

IASI is turning ten!

Cathy Clerbaux¹, Maya George¹, Sophie Bauduin², Anne Boynard¹, Pierre Coheur², Lieven Clarisse², Cyril Crevoisier³, Stamatia Doniki², Juliette Hadji-Lazaro¹, Daniel Hurtmans², Jean-Lionel Lacour¹, Gaétane Ronsmans², Sarah Safieddine¹, Martin Van Damme², Catherine Wespes², Simon Whitburn²

¹ *UPMC Univ. Paris 06; CNRS/INSU, LATMOS-IPSL, Paris, France*
Email : cathy.clerbaux@latmos.ipsl.fr

² *Spectroscopie de l'Atmosphère, Service de Chimie Quantique et Photophysique, Université Libre de Bruxelles (ULB), Brussels, Belgium*

³ *Laboratoire de Météorologie Dynamique, CNRS, IPSL, Ecole Polytechnique, Palaiseau*

ABSTRACT

The IASI mission is a versatile mission that fulfills the needs of three different communities: numerical weather prediction, climate research and atmospheric composition monitoring. In order to converge on the design of such an instrument all three communities had to make reasonable accommodations 20 years ago, and it turns out that this mission is now recognized as essential for weather forecasting, and for tracking pollutants/greenhouse gases from space. With the launch of MetOp-B and -C and the continuity and new challenges offered by IASI-NG, an exceptional data record will be available in the next few years.

The presentation (movie) illustrates some of the major findings related to atmospheric composition changes as monitored by IASI during the last 10 years. It relies on accurate data available in near real time along with an excellent horizontal coverage. We will show the global scale mapping of gases, along with the detection of dust and ash particles, as well as the potential of the mission to catch special events such as volcanic eruptions, large fires and pollution peaks.

Tropospheric Ozone Variability during the East Asian Summer Monsoon as Observed by Satellite (IASI), Aircraft (MOZAIC) and Ground Stations

S. Safieddine^{1*}, A. Boynard¹, N. Hao², F. Huang³, L. Wang⁴, D. Ji⁴, B. Barret⁵, S. D. Ghude⁶, P.-F. Coheur⁷, D. Hurtmans⁷, and C. Clerbaux^{1,7}

¹Sorbonne Universités, UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU, LATMOS-IPSL, Paris, France

**Now at* Department of Civil and Environmental Engineering, Massachusetts Institute of Technology, Cambridge, MA, USA

²German Aerospace Center (DLR), Remote Sensing Technology Institute (IMF), Oberpfaffenhofen, Germany

³National Satellite Meteorological Center, China Meteorological Administration, Beijing, China

⁴Institute of Atmospheric Physics, Chinese Academy of Sciences, Beijing, China

⁵Laboratoire d'Aérodynamique, Observatoire Midi-Pyrénées, Université Paul Sabatier, CNRS, Toulouse, France

⁶Indian Institute of Tropical Meteorology; Pashan Rd., Pune-411 008, India

⁷Spectroscopie de l'Atmosphère, Chimie Quantique et Photophysique, Université Libre de Bruxelles (U.L.B.), Brussels, Belgium

Abstract

Satellite measurements from the thermal Infrared Atmospheric Sounding Interferometer (IASI), the Measurements of Ozone and water vapor by in-service Airbus airCRAFT (MOZAIC), as well as observations from ground based stations, are used to assess the tropospheric ozone (O₃) variability during the East Asian Summer Monsoon (EASM). Six years [2008-2013] of IASI data analysis reveals the ability of the instrument to detect the onset and the progression of the monsoon reflected by a decrease in the tropospheric [0-6] km O₃ column due to the EASM, and to reproduce this decrease from one year to the other. Focusing on the period of May-August 2011, taken as an example year, IASI data show clear inverse relationship between tropospheric [0-6] km O₃ on one hand and meteorological parameters such as cloud cover, relative humidity and wind speed, on the other hand. Aircraft data from the MOZAIC project at Hyderabad, Nanjing and Guangzhou are used to validate the IASI data and to assess the effect of the monsoon on the vertical distribution of the tropospheric O₃ at different locations. Results show good agreement with a correlation coefficient of 0.74 between the [0-6] km O₃ column derived from IASI and MOZAIC. The aircraft data show a decrease in the tropospheric O₃ that is more important in the free troposphere than in the boundary layer and at Hyderabad than at the other two Chinese cities. Ground station data at different locations in India and China show a spatiotemporal dependence on meteorology during the monsoon, with decrease up to 22 ppbv in Hyderabad, and up to 5 ppbv in the North China Plain.

Towards a 10 years reanalysis of tropospheric and lower-stratospheric ozone with assimilation of IASI and MLS observations.

Hélène PEIRO ⁽¹⁾, Emanuele EMILI ⁽¹⁾, Eric LE FLOCHMOEN ⁽²⁾, Brice BARRET ⁽²⁾,
Daniel CARIOLLE ⁽¹⁾

⁽¹⁾ **CERFACS/CNRS**

42, avenue G. Coriolis, 31057 Toulouse cedex 01, FRANCE

EMail: peiro@cerfacs.fr ; emili@cerfacs.fr ; cariolle@cerfacs.fr

⁽²⁾ **Laboratoire d'Aérodynamique/CNRS/Université de Toulouse**

14, avenue Edouard Belin, 31400 Toulouse, FRANCE

EMail: Eric.Leflochmoen@aero.obs-mip.fr ; brice.barret@aero.obs-mip.fr

ABSTRACT

Tropospheric ozone is a trace gas involved in the global greenhouse effect. During the last ten years, the tropospheric ozone increase has stopped in North mid-latitudes. Ozone variability in the troposphere is largely driven by stratosphere to troposphere exchanges (STE). An accurate determination of tropospheric and lower stratospheric ozone is necessary to quantify accurately the ozone dynamics.

