# Quantifying nitrous oxide emissions from agriculture in the Midwest of the U.S.

Maximilian Eckl<sup>1</sup>, Anke Roiger<sup>1</sup>, Julian Kostinek<sup>1</sup>, Alina Fiehn<sup>1</sup>, Heidi Huntrieser<sup>1</sup>, Christoph Knote<sup>2</sup>, Zachary Barkley<sup>3</sup>, Bianca Baier<sup>4</sup>, Colm Sweeney<sup>5</sup>, and Kenneth Davis<sup>3</sup>

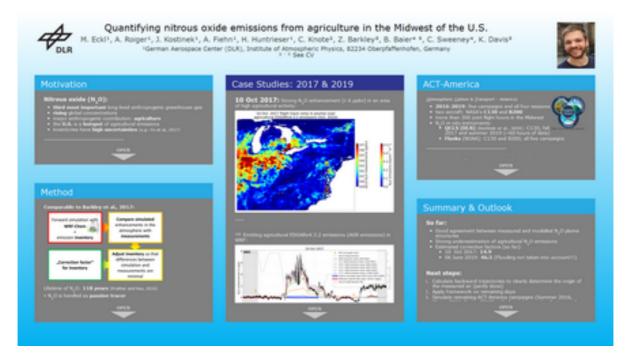
<sup>1</sup>German Aerospace Center (DLR) Oberpfaffenhofen <sup>2</sup>Ludwig-Maximilians-University (LMU) <sup>3</sup>The Pennsylvania State University <sup>4</sup>Cooperative Institute for Research in Environmental Sciences <sup>5</sup>NOAA ESRL

November 22, 2022

#### Abstract

Atmospheric nitrous oxide (N2O) is, after carbon dioxide and methane, the third most important long-lived anthropogenic greenhouse gas in terms of radiative forcing. Since preindustrial times a rising trend in the global N2O concentrations is observed. Anthropogenic emissions of N2O, mainly from agricultural activity, contribute considerably to this trend. Sparse observational constraints have made it difficult to quantify these emissions. The few studies on top-down approaches in the U.S. that exist are mainly based on Lagrangian models and ground-based measurements. They all propose a significant underestimation of anthropogenic N2O emission sources in established inventories, such as the Emissions Database for Global Atmospheric Research (EDGAR). In this study we quantify anthropogenic N2O emissions in the Midwest of the U.S., an area of high agricultural activity. In the course of the Atmospheric Carbon and Transport - America (ACT-America) campaign spanning from summer 2016 to summer 2019, an extensive dataset over four seasons has been collected including in-situ N2O aircraft based measurements in the lower and middle troposphere onboard NASA's C-130 and B-200 aircraft. During fall 2017 and summer 2019 we conducted measurements onboard the NASA-C130 with a Quantum-Cascade-Laser-Spectrometer (QCLS) and on both aircraft over the whole campaign flask measurements (NOAA) were collected. More than 300 joint flight hours were conducted and more than 500 flask samples were collected over the U.S. Midwest. The QCLS system collected continuous N2O data for approximately 60 flight hours in this region. The Eulerian Weather Research and Forecasting model with chemistry enabled (WRF-Chem) is being used to quantify regional agricultural N2O emissions using the spatial characteristics of these atmospheric N2O mole fraction observations. The numerical simulations enable potential surface emission distributions to be compared to our airborne measurements, and source estimates can be adjusted to minimize the differences, thus quantifying N2O sources. These results are then compared to emission rates in the EDGAR inventory.

# Quantifying nitrous oxide emissions from agriculture in the Midwest of the U.S.



M. Eckl<sup>1</sup>, A. Roiger<sup>1</sup>, J. Kostinek<sup>1</sup>, A. Fiehn<sup>1</sup>, H. Huntrieser<sup>1</sup>, C. Knote<sup>2</sup>, Z. Barkley<sup>3</sup>, B. Baier<sup>4 5</sup>, C. Sweeney<sup>4</sup>, K. Davis<sup>3</sup>

<sup>1</sup>German Aerospace Center (DLR), Institute of Atmospheric Physics, 82234 Oberpfaffenhofen, Germany <sup>2 - 5</sup> See CV



PRESENTED AT:



