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RESEARCH ARTICLE

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Key Points:

- Consistency between GOSAT N₂O measurements and LMDz-OR-INCA model N₂O results
- Evidence of a link between summertime Asian N₂O Emissions and the Mediterranean tropospheric N₂O

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Summertime upper tropospheric nitrous oxide over the Mediterranean as a footprint of Asian emissions

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Abstract The aim of this paper is to study the transport of nitrous oxide (N₂O) from the Asian surface to the eastern Mediterranean Basin (MB). We used measurements from the spectrometer Thermal and Near infrared Sensor for carbon Observation Fourier transform spectrometer on board the Greenhouse gases Observing SATellite (GOSAT) over the period of 2010–2013. We also used the outputs from the chemical transport model LMDz-OR-INCA over the same period. By comparing GOSAT upper tropospheric retrievals to aircraft measurements from the High-performance Instrumented Airborne Platform for Environmental Research Pole-to-Pole Observations, we calculated a GOSAT High-performance Instrumented Airborne Platform for Environmental Research standard deviation (SD error) of ~2.0 ppbv for a single pixel and a mean bias of approximately –1.3 ppbv (approximately –0.4%). This SD error is reduced to ~0.1 ppbv when we average the pixels regionally and monthly over the MB. The use of nitrogen fertilizer coupled with high soil humidity during the summer Asian monsoon produces high N₂O emissions, which are transported from Asian surfaces to the eastern MB. This summertime enrichment over the eastern MB produces a maximum in the difference between the eastern and the western MB upper tropospheric N₂O (east-west difference) in July in both the measurements and the model. N₂O over the eastern MB can therefore be considered as a footprint of Asian summertime emissions. However, the peak-to-peak amplitude of the east-west difference observed by GOSAT (~1.4 ± 0.3 ppbv) is larger than that calculated by LMDz-OR-INCA (~0.8 ppbv). This is due to an underestimation of N₂O emissions in the model and to a relatively coarse spatial resolution of the model that tends to underestimate the N₂O accumulation into the Asian monsoon anticyclone.

1. Introduction

Nitrous oxide (N₂O) is produced in oceans and soils by microbial processes of nitrification and denitrification [Butterbach-Bahl *et al.*, 2013]. With a 120 year lifetime, N₂O is the third anthropogenic greenhouse gas after carbon dioxide (CO₂) and methane (CH₄) in terms of radiative forcing [Ciais *et al.*, 2014]. The N₂O radiative forcing contribution accounts for 6–15% of the greenhouse gas emission contribution to global warming. In spite of its far lower concentration in the atmosphere than CO₂, N₂O is a much more efficient greenhouse gas with a global warming potential of 265 over a 100 year time span. Although its main sink is photolysis in the stratosphere, an additional loss of N₂O comes from its reaction with excited atomic oxygen O(¹D) in the stratosphere. This reaction is the main source of stratospheric nitrogen oxides [Minschwaner *et al.*, 1993] that are involved in stratospheric ozone destruction. It is therefore the main emitted ozone-depleting substance of the 21st century [Ravishankara *et al.*, 2009]. Moreover, N₂O volume mixing ratio (vmr) has increased at an average rate of 0.75 ppbv yr⁻¹ since the late 1970s and is expected to increase until 2100 according to the Intergovernmental Panel on Climate Change climatic scenarios due to the increasing use of nitrogen fertilizers in agricultural activities [Ciais *et al.*, 2014]. In Asia, the use of nitrogen fertilizers in agriculture is increasing to meet the growing demand for food production, making China and India the first and second largest consumers and producers of nitrogen fertilizers (<https://www.tfi.org/statistics/statistics-faqs>), respectively.

At a global scale, the strongest N₂O sources are natural soil emissions with an emission rate of 6–7 TgN yr⁻¹, followed by emissions from agricultural sources (4.3–5.8 TgN yr⁻¹) [Syakila and Kroeze, 2011]. Soil moisture is the major driver of N₂O emissions whose maxima occur around 55% soil water content (SWC) [Oswald *et al.*, 2013].

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The second main control variable for N₂O emissions by soils is the nitrogen content of the soil, which increases when fertilizers are added in agro systems [Butterbach-Bahl and Dannenmann, 2011].

Contrary to surface N₂O measurements from the World Meteorological Organization Global Atmosphere Watch Programme [Tsumi *et al.*, 2009], tropospheric N₂O measurements at global or regional scale have neither been widely performed nor widely used up to now like those of CH₄ or CO₂. Remote Fourier transform infrared measurements of total column N₂O are regularly performed within the Network for the Detection of Atmospheric Composition Change [Zander *et al.*, 2005]. There are also sporadic aircraft campaigns including the High-performance Instrumented Airborne Platform for Environmental Research Pole-to-Pole Observations (HIPPO) [Kort *et al.*, 2011] used in this paper. However, satellite observations of tropospheric N₂O profiles and N₂O total columns have been performed by using respectively the Atmospheric Infrared Sounder (AIRS) on board the AQUA platform and the Infrared Atmospheric Sounder Interferometer (IASI) on board the METOP A/B platforms. Consistent interannual trends of N₂O with surface measurements can be derived from AIRS measurements [Xiong *et al.*, 2014], and the African equatorial maximum during March–May can be observed by using IASI total column measurements [Ricaud *et al.*, 2009]. In addition, N₂O is retrieved simultaneously with methane from the thermal infrared measurements of the Greenhouse gases Observing SATellite (GOSAT) at a global scale. Since this study is the first use of GOSAT N₂O data, this data set is introduced in this paper.

In the context of monitoring pollution effects on regional climate, the Chemistry and Aerosol Mediterranean Experiment (ChArMEx) project aims, among other objectives, to understand the spatial and temporal evolutions of chemical compounds over the Mediterranean Basin (MB). In this framework and despite the fact that well-mixed greenhouse gases were not expected to have significant regional variations in the troposphere, a summertime peak in the difference between the eastern and the western MB of middle-to-upper tropospheric CH₄ has been attributed to the transport of pollutants from Asia via the Asian monsoon anticyclone to the eastern MB [Ricaud *et al.*, 2014]. Therefore, variations of long-lived species over the MB can probably be used as footprints of Asian emissions of pollutants. Following this previous work, the aim of this paper is to demonstrate the impact of the Asian N₂O emissions on the upper tropospheric N₂O over the eastern MB. First, we present the N₂O data from the GOSAT Thermal and Near infrared Sensor for carbon Observation Fourier transform spectrometer (TANSO-FTS) together with our validation of GOSAT upper tropospheric N₂O by using the HIPPO aircraft measurements and introduce the chemical transport model LMDz-OR-INCA. Second, we study the N₂O seasonal variability in the eastern and the western MBs as well as the east-west difference of N₂O seasonal variability. Then, we analyze the link between these variabilities and the Asian N₂O emissions via the summer Asian monsoon and its associated anticyclone. Before concluding, we discuss first the correlation between high precipitation rates associated with the monsoon period and high N₂O emissions over Asia. Then, we discuss the influence of horizontal resolution in the model and emission modeling as well as the vertical resolution of measurements on the assessed east-west difference of the MB upper tropospheric N₂O seasonal variability in the model and in the observations, respectively.

