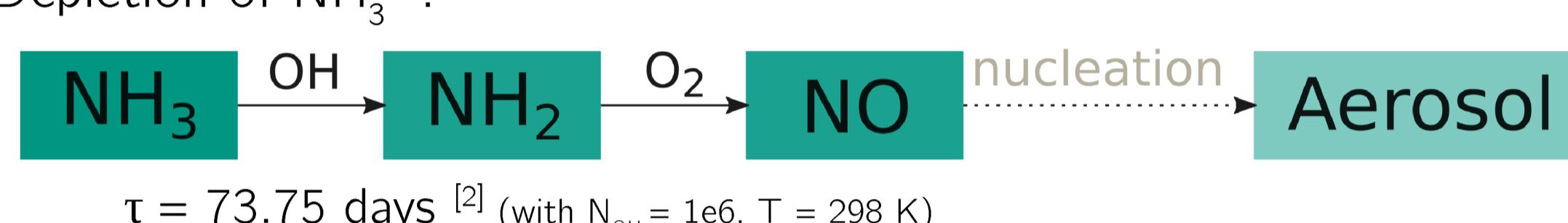


# Investigation of the distribution of aerosol-forming trace gases with ICON-ART

C. Ullwer<sup>1</sup>, R. Ruhnke<sup>1</sup>, M. Höpfner<sup>1</sup>, S. Johansson<sup>1</sup>, J. Schröter<sup>1</sup>, H. Vogel<sup>1</sup>, M. Weimer<sup>2</sup>, S. Werchner<sup>1</sup>, P. Braesicke<sup>1</sup>

## 1. Aerosol-forming trace gases

- Trace gases (e.g. NH<sub>3</sub>) come from different sources (anthropogenic, biogenic, biomass burning). They are oxidized to acids in the atmosphere. Some oxidized trace gases can nucleate (e.g. H<sub>2</sub>SO<sub>4</sub>, HNO<sub>3</sub> and NH<sub>3</sub>).
- Aerosols are important for cloud formation and radiative forcing
- Stratospheric aerosol<sup>[3]</sup>: mainly sulfate (60 – 80 %)
- Tropospheric aerosol<sup>[3]</sup>: Sulfate, nitrate, ammonium, organic and black carbon
- Depletion of NH<sub>3</sub><sup>[1]</sup>:



## 2. ICON-ART

- Global weather and composition model
- Gasphase reactions and aerosol formation
- Model settings:
  - ICON-ART 2.2
  - Horizontal resolution: R2B06 (~ 40 km), Vertical: 90 levels
  - Integration timestep: 300 s
  - Output: regular lat-lon grid: 0.5°

## 3. StratoClim campaign

- Duration: 20.07. - 11.08.2017
- Location: Kathmandu (Nepal)
- 8 successful flights with ‚Geophysica‘ in 21 km altitude (above the asian monsoon)
- Measurements of transported airmasses (UTLS-region) with GLORIA (Gimballed Limb Observer for Radiance Imaging of the Atmosphere; unique imaging Fourier-Transform-Spectrometer)