### MOTIVATION

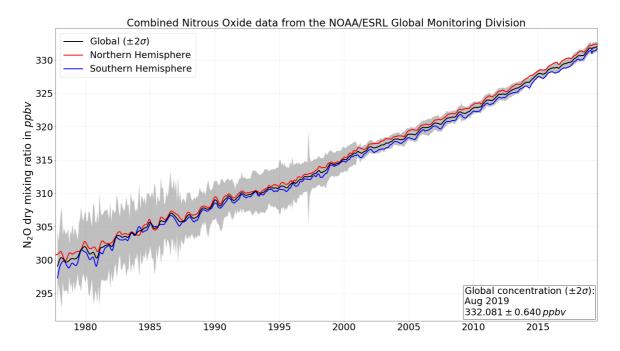
#### Nitrous oxide (N<sub>2</sub>O):

- third most important long-lived anthropogenic greenhouse gas
- rising global concentrations
- major anthropogenic contribution: agriculture
- the **U.S.** is a **hotspot** of agricultural emissions
- inventories have high uncertainties (e.g.: Fu et al., 2017)

N<sub>2</sub>O in the atmosphere:

- third most important long-lived anthropogenic greenhouse gas in terms of radiative forcing
- accounts for ~ 7.5 % of the total anthropogenic forcing (IPCC, AR5)
- <u>Global Warming Potential</u> on a 100 years horizon (GWP<sub>100</sub>) is **265** (Myhre et al., 2013)
- nowadays the dominant ozone depleting species (Ravishankara et al., 2009)

Global Concentrations of N<sub>2</sub>O:

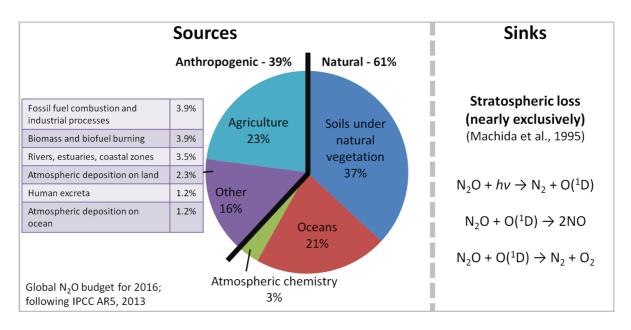


Global concentrations are rising: (https://www.n2olevels.org/) (https://www.n2olevels.org/)

- preindustrial era (i.e. before 1750): 270 ppbv (MacFarling Meure et al., 2006)
- August 2019: **332** *ppbv* (Combined Nitrous Oxide data from the NOAA/ESRL Global Monitoring Division (ftp.cmdl.noaa.gov/hats/n2o/combined/HATS\_global\_N2O.txt); last accessed: 20 Nov 2019)
- current growth: ~ **0.8** *ppbv year*<sup>-1</sup> (WMO, 2011)

#### Lifecycle of N<sub>2</sub>O:

Most important anthropogenic contribution: Agriculture.

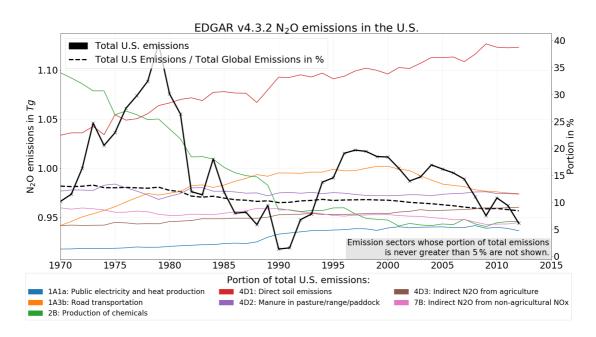


⇒ lifetime: **118** *years* (Prather and Hsu, 2010)

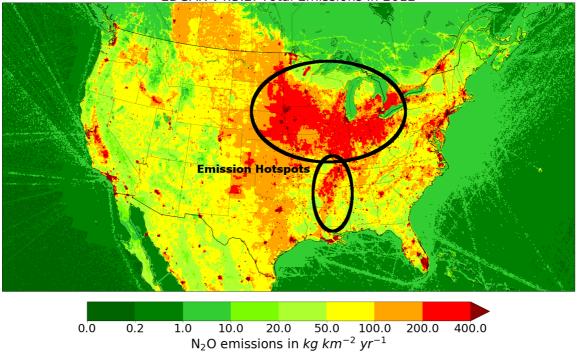
#### N<sub>2</sub>O emissions in the U.S.:

