

Retrieval of SO₂ from high spectral resolution measurements: AIRS and IASI

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Why measure SO₂?

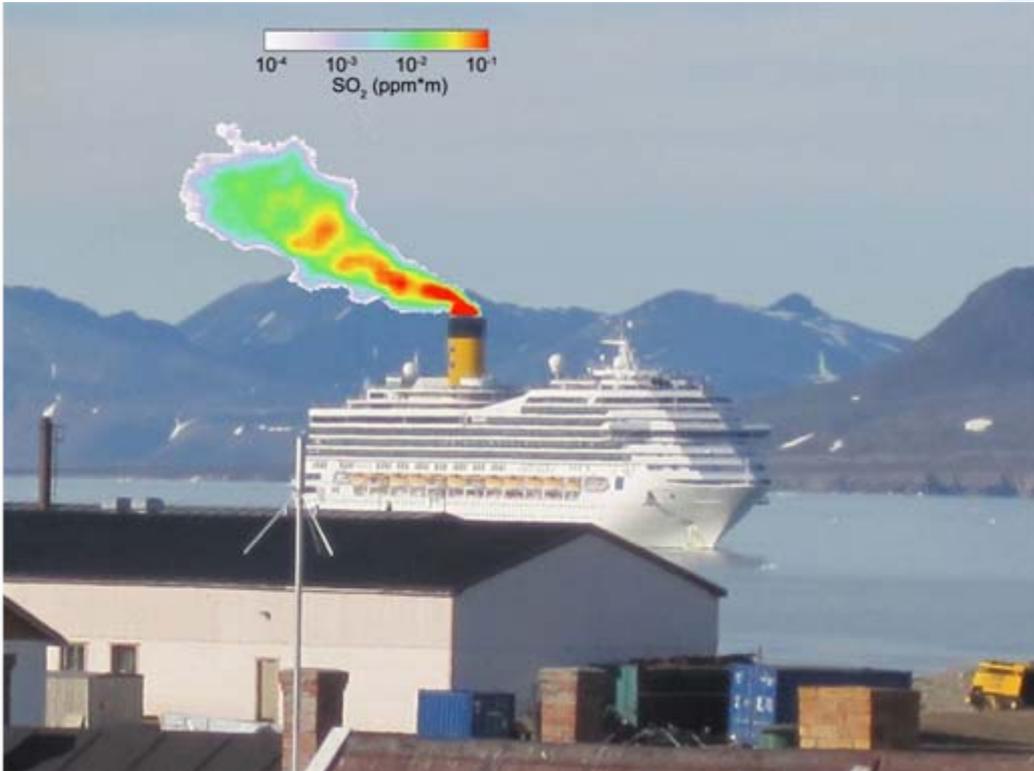
Atmospheric pollution

Effects on climate

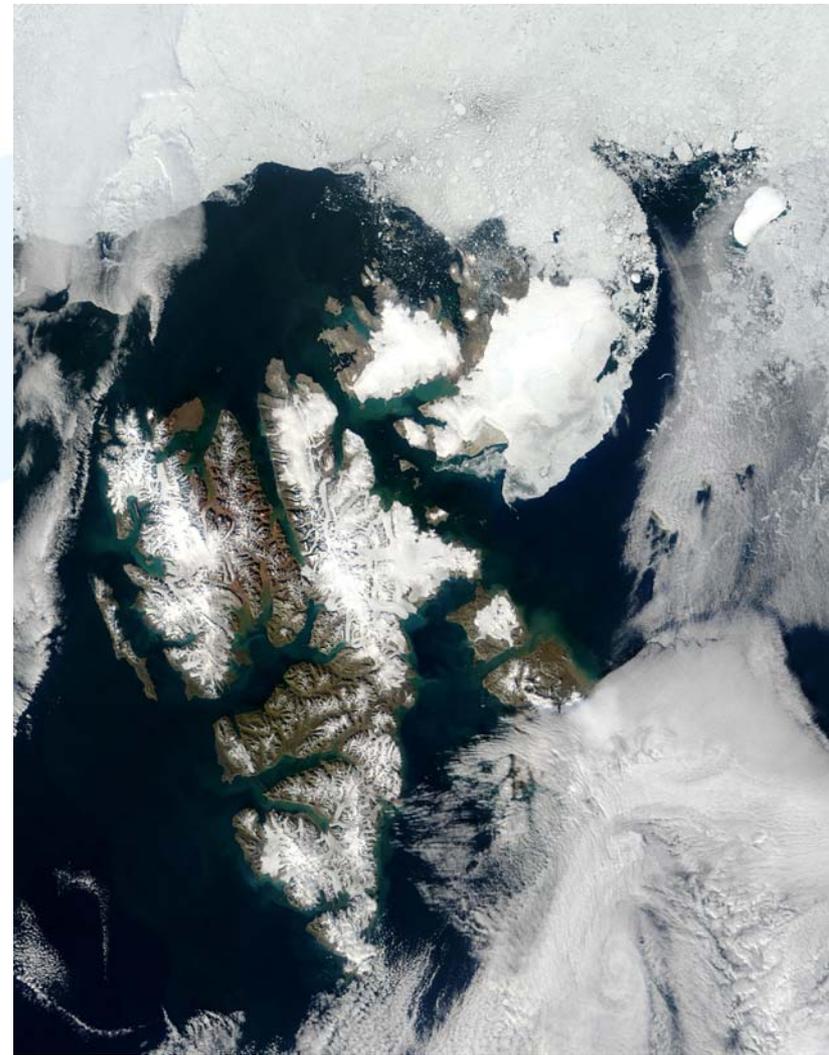
Hazards to aviation

- Background levels generally low $<10^{-4}$ ppmV
- Man-made emissions from industry, mining operations, shipping
 - emission rates from 10 gs^{-1} to $>1000 \text{ kgs}^{-1}$
- Large natural emissions from volcanoes
 - passive degassing $>10 \text{ kgs}^{-1}$
 - explosive volcanism $>10^3 \text{ kgs}^{-1}$

Atmospheric Pollution



Measurements from NILU's ground-based UV camera – EnviCam



SO₂ ship emissions from a large cruise ship entering Kongsfjorden, Ny Ålesund, Svalbard.

Volcanic emissions



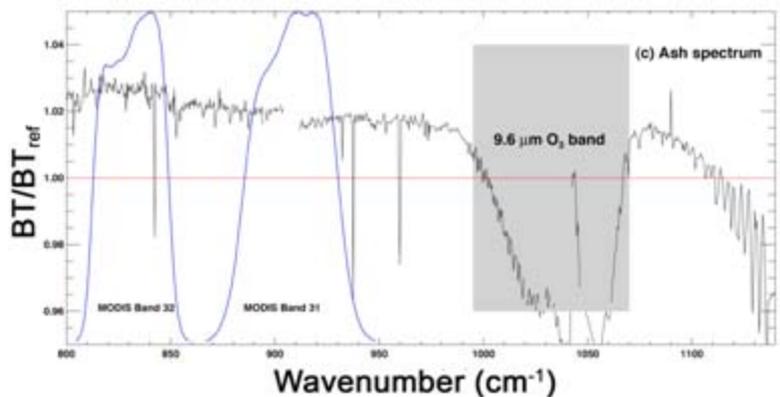
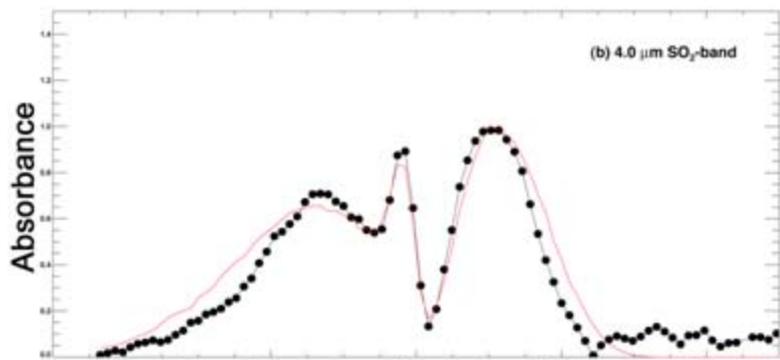
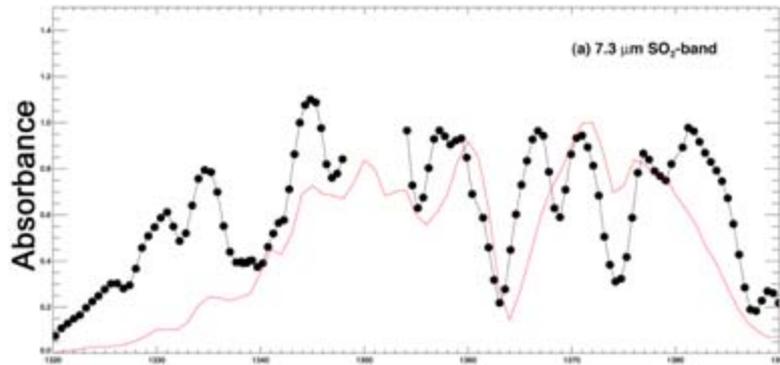
Villarica volcano, Chile.
Passive degassing



Soufriere Hills, Montserrat explosive eruption, 11 February 2010



Ground-based IR camera (CyClops) at Tavurvur volcano, New Britain.



AIRS and IASI can measure SO_2 absorption in the strong ν_3 ($1363 \text{ cm}^{-1}/7.3 \mu\text{m}$) band of SO_2

AIRS and IASI can also measure SO_2 absorption in the weak combination band $\nu_1 + \nu_3$ ($2500 \text{ cm}^{-1}/4 \mu\text{m}$) band of SO_2

Ash can be measured between $800\text{--}1130 \text{ cm}^{-1}$

Retrieval scheme

$$\begin{aligned}
 I_\nu &\approx \int_0^{z_1} B_\nu[T(z)] \left(\frac{\partial \tau_\nu[z, q_2(z) \dots q_n(z)]}{\partial z} \right) dz \\
 &+ \int_{z_1}^{z_2} B_\nu[T(z)] \left(\frac{\partial \tau_\nu[z, q_1(z), q_2(z) \dots q_n(z)]}{\partial z} \right) dz \\
 &+ \int_{z_2}^{\infty} B_\nu[T(z)] \left(\frac{\partial \tau_\nu[z, q_2(z) \dots q_n(z)]}{\partial z} \right) dz.
 \end{aligned}$$