The aim of this study is to understand and to assess STE and the variability of tropospheric ozone during the available years (2008-2013) of IASI observations from satellite Metop-A. IASI instrument allows global coverage with very high spatial resolution and its measurements are promising for long term ozone monitoring.

The global chemical transport model MOCAGE (MOdèle de Chimie Atmosphérique à Grande Echelle) has been used with a linear ozone chemistry scheme and meteorological forcing fields from ERA-Interim (ECMWF global reanalysis) to assimilate Microwave Limb Sounder (MLS) and IASI data. The 2°x2° global version of MOCAGE, with 60 sigma-hybrid vertical levels has been used. MLS observations and IASI-SOFRID O₃ retrievals have been assimilated with a 4D-VAR variational algorithm to constrain stratospheric and tropospheric ozone respectively.

The MLS analysis, computed for the period 2004 to 2014, is validated against ozonesonde measurements and, for instance, shows episodes of ozone transport between stratosphere and troposphere. This validation confirms the effectiveness of MLS profiles assimilation in the stratosphere. Comparisons between MLS analysis and IASI-SOFRID O₃ tropospheric column show a good consistency. This comparison shows that ozone transport from the upper-troposphere / lower-stratosphere (UTLS) detected by IASI-SOFRID data is consistent with the MLS analysis. As some previous research studies showed, biases between IASI-SOFRID and model (as high as 30%) in high latitudes are present but the ozone variability has been improved by the assimilation (standard deviation in the range of 15%). Overall the assimilation of IASI-SOFRID ozone retrievals improves the model ozone columns in the free troposphere and in the UTLS.

Global tropospheric budget using such long term reanalysis will be discussed. In the future, similar studies could be performed with the IASI-NG instruments with a first launch scheduled in 2018. The objective will be to extend the studied period to several decades.

Ozone and CO upper-tropospheric budget and variability during the Asian Summer Monsoon for 2008-2015: IASI and IAGOS observations and GEOS-Chem simulations.

Brice Barret^(1,2), **Bastien Sauvage**⁽¹⁾, Yasmine Bennouna^(1,2) and Eric Le Flochmoën^(1,2)

⁽¹⁾ Université de Toulouse, Laboratoire d'Aérodynamique, Toulouse, France

⁽²⁾ CNRS UMR 5560, Toulouse, France

Brice Barret

Observatoire Midi-Pyrénées - Laboratoire d'Aérodynamique

14 avenue Edouard Belin

31400 TOULOUSE - FRANCE

Email: brice.barret@aero.obs-mip.fr

Phone: +33 (0)5 61 33 27 23

Topic 2 : Science and application

Atmospheric composition

Oral presentation

Abstract :

During the Asian Summer Monsoon (ASM), primary pollutants (CO, HCN, aerosols) accumulate into the large scale Asian Monsoon Anticyclone (AMA) that extends from the upper troposphere to the lower stratosphere. This accumulation is mostly due to South and East Asian pollution uplifted by monsoon convection. This lower tropospheric air uplift has a potentially strong impact upon the UTLS O₃, aerosols and water vapour budget and therefore also on the surface radiative forcing. Boundary layer precursors (NO_x, COV's) uplifted from the densely populated regions of Asia are responsible for the photochemical production of O₃. Other sources have a potentially large impact on UT O₃ enhancement such as the production of NO_x by lightning's and stratospheric intrusions from the subtropical jet.

The first objective of our study is to quantify the relative impact of these different sources on the Asian UT composition during the pre- to post-monsoon period. We have therefore performed a reference simulation with the GEOS-Chem global chemistry transport model for the 2008-2013 period. Distributions of CO and O₃ in the Asian troposphere are evaluated versus IAGOS observations from flights between Asia and Europe or the Middle-East and versus IASI-SOFRID retrievals. For chemical budget assessment we have performed sensitivity simulations with CO and NO_x emissions from different types and regions impacting the AMA switched off. Our second objective is to characterize the intra-seasonal and interannual variabilities of the AMA composition. We will therefore present the departures of the CO and O₃ distributions from their climatological means for the 2008-2015 period. The analyses of these departures in light of different climate indices, meteorological parameters and sensitivity simulations will help to understand the dynamical and chemical factors controlling them.

CO monitoring with IASI: global and local variability

Maya George^{1,2}, C. Clerbaux^{1,2}, J. Hadji-Lazaro¹, D. Hurtmans², S. Bauduin², S. Whitburn², I. Bouarar³, A. Inness⁴, M. Krol⁵ and P.-F. Coheur²

¹ *Sorbonne Universités, UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU, LATMOS-IPSL, Paris, France*

Email : maya.george@latmos.ipsl.fr

Email : cathy.clerbaux@latmos.ipsl.fr

Email : juliette.hadji-lazaro@latmos.ipsl.fr

² *Spectroscopie de l'Atmosphère, Service de Chimie Quantique et Photophysique, Université Libre de Bruxelles, Brussels, Belgium*

Email : dhurtma@ulb.ac.be

Email : sbauduin@ulb.ac.be

Email : swhitbur@ulb.ac.be

Email : pfcoheur@ulb.ac.be

³ *Max Planck Institute for Meteorology, Hamburg, Germany*

Email : Idir.Bouarar@latmos.ipsl.fr

⁴ *European Centre for Medium-Range Weather Forecasts, Reading, UK*

Email : Antje.Inness@ecmwf.int

⁵ *Institute for Marine and Atmospheric Research Utrecht (IMAU), University of Utrecht, Utrecht, The Netherlands*

Email : m.c.krol@uu.nl

ABSTRACT

Carbon monoxide (CO) is an important trace gas for understanding air quality and atmospheric composition. It is a good tracer of pollution plumes and atmospheric dynamics. Since the launch of the MetOp satellite, IASI CO concentrations are retrieved on a daily base and in near real time from the radiance data, using the Fast Operational/Optimal Retrievals on Layers for IASI (FORLI) algorithm. These products are now generated and operationally distributed by EUMETCAST under the O3MSAF auspices.

