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The vertical distribution of volcanic SO₂ plumes measured by IASI

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Discussion Paper

Abstract

Sulphur dioxide (SO_2) is an important atmospheric constituent that plays a crucial role in many atmospheric processes. Volcanic eruptions are a significant source of atmospheric SO₂ and its effects and lifetime depend on the SO₂ injection altitude. The Infrared Atmospheric Sounding Interferometer (IASI) on the METOP satellite can be used to 5 study volcanic emission of SO₂ using high-spectral resolution measurements from 1000 to 1200 cm⁻¹ and from 1300 to 1410 cm⁻¹ (the 7.3 and 8.7 μ m SO₂ bands) returning both SO₂ amount and altitude data. The scheme described in Carboni et al. (2012) has been applied to measure volcanic SO₂ amount and altitude for fourteen explosive eruptions from 2008 to 2012. The work includes a comparison with independent measurements: (i) the SO₂ column 10 amounts from the 2010 Eyjafjallajökull plumes have been compared with Brewer ground measurements over Europe; (ii) the SO₂ plumes heights, for the 2010 Eyjafjallajökull and 2011 Grimsvötn eruptions, have been compared with CALIPSO backscatter profiles. The results of the comparisons show that IASI SO₂ measurements are not affected by underlying cloud and are consistent (within the retrieved errors) with the other measurements. The 15

series of analysed eruptions (2008 to 2012) show that the biggest emitter of volcanic SO₂ was Nabro, followed by Kasatochi and Grímsvötn. Our observations also show a tendency for volcanic SO₂ to reach the level of the tropopause during many of the moderately explosive eruptions observed. For the eruptions observed, this tendency was independent of the maximum amount of SO₂ (e.g. 0.2 Tg for Dalafilla compared with 1.6 Tg for Nabro) and of the volcanic explosive index (between 3 and 5).

1 Introduction

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Sulphur dioxide (SO_2) is an important atmospheric constituent, important in many atmospheric processes (Stevenson et al., 2003; Seinfeld and Pandis, 1998; Schmidt et al., 2012). Volcanic eruptions are a significant source of atmospheric SO_2 , with its effects and lifetime depending on the SO_2 injection altitude. In the troposphere these include acidifi-

cation of rainfall, modification of cloud formation and impacts on air quality and vegetation (Ebmeier et al., 2014; Delmelle et al., 2002; Delmelle, 2003; Calabrese et al., 2011). In the stratosphere SO₂ oxidises to form a stratospheric H₂SO₄ aerosol that can affect climate for several years (Robock, 2000). Volcanoes contribute about one third of the tropospheric sulphur burden $(14 \pm 6 \text{ Tg S yr}^{-1}; \text{ Graf et al., 1997}; \text{ Textor et al., 2003})$. The annual amount 5 of volcanic SO₂ emitted is both poorly constrained and highly variable. The uncertainty in released SO₂ arises from the stochastic nature of volcanic processes, very little or no surface monitoring of many volcanoes, and from uncertainties in the contribution of volcanic sulphur emitted by guiescent (non-explosive) degassing. The effects of SO₂ in the atmosphere depend not only on the amount released but also the altitude of the plume. Altitude 10 information is important for atmospheric chemistry, as SO₂ reactions and depletion times change with height and atmospheric composition, particularly as a function of water vapour concentration (Kroll et al., 2015; von Glasow, 2010; McGonigle et al., 2004; Mather et al.,

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2003).

Most volcanic eruptions are accompanied by the release of SO₂ to the atmosphere, and data both on the quantity emitted, and the height at which it is injected into the atmosphere are valuable indicators of the nature of the eruption.

It is important to monitor volcanic SO₂ plumes for air safety (Brenot et al., 2014; Schmidt et al., 2014) as sulphidation of nickel alloys cause rapid degeneration of an aircraft engine (Encinas-Oropesa et al., 2008). SO₂ is also used as a proxy for the presence of volcanic 20 ash, although the co-location of SO_2 and ash depends upon the eruption so this approach is not always reliable for hazard avoidance (Sears et al., 2013). Mitigation strategies to avoid volcanic SO₂ and ash are currently based on ground monitoring and satellite measurements to assess the location and altitude of the proximal volcanic plume, followed by the use of dispersion models to forecast the future position and concentration of the plume 25 (Tupper and Wunderman, 2009: Bonadonna et al., 2012; Flemming and Inness, 2013). The outputs of the models are strongly dependent on the assumed initial plume altitude mainly because of the variability of the wind fields with altitude. The dispersion models may use an ensemble of possible initial altitudes to identify all places where the plume can potentially arrive. Hence improving the accuracy of the initial plume altitude reduces the uncertainty in the location of the transported volcanic SO₂.

Limb viewing satellite instruments, such as the Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) and the Microwave Limb Sounder (MLS), can estimate the vertical profile of SO₂ (Höpfner et al., 2013; Pumphrey et al., 2014) in the upper troposphere and lower stratosphere (UTLS), but generally cannot observe in the lower troposphere. Nadir measurements, by both spectrometers and radiometers, have been used to retrieve SO₂ column amounts, assuming the altitude of the plume, e.g., the Total Ozone Monitoring Spectrometer (TOMS, Carn et al., 2003), the Moderate Resolution Imaging Spectroradiometer (MODIS, Watson et al., 2004; Corradini et al., 2009), the Ozone Monitoring Instrument (OMI, Krotkov et al., 2006; Yang et al., 2009; Carn et al., 2009), the Atmospheric Infrared Sounder (AIRS, Prata and Bernardo, 2007), the Advanced Spaceborne Thermal

Emission and Reflection Radiometer (ASTER, Pugnaghi et al., 2006; Campion et al., 2010), the Global Ozone Monitoring Experiment 2 (GOME-2, Rix et al., 2012), and the Spinning 15 Enhanced Visible and Infrared Imager (SEVIRI, Prata and Kerkmann, 2007).

Nadir spectrometer measurements in the ultraviolet (OMI, GOME-2) and infrared (AIRS, the Tropospheric Emission Sounder, TES, IASI) can be used to infer information on the SO₂ plume altitude. This is based on the fact that the top-of-atmosphere radiance of a plume, with the same amount of SO₂ but at another altitude, will have a different spectral shape within the SO₂ absorption bands.

UV spectra have been used to retrieve information on SO_2 altitude by Nowlan et al. (2011) using an optimal estimation retrieval applied to GOME-2 measurements. Yang et al. (2010) used a direct fitting algorithm to retrieve the SO_2 amount and altitude from OMI data. Both of these techniques were applied to the 2008 Kasatochi eruption which injected a relatively

²⁵ large amount of SO₂ high (circa 12 km) in to the atmosphere. A new scheme to retrieve altitude from GOME-2 has been developed by Van Gent et al. (2015) and applied to Icelandic eruption case studies (Koukouli et al., 2014).

Volcanic SO₂ retrievals from satellite data in the thermal infra-red (TIR) part of the spectrum are based on two regions of SO₂ absorption around 7.3 and $8.7 \,\mu$ m. The strongest

 SO_2 band is at 7.3 µm. This is in a strong water vapour (H₂O) absorption band and is not very sensitive to emission from the surface and lower atmosphere. Above the lower atmosphere this band contains information on the vertical profile of SO_2 . Fortunately differences between the H₂O and SO₂ emission spectra allow the signals from the two gases

- to be decoupled in high resolution measurements. The $8.7 \,\mu\text{m}$ absorption feature is in an atmospheric window so it contains information on SO₂ from throughout the column. In Clerbaux et al. (2008) the SO₂ absorbing feature around $8.7 \,\mu\text{m}$ was used to retrieve the total SO₂ amount and profile from TES data by exploiting this instrument's ability to resolve the change in SO₂ line-width with pressure. The 7.3 μm spectral range has been used by
- ¹⁰ Clarisse et al. (2014) in a fast retrieval of SO₂ altitude and amount. Their approach, based on Walker et al. (2012), assumes that the SO₂ is located at different altitudes, choosing the altitude that gives the best spectral fit.

In this paper, the Carboni et al. (2012) retrieval scheme (hereafter C12) has been applied to study the vertical distribution of SO_2 for the major eruptions during the period 2008–2012 and for some minor low tropospheric eruptions such as an Etna lava fountain.

The aims of this paper are to test the retreived SO_2 amount and altitude against other datasets (we used CALIPSO backscattering profile to test the altitudes and Brewer ground measurements for the column amounts) and to study the volcanic plumes from different eruption types and in different locations.

20 2 IASI data

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There are currently IASI sensors on-board the European weather satellites METOP-A and METOP-B. METOP-A has been operating since October 2006 and was the first of three satellites scheduled to operate for a total of 14 years. METOP-B has been operating since September 2012. The satellites cross the Equator on the descending node at a local time of around 9:30 a.m.

IASI is a Fourier transform spectrometer covering the spectral range $645-2760 \text{ cm}^{-1}$ (3.62 to $15.5 \,\mu\text{m}$) with spectral sampling of $0.25 \,\text{cm}^{-1}$ and an apodized spectral resolution

of $0.5 \,\mathrm{cm}^{-1}$ (Blumstein et al., 2004). It has a nominal radiometric noise of 0.1-0.3K in the SO₂ spectral range considered, according to Hilton et al. (2012).

The field-of-view (FOV) consists of four circular footprints of 12 km diameter (at nadir) inside a square of $50 \,\mathrm{km} \times 50 \,\mathrm{km}$, step-scanned across track (with 30 steps). The swath is

- 2100 km wide and the instrument can nominally achieve global coverage in twelve hours. 5 Observations are co-located with the Advanced Very High Resolution Radiometer (AVHRR), providing complementary visible/near infrared measurements. IASI carries out nadir observation of the Earth simultaneously with the Global Ozone Monitoring Experiment (GOME-2) also on-board METOP. GOME-2 is a UV spectrometer measuring SO₂ in
- the UV absorption band and is used for both Differential Optical Absorption Spectroscopy 10 (DOAS) (Rix et al., 2012) and optimal estimation retrievals (Nowlan et al., 2011) of SO₂. More information about IASI can be found in Clerbaux et al. (2009). The IASI level 1C data (geolocated with apodized spectra) used here were obtained from both the British Atmospheric Data Centre (BADC) archive and EUMETSAT Unified Meteorological Archive

Facility (UMARF) archive. 15

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3 SO₂ retrieval scheme

The retrieval scheme follows C12 (Carboni et al., 2012) where the SO₂ concentration is parametrised as a Gaussian profile in pressure (with 100 mb spread). Using IASI measurements from 1000 to 1200 cm⁻¹ and from 1300 to 1410 cm⁻¹, an optimal estimation retrieval (Rodgers, 2000) is employed to estimate the SO₂ column amount, the height and spread of the SO₂ profile, and the surface skin temperature.

The forward model, based on Radiative Transfer model for TOVS (RTTOV) (Saunders et al., 1999) but extended to include SO₂ explicitly, uses ECMWF temperatures interpolated to the measurement time and location. The retrieval technique uses an error covariance matrix, based on a climatology of differences between the IASI measurements and SO₂-25 free forward modelled spectra. Any differences not related to SO₂ between IASI spectra and those simulated by a forward model are included in the covariance matrix. Note that

the SO_2 retrieval is not affected by underlying cloud. If the SO_2 is within or below an ash or cloud layer its signal will be masked and the retrieval will underestimate the SO_2 amount, in the case of ash this is indicated by a cost function value greater than two.

- The retrieval is performed for every pixel where the SO₂ detection result is positive (Walker et al., 2011, 2012). The scheme iteratively fits the forward model (simulations) with the measurements, to seek a minimum of a cost function. The solution, when the measurements do not contain enough information to retrieve all the parameters in the state vector, is strongly affected by the assumed a priori values. When the SO₂ amount decreases, the spectral information decreases, and it is not possible to retrieve both SO₂ amount and altitude. This often happens at the edge of the plume. In this work we use 400 ± 500 mb as
- the a priori value for plume altitude and $0.5 \pm 100 \text{ DU}$ for column amount. In addition, only quality controlled pixels are considered; these are values where the minimisation routine converges within 10 iterations, the SO₂ amount is positive, the plume pressure is below 1000 mb and the cost function is less then 10.
- ¹⁵ Rigorous error propagation, including the incorporation of forward model and forward model parameter error, is built into the system, providing quality control and error estimates on the retrieved state. Retrieved uncertainties increase with decreasing altitude, nevertheless it is possible to retrieve information in the lower troposphere and monitor volcanic degassing. In the case of two or multiple SO₂ layers the forward model assumption of a sin-
- ²⁰ gle Gaussian layer is a source of error (and this error is not included in the pixel by pixel error estimate). In this case the retrieved altitude is an effective altitude between the two (or more) plume layers; in particular the altitude will be the one that is radiatively closer to the measured spectra and the altitude will be the radiatively equivalent altitude.

The altitude of the SO₂ plume strongly modulates the retrieval error as the contrast between plume temperature and surface temperature is a critical factor. The error in SO₂ amount decreases with an increase in plume altitude. Typical uncertainties are 2 DU for a plume centred at 1.5 km and less then 1 DU for plumes above 3 km.

When using the result from this setting of the retrieval scheme the following caveats should be considered:

- The retrieval is valid for SO₂ column amounts less then 100 DU (limit used in the computation of RTTOV coefficients) and altitudes less than 20 km (due to the used Gaussian profile with 100 mb spread). For higher eruptions one should use a thinner Gaussian profile, for column amounts bigger then 100 DU new RTTOV coefficients will be needed. At present both of these conditions produce results that do not pass the quality control.
- 2. Quality control and orbit gaps can produce artefacts (see Sect. 6).