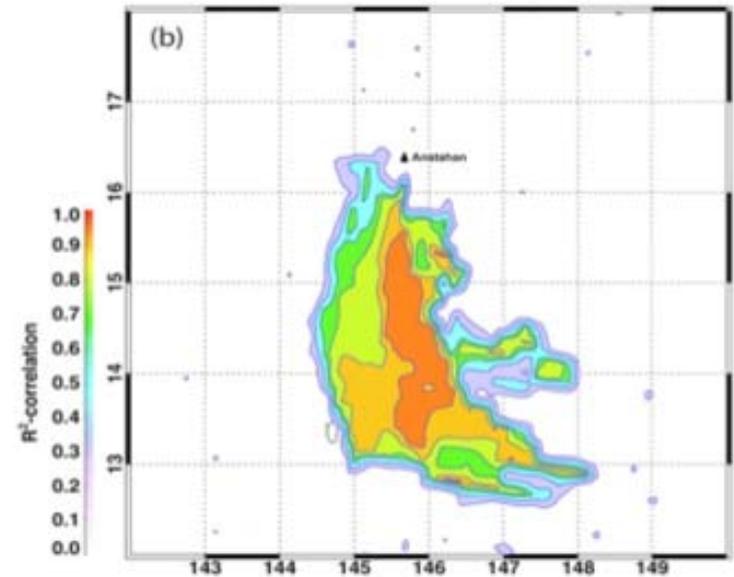
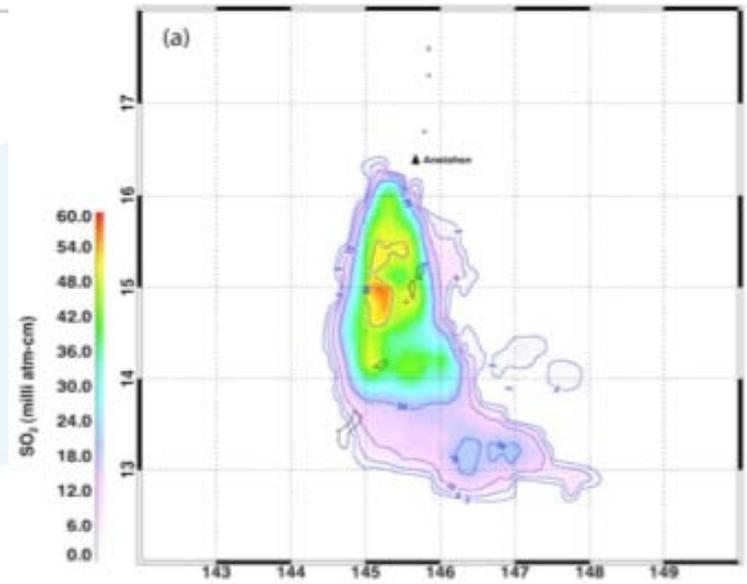
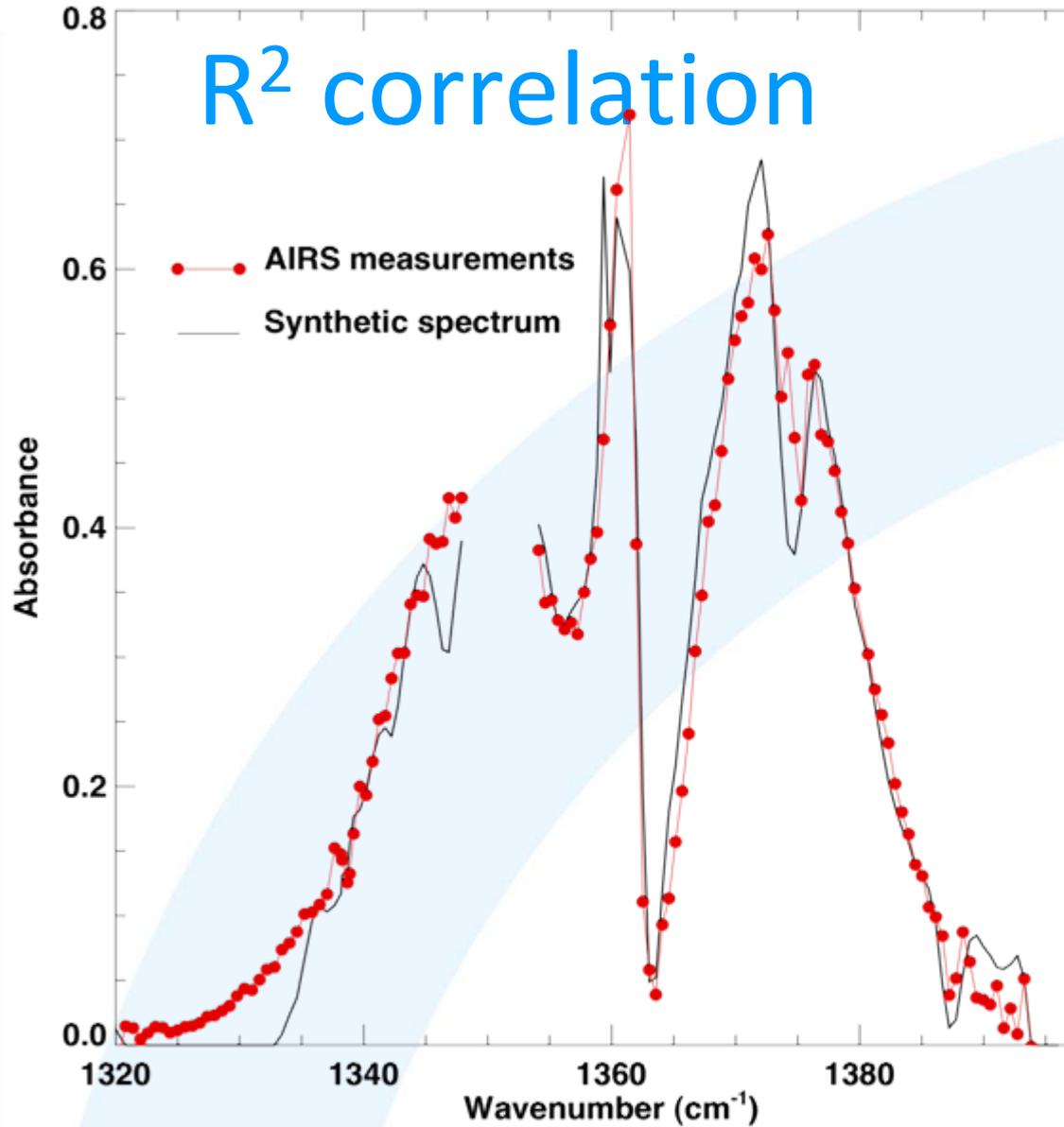
$$I'_\nu = I_{\nu,0} \exp \left\{ - \int_0^{\infty} k_\nu(z) q(z) dz \right\}; \quad A_\nu = -\ln \left\{ \frac{I'_\nu}{I_{0,\nu}} \right\} = \int_0^{\infty} k_\nu(z) q(z) dz.$$

Absorbing layer only

$$A_\nu = -Ln \left\{ \frac{I_{pr,lr}}{I_{pr,lr}} \right\} \quad S_\nu = -Ln \left\{ \frac{I_s}{I_0} \right\}$$

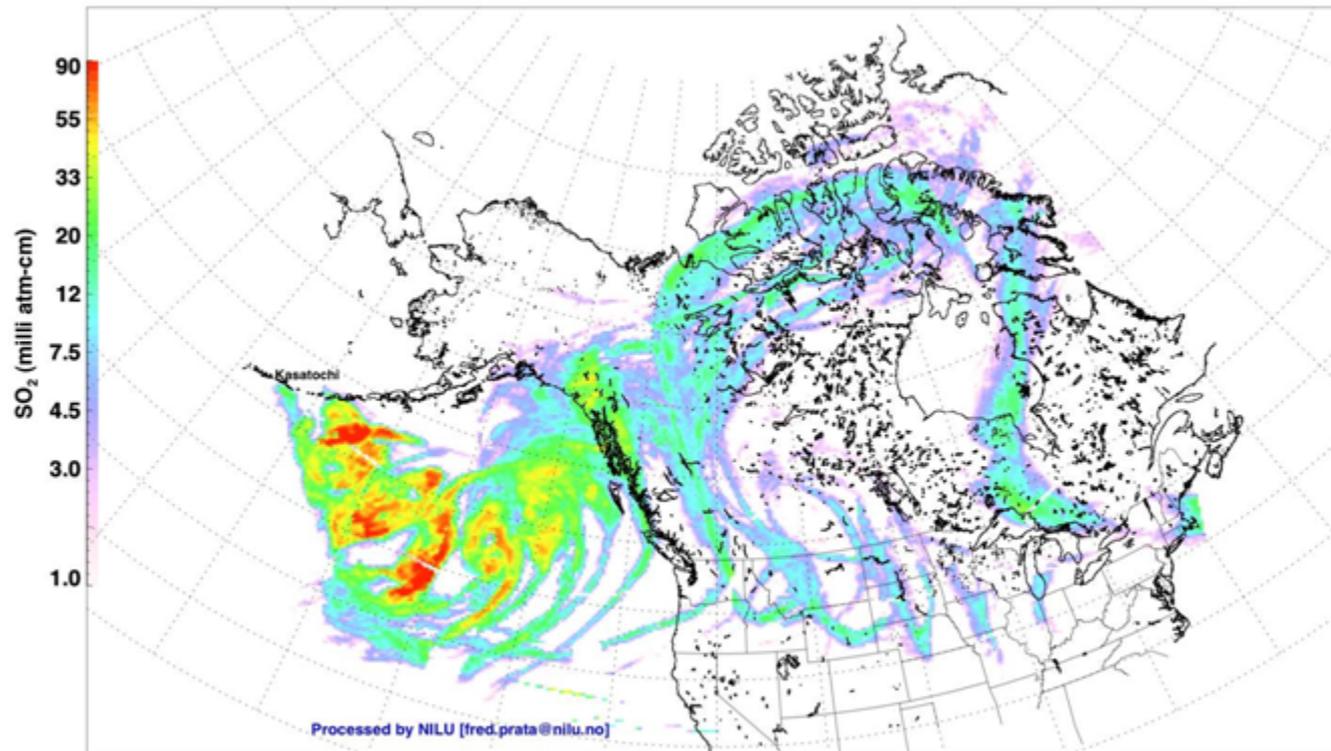
$$R = \frac{\frac{1}{n-1} \sum_{i=0}^{n-1} \tilde{A}_i \tilde{S}_i}{\sqrt{\frac{1}{n-1} \sum_{i=0}^{n-1} \tilde{A}_i^2} \sqrt{\frac{1}{n-1} \sum_{i=0}^{n-1} \tilde{S}_i^2}}.$$

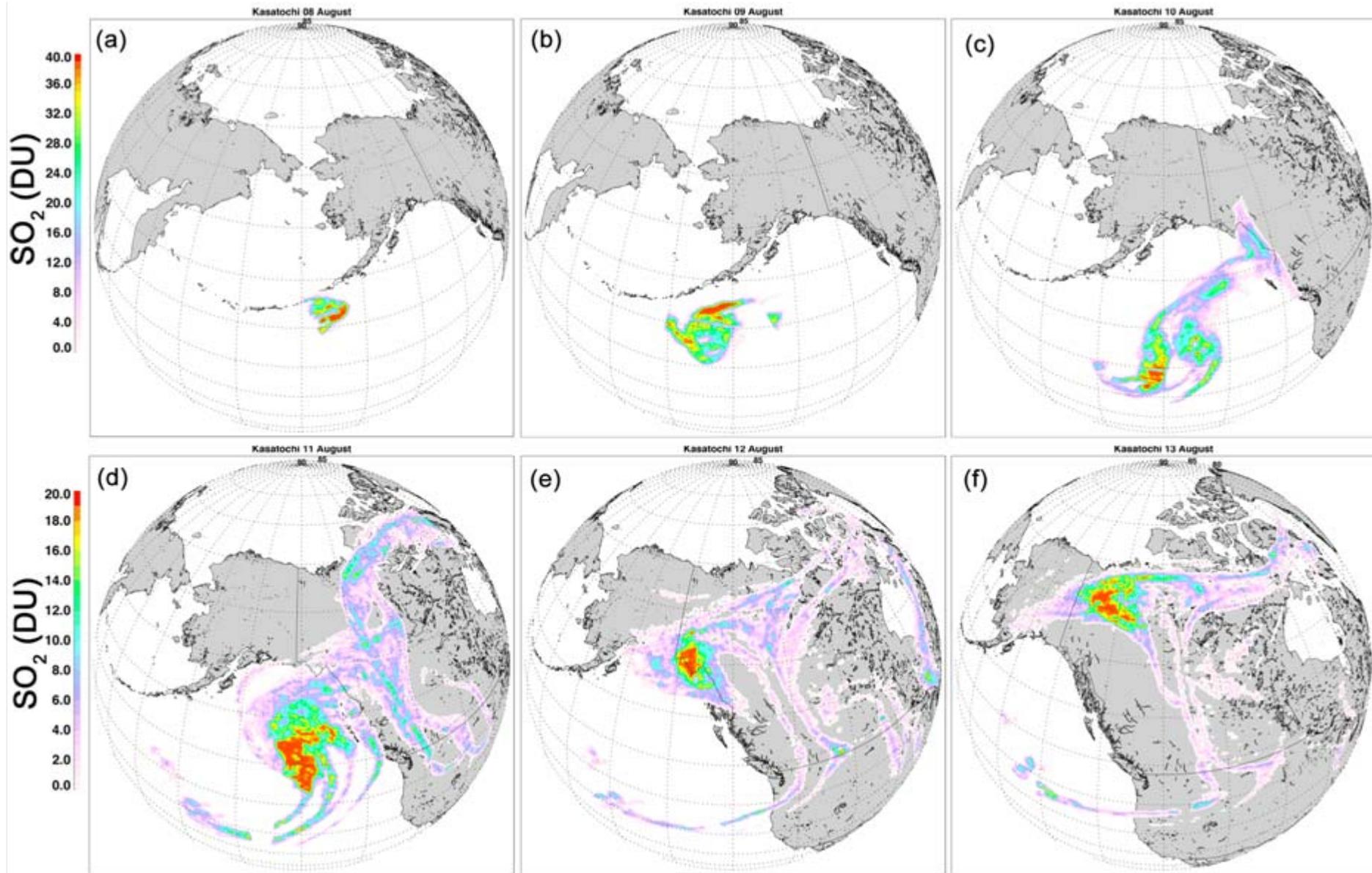
R² correlation



AIRS SO₂ using the strong ν_3 (7.3 μm band)