Global scale CO distributions will be shown, and the seasonal/annual variabilities over 8 years will be analyzed. A special focus will be put on the record concentrations observed over the islands of Borneo and Sumatra in the fall 2015, due to the worst forest fires in nearly two decades. This pollution excess is associated with the predictable impacts of the strong El Niño currently underway. We will provide estimates of the carbon emissions using a data-assimilation system that derives CO emissions from IASI data. Finally, we will show how this event is taken into account by the Copernicus Atmospheric Monitoring Service, and the usefulness of the IASI data in this context.

IASI 2016 Abstract

Assessing the Impacts of Assimilating IASI and MOPITT CO Retrievals using CESM-CAM-chem and DART

D. Edwards ⁽¹⁾, J. Barré ⁽¹⁾, B. Gaubert ⁽¹⁾, A. Arellano ⁽²⁾, H. Worden ⁽¹⁾, M. Deeter ⁽¹⁾, J. Anderson ⁽¹⁾, K. Raeder ⁽¹⁾, N. Collins ⁽¹⁾, S. Tilmes ⁽¹⁾, G. Francis ⁽¹⁾, C. Clerbaux ^(3,4), L. Emmons ⁽¹⁾, G. Pfister ⁽¹⁾, P.-F. Coheur ⁽⁴⁾, D. Hurtmans ⁽⁴⁾

⁽¹⁾ National Center for Atmospheric Research

Boulder, CO, USA

Email: edwards@ucar.edu

⁽²⁾ University of Arizona

Tucson, AZ, USA

⁽³⁾ UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU, LATMOS-IPSL

Paris, France

⁽⁴⁾ Spectroscopie de l'Atmosphère, Service de Chimie Quantique et Photophysique, Université Libre de Bruxelles (ULB)

Brussels, Belgium

ABSTRACT

We show the results and evaluation with independent measurements from assimilating both MOPITT (Measurements Of Pollution In The Troposphere) and IASI (Infrared Atmospheric Sounding Interferometer) retrieved profiles into the Community Earth System Model (CESM). We used the Data Assimilation Research Testbed ensemble Kalman filter technique, with the full atmospheric chemistry CESM component Community Atmospheric Model with Chemistry. We first discuss the methodology and evaluation of the current data assimilation system with coupled meteorology and chemistry data assimilation. The different capabilities of MOPITT and IASI retrievals are highlighted, with particular attention to instrument vertical sensitivity and coverage and how these impact the analyses. MOPITT and IASI CO retrievals mostly constrain the CO fields close to the main anthropogenic, biogenic, and biomass burning CO sources. In the case of IASI CO assimilation, we also observe constraints on CO far from the sources. During the simulation time period (June and July 2008), CO assimilation of both instruments strongly improves the atmospheric CO state as compared to independent observations, with the higher spatial coverage of IASI providing better results on the global scale. However, the enhanced sensitivity of multispectral MOPITT observations to near surface CO over the main source regions provides synergistic effects at regional scales.

Spatio-temporal variability of three anthropogenic greenhouse gases CO₂, CH₄ and N₂O in the mid-troposphere as seen from IASI onboard Metop-A and Metop-B

Cyril Crevoisier ⁽¹⁾, Nicolas Meilhac⁽¹⁾, Olivier Membrive ⁽¹⁾, Laurent Crépeau ⁽¹⁾, Raymond Armante ⁽¹⁾, Noëlle Scott ⁽¹⁾, Alain Chédin ⁽¹⁾

⁽²⁾ *Laboratoire de Meteorologie Dynamique, CNRS, IPSL*

Ecole polytechnique, Université Paris-Saclay, 91128 Palaiseau Cedex, France

Email: cyril.crevoisier@lmd.polytechnique.fr

ABSTRACT

Thanks to its continuous spectral coverage of the whole thermal infrared domain, the IASI instrument offers the possibility to retrieve mid-tropospheric columns of the 3 major greenhouse gases influenced by human activities. These are: carbon dioxide (CO₂), methane (CH₄) and nitrous oxide (N₂O). The very small seasonal variability of these gases compared to their background values, combined to the strong dependence of IR radiances to atmospheric temperature and the simultaneous sensitivity of the channels to several gases, makes their retrieval challenging.

The non-linear inference scheme developed at LMD delivers mid-tropospheric columns of the 3 gases for both day and night condition, over land and over sea. It strongly relies on careful validation of levelle spectra, characterization of systematic radiative biases and severe cloud and aerosol screening. Almost 9 years of observations from Metop-A and 2 years from Metop-B have now been processed. CH₄ and CO₂ atmospheric columns are delivered on “near real time” (D-1) basis to the Copernicus-Atmosphere Service. Owing to its 20 year-program, IASI also participates to the establishment of long time series in the ESA-Climate Change Initiative-GHG. The retrievals are thus used for a variety of purpose: assimilation to produce CH₄/CO₂ profile forecasts; estimation of surface fluxes using “top-down” atmospheric inversions; characterization of specific emissions such as biomass burnings.