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- 3. Care should be taken when using the altitude to infer if a plume is above or below the tropopause. It is therefore desirable to combine or confirm these measurements using other measurements. This is because there are altitudes with the same temperature above and below the tropopause, and the SO₂ signal could be similar. Nevertheless, the IASI spectrum contains altitude information from the overlapping of SO₂ with other gas absorption.
- 4. The global annual mean variability between real atmospheric profiles and ECMWF profiles is included in the error covariance matrix, and then propagated into the retrieval errors. However we cannot exclude some extreme events where the difference between ECMWF profiles and the real profile is at the tail of the annual global statistical distribution. In this case, local and seasonal covariance matrices could be used. For the retrievals presented here a global covariance matrix has been used, computed using days from all four seasons.
 - 5. Volcanoes also emit water vapour. If this emission is larger than the water vapour variability considered in the error covariance matrix, the results could be affected by errors that are not included in the output errors.

Comparisons with the Universite libre de Bruxelles (ULB) IASI SO₂ dataset, as well as ²⁵ UV-Vis instruments such as GOME2/MetOp-A and OMI/Aura, have been performed and are cited in the relevant eruptions sections. We simply note here that for the Grímsvötn eruption our data were compared to the ULB IASI/MetOp-A SO₂ and the Belgian Institute for Space Aeronomy (BIRA-IASB) GOME2/MetOp-A SO₂ retrievals (Koukouli et al., 2014). The Etna continuous outflow measurement were compared with Istituto Nazionale di Geofisica e Vulcanologia (INGV) MODIS/Terra, Rutherford Appleton Laboratory (RAL) MODIS/Terra, ULB IASI/METOP-A, German Aerospace Center (DLR) GOME2/MetOpA and ground-based Flame network measurements (Spinetti et al., 2015) and finally the data from the Eyjafjallajökull eruptions, with the DLR GOME2/MetOpA, the BIRA OMI/Aura and the AIRS data (Carboni et al., 2012; Koukouli et al., 2014). More recently data for the Bárdarbunga eruption, were compared with the BIRA-IASB OMI/Aura data (Schmidt et al., 2015).

4 Altitude comparison with CALIOP

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Vertical profiles of aerosol and clouds are provided by NASA's Cloud Aerosol Lidar with Orthogonal Polarization instrument (CALIOP, Winker et al., 2009) carried on the CALIPSO satellite. CALIOP's vertical/horizontal resolution is 0.06/1.0 km at altitudes between 8.2 and 20.2 km. CALIPSO is part of the A-train with an equatorial overpass time of ~ 13.30 .

¹⁵ and 20.2 km. CALIPSO is part of the A-train with an equatorial overpass time of ~ 13.30. CALIOP and IASI have coincident measurements around $\pm 70^{\circ}$ latitude. Moving from this latitude towards the equator causes the time difference between the two measurements to increase.

CALIOP backscatter profiles have been averaged (to 3 km along-track, 250 m vertical) and CALIOP observations of volcanic plumes have been identified using SEVIRI false colour images based on the infrared channel at 8.7, 11 and 12 μ m (Thomas and Siddans, 2015). CALIOP is sensitive to aerosol and water droplets that scatter sunlight; in the case of volcanic eruptions these aerosols are H₂SO₄ and ash. SEVIRI is sensitive to both ash and SO₂ since its channel around 8.7 μ m is within the SO₂ absorption band.

The criteria used to define a coincidence between CALIOP and IASI pixels were: a distance of < 100 km and a time difference of < 2 h. With these relatively strict criteria, the selected coincident data are only from the Icelandic Eyjafallajökull and Grimsvötn eruptions.

Figure 1 shows the comparison between CALIOP and IASI for the Eyjafjallajökull plume on the 7 May 2010. This is the CALIOP track presented by Thomas and Prata (2011) in which they identify both ash and SO₂ plumes in the southern part of the track (less then 55° N) and SO₂ only plumes in the northern part of the track (the scattering feature around 5 km between 55–60° N where the scattering signal is possibly from H₂SO₄ aerosol result ing from the oxidation of volcanic SO₂).

Ash and SO₂ are not necessarily advected together (Thomas and Prata, 2011; Sears et al., 2013), but in these case studies (Figs. 1-4), there is good agreement between IASI and the backscattering features within the IASI error bars. This may be due to the presence of H_2SO_4 associated with the SO₂ plume. An important aspect to note is that in nearly all of the coincidences for the Eyjafallajökull case (with the exception of a few pixels on 15 14 May) the volcanic plume is above a lower meteorological cloud. This is identifiable as the scattering layer at around 1 km height in Fig. 1 and between 1 and 4 km in Figs. 2 and 3. This comparison against CALIOP in the presence of water cloud confirms that the retrieval is not affected by underlying cloud. This is because the variability that such cloud introduces is represented by the error covariance matrix which includes radiance differences between 20 the clear forward model and cloudy IASI spectra. An accurate retrieval of altitude is also an important factor for the estimate of SO_2 amount, this is because the thermal SO_2 signal can be up to one order of magnitude different between different altitudes. A good comparison of altitude with CALIPSO then gives confidence in both altitude and column amount.

Figure 4 shows the Grimsvötn plume at the beginning of the eruption. There is separation between the higher part of the plume (dominated by SO₂) moving higher and spreading north, and the lower part of the plume (dominated by ash, but still with some SO₂ associated with it) moving south-west. It was this part of the plume that moved over Europe in the following days and caused airspace closure.

5 Comparison with Brewer ground data

During the Eyjafjallajökull eruption in April and May 2010, the volcanic plume overpassed several European ground measurement stations equipped with Brewer instruments. Brewer instruments are UV spectrophotometers, principally dedicated to measuring the total ozone column but also capable of determining total column SO₂ (Kerr et al., 1981). Extraction of 5 the SO₂ signal from the UV measurements is performed as a second step after the ozone quantity retrieval due to the much lower SO₂ absorption feature strength (De Muer and De Backer, 1992). Depending on the amount of atmospheric SO₂ affecting the instruments, the Brewer spectrophotometer has been shown to be sensitive both to anthropogenic SO_2 loading in the lower troposphere (Zerefos et al., 2000) as well as to an overpass of a vol-10 canic plume, which produces a strong SO₂ signal within the ozone absorption bands. These ground-based measurements have lately been used in Rix et al. (2012) in a comparison with the GOME-2/METOP-A SO₂ retrievals and in the validation work of the European Space Agency, ESA, projects (SMASH and SACS2¹) on the 2010 and 2011 Icelandic eruptions (Koukouli et al., 2014). The World Ozone and Ultraviolet Radiation Data Centre (WOUDC) 15

is an active archive facility that includes quality-assured Brewer ground-based O_3 measurements and also provides SO_2 daily averages. The dataset for April and May 2010 was downloaded from the WOUDC archive (http://www.woudc.org/) for all European sites with Brewer instruments.

The reported SO₂ amount in non-volcanic conditions (e.g. over Europe before 15 April 2010) from Brewer stations varies considerably which can mostly be attributed to the small signal-to-noise ratio of the SO₂ signal compared to the ozone signal. This variability may also be due to insufficient reported data quality control with respect to SO₂ retrieval, since most stations focus on the quality of the ozone measurements and not their by-products.

²⁵ Negative values can also be present in the daily average datasets, a result of the nominal Brewer algorithm's inability to resolve small atmospheric SO₂ amounts. Therefore, only

¹"Study on an End-to-End system for volcanic ash plume monitoring and prediction" (SMASH) and "Support to aviation control service 2" (SMASH2).

a subset of the available Brewer sites in the WOUDC database were selected for this study. Stations were not selected if they reported a majority of negative values (such as Reading, UK; La Coruna, Spain, and so on) for the two months considered (April and May 2010), or had negative values less than -1 DU (such as Madrid, Spain; Poprad-Ganovce, Slovakia). These negative values point to the small amount of the volcanic gas reaching the specific locations. Despite recording several negative values, the Valencia, Spain, site was considered since it was presented in Rix et al. (2012), after checking that the measurements coincident with the presence of the IASI plume were statistically significantly larger than the average background measurements.

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- All the positive values of the selected ground stations (listed in Fig. 5) have been com-10 pared with the IASI measurements of the SO₂ plume. The SO₂ estimates available from WOUDC are daily averages. In order to compare these datasets with the IASI observations, all the satellite pixels of the morning and evening orbits within a 200 km radius from the Brewer site have been averaged. The choice of radius was made since at an average wind speed of 6 m s^{-1} an air parcel will travel 250 km in 12 h, so with this spatial criteria 15 we are including in the satellite averaging all pixels that might be overpassing the Brewer location within a temporal frame of 12h (daytime). At the same time, the averaged IASI error and standard deviation have been computed. The results are presented in the scatter plot in Fig. 5. The linear fit is computed considering the IASI average error and a fixed error of 0.5 DU for the Brewer measurements. Note that the variability of the IASI SO₂ amount 20 within 200 km is often much bigger than the IASI error bar. A correlation coefficient of 0.76 with root mean square differences of 1.16 DU has been found. This result is encouraging for the IASI retrieval, even if this initial comparison alone cannot be considered a comprehen-
- sive validation exercise because (1) it is difficult to assess the quality of Brewer SO₂ daily
 ²⁵ average values; (2) the comparison is restricted to the Eyjafjallajökull eruption, an eruption where the SO₂ plume covered a small range of altitudes (between 2 and 5 km) and a relative small loading amount.

From Fig. 5 it can be noted that for loadings up to and around 2 DU both types of observation appear to depict the same atmospheric SO₂ loading, which, depending on the

location of the site, might be both of anthropogenic and volcanic provenance. For values between 2 and 3 DU there appears to be a slight underestimation by IASI of around 0.5 DU, well within the statistical uncertainty. For higher loadings still, the Brewer instruments report higher SO₂ values than the satellite.

- ⁵ The overestimation of low amounts can be explained by the a priori SO₂ amount, which is 0.5 DU. Operationally, if there is insufficient information from the measurements (as is the case when there is less than 0.5 DU), the output of the optimal estimation retrieval tends to the a priori value. One factor that can explain the underestimation (by IASI of SO₂ > 3 DU) is that this IASI retrieval is sensitive to SO₂ values higher than the climatological SO₂ amount
- considered in the IASI data ensemble used to compute the error covariance matrix. In the case of the Eyjafallajökull eruption the north Atlantic and European region of April and May 2009 were used to compute the error covariance matrix, so the retrieval could be insensitive (biased) to the average values of SO₂ amount in that region, which includes the European background value of SO₂.

15 6 Total mass and vertical distribution

The SO₂ retrieval algorithm C12 has been applied to several volcanic eruptions in the period 2008–2012. For each eruption the IASI orbits are grouped into twelve hour intervals in order to have two maps, each day, of IASI retrieved SO₂ amount and altitude. IASI pixels of overlapping orbits are averaged together. These maps are gridded into 0.125° lat/lon boxes.

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- An example of maps for the Nabro eruption is shown in Fig. 6. Note that with only METOP-A the IASI data have a gap between the orbits at tropical latitudes, this gap is filled with the METOP-B launch in September 2012. It is also possible to have gaps in SO₂ coverage due to pixels that did not pass quality control. The regridding routine fills gaps by a triangular interpolation of neighbouring pixels.
- ²⁵ Figure 6 shows an example of how the regridding routine fills in missing data. However, because of the possible creation of artefacts, the regridding should be used carefully. For example, in the case of a plume covering the edge of one orbit but with no plume present

in the adjacent orbit, regridding can fill up the gaps between the orbits with a bigger plume than would be reasonable to expect. For the case studies presented here, all the regridded IASI maps have been inspected "by eye" to check that no particularly significant artefacts have been introduced.

- The total SO_2 mass present in the atmosphere is obtained by summing all the values of the regularly gridded map of SO_2 amounts. In particular every grid-box column amount is multiplied by the grid-box area to obtain the SO_2 mass, and all the grid-box masses are summed together to obtain the total mass of SO_2 for each IASI image. The total mass errors are obtained in the same way from the grid box errors, i.e. all the box errors are summed
- to produce the total mass error. This is an overestimation of the error, but considering the mean squared error as total error will be an underestimation. This is due to the presence of systematic errors within the retrieval. The systematic errors are included within the error estimate but cannot be considered independent of each other. It is more likely that, if present, the systematic error will become a bias in the region and time considered. The time series of these total masses, together with the errors, are presented in Fig. 7 for the studied volcanic eruptions.

Using the IASI dataset it is possible to follow the plume evolution of several volcanic eruptions. Within the eruptions considered Nabro produced a maximum load of 1.6 Tg of SO₂, followed by Kasatochi (0.9 Tg), Grímsvötn (0.75 Tg), Copahue (0.72 Tg) and Sarychev (0.60 Tg). Using the time series created by this dataset it might be possible to estimate the SO₂ lifetime but this is beyond the scope of this current work.

From the eruptions presented in Fig. 7, there is a wide spread of error-bars, depending on the SO_2 amount, altitude and atmospheric conditions. In general, plumes tend to spread and dissipate with time, covering a larger area (i.e. more IASI pixels) with smaller quantities. This produces smaller SO_2 signals and consequently bigger retrieval errors. This is the main cause of the general tendency of increasing error bars with time.

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It is also possible to estimate the SO₂ mass present between two altitude levels. Doing this every 0.5 km between 0 and 20 km gives a vertical distribution of SO₂ every \sim 12 h. These results are shown, in chronological order, in Figs. 8–11. Note that the colour-bars

for different volcanic eruptions have different values, going from smaller values for Etna and Llaima eruptions to a maximum for Nabro. From these figures, it is possible to observe the temporal evolution of the SO₂ plume as a function of altitude. These plots have to be interpreted carefully and studied together with the maps of the amount and altitude (values and errors) because here the retrieved errors in altitude are not accounted for, and for low amounts of SO₂, error in altitude can be significant. Within the IASI spectra there is enough information to retrieve altitude from small/medium eruptions such as Etna and Nyamuragira/Nyiragongo.