Kasatochi AIRS 7.3 μm Cumulative SO₂ 8-13 August, 2008



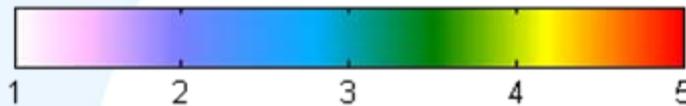
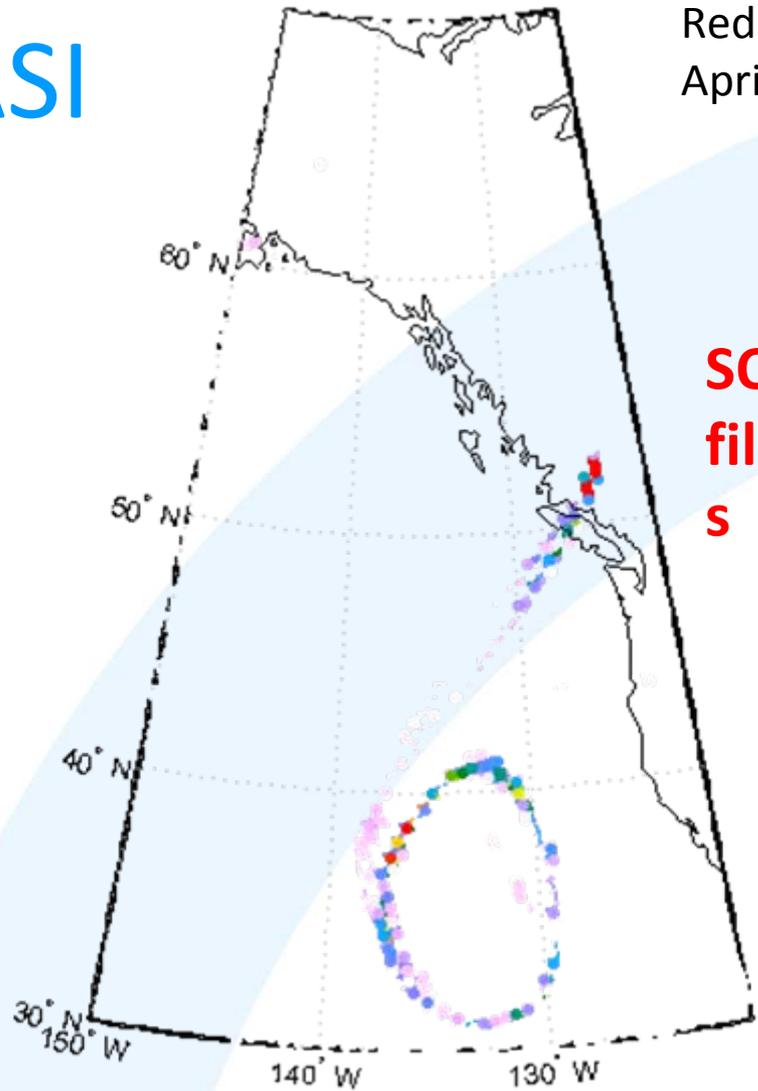


Kasatochi AIRS retrievals: 8–13 August 2008. >1.2 Tg (SO₂)

IASI

Redoubt, Alaska:
April 5 2009

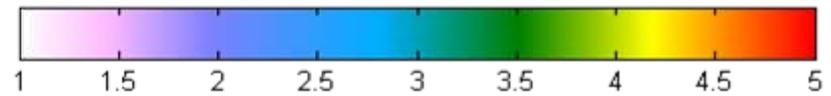
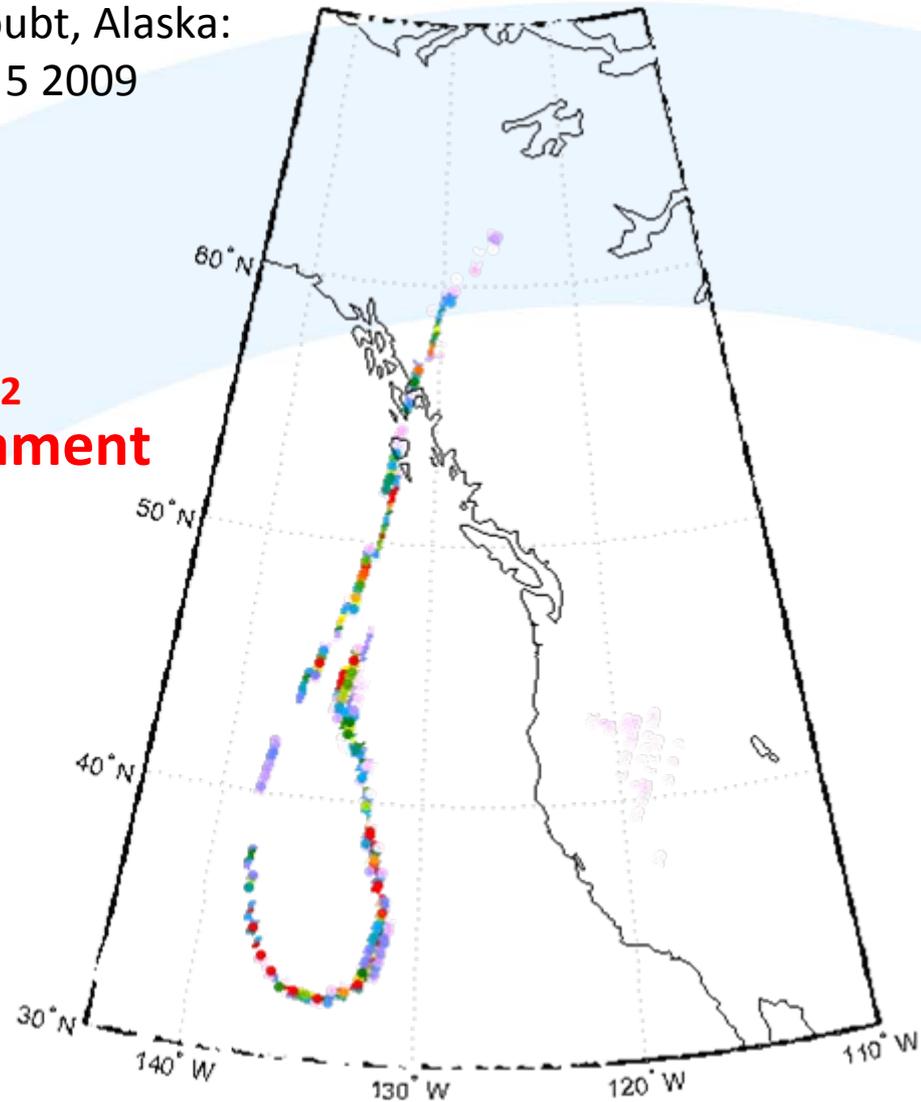
**SO₂
filament
s**



14 April 2010

ΔBT (K)

ITSC-17 Monterey, California

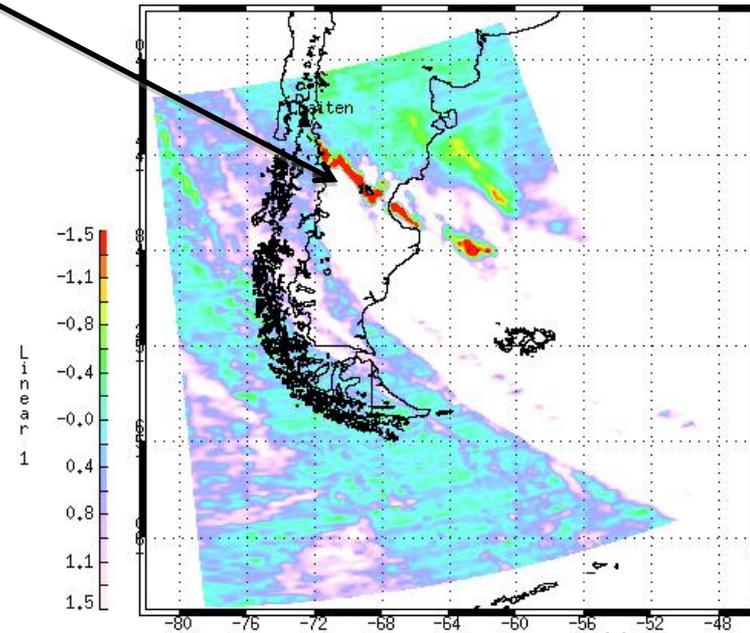
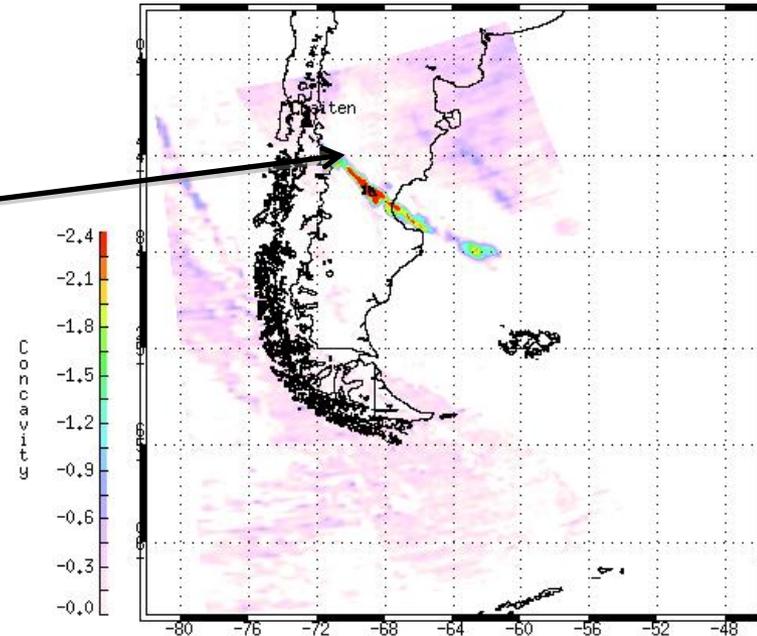
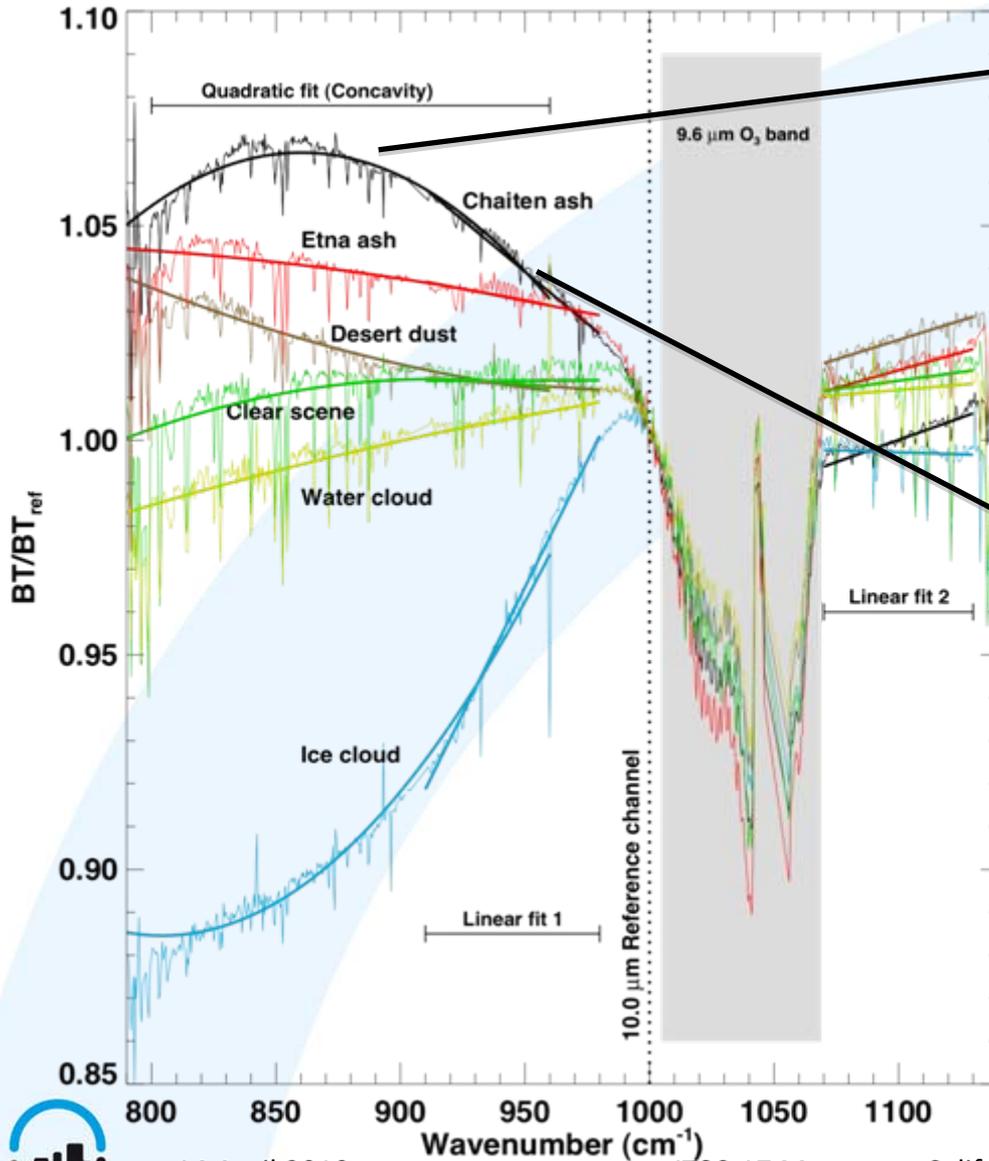