In this talk we will present the latest development of the retrievals of these 3 gases. In particular, recent achievements include: (i) extension of the spatial coverage of methane from the tropical region to mid and high latitudes; (ii) improvement of CO₂ retrieval in the tropical region by a combination of hundreds of 15μm channels; (iii) processing of Metop-B; (iv) characterization of the retrieval uncertainty based on the ARSA database; (v) near-real-time delivery of the retrievals (D-1). We will conclude by presenting the first results on spatio-temporal distribution of N₂O obtained from IASI observation.

Global monitoring of atmospheric CO₂ and CH₄ through the assimilation of level 2 IASI data within the Copernicus Atmosphere Monitoring Service

Sébastien Massart ⁽¹⁾, Cyril Crevoisier ⁽²⁾, Anna Agusti-Panareda ⁽³⁾, Mark Parrington ⁽⁴⁾

⁽¹⁾ *European Centre for Medium-Range Weather Forecasts
Shinfield Park, RG2 9AX, Reading, United Kingdom
EMail: sebastien.massart@ecmwf.int*

⁽²⁾ *Laboratoire de Meteorologie Dynamique, CNRS, IPSL
Ecole Polytechnique, 91128 Palaiseau Cedex, France
EMail: cyril.crevoisier@lmd.polytechnique.fr*

⁽³⁾ *European Centre for Medium-Range Weather Forecasts
Shinfield Park, RG2 9AX, Reading, United Kingdom
EMail: anna.agusti-panareda@ecmwf.int*

⁽⁴⁾ *European Centre for Medium-Range Weather Forecasts
Shinfield Park, RG2 9AX, Reading, United Kingdom
EMail: mark.parrington@ecmwf.int*

ABSTRACT

As part of a delegation agreement with the European Union, the European Centre for Medium-Range Weather Forecasts (ECMWF) is managing the Copernicus Atmosphere Monitoring Service (CAMS: <http://atmosphere.copernicus.eu/>). CAMS provides daily near-real-time analyses and forecasts of global atmospheric composition including greenhouse gases (GHGs), reactive gases, and aerosols. We will focus the presentation on the analysis and forecast of carbon dioxide (CO₂) and methane (CH₄), two of the key greenhouse gases that are currently available in the CAMS system.

The present GHG analysis relies on the assimilation of the column-average dry-air mole fractions of CO₂ and CH₄ from the Greenhouse gases Observing Satellite (GOSAT). The assimilated GOSAT retrievals have a high near-surface sensitivity. They are, nevertheless, limited to daytime observations and sufficiently cloud-free conditions are also needed for accurate retrievals. These limitations combined with the design of the instrument reduce the number of available observations to only a few within the 12 hour assimilation window of the CAMS system.

Retrievals of CO₂ and CH₄ from the Infrared Atmospheric Sounding Interferometer (IASI) have a much better spatial and temporal coverage, with availability during daytime and nighttime, but with limited sensitivity to the lower troposphere due to the low thermal contrast between the Earth's surface and the air masses above. However, the IASI measurements have higher sensitivity to the mid-troposphere and retrieved GHG products from IASI are highly complementary to the GOSAT products.

We have already demonstrated the positive impact of the assimilation of IASI CH₄ retrievals on the analysis when we limited the data to the tropics. We will demonstrate the impact of the assimilation of IASI CO₂ retrievals and the assimilation of extra-tropical IASI data on the analysis. To assess this, we will use independent measurements such as in situ profiles from HIAPER Pole-to-Pole Observations (HIPPO) and from Comprehensive Observation Network for TRace gases by AIRLiner (CONTRAIL) or column-average remote sensing measurements from the Total Carbon Column Observing Network (TCCON).

O₃ variability in the troposphere and the stratosphere from IASI observations in 2008-2015

Catherine Wespes¹, Pierre-François Coheur¹, Daniel Hurtmans¹, and Cathy Clerbaux^{1,2}

¹ Spectroscopie de l'Atmosphère, Service de Chimie Quantique et Photophysique, Université Libre de Bruxelles (ULB), Brussels, Belgium

² Sorbonne Universités, UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU, LATMOS-IPSL, Paris, France

In this study, we present geographical patterns of ozone (O₃) variability in the troposphere and the stratosphere derived from 8 years of IASI observations (2008-2015). The instrument provides a unique dataset of global vertically-resolved O₃ profiles with a high temporal sampling and a fairly good vertical resolution in the troposphere and the stratosphere allowing us to monitor the year-to-year variability in these layers. The retrievals are performed using the FORLI software, a fast radiative transfer model based on the optimal estimation method, set up for near real time and large scale processing of IASI data.

Multivariate regressions which include important geophysical drivers of O₃ variation (e.g. solar flux - SF, quasi biennial oscillations – QBO, El Niño/Southern Oscillation – ENSO) and a linear trend term have been performed on time series of spatially averaged O₃ on spatial grids. The resulting covariates and trend spatial structures are analyzed. In particular, we focus on O₃ variation and trends in the troposphere over and downwind anthropogenic polluted areas characterized either by increased (e.g. over Asia) or decreased (e.g. over Europe and the US) O₃ precursor emissions. Direct effects of positive (or negatives) ENSO indexes measured during moderate to intense El Niño (or La Niña) episodes in 2009 and 2014 (or 2010) observed in the ozone columns in the tropics will be also discussed.