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ing ECMWF temperature and pressure profiles. The hydostatic equation is used to obtain height from these. The troppquse is defined as the lowest layer with a lapse rate below $2 \,\mathrm{K \, km^{-1}}$ and lapse rate of $2 \,\mathrm{K \, km^{-1}}$ in all layers within $2 \,\mathrm{km}$ above this. (The hydrostatic equation is also used to convert the retrieved SO₂ pressure into height, so any error in the pressure / height conversion will be common to both plume and tropopause heights.)

The tropopause heights have been computed for every pixel of the volcanic plume us-

- The three tropopause lines, shown in Figs. 8–11, are the mean, and the mean plus or 15 minus the standard deviations, within the IASI plume pixels in the 12 h maps. The reported values of the tropopause are computed using the location of the volcanic pixels only. Eyjafjallajökull and Puyehue eruptions cover the latitude range between 30-80° N and -20 -60° S respectively, so spanning a large range in tropopause heights. Day to day variation are sometimes large due to small amounts of SO₂ being detected or not from one day to the 20 next (coupled to the wide range of latitudes spanned by the plumes). Within the plume, the
- tropopause heights can differ by many kilometres especially for plumes that cover a wide latitude range. As an example, the Kasatochi plume has been analysed between 30 and 90° N. Over this range of latitudes the tropopause height varies between 8 and 18 km.
- Given this caveat the lines of tropopause mean and standard deviation heights are indica-25 tive, e.g. a plume that is below the three lines is likely, but not necessary, confined to the troposphere, a plume that is above the three lines is likely confined in the stratosphere, and SO_2 that is between the lines could be in either the troposphere or stratosphere.

In the following Sects. 5.1–5.5 we describe the eruptions shown in Figs. 8–11 grouped by geographic area. A summary of these comments together with other relevant literature and volcanic explosivity index (VEI) estimates from the Smithsonian is given in Table 1.

Three principal factors affect plume height: (i) the energetics of the eruption, (ii) the dy-⁵ namic effect, (iii) retrieval artefacts in the case of multilayer plumes. Here we attempt to discriminate between these factors as follows:

a. The energetics of the eruption. A crude parameter for eruption intensity and plume height is the "volcanic explosivity index" or VEI. The VEI is a semi-quantitative index of eruption size, which for contemporary eruptions can be used as a 'threshold' to determine the likelihood of stratospheric injection (Newhall and Self, 1982) Eruptions of VEI 4 and larger are expected to have strong plumes and be associated with significant stratospheric injections. Eruptions of VEI 3 are intermediate in size, with eruptive ash plumes that rise 5 – 15 km above the vent. Based on analysis of eruption statistics, 25 – 30% of VEI 3 eruptions may reach the stratosphere (Pyle et al., 1996). Eruptions of VEI 2 and smaller are not expected to reach the stratosphere. We report these, when available, in Table 1.

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- b. The atmospheric effect. In the following sections we group the eruptions by geographic area in order to consider together eruptions that may have similar conditions of water vapour. Moreover we consider the tropopause altitude together with the plume altitude.
- c. Retrieval artefact in the case of multilayer plumes. We do not have a way to identify this in a fresh plume but we indicate (with a black triangle) when the old plume overpasses near the volcano again (with the possibility of presence of both the old overpassing plume and the new emitted plume, at two different altitudes). The vertical distribution plots in Figs. 8–11 present the studied eruptions in chronological order and indicate the presence of a new plume connected with the volcano with red triangular symbols at the bottom of the column.

6.1 Southern Chile: Llaima, Puyuhue-Cordon-Caulle and Copahue

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In the Southern Hemisphere we analysed three eruptions that originated from volcanic activity in Chile. The wind direction was similar for each eruption so the SO₂ was transported to the east (towards South Africa).

⁵ Llaima (period analysed: 2–6 January 2008), is presented in Fig. 8. This is the smallest eruption in terms of SO₂ amount (with a maximum atmospheric load of 0.04 Tg) and after the first day, the plume was disconnected from the volcano.

Copahue (22–27 December 2012), in Fig. 11, released a higher amount of SO_2 (up to 0.72 Tg of atmospheric load) and there was a continuous plume from the volcano, indicating that the volcano emitted SO_2 for at least seven days. Both of these eruptions remained confined to the troposphere.

The Puyehue-Cordon-Caulle eruption of 5–30 June 2011 was the most significant eruption in southern Chile since 1991. The eruption commenced with the formation of a very significant eruption plume, that was tracked around the globe, and sustained, lower-level activity that continued intermittently during the observation period. This eruption was between Llaima and Copahue in terms of SO₂ amount (maximum load of 0.13 Tg), but higher than both in terms of VEI (with a value of 5), and with a much higher altitude (Fig. 11). The first part of the eruption produced a higher plume in terms of amount and altitude, and a lower plume connected with the volcano was present intermittently for all of the period observed. The maximum amount of SO₂ loading found was 0.13±0.06 Tg on 8 and 9 June. Theys et al. (2013) present the SO₂ amount from the Clarisse et al. (2012) IASI scheme, assuming an altitude of 13 km, and reported values around 0.14 Tg (and higher) for the periods 6–9 June, with a maximum on 7 June. The fresh part of the plume (over south America and the western Atlantic ocean) is found to be significantly lower using C12 (between 2 and

5 km of altitude) than the altitude of 13 km assumed by Theys et al. (2013), and the SO₂ amount for a plume at 5 km will be underestimated using the assumption of higher altitude. Due to the opposite effect, the presence of SO₂ higher than 13 km in the first two days may explain the discrepancy in the SO₂ load (Fig. 2 from Theys et al., 2013 reports higher SO₂

than C12 but assumes lower altitude than C12). This shows how important it is to have good altitude information for the SO_2 retrieval as assumptions regarding plume altitude may lead to significant differences in SO_2 total mass estimates.

6.2 Northern Pacific Ocean: Okmok, Kasatochi and Sarychev

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⁵ Okmok, Kasatochi and Sarychev eruptions (Figs. 8 and 9) all affected the boreal atmosphere and injected SO₂ to altitudes around the tropopause (and higher).

Okmok (12–20 July 2008) injected SO_2 around 12 km and the plume spread south and east from the volcano, over the Pacific Ocean. The main plume reached the US and Canada, and there was an intermittent presence of a small and low plume connected with the volcano.

Kasatochi (7–22 August 2008) injected the majority amount of SO₂ within the first three days and the plume spread around the Northern Hemisphere, going east, north and west from the volcano. The SO₂ amount reached a maximum seven days after the start of the eruption, indicating continuous injection from the volcano. Within the first ten days of the eruption it affected the latitude bands 30 to 90° N. The SO₂ amounts for the first days of the eruption are in the ranges reported from Corradini et al. (2010) for AIRS and MODIS estimates. (Corradini et al., 2010 report AIRS and MODIS 7.3 μm SO₂ mass loadings between 0.3 and 1.2 Tg, while the MODIS ash corrected 8.7 μm SO₂ masses vary between 0.4 and 2.7 Tg). The altitude retrieved using C12 for Kasatochi is consistent with 12.5±4 km reported by Karagulian et al. (2010) but the total SO.

²⁰ reported by Karagulian et al. (2010), but the total SO₂ load is around 30 % lower. Karagulian et al. (2010) use an IASI retrieval from ν_3 band (1362 cm⁻¹) and $\nu_1 + \nu_3$ band (2500 cm⁻¹). The altitudes from C12 are also comparable with the range reported in CALIPSO and OMI data from Wang et al. (2013) (Wang et al., 2013 used the OMI retrieval of altitude and amount and GEOS-Chem models to estimate that the forcing by Kasatochi volcanic sulphate aerosol became negligible six months after the eruption).

Sarychev is located at a similar latitude as Kasatochi, but in the Kurile islands south of Kamchatka. The IASI SO₂ scheme, C12, (similar to that previously reported by the Université Libre de Bruxelles, IASI-ULB, near real-time SO₂ alert system, http://cpm-ws4.ulb.ac.

be/Alerts/index.php, Rix et al., 2009; Haywood et al., 2010), retrieves a small tropospheric plume in the two images from 11 June 2009, followed by a higher plume on 12 June. The SO₂ loading increased in the following days (with a big injection the 15 June 2009) reaching a maximum on 16 June (0.6 Tg). Then the SO₂ load remained approximately constant for 16–18 June, before decreasing after that. The plume went in two directions, one branch spreading across the Pacific Ocean to North America and crossing Canada, reaching the Atlantic Ocean on 22 June and the Spanish coast of Europe on 24 June; the second branch went north, crossing Siberia up to the Siberian Sea and turning east to Greenland.

6.3 Africa: Dalafilla, Nabro

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- Two volcanic eruptions from the the Ethiopian Rift, Dalafilla in November 2008 and Nabro in June 2011, had a lower plume around 4–6 km and a higher part of the plume around the tropopause, but they were nearly one order of magnitude different in terms of SO₂ amount. Dalafilla produced a maximum SO₂ atmospheric loading of 0.2 Tg while Nabro produced 1.6 Tg. This could be the result of volcanic effects (for example Dalafilla was a very short-
- lived and vigorous fire-fountaining eruption from an extended fissure that produced, in some parts low mass eruption rates and in other parts much higher eruption rates. Another possible explanation of how a medium/small eruption such as Dalafilla (VEI 3) reached the tropopause altitude is illustrated in Tupper et al. (2009) and implicated the effect of a moist atmosphere. Tupper et al. (2009) showed that volcanic emissions with different eruption rates in moist atmosphere both reached the tropopause; in dry atmosphere they reach different altitudes.

The first IASI observation showing the Dalafilla eruption was at 05:00 UTC on 4 November 2008. The plume is divided into a lower and a higher part. These spread north and east from Ethiopia, covering the Arabic peninsula, and arrived over north India and the ²⁵ Himalayas on the 5 November. They spread then over China and a diluted plume arrived over south Japan on 6 November. IASI detected SO₂ near the volcano in every image in this period. According to Meteosat-9 false color RGB images, the eruption started between

12:45 UTC and 13:00 UTC on 3 November 2008, with high-level SO₂ plumes and an ash plume (mixed with ice). The low-level SO₂ plume, started three hours later².

The IASI images of SO_2 amount and altitude are consistent with the measurements over northern India reported by Mallik et al. (2013). In particular Mallik et al. (2013) report: (i) high

- ⁵ concentration of column SO₂ from ground measurements where the time of exceptionally high SO₂ amount is consistent with the IASI plume arriving over the Indian location; (ii) CALIPSO observations of a scattering feature around 4–5 km altitude on 6 November that is consistent with the lower part of the plume reported here by IASI; (iii) OMI maps, obtained assuming a fixed altitude, with a position of the SO₂ plume similar to IASI. With the use of
- ¹⁰ C12 IASI scheme it is possible to discern that there is both a lower plume in the troposphere (as reported by Mallik et al., 2013) and a significant part of the plume at altitudes around the tropopause.

Nabro produced the largest amount of SO_2 in any volcanic plume observed by IASI with a maximum of up to $1.6\pm0.3\,Tg$ of SO_2.

Numerous previous studies have documented this eruption, with many of these focusing 15 on whether Nabro injected SO₂ directly into the stratosphere or whether the injection was associated with monsoon circulation (Bourassa et al., 2012, 2013; Sawamura et al., 2012; Vernier et al., 2013; Fromm et al., 2013, 2014; Clarisse et al., 2014; Biondi et al., 2015). A more detailed study of the Nabro eruption, also using the C12 IASI retrieval scheme, is reported in Fromm et al. (2014), and concluded that "Nabro injected sulphur directly to or 20 above the tropopause upon the initial eruption on 12/13 June, and again on 16 June 2011". Here we include the Nabro summary of the C12 IASI dataset. The Nabro plume is retrieved on 13 June, with a plume over north-east Africa between 14 and 18 km height. On the 14 the plume arrived over the Middle East and over central Asia on 15 June. The eruption formed two plumes at different altitudes, the higher one that reached the stratosphere and 25 a lower one that remained confined to the troposphere with less than 10 km altitude. The higher plume is further separated into two segments, a "north" one (15 km and above) and a "south" one, a bit lower. Over all these days the plume was still attached to the volcano,

²http://www.eumetsat.int/website/home/Images/ImageLibrary/DAT_IL_08_11_03_A.html

indicating continuous injection. On 17 June there is a lower altitude plume and a new high altitude part going over north-east Africa.

In Fromm et al. (2014) two comparisons have demonstrated the consistency of IASI altitude with other measurements: (i) morning and afternoon IASI data of the 14 June are compared with the lidar ground data at Sede Boker (Israel) and an SO₂ profile from MLS (Fromm et al., 2014; Fig. 8); (ii) the night time IASI measurements of 17 June are compared with the CALIOP and MLS measurements (Fromm et al., 2014; Fig. 9).

6.4 Iceland: Eyjafjallajökull and Grímsvötn

Eruptions of Eyjafjallajökull in April and May 2010 and Grímsvötn in May 2011 have been examined. A deeper analysis of the IASI SO₂ plume from the Eyjafjallajökull eruption is presented in Carboni et al. (2012) and here we only report the time series of vertical distribution.