ΔBT (K)

10



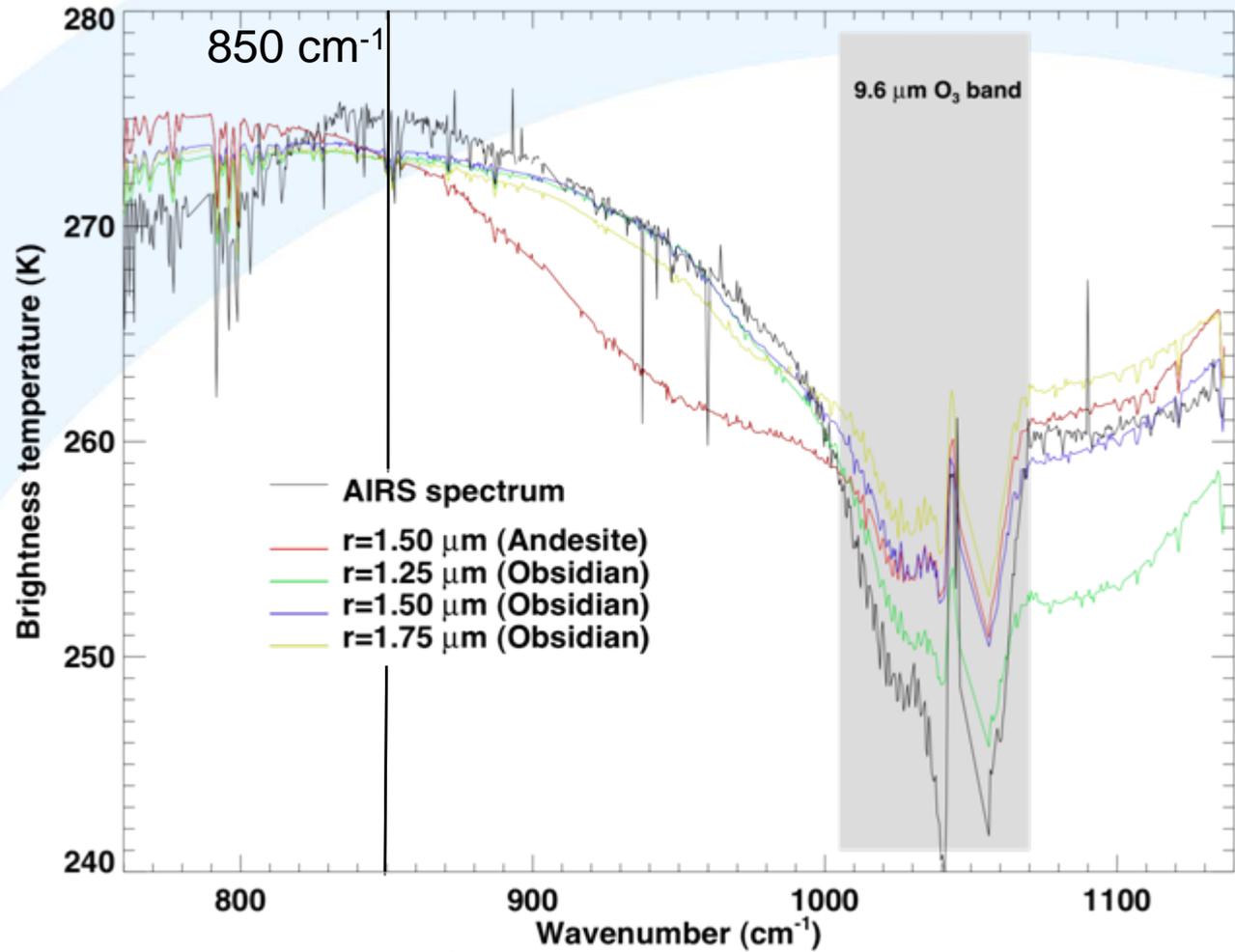
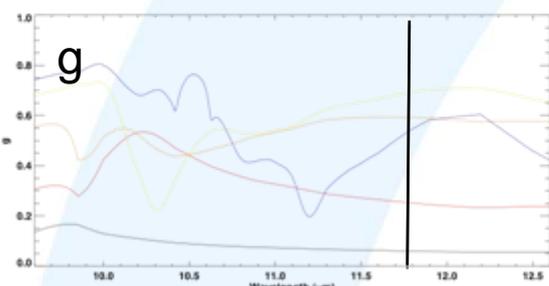
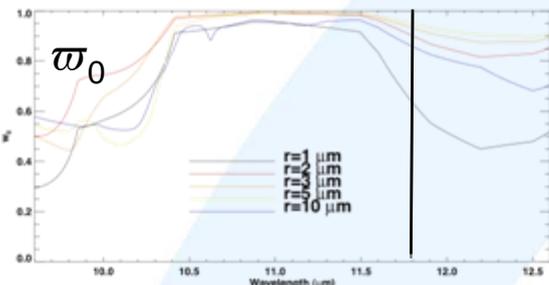
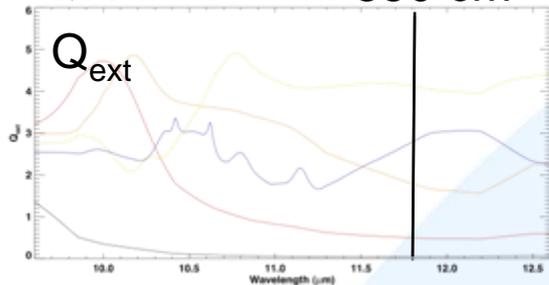
Spectral Signatures

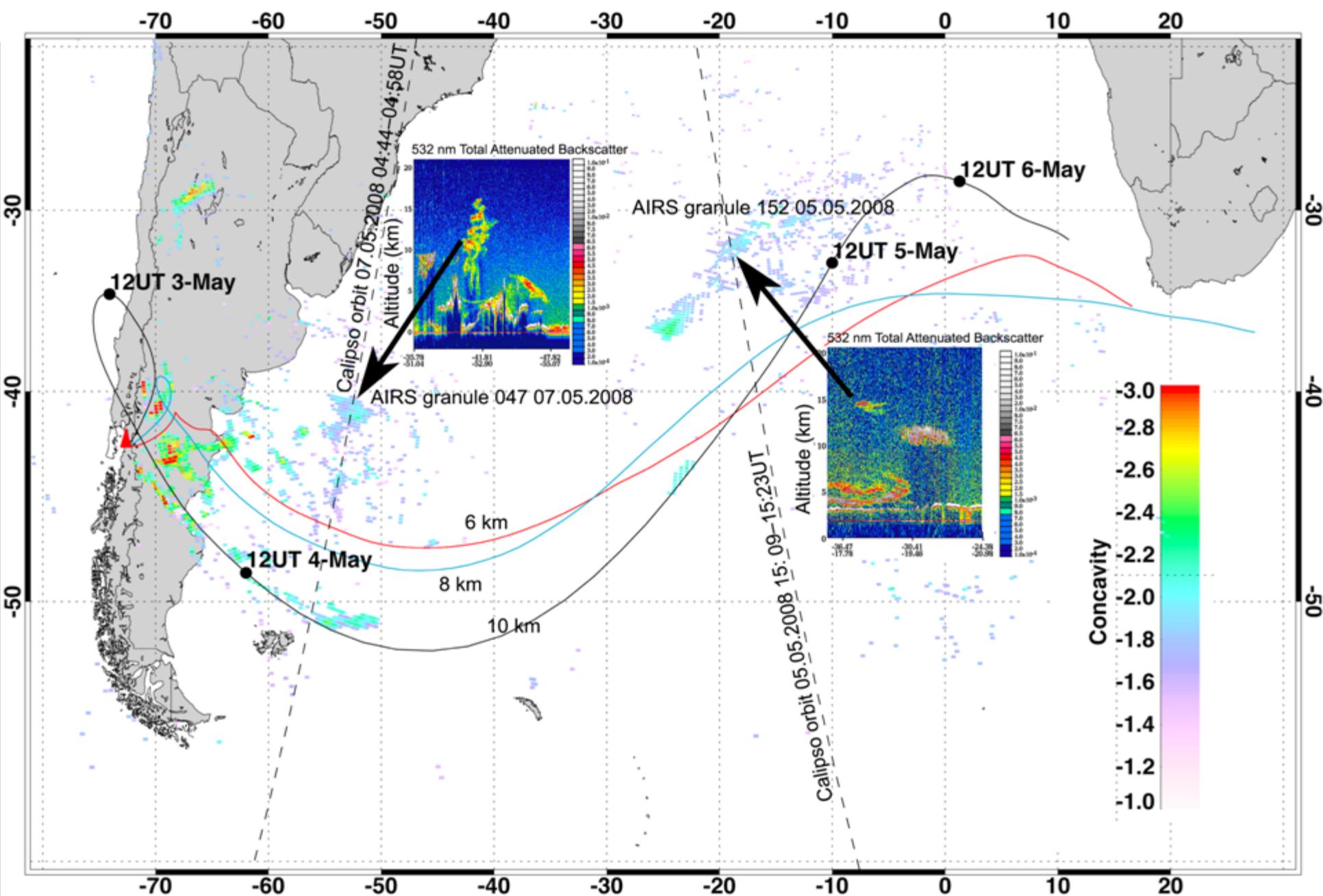


Microphysics retrieval

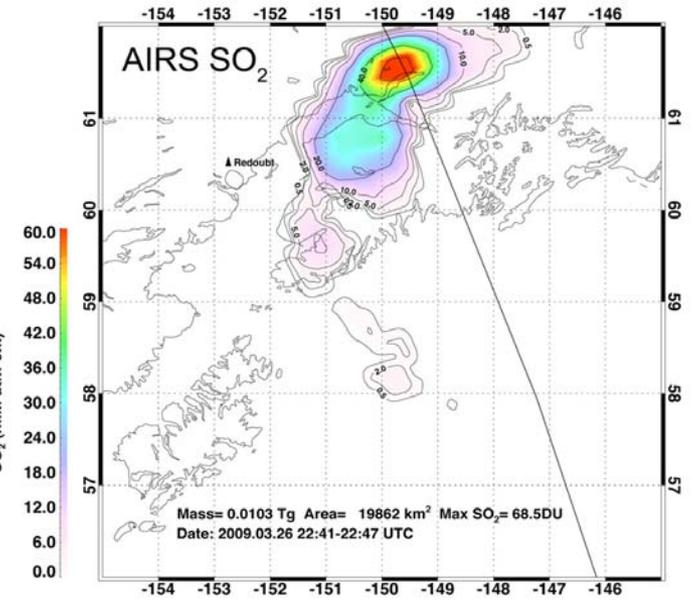
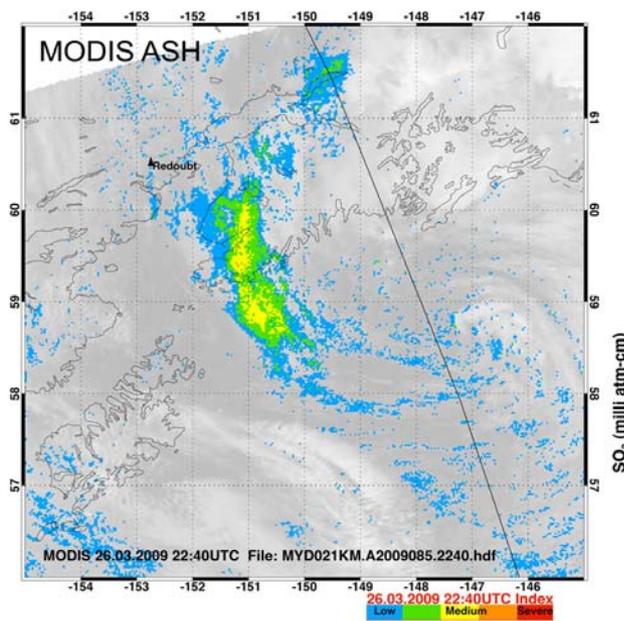
r_e – effective particle radius
 τ – infrared optical depth
 M – mass loading
 Si – silicate composition

