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Satellite-based detection of volcanic sulphur dioxide from recent eruptions in Central and South America

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Abstract. Volcanic eruptions can emit large amounts of rock fragments and fine particles (ash) into the atmosphere, as well as several gases, including sulphur dioxide (SO₂). These ejecta and emissions are a major natural hazard, not only to the local population, but also to the infrastructure in the vicinity of volcanoes and to aviation. Here, we describe a methodology to retrieve quantitative information about volcanic SO₂ plumes from satellite-borne measurements in the UV/Visible spectral range. The combination of a satellite-based SO₂ detection scheme and a state-of-the-art 3D trajectory model enables us to confirm the volcanic origin of trace gas signals and to estimate the plume height and the effective emission height. This is demonstrated by case-studies for four selected volcanic eruptions in South and Central America, using the GOME, SCIAMACHY and GOME-2 instruments.

1 Introduction

Volcanic eruptions and outgassing are the most important natural sources of SO₂ (Graf et al., 1997). Most volcanic SO₂ emissions and aerosols remain in the troposphere. There, the lifetime of SO₂ depends strongly on meteorological conditions but it is typically in the order of a few days. In the troposphere, the major sink of SO₂ is its transformation into sulphate aerosol and sulphuric acid, the latter being responsible for the “acid rain” phenomenon. Since SO₂ is also toxic it has to be considered a possible hazard to human health (e.g. EPA, 2007).

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Severe volcanic eruptions, such as those of El Chichon (1982) and Pinatubo (1991), can bring a considerable amount of sulphur dioxide and particles into the stratosphere. Here, the SO₂ is mainly transformed into sulphuric droplets (aerosols). These aerosol droplets may persist for weeks in the stratosphere and may be transported on intercontinental scales (Read et al., 1993). Sulphate aerosol has a cooling effect on the atmosphere and leads to enhanced rates of ozone loss in the lower stratosphere, as it was observed after the Pinatubo eruption (Self et al., 1996).

Satellite remote sensing of volcanic SO₂ provides a unique global and systematic way of detecting and tracking volcanic gas plumes and their evolution in near-real time, thus delivering additional information to authorities to take appropriate actions. The new generation of atmospheric remote sensing sensors allows the accurate detection of volcanic SO₂ plumes and ash clouds. Especially the latter are also a hazard to aviation: the ash can limit the view of aircraft pilots and it can melt inside the engines, as a result of which the engines may fail. Since SO₂ is also dissolved in water droplets and forms corrosive sulphuric acid, it can create sulphate deposits in the engines (van Geffen et al., 2007).

The main focus of this paper is the retrieval of SO₂ using satellite measurements in the UV spectral range from the Global Ozone Monitoring Experiment (GOME) (Burrows et al., 1999) on-board ERS-2 and the Scanning Imaging Absorption Spectrometer for Atmospheric CHartography (SCIAMACHY) (Bovensmann et al., 1998; Gottwald et al., 2006) on-board ENVISAT. We focus on major eruptions of four volcanoes in South and Central America during the last decade. Finally, the perspectives for global SO₂ monitoring using GOME-2 data on-board MetOp and the issues related to the near-real time retrieval and delivery of SO₂ data are addressed.