(From the EDGARv4.3.2 (http://edgar.jrc.ec.europa.eu/overview.php?v=432&SECURE=123) dataset ranging from 1970 to 2012)

- approximatelly 9% of the global N<sub>2</sub>O emissions in 2012 were emitted in the U.S.
- agricultural emissions are rising since 1970 (not shown)
- the dominant anthropogenic emission sector is 4D1 (direct agricultural soil emissions)
- in 2012 nearly 40% of the total emissions were 4D1 emissions



• emission hotspots: Cornbelt and Mississippi area, regions of high agricultural activity



EDGAR v4.3.2: Total Emissions in 2012

#### High uncertainties in N<sub>2</sub>O inventories:

- limited amount of top-down studies
- most studies are based on tall tower measurements and Lagrangian models
- common inventories significantly underestimate anthropogenic agricultural emissions

Table: Correction factors foragricultural emissions in theU.S. Midwest for variousemission inventories;parenthesis indicate theinvestigated time period

tall tower measurements + Lagrangian model (STILT+WRF):

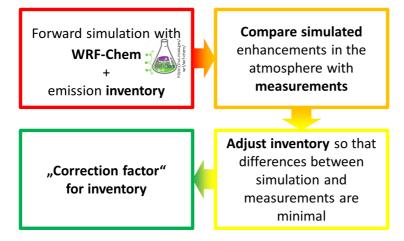
tall tower measurements + Eulerian model (WRF-Chem)

aircraft measurements + Lagrangian model (STILT+WRF)

ne						
	Inventory: Publication:	EDGAR32 FT2000	EDGAR4	EDGAR42	GEIA	DLEM
[	Miller et al., 2012 (2008, June)	5.4	10.1		4.5	2.1
+	Chen et al., 2016 (2010-2011)			1.9 - 4.6		
_	Griffis et al., 2013 (2010)			2.6	8.8	
+	Fu et al., 2017 (2010, June)			19.0 - 28.1		
	Kort et al., 2008 (2003, May-June)	2.62			3.05	
	Xiang et al., 2013 (2012, California)	1.14	1.62		1.62	

## METHOD

Comparable to Barkley et al., 2017:



Lifetime of N<sub>2</sub>O: **118** years (Prather and Hsu, 2010)

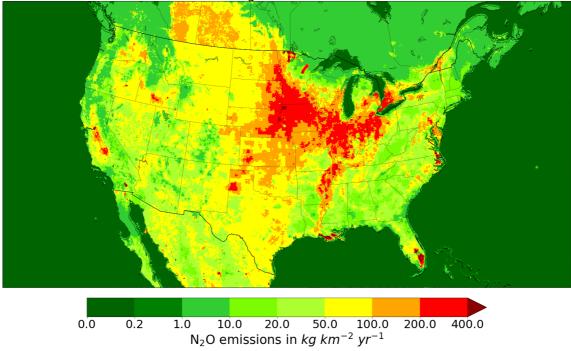
 $\Rightarrow$  N<sub>2</sub>O is handled as **passive tracer** 

#### Inventory:

Only incorporated emission inventory (so far): **EDGAR v4.3.2** (http://edgar.jrc.ec.europa.eu/overview.php?v=432&SECURE=123) (<u>E</u>missions <u>D</u>atabase for <u>G</u>lobal <u>A</u>tmospheric <u>R</u>esearch; https://data.europa.eu/doi/10.2904/JRC\_DATASET\_EDGAR (https://data.europa.eu/doi/10.2904/JRC\_DATASET\_EDGAR); Janssens-Maenhout et al., 2017)