Quartz 850 cm^{-1}

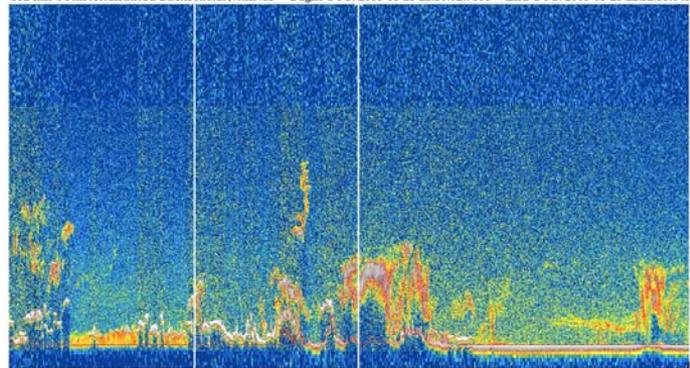




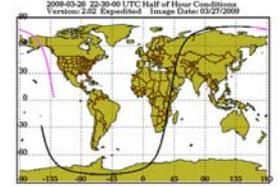
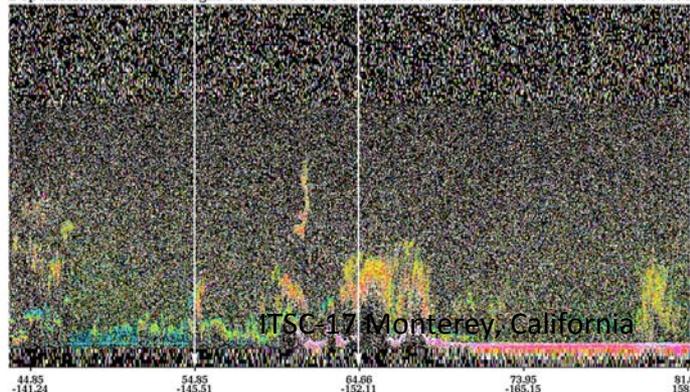
MODIS AIRS CALIOP Comparisons



532 nm Total Attenuated Backscatter, /km /sr Begin UTC: 2009-03-26 22:30:02.7801 End UTC: 2009-03-26 22:52:36.0431



Depolarization Ratio Begin UTC: 2009-03-26 22:30:02.7801 End UTC: 2009-03-26 22:52:36.0431



14 April 2010

IASI H_2SO_4 detection

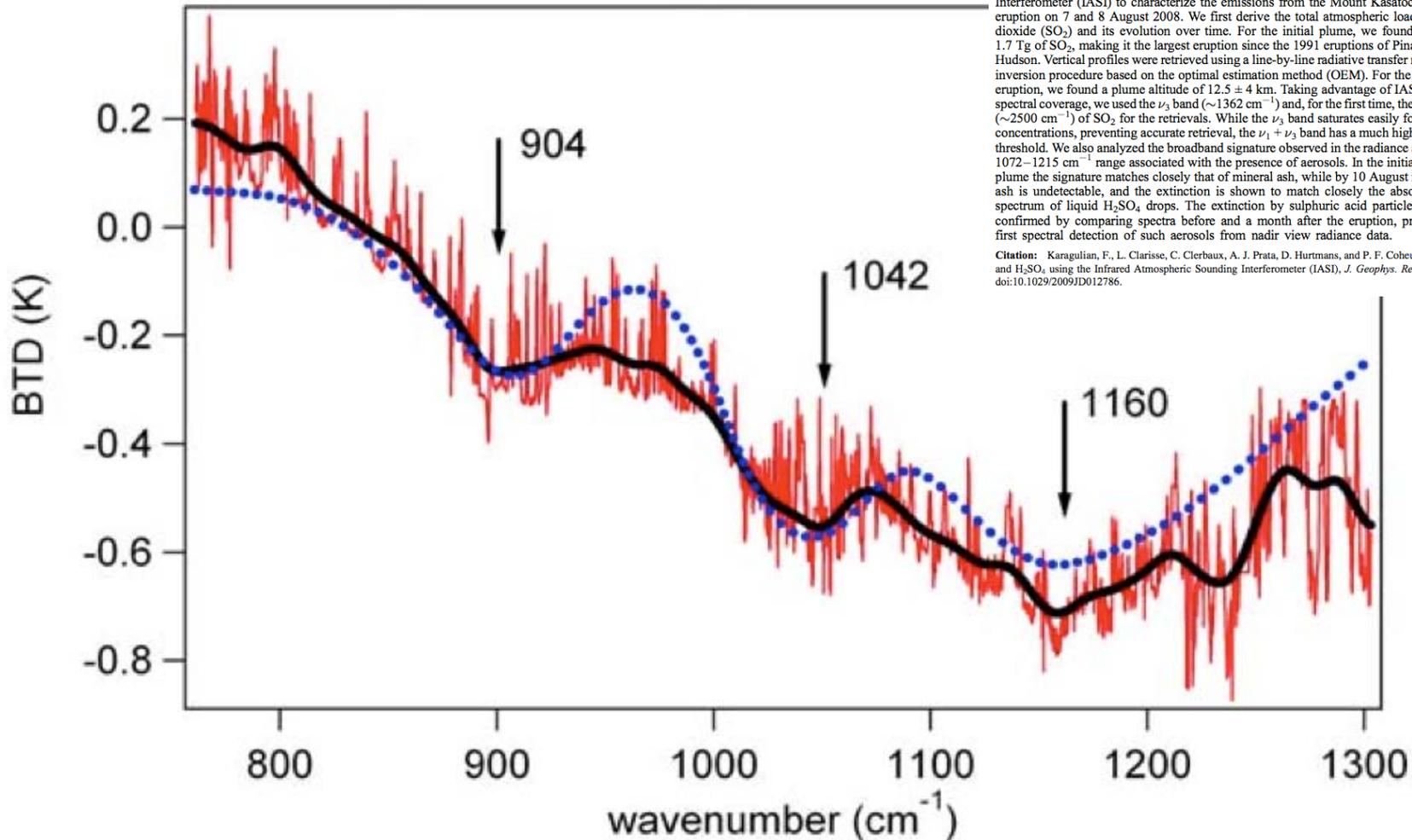
Detection of volcanic SO_2 , ash, and H_2SO_4 using the Infrared Atmospheric Sounding Interferometer (IASI)

F. Karagulian,¹ L. Clarisse,¹ C. Clerbaux,^{1,2} A. J. Prata,³ D. Hurtmans,¹ and P. F. Coheur¹

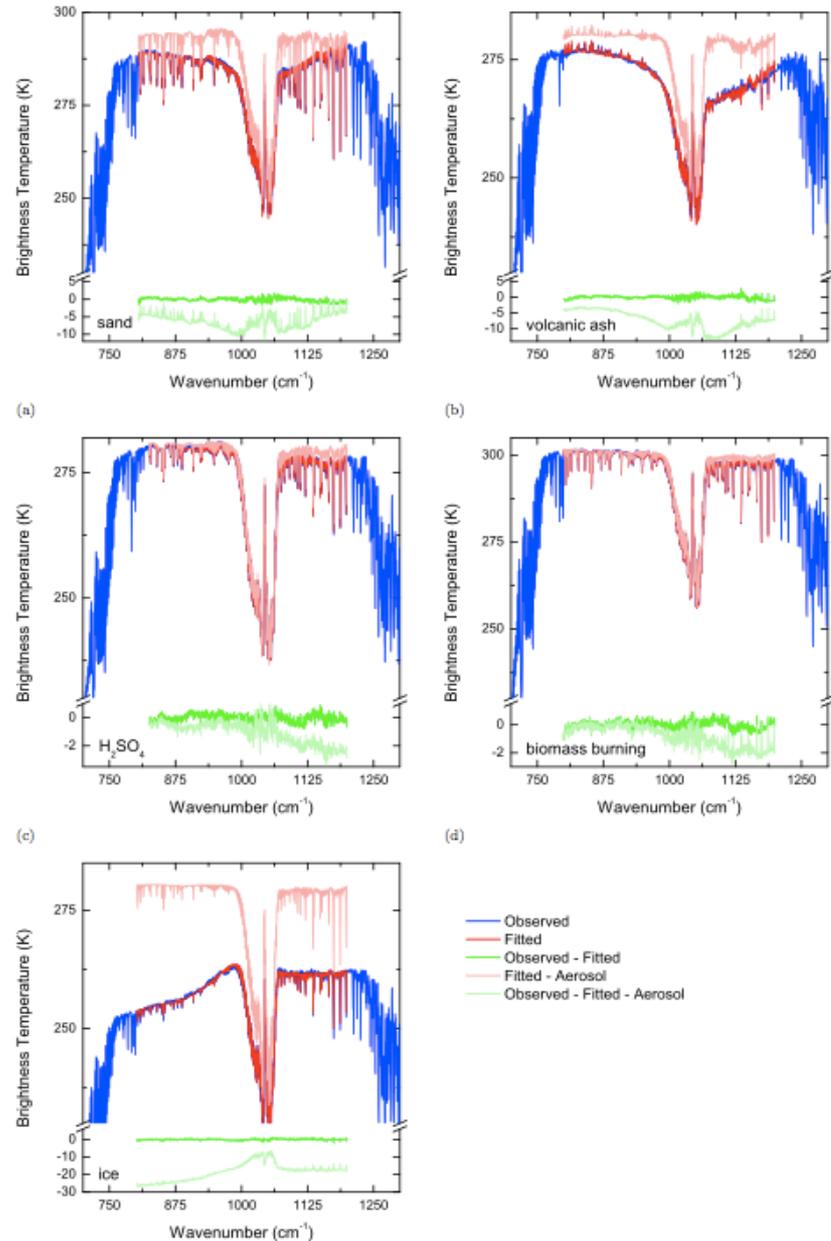
Received 6 July 2009; revised 24 September 2009; accepted 2 October 2009; published 27 February 2010.

[1] In this work we use infrared spectra recorded by the Infrared Atmospheric Sounding Interferometer (IASI) to characterize the emissions from the Mount Kasatochi volcanic eruption on 7 and 8 August 2008. We first derive the total atmospheric load of sulfur dioxide (SO_2) and its evolution over time. For the initial plume, we found values over 1.7 Tg of SO_2 , making it the largest eruption since the 1991 eruptions of Pinatubo and Hudson. Vertical profiles were retrieved using a line-by-line radiative transfer model and an inversion procedure based on the optimal estimation method (OEM). For the Kasatochi eruption, we found a plume altitude of 12.5 ± 4 km. Taking advantage of IASI's broad spectral coverage, we used the ν_3 band ($\sim 1362 \text{ cm}^{-1}$) and, for the first time, the $\nu_1 + \nu_3$ band ($\sim 2500 \text{ cm}^{-1}$) of SO_2 for the retrievals. While the ν_3 band saturates easily for high SO_2 concentrations, preventing accurate retrieval, the $\nu_1 + \nu_3$ band has a much higher saturation threshold. We also analyzed the broadband signature observed in the radiance spectra in the $1072\text{--}1215 \text{ cm}^{-1}$ range associated with the presence of aerosols. In the initial volcanic plume the signature matches closely that of mineral ash, while by 10 August most mineral ash is undetectable, and the extinction is shown to match closely the absorption spectrum of liquid H_2SO_4 drops. The extinction by sulphuric acid particles was confirmed by comparing spectra before and a month after the eruption, providing the first spectral detection of such aerosols from nadir view radiance data.

Citation: Karagulian, F., L. Clarisse, C. Clerbaux, A. J. Prata, D. Hurtmans, and P. F. Coheur (2010), Detection of volcanic SO_2 , ash, and H_2SO_4 using the Infrared Atmospheric Sounding Interferometer (IASI), *J. Geophys. Res.*, 115, D00L02, doi:10.1029/2009JD12786.