2. Data Sets

2.1. GOSAT/TANSO-FTS Data

The Japanese Ministry of the Environment, the Japanese National Institute for Environmental Studies, and the Japan Aerospace eXploration Agency developed the Sun-synchronous orbital satellite GOSAT platform [Hamazaki *et al.*, 2005] dedicated to the study of two greenhouse gases: CO₂ and CH₄. The platform was launched on 23 January 2009 carrying two sensors: a Fourier transform spectrometer (TANSO-FTS) and a Cloud and Aerosol Imager (TANSO-CAI) [Yokota *et al.*, 2009]. The TANSO-FTS is a nadir-viewing instrument using four spectral bands covering the thermal infrared (TIR) and the shortwave infrared domains (0.76, 1.6, 2.0, and 5.5–14.3 μm) [Kuze *et al.*, 2009] with a 10.5 km diameter of instrumental field of view and a spectral resolution of ~0.2 cm⁻¹. Along with CH₄ retrievals in the TIR, N₂O is also processed as a by-product by using a nonlinear optimal estimation retrieval method [Saitoh *et al.*, 2009] from the latest V1 retrieval algorithm [Saitoh *et al.*, 2016]. The retrieval has been performed in clear-sky conditions [Eguchi and Yokota, 2008] in which clouds are filtered by using cloud flags provided by the TANSO-CAI during daytime [Ishida and Nakajima, 2009; Ishida *et al.*, 2011] and on the basis of the TANSO-FTS TIR spectrum during nighttime [Saitoh *et al.*, 2016]. The N₂O-retrieved profiles have been obtained by

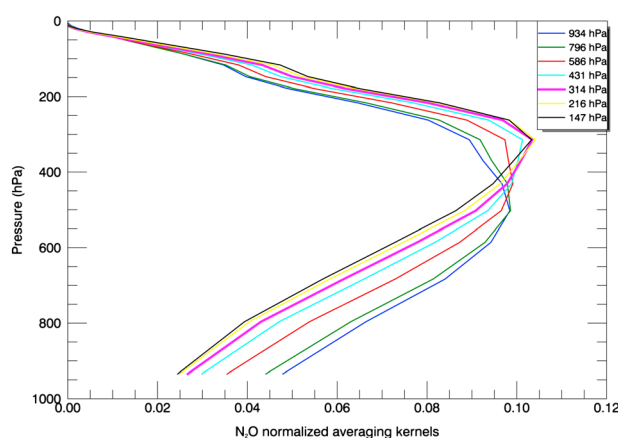


Figure 1. Normalized N_2O averaging kernels from GOSAT/TANSO-FTS at 35°N and 20°E and monthly averaged over July 2010. The different colors represent different retrieval representative levels as shown in the caption.

over the MB. They characterize the sensitivity of GOSAT N_2O measurements at each of the 22 retrieval layers to the true atmospheric N_2O profile [Rodgers, 2000] (only seven of them are represented in Figure 1). The GOSAT N_2O -retrieved profiles reach a maximum of sensitivity between 500 and 200 hPa peaking at the pressure level of 314 hPa. Since we are studying upper tropospheric N_2O transport processes, 314 hPa will be the reference pressure level for our analyses.

Individual retrievals have total theoretical random errors less than 2% (~ 6.4 ppbv for 320 ppbv) at each retrieval level. In this study, we took averages (spatially and temporally) (monthly/interannually) of GOSAT N_2O data (see section 3.1) over the MB at 314 hPa instead of using the individual retrievals. The random errors on the N_2O averaged values are less than 0.1% (~ 0.3 ppbv). TIR measurements are highly affected by surface parameters such as surface emissivity and temperature [Claeyman *et al.*, 2011]. The errors in the surface parameters can impact the N_2O retrieval as systematic errors. Because surface parameters have generally high variability (e.g., diurnal amplitude of surface temperature), these errors may vary in time and space. Thus, to minimize the impact of systematic errors caused by surface parameters, we only used maritime pixels where these parameters exhibit much smaller variations than over land. In addition, consistent with the methodology presented in Ricaud *et al.* [2014] to minimize any systematic errors, we mostly analyze seasonal variability of differences such as anomalies and east-west differences over the MB (see section 3.1), assuming that systematic errors are of the same order of magnitude in the eastern and the western MBs.

Since we are interested in the N_2O variability, we determined the anomaly from its respective mean of all N_2O data over the latitude band 15°N – 60°N for each year over the period of 2010–2013. This approach has the main advantage of facilitating the comparison between model results and measurements.

2.2. GOSAT N_2O Data Validation

In addition to the theoretical errors analysis presented in section 2.1, we used the HIPPO airborne campaigns to assess the quality of the GOSAT N_2O retrievals. The HIPPO campaigns performed in situ measurements over the Pacific Ocean by using the National Science Foundation's Gulfstream V aircraft. Several aerosols and chemical species including N_2O were measured over a wide range of latitudes (67°S – 87°N) and altitudes (600–15,000 m) with vertical profiles every 2.2° of latitude [Wofsy, 2011, Wofsy *et al.*, 2012]. HIPPO encompasses five campaigns: HIPPO 1 (January 2009), HIPPO 2 (October–November 2009), HIPPO 3 (March–April 2010), HIPPO 4 (June–July 2011), and HIPPO 5 (August–September 2011). Since our GOSAT data cover the period of 2010–2013, we used the HIPPO campaigns 3–5 to assess the quality of the GOSAT N_2O retrievals (Figure 2). For our comparison, we used the measurements from the Harvard/Aerodyne Quantum Cascade Laser Spectrometer, one of the airborne instruments of HIPPO. This instrument has a 1σ precision of 0.09 ppbv and an accuracy of 0.2 ppbv [Kort *et al.*, 2011].

Each vertical profile from HIPPO is identified by its mean coordinates (latitude, longitude, and time) and is interpolated on the 22 retrieval representative levels of GOSAT. Since HIPPO profiles do not cover the

assuming a fixed a priori profile derived from the Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) reference atmosphere (V3) climatology from 15°N to 60°N latitude (MIPAS reference atmosphere profiles can be downloaded from <http://www.atm.ox.ac.uk/RFM/atm/>).

GOSAT N_2O retrievals are performed on 22 retrieval grid layers, and each grid layer has a representative pressure level. The normalized averaging kernels shown in Figure 1 have been calculated at 35°N and 20°E and averaged over July 2010. They are representative of the MB since the N_2O averaging kernels do not have significant spatial variations

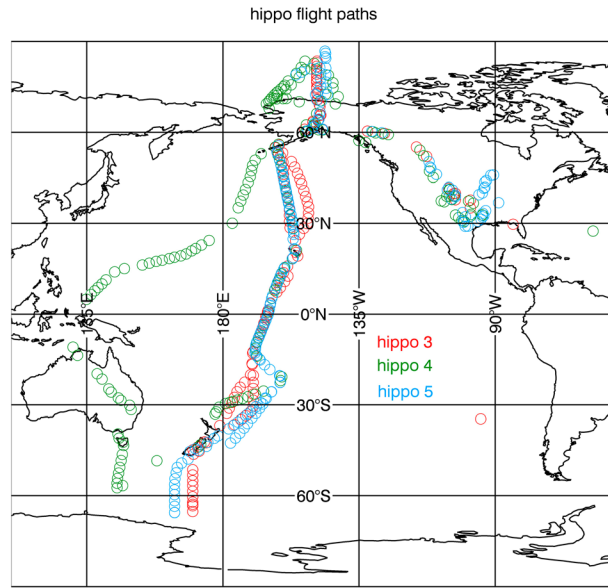


Figure 2. Flight paths of HIPPO campaigns used for GOSAT N₂O validation: HIPPO 3 (March–April 2010, red), HIPPO 4 (June–July 2011, green), and HIPPO 5 (August–September 2011, blue).

the identity matrix, x_a the a priori profile (the same used for GOSAT retrievals), x the HIPPO profile, and \hat{x} the HIPPO convolved profile. Each HIPPO convolved profile is then compared to the GOSAT measurements in time and space coincidence considering GOSAT maritime profiles within a distance of ± 300 km of the HIPPO profile and a temporal window of ± 12 h. The standard deviation of the HIPPO N₂O data selected from all of the three campaigns at 314 hPa is ~ 3.3 ppbv. The standard deviation of the convolved HIPPO N₂O data at 314 hPa is ~ 2.0 ppbv. Therefore, the vertical sensibility of the GOSAT instrument represented by the averaging kernels tends to underestimate the variability of N₂O at 314 hPa.