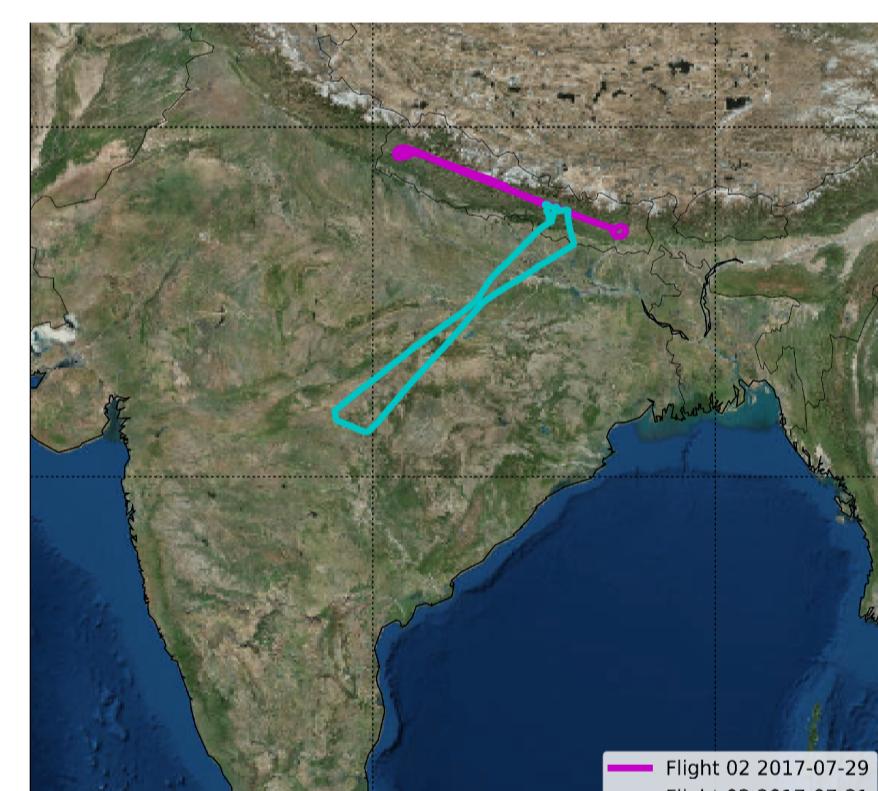