Unaccounted variability in NH₃ agricultural sources detected by IASI contributing to European spring haze episode

A. Fortems-Cheiney¹, G. Dufour¹, L. Hamaoui-Laguel², G. Foret¹, G. Siour¹, M. Van Damme³, F. Meleux², P.-F. Coheur³, C. Clerbaux^{3,4}, L. Clarisse³, M. Wallasch⁵, and M. Beekmann¹

¹Laboratoire Interuniversitaire des Systèmes Atmosphériques, CNRS/INSU UMR7583, Université Paris-Est Créteil et Université Paris Diderot, Institut Pierre Simon Laplace, Créteil, France.

audrey.cheiney@lisa.u-pec.fr

gaelle.dufour@lisa.u-pec.fr

gilles.foret@lisa.u-pec.fr

guillaume.siour@lisa.u-pec.fr

matthias.beekmann@lisa.u-pec.fr

²Institut National de l'Environnement et des RISques industriels (INERIS), Verneuil-en-Halatte, France.

Lynda.Hamaoui-Laguel@ineris.fr

Frederik.Meleux@ineris.fr

³Spectroscopie de l'atmosphère, Chimie Quantique et Photophysique, Université Libre de Bruxelles (ULB), Brussels, Belgium.

Martin Van Damme <Martin.Van.Damme@ulb.ac.be>

[Pierre Coheur <pfcoheur@ulb.ac.be>](mailto:pfcoheur@ulb.ac.be)

Lieven Clarisse <lclariss@ulb.ac.be>

⁴UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU, LATMOS-IPSL, Paris, France.

Cathy Clerbaux <Cathy.Clerbaux@latmos.ipsl.fr>

⁵Umweltbundesamt, Paul-Ehrlich-Str; 29, 63225, Langen, Germany.

[Wallasch, Markus <markus.wallasch@uba.de>](mailto:markus.wallasch@uba.de)

ABSTRACT

Ammonia (NH₃), whose main source is agriculture, is an important gaseous precursor of atmospheric particulate matter (PM). For the first time, we derived daily ammonia emissions using NH₃ total columns from the Infrared Atmospheric Sounding Interferometer (IASI) onboard Metop-A, at a relatively high spatial resolution (grid-cell of 0.5° x 0.5°). During the European spring haze episode of March 2014 8th to 15th, IASI reveals NH₃ total column magnitudes highlighting higher NH₃ emissions over Central Europe (especially over Germany, Czech Republic and eastern France) from the ones provided by the European reference EMEP inventory. These ammonia emissions exhibit in addition a large day-to-day variability, certainly due to spreading practices.

The increase of NH₃ emissions in the model, that reaches +300% locally, leads to an increase of both NH₃ and PM₁₀ surface concentrations and allows for a better comparison with independent measurements (in terms of bias, root mean square error and correlation). This preliminary study suggest that there are good promises for better quantifying NH₃ emissions by atmospheric inversions.

Ozone pollution outbreaks analysed by synergism of novel IASI+GOME2 multispectral satellite observations, chemistry-transport models and in situ measurements

Juan Cuesta⁽¹⁾, Yugo Kanaya⁽²⁾, Emanuele Emili⁽³⁾, Gaëlle Dufour⁽¹⁾, Adriana Coman⁽¹⁾, Masayuki Takigawa⁽²⁾, Kazuyuki Miyazaki⁽²⁾, Maxim Eremenko⁽¹⁾, Gilles Foret⁽¹⁾, Matthias Beekmann⁽¹⁾

⁽¹⁾ *LISA, UMR CNRS 7583, Université Paris Est Créteil and Université Paris Diderot
61 Av. Général de Gaulle, 94010, Créteil, France*

Email: cuesta@lisa.u-pec.fr

⁽²⁾ *JAMSTEC*

2-15 Natsushimacho, Yokosuka, 237-0061, Kanagawa Prefecture, Japan

⁽³⁾ *CERFACS*

42 Av. G. Coriolis, 31057, Toulouse, France

ABSTRACT

Ozone pollution is a major worldwide-shared concern, which poses great threats to human health and ecosystems. Assessment of the origin of low-level ozone is complex, due to a great diversity of sources (local photochemical production, transboundary transport and exchanges with the stratosphere) and transport mechanisms determining its three-dimensional distribution. Satellite observations play a key role to monitor tropospheric ozone from regional to global scales and daily basis. Particularly, the capacity to observe ozone pollution has been recently enhanced by the novel multispectral synergism so-called IASI+GOME2 of IASI and GOME-2 measurements (in the IR and UV respectively) that has enabled the first spaceborne observation of ozone plumes in the lowermost troposphere (below 3 km of altitude). However, one needs to combine these observations with chemistry-transport models in order to describe the hourly evolution and fine vertical distribution of ozone pollution down to the surface. Moreover, such combination of satellite observations and models is a powerful approach for their inter-validation and for correcting the model by data assimilation.

This research work presents an analysis of ozone pollution combining IASI+GOME2 satellite observations, state-of-the-art chemistry transport models and in situ measurements over two key regions: East Asia and Europe. In both cases, ozone pollution outbreaks are analysed in term of their origin, their three-dimensional pathways and the impact on regional air quality. We also use pollution tracers such as CO co-located observations derived from IASI measurements and stratospheric air indicators such as potential vorticity fields derived from the ECMWF meteorological model. The analysis over East Asia focuses on the springtime of 2009 and that over Europe in July 2010. The following chemistry-transport simulations are used i) for East Asia: WRF-CHEM, WRF-CMAQ and CHASER, this last one having assimilated numerous satellite observations and ii) for Europe: CHIMERE and MOCAGE, with assimilation in this last case of IASI and surface datasets. Moreover, we study the contribution of the assimilation IASI+GOME2 observations into the MOCAGE and CHIMERE models, for better representing the 3D distribution of tropospheric ozone over Europe.