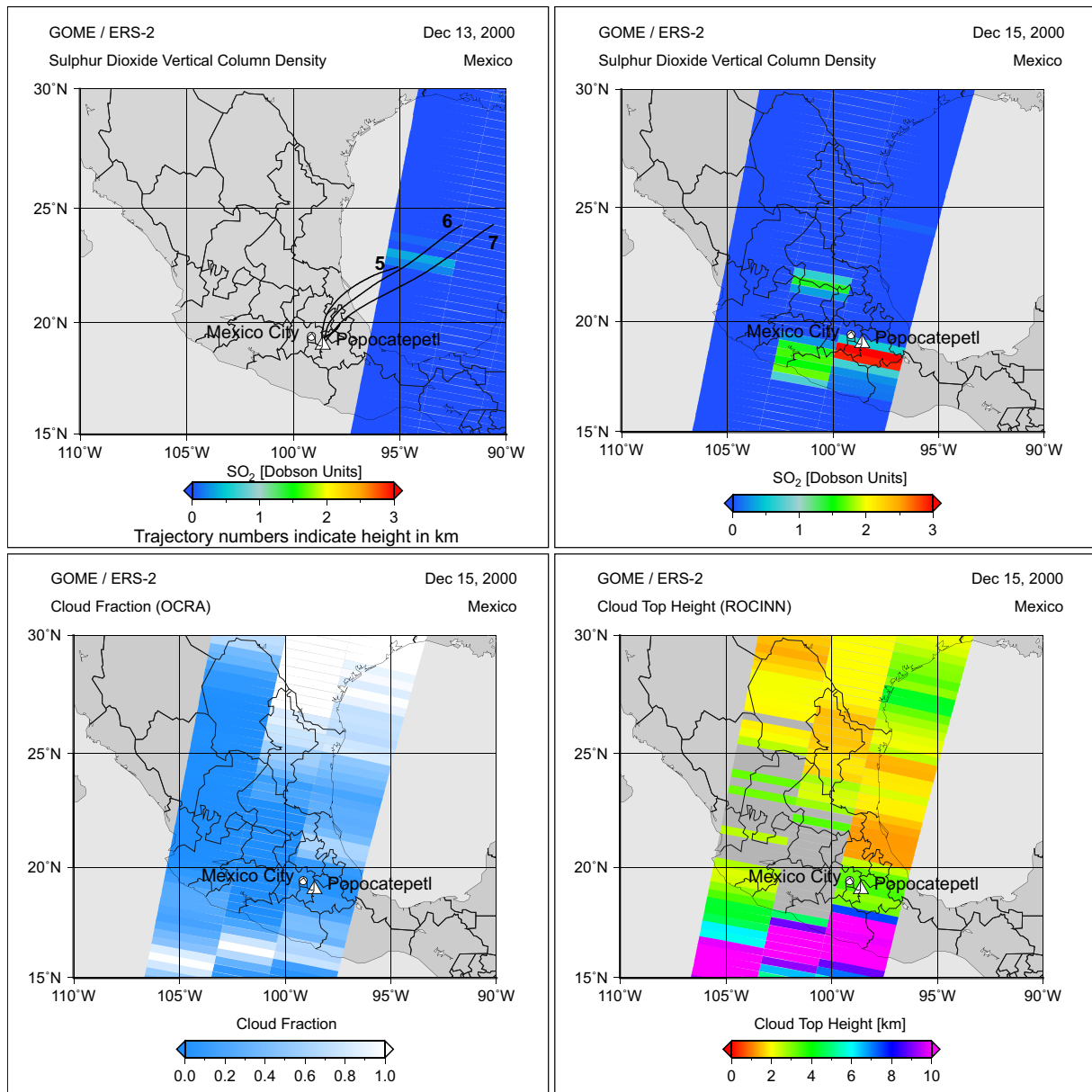


Fig. 1. SO_2 vertical column densities as retrieved from GOME for 13 December 2000 (top-left). Representing a coherent ensemble of matching forward trajectories, three trajectories are plotted released at the Popocatepetl at altitudes from 5 to 7 km and durations of 24 h. The other panels show SO_2 vertical column densities, cloud fraction and cloud top height on 15 December 2000.

2 SO_2 column retrieval

The SO_2 column along slant absorption paths is retrieved from the UV backscatter spectra by GOME and SCIAMACHY, applying the well-established Differential Optical Absorption Spectroscopy (DOAS) approach (Platt, 1994). The non-linear least square fit is performed in the 315–326 nm wavelength range and provides the SO_2 slant column (Eisinger et al., 1998). Absorption cross-sections of sulphur dioxide, ozone, nitrogen dioxide and a correction for the Ring effect (rotational Raman scattering) are fitted

simultaneously. Furthermore, a SO_2 background correction is applied to account for systematic artefacts introduced by the interference between ozone and SO_2 in the given spectral window.