Figure 3 shows the comparison between the GOSAT N₂O data retrievals and the HIPPO N₂O at 314 hPa by using 200 coincident pairs with our selection criteria. $P_0 = 318.4$ ppbv is the intersection point between the regression line and the first bisector (the line $y = x$). The Pearson linear correlation coefficient is about 0.44. The slope of the regression line is about ~ 0.22 ppbv ppbv⁻¹, which means that the GOSAT N₂O variations are smaller than HIPPO N₂O variations at 314 hPa. GOSAT data underestimate N₂O for volume mixing ratio (vmr) larger than P_0 and conversely for vmr smaller than P_0 . This confirms the fact that, in general, the GOSAT retrieval system tends to reduce the amplitude of N₂O variations. Since N₂O is retrieved simultaneously with CH₄ in the GOSAT retrieval system, a strong constraint has been applied on N₂O in order to avoid large oscillations that could impact the CH₄ retrieval product. This strong constraint artificially reduces the GOSAT N₂O variability at 314 hPa.

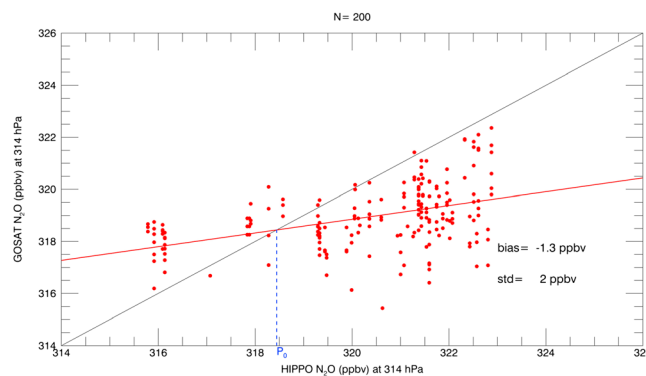


Figure 3. HIPPO (3–5) N₂O airborne measurements versus GOSAT N₂O at 314 hPa in time and space coincidence within a box of ± 300 km and ± 12 h. $N (=200)$ is the total number of coincident pairs selected. The black and red lines represent the first bisector ($y = x$) and the linear regression line, respectively. The dashed blue line represents the intersection point ($P_0 = 318.4$ ppbv) of the regression line and the first bisector. Note that we applied the GOSAT averaging kernels on HIPPO observations.

entire vertical range of GOSAT retrievals, we extended the HIPPO profiles to the bottom and to the top of the atmosphere by using monthly averaged profiles from LMDz-OR-INCA. However, to minimize the impact of such artificial extension on our validation results and since we are interested in the 314 hPa level, we only took HIPPO profiles with a ceiling pressure of less than 250 hPa and a bottom pressure level of greater than 400 hPa.

To evaluate only the retrieval system and not the instrumental performance of the TANSO-FTS sounder, we applied the GOSAT N₂O averaging kernel to the HIPPO measurements by using the following equation [Rodgers, 2000]:

$$\hat{x} = Ax + (I - A)x_a \quad (1)$$

where A is the averaging kernel matrix, I the identity matrix, x_a the a priori profile (the same used for GOSAT retrievals), x the HIPPO profile, and \hat{x} the HIPPO convolved profile. Each HIPPO convolved profile is then compared to the GOSAT measurements in time and space coincidence considering GOSAT maritime profiles within a distance of ± 300 km of the HIPPO profile and a temporal window of ± 12 h. The standard deviation of the HIPPO N₂O data selected from all of the three campaigns at 314 hPa is ~ 3.3 ppbv. The standard deviation of the convolved HIPPO N₂O data at 314 hPa is ~ 2.0 ppbv. Therefore, the vertical sensibility of the GOSAT instrument represented by the averaging kernels tends to underestimate the variability of N₂O at 314 hPa.

The mean N₂O bias (GOSAT-HIPPO) at 314 hPa is about -1.3 ppbv ($\sim 0.4\%$) because tropospheric N₂O vmr are mostly greater than P_0 . Furthermore, since HIPPO measurements have a very good precision ($\sim 0.03\%$ at 320 ppbv), the standard deviation of the

GOSAT-HIPPO differences (called hereafter SD error), which is about 2.0 ppbv (~0.6%), can be considered as an estimation of the real random error on an individual GOSAT measurement. This value is small and much less than the theoretical value of 6.4 ppbv (~2%) (see section 2.1) associated with the total random error on a single retrieval.

To conclude, the comparisons with HIPPO data show that the GOSAT N₂O maritime pixels at 314 hPa are of sufficient quality to be used in order to analyze regional and seasonal N₂O variabilities. However, we have to keep in mind that the GOSAT retrieval system tends to underestimate the amplitude of the upper tropospheric N₂O variations. Furthermore, averaging spatially and temporally (between 400 and 900 profiles) over the eastern and the western MBs results in a very small random errors: less than ~0.3 ppbv (~0.09%) for the theoretical error and ~0.1 ppbv for the SD error. Thus, we also have to keep in mind that the error values used in our analysis of N₂O seasonal variabilities are calculated by using the theoretical total random errors which are much more pessimistic than the SD error from the comparison with HIPPO measurements.

2.3. LMDz-OR-INCA

The Interaction between Chemistry and Aerosol (INCA) model coupled with the general circulation model LMDz GCM is used to simulate the distribution of aerosols and gaseous reactive species in the troposphere [Hauglustaine *et al.*, 2004]. In addition, soil/atmosphere exchanges of water and energy are simulated by coupling LMDz with the Organizing Carbon and Hydrology in Dynamic Ecosystems dynamic global vegetation model [Krinner *et al.*, 2005]. Together, these three models form the LMDz-OR-INCA model. The model has a horizontal resolution of 1.26° in latitude and 2.5° in longitude and 39 vertical levels from the surface up to 80 km. The N₂O surface emissions are monthly averages based on Thompson *et al.* [2014] for anthropogenic and natural emissions and on the Global Fire Emissions Database version 4.1 (GFEDv4.1) inventory for biomass burning [Giglio *et al.*, 2013]. The total annual emission rate is 16.08 TgN yr⁻¹ in our simulations. Our model is forced dynamically by the meteorological analysis from the European Centre for Medium-Range Weather Forecasts. We used monthly averaged N₂O outputs covering the period of 2010–2013 together with GOSAT retrievals to assess the impact of Asian summertime emissions on eastern MB upper tropospheric N₂O.