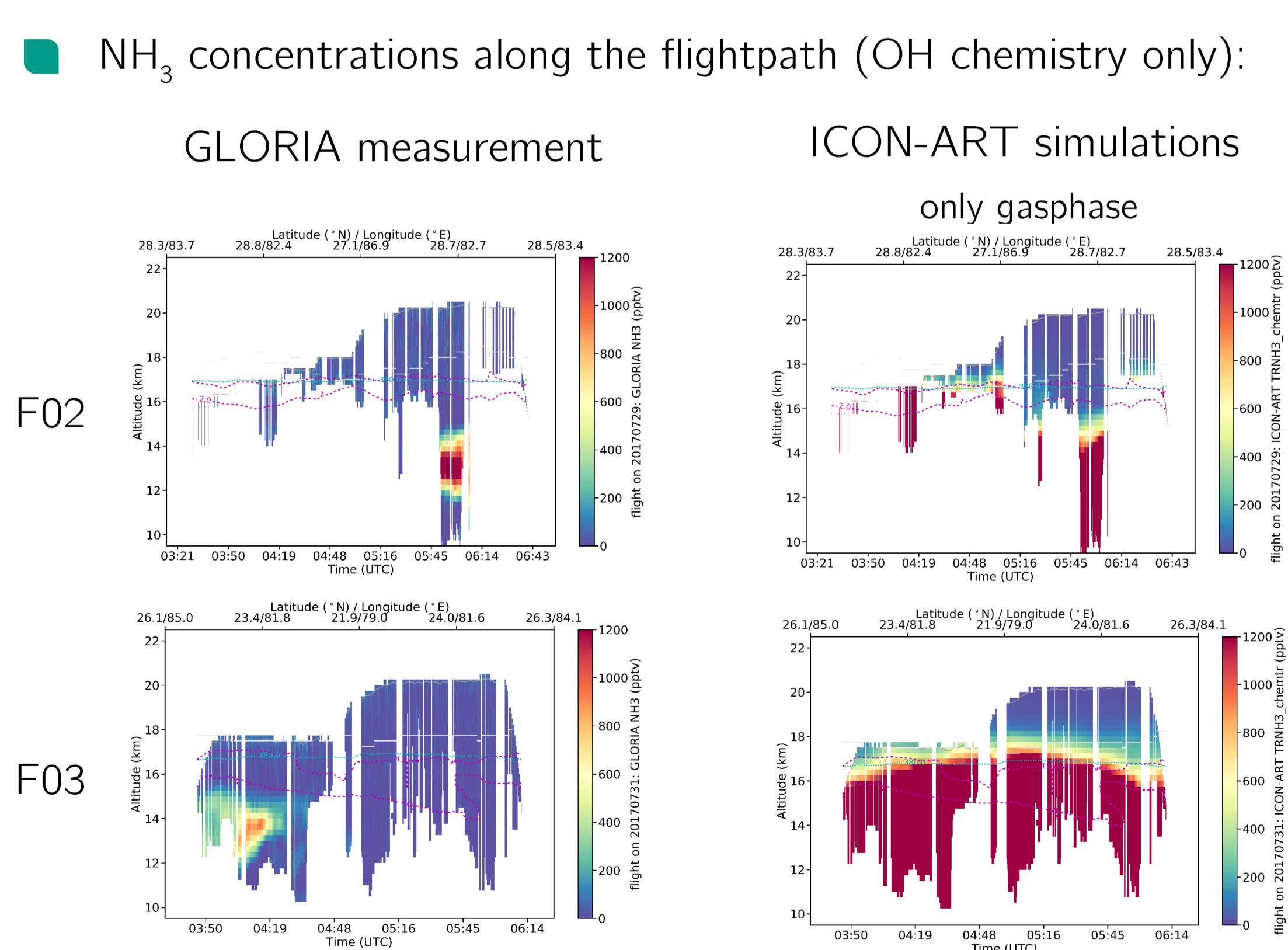