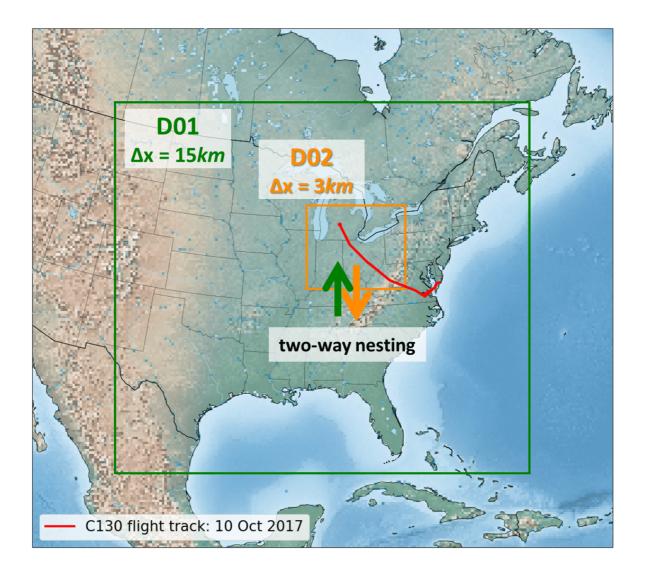
- temporal resolution: yearly (1970-2012) and monthly (2010)
- spatial resolution: 0.1 ° x 0.1 °
- coverage: global



#### EDGAR v4.3.2: Agricultural emissions in 2012

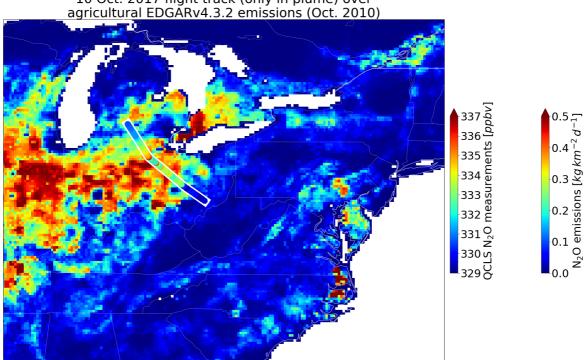
#### WRF-Chem setup:

- version: 4.0.2 (http://www2.mmm.ucar.edu/wrf/users/)
- initial conditions: **ERA5** reanalysis (https://www.ecmwf.int/en/forecasts/datasets/reanalysisdatasets/era5)
  - 30 x 30 *km*
  - 137 vertical layers
  - hourly
- FDDA:
  - **D01**: analysis nudging, surface analysis nudging, obs. nudging
  - D02: obs. nudging
  - observations: NCEP ADP global surface/upper air observations (https://rda.ucar.edu/) + OBSGRID (https://github.com/wrf-model/OBSGRID)
- Chemistry:
  - passive tracer (chem\_opt = 14)
  - emissions: EDGAR (http://edgar.jrc.ec.europa.eu/overview.php?v=432&SECURE=123) + anthro\_emiss (https://www2.acom.ucar.edu/wrf-chem/wrf-chem-tools-community)
- Simulation performance: Comparison of in-flight measurements of meterological parameters (wind in the first place) with corresponding simulated values
- Example domain setup for 10 Oct 2017:



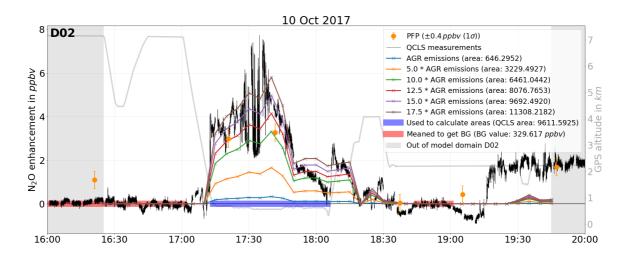
# CASE STUDIES: 2017 & 2019

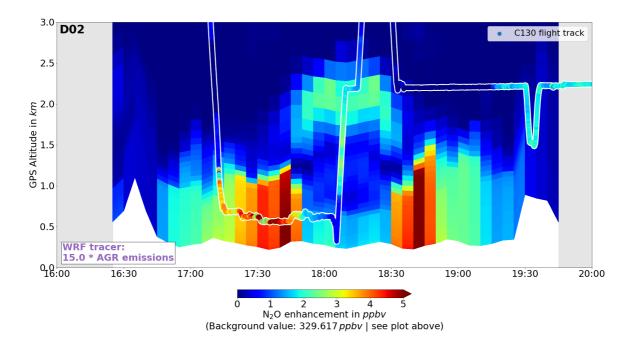
#### **10 Oct 2017:** Strong N<sub>2</sub>O enhancement ( $\leq 6 \text{ ppbv}$ ) in an area of high agricultural activity:



10 Oct. 2017 flight track (only in plume) over agricultural EDGARv4.3.2 emissions (Oct. 2010)

 $\Rightarrow$  Emitting agricultural EDGARv4.3.2 emissions (*AGR emissions*) in WRF:





#### Qualitatively:

- + Simulated plumes spatially coincide with measured N<sub>2</sub>O enhancements
- Simulated enhancements are much too low

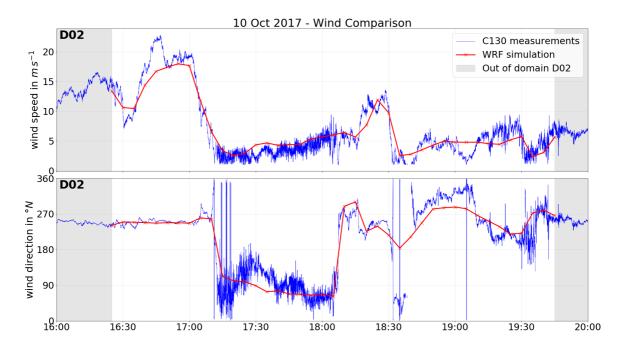
#### Quantitatively:

Increasing strength of emissions by multiplying with factor:

- linear relationship between plume strength and correction factor (compare areas)
- estimated correction factor from linear fit: 14.9

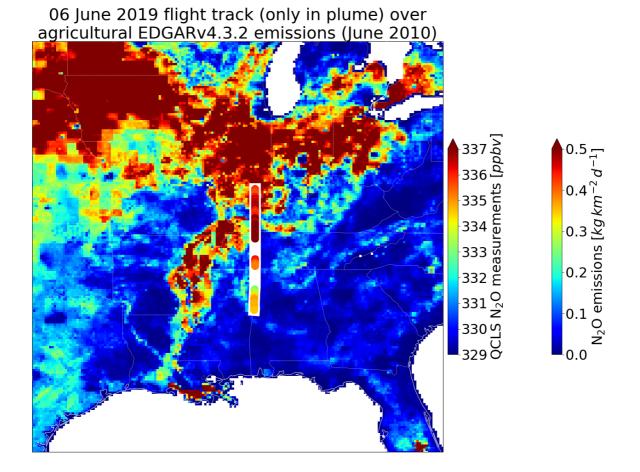
\_\_\_\_

Simulation performance:

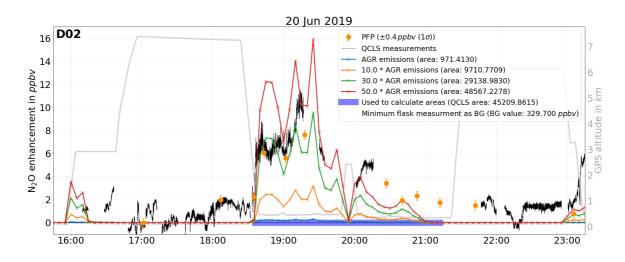


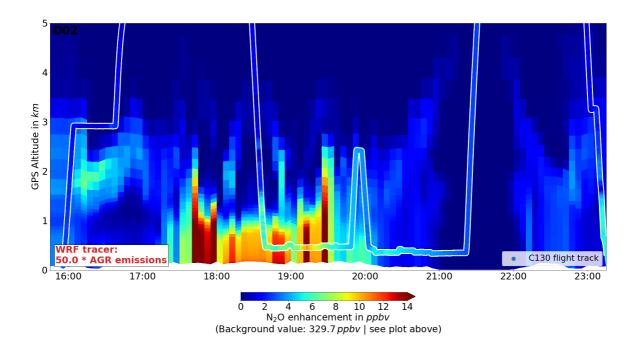
**Good agreement** between onboard wind measurments and model simulations  $\Rightarrow$  The N<sub>2</sub>O transport is assumed to be well represented in the model.