However, in order to assess the impact of improving the horizontal resolution and the surface emission inventories on the modeled N₂O transport, we used in section 4 another configuration of LMDz-OR-INCA referred to as LMDz-OR-INCA low resolution (LMDz-OR-INCA *low res*). LMDz-OR-INCA *low res* has been performed for climate change impact investigations by using different Representative Concentration Pathway scenarios [Hauglustaine *et al.*, 2014]. The horizontal resolution in this configuration is 1.9° in latitude and 3.75° in longitude. This configuration used monthly averaged N₂O surface emissions based on Bouwman and Taylor [1996] for anthropogenic and natural emissions and on the Global Fire Emissions Database version 3 inventory for biomass burning [Van der Werf *et al.*, 2010]. The total annual emission rate in this configuration is 13.27 TgN yr⁻¹. The vertical levels are the same in the two configurations.

3. Results

3.1. N₂O Variability Over the Mediterranean Basin

GOSAT and LMDz-OR-INCA anomalies have been calculated for each year from their annual mean over the latitude band 15°N–60°N. These anomalies are averaged monthly from 2010 to 2013 and selected within two boxes (36–45°N, 1–12°E) and (30–37°N, 26–37°E) to represent the western and eastern MBs (black boxes in Figure 4), respectively. Figure 4 shows the N₂O anomaly fields in summer (June, July, and August) and winter (December, January, and February) at 314 hPa observed by GOSAT and calculated by LMDz-OR-INCA. In winter, a very weak positive east-west N₂O difference is measured by GOSAT (~0.7 ppbv) and calculated by the model (~0.15 ppbv) with a larger anomaly over the eastern MB.

In summer, this gradient is enhanced within the two data sets, although it remains less intense in the model (~0.5 ppbv) compared to GOSAT (~1.5 ppbv). Figure 5 shows the seasonal variation of the N₂O anomaly observed by GOSAT and calculated by LMDz-OR-INCA for the two basins. In the GOSAT data, a seasonal cycle is observed in the eastern MB with a maximum in July and a peak-to-peak amplitude of ~1.8 ± 0.4 ppbv, whereas in the western MB, a maximum with a peak-to-peak amplitude of ~0.8 ± 0.4 ppbv is observed in August. The GOSAT peak in July over the eastern MB is sharper than the peak in August over the western

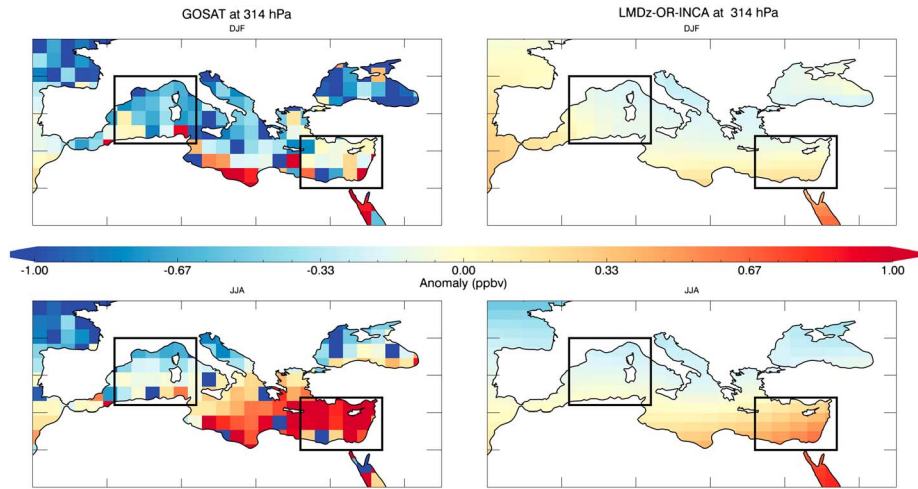


Figure 4. N₂O volume mixing ratio anomaly (ppbv) from (left) GOSAT TANSO-FTS (right) and LMDz-OR-INCA at 314 hPa for (top) December-January-February (DJF) and for (bottom) June-July-August (JJA) averaged over the period of 2010–2013. The satellite measurements and the model outputs are selected within the black boxes to represent the eastern and the western Mediterranean Basins.

MB. The error bars represent the random total errors on monthly averaged N₂O retrievals and indicate the confidence level on the mean value. Despite the small values of the seasonal amplitude, the seasonal cycle is statistically significant regarding measurements errors. Consistently with GOSAT anomalies, a seasonal cycle is also detected in LMDz-OR-INCA data but with a maximum from August to November in the eastern MB and from September to December in the western MB (peak-to-peak amplitude of ~1.5 ppbv in the two basins).

Figure 6 shows the N₂O difference between the eastern and the western MBs at 314 hPa called hereafter east-west difference. In GOSAT data, the monthly averaged east-west difference has no significant variations ($\sim 0.4 \pm 0.8$ ppbv) from January to May and starts to increase in June reaching a peak-to-peak amplitude of $\sim 1.4 \pm 0.3$ ppbv in July before decreasing until November. The east-west difference in LMDz-OR-INCA follows almost the same seasonal variation but with a peak-to-peak amplitude less intense than in GOSAT (~ 0.8 ppbv). Although the two data sets are consistent with each other in terms of the seasonal variation of the east-west difference, the amplitude of the seasonal variations and the absolute value of the difference show inconsistencies. These differences are discussed in section 4. We note that the maximum of the east-west difference occurs in July in both the model results and in the GOSAT observations. The processes

that govern this summertime N₂O increase over the eastern MB are investigated in the next section.

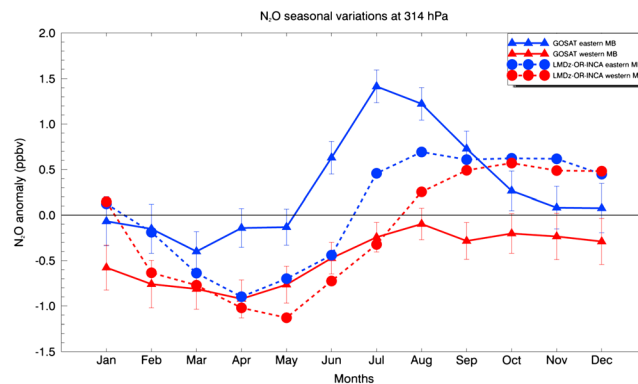


Figure 5. Seasonal variation of the monthly averaged N₂O anomalies (ppbv) from GOSAT TANSO-FTS (solid lines and triangles) and LMDz-OR-INCA (dashed lines and circles) at 314 hPa on the eastern (blue) and western (red) Mediterranean Basins averaged over the period of 2010–2013. The error bars represent the error on the mean considering the theoretical random error of the retrieval and the number of monthly averaged pixels.

3.2. Impact of Asian Summer Monsoon

Due to its particular location, the MB is influenced by several meteorological patterns and air masses originating from Europe, North Africa, the Arabian peninsula, North America, and Asia [Ricaud et al., 2014]. Figure 7 shows the LMDz-OR-INCA northern hemisphere N₂O anomaly and wind patterns representative of the months of July and August (bottom) and the other months (top) over the period of 2010–2013 at 314 hPa.

