485 the WRF model. *Atmospheric Environment*, 39(37), 6957–6975. doi:
- 486 10.1016/j.atmosenv.2005.04.027
- 487 Griffis, T. J., Lee, X., Baker, J. M., Russelle, M. P., Zhang, X., Venterea, R., & Mil-
- 488 let, D. B. (2013). Reconciling the differences between top-down and bottom-up
- 489 estimates of nitrous oxide emissions for the U.S. Corn Belt. *Global Biogeo-*
- 490 *chemical Cycles*, 27(3), 746–754. doi: 10.1002/gbc.20066
- 491 Hall, B. D., Dutton, G. S., & Elkins, J. W. (2007). The NOAA nitrous oxide stan-
- 492 dard scale for atmospheric observations. *Journal of Geophysical Research: At-*
- 493 *mospheres*, 112, D09305. doi: 10.1029/2006JD007954
- 494 Hersbach, H., Bell, B., Berrisford, P., Hirahara, S., Horányi, A., Muñoz Sabater, J.,
- 495 ... Thépaut, J.-N. (2020). The ERA5 global reanalysis. *Quarterly Journal of*
- 496 *the Royal Meteorological Society*, 1–51. doi: 10.1002/qj.3803
- 497 Janssens-Maenhout, G., Crippa, M., Guizzardi, D., Muntean, M., Schaaf, E., Den-
- 498 tener, F., ... Oreggioni, G. D. (2019). EDGAR v4.3.2 Global Atlas of the
- 499 three major greenhouse gas emissions for the period 1970–2012. *Earth System*
- 500 *Science Data*, 11(3), 959–1002. doi: 10.5194/essd-11-959-2019
- 501 Kanter, D. R., Ogle, S. M., & Winiwarter, W. (2020). Building on Paris: integrat-
- 502 ing nitrous oxide mitigation into future climate policy. *Current Opinion in En-*
- 503 *vironmental Sustainability*, 47, 1–6. doi: 10.1016/j.cosust.2020.04.005
- 504 Kort, E. A., Eluszkiewicz, J., Stephens, B. B., Miller, J. B., Gerbig, C., Nehr Korn,
- 505 T., ... Wofsy, S. C. (2008). Emissions of CH₄ and N₂O over the United States
- 506 and Canada based on a receptor-oriented modeling framework and COBRA-
- 507 NA atmospheric observations. *Geophysical Research Letters*, 35, L18808. doi:
- 508 10.1029/2008GL034031

- 509 Kostinek, J., Roiger, A., Davis, K. J., Sweeney, C., DiGangi, J. P., Choi, Y., ...
 510 Butz, A. (2019). Adaptation and performance assessment of a quantum and
 511 interband cascade laser spectrometer for simultaneous airborne in situ observa-
 512 tion of CH₄, C₂H₆, CO₂, CO and N₂O. *Atmospheric Measurement Techniques*,
 513 12(3), 1767–1783. doi: 10.5194/amt-12-1767-2019
- 514 MacFarling Meure, C., Etheridge, D., Trudinger, C., Steele, P., Langenfelds, R., van
 515 Ommen, T., ... Elkins, J. (2006). Law Dome CO₂, CH₄ and N₂O ice core
 516 records extended to 2000 years BP. *Geophysical Research Letters*, 33(14). doi:
 517 10.1029/2006GL026152
- 518 Miller, S. M., Kort, E. A., Hirsch, A. I., Dlugokencky, E. J., Andrews, A. E., Xu,
 519 X., ... Wofsy, S. C. (2012). Regional sources of nitrous oxide over the United
 520 States: Seasonal variation and spatial distribution. *Journal of Geophysical*
 521 *Research: Atmospheres*, 117, D06310. doi: 10.1029/2011JD016951
- 522 Myhre, G., Shindell, D., Bréon, F.-M., Collins, W., Fuglestedt, J., Huang, J.,
 523 ... Zhang, H. (2013). Anthropogenic and Natural Radiative Forcing. In
 524 T. F. Stocker et al. (Eds.), *Climate Change 2013: The Physical Science Basis.*
 525 *Contribution of Working Group I to the Fifth Assessment Report of the Inter-*