Atmospheric Trace Gas Anomaly Detection using IASI

Lucy J. Ventress ⁽¹⁾, Roy G. Grainger ⁽¹⁾, Catherine Hayer ⁽¹⁾, Elisa Carboni ⁽¹⁾

⁽¹⁾ *Atmospheric Oceanic and Planetary Physics, University of Oxford
Clarendon Laboratory, Parks Road, Oxford, OX1 3PU, U.K.*

Email: lucy.ventress@physics.ox.ac.uk

ABSTRACT

In order to provide near-real-time monitoring of atmospheric contaminants, fast and reliable methods are required to detect anomalies in the atmospheric state. Full optimal estimation retrievals are computationally expensive; therefore, faster methods are needed to identify such anomalous events. The Infrared Atmospheric Sounding Interferometer (IASI), on board both the MetOp-A and MetOp-B platforms, is a Fourier transform spectrometer covering the mid-infrared from 645-2760 cm^{-1} (3.62-15.5 μm) with a spectral resolution of 0.5 cm^{-1} (apodised) and a pixel diameter at nadir of 12km. These characteristics allow global coverage to be achieved twice daily for each instrument, making IASI a very useful tool for the observation and tracking of atmospheric pollutants, large aerosol particles (such as desert dust) and volcanic plumes. The method shown makes full use of the spectral information from hyperspectral sounders and allows the presence of the target species to be determined in near-real-time (NRT), if required.

The analysis algorithms developed at the University of Oxford are well established for the flagging of volcanic ash and SO_2 . A linear retrieval is used to detect departures in the atmospheric state from a generalised background covariance matrix created from IASI data containing no enhancement. These procedures have been extended to detect enhancements in additional atmospheric species, such as NH_3 and CO , which are important contaminants in pollution monitoring and forest fire detection. The data from the anomaly detection tests are available online via the IASI NRT website within 3 hours of measurement.

Ammonia emissions in biomass burning regions: Estimation from satellite-derived measurements and comparison with bottom-up inventories

S. Whitburn⁽¹⁾, M. Van Damme⁽¹⁾, J.W. Kaiser⁽²⁾, G.R. van der Werf⁽³⁾, S. Turquety⁽⁴⁾, L. Clarisse⁽¹⁾, C. Clerbaux^(1,5) and P.-F. Coheur⁽¹⁾

⁽¹⁾ *Spectroscopie de l'Atmosphère, Service de Chimie Quantique et Photophysique, Université Libre de Bruxelles,*

Avenue Franklin Roosevelt 50, 1050, Brussels, Belgium

Email: simon.whitburn@ulb.ac.be

⁽²⁾ *Max Planck Institute for Chemistry,*

Mainz, Germany

⁽³⁾ *Faculty of Earth and Life Sciences, VU University Amsterdam,*

Amsterdam, The Netherlands

⁽⁴⁾ *UPMC Univ. Paris 06; Ecole Polytechnique, CNRS/INSU, LMD-IPSL,*

Palaiseau, France

⁽⁵⁾ *Sorbonne Universités, UPMC Univ. Paris 06; Université Versailles St-Quentin; CNRS/INSU,*

LATMOS-IPSL,

Paris, France

Email: Martin.Van.Damme@ulb.ac.be, j.kaiser@mpic.de, guido.vander.werf@vu.nl,

solene.turquety@lmd.polytechnique.fr, lclariss@ulb.ac.be, cathy.clerbaux@latmos.ipsl.fr, pfcoheur@ulb.ac.be

ABSTRACT

Vegetation fires emit large amounts of ammonia in the atmosphere (NH₃). Excess NH₃ can affect the environment in many ways: eutrophication of the ecosystem, soil acidification and loss of plant diversity. Despite the numerous studies achieved in the past decades to better quantify NH₃ emissions from biomass burning (mainly from bottom-up approaches), the uncertainties remain still large. NH₃ satellite measurements are now available for a few years and offer the possibility to better assess the NH₃ atmospheric budget, its spatial distribution and long-term trends.

In this presentation we first analyze monthly NH₃ total columns (molec cm⁻²) from IASI on MetOp-A satellite over four large tropical regions known for their important seasonal fire activity. We next derive for these regions the monthly NH₃ emissions using a simplified box model and perform a top-down evaluation for NH₃ of the most widely used fire emission inventories (GFEDv3.1 and the GFASv1.2).

We then focus our attention on the NH₃ emissions from the intense fires which were devastating Indonesia in fall 2015. The region of Indonesia is characterized by peat soils, whose burning is thought to emit large amounts of NH₃. To assess the importance of NH₃ emissions from peat fires compared to other biomes, we derive NH₃ enhancement ratios (ERs) relative to CO and we compare them to ERs derived from other biomes in tropical biomass burning regions. We also compare these results to the ERs reported in the literature.

IASI 2016 Abstract

Detection of tropospheric ozone from IASI during exceptional events

**Anne Boynard ⁽¹⁾, Cathy Clerbaux ^(1,2), Sarah Safieddine ⁽³⁾, Catherine Wespes ⁽²⁾,
Juliette Hadji-Lazaro ⁽¹⁾, Pierre-Francois Coheur ⁽²⁾, and Daniel Hurtmans ⁽²⁾**

*⁽¹⁾ Sorbonne Universités, UPMC Univ. Paris 06; Université Versailles St-Quentin;
CNRS/INSU, LATMOS-IPSL
4 place Jussieu, 75252 Paris Cedex 05, France
EMail: anne.boynard@latmos.ipsl.fr, cathy.clerbaux@latmos.ipsl.fr, juliette.hadji-
lazaro@latmos.ipsl.fr*

*⁽²⁾ Spectroscopie de l'Atmosphère, Chimie Quantique et Photophysique, Université Libre
de Bruxelles (U.L.B.)
CP160/09, avenue F.D. Roosevelt 50, 1050 Brussels, Belgium
EMail: wespes@ulb.be, coheur@ulb.be, hurtmans@ulb.be*

*⁽³⁾ Department of Civil and Environmental Engineering, Massachusetts Institute of
Technology
77 Massachusetts Avenue, Cambridge, MA 02139-4307, USA
EMail: sarahsaf@mit.edu*

ABSTRACT

As an important greenhouse gas and air pollutant, tropospheric ozone (O₃) has become a significant environmental and climate concern. Both strong local photochemical production within the troposphere and transport from the stratosphere can lead to elevated tropospheric O₃ concentrations. These have increased globally over the last century due to enhanced anthropogenic precursor emissions. Monitoring of tropospheric O₃ is therefore essential to understand its impact on air quality, chemical composition and climate change.