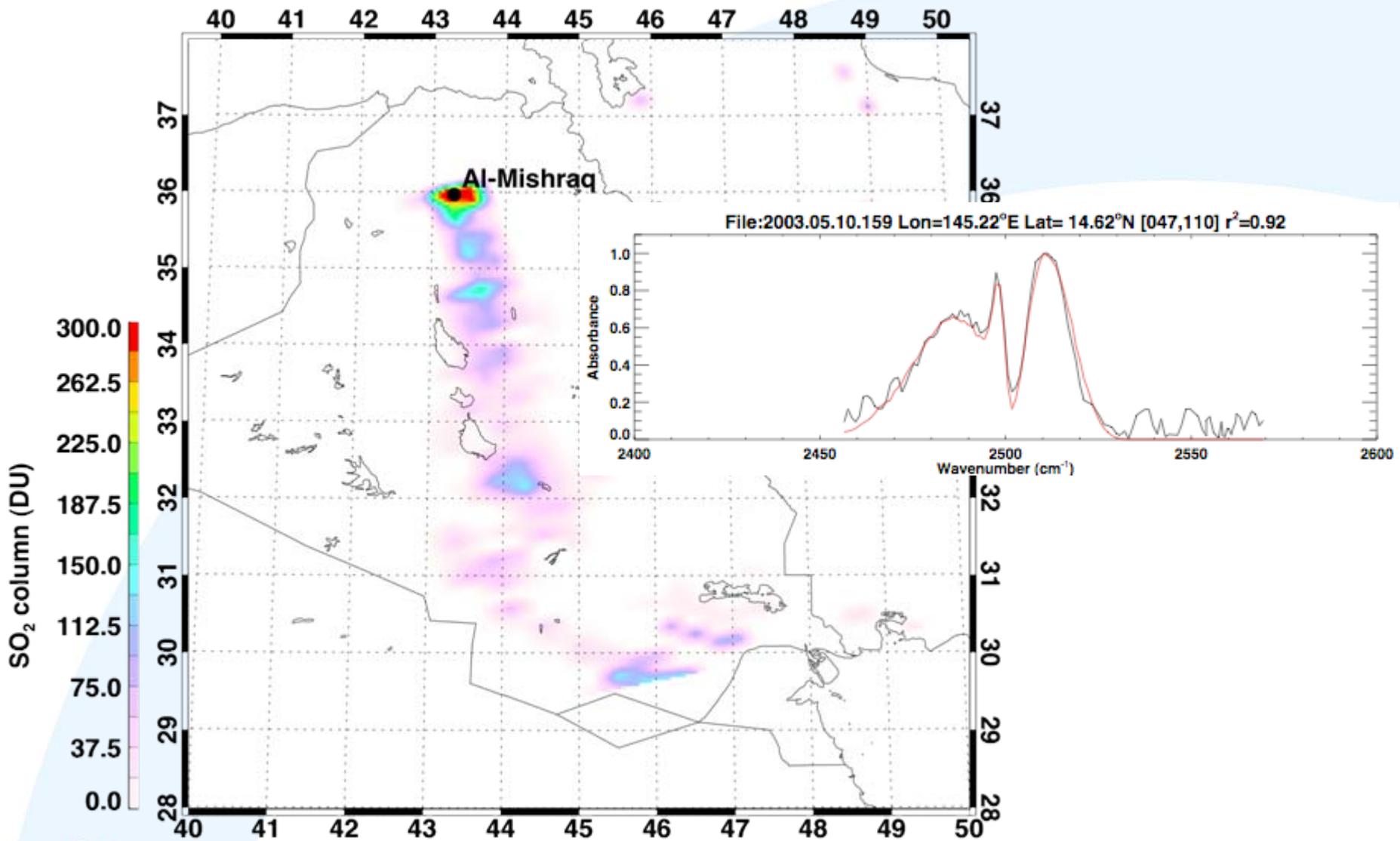


Detection of other species



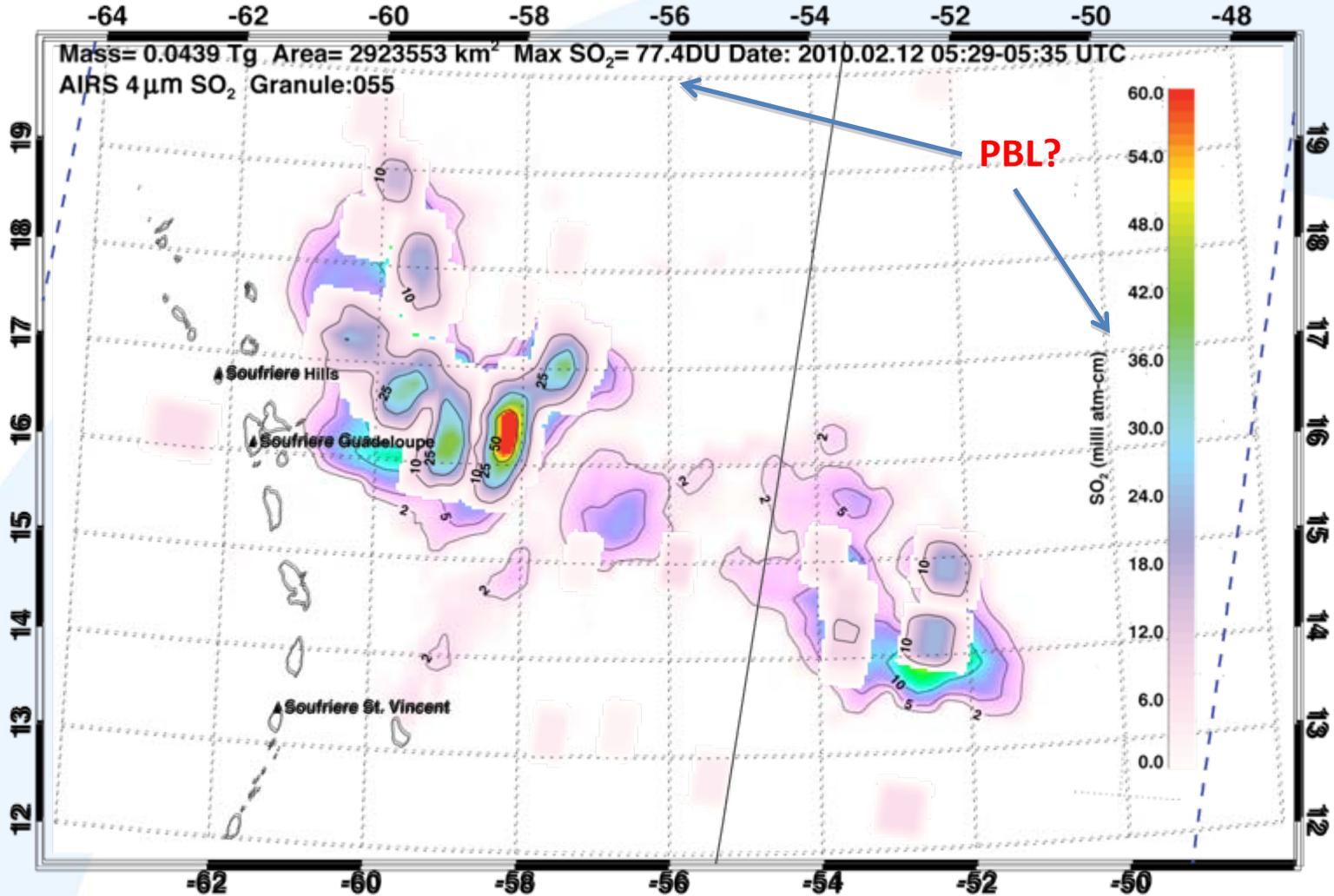
L. Clarisse, D. Hurtmans, A.J. Prata, F. Karagulian, C. Clerbaux, M. De Mazière and P.-F. Coheur, Retrieving radius, concentration, optical depth and mass of different types of aerosols from high resolution infrared nadir spectra, *Submitted to Applied Optics, 2010*.

Boundary layer SO₂ at 4 μm



Mass= 0.0335 Tg Area= 257792 km² Max SO₂= 29.0DU Date: 2010.02.12 05:29:24.000UTC

UTLS



PBL?

Soufriere Hills, 12 February 2010



Combining satellite data with a Lagrangian dispersion model

- Volcanic ash and SO_2 can be placed into the atmosphere at different levels
- Wind shear ensures that ash and SO_2 travel in different directions and at different speeds
- Is there a better way to utilise dispersion models and satellite data to provide predictions for aviation?

Inverse Modelling- Analytic method

Source-receptor matrix calculation with a Lagrangian particle dispersion model in backward mode, P. Seibert and A. Frank, ACP, 4, 51-63, 2004.

Sources x (1..n) x^a a priori profile

Satellite observation y^o (1..m)

M Emission sensitivity Matrix ($m \times n$), as obtained from FLEXPART

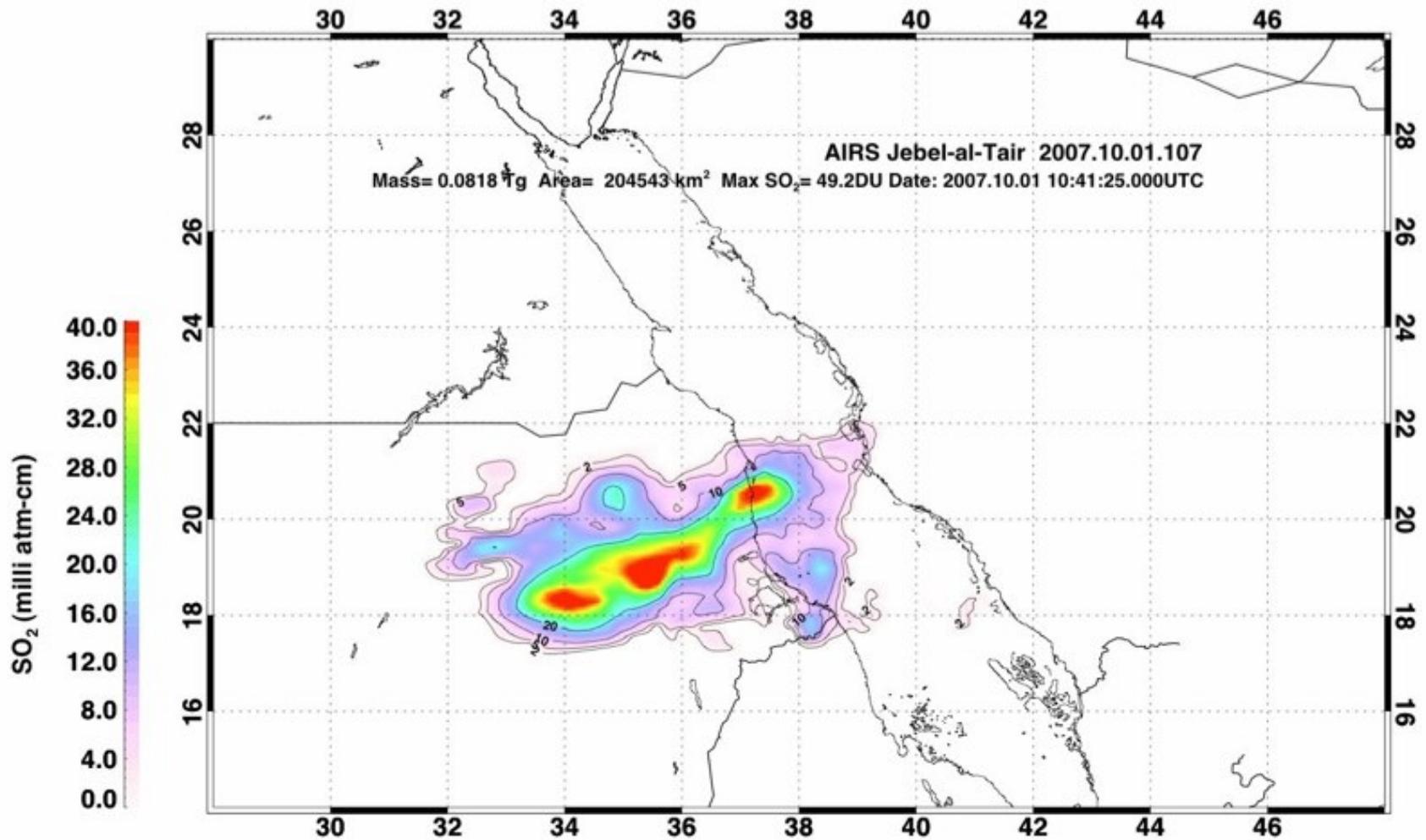
σ standard error of observation

$$M(x - x^a) \approx y^o - Mx^a$$

$$M\tilde{x} \approx \tilde{y}.$$

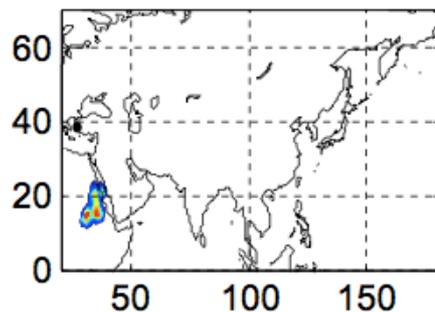
$$J = \underbrace{(M\tilde{x} - \tilde{y})^T \text{diag}(\sigma_o^{-2})(M\tilde{x} - \tilde{y})}_{\text{I) misfit model - observation}} + \underbrace{\tilde{x}^T \text{diag}(\sigma_x^{-2}) \tilde{x}}_{\text{II) deviation from first guess}} + \underbrace{\epsilon (D\tilde{x})^T D\tilde{x}}_{\text{III) smoothness condition}}$$

Satellite Data–AIRS

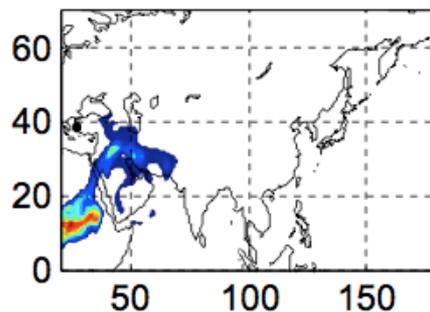


Model-FLEXPART

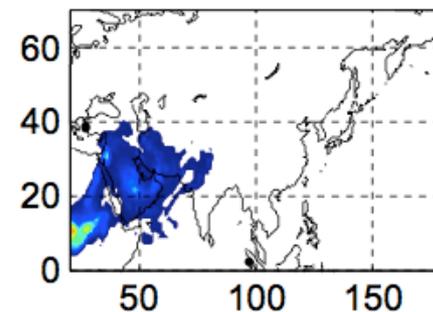
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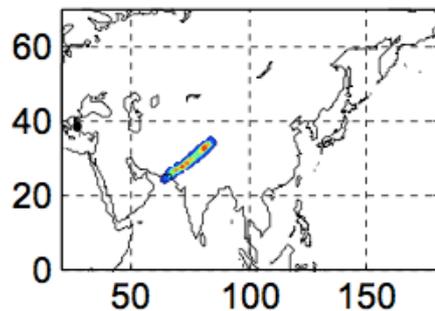
10.06. 08:00 height: 4000