**20 June 2019:** Strong N<sub>2</sub>O enhancement ( $\lesssim$  10 *ppbv*) downwind of the Mississippi area:



 $\Rightarrow$  Emitting agricultural EDGARv4.3.2 emissions (*AGR emissions*) in WRF:





Qualitatively (again, like 10 Oct 2017):

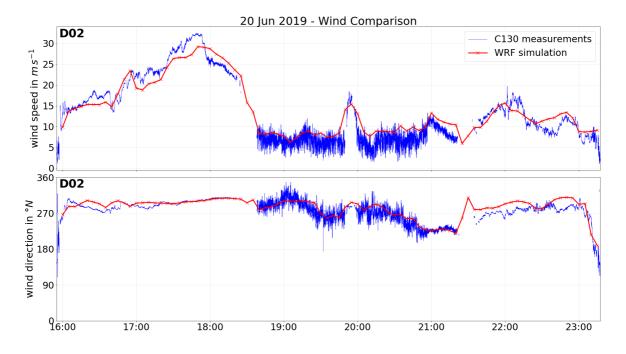
- + Simulated plumes spatially coincide with measured N<sub>2</sub>O enhancements
- Simulated enhancements are much too low

#### Quantitatively:

Increasing strength of emissions by multiplying with factor:

- linear relationship between plume strength and correction factor (compare areas)
- estimated correction factor from linear fit: 46.5
- **BUT:** Flooded Mississippi area most probably influences N<sub>2</sub>O emissions ⇒ **Further analysis** necessary!

Simulation performance:



**Good agreement** between onboard wind measurments and model simulations  $\Rightarrow$  The N<sub>2</sub>O transport is assumed to be well represented in the model.

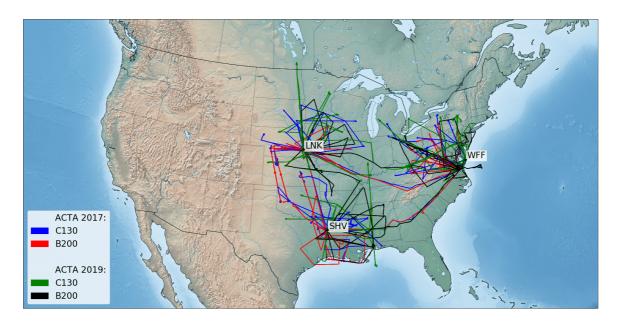
# ACT-AMERICA

(Atmospheric Carbon & Transport - America)

- 2016-2019: five campaigns and all four seasons
- two aircraft: NASA's C130 and B200
- more than 300 joint flight hours in the Midwest
- N<sub>2</sub>O *in-situ* instruments:
  QCLS (DLR) (Kostinek et al., 2019): C130; fall 2017 and summer 2019 (~60 hours of data)
  - Flasks (NOAA): C130 and B200; all five campaigns



C130 and B200 flight tracks during fall 2017 and summer 2019 (continous  $N_2O$  data available (QCLS)):

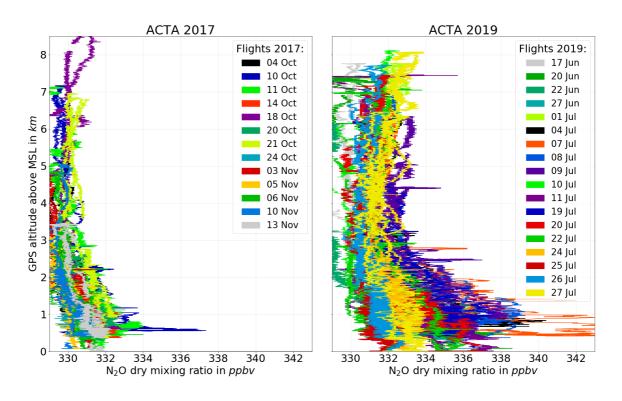


During each campaign the team was stationed for two weeks in:

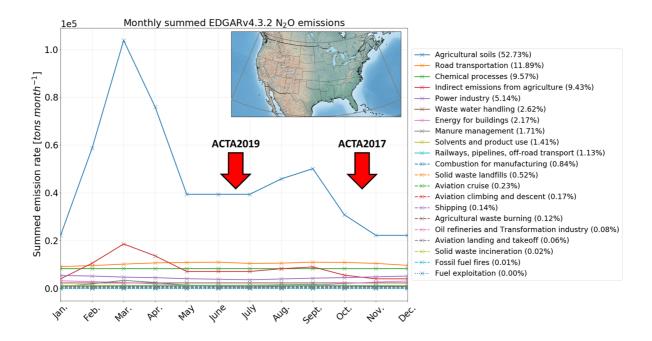
- WFF: Wallops Flight Facility, Virginia
- LNK: Lincoln, Nebraska
- SHV: Shreveport, Louisiana

#### Overview - N<sub>2</sub>O during ACTA 2017 & 2019:

- strong enhancements in the lower troposphere observed
- more and stronger enhancements in summer 2019 than in fall 2017



**Expected emission strengths during ACTA 2017 & 2019:** Throughout the year, anthropogenic N<sub>2</sub>O emissions in the U.S. are dominated by agricultural emissions:



# SUMMARY & OUTLOOK

#### So far:

- Good agreement between measured and modelled  $N_2O$  plume structures
- Strong underestimation of agricultural N<sub>2</sub>O emissions
- Estimated correction factors (so far):
  10 Oct 2017: 14.9
  - 06 June 2019: 46.5 (Flooding not taken into account!!!)

#### Next steps:

- 1. Calculate backward trajectories to clearly determine the origin of the measured air (partly done)
- 2. Apply framework on remaining days
- 3. Simulate remaining ACT-America campaigns (Summer 2016, Winter 2016, Spring 2018) with derived correction factors and compare results to flask measurements
- 4. Investigate different inventories (at best process-based like DAYCENT (https://www2.nrel.colostate.edu/projects/daycent/))

# CV

<sup>2</sup>Ludwig-Maximilians-University (LMU), Meteorological Institute, 80333 Munich, Germany

<sup>3</sup>The Pennsylvania State University, Department of Meteorology and Atmospheric Science, University Park, PA 16802, USA

<sup>4</sup>NOAA ESRL Global Monitoring Division, Boulder, CO 80305-3328, USA

<sup>5</sup>Cooperative Institute for Research in Environmental Sciences, University of Colorado Boulder, Boulder, CO 80503

Every shown dataset of the QCLS is Revision RA.

# ABSTRACT

Atmospheric nitrous oxide (N2O) is, after carbon dioxide and methane, the third most important long-lived anthropogenic greenhouse gas in terms of radiative forcing. Since preindustrial times a rising trend in the global N2O concentrations is observed. Anthropogenic emissions of N2O, mainly from agricultural activity, contribute considerably to this trend. Sparse observational constraints have made it difficult to quantify these emissions. The few studies on top-down approaches in the U.S. that exist are mainly based on Lagrangian models and ground-based measurements. They all propose a significant underestimation of anthropogenic N2O emission sources in established inventories, such as the Emissions Database for Global Atmospheric Research (EDGAR). In this study we quantify anthropogenic N2O emissions in the Midwest of the U.S., an area of high agricultural activity. In the course of the Atmospheric Carbon and Transport – America (ACT-America) campaign spanning from summer 2016 to summer 2019, an extensive dataset over four seasons has been collected including in-situ N2O aircraft based measurements in the lower and middle troposphere onboard NASA's C-130 and B-200 aircraft. During fall 2017 and summer 2019 we conducted measurements onboard the NASA-C130 with a Quantum-Cascade-Laser-Spectrometer (QCLS) and on both aircraft over the whole campaign flask measurements (NOAA) were collected. More than 300 joint flight hours were conducted and more than 500 flask samples were collected over the U.S. Midwest. The QCLS system collected continuous N2O data for approximately 60 flight hours in this region. The Eulerian Weather Research and Forecasting model with chemistry enabled (WRF-Chem) is being used to guantify regional agricultural N2O emissions using the spatial characteristics of these atmospheric N2O mole fraction observations. The numerical simulations enable potential surface emission distributions to be compared to our airborne measurements, and source estimates can be adjusted to minimize the differences, thus quantifying N2O sources. These results are then compared to emission rates in the EDGAR inventory.