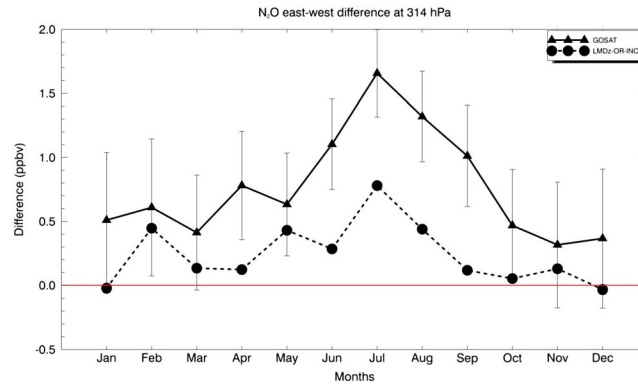


Figure 6. Seasonal variation of the east-west difference of N₂O (ppbv) measured by GOSAT TANSO-FTS (solid line) and calculated by LMDz-OR-INCA (dashed line) at 314 hPa. The error bars represent the error on the mean considering the theoretical random error of the retrieval and the number of monthly averaged pixels.

Upper tropospheric air masses over the eastern and western MBs mainly come from North America and the northern Atlantic throughout the year. But in summer, over the eastern MB, upper tropospheric air masses are mostly transported from Asia through the Asian monsoon anticyclone [Scheeren *et al.*, 2003; Rodwell and Hoskins, 1996] and are also influenced by the Arabian Peninsula and North Africa [Dayan, 1986; Tyrllis *et al.*, 2013].

The impact of long-range transport of air masses from Asia on the MB explains the maximum in the east-west CH₄ difference in the upper tro-

posphere observed in August [Ricaud *et al.*, 2014]. Westerly circulation brings air masses from the northern Atlantic and North America from January until May. At the beginning of the Asian monsoon period in June, a southerly circulation takes place over the eastern MB, bringing tropical air masses from Asia to the eastern MB. These air masses are enriched in pollutants trapped by the monsoon anticyclone. Thus, pollutants emitted in Asia are transported to the upper troposphere by updrafts associated with the monsoon region and redistributed westward to the eastern MB by the monsoon anticyclone. In the next section, we investigate the different regions of N₂O emissions in Asia.

3.3. Asian N₂O Surface Emissions

The eastern MB air masses come from the Asian anticyclone in summer and are therefore impacted by the N₂O surface emissions from different Asian regions. Figure 8 synthesizes the pressure levels of air masses from 10 day back trajectories covering the period of 2001–2011 (two trajectories per day at 00:00 and 12:00 UTC) in July and in August from the point located at 33°N, 35°E and 300 hPa in the eastern MB. This back trajectory climatology has been performed by using the British Atmospheric Data Centre trajectory service

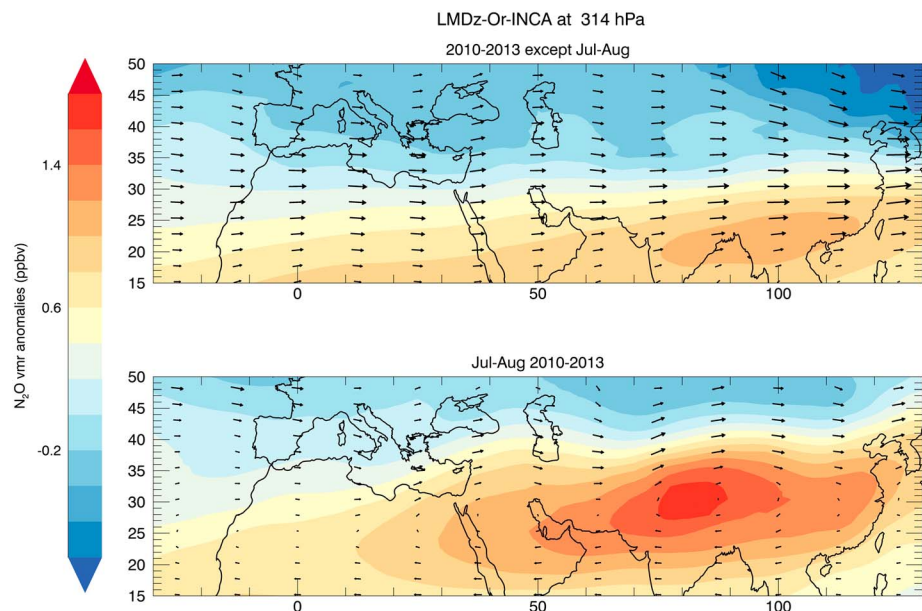


Figure 7. N₂O volume mixing ratio anomaly (ppbv) from LMDz-OR-INCA and horizontal wind from IFS at 314 hPa averaged over (bottom) July and August 2010–2013 and over (top) the other months.

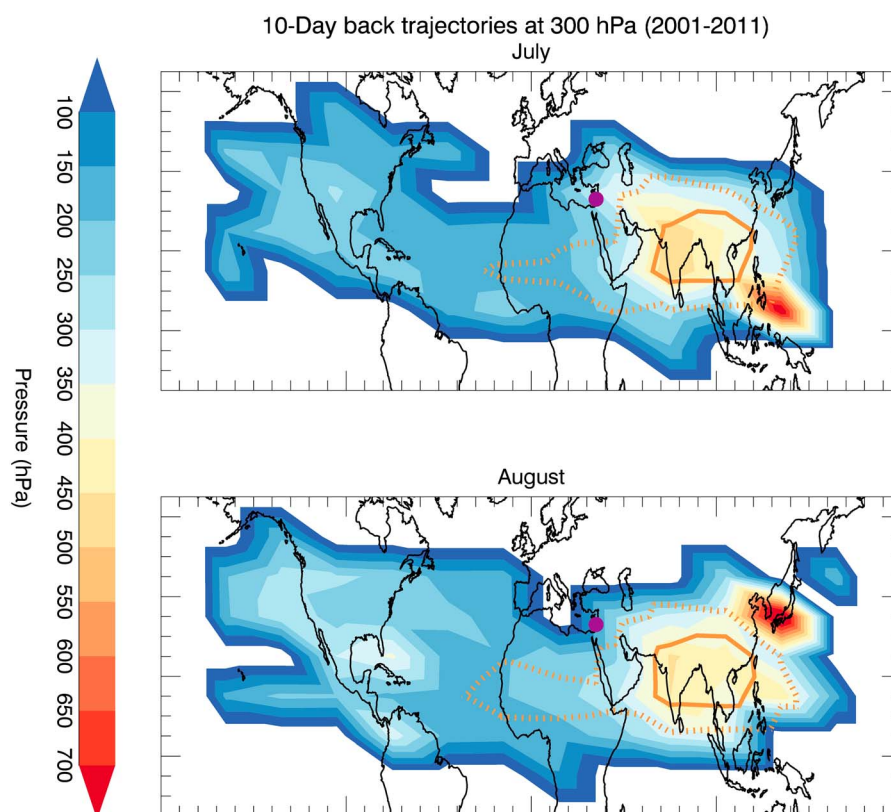


Figure 8. Pressure levels of air masses after 10 day back trajectories from the point located at 33°N, 35°E (purple filled circle) and 300 hPa in the eastern MB from the British Atmospheric Data Centre trajectory service (<http://badc.nerc.ac.uk/community/trajectory/>) in (top) July and in (bottom) August over the period of 2001–2011. The contour lines superimposed to the maps delimit the percentage of air masses reaching the eastern MB at 314 hPa after 10 days of transport: 55% (solid orange line) and 80% (dotted orange line).