In the next step, the SO_2 slant columns are converted to geometrically independent vertical column amounts through division mass factors (AMF). The AMFs are calculated with the radiative transfer model LIDORT (Linearized Discrete Ordinate Radiative Transfer model) (Spurr et al., 2001) and depend on the viewing geometry, the reflection properties of the underlying surface, the SO_2 plume height, and the pres-

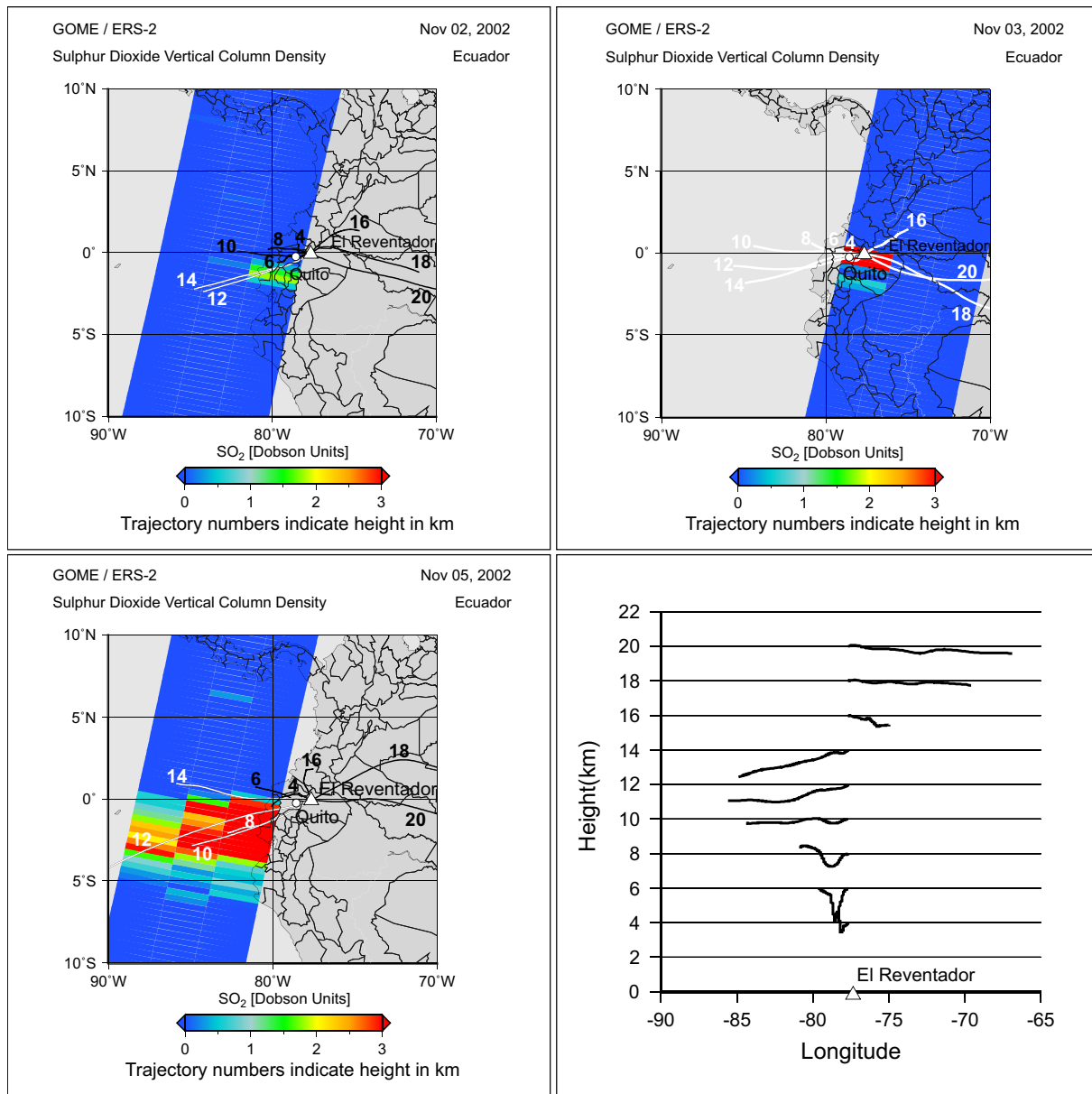


Fig. 2. SO₂ vertical column densities of the eruption of El Reventador, Ecuador for 2, 3 and 5 November 2002, as measured by GOME. Trajectories were released from 4 km to 20 km in steps of 2 km using durations of 24 h. The trajectories matching the plume are coloured in white. Bottom-right: Longitudinal crosssection of the forward trajectories released for the El Reventador eruption at 3 November 2002. Trajectories were released at 06:00 UTC from 4 to 20 km above sea level in steps of 2 km using durations of 24 h. The vertical wind shear between 14 km and 16 km can be well recognized.

ence and optical properties of clouds and aerosols (Thomas et al., 2005).

