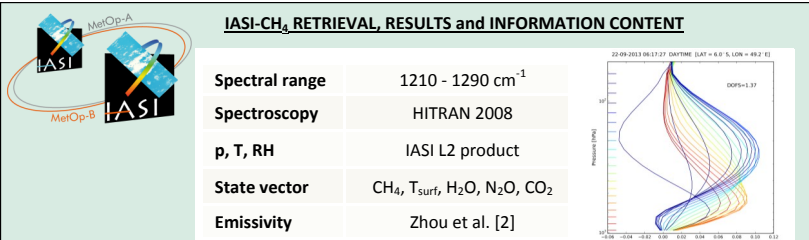


ABSTRACT : The IR team of BIRA has developed an automatized processing chain for the fast retrieval of tropospheric IASI CH₄ profiles with the ASIMUT Optimal Estimation Method (OEM) retrieval software [1]. Here we give a quality assessment of this new BIRA IASI CH₄ product and results of its validation with co-located NDACC ground-based observations. In addition first results will be shown of a comparison, on a global scale, with the IASI mid-tropospheric CH₄ product from Laboratoire de Météorologie Dynamique (LMD), that have been obtained as part of the ESA Climate Change Initiative Greenhouse Gas (GHG-CCI) project.



Tbl.1 : BIRA IASI-CH₄ retrieval set-up and typical IASI-CH₄ averaging kernel (AK). Sensitivity of the retrievals lies in the 4-17 km altitude range.

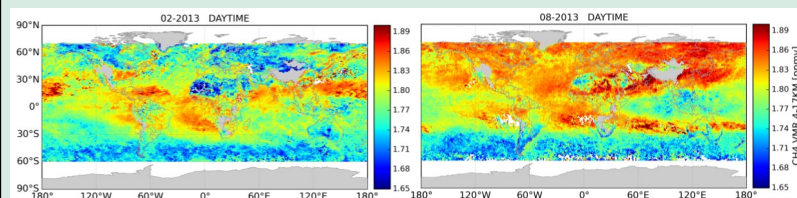


Fig. 1 : Monthly mean column averaged BIRA IASI-CH₄ between 4 and 17 km for February and August 2013. Only daytime retrievals are shown. We see a clear increase in CH₄ at high latitudes in NH summer probably due to increased wetland emissions.

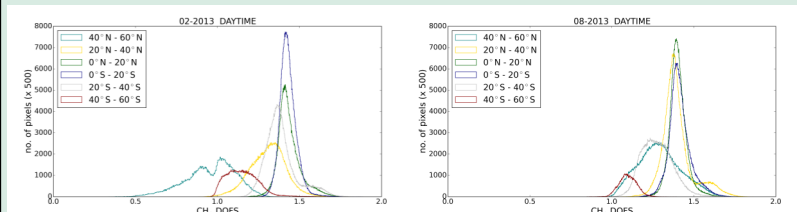


Fig. 2 : Degrees Of Freedom for Signal (DOFS) for different latitudinal bands for February and August 2013 (daytime retrievals). On the global scale, the values range between 1 and 1.7 for NH summer, and between 0.5 and 1.7 for NH winter, when values can become less than 1 for latitudes > 40° N (between 0.9 and 1.6, and between 0.35 and 1.6 respectively for nighttime retrievals; not shown here). In the tropics, DOFS are typically around 1.4, hence 1 atmospheric column is retrieved.

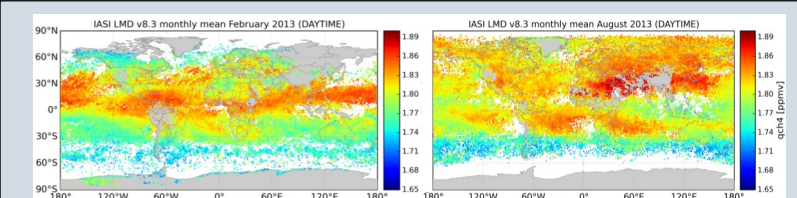


Fig. 7 : Monthly means of the LMD v8.3 IASI-CH₄ daytime product for February and August 2013. Given is the so-called qCH₄ in vmr, a mid-tropospheric column with peak sensitivity at about 230 hPa (~11 km) and half the peak sensitivity at 100 and 500 hPa (~6 and 16 km). The LMD IASI-CH₄ retrievals are based upon a non-linear regression inverse radiative transfer model using Multi-Layer Perceptrons [3]. Comparing Fig. 7 to Fig. 1 we see that similar features in the CH₄ distribution are captured by the two algorithms.