The first IASI observation of the Eyjafjallajökull plume is in the evening of 14 April 2010. The SO_2 plume altitudes retrieved in successive observations up to 20 April are below

- 10 km, mainly confined below 5 km. Small amounts of SO₂ between 16 and 20 April are between 5 and 10 km in altitude. From 20 to 30 April SO₂ is always below 5 km. There is a little increase of SO₂ amount and altitude on the 1 and 2 May, and a more pronounced increase of both SO₂ amount and altitude from the 5 May. The retrieved altitude has a maximum during the 14–17 May period, where values reach around 10 km.
- ²⁰ Grímsvötn's plume was first observed by IASI on the morning of 22 May 2011. From the afternoon of 22 May it is possible to see an SO₂ rich plume going north, with a segment going north-east, and a lower plume going south/south-east. The denser part of the higher SO₂ plume moved west and arrived over Greenland, and out of the analysed area on 24 May. This segment re-entered the analysed area on 26 May in the afternoon (this is why the last column of the plot has an apparent increase in SO₂). The lower altitude plume,
 - moving in south/south-east direction, is reported to be more ash rich (Moxnes et al., 2014) and travelled towards Europe, arriving over the northern UK on 24 May in the morning, and over Scandinavia in the afternoon. These observations of the Grímsvötn eruption do not

completely represent the total atmospheric SO_2 due to the fact that a part of the plume is missing.

6.5 Minor frequent events: Etna and Congo

The activity of Etna and Nyamuragira/Nyiragongo during the period was limited to smaller
eruptions and lava-fountains. Here we do not report an exhaustive list of these episodes, but a few examples to show how these emissions can be observed by IASI. The Smithsonian Institution reported that Nyamuragira erupted on 2 January 2010 and 6 November 2011. From IASI data it is not possible to distinguish between Nyamuragira and Nyriagongo emissions, but SO₂ plumes are observed from 3 to 12 January 2010 and from 7 to 15 November 2011. The SO₂ plumes spread over the equatorial area of Africa from Congo up to Sudan and Chad, with decreasing SO₂ loading away from the volcano. The main plumes were confined to the troposphere and disappeared after a few days.

The Etna events considered are the 23–25 November 2007 eruption and the 2011 lavafountains: 11–13 January, 17–19 February, 9–12 April, 11–13 May, 8–10 July, 18–20 July,

- ¹⁵ 24–26 July, 29 July, 1 August, 4–7 August, 11–13 August, 19–21 August, 28–30 August, 7–10 September, 18–20 September, 27–30 September, 12–17 November. Analysis was performed for the lava fountain period plus one day either side. The Etna eruption in 2007 produced the highest SO₂ atmospheric loading for the period analysed (more then 0.1 Tg). The lava-fountains are associated with smaller amounts of SO₂ and with large error. Etna emissions are appreximately ton times amounts from the Conse. In particular.
- emissions are approximately ten times smaller then those from the Congo. In particular, the SO₂ for 2011 Etna fountains have been validated within the SAMSH-SACS2 project and the SO₂ amount measured by IASI are consistent with ground observation (Spinetti et al., 2014).
- ²⁵ We now examine, for these eruptions, whether the VEI and SO₂-plume height are correlated or not. The definition of eruption altitude is not straightforward due to the change in plume altitude (in both space and time). The values of VEI in this study were taken from the Global Volcanism Program (2013) updated through 2015. VEI estimates may be subject to

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revision, but for contemporary eruptions typically refer to the 'peak' phase of the eruption; they are not good descriptors for long-lived eruptions (e.g. Montserrat), or eruptions of variable intensity. Fig. 12 shows different plume altitudes, and altitude ranges, for the eruptions with VEI 3, 4 and 5. The altitudes where the 50% SO₂ mass is contained and where the maximum mass is retrieved are obtained by averaging our vertical distributions (in Fig. 8-11) over time, except for Puyehue where only the first 48 hours are considered, as the rest of the emissions during this eruption were mainly injected lower than 10 km. Fig. 12 shows that some vigorous phases of eruptions like Montserrat and Dalafilla, which are both VEI 3, nonetheless injected SO₂ higher in the atmosphere than the VEI 5 Puyehue eruption, and that VEI and SO₂-plume height are not straightforwardly correlated.

7 Conclusions

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In this paper, the IASI SO₂ volcanic plume altitudes and amount from a recently developed SO₂ retrieval algorithm (Carboni et al., 2012) are presented. IASI has significant advantages for the monitoring of extended volcanic eruptions, because it can be used to track the intensity of the eruption, both in terms of gas amount and gas plume height twice a day.

- Comparisons of IASI altitudes against CALIPSO profiles and IASI SO₂ column amounts against Brewer ground measurements have been performed. These show that IASI retrieved values are consistent with these satellite and ground measurements. Despite the ability of IASI to retrieve the plume altitude, it could be difficult to distinguish, with only IASI data, if the eruption injects into the stratosphere or into the high troposphere. Further work integrating and comparing IASI with other measurements, such as lidar or limb mea
 - surements (e.g. Michelson Interferometer for Passive Atmospheric Sounding MIPAS), are needed to better assess the ability of IASI to determine injection into the stratosphere.

The IASI scheme has been used to follow the vertical distribution of SO₂ as a function of time (twice daily), for different eruption types (e.g. VEI ranging between 1 and 5) and different latitudes. This work suggest that while VEI is a convenient and rapidly estimated proxy for eruption "scale", it is a poor index of the potential height to which volcanic SO₂ is injected.

There is a tendency for volcanic SO_2 plumes to reach a point of buoyancy near the tropopause. All of the eruptions in the tropics (except Nyamuragira), reached the tropopause. In the mid latitudes, the eruptions of Eyjafjallajökull, Llaima, Copahue and Etna remained confined in the troposphere.

The dataset can be made available by contacting the author Elisa Carboni <elisa.carboni@physics.ox.ac.uk>.

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References

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Aiuppa, A., Cannata, A., Cannavo, F., Di Grazia, G., Ferrari, F., Giudice, G., Gurrieri, S., Liuzzo, M., Mattia, M., Montalto, P., Patan, D., and Puglisi, G.: Patterns in the recent 2007–2008 activity of Mount Etna volcano investigated by integrated geophysical and geochemical observations, Geochem. Geophy. Geosy., 11, Q09008, doi:10.1029/2010GC003168, 2010.

Behncke, B., Branca, S., Corsaro, R., De Beni, E., Miraglia, L., and Proietti, C.: The 2011–2012

- summit activity of Mount Etna: birth, growth and products of the new SE crater, J. Volcanol. Geoth. Res., 270, 10–21, doi:10.1016/j.jvolgeores.2013.11.012, 2014.
 - Bignami, C., Corradini, S., Merucci, L., De Michele, M., Raucoules, D., De Astis, G., Stramondo, S., and Piedra, J.: Multisensor satellite monitoring of the 2011 Puyehue-Cordon Caulle Eruption, IEEE J. Sel. Top. Appl., 7, 2786, doi:10.1109/JSTARS.2014.2320638, 2014.
- ²⁵ Biondi, R., Steiner, A., Kirchengast, G., Brenot, H., and Rieckh, T.: A novel technique including GPS radio occultation for detecting and monitoring volcanic clouds, Geophys. Res. Lett., in revision, 2015.

Blumstein, D., Chalon, G., Carlier, T., Buil, C., Hebert, P., Maciaszek, T., Ponce, G., Phulpin, T.,

- ⁵ Tournier, B., Simeoni, D., Astruc, P., Clauss, A., Kayal, G., and Jegou, R.: IASI instrument: technical overview and measured performances, Proceedings of SPIE – The International Society for Optical Engineering, 5543, 196–207, doi:10.1117/12.560907, 2004.
 - Bluth, G. and Carn, S.: Exceptional sulfur degassing from Nyamuragira volcano, 1979–2005, Int. J. Remote Sens., 29, 6667–6685, doi:10.1080/01431160802168434, 2008.
- Boichu, M., Menut, L., Khvorostyanov, D., Clarisse, L., Clerbaux, C., Turquety, S., and Coheur, P.-F.: Inverting for volcanic SO₂ flux at high temporal resolution using spaceborne plume imagery and chemistry-transport modelling: the 2010 Eyjafjallajökull eruption case study, Atmos. Chem. Phys., 13, 8569–8584, doi:10.5194/acp-13-8569-2013, 2013.

- monitoring of volcanic ash clouds: outcomes from the first IAVCEI-WMO workshop on ash dispersal forecast and civil aviation, B. Volcanol., 74, 1–10, doi:10.1007/s00445-011-0508-6, 2012.
 Bonali, F.: Earthquake-induced static stress change on magma pathway in promoting the 2012 Copahue eruption, Tectonophysics, 608, 127–137, doi:10.1016/j.tecto.2013.10.006, 2013.
- Bourassa, A., Robock, A., Randel, W., Deshler, T., Rieger, L. A., Lloyd, N., Llewellyn, E., and De genstein, D.: Large volcanic aerosol load in the stratosphere linked to asian monsoon transport, Science, 337, 78–81, doi:10.1126/science.1219371, 2012.
 - Bourassa, A., Robock, A., Randel, W., Deshler, T., Rieger, L., Lloyd, N., Llewellyn, E., and Degenstein, D.: Response to comments on "large volcanic aerosol load in the stratosphere linked to Asian monsoon transport", Science, 339, 647, doi:10.1126/science.1227961, 2013.
- Brenot, H., Theys, N., Clarisse, L., van Geffen, J., van Gent, J., Van Roozendael, M., van der A, R., Hurtmans, D., Coheur, P.-F., Clerbaux, C., Valks, P., Hedelt, P., Prata, F., Rasson, O., Sievers, K., and Zehner, C.: Support to Aviation Control Service (SACS): an online service for near-realtime satellite monitoring of volcanic plumes, Nat. Hazards Earth Syst. Sci., 14, 1099–1123, doi:10.5194/nhess-14-1099-2014, 2014.
- Budi-Santoso, A., Lesage, P., Dwiyono, S., Sumarti, S., Subandriyo, Surono, Jousset, P., and Metaxian, J.-P.: Analysis of the seismic activity associated with the 2010 eruption of Merapi Volcano, Java, J. Volcanol. Geoth. Res., 261, 153–170, doi:10.1016/j.jvolgeores.2013.03.024, 2013.

Bonadonna, C., Folch, A., Loughlin, S., and Puempel, H.: Future developments in modelling and monitoring of volcanic ash clouds: outcomes from the first IAVCEI-WMO workshop on ash disper-

Calabrese, S., Aiuppa, A., Allard, P., Bagnato, E., Bellomo, S., Brusca, L., D'Alessandro, W., and Parello, F.: Atmospheric sources and sinks of volcanogenic elements in a basaltic volcano (Etna, Italy), Geochim. Cosmochim. Ac., 75, 7401–7425, doi:10.1016/j.gca.2011.09.040, 2011.

Campion, R.: New lava lake at Nyamuragira volcano revealed by combined ASTER and OMI SO2

- measurements, Geophys. Res. Lett., 41, 7485–7492, doi:10.1002/2014GL061808, 2014. Campion, R., Salerno, G., Coheur, P.-F., Hurtmans, D., Clarisse, L., Kazahaya, K., Burton, M., Caltabiano, T., Clerbaux, C., and Bernard, A.: Measuring volcanic degassing of SO₂ in the lower troposphere with ASTER band ratios, J. Volcanol. Geoth. Res., 194, 42–54, doi:10.1016/j.jvolgeores.2010.04.010, 2010.
- ¹⁰ Carboni, E., Grainger, R., Walker, J., Dudhia, A., and Siddans, R.: A new scheme for sulphur dioxide retrieval from IASI measurements: application to the Eyjafjallajökull eruption of April and May 2010, Atmos. Chem. Phys., 12, 11417–11434, doi:10.5194/acp-12-11417-2012, 2012.

Carn, S. A. and Bluth, G. J. S.: Prodigious sulfur dioxide emissions from Nyamuragira volcano, D. R. Congo, Geophys. Res. Lett., 30, 2211, doi:10.1029/2003GL018465, 2003.

- ¹⁵ Carn, S. A., and Prata, F. J.: Satellite-based constraints on explosive SO₂ release from Soufriere Hills Volcano, Montserrat, Geophys. Res. Lett., 37, L00E22, doi:10.1029/2010GL044971, 2010. Carn, S., Krueger, A., Krotkov, N., Yang, K., and Evans, K.: Tracking volcanic sulfur dioxide clouds for aviation hazard mitigation, Nat. Hazards, 51, 325–343, doi:10.1007/s11069-008-9228-4, 2009. Carn, S., Krotkov, N., Yang, K., and Krueger, A.: Measuring global volcanic degassing with the ozone
- ²⁰ monitoring instrument (OMI), Geol. Soc. Sp., 380, 229–257, doi:10.1144/SP380.12, 2013. Carn, S. A. and Lopez, T. M.: Opportunistic validation of sulfur dioxide in the Sarychev Peak volcanic eruption cloud, Atmos. Meas. Tech., 4, 1705–1712, doi:10.5194/amt-4-1705-2011, 2011.
 - Carn, S. A., Krueger, A. J., Bluth, G. J. S., Schaefer, S. J., Krotkov, N. A., Watson, I. M., and Datta, S.: Volcanic eruption detection by the Total Ozone Mapping Spectrometer (TOMS) instrument: a
- 22-year record of sulfur dioxide and ash emissions, in: Volcanic Degassing, edited by: Oppenheimer, C., Pyle, D. M., and Barclay, J., vol. 213, Geological Society, London, 177–203, 2003.
 Christopher, T., Edmonds, M., Humphreys, M., and Herd, R.: Volcanic gas emissions from Soufriere Hills Volcano, Montserrat 1995–2009, with implications for mafic magma supply and degassing,
 - Geophys. Res. Lett., 37, L00E04, doi:10.1029/2009GL041325, 2010.
- ³⁰ Christopher, T., Edmonds, M., Taisne, B., Odbert, H., Costa, A., Hards, V., and Wadge, G.: Periodic sulphur dioxide degassing from the Soufriere Hills Volcano related to deep magma supply, Geological Society, London, Special Publications, 410, doi:10.1144/SP410.11, 2014.