3 Trajectory analysis

In order to attribute increased SO₂ values as found in satellite-based observations to a particular volcano or an active volcanic region, we apply the trajectory matching tech-

nique (Thomas et al., 2005). By means of calculating ensembles of backward trajectories starting from enhanced SO₂ column observations, the volcanic origin of such SO₂ plumes can be identified or rejected. In the latter case the emission can be attributed e.g., to anthropogenic processes or other natural sources. The quantitative retrieval of SO₂ column densities strongly depends on the SO₂ plume height, i.e. the sulphur dioxide profile. We determine the unknown plume height and the emission height of SO₂ at a volcano using the

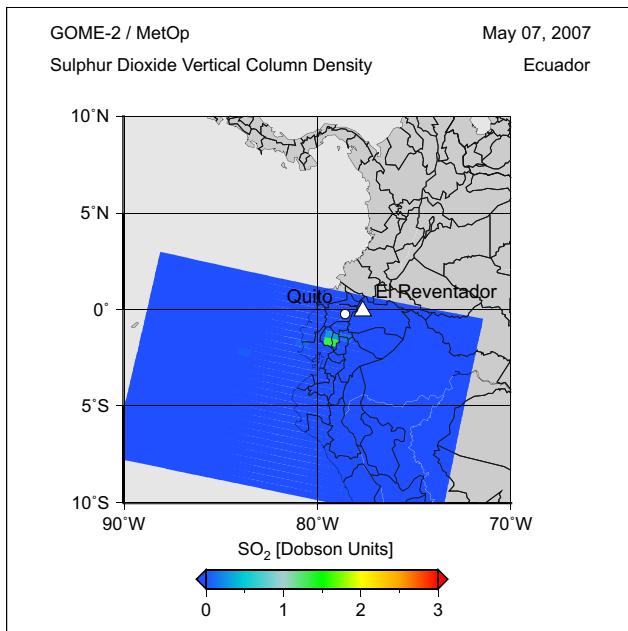


Fig. 3. SO₂ vertical column densities derived from GOME-2 for the eruption of El Reventador/Ecuador for 7 May 2007. Backward trajectories calculated for this case indicate an emission height of 6 km.

3-D trajectory model FLEXTRA (Stohl et al., 1999). This model is driven by meteorological analyses of the European Centre for Medium Range Weather Forecast (ECMWF). Ensembles of backward trajectories were released at different pressure levels from 2 km to 20 km at the geolocation of satellite-based observations with elevated SO₂ columns. This allows confirming the volcanic origin of the SO₂, determining the effective emission height over the volcano and revealing the spatial and temporal evolution of the SO₂ plume. In order to reconfirm an estimated volcanic source, ensembles of 3-D forward trajectories were released at volcanoes (see Sect. 4) at different levels ranging from 2 km to 20 km above mean sea level (AMSL). By matching the parcel trajectories with the first guess satellite-based SO₂ retrievals the height of the SO₂ plume is determined. The accuracy of the combined retrieval and trajectory matching approach was evaluated in Thomas et al. (2005) using ground-based measurements of SO₂ and aerosols.

4 Case studies

This section presents the SO₂ and trajectory analysis for selected eruptions of four volcanoes in South and Central America. We used observations from the GOME and SCIAMACHY sensors, as well as data from the recently launched GOME-2 instrument. Important differences between the instruments are the spatial resolution and the temporal coverage. The spatial resolution for GOME is 320×40 km², pro-

viding global coverage within 3 days, SCIAMACHY has a spatial resolution of 60×30 km² and global coverage is reached within 6 days only. The new GOME-2 instrument offers nearly global coverage within one day at a spatial resolution of about 80×40 km².

