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HCOOH distributions from IASI with updated retrieval parameters: comparison with ground-based FTIR measurements and IMAGESv2 model

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European Space Agency

Tropospheric formic acid (HCOOH)

- HCOOH is **one among the most abundant volatile organic compounds** (VOCs) present in the atmosphere.

- With acetic acid it is a major **contributor to the acidity of precipitation**

• Many sources:

secondary product from other organic precursors + small direct emissions by vegetation, ants, biomass burning, soils, agriculture, motor vehicles.

• Sinks:

mainly removed through wet and dry deposition + (lesser extent) oxidation by the OH radical

HCOOH = short-lived species

lifetime is conditioned by the ratio of precipitation: in the boundary layer 2 days (rainy period) \rightarrow 6 days (the dry season). **Global lifetime in the troposphere=3–4 days** Photochemical loss is relatively slow ($\tau \sim 25$ days), so that any HCOOH formed or vented outside of the boundary layer **can be transported for long distances in the free**

troposphere.



Recent studies highlight:

A misrepresentation of emission from tropical and boreal forest in models (Stavrakou et al., Nature 2011)

A possible source of HCOOH over the Arctic Ocean (Jones et al., Atmos. Env 2014).

One or more large missing sources (Millet et al., ACPD 2015) \rightarrow suggest a gap in our current understanding of hydrocarbon oxidation or the existence of an unknown direct flux

HCOOH: observations from space



Space missions allow getting global observations of the atmosphere



Their spatial coverage allows observing remote regions which are sparsely studied by field campaigns



Limited vertical sensitivity

few satellites provide tropospheric HCOOH observations

nadir-viewing instruments: Infrared Atmospheric Sounding Interferometer (IASI) (e.g. Coheur et al., 2009), Tropospheric Emission Spectrometer (TES) (e.g. Cady-Pereira et al., 2014)

solar-occultation instrument: Atmospheric Chemistry Experiment (ACE) (e.g. González Abad, 2009)

limb instrument: Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) \rightarrow global distribution of HCOOH at 10 km (Grutter et al., 2010).

HCOOH: observations with IASI





 \rightarrow signal from HCOOH expressed in Δ BT

HCOOH: observations with IASI





First global distribution by IASI (Razavi et al., ACP 2011)

Detection of extreme events: forest fires in Russia on 2010 (R'Honi et al., ACP 2013)

Retrieval relevant for huge amount of HCOOH



HCOOH: Retrieval approach (1/2)



MEGAN-MACC HCOOH 40 emissions for the period 20 between 2008 and 2010 on a 0.5°×0.5° grid. (Sindelarova, ACP 2014)

FTIR sites location



Region	Localization	Number of retrieved spectra
Africa N	6-7°N 18-22°E	265
Africa S	12-14°S 20-24°E	788
Amazonia	6-10°S 43-45°W	682
Atlantic	22-24°N 42-45°W	675
Australia	14-15°S 131-133°E	218
Pacific	20-22°S 140-142°E	472
Russia	50-54°N 60-62°E	538

Data retrieved over 7 regions → emission sources, remote area, areas influenced by longrange transport + land and sea scenes

Seasonal variation is taken account: 5 first days of each month on 2009 are retrieved

HCOOH: Retrieval approach (2/2)

Parameters:

- 1095-1114 cm⁻¹
- Cloud fraction < 2%
- Same a priori than Razavi et al. (2011)
- Thermal contrast > 0 K (T _{surf} T _{atm 1st layer}) (Razavi et al. used TC >5K)
- day-time and night-time data used



Good correlation between retrieved columns and ΔBT

But conversion is dependent on the TC

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Establishment of a new dataset

Reduction of the thermal contrast dependence: Mean over 7 regions



Conversion using both relationships: Column = 1.742 × (ΔBT – TC × 0.0361 - 0.1969) + 0.646

Error estimation



3 terms:

- 1) the instrumental error
- 2) the error caused by the conversion between the ΔBT and the total column
- 3) the error from the retrievals.

total error by forward simulation (>3000 spectra):

- Gaussian distributed random noise (σ = 0.15K) added to the BT channels
- the conversion formula is applied on the calculated ΔBT .



Comparison with FTIR (± 0.5°, daily mean)





Seasonal Variation (global distribution) Cesa

IASI







Conclusions



- Global overview of HCOOH
- 7 years of data to analyze
- Detection hotspots (MiddleEast US, Asia...) & seasonal cycle
- No averaging kernels → difficulties to compare IASI total columns to FTIR and simultations
- Challenge in the interpretation of the data
 → misrepresentation of sources





?



Extra slides







IASI

Seasonal Variation (global distribution) 🏼 🖉 esa

MAM 2008-2013





-0.8 -0.6 -0.4 -0.2 0 0.2 0.4 0.6 0.8 IASI - IMAGESv2 (10¹⁶ molec.cm⁻²)

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Detection of extreme events: fires over Russia

0.2°×0.2° 27 July – 27 August 2010 over Russia



HCOOH total column

CO total column

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+60°

2.5

2

3

+90°

3.5

Detection of extreme events: fires over Russia

0.2°×0.2° 27 July – 27 August 2010 over Russia



HCOOH total column

MODIS hotspot

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3

+90°

5