(<http://badc.nerc.ac.uk/community/trajectory/>). The contour lines superimposed to the maps delimit the percentage of air masses reaching the eastern MB at 314 hPa after a transport of 10 days: 55% (solid orange line) and 80% (dotted orange line) of all trajectories (682 trajectories). Since 6 day back trajectories were sufficient to highlight the impact of the Indian subcontinent air masses on the MB [Ricaud *et al.*, 2014], we used 10 day back trajectories to detect the influence of planetary boundary layer sources.

The eastern MB air masses during summer come mostly from southeastern Asian and the Indian/Tibetan Plateau regions in July and August, as shown by the contour lines in Figure 8. However, air masses from the Indian subcontinent come from lower levels (500–550 hPa), whereas those from southeastern Asia come from upper levels (350–400 hPa). The so-called Asian monsoon anticyclone is actually an upper level divergence zone linked to the monsoon convective system and causes strong updrafts in the troposphere [Randel and Park, 2006]. Thus, air masses that have their origins in the Indian subcontinent are likely influenced by the boundary layer in the short term (less than 1 month) via the updrafts associated with the upper level divergence zone (referred to as monsoon anticyclone). Figure 9 represents the monthly averaged Asian N₂O emission fluxes used in LMDz-OR-INCA simulations superimposed with the contour lines in Figure 8. The peak of Asian N₂O emissions occurs in July, consistent with the peak of the east-west difference measured by GOSAT and calculated by LMDz-OR-INCA (Figure 6). There are two main regions of emissions: in northern India and in eastern China. The main emission region from eastern China is outside, while the northern Indian region is well inside the contour lines delimiting the main regions of origin of eastern MB air masses. This confirms the fact that upper tropospheric N₂O over the eastern MB in July and August comes mostly from Indian subcontinent N₂O emissions. This hypothesis is consistent with previous work on the Asian monsoon anticyclone composition in July and in August. First, Vogel *et al.* [2015] show that, in July 2012, the contribution of boundary layer sources to the Asian monsoon composition is between 35% and 55% and the

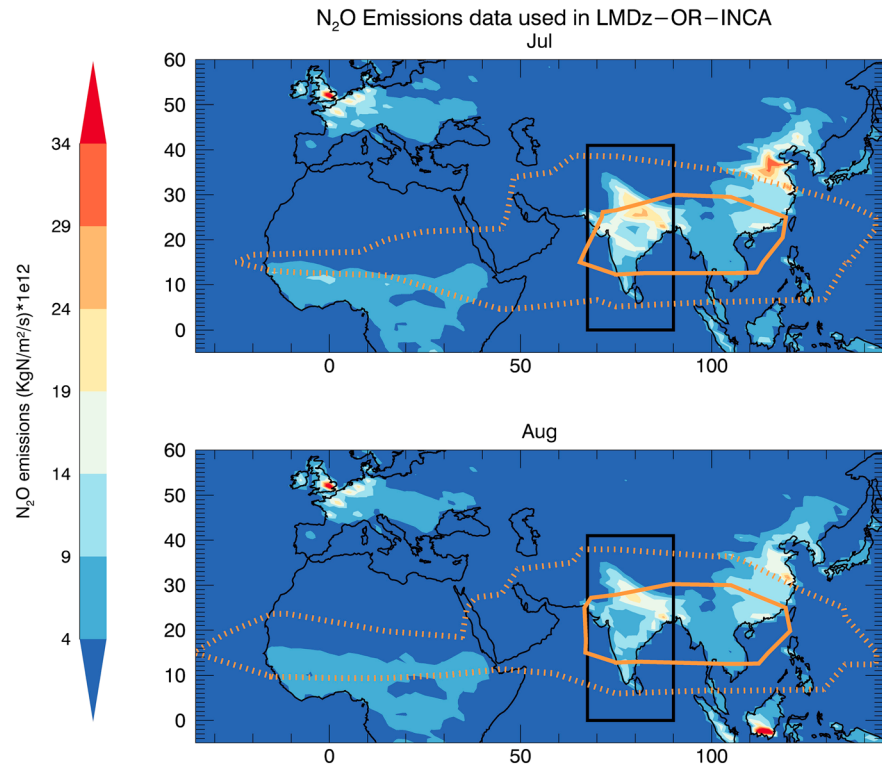


Figure 9. Monthly averaged Asian N_2O surface emission fluxes ($kg\ m^{-2}\ s^{-1}$) used in LMDz-OR-INCA simulations in (top) July and in (bottom) August. The black rectangle ($0\text{--}41^\circ N$, $67.5\text{--}90^\circ E$) delimits the Indian subcontinent N_2O emission region. The contour lines (see Figure 8) delimit the percentage of air masses reaching the eastern MB at 314 hPa after 10 days of transport: 55% (solid orange line) and 80% (dotted orange line).

contributions of the Indian/Tibetan Plateau region and eastern China are 24% and 10%, respectively. Consequently, in July, the Indian/Tibetan Plateau and eastern China represent 43–68% and 18–28% of the contribution from all boundary layer sources to the Asian monsoon anticyclone, respectively. Second, Bergman *et al.* [2013] show that, in August 2011, the main contributions from the boundary layer sources to the Asian monsoon anticyclone composition are from India/Southeast Asia and the Tibetan Plateau regions (70–80%).

In summary, we showed the predominant contribution of the Indian subcontinent N_2O emissions on the composition of upper tropospheric N_2O over the eastern MB in July and in August. In the next section, we

explain the biochemical processes involved in these N_2O emissions.

3.4. Emission Processes

The east-west difference of N_2O from GOSAT and LMDz-OR-INCA exhibits a maximum in summer peaking in July, namely, 1 month earlier than the occurrence of a CH_4 maximum analyzed by Ricaud *et al.* [2014]. This is due to environmental parameters such as soil moisture that influence N_2O emissions. Soil ability to emit N_2O can be characterized by analyzing the nitrification and the denitrification processes [Qu *et al.*, 2014]. Since intensified agriculture, characterized by high nitrogen fertilizer

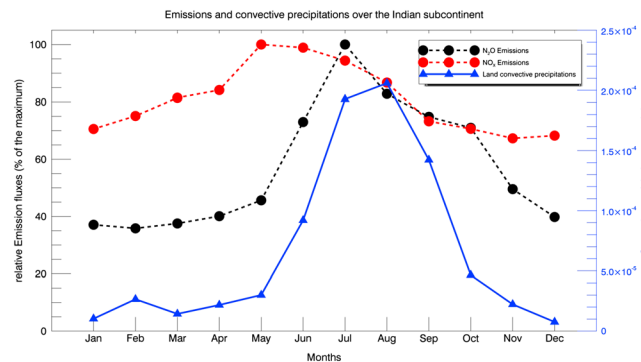


Figure 10. Seasonal variation of monthly averaged N_2O (black) and NO_x (red) relative to surface emission fluxes (emission fluxes divided by the maximum fluxes) used in LMDz-OR-INCA and the convective precipitation amount from the reanalysis ERA-Interim (blue) averaged over the Indian subcontinent (land pixels within the black box of Figure 9). The convective precipitation data are monthly averaged over the period of 2010–2013.