4.1 Popocatepetl, 12–15 December 2000

Popocatepetl (19°1' N, 98°37' W), located 70 km south-east from Mexico City, erupted from 12–15 December 2000. Popocatepetl is one of the most active volcanoes in Mexico. The SO₂ vertical column densities as retrieved from GOME for 13 December 2000 are shown in Fig. 1. The enhanced SO₂ values over the Gulf of Mexico were the first signal detected for this eruptive period and hence they were used to determine the origin of the SO₂ and the effective emission height. The trajectory analysis revealed an effective emission height at the Popocatepetl between 5 km and 7 km, which is around its summit height of 5462 m. The three trajectories overlaid in Fig. 1 indicate the main path of the SO₂ plume and are a subset of a coherent ensemble of matching forward trajectories (not shown here). They were released at 00:00 UTC at the geolocation of Popocatepetl and represent a travel time of 24 h at altitudes from 5 km and 7 km. The retrieved SO₂ vertical column densities, the cloud fraction and the cloud top height for 15 December 2000 are also depicted in Fig. 1. The low cloud fraction above the volcano provided good observation conditions. On this day, maximum SO₂ column densities of ~7 Dobson Units were detected with GOME southwards from the volcano (1 DU=2.69×10¹⁶ molec/cm²).

4.2 El Reventador, 2–5 November 2002 and 7 May 2007

El Reventador (0°4' S, 77°39' W) is located 90 km north-east from Quito/Ecuador and erupted from 2–5 November 2002. Due to a strong wind shear between 14 km and 16 km altitude, the plume split in two, one part moving east and reaching 20 km height, and the other part moving south-west and reaching 14 km height. The GOME-based SO₂ vertical column densities of this eruptive period are presented in Fig. 2 for 2, 3 and 5 November 2002. The trajectory analysis revealed an effective emission height at the Reventador between 4 km and 14 km in south-westerly direction and between 14 km to 20 km in easterly direction. This is corroborated by airplane observations (VAAC Washington, 2002). The trajectories depicted in Fig. 2 confirm the observed transport of the SO₂ plume. They were released at 15:00 UTC at the Reventador (06:00 UTC for 3 November 2002) and represent a travel time of 24 h at altitudes ranging from 4 km to 20 km. Trajectories matching the plume are coloured in white while others are black. For 2 November 2002 the matching trajectories can be associated with heights between 12 km and 14 km, respectively. At 3 November 2002 all released trajectories cover the SO₂ plume and at 5 Novem-