Although quantifying tropospheric O₃ from space is difficult because of the abundance of ozone in the stratosphere, over the past decade satellite instruments have provided valuable information on its distribution and temporal evolution. Here, we use the IASI instrument to retrieve tropospheric O₃ from space and assess its ability to detect ozone in the lower troposphere during exceptional events, such as photochemical smogs. We use the WRF-Chem chemical transport model to interpret observations of tropospheric O₃ from IASI, and to assess the meteorological and dynamical processes involved in such extreme events.

Variability and short-term trends of lower and upper tropospheric ozone over the North China Plain from IASI observations.

G. Dufour⁽¹⁾, M. Eremenko⁽¹⁾, J. Cuesta⁽¹⁾, G. Forêt⁽¹⁾, M. Beekmann⁽¹⁾, A. Cheiney⁽¹⁾, Z. Cai⁽²⁾, Y. Liu⁽²⁾, M. Takigawa⁽³⁾, Y. Kanaya⁽³⁾

⁽¹⁾*Laboratoire Inter-universitaire des Systèmes Atmosphériques (LISA/CNRS)
Universités Paris-Est Créteil, 61 avenue du Général de Gaulle, 94000, Créteil, France
Email: Gaelle.Dufour@lisa.u-pec.fr*

⁽²⁾*Key Laboratory of middle Atmosphere and Global Environment Observation
Institute of Atmospheric Physics, Chinese Academy of Sciences, Beijing, China*

⁽³⁾*Japan Agency for Marine-Earth Science and Technology
Yokohama, Japan*

Important progresses in the field of atmospheric ozone sounding from space have been accomplished during the last decade. The lower troposphere is now available from IASI (Infrared Atmospheric Sounding Interferometer) with a maximum of sensitivity between 3 and 4 km. We use satellite observations from IASI on board the MetOp satellites to evaluate the short-term trends of tropospheric ozone over the North China Plain (NCP) for the period 2008-2015. The availability of two semi-independent columns of ozone from the surface up to 12 km allow ones to derive ozone trends for the lower (surface to 6 km a.s.l) and the upper troposphere (6 to 12 km a.s.l). Monthly variations show a maximum in late spring/early summer (May, June) in the lower troposphere. Short-term trends are calculated from deseasonalized monthly variations. Preliminary results show a negative trend of about -0.50%/yr in the lower troposphere whereas a positive trend ($\sim +0.7\%/yr$) is observed in the upper troposphere. If the negative trend appears to be significant in the lower troposphere, the positive trend in the upper troposphere remains poorly significant over such a short period and seems moderated by the stratosphere-troposphere exchanges. The consistency between the two in-flight IASI instruments aboard MetOp-A and MetOp-B is discussed. The IASI derived trends are compared to independent in situ observations provided by ozone sounding at different sites of East Asia for evaluation. The different processes driving the variability and short-term trends of lower tropospheric ozone during the 2008-2015 period are discussed.

Seasonal and interannual variabilities of tropospheric Ozone and CO over the Indian Ocean with IASI-SOFRID 2008-2013 data

Eric Le Flochmoën^(1,2), Brice Barret^(1,2), Bastien Sauvage⁽¹⁾

⁽¹⁾ Université de Toulouse, Laboratoire d'Aérodynamique, Toulouse, France

⁽²⁾ CNRS UMR 5560, Toulouse, France

Eric LE FLOCHMOËN

Observatoire Midi-Pyrénées - Laboratoire d'Aérodynamique

14 avenue Edouard Belin

31400 TOULOUSE - FRANCE

Email: eric.leflochmoen@aero.obs-mip.fr

Phone: +33 - 5 61 33 27 14

Topic 2 : Science and application

Atmospheric composition

Poster presentation

Abstract :

Tropospheric O₃ over the Southern Indian Ocean is controlled by a number of sources such as Stratosphere to Troposphere Exchanges, LiNO_x and Biomass Burning from Africa. From multi annual (2005-2009) observations from the TES space borne sensor, Zhang et al. (2012) recently highlighted an O₃ maximum in May. This maximum was shown to be caused by the production of LiNO_x from Africa. Important interannual variability was observed with larger maxima in 2006 and 2008 related to anomalous anti-cyclonic circulation over central Africa. Nevertheless, many aspects concerning the factors controlling the variability and evolution of O₃ over the Southern Indian Ocean region remain unclear. In this study we use 6 years (2008-2013) of O₃ and CO data retrieved from the Metop-A/IASI sensor with the Software for a Fast Retrievals of IASI Data (SOFRID) to document the seasonal and interannual variability of tropospheric O₃ and CO over the Indian Ocean region. In order to understand the observed seasonal and interannual variabilities, we have performed simulations with the GEOS-Chem chemistry transport model. We present comparisons between the simulated and observed O₃ and CO distributions and a preliminary discussion concerning the source regions and transport processes responsible for the observed variabilities.