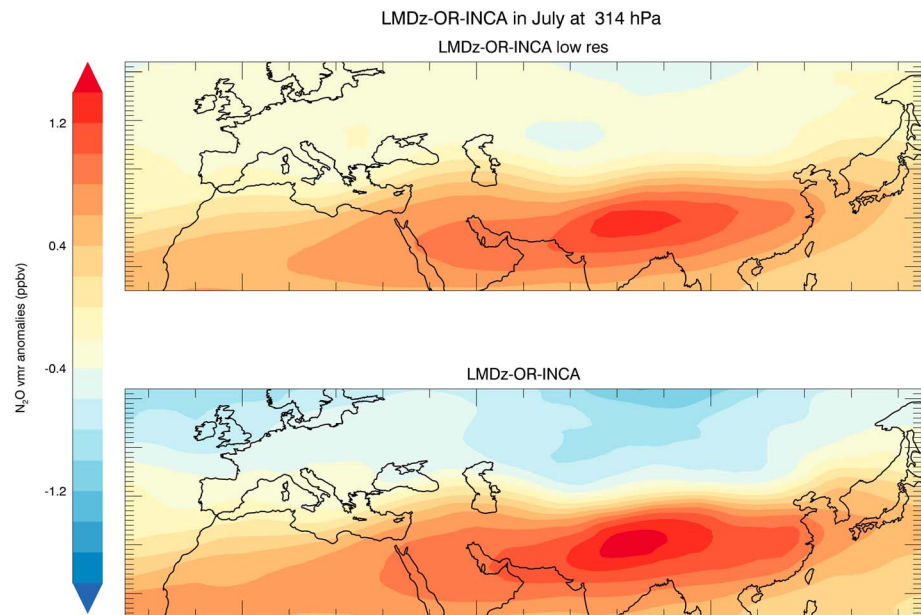


Figure 11. Monthly averaged N_2O volume mixing ratio anomaly (ppbv) from (top) LMDz-OR-INCA *low res* and (bottom) LMDz-OR-INCA at 314 hPa in July over the period of 2010–2013.

levels, are replacing traditional farming in many South Asian countries [Rasul and Thapa, 2003], the denitrification process is enhanced by the use of nitrogen fertilizers leading to an increase of N_2O emissions to the atmosphere [Solomon *et al.*, 2007]. N_2O emissions are positively correlated with soil moisture [Raut *et al.*, 2015], and the relative soil nitrogen emissions (NO , HONO , and N_2O) critically depend on the SWC [Oswald *et al.*, 2013] favoring NO and HONO when SWC ranges from 5 to 40% and favoring N_2O when SWC ranges from 40 to 70%. Thus, the South Asian region is a hot spot of N_2O emissions, especially during summertime when high precipitation events occur. Figure 10 shows the convective precipitation from the reanalysis ERA-Interim [Dee *et al.*, 2011] over the Indian subcontinent (land pixels within the black box in Figure 9) and the monthly averaged N_2O and nitrogen oxide (NO_x) emission fluxes used in LMDz-OR-INCA in this region. In addition to the rainfall information, the convective precipitation can be considered as a footprint of the presence and intensity of the updrafts associated with the Indian monsoon. The Indian subcontinent convective precipitation shows a maximum in July–August. The N_2O surface emissions over this region are highly correlated with precipitation with a maximum in summer and peaking in July. Actually, NO_x emissions peak in May–June, while N_2O emissions peak 1 month later in July (Figure 10). This time lag results from the processes occurring in the soil. Prior to the monsoon period, when SWC is low, a large amount of fertilizer is used in the Indian subcontinent and the surplus of nitrogen in soil mostly produces emissions of the nitrogen compounds NO and HONO . The emissions of NO are represented by NO_x emissions since NO is predominant in emissions of NO_x from soil [Bertram *et al.*, 2005]. During the Indian monsoon period, intense precipitation increases the soil humidity, and once the SWC reaches $\sim 40\%$ [Oswald *et al.*, 2013], N_2O becomes the dominant nitrogen compound emitted. Thus, if a peak in NO_x emissions is reached in June [Ghude *et al.*, 2010] through nitrification processes, it takes few days or weeks for the soil to have enough humidity over the entire Indian subcontinent and for microbial processes to get involved in the production and release of N_2O in the atmosphere through nitrification and denitrification processes [Smith and Tiedje, 1979]. This explains the July peak in N_2O emissions. Consequently, the measurements of N_2O in the upper troposphere over the eastern MB in summer (June–July–August, JJA) can be considered as a footprint of the N_2O emissions over the Asian and especially the Indian subcontinent regions. This is all the more interesting, as this region is difficult to observe from space in summer because of cloudy conditions associated with the monsoon system. It is valuable to examine another source of information to estimate the accumulation of N_2O in the Asian monsoon anticyclone and then to improve the estimation of N_2O emissions from agriculture over Asia.

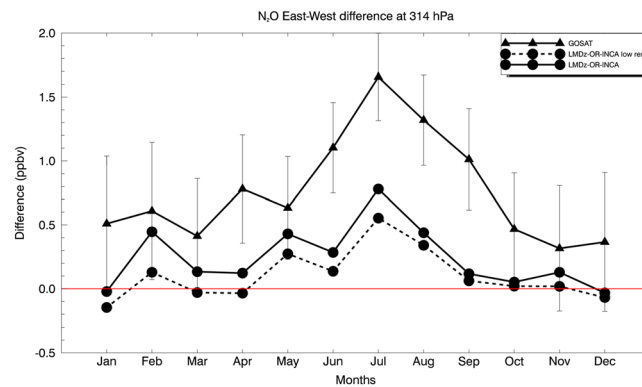


Figure 12. Seasonal variation of the east-west difference of N_2O (ppbv) measured by GOSAT/TANSO-FTS (solid line with triangles) and calculated by LMDz-OR-INCA (solid line with circles) and LMDz-OR-INCA *low res* (dashed line with circles) at 314 hPa over the period of 2010–2013. The error bars represent the error on the mean considering the total random errors of the retrieval and the number of monthly averaged values.

To illustrate the impact of the horizontal resolution and the emission sources on the estimated east-west difference, we have compared the results from two different configurations of the model (LMDz-OR-INCA and LMDz-OR-INCA *low res*) as described in section 2.3. The global total emission rate in LMDz-OR-INCA is about 23% greater than in LMDz-OR-INCA *low res*. Over the Indian subcontinent (black box in Figure 9) in July, the emission rate in LMDz-OR-INCA is about 7% higher than in LMDz-OR-INCA *low res*. Figure 11 shows the northern hemisphere N_2O anomaly in July 2010–2013 at 314 hPa for LMDz-OR-INCA and LMDz-OR-INCA *low res*, and Figure 12 shows the east-west difference of GOSAT, LMDz-OR-INCA, and LMDz-OR-INCA *low res*. The N_2O vmr maxima and gradient in the Asian monsoon are more intense in LMDz-OR-INCA than in LMDz-OR-INCA *low res*. Consistently, the peak-to-peak amplitude of the east-west difference (Figure 12) is more intense in LMDz-OR-INCA (~0.8 ppbv) than in LMDz-OR-INCA *low res* (0.7 ppbv).