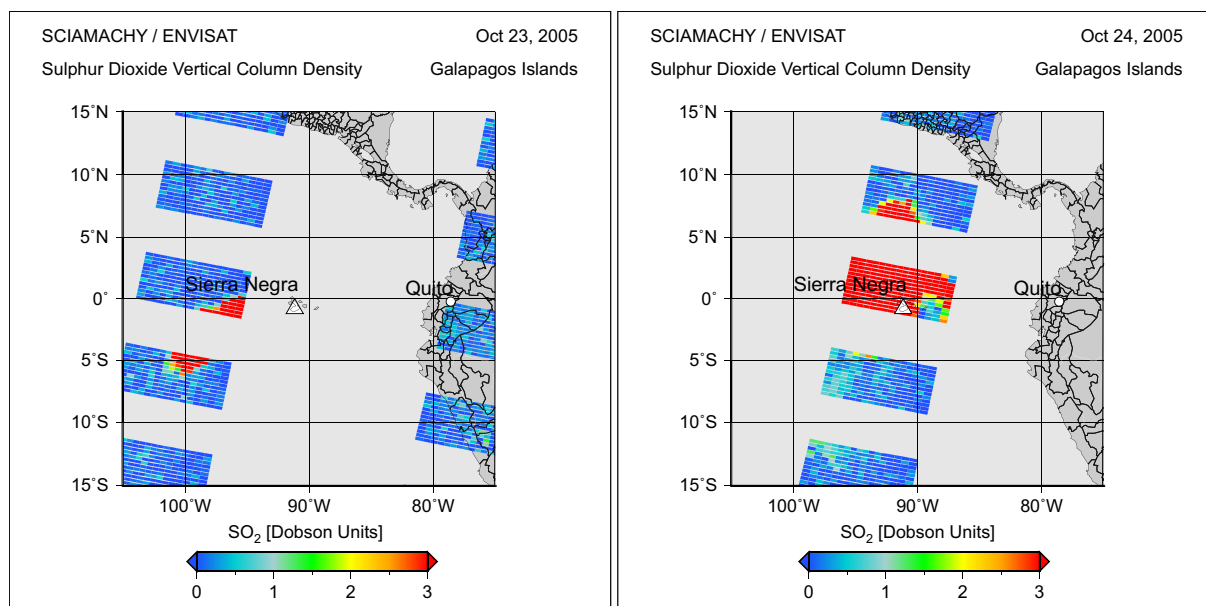


Fig. 4. SO₂ vertical column densities retrieved from SCIAMACHY for the eruption of Sierra Negra on the Galapagos Island for 23 and 24 October 2005.

ber 2002 the trajectories from levels between 8 km to 14 km match the SO₂ plume. Maximum SO₂ columns of ~10 DU were measured with GOME on 3 November 2002, close to the volcano.

More recent eruptions of the El Reventador took place during March and May 2007 and were observed by GOME-2. Backward trajectories calculated for the eruption on 7 May 2007 revealed an emission height of 6 km. Figure 3 shows the SO₂ vertical columns derived from preliminary GOME-2 data for 7 May 2007. A relatively small SO₂ amount of ~1.7 DU was measured for the plume south-westerly of the volcano. Figure 3 shows also the smaller GOME-2 ground-pixels and the larger swath-width of the GOME-2 instrument compared to GOME.

4.3 Sierra Negra, 23–24 October 2005

The Sierra Negra volcano (0°49' S, 91°10' W), located on the Isla Isabela of the Galapagos archipelago (Ecuador), erupted on 23 October 2005. The eruption produced a large amount of volcanic smog (vog), which is typically a mixture of SO₂, oxygen, water, and aerosols. The vog spread over several hundred square-kilometres of the Pacific Ocean. Figure 4 shows the SO₂ vertical column densities for 23 and 24 October 2005 retrieved from SCIAMACHY observations. Backward trajectories calculated for 23 October 2005 indicate a plume height of 12 km. While the moving direction of the plume on 23 October 2007 was to west-southwest direction, we could not find a preferred direction for the 24 October 2007. We therefore found elevated SO₂ values in nearly the whole region around Sierra Negra. On 24 October 2007, very

large SO₂ column densities of more than 60 Dobson Units were detected with SCIAMACHY, close to the volcano.

5 Conclusions

We have shown the unique possibilities of satellite remote sensing in detecting and tracking volcanic SO₂ plumes. The GOME (ERS-2), SCIAMACHY (ENVISAT) and GOME-2 (MetOp) satellite instruments allow retrieving atmospheric SO₂ column densities on a global scale and in near-real time. By combining the satellite-based SO₂ retrieval with state-of-art 3-D trajectory models, the volcanic origin can be confirmed, and the plume and effective emission heights can be estimated. The maximum SO₂ column densities of the four eruptions analysed in this study, range from ~1.7 Dobson Units for the El Reventador eruption in May 2007, to more than 60 DU for the Sierra Negra eruption in October 2005.

Volcanic eruptions are a major hazard to aviation, and the only safe solution for aircrafts is to avoid flying through such volcanic clouds. The Volcanic Ash Advisory Centres (VAACs) have been set up by the International Civil Aviation Organisation (ICAO) to gather information on volcanic activity and to issue alerts on the possible danger of volcanic eruptions to aviation. To assist the VAACs in their task, the Support to Aviation Control Service (SACS) of the PROMOTE project intends to deliver in near-real-time SO₂ data and, in the event of high SO₂ concentrations, notifications by email to the VAACs and other interested parties (van Geffen et al., 2007). The SO₂ data of SACS can be accessed

via <http://sacs.aeronomie.be/> or via the PROMOTE website <http://www.gse-promote.org/> under “Special Services”.

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