Clarisse, L., Coheur, P.-F., Theys, N., Hurtmans, D., and Clerbaux, C.: The 2011 Nabro eruption,

- a SO₂ plume height analysis using IASI measurements, Atmos. Chem. Phys., 14, 3095–3111, 5 doi:10.5194/acp-14-3095-2014, 2014.
 - Clerbaux, C., Coheur, P.-F., Clarisse, L., Hadji-Lazaro, J., Hurtmans, D., Turquety, S., Bowman, K., Worden, H., and Carn, S.: Measurements of SO₂ profiles in volcanic plumes from the NASA Tropospheric Emission Spectrometer (TES), Geophys. Res. Lett., 35, L22807, doi:10.1029/2008GL035566, 2008.
- 10
 - Clerbaux, C., Boynard, A., Clarisse, L., George, M., Hadji-Lazaro, J., Herbin, H., Hurtmans, D., Pommier, M., Razavi, A., Turguety, S., Wespes, C., and Coheur, P.-F.: Monitoring of atmospheric composition using the thermal infrared IASI/MetOp sounder, Atmos. Chem. Phys., 9, 6041-6054, doi:10.5194/acp-9-6041-2009, 2009.
- Corradini, S., Merucci, L., and Prata, A. J.: Retrieval of SO₂ from thermal infrared satellite measure-15 ments: correction procedures for the effects of volcanic ash, Atmos. Meas. Tech., 2, 177-191, doi:10.5194/amt-2-177-2009. 2009.
 - Corradini, S., Merucci, L., Prata, A. J., and Piscini, A.: Volcanic ash and SO₂ in the 2008 Kasatochi eruption: retrievals comparison from different IR satellite sensors, J. Geophys. Res., 115, D00L21,
- doi:10.1029/2009JD013634, 2010. 20
 - Costa, F., Andreastuti, S., Bouvet de Maisonneuve, C., and Pallister, J.: Petrological insights into the storage conditions, and magmatic processes that yielded the centennial 2010 Merapi explosive eruption, J. Volcanol. Geoth. Res., 261, 209-235, doi:10.1016/j.jvolgeores.2012.12.025, 2013. Cronin, S., Lube, G., Davudi, D., Sumarti, S., Subrandivo, S., and Surono: Insights into the October-
- November 2010 Gunung Merapi eruption (Central Java, Indonesia) from the stratigraphy, vol-25 ume and characteristics of its pyroclastic deposits, J. Volcanol. Geoth. Res., 261, 244-259,
 - doi:10.1016/j.jvolgeores.2013.01.005, 2013.
 - Cuoco, E., Tedesco, D., Poreda, R., Williams, J., De Francesco, S., Balagizi, C., and Darrah, T.: Impact of volcanic plume emissions on rain water chemistry during the January 2010 Nyamuragira
- eruptive event: implications for essential potable water resources, J. Hazard. Mater., 244-245, 30 570-581, doi:10.1016/j.jhazmat.2012.10.055, 2013.

27

De Muer, D. and De Backer, H.: Revision of 20 years of Dobson total ozone data at Uccle (Belgium): fictitious Dobson total ozone trends induced by sulfur dioxide trends, J. Geophys. Res.-Atmos., 97, 5921–5937, doi:10.1029/91JD03164, 1992.

Delmelle, P.: Environmental impacts of tropospheric volcanic gas plumes, Geol. Soc. Sp., 213, 381– 399. doi:10.1144/GSL.SP.2003.213.01.23, 2003.

5

20

 Delmelle, P., Stix, J., Baxter, P. J., Garcia-Alvarez, J., and Barquero, J.: Atmospheric dispersion, environmental effects and potential health hazard associated with the low-altitude gas plume of Masaya volcano, Nicaragua, B. Volcanol., 64, 423–434, doi:10.1007/s00445-002-0221-6, 2002.
 Doeringer, D., Eldering, A., Boone, D., Gonzalez Abad, G., and Bernath, P.: Observation of sulfate

- aerosols and SO₂ from the Sarychev volcanic eruption using data from the Atmospheric Chemistry Experiment (ACE), J. Geophys. Res., 117, D03203, doi:10.1029/2011JD016556, 2012.
 - Dzierma Y. and Wehrmann H.: On the likelihood of future eruptions in the Chilean Southern Volcanic Zone: interpreting the past century?s eruption record based on statistical analyses, Andean Geology 39, 380-393, doi: 10.5027/andgeoV39n3-a02, 2012.
- Ebmeier, S. K., Sayer, A. M., Grainger, R. G., Mather, T. A., and Carboni, E.: Systematic satellite observations of the impact of aerosols from passive volcanic degassing on local cloud properties, Atmos. Chem. Phys., 14, 10601–10618, doi:10.5194/acp-14-10601-2014, 2014.

Encinas-Oropesa, A., Drew, G., Hardy, M., Leggett, A., Nicholls, J., and Simms, N.: Effects of oxidation and hot corrosion in a nickel disc alloy, in: Superalloys Conference Proceedings, 609–618, 2008.

Flemming, J. and Inness, A.: Volcanic sulfur dioxide plume forecasts based on UV satellite retrievals for the 2011 Grimsvotn and the 2010 Eyjafjallajökull eruption, J. Geophys. Res.-Atmos., 118, 10172–10189, doi:10.1002/jgrd.50753,2, 2013.

Fromm, M., Nedoluha, G., and Charvát, Z.: Comment on 'Large volcanic aerosol load in the strato-

- sphere linked to Asian monsoon transport", Science, 339, 647, doi:10.1126/science.1228605, 2013.
 - Fromm, M., Kablick, G., Nedoluha, G., Carboni, E., Grainger, R., Campbell, J., and Lewis, J.: Correcting the record of volcanic stratospheric aerosol impact: Nabro and Sarychev Peak, J. Geophys. Res., 119, 10343–10364, doi:10.1002/2014JD021507, 2014.
- von Glasow, R.: Atmospheric chemistry in volcanic plumes, P. Natl. Acad. Sci. USA, 107, 6594– 6599, doi:10.1073/pnas.0913164107, 2010.
 - Global Volcanism Program. Volcanoes of the World, v. 4.4.2. Venzke, E (ed.). Smithsonian Institution. doi:10.5479/si.GVP.VOTW4-2013, 2013. Downloaded 02 Mar 2016.

- Graf, H.-F., Feichter, J., and Langmann, B.: Volcanic sulfur emissions: estimates of source strength and its contribution to the global sulfate distribution, J. Geophys. Res.-Atmos., 102, 10727–10738, doi:10.1029/96JD03265, 1997.
- Guerrieri, L., Merucci, L., Corradini, S., and Pugnaghi, S.: Evolution of the 2011 Mt. Etna ash and
- SO₂ lava fountain episodes using SEVIRI data and VPR retrieval approach, J. Volcanol. Geoth. Res., 291, 63–71, doi:10.1016/j.jvolgeores.2014.12.016, 2015.

5

10

30

- Haywood, J., Jones, A., Clarisse, L., Bourassa, A., Barnes, J., Telford, P., Bellouin, N., Boucher, O., Agnew, P., Clerbaux, C., Coheur, P., Degenstein, D., and Braesicke, P.: Observations of the eruption of the Sarychev volcano and simulations using the HadGEM2 climate model, J. Geophys. Res.-Atmos., 115, D21212, doi:10.1029/2010JD014447, 2010.
- Hilton, F., R. Armante, T. August et al., Hyperspectral earth observation from IASI: Five years of accomplishments. Bulletin of the American Meteorological Society, 93, 347–370, doi:10.1175/BAMS-D-11-00027.1, 2012.
- Höpfner, M., Glatthor, N., Grabowski, U., Kellmann, S., Kiefer, M., Linden, A., Orphal, J., Stiller, G.,
- von Clarmann, T., Funke, B., and Boone, C. D.: Sulfur dioxide (SO₂) as observed by MI-PAS/Envisat: temporal development and spatial distribution at 15–45 km altitude, Atmos. Chem. Phys., 13, 10405–10423, doi:10.5194/acp-13-10405-2013, 2013.
 - Innocenti, S., Andreastuti, S., Furman, T., del Marmol, M.-A., and Voight, B.: The pre-eruption conditions for explosive eruptions at Merapi volcano as revealed by crystal texture and mineralogy, J.
- Volcanol. Geoth. Res., 261, 69–86, doi:10.1016/j.jvolgeores.2012.12.028, 2013a.
 Innocenti, S., del Marmol, M.-A., Voight, B., Andreastuti, S., and Furman, T.: Textural and mineral chemistry constraints on evolution of Merapi Volcano, Indonesia, J. Volcanol. Geoth. Res., 261, 20–37, 2013b.
 - Jenkins, S., Komorowski, J.-C., Baxter, P., Spence, R., Picquout, A., Lavigne, F., and
- Surono: The Merapi 2010 eruption: an interdisciplinary impact assessment methodology for studying pyroclastic density current dynamics, J. Volcanol. Geoth. Res., 261, 316–329, doi:10.1016/j.jvolgeores.2013.02.012, 2013.
 - Karagulian, F., Clarisse, L., Clerbaux, C., Prata, A. J., Hurtmans, D., and Coheur, P. F.: Detection of volcanic SO₂, ash, and H₂SO₄ using the Infrared Atmospheric Sounding Interferometer (IASI), J. Geophys. Res., 115, D00L02, doi:10.1029/2009JD012786, 2010.
- Kerr, J., McElroy, C., and Olafson, R.: Measurements of ozone with the Brewer spectrophotometer, in: Proceedings of the Quadrennial International Ozone Symposium, edited by: London, J., Natl. Cent. for Atmos. Res., Boulder, Colo., vol. 1, 74–79, 1981.

Koukouli, M. E., Clarisse, L., E., C., van Gent, J., C., S., Balis, D., Dimopoulos, S., R. G.3, G., Theys, N., Tampellini, L., and Zehner, C.: Intercomparison of Metop-A SO₂ measurements during the 2010–2011 Icelandic eruptions, Ann. Geophys.-Italy, 57, doi:10.4401/ag-6613, 2014.

Koukouli, M. E., Balis, D., Dimopoulos, S., Siomos, N.,: SACS2/SMASH Validation Report on the Eyjafjallajökull and Grímsvötn eruptions, ESA report (last accessed: 17 December 2015), 2014.

- Eyjafjallajökull and Grimsvötn eruptions, ESA report (last accessed: 17 December 2015), 2014. Kristiansen, N., Stohl, A., Prata, A., Richter, A., Eckhardt, S., Seibert, P., Hoffmann, A., Ritter, C., Bitar, L., Duck, T., and Stebel, K.: Remote sensing and inverse transport modeling of the Kasatochi eruption sulfur dioxide cloud, J. Geophys. Res.-Atmos., 115, D00L16, doi:10.1029/2009JD013286, 2010.
- Kroll, J., Cross, E., Hunter, J., Pai, S., TREX XII, TREX XI, Wallace, L., Croteau, P., Jayne, J., Worsnop, D., Heald, C., Murphy, J., and Frankel, S.: Atmospheric evolution of sulfur emissions from Kilauea: real-time measurements of oxidation, dilution, and neutralization within a volcanic plume, Environ. Sci. Technol., 49, 4129–4137, doi:10.1021/es506119x, 2015.

Krotkov, N., Cam, S., Krueger, A., Bhartia, P., and Yang, K.: Band residual difference algorithm for

- retrieval of SO₂ from the aura Ozone Monitoring Instrument (OMI), IEEE T. Geosci. Remote, 44, 1259–1266, doi:10.1109/TGRS.2005.861932, 2006.
 - Krotkov, N., Schoeberl, M., Morris, G., Carn, S., and Yang, K.: Dispersion and lifetime of the SO₂ cloud from the August 2008 Kasatochi eruption, J. Geophys. Res.-Atmos., 115, D00L20, doi:10.1029/2010JD013984, 2010.
- Luehr, B.-G., Koulakov, I., Rabbel, W., Zschau, J., Ratdomopurbo, A., Brotopuspito, K., Fauzi, P., and Sahara, D.: Fluid ascent and magma storage beneath Gunung Merapi revealed by multiscale seismic imaging, J. Volcanol. Geoth. Res., 261, 7–19, doi:10.1016/j.jvolgeores.2013.03.015, 2013.

Mallik, C., Lal, S., Naja, M., Chand, D., Venkataramani, S., Joshi, H., and Pant, P.: Enhanced SO₂

- concentrations observed over northern India: Role of long-range transport, Int. J. Remote Sens.,
 34, 2749–2762, doi:10.1080/01431161.2012.750773, 2013.
 - Marzano, F. S., Lamantea, M., Montopoli, M., Di Fabio, S., and Picciotti, E.: The Eyjafjöll explosive volcanic eruption from a microwave weather radar perspective, Atmos. Chem. Phys., 11, 9503–9518, doi:10.5194/acp-11-9503-2011, 2011.
- Mather, T., Pyle, D., and Oppenheimer, C.: Tropospheric volcanic aerosol, in: Volcanism and the Earth's Atmosphere, Geophysical Monograph 139, edited by: Robock, A. and Oppenheimer, C., Am. Geophys. Union, Washington, DC, 189–212, doi:10.1029/139GM12, 2003.