# REFERENCES

Fu, C., Lee, X., Griffis, T. J., Dlugokencky, E. J., & Andrews, A. E. (2017). Investigation of the N2O emission strength in the US Corn Belt. Atmospheric research, 194, 66-77.

IPCC, 2013: Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change [Stocker, T.F., D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA, 1535 pp, doi:10.1017/CB09781107415324.

Ravishankara, A. R., Daniel, J. S., & Portmann, R. W. (2009). Nitrous oxide (N2O): the dominant ozone-depleting substance emitted in the 21st century. science, 326(5949), 123-125.

Myhre, G., D. Shindell, F.-M. Bréon, W. Collins, J. Fuglestvedt, J. Huang, D. Koch, J.-F. Lamarque, D. Lee, B. Mendoza, T. Nakajima, A. Robock, G. Stephens, T. Takemura and H. Zhang, 2013: Anthropogenic and Natural Radiative Forcing. In: Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change [Stocker, T.F., D. Qin, G.-K. Plattner, M. Tignor, S.K. Allen, J. Boschung, A. Nauels, Y. Xia, V. Bex and P.M. Midgley (eds.)]. Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA

MacFarling Meure, C., Etheridge, D., Trudinger, C., Steele, P., Langenfelds, R., Van Ommen, T., ... & Elkins, J. (2006). Law Dome CO2, CH4 and N2O ice core records extended to 2000 years BP. Geophysical Research Letters, 33(14).

Machida, T., Nakazawa, T., Fujii, Y., Aoki, S., & Watanabe, O. (1995). Increase in the atmospheric nitrous oxide concentration during the last 250 years. Geophysical Research Letters, 22(21), 2921-2924.

Barkley, Z. R., Lauvaux, T., Davis, K. J., Deng, A., Miles, N. L., Richardson, S. J., ... & Kort, E. A. (2017). Quantifying methane emissions from natural gas production in north-eastern Pennsylvania. Atmospheric Chemistry and Physics (Online), 17(22).

Prather, M. J., & Hsu, J. (2010). Coupling of nitrous oxide and methane by global atmospheric chemistry. Science, 330(6006), 952-954.

Miller, S. M., Kort, E. A., Hirsch, A. I., Dlugokencky, E. J., Andrews, A. E., Xu, X., ... & Wofsy, S. C. (2012). Regional sources of nitrous oxide over the United States: Seasonal variation and spatial distribution. Journal of Geophysical Research: Atmospheres, 117(D6).

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 US Corn Belt using an inverse modeling approach. Global Biogeochemical Cycles, 30(8), 1192-1205.

Kort, E. A., Eluszkiewicz, J., Stephens, B. B., Miller, J. B., Gerbig, C., Nehrkorn, T., ... & Wofsy, S. C. (2008). Emissions of CH4 and N2O over the United States and Canada based on a receptor-oriented modeling framework and COBRA-NA atmospheric observations. Geophysical Research Letters, 35(18).

Janssens-Maenhout, G., Crippa, M., Guizzardi, D., Muntean, M., Schaaf, E., Dentener, F., ... & Aardenne, J. A. V. (2019). EDGAR v4. 3.2 Global Atlas of the three major Greenhouse Gas Emissions for the period 1970–2012. Earth System Science Data, 11(3), 959-1002.

Griffis, T. J., Lee, X., Baker, J. M., Russelle, M. P., Zhang, X., Venterea, R., & Millet, D. B. (2013). Reconciling the differences between top-down and bottom-up estimates of nitrous oxide emissions for the US Corn Belt. Global Biogeochemical Cycles, 27(3), 746-754.

Xiang, B., Miller, S. M., Kort, E. A., Santoni, G. W., Daube, B. C., Commane, R., ... & Nehrkorn, T. (2013). Nitrous oxide (N2O) emissions from California based on 2010 CalNex airborne measurements. Journal of Geophysical Research: Atmospheres, 118(7), 2809-2820.

Kostinek, J., Roiger, A., Davis, K. J., Sweeney, C., DiGangi, J. P., Choi, Y., ... & Klausner, T. (2019). Adaptation and performance assessment of a quantum and interband cascade laser spectrometer for simultaneous airborne in situ observation of CH 4, C 2 H 6, CO 2, CO and N 2 O. Atmospheric Measurement Techniques, 12(3), 1767-1783.