ACKNOWLEDGEMENT : The work presented is funded by the IASI.Flow PRODEX program and by the ESA GHG-CCI project.

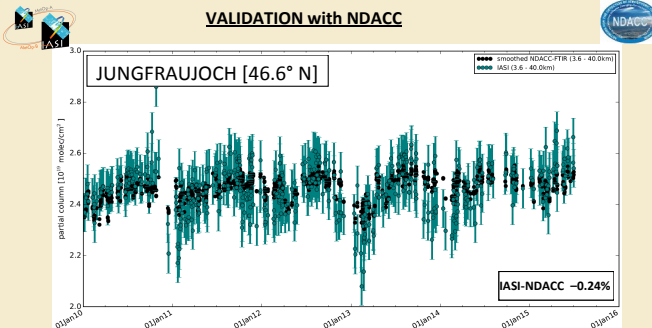


Fig. 4 : Timeseries of smoothed NDACC FTIR CH₄ [black] and BIRA IASI-CH₄ [blue] partial columns (surface-40 km) at Jungfraujoch. The vertical bars represent the retrieval error. The NDACC FTIR profiles are smoothed with the IASI AK and a common a priori profile is used for both IASI and NDACC retrievals. We see a good representation of the seasonal cycle by IASI and a mean relative difference of -0.24%.

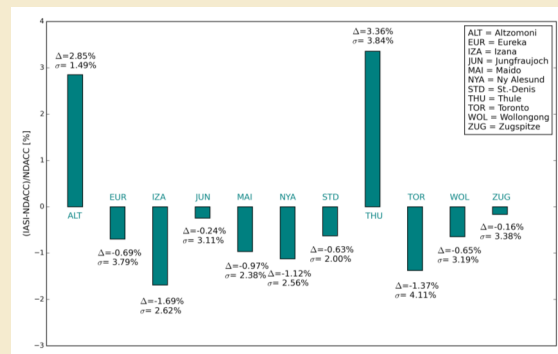


Fig. 5 : Barchart of smoothed NDACC-IASI CH₄ partial columns (surface-40 km) at 11 NDACC sites for the timeperiod 2010-2015. The mean (IASI-NDACC)/NDACC (Δ) and standard deviation (σ) of the relative differences is given for each site. We have an overall negative bias for IASI < 1.7% with exception of the sites Altimoni and Thule, who show a positive bias.

COMPARISONS with IASI-LMD CH₄

As part of ESA GHG-CCI the BIRA IASI CH₄ fields are compared to the LMD CH₄ product on a global scale.

For the comparison the BIRA qCH₄ is calculated as : $q_{BIRA} = \frac{\sum_{i=1}^n H_i \Delta p_i x_i}{\sum_{i=1}^n H_i \Delta p_i}$ where H_i is the LMD weighting function interpolated to the BIRA pressure grid. n is the number of BIRA pressure layers Δp_i and x_i the BIRA CH₄ profile in vmr.

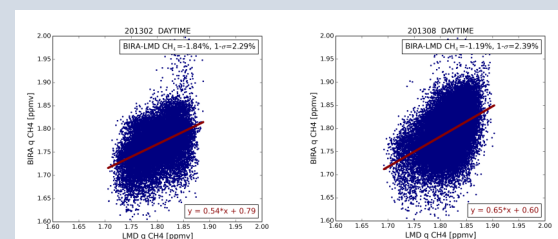


Fig. 6 : Correlation plots of LMD qCH₄ and BIRA qCH₄ for February and August 2013. We see overall lower values of BIRA qCH₄ wrt LMD qCH₄ with relative differences of -1.84% for February and -1.19% for August, and a standard deviation (1-σ) of the difference below 2.4%.

[1] Vandaele, A.C., et al., Modelling and retrieval of Atmospheric spectra using ASIMUT in Proc. of the First 'Atmospheric Science Conference', ESIRIN, Frascati, Italy, 8 – 12 May 2006.

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[3] Crevoisier, C., Nobileau, D., Fiore, A., Armande, R., Chédin, A., and Scott, N. A.: Tropospheric methane in the tropics - first year from IASI hyperspectral infrared observations, Atmos. Chem. Phys., 9, 6337-6350, 2009.