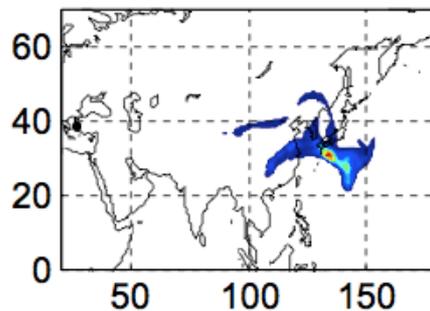
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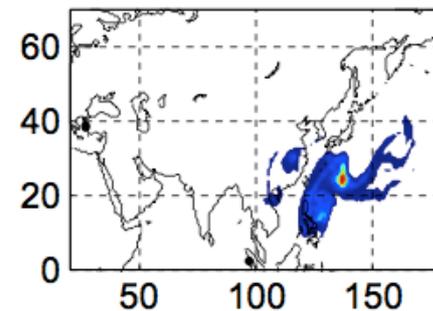
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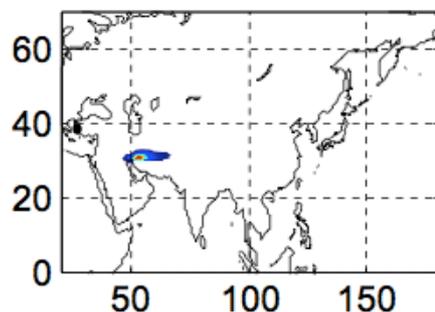
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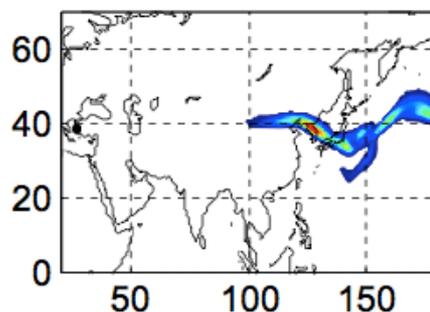
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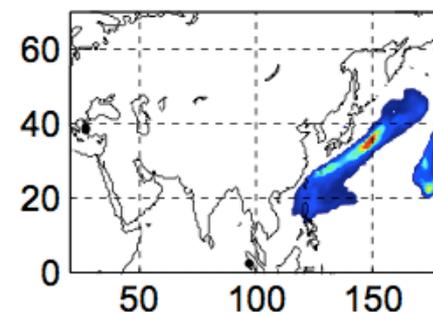
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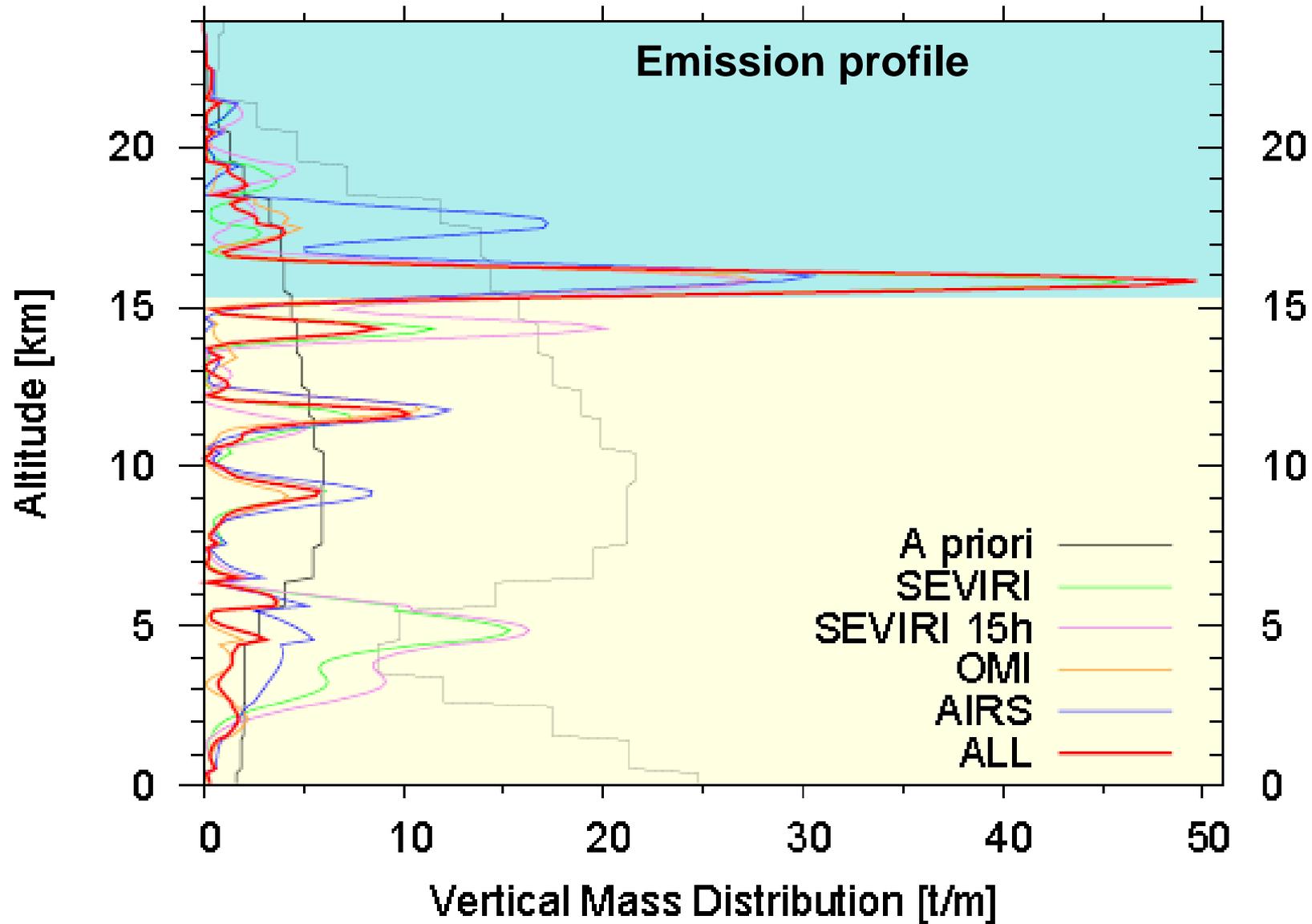


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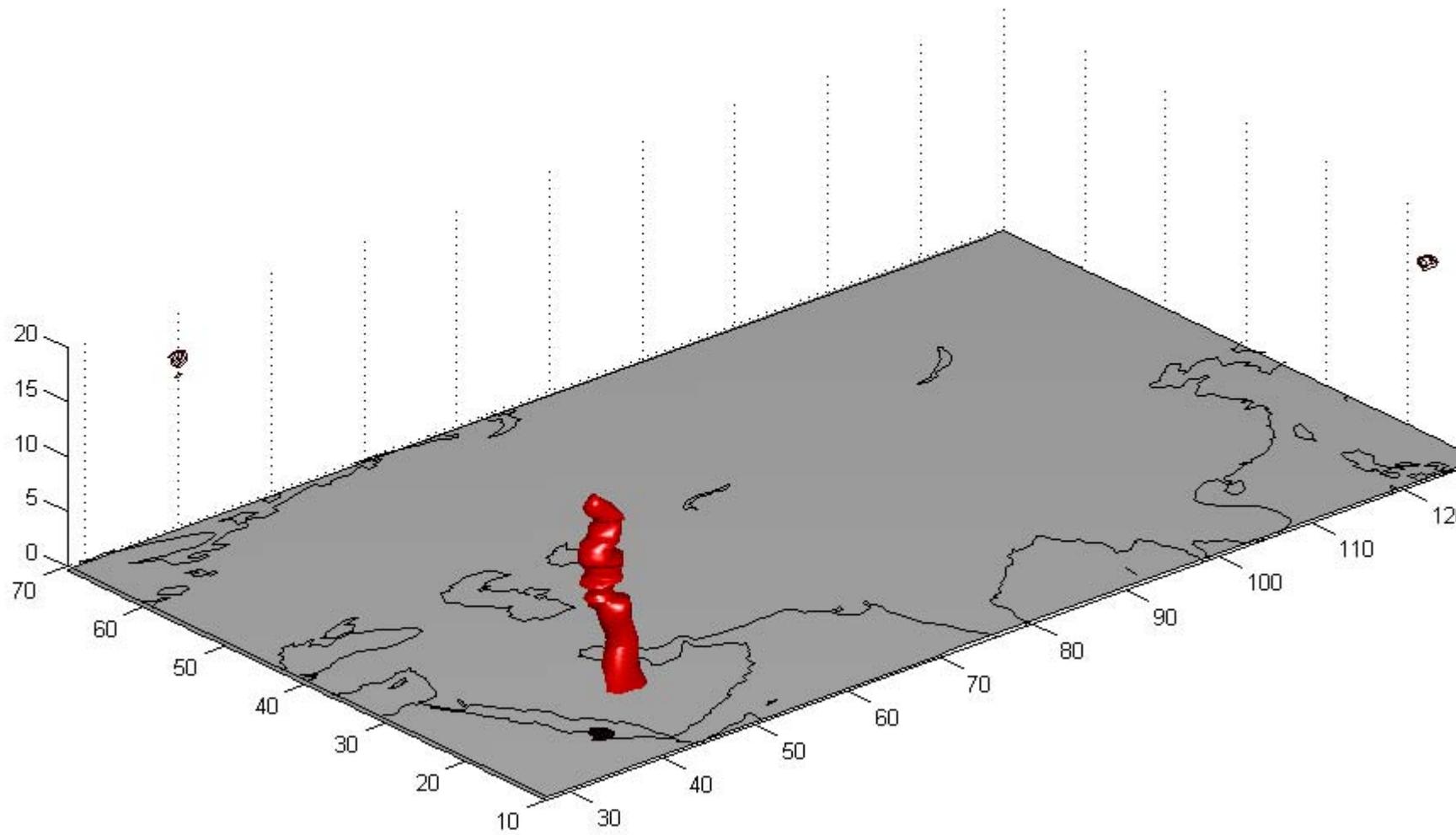


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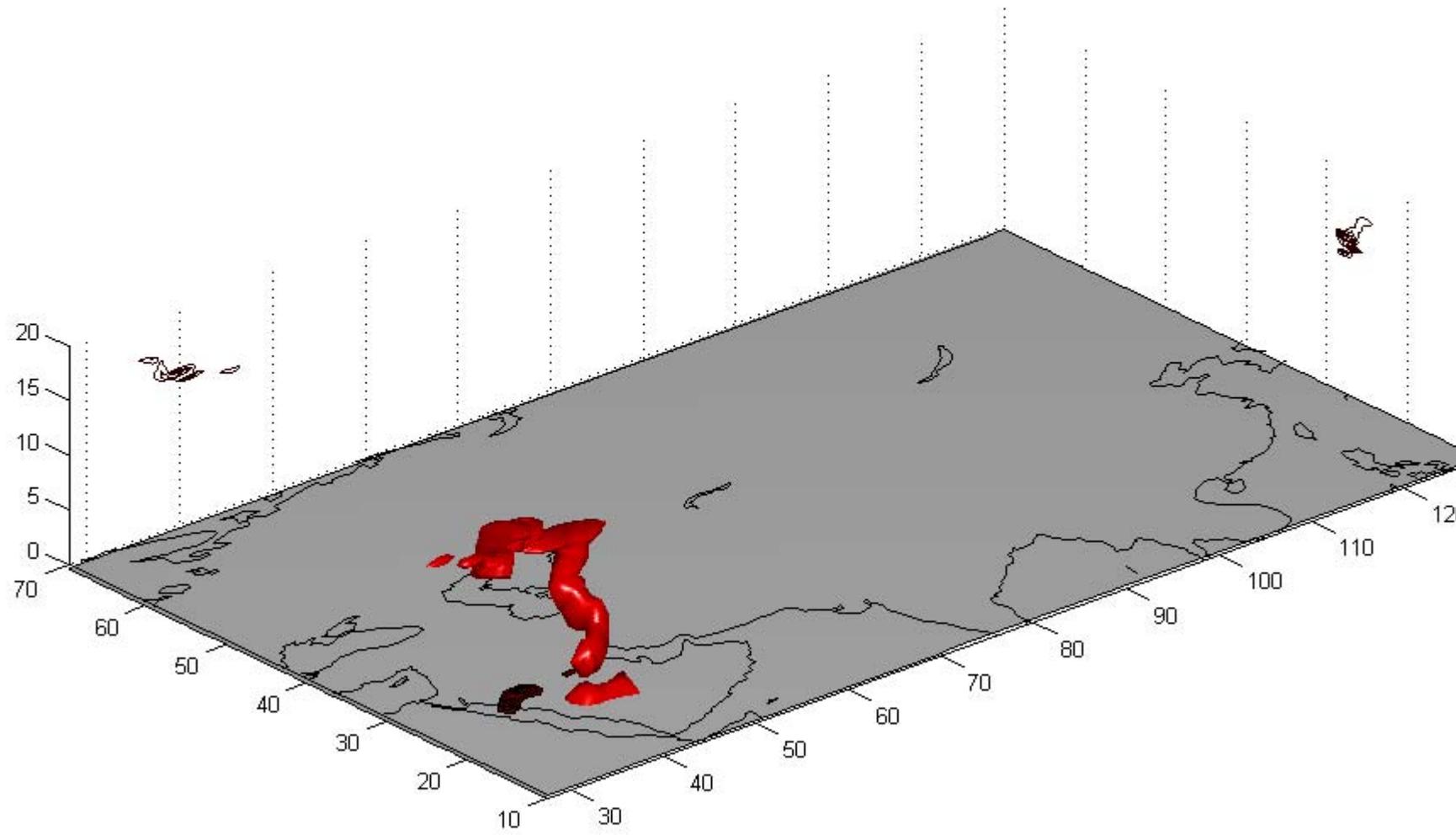