Using IASI to characterise CO and other volatile organic compound emissions

David Moore ⁽¹⁾, Michael Barkley ⁽²⁾, Jeremy Harrison ⁽²⁾, and John Remedios ⁽²⁾

⁽¹⁾ *National Centre for Earth Observation*

University of Leicester, Dept. Physics and Astronomy, LE1 7RH, UK.

EMail: david.moore@le.ac.uk

⁽²⁾ *Dept. Physics and Astronomy*

University of Leicester, LE1 7RH, UK

ABSTRACT

It is well established that terrestrial vegetation emits a diverse range of volatile organic compounds (VOCs) into the atmosphere either through direct emission or as a consequence of burning through fire events. VOCs serve important roles in the biosphere and influence global atmospheric chemistry and affect climate. Photochemical reactions of VOCs influence the oxidation capacity of the atmosphere and, as a direct consequence, the lifetimes and distributions of other key climatic gases, such as methane (CH₄) and carbon monoxide (CO)

The University of Leicester IASI Retrieval Scheme (ULIRS) is an algorithm which has been developed to retrieve CO and VOC concentrations from IASI measured top of atmosphere radiances on a local scale, using an Optimal Estimation Method. The scheme was developed to incorporate a line-by-line forward model and the calculation of the extinction coefficient is one of the most time-consuming steps of the ULIRS. We investigate the use of pre-calculated look-up tables for CO retrievals and discuss the effects on retrieval accuracy and significantly reduced computation times which allow the scheme to be used for near-real time applications.

The CO and VOC signatures in IASI spectra in the immediate vicinity of fires and within (boreal) biomass plumes, has been examined. For example, HCOOH contributes significantly to acidity of precipitation and it is shown that a state-of-the-art chemical transport model significantly underestimates the enhancement ratios for $\Delta\text{HCOOH}/\Delta\text{CO}$ from fires over North America implying that the model lacks important primary and secondary sources of atmospheric HCOOH.

Emissions from tropical forest vegetation is an area where remotely-sensed products from satellite measurements are helping us to better understand local chemical processes and complement the sporadic aircraft campaign measurements and sparse observing networks over these regions. We show successful detection of very short-lived, weakly absorbing, biogenic VOCS (BVOCs) over the Amazon by using co-addition of IASI spectra to reduce the spectral noise and enhance the absorption signal. Although some information is lost by spatially averaging data, the large number of IASI measurements each day means that this method still provides valuable information to determine regional emissions on weekly to monthly scales. Regional positive correlations of up to 0.5 are found between IASI BVOCs and GEOS-Chem model data. Seasonal comparisons show some significant temporal differences between the maxima in the datasets, peaking around one or two months later in the satellite data over the Amazon region, for example.

**Main results of Carbon Dioxide comparison
between IASI, TANSO-FTS and HIPPO products.**

Cecilia Tirelli ⁽¹⁾, Ugo Cortesi ⁽¹⁾, Samuele Del Bianco ⁽¹⁾, Lucia Laurenza ⁽²⁾, Naoko Saitoh ⁽³⁾

⁽¹⁾ ***Institute for Applied Physics “Nello Carrara” (IFAC-CNR)***
Via Madonna del Piano 10, 50019 Sesto Fiorentino, Firenze, Italy

Email: c.tirelli@ifac.cnr.it, u.cortesi@ifac.cnr.it, s.delbianco@ifac.cnr.it

⁽²⁾ ***Institute for Electromagnetic Sensing of the Environment (IREA-CNR)***

Via Bassini, 15, 20133 Milano, Italy

Email: laurenza.l@irea.cnr.it

⁽³⁾ ***Center for Environmental Remote Sensing, Chiba University***

1-33 Yayoi-cho, Inage-ku, Chiba-shi, Japan

Email: nsaitoh@faculty.chiba-u.jp

ABSTRACT

A great effort is undertaken by the scientific community to investigate carbon dioxide amounts and vertical distribution and to characterize its sources and sinks. In this background, the development of CO₂ retrieval models in the Thermal InfraRed (TIR) spectral range is extremely important for the characterization of the CO₂ distribution in the mid-to-upper troposphere with a high accuracy level. The research project “Application of KLIMA Algorithm to CO₂ Retrieval from IASI/METOP-A Observations and Comparison with TANSO-FTS/GOSAT Products” had the main purpose to investigate the performances of KLIMA algorithm applied to the Infrared Atmospheric Sounding Interferometer (IASI) observations. The Thermal And Near infrared Sensor for carbon Observation (TANSO)-FTS on board of the Greenhouse gases Observing SATellite (GOSAT) can measure CO₂ column amounts from the ShortWave InfraRed band (SWIR) and CO₂ profiles from the TIR band, simultaneously. SWIR observations are more sensitive to CO₂ near the surface, but less sensitive to the vertical profiles than TIR observations. Thus, the use of the SWIR and TIR combined data provide the possibility to estimate the amounts of CO₂ in the boundary layer accurately and consequently may produce a useful dataset for the study of CO₂ sources and sinks. The main goal of the HIAPER Pole-to-Pole Observations (HIPPO) project is to provide the first high-resolution vertically resolved global survey of trace gases and aerosol investigating the Carbon Cycle and greenhouse gases annual cycle throughout various altitudes of the western hemisphere. In order to provide additional evidence to the ongoing discussion on the measurement of carbon dioxide from space, we performed a comparison between temporally reduced dataset of collocated IASI, GOSAT TIR and HIPPO data. In this work, we present the main results from this activity, developed as part of the collaboration between IFAC-CNR and the GOSAT team.