- McCormick, B., Edmonds, M., Mather, T., Campion, R., Hayer, C., Thomas, H., and Carn, S.: Volcano monitoring applications of the ozone monitoring instrument, Geol. Soc. Sp., 380, 259–291, doi:10.1144/SP380.11, 2013.
- McGonigle, A. J. S., Delmelle, P., Oppenheimer, C., Tsanev, V. I., Delfosse, T., Williams-Jones, G.,
- ⁵ Horton, K., and Mather, T. A.: SO₂ depletion in tropospheric volcanic plumes, Geophys. Res. Lett., 31, L13201, doi:10.1029/2004GL019990, 2004.
 - Moxnes, E., Kristiansen, N., Stohl, A., Clarisse, L., Durant, A., Weber, K., and Vogel, A.: Separation of ash and sulfur dioxide during the 2011 Grímsvötn eruption, J. Geophys. Res.-Atmos-, 119, 7477–7501, doi:10.1002/2013JD021129, 2014.
- Newhall, C.G., and Self, S.: The volcanic explosivity index (VEI) an estimate of explosive magnitude for historical volcanism. J. Geophys. Res. 87, 1231?1238, dai: 10.1029/JC087iC02p01231, 1982.
 - Nicholson, E., Mather, T., Pyle, D., Odbert, H., and Christopher, T.: Cyclical patterns in volcanic degassing revealed by SO₂ flux timeseries analysis: an application to Soufriére Hills Volcano, Montserrat, Earth Planet. Sc. Lett., 375, 209–221, doi:10.1016/j.epsl.2013.05.032, 2013.
- Montserrat, Earth Planet. Sc. Lett., 375, 209–221, doi:10.1016/j.epsl.2013.05.032, 2013.
 Nowlan, C. R., Liu, X., Chance, K., Cai, Z., Kurosu, T. P., Lee, C., and Martin, R. V.: Retrievals of sulfur dioxide from the Global Ozone Monitoring Experiment 2 (GOME-2) using an optimal estimation approach: algorithm and initial validation, J. Geophys. Res., 116, D18301, doi:10.1029/2011JD015808, 2011.
- Pagli, C., Wright, T., Ebinger, C., Yun, S.-H., Cann, J., Barnie, T., and Ayele, A.: Shallow axial magma chamber at the slow-spreading Erta Ale Ridge, Nat. Geosci., 5, 284–288, doi:10.1038/ngeo1414, 2012.
 - Pallister, J., Schneider, D., Griswold, J., Keeler, R., Burton, W., Noyles, C., Newhall, C., and Ratdomopurbo, A.: Merapi 2010 eruption-Chronology and extrusion rates monitored with
- satellite radar and used in eruption forecasting, J. Volcanol. Geoth. Res., 261, 144–152, doi:10.1016/j.jvolgeores.2012.07.012, 2013.
 - Patane, D., Aiuppa, A., Aloisi, M., Behncke, B., Cannata, A., Coltelli, M., Di Grazia, G., Gambino, S., Gurrieri, S., Mattia, M., and Salerno, G.: Insights into magma and fluid transfer at Mount Etna by a multiparametric approach: A model of the events leading to the 2011 eruptive cycle, J. Geophys.
- ³⁰ Res.-Solid, 118, 3519–3539, doi:10.1002/jgrb.50248, 2013.
 - Petersen, G. N., Bjornsson, H., and Arason, P.: The impact of the atmosphere on the Eyjafjallajökull 2010 eruption plume, J. Geophys. Res., 117, D00U07, doi:10.1029/2011JD016762, 2012.

Picquout, A., Lavigne, F., Mei, E., Grancher, D., Noer, C., Vidal, C., and Hadmoko, D.: Air traffic disturbance due to the 2010 Merapi volcano eruption, J. Volcanol. Geoth. Res., 261, 366–375, doi:10.1016/j.jvolgeores.2013.04.005, 2013.

Pyle, D.M., Beattie, P.D. and G.J.S. Bluth: Sulphur emissions to the stratosphere from explosive volcanic eruptions, Bulletin of Volcanology, 57, 663-671, dai:10.1007/s004450050119, 1996.

5

20

Prata, A. and Bernardo, C.: Retrieval of volcanic SO₂ column abundance from Atmospheric Infrared Sounder data, J. Geophys. Res.-Atmos., 112, D20204, doi:10.1029/2006JD007955, 2007.

Prata, A. J. and Kerkmann, J.: Simultaneous retrieval of volcanic ash and SO₂ using MSG-SEVIRI measurements, Geophys. Res. Lett., 34, L05813, doi:10.1029/2006GL028691, 2007.

- Prata, A. J., Carn, S. A., Stohl, A., and Kerkmann, J.: Long range transport and fate of a stratospheric volcanic cloud from Soufrière Hills volcano, Montserrat, Atmos. Chem. Phys., 7, 5093– 5103, doi:10.5194/acp-7-5093-2007, 2007.
 - Prata, A. J., Gangale, G., Clarisse, L., and Karagulian, F.: Ash and sulphur dioxide in the 2008 eruptions of Okmok and Kasatochi: insights from high spectral resolution satellite measurements, J. Geophys. Res., 115, D00L18, doi:10.1029/2009JD013556, 2010.
- Geophys. Res., 115, D00L18, doi:10.1029/2009JD013556, 2010.
 Pugnaghi, S., Gangale, G., Corradini, S., and Buongiorno, M.: Mt. Etna sulfur dioxide flux monitoring using ASTER-TIR data and atmospheric observations, J. Volcanol. Geoth. Res., 152, 74–90, doi:10.1016/j.jvolgeores.2005.10.004, 2006.

Pumphrey, H. C., Read, W. G., Livesey, N. J., and Yang, K.: Observations of volcanic SO₂ from MLS on Aura, Atmos. Meas. Tech., 8, 195–209, doi:10.5194/amt-8-195-2015, 2015.

- Rix, M., Valks, P., Hao, N., Loyola, D. G., Schlager, H., Huntrieser, H. H., Flemming, J., Koehler, U., Schumann, U., and Inness, A.: Volcanic SO₂, BrO and plume height estimations using GOME-2satellite measurementsduring the eruption of Eyjafjallajökull in May 2010, J. Geophys. Res., 117, D00U19, doi:10.1029/2011JD016718, 2012.
- ²⁵ Robock, A.: Volcanic eruptions and climate, Rev. Geophys., 38, 191–219, 2000.
- Rodgers, C. D.: Inverse Methods for Atmospheric Sounding: Theory and Practice, World Scientific, River Edge, N. J., 2000.

Saepuloh, A., Urai, M., Aisyah, N., Sunarta, Widiwijayanti, C., Subandriyo, and Jousset, P.: Interpretation of ground surface changes prior to the 2010 large eruption of Merapi volcano using

³⁰ ALOS/PALSAR, ASTER TIR and gas emission data, J. Volcanol. Geoth. Res., 261, 130–143, doi:10.1016/j.jvolgeores.2013.05.001, 2013.

Saunders, R. W., Matricardi, M., and Brunel, P.: An improved fast radiative transfer model for assimilation of satellite radiance observations, Q. J. Roy. Meteor. Soc., 125, 1407–1425, doi:10.1002/qj.1999.49712555615, 1999.

Sawamura, P., Vernier, J., Barnes, J., Berkoff, T., Welton, E., Alados-Arboledas, L., Navas-

- ⁵ Guzmán, F., Pappalardo, G., Mona, L., Madonna, F., Lange, D., Sicard, M., Godin-Beekmann, S., Payen, G., Wang, Z., Hu, S., Tripathi, S., Cordoba-Jabonero, C., and Hoff, R.: Stratospheric AOD after the 2011 eruption of Nabro volcano measured by lidars over the Northern Hemisphere, Environ. Res. Lett., 7, 034013, doi:10.1088/1748-9326/7/3/034013, 2012.
- Schmidt, A., Carslaw, K. S., Mann, G. W., Rap, A., Pringle, K. J., Spracklen, D. V., Wilson, M., and
 Forster, P. M.: Importance of tropospheric volcanic aerosol for indirect radiative forcing of climate, Atmos. Chem. Phys., 12, 7321–7339, doi:10.5194/acp-12-7321-2012, 2012.
 - Schmidt, A., Witham, C. S., Theys, N., Richards, N. A. D., Thordarson, T., Szpek, K., Feng, W., Hort, M. C., Woolley, A. M., Jones, A. R., Redington, A. L., Johnson, B. T., Hayward, C. L., and Carslaw, K. S.: Assessing hazards to aviation from sulfur dioxide emitted by explosive Icelandic eruptions, J. Geophys. Res.-Atmos., 119, 14180–14196, doi:10.1002/2014JD022070, 2014.
- eruptions, J. Geophys. Res.-Atmos., 119, 14180–14196, doi:10.1002/2014JD022070, 2014.
 Schmidt, A., Leadbetter, S., Theys, N., Carboni, E., Witham, C.S., Stevenson, J.A., Birch, C.E., Thordarson, T., Turnock, S., Barsotti, S., Delaney, L., Feng, W., Grainger, R.G., Hort, M.C., Hoskuldsson, A., Ialongo, I., Ilyinskaya, E., Johannsson, T., Kenny, P., Mather, T.A., Richards, N.A.D., Richards and Shepherd, J., Satellite detection, long-range transport, and air quality impacts of
- volcanic sulfur dioxide from the 2014-2015 flood lava eruption at Bárdarbunga (Iceland), Journal of Geophysical Research, 120, 9739?9757, doi:10.1002/2015JD023638, 2015.
 - Scollo, S., Prestifilippo, M., Pecora, E., Corradini, S., Merucci, L., Spata, G., and Coltelli, M.: Eruption column height estimation of the 2011-2013 Etna lava fountains, Ann. Geophys.-Italy, 57, S0214, doi:10.4401/ag-6396, 2014.
- ²⁵ Sears, T., Thomas, G., Carboni, E., Smith, A., and Grainger, R.: SO₂ as a possible proxy for volcanic ash in aviation hazard avoidance, J. Geophys. Res.-Atmos., 118, 5698–5709, doi:10.1002/jgrd.50505, 2013.
 - Seinfeld, J. H. and Pandis, S. N.: Atmospheric Chemistry and Physics: From Air Pollution to Climate Change, J. Wiley, New York, 1998.
- Smets, B., d'Oreye, N., Kervyn, F., Kervyn, M., Albino, F., Arellano, S., Bagalwa, M., Balagizi, C., Carn, S., Darrah, T., Fernandez, J., Galle, B., Gonzalez, P., Head, E., Karume, K., Kavotha, D., Lukaya, F., Mashagiro, N., Mavonga, G., Norman, P., Osodundu, E., Pallero, J., Prieto, J., Samsonov, S., Syauswa, M., Tedesco, D., Tiampo, K., Wauthier, C., and Yalire, M.: Detailed multi-

disciplinary monitoring reveals pre- and co-eruptive signals at Nyamulagira volcano (North Kivu, Democratic Republic of Congo), B. Volcanol., 76, 1–35, doi:10.1007/s00445-013-0787-1, 2014.

- Spinei, E., Carn, S., Krotkov, N., Mount, G., Yang, K., and Krueger, A.: Validation of ozone monitoring instrument SO₂ measurements in the Okmok volcanic cloud over Pullman, WA, July 2008, J. Geophys. Res.-Atmos., 115, D00L08, doi:10.1029/2009JD013492, 2010.
- Spinetti, C., Salerno, G., Caltabiano, T., Carboni, E., Clarisse, L., Corradini, S., Grainger, R., Hedelt, P., Koukouli, M., Merucci, L., Siddans, R., Tampellini, L., Theys, N., Valks, P., and Zehner, C.: Volcanic SO₂ by UV-TIR satellite retrievals: validation by using ground-based network at Mt. Etna, Ann. Geophys.-Italy, 57, doi:10.4401/ag-6641, 2014.

5

15

- ¹⁰ Stevenson, D., Johnson, C., Collins, W., and Derwent, R.: The tropospheric sulphur cycle and the role of volcanic SO₂, Geol. Soc. Sp., 213, 295–305, 2003.
 - Stevenson, J. A., Loughlin, S., Rae, C., Thordarson, T., Milodowski, A. E., Gilbert, J. S., Harangi, S., Lukács, R., Hojgaard, B., Árting, U., Pyne-O'Donnell, S., MacLeod, A., Whitney, B., and Cassidy, M.: Distal deposition of tephra from the Eyjafjallajökull 2010 summit eruption, J. Geophys. Res., 117, B00C10, doi:10.1029/2011JB008904, 2012.
- Stohl, A., Prata, A. J., Eckhardt, S., Clarisse, L., Durant, A., Henne, S., Kristiansen, N. I., Minikin, A., Schumann, U., Seibert, P., Stebel, K., Thomas, H. E., Thorsteinsson, T., Tørseth, K., and Weinzierl, B.: Determination of time- and height-resolved volcanic ash emissions and their use for quantitative ash dispersion modeling: the 2010 Eyjafjallajökull eruption, Atmos. Chem. Phys., 11, 4333–4351, doi:10.5194/acp-11-4333-2011, 2011.
- Surono, Jousset, P., Pallister, J., Boichu, M., Buongiorno, M., Budisantoso, A., Costa, F., Andreastuti, S., Prata, F., Schneider, D., Clarisse, L., Humaida, H., Sumarti, S., Bignami, C., Griswold, J., Carn, S., Oppenheimer, C., and Lavigne, F.: The 2010 explosive eruption of Java's Merapi volcano-A "100-year" event, J. Volcanol. Geoth. Res., 241–242, 121–135, doi:10.1016/j.jvolgeores.2012.06.018, 2012.
 - Textor, C., Graf, H.-F., Herzog, M., and Oberhuber, J.: Injection of gases into the stratosphere by explosive volcanic eruptions, J. Geophys. Res.-Atmos., 108, 4606, doi:10.1029/2002JD002987, 2003.
- Theys, N., Van Roozendael, M., Dils, B., Hendrick, F., Hao, N., and De Maziere, M.: First satellite detection of volcanic bromine monoxide emission after the Kasatochi eruption, Geophys. Res. Lett., 36, L03809, doi:10.1029/2008GL036552, 2009.
 - Theys, N., Campion, R., Clarisse, L., Brenot, H., van Gent, J., Dils, B., Corradini, S., Merucci, L., Coheur, P.-F., Van Roozendael, M., Hurtmans, D., Clerbaux, C., Tait, S., and Ferrucci, F.: Volcanic