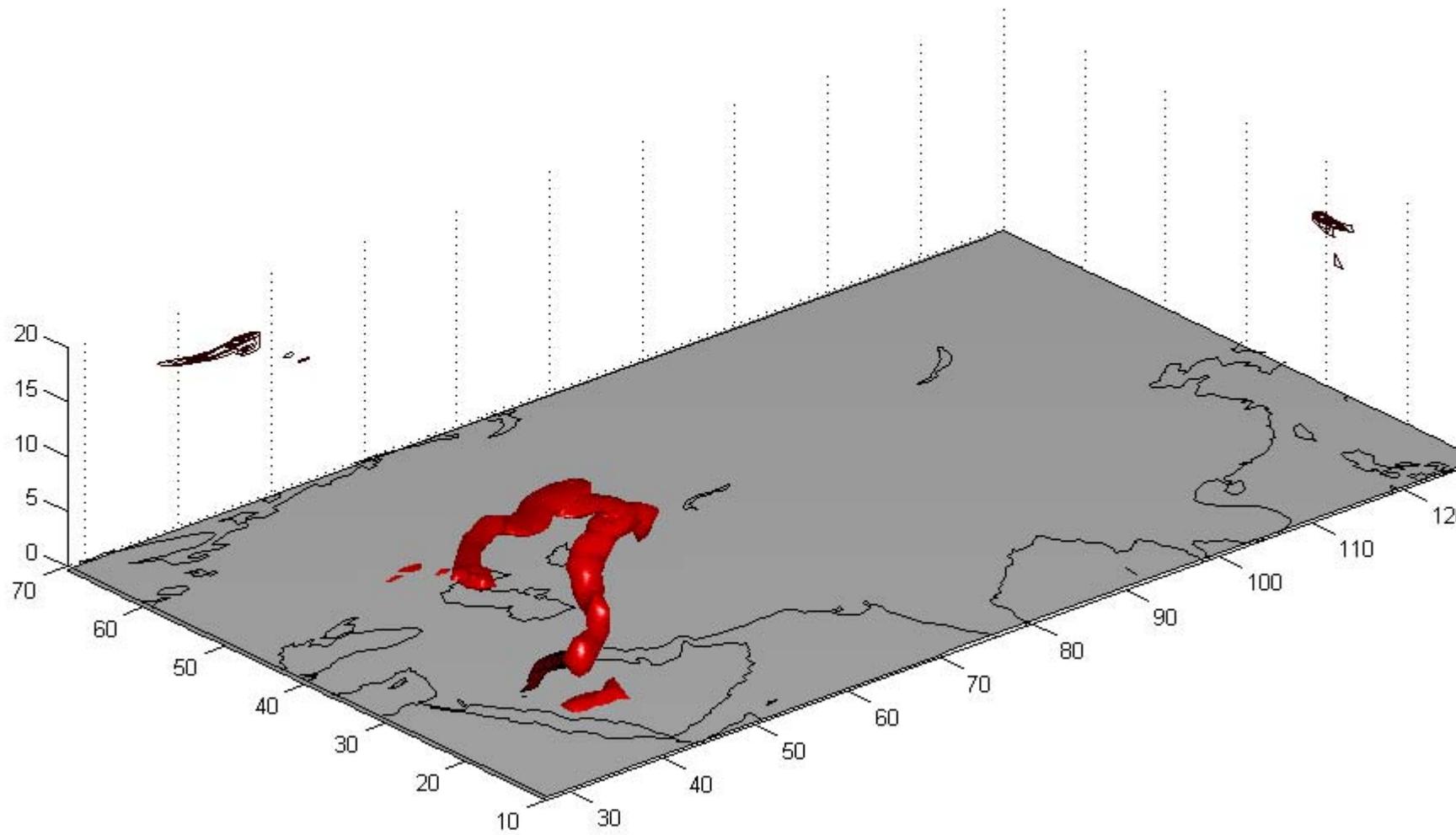
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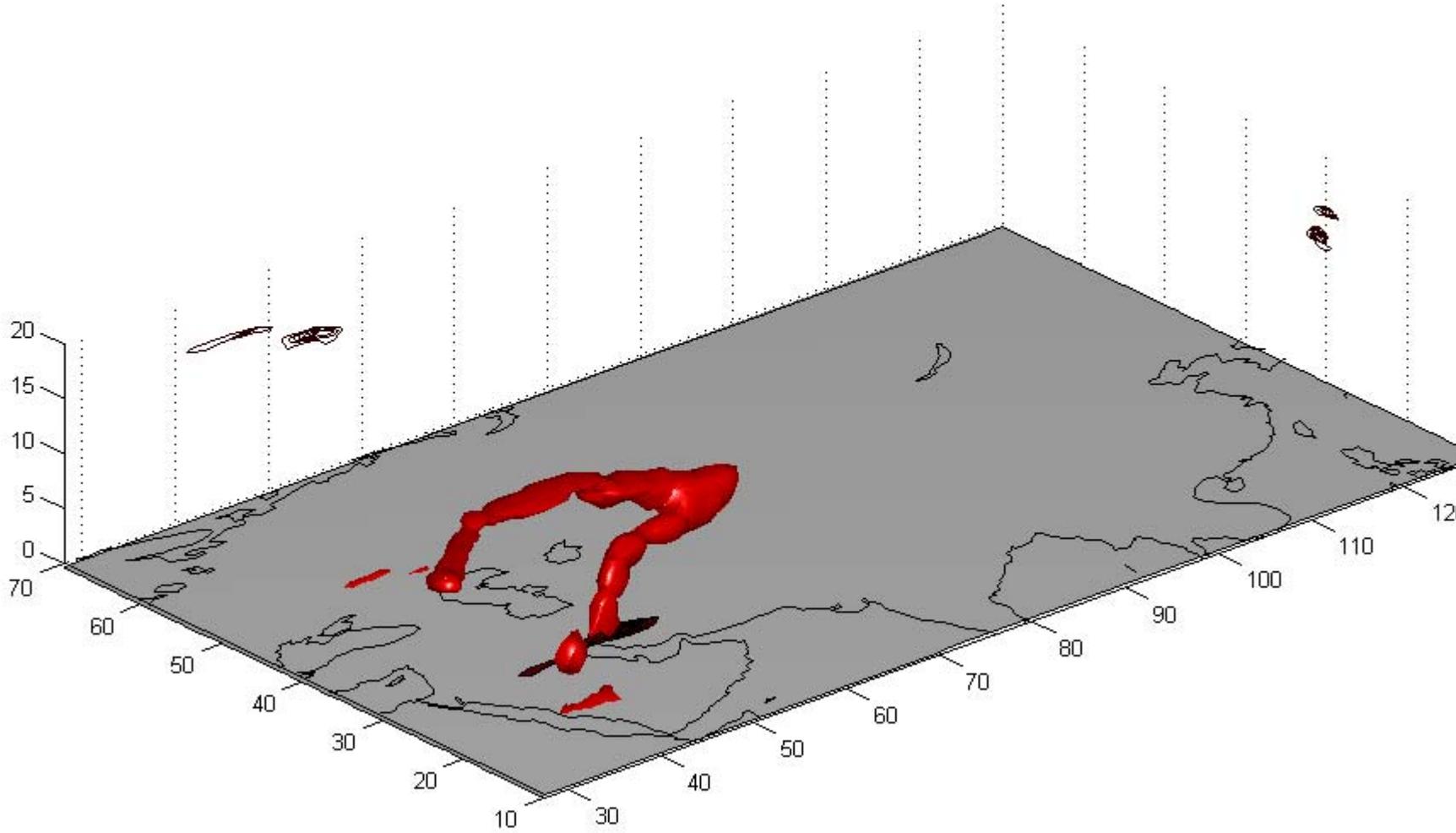
10.02. 04:00



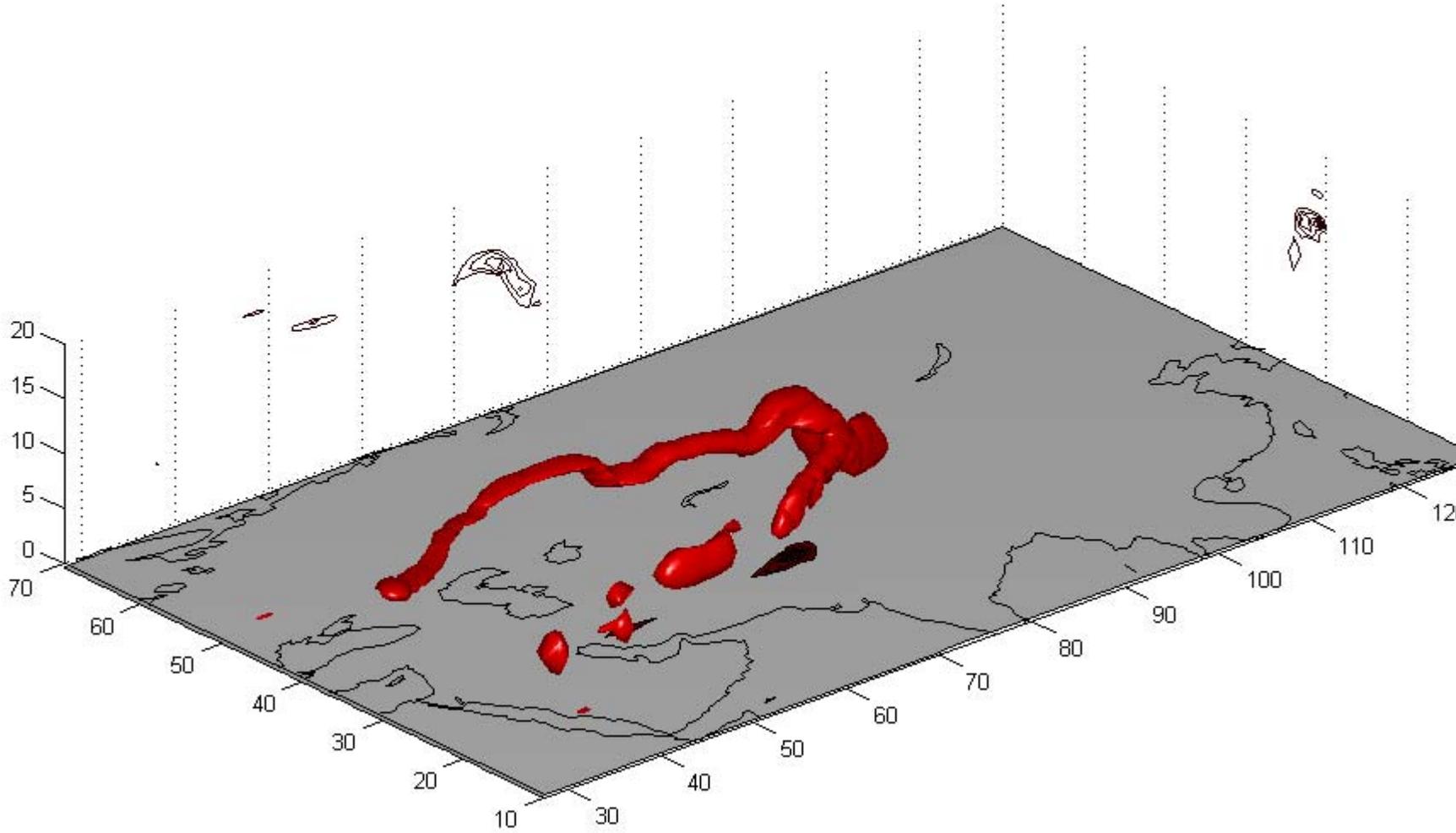
10.02. 12:00



10.02. 20:00



10.03. 12:00



10.04. 04:00