Fig.1: Flightpaths F02 (July 29, 2017) and F03 (July 31, 2017)

## 4. Proof of concept



## 5. Sources of NH<sub>3</sub>

- NH<sub>3</sub> from the MACCity inventory:

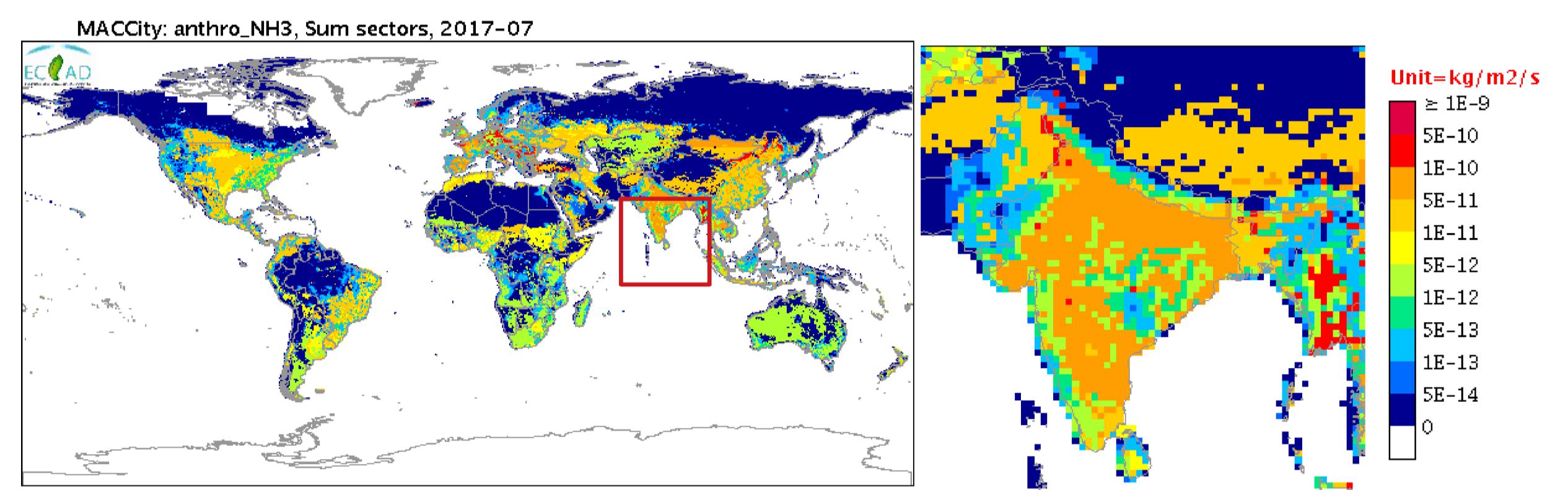


Fig.3: accumulated NH<sub>3</sub> emissions from the ECCAD-database (July 2017) <sup>[4,5,6,7]</sup>

- Emissions by sectors: The main surface source of NH<sub>3</sub> is agriculture:

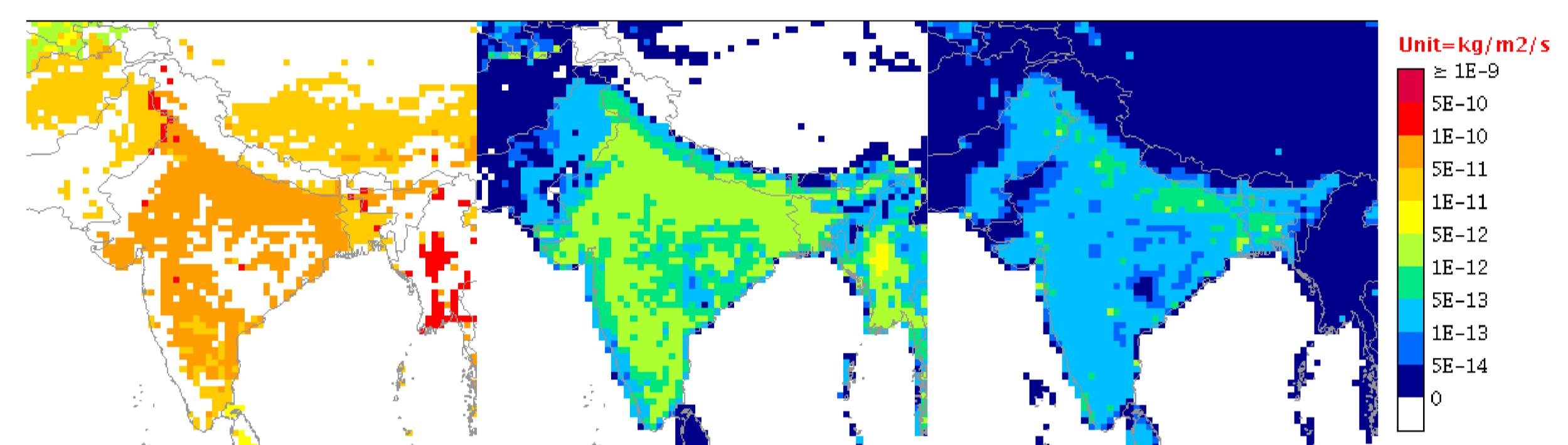


Fig.4: regional NH<sub>3</sub> emissions (July 2017) split up by their sectoral origin provided by the ECCAD-database from left to right: agriculture, agricultural waste, industries <sup>[4,5,6,7]</sup>

## 6. Results

- Accumulated NH<sub>3</sub>:

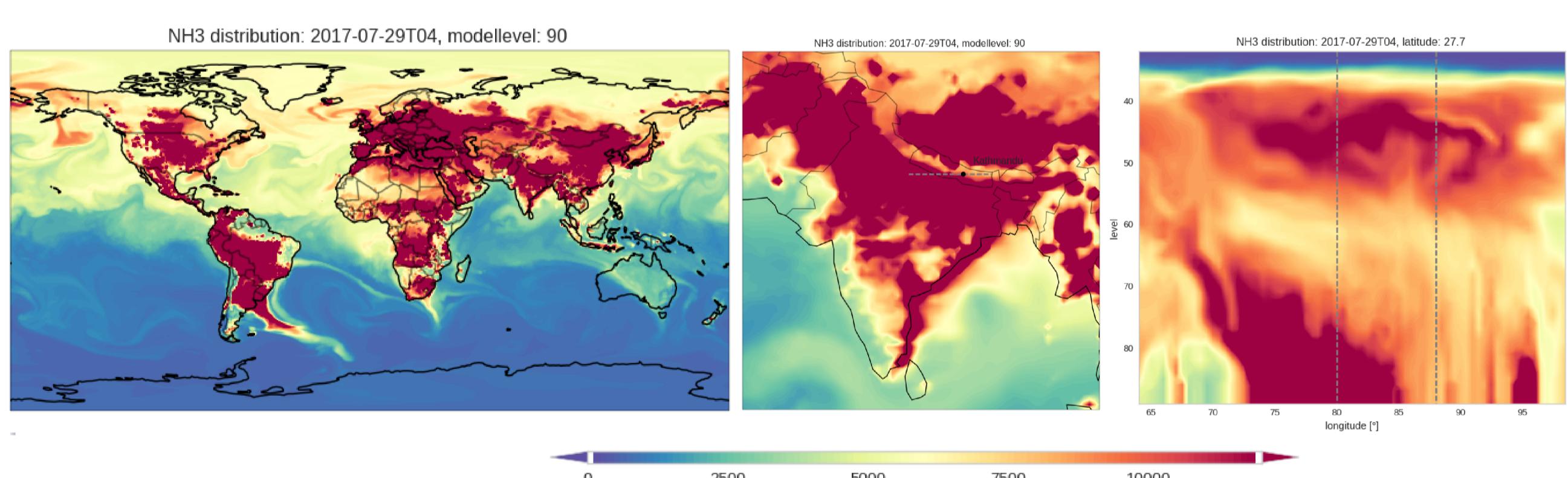


Fig.5: sum of NH<sub>3</sub> emissions from model output; from left to right: global and regional distribution in the lowest model level, vertical cross-section along the dotted line in regional plot

- Sectors with the largest contribution to the total emissions of NH<sub>3</sub>:

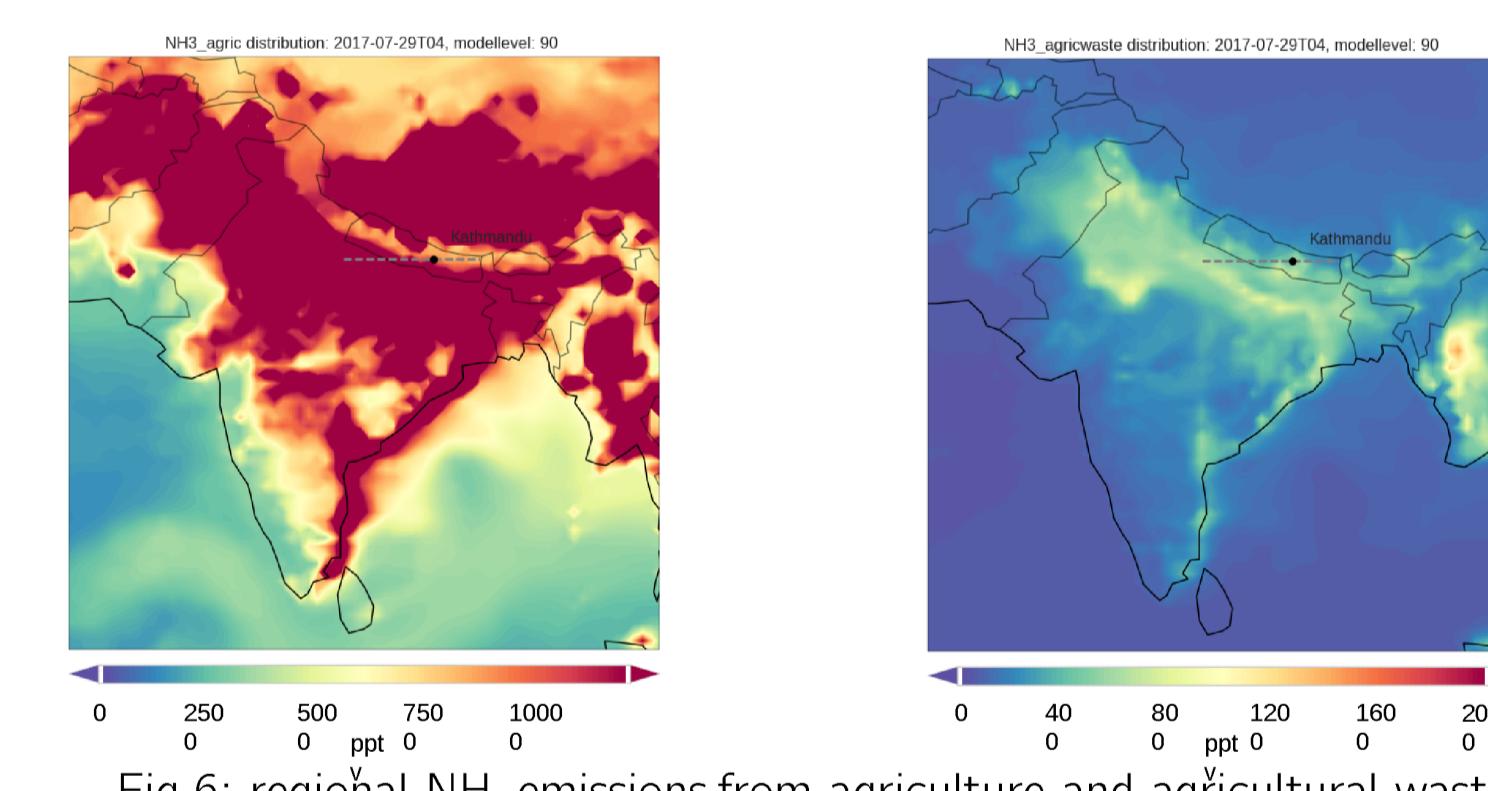


Fig.6: regional NH<sub>3</sub> emissions from agriculture and agricultural waste

## 7. Conclusion and Outlook

- As expected the model run shows too high values in comparison to the measurements, because NH<sub>3</sub> is only reduced by OH, the uptake of NH<sub>3</sub> by the aerosol phase and rainout/washout is yet to be implemented.
- Distinguishing NH<sub>3</sub> sources by sectors helps to understand how transport processes contribute to regional NH<sub>3</sub> concentration differences.
- Improved integrations using a comprehensive gas phase chemistry with coupling to the aerosol module are in preparation.
- Air mass origins will be diagnosed using artificial tracers

[1] D. A. Hauglustaine et al. (2014). *Atmos. Chem. Phys.*, 14, 11031-11063 [2] S. Sander et al. (2015). Evaluation No.18, JPL Publications 15-10 [3] John H. Seinfeld, Spyros N. Pandis. *Atmospheric Chemistry and Physics: From Air Pollution to Climate Change* (3rd Edition) [4] C. Grainger et al. (2011). *Climate Change* 109 (1-2) : 163-190 [5] T. Diehl et al. (2012). *Atmospheric Chemistry and Physics Discussion* 12 : 24895-24954 [6] J.-F. Lamarque et al. (2010). *Atmospheric Chemistry and Physics* 10 : 7017-7039 [7] G.R. van der Werf (2006). *Climate Change* 6 (Atmospheric Chemistry and Physics) : 3423-3441 [8] B. Vogel et al. (2015). *Atmospheric chemistry and physics*, 15, 13699-13716