- Theys, N., De Smedt, I., Van Roozendael, M., Froidevaux, L., Clarisse, L., and Hendrick, F.: First satellite detection of volcanic OCIO after the eruption of Puyehue-Cordon Caulle, Geophys. Res.
- Lett., 41, 667-672, doi:10.1002/2013GL058416, 2014. 5
 - Thomas, G. E. and Siddans, R.: Development of OCA type processors to volcanic ash detection and retrieval. Final Report, Tech. rep., EUMETSAT RFQ 13/715490. available at: ftp://ftp.rsg.rl.ac. uk/eumetsat ash/EUMETSAT RFQ 13-715490 final report v2.0.pdf (last access: 8 September 2015), 2015.
- Thomas, H. E. and Prata, A. J.: Sulphur dioxide as a volcanic ash proxy during the April-10 May 2010 eruption of Eyjafjallajökull Volcano, Iceland, Atmos. Chem. Phys., 11, 6871-6880, doi:10.5194/acp-11-6871-2011, 2011.
 - Tupper, A. and Wunderman, R.: Reducing discrepancies in ground and satellite-observed eruption heights, J. Volcanol. Geoth. Res., 186, 22–31, doi:10.1016/j.jvolgeores.2009.02.015, 2009.
- Tupper, A., Textor, C., Herzog, M., Graf, H.-F., and Richards, M.: Tall clouds from small eruptions: 15 the sensitivity of eruption height and fine ash content to tropospheric instability, Nat. Hazards, 51, 375-401. doi:10.1007/s11069-009-9433-9. 2009.
 - Van Gent, J., Spurr, R., Theys, N., and et al.: Towards operational retrieval of SO₂ plume height from GOME-2 radiance measurements, Atmos. Meas. Tech. Discuss., in preparation, 2015.
- Velez, M., Euillades, P., Caselli, A., Blanco, M., and Diaz, J.: Deformation of Copahue volcano: 20 inversion of InSAR data using a genetic algorithm, J. Volcanol. Geoth. Res., 202, 117-126, doi:10.1016/j.jvolgeores.2011.01.012, 2011.
 - Vernier, J.-P., Thomason, L., Fairlie, T., Minnis, P., Palikonda, R., and Bedka, K.: Comment on "large volcanic aerosol load in the stratosphere linked to Asian monsoon transport", Science, 339, 647d, doi:10.1126/science.1227817, 2013.
- 25
 - Viccaro, M., Garozzo, I., Cannata, A., Di Grazia, G., and Gresta, S.: Gas burst vs. gasrich magma recharge: a multidisciplinary study to reveal factors controlling triggering of the recent paroxysmal eruptions at Mt. Etna, J. Volcanol. Geoth. Res., 278-279, 1-13, doi:10.1016/j.jvolgeores.2014.04.001, 2014.
- Wadge, G. and Burt, L.: Stress field control of eruption dynamics at a rift volcano: Nyamuragira, D.R. 30 Congo, J. Volcanol. Geoth. Res., 207, 1–15, doi:10.1016/j.jvolgeores.2011.06.012, 2011.

- Walker, J. C., Dudhia, A., and Carboni, E.: An effective method for the detection of trace species demonstrated using the MetOp Infrared Atmospheric Sounding Interferometer, Atmos. Meas. Tech., 4, 1567–1580, doi:10.5194/amt-4-1567-2011, 2011.
- Walker, J. C., Carboni, E., Dudhia, A., and Grainger, R. G.: Improved detection of sulphur dioxide
- in volcanic plumes using satellite-based hyperspectral infra-red measurements: application to the Eyjafjallajökull 2010 eruption, J. Geophys. Res., 117, D00U16, doi:10.1029/2011JD016810, 2012.
 - Wang, J., Park, S., Zeng, J., Ge, C., Yang, K., Carn, S., Krotkov, N., and Omar, A. H.: Modeling of 2008 Kasatochi volcanic sulfate direct radiative forcing: assimilation of OMI SO₂ plume height data
- and comparison with MODIS and CALIOP observations, Atmos. Chem. Phys., 13, 1895–1912, doi:10.5194/acp-13-1895-2013, 2013.
 - Watson, I., Realmuto, V., Rose, W., Prata, A., Bluth, G., Gu, Y., Bader, C., and Yu, T.: Thermal infrared remote sensing of volcanic emissions using the moderate resolution imaging spectroradiometer, J. Volcanol. Geoth. Res., 135, 75–89, doi:10.1016/j.jvolgeores.2003.12.017, 2004.
- ¹⁵ Winker, D., Vaughan, M., Omar, A., Hu, Y., Powell, K., Liu, Z., Hunt, W., and Young, S.: Overview of the CALIPSO mission and CALIOP data processing algorithms, J. Atmos. Ocean. Tech., 26, 2310–2323, doi:10.1175/2009JTECHA1281.1, 2009.
 - Yang, K., Krotkov, N., Krueger, A., Carn, S., Bhartia, P., and Levelt, P.: Improving retrieval of volcanic sulfur dioxide from backscattered UV satellite observations L03102, Geophys. Res. Lett., 36,
- 20 L03102, doi:10.1029/2008GL036036, 2009.
 - Yang, K., Liu, X., Bhartia, P., Krotkov, N., Carn, S., Hughes, E., Krueger, A., Spurr, R., and Trahan, S.: Direct retrieval of sulfur dioxide amount and altitude from spaceborne hyperspectral UV measurements: theory and application, J. Geophys. Res.-Atmos., 115, D00L09, doi:10.1029/2010JD013982, 2010.
- Zerefos, C., Ganev, K., Kourtidis, K., Tzortziou, M., Vasaras, A., and Syrakov, E.: On the origin of SO₂ above Northern Greece, Geophys. Res. Lett., 27, 365–368, doi:10.1029/1999GL010799, 2000.

Table 1. Summary of studied eruptions, in chronological order, together with other relevant literature, volcanic explosivity index (VEI) and a short descriptions of the events from the Smithsonian Institution Global Volcanism Programme website (http://www.volcano.si.edu).

Volcano name	Analysed dates	Lat/Lon/ Ele- vation	Comments on IASI SO ₂ plume	References to other work	Volcanic event description (Smithsonian weekly reports)
Llaima (Chile)	2–6 Jan 2008	38.692° S 71.729° W 3125 m	$ \begin{array}{l} \mbox{Small eruption (maximum of 0.04 Tg of O_2).} \\ \mbox{Plume connected to the volcano only on the first day.} \\ \mbox{Confined in troposphere.} \end{array} $	McCormick et al. (2013): OMI SO ₂ study.	VEI 3. Strombolian eruption began 1 Jan; ash plumes to 12.5 km on 2 Jan, and to 3.7 km on 3 Jan.
Okmok (Aleutian islands)	12–20 Jul 2008	53.43° N 168.13° W 1073 m	SO ₂ injected around tropopause and the plume spreads south (pacific ocean) and east (north America) from the volcano. Presence of intermittent smaller and lower plume connected with the volcano. Main plume altitude increases with time following the tropopause altitude.	Spinei et al. (2010): OMI SO ₂ validation against ground measurements over north America; Prata et al. (2010): SO ₂ and ash from AIRS.	VEI 4. Strong explosive eruption began 12 Jul, with ash to at least 15 km. Erup- tion intensity declined, with ash plumes to 6 km on 17 Jul. Eruptions on 19 Jul sent ash to 9 km.
Kasatochi (Aleutian islands)	7–22 Aug 2008	52.177° N 175.508° W 314 m	SO ₂ amount reached the maximum after seven days indicating continuous injec- tion from volcano. Within ten days SO ₂ plume affected all latitudes between 30–90° N. Reached to tropopause and strato- sphere.	Krotkov et al. (2010): Dispersion and life- time of the SO ₂ using OMI; Yang et al. (2010): OMI retrieval of SO ₂ amount and altitude; Karagulian et al. (2010): SO ₂ and ash from IASI; Corradini et al. (2010): Comparison of different IR instruments (AIRS, MODIS); Kristiansen et al. (2010): inverse trans- port modelling; Bitar et al. (2010): lidar measurements, injected material into the troposphere and lower stratosphere; Prata et al. (2011): GOME-2 optimal estimation retrieval of amount and alti- tude; Wang et al. (2013): Volcanic sulfate forc- ing using OMI SO ₂ .	VEI 4. Explosive eruption began on 7 Aug, with ash plumes to 14 km; con- tinuous eruptions from 8–9 Aug, with ash plumes to 9–14 km. Subsequent obser- vations obscured by cloud.
Alu-Dalaffilla (Ethiopia)	4–7 Nov 2008	13.792° N 40.55° E 613 m	Plume is divided in two parts from the beginning (one lower in troposphere and one higher into tropopause /strato- sphere). SO ₂ near the volcano in every image (continuous emission). Nearly one order of magnitude smaller than Nabro (in amount of SO ₂) but reached comparable height.	Pagli et al. (2012): Ethiopian Rift defor- mation paper, studying the source of Alu- Dalafilla Eruption. Mallik et al. (2013): Enhanced SO ₂ con- centrations observed over northern India in Nov 2008.	VEI 3 – Mainly effusive eruption. Strong fissure eruption began on 3 Nov, for a few hours, waned rapidly and ended on 6 Nov. (Pagli et al., 2012). No direct ob- servations of the eruption.

Table 1. Continued.

Volcano name	Analysed dates	Lat/Lon/ Ele- vation	Comments on IASI SO ₂ plume	References to other work	Volcanic event description (Smithsonian weekly reports)
Sarychev Peak (Kuril islands)	11–26 Jun 2009	48.092° N 153.2° E 1496 m	Started with a small tropospheric plume building up with increasing SO ₂ load, maximum on IG Jun (0.6 Tg). Reached tropopause and stratosphere.	Theys et al. (2009): Volcanic BrO from GOME-2. Haywood et al. (2010): Climate model (HadGEM2), IASI SO ₂ , OSIRIS limb sounder and CALIPSO lidar measurements to investigate the distributions of SO ₂ and radiative impact of sulphate aerosol; Carn and Lopez (2011): validation of OMI SO ₂ ; Doeringer et al. (2012): sulphate aerosols and SO ₂ from the Atmospheric Chemistry Experiment (ACE) measurements; Fromm et al. (2014); on correction for 20	VEI 4. Eruption began 11 Jun, with ash plumes to 7.5 km on 12 Jun, and a strong ash plume to 12 km on 14 Jun. Further explosions with ash to 10–14 km contin- ued through 18 Jun.
Soufrière Hills (Montserrat)	10–15 Feb 2010	16.72° N 62.18° W 915 m	When it was first observed the SO_2 plume was divided into two parts: a higher one around 16–19 km (tropopause/stratosphere) and a lower one in the troposphere. The lower plume disappeared (non de- tectable) after one day, while the higher one spread east, south east.	Prata et al. (2007) on 2006 eruption. Carn and Prata (2010): satellite mea- surements for 1995–2009; Christopher et al. (2010) include SO_2 measurments for 1995–2009. Wadge 2010: Lava production for the pe- riod 1995–2009. Christopher et al. (2014): SO ₂ degassing, for 1995–2013, and relation with magma supply; Nicholson et al. (2013), include daily mean SO ₂ flux.	VEI 3. Large dome collapse on 11 Feb, with pyroclastic flows and an ash plume to 15 km.
Eyjafjallajökull (Iceland)	14 Apr-24 May 2010	63.63° N 19.62° W 1666 m	SO ₂ plume confined in troposphere. Up to 20 Apr the plume was below 10 km (mainly confined within 5 km); From 20 to 30 Apr plume < 5 km. Increase in both (amount and altitude) from 5 May, with maximum altitude (around 10 km) from 14 to 17 May.	Stohl et al. (2011): ash emission height from inversion with dispersion model FLEXPART. Marzano et al. (2011): radar measure- ments of plume altitude. Stevenson et al. (2012): tephra deposit. Petersen et al. (2012): tipmact in atmo- phere and plume altitude. Carboni et al. (2012): IASI SO ₂ . Boichu et al. (2013): chemistry-transport model + IASI SO ₂ to invert SO ₂ flux.	VEI 4. Explosive eruptions from the sum- mit, with ash plumes from 3–10 km (Gud- mundsson et al., 2012).
Merapi (Java, Indonesia)	4–11 Nov 2010	7.542° S 110.442° E 2968 m	High plume up to tropopause and strato- sphere. (Problems: old plume overpass- ing the volcano).	Surono et al. (2012): estimate a total of 0.44 Tg of SO ₂ , using IASI -AIRS assum- ing a plume attitude of 16 km. Saepuloh et al. (2013): Picquout et al. (2013); Pallister et al. (2013): Luehr et al. (2013); Janins et al. (2013): Innocenti et al. (2013), a); Cronin et al. (2013); Costa et al. (2013); Budi-Santoso et al. (2013)	VEI 4. Major explosive eruption, with a climactic phase that began on 3 Nov with ash plumes to 18 km. Strong explosions continued through 5–6 Nov (17 km ash plumes), declining to $<8km$ by 12 Nov.

Table 1. Continued.