 526 *governmental Panel on Climate Change* (pp. 659–740). Cambridge, United
 527 Kingdom and New York, NY, USA: Cambridge University Press.
- 528 NARR. (2005). *National Centers for Environmental Prediction, National Weather*
 529 *Service, NOAA, U.S. Department of Commerce: NCEP North American Re-*
 530 *gional Reanalysis, updated monthly.* Research Data Archive at the National
 531 Center for Atmospheric Research, Computational and Information Systems
 532 Laboratory. Retrieved from <https://rda.ucar.edu/datasets/ds608.0> (last
 533 accessed: 27 May 2020)
- 534 NOAA. (2020). *National Centers for Environmental Information: Climate at a*
 535 *Glance - Rankings.* Retrieved from <https://www.ncdc.noaa.gov/cag/> (last
 536 accessed: 20 Jul 2020)
- 537 NOAA-ESRL. (2020). *Combined Nitrous Oxide data from the NOAA/ESRL Global*
 538 *Monitoring Division.* Retrieved from [https://www.esrl.noaa.gov/gmd/hats/](https://www.esrl.noaa.gov/gmd/hats/combined/N20.html)
 539 [combined/N20.html](https://www.esrl.noaa.gov/gmd/hats/combined/N20.html) (last accessed: 20 Jul 2020)
- 540 NOAA-NCEI. (2020). *U.S. Billion-Dollar Weather and Climate Disasters.* NOAA
 541 National Centers for Environmental Information (NCEI). Retrieved from
 542 <https://www.ncdc.noaa.gov/billions/> doi: 10.25921/stkw-7w73
- 543 Ogle, S. M., Butterbach-Bahl, K., Cardenas, L., Skiba, U., & Scheer, C. (2020).
 544 From research to policy: optimizing the design of a national monitoring system
 545 to mitigate soil nitrous oxide emissions. *Current Opinion in Environmental*
 546 *Sustainability*, 47, 28–36. doi: 10.1016/j.cosust.2020.06.003
- 547 Olivier, J. G. J., Bouwman, A. F., Berdowski, J. J. M., Veldt, C., Bloos, J. P. J.,
 548 Visschedijk, A. J. H., ... Zandveld, P. Y. J. (1999). Sectoral emis-
 549 sion inventories of greenhouse gases for 1990 on a per country basis as
 550 well as on 1°×1°. *Environmental Science & Policy*, 2(3), 241–263. doi:
 551 10.1016/s1462-9011(99)00027-1
- 552 Olivier, J. G. J., Bouwman, A. F., van der Maas, C. W. M., Berdowski, J. J. M.,
 553 Veldt, C., Bloos, J. P. J., ... Haverlag, J. L. (1996). Description of EDGAR
 554 Version 2.0: A set of global emission inventories of greenhouse gases and
 555 ozone-depleting substances for all anthropogenic and most natural sources
 556 on a per country basis and on 1°×1° grid. *National Institute of Public Health*
 557 *and the Environment (RIVM) report no. 771060 002 / TNO-MEP report no.*
 558 *R96/119.* Retrieved from <http://hdl.handle.net/10029/10497>
- 559 Park, S., Croteau, P., Boering, K. A., Etheridge, D. M., Ferretti, D., Fraser, P. J.,
 560 ... M., T. C. (2012). Trends and seasonal cycles in the isotopic compo-
 561 sition of nitrous oxide since 1940. *Nature Geoscience*, 5(4), 261–265. doi:
 562 10.1038/ngeo1421
- 563 Parton, W. J., Hartman, M., Ojima, D., & Schimel, D. (1998). DAYCENT and its

- 564 land surface submodel: description and testing. *Global and Planetary Change*,
565 19(1), 35–48. doi: 10.1016/S0921-8181(98)00040-X
- 566 Paustian, K., Lehmann, J., Ogle, S. M., Reay, D., Robertson, G. P., & Smith, P.