Since there is still a significant difference between the peak-to-peak amplitudes observed by GOSAT and calculated by LMDz-OR-INCA (1.8 ppbv and 1.5 ppbv, respectively), the N_2O emission fluxes could be larger over Asia than the fluxes fixed in the emission inventory used by LMDz-OR-INCA. The inaccurate understanding of some processes such as soil acidification involved in soil N_2O emission leads generally to an underestimation of the emission factor used to estimate soil emissions from agriculture [Raut *et al.*, 2015]. Consequently, this generates an underestimation of N_2O emission fluxes, which are linearly affected by the emission factors [Dobbie *et al.*, 1999]. There is also evidence of an underestimation of N_2O emissions at very high N-fertilizer application rates due to a nonlinear relationship between N-fertilizer and N_2O emissions [Shcherbak *et al.*, 2014].

In addition to the emission sources, the horizontal resolution of the model could likely play a role in the model east-west difference. Bergman *et al.* [2013] showed that the contribution of Indian boundary layer sources to the Asian monsoon anticyclone composition in August is less intense for a lower resolution of the dynamical model. This is due to the fact that the finer the horizontal resolution, the better the representation of mesoscale convection and therefore the more efficient the vertical transport processes to bring N_2O -enriched air masses into the upper troposphere. Moreover, the horizontal transport of air masses is also improved and better represented in the high-resolution model than in the coarser resolution model.

In the GOSAT data, the east-west difference is affected by the vertical sensitivity of the measurements. As we have already noted in section 2.2, applying GOSAT optimal estimation formalism (cf. equation (1)) to HIPPO measurements significantly reduces N_2O variability. Thus, to assess the impact of the vertical sensitivity of the measurements (represented by the averaging kernels in Figure 1) on the observed east-west difference, we applied the GOSAT averaging kernels to LMDz-OR-INCA outputs by using equation (1). The result in terms of east-west difference is shown in Figure 13. The smoothing of the model's vertical resolution by the application of GOSAT averaging kernels tends to reduce the variations in east-west difference. Although the difference still peaks in July, the peak-to-peak amplitude decreases from 0.8 to 0.4 ppbv. Actually, the east-west

4. Discussion

Although measured and modeled east-west differences peak in July (Figure 6), the amplitude is twice as large in the GOSAT measurements than in the model outputs. To understand the possible reasons of these differences, we studied some parameters that impact the east-west difference in the model and in the GOSAT data sets.

In the model data, the east-west difference is affected by at least two factors: the emission sources, including the magnitude of emissions as well as the spatial distribution, and the horizontal resolution of the transport model.

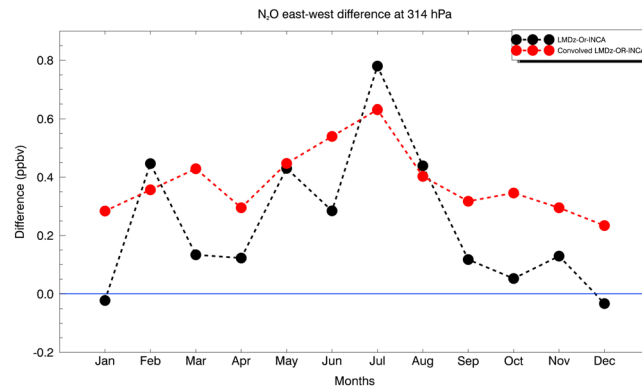


Figure 13. Seasonal variations of the east-west differences calculated by LMDz-OR-INCA (black) and LMDz-OR-INCA convolved by the GOSAT averaging kernels (red) monthly averaged over the period of 2010–2013.

differences are higher all year long in the convolved model than in the original, but the peak in July is less marked in the convolved model than in the original. These changes in the seasonal variability of the east-west difference can be understood when analyzing the broad width of the 314 hPa averaging kernel peak (Figure 1). The convolved LMDz-OR-INCA N_2O data at 314 hPa are also sensitive to higher and lower levels. Since the N_2O east-west difference variability is smaller in LMDz-OR-INCA at lower levels (e.g., a peak-to-peak amplitude of ~ 0.2 ppbv at 502 hPa, not shown) than in the upper troposphere, the impact of lower levels tends to decrease the variability of the east-west difference at 314 hPa. Moreover, because the western MB is located at higher latitudes than the eastern MB (see Figure 4), the convolved model at 200 hPa is influenced more by stratospheric N_2O in the western MB than in the eastern MB. Since N_2O vmr decreases with height in the stratosphere, the impact of the influence of stratospheric N_2O tends to increase the east-west difference at 314 hPa all year long. Similarly, we can consider that the amplitude of the east-west difference variations observed in GOSAT underestimates the real variations in amplitude and the positive east-west difference observed all year long is also likely due to the vertical sensitivity of the GOSAT measurements.

5. Conclusion

We used global-scale measurements of upper tropospheric N_2O from the GOSAT TANSO-FTS satellite instrument and outputs from the chemical transport model LMDz-OR-INCA to evaluate the impact of the Asian N_2O emissions on the N_2O distribution in the MB during summer. We assessed the quality of the GOSAT upper tropospheric N_2O retrievals by using the HIPPO airborne campaigns. This validation indicated an SD error of about 2.0 ppbv for a single pixel and a mean bias of about -1.3 ppbv (approximately -0.4%). Since we used monthly and regional means, this SD error is reduced to ~ 0.1 ppbv. The SD error deduced from the comparisons with HIPPO data is about 3 times less than the theoretical random errors used in this paper to analyze the significance of the seasonal variations.

The GOSAT and LMDz-OR-INCA monthly averaged upper tropospheric N_2O (2010–2013) over the MB have a significant seasonal cycle with a strong increase in summer. The eastern MB is highly enriched in N_2O producing a peak in July in the east-west difference of N_2O over the MB as calculated by the model and as observed by GOSAT. This is due to the impact of air masses originated from Asia and transported over the eastern MB through the Asian monsoon anticyclone. Back trajectory calculations also showed that the main sources of eastern MB upper tropospheric N_2O in summer come from the Indian subcontinent region where high emissions occur during the monsoon period. These high N_2O emissions are due to the high precipitation associated with the monsoon and combined with heavy use of fertilizers, since N_2O emissions occur under conditions of high SWC.

Since the summertime maximum over the eastern MB is linked to the maximum of N_2O surface emissions over Asia and mainly the Indian subcontinent, the eastern MB can therefore be used as a footprint of summertime N_2O emissions over this region. However, the east-west difference peak in July is almost twice as large in GOSAT (~ 1.4 ppbv) than in LMDz-OR-INCA (~ 0.8 ppbv). This is likely due to an underestimation of the N_2O Indian subcontinent emission sources and a relatively coarse horizontal resolution in the model that affects the efficiency of the vertical transport within the Asian monsoon Anticyclone. In addition, the application of the averaging kernels from GOSAT to the model tends to reduce the east-west N_2O difference variations likely due to the low vertical sensitivity of the satellite measurements (represented by the averaging kernel).

One major outcome of this study is that the use of current GOSAT N₂O measurements over the MB can help to improve the estimation of N₂O emissions over Asia and especially the Indian subcontinent in a top-down emission estimation strategy. Moreover, tropospheric N₂O satellite measurements are made difficult in summer over Asia due to cloudy conditions associated with the monsoon period. In such case, upper tropospheric N₂O measurements over the MB could be the best alternative to estimate the N₂O accumulation in the Asian summer monsoon anticyclone and then to assess the surface emissions that caused this accumulation.

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