Volcano name	Analysed dates	Lat/Lon/ Ele- vation	Comments on IASI SO ₂ plume	References to other work	Volcanic event description (Smithsonian weekly reports)
Etna (Sicily, Italy)	23–25 Nov 2007 + 2011 fountains	37.734° N 15.004° E 3330 m	All episodes considered are confined to the troposphere.	Aiuppa et al. (2010); combine seismic (volcanic tremor and long-period seis- micity), deformation (GPS), and geo- chemical (volcanic gas plume C0 ₂ /S0 ₂ ratios) measurements to interpret trends in the recent (2007–2008) activity of Etna volcano. Patane et al. (2013); study the 2011 Etna episodes using multi-parameter ap- proach (sesmicity, ground deformations and geochemistry, S0 ₂ flux, CO ₂ flux, CO ₂ /SO ₂ ratio). Viccaro et al. (2014); Behncke et al. (2014); Birth, growth and products of the new SE crater. Scoilo et al. (2011-2013 Etna lava fountains. Guerrieri et al. (2015): ash and SO ₂ , re- rieval and flux estimate, from SEVIRI. Spinetti et al. (2014), satellite SO ₂ valida- tion with ground measurements (includ- ing this JASI SO ₂ scheme).	
Nyamuragira/ Nyiragongo (DR Congo)	3–12 Jan 2010 and 7–15 Nov 2011	1.408° S 29.2° E 3058 m/ 1.52° S 29.25° E 3470 m	Confined to the troposphere.	Carn and Bluth (2003); Nyamuragira SO ₂ from TOMS period: 1978–2002. Bluth and Carn (2008); Exceptional sub- phur degassing from Nyamuragira vol- cano using TOMS, 1979–2005. Wadge and Burt (2011). Carn et al. (2013): monitored Nyiragongo volcano degassing with OMI, including monthly means SO ₂ for 2004–2008. Cuoco et al. (2013): impact of Nyamuragira emissions of SO ₂ using OMI and ASTER. Smets et al. (2014): multidisciplinary monitoring reveals pre- and co-eruptive signals at Nyamuragira volcano associ- ated with the Jan 2010 volcanic eruption (seismic and OMI data).	Jan 2010, VEI 1. (Nyamuragira). A flank fissure eruption began on 2 Jan 2010, and continued intermittently for 3-4 weeks. Nov 2011, VEI 2. (Nyamuragira): ma- jor flank fissure eruption began with fire-fountaining and the emplacement of a long lava flow on 6 Nov 2011. The erup- tion is thought to have ceased by 18 Nov. During both Jan 2010 and Nov 2011, the lava lake at Nyiragongo remained active (based on thermal alerts).

d.

Volcano name	Analysed dates	Lat/Lon/ Ele- vation	Comments on IASI SO_2 plume	References to other work	Volcanic event description (Smithsonian weekly reports)
Grímsvötn	21–26 May 2011	64.42° N 17.33° W 1725 m	Higher part of the plume moved west and spread north reaching the tropopause/stratosphere. Lower plume travelled towards Europe (together with ash), confined to the tro- posphere.	Flemming and Inness (2013): Assimila- tion of UV satelilite measurements for forecast. Moxnes et al. (2014): FLEXPART + IASI SO ₂ and ash. Koukouli et al. (2014): Intercomparison of SO ₂ from IASI (including C12), GOME-2 and ground data.	VEI 4. Strong explosive eruption, with an ash plume to 20 km on 21 May, declining to 10–15 km on the morning of 22 May. Minor activity on 24 May, with ash to 5–7 km.
Puyehue- Cordon- Caulle	5–30 Jun 2011	40.59° S 72.117° W 2236 m	Intermediate magnitude (between Llaima and Copahue) Reached the tropopause/stratosphere and we can follow the plume going 3 times around the Southern Hemisphere in 30 days. First part of the eruption is higher in SO ₂ amount and altitude. Intermittent low plume connected with the volcano for all the period considered.	Theys et al. (2014): detection of volcanic OCIO. Bignami et al. (2014): SAR deformation, MODIS ash and $\rm SO_2$.	VEI 5. Major sequence of explosive and effusive eruptions, that began on 4 Jun with ash plumes to 11-14 km. For the rest of Jun, lower-level eruptive activity persisted, with intermitten tash plumes to 5–8 km. The eruption continued until Apr 2012.
Nabro	12–23 Jun 2011	13.37° N 41.7° E 2218 m	The highest emission of SO ₂ for the pe- riod considered (2008–2012). Two plumes at different altitudes, the highest one reached the to stratosphere, the lower remained confined to the tropo- sphere.	Fromm et al. (2014): Nabro injected sul- phur directly to or above the tropopause upon the initial eruption on 12/13 Jun, and again on 16 Jun 2011. Bourassa et al. (2012); Sawamura et al. (2012); Vernier et al. (2013); Grarisse et al. (2014); Boindi et al. (2015).	VEI 4. The eruption began with a major ash plume, to 9–14 km on 12 Jun. The activity continue dominated by lava flow, and with ash plumes to 6–8 km from 15– 19 Jun.
Copahue	22–27 Dec 2012	37.856° S 71.183° W 2953 m	Maximum of 0.72 Tg of SO_2 present in at- mosphere. Plume connected with volcano every day. Confined to the troposphere.	Velez et al. (2011): ground deformation using InSAR; Bonali (2013): earthquake and stress study.	VEI 2. Moderate explosive eruptions, with a mixture of Strombolian fire- fountaining and steam-driven phreatic explosions, producing frequent low-level ash plumes rising 1–2 km above the cratter rim.

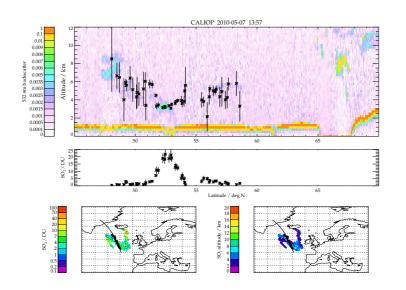


Figure 1. CALIOP/ IASI coincidences for the Eyjafjallajökull plume on 7 May 2010 for overpasses within 2 h off each other. Top plot: CALIOP backscatter profile with IASI over-plotted retrieval altitude (black stars) and error-bar (black line); middle plot: the IASI SO₂ amount and error-bars corresponding to the altitude plotted above; bottom plots: map of IASI SO₂ amount (left) and altitude (right) with CALIPSO track (black line) and identifying the IASI pixel plotted above with black stars.

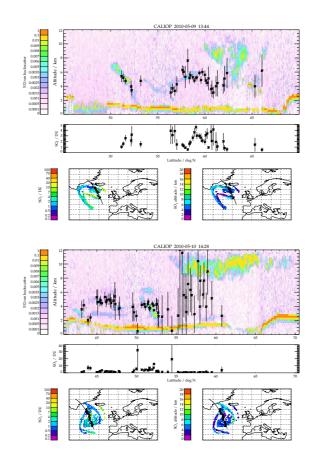


Figure 2. CALIOP-IASI coincidences for the 9 and 10 May 2010, Eyjafjallajökull plume. As Fig. 1.



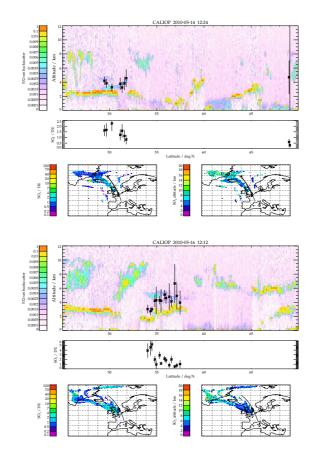


Figure 3. CALIOP-IASI coincidences for the 14 and 16 May 2010, Eyjafjallajökull plume. As Fig. 1.

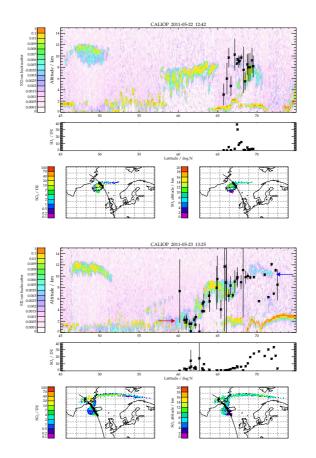


Figure 4. CALIOP/IASI coincidences for the Grímsvötn eruption on 22 and 23 May 2011. Blue arrow indicate the higer part of the plume, red one the lower part. As Fig. 1.

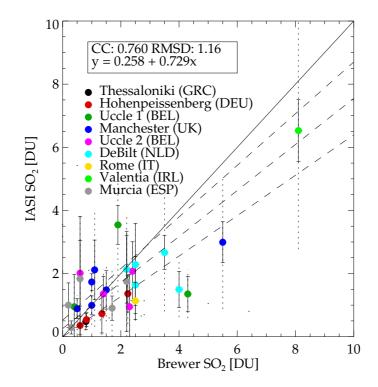


Figure 5. Scatter plot of IASI SO₂ measurements, averaged within a distance of 200 km from the ground station, vs. the daily SO₂ column amount, measured from Brewer spectrometers. Different colours correspond to different Brewer ground stations. Black error-bars are the IASI average errors; dotted error-bars are the standard deviation of the IASI data within the selected distance. Black lines represent the ideal line y = x; dotted lines are the best fits with error in the best fit. The legend box shows the correlation coefficient (CC), root mean square difference (RMSD) and the best fit line.

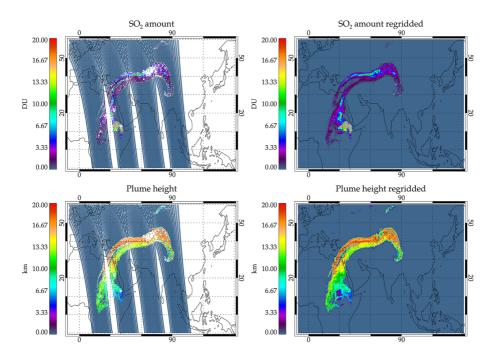


Figure 6. Maps of IASI SO₂ amount (top left) and height (bottom left) and the equivalent maps of SO₂ amount (top right) and height (bottom right) obtained after regridding. Grey colour indicate values higher than the colour bar. The IASI data are a combination of four orbits on the 15 June 2011 from 13:00 to 18:00 UTC during Nabro eruption.

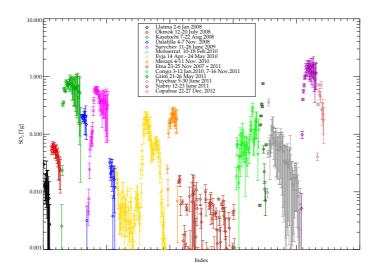


Figure 7. SO₂ mass present in the atmosphere as retrieved from IASI data. The values are the measured amount every half day and vary with volcanic emission, gas removal and satellite sampling. Points are separated by ~ 12 h. Data are presented in temporal order along the ordinate (*x* axis) but eruptions are plotted consecutively one after the other without a gap between them. The total SO₂ amount reported here is computed using the geographic area associated with the eruption. For eruptions which overlap in time (e.g Grimsvötn, Puyehue, and Nabro in May and June 2011) the SO₂ loads within each respective area are considered and plotted separately.

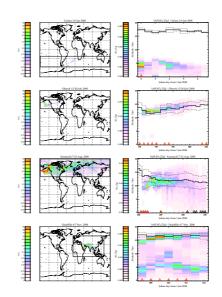


Figure 8. The left column shows maps of the maximum SO_2 amount retrieved within the considered area (black rectangle). The right column shows the SO_2 vertical distribution for the considered volcanic eruption. In each plot the *y* axes are the vertical levels in km. The colour represents the total mass of SO_2 in Tg, dark-red represents values higher than the colour-bar. Every column of the plots come from an IASI map (one every 12 h). The black lines are the mean, and the mean plus or minus the standard deviations, of tropopause computed from ECMWF profiles at the location of plume pixels. Red triangles in the bottom line indicate the presence of a fresh plume connected with the volcano, black triangles indicate the presence of an old plume overpassing the volcano (this may eventually mask a newer plume). Note that the plots for different eruptions have different colour-scales and cover different time ranges.

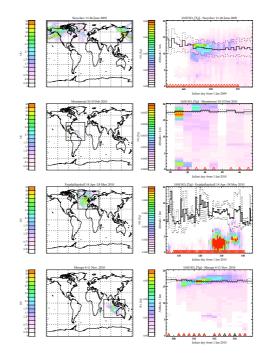


Figure 9. As for Fig. 8.

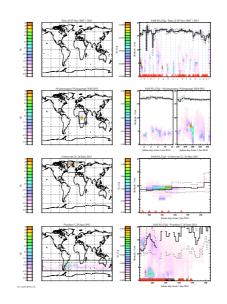


Figure 10. As for Fig. 8. The Etna plots show different eruptive episodes corresponding to x axes labels: (a) 23–25 November 2007 eruption; and the 2011 lava-fountains: (b) 11–13 January, (c) 17–19 February, (d) 10–12 April, (e) 11–13 May, (f) 8–10 July, (g) 18–20 July, (h) 24–26 July, (i) 29 July–1 August, (j) 4–7 August, (k) 11–13 August, (l) 19–21 August, (m) 28–30 August, (n) 7–10 September, (o) 18–20 September, (p) 28–30 September, (q) 11–14 October, (r) 12–17 November.

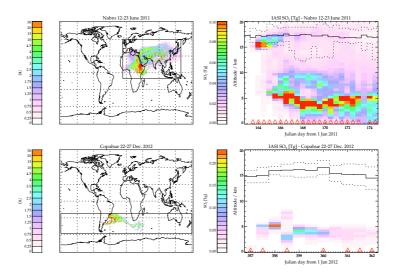


Figure 11. As for Fig. 8.

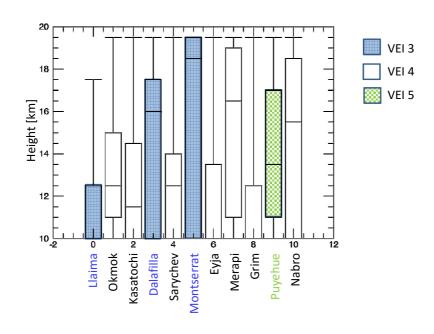


Figure 12. Eruption SO_2 -plume altitudes from this study. The black lines show the range of altitudes retrieved, the boxes depict the altitude range into which the 50% of the SO_2 mass falls, the horizontal lines within each box are the altitudes of the maximum of SO_2 mass. Different colors represent eruptions of different VEI (Source: Global Volcanism Program, 2013).