567 (2016). Climate-smart soils. *Nature*, 532, 49–57. doi: 10.1038/nature17174
- 568 Prather, M. J., Hsu, J., DeLuca, N. M., Jackman, C. H., Oman, L. D., Douglass,
569 A. R., ... Funke, B. (2015). Measuring and modeling the lifetime of nitrous
570 oxide including its variability. *Journal of Geophysical Research: Atmospheres*,
571 120(11), 5693–5705. doi: 10.1002/2015jd023267
- 572 Ravishankara, A. R., Daniel, J. S., & Portmann, R. W. (2009). Nitrous Oxide
573 (N₂O): The Dominant Ozone-Depleting Substance Emitted in the 21st Cen-
574 tury. *Science*, 326(5949), 123–125. doi: 10.1126/science.1176985
- 575 Reay, D. S., Davidson, E. A., Smith, K. A., Smith, P., Melillo, J. M., Dentener, F.,
576 & Crutzen, P. J. (2012). Global agriculture and nitrous oxide emissions.
577 *Nature Climate Change*, 2(6), 410–416. doi: 10.1038/nclimate1458
- 578 Stehfest, E., & Bouwman, L. (2006). N₂O and NO emission from agricultural fields
579 and soils under natural vegetation: summarizing available measurement data
580 and modeling of global annual emissions. *Nutrient Cycling in Agroecosystems*,
581 74(3), 207–228. doi: 10.1007/s10705-006-9000-7
- 582 Sweeney, C., Baier, B. C., Miller, J. B., Lang, P., Miller, B. R., Lehman, S., ...
583 Yang, M. M. (2018). *ACT-America: L2 In Situ Atmospheric Gas Concentra-*
584 *tions from Flasks, Eastern USA*. ORNL Distributed Active Archive Center.
585 Retrieved from https://daac.ornl.gov/cgi-bin/dsviewer.pl?ds_id=1575
586 doi: 10.3334/ORNLDAAC/1575
- 587 Sweeney, C., Karion, A., Wolter, S., Newberger, T., Guenther, D., Higgs, J. A., ...
588 Tans, P. P. (2015). Seasonal climatology of CO₂ across North America from
589 aircraft measurements in the NOAA/ESRL Global Greenhouse Gas Reference
590 Network. *Journal of Geophysical Research: Atmospheres*, 120(10), 5155–5190.
591 doi: 10.1002/2014jd022591
- 592 Thompson, R. L., Lassaletta, L., Patra, P. K., Wilson, C., Wells, K. C., Gressent,
593 A., ... Canadell, J. G. (2019). Acceleration of global N₂O emissions seen from
594 two decades of atmospheric inversion. *Nature Climate Change*, 9(12), 993–998.
595 doi: 10.1038/s41558-019-0613-7
- 596 Tian, H., Xu, R., Canadell, J. G., Thompson, R. L., Winiwarter, W., Sun-
597 tharalingam, P., ... Yao, Y. (2020). A comprehensive quantification
598 of global nitrous oxide sources and sinks. *Nature*, 586, 248–256. doi:
599 10.1038/s41586-020-2780-0
- 600 USDA. (2020). *Economics, Statistics and Market Information System - Crop*
601 *Progress*. Retrieved from [https://usda.library.cornell.edu/concern/](https://usda.library.cornell.edu/concern/publications/8336h188j?locale=en#release-items)
602 [publications/8336h188j?locale=en#release-items](https://usda.library.cornell.edu/concern/publications/8336h188j?locale=en#release-items) (last accessed: 20 Jul
603 2020)
- 604 USDA-NASS. (2020). *United States Department of Agriculture - National Agricul-*
605 *tural Statistics Service - Statistics by State*. Retrieved from [https://www.nass](https://www.nass.usda.gov/Statistics_by_State/index.php)
606 [.usda.gov/Statistics_by_State/index.php](https://www.nass.usda.gov/Statistics_by_State/index.php) (last accessed: 23 Jul 2020)