

A Sulfur Dioxide Covariance-Based Retrieval Algorithm (COBRA): application to TROPOMI reveals new emission sources

Nicolas Theys¹, Vitali Fioletov², Can Li^{3,4}, Isabelle De Smedt¹, Christophe Lerot¹, Chris McLinden², Nickolay Krotkov³, Debora Griffin², Lieven Clarisse⁵, Pascal Hedelt⁶, Diego Loyola⁶, Thomas Wagner⁷, Vinod Kumar⁷, Antje Innes⁸, Roberto Ribas⁸, François Hendrick¹, Jonas Vlietinck¹, Hugues Brenot¹, Michel Van Roozendael¹

¹ Royal Belgian Institute for Space Aeronomy (BIRA-IASB), Brussels, Belgium.

² Air Quality Research Division, Environment and Climate Change Canada, Toronto, Canada.

³ Atmospheric Chemistry and Dynamics Laboratory, NASA Goddard Space Flight Center, Greenbelt, MD, USA.

⁴ Earth System Science Interdisciplinary Center, University of Maryland, College Park, MD, USA.

⁵ Université libre de Bruxelles (ULB), Spectroscopy, Quantum Chemistry and Atmospheric Remote Sensing (SQUARES), C. P. 160/09, Brussels, Belgium.

⁶ Institut für Methodik der Fernerkundung (IMF), Deutsches Zentrum für Luft und Raumfahrt (DLR), Oberpfaffenhofen, Germany.

⁷ Max Planck Institute for Chemistry (MPIC), Hahn-Meitner-Weg 1, 55128 Mainz, Germany.

⁸ European Centre for Medium-Range Weather Forecast (ECMWF), Shinfield Park, Reading, RG2 9AX, UK.

Correspondence to: N. Theys (theys@aeronomie.be)

ABSTRACT

Sensitive and accurate detection of sulfur dioxide (SO_2) from space is important for monitoring and estimating global sulfur emissions. Inspired by detection methods applied in the thermal infrared, we present here a new scheme to retrieve SO_2 columns from satellite observations of ultraviolet back-scattered radiances. The retrieval is based on a measurement error covariance matrix to fully represent the SO_2 -free radiance variability, so that the SO_2 slant column density is the only retrieved parameter of the algorithm. We demonstrate this approach, named COBRA, on measurements from the TROPospheric Monitoring Instrument (TROPOMI) aboard the Sentinel-5 Precursor (S-5P) satellite. We show that the method reduces significantly both the noise and biases present in the current TROPOMI operational DOAS SO_2 retrievals. The performance of this technique is also benchmarked against that of the Principal Component Algorithm (PCA) approach. We find that the quality of the data is similar and even slightly better with the proposed COBRA approach. The ability of the algorithm to retrieve SO_2 accurately is also further supported by comparison with ground-based observations. We illustrate the great sensitivity of the method with a high-resolution global SO_2 map, considering two and a half years of TROPOMI data. In addition to the known sources, we detect many new SO_2 emission hotspots worldwide. For the largest sources, we use the COBRA data to estimate SO_2 emission rates. Results are comparable to other recently published TROPOMI-based SO_2 emissions estimates, but the associated uncertainties are significantly lower than with the operational data. Next, for a limited number of weak sources, we demonstrate the potential of our data for quantifying SO_2 emissions with a detection limit of about 8 kt yr^{-1} , a factor of 4 better than the emissions derived from the Ozone Monitoring Instrument (OMI). We anticipate that the systematic use of our TROPOMI COBRA SO_2 column data set at a global scale will allow identifying and quantifying missing sources, and help improving SO_2 emission inventories.

1. INTRODUCTION

Sulfur dioxide (SO_2) in the atmosphere rapidly oxidizes into sulfuric acid and sulfate aerosols, which have environmental effects ranging from local and long-range air pollution to global climate impact. SO_2 is released into the atmosphere from anthropogenic activities, due to fossil fuel

burning (coal, oil and gas) and smelting, and from natural sources, mainly volcanoes. Satellites provide a viable means to monitor global SO₂ emissions and assess their environmental impacts. Since the late seventies, SO₂ vertical column densities (VCD) are provided by several ultraviolet (UV) polar-orbiting nadir instruments, namely the Total Ozone Monitoring Spectrometer (TOMS; Krueger, 1983), Global Ozone Monitoring Experiment (GOME; Eisinger and Burrows, 1998; Khokhar et al., 2005), SCanning Imaging Absorption spectroMeter for Atmospheric CHartographY (SCIAMACHY; Afe et al., 2004), Ozone Monitoring Instrument (OMI; Krotkov et al., 2006; Yang et al., 2007, 2010; Li et al., 2013; Theys et al., 2015), Global Ozone Monitoring Experiment-2 (GOME-2; Nowlan et al., 2011; Rix et al., 2012; Hörmann et al., 2013), Ozone Mapping and Profiler Suite (OMPS; Yang et al., 2013, Zhang et al., 2017) and TROPospheric Monitoring Instrument (TROPOMI; Theys et al., 2017). From the various datasets, a remarkable trend emerges in the ability of successive sensors to detect weaker and more localized emissions. This is in part due to the better spatial resolution and signal-to-noise of the modern UV spectrometers (see e.g. Fioletov et al., 2013; Theys et al., 2019), but also from advances in retrieval techniques. In particular, the Principal Component Algorithm (PCA) applied to OMI (Li et al., 2013, 2020a) and OMPS (Zhang et al., 2017) proved to be a very efficient method to reduce retrieval noise and biases and thus to increase the sensitivity of the retrievals to weak SO₂ emissions to 30-40 kt yr⁻¹. This enabled major improvements in bottom-up emissions inventories (Liu et al., 2018) and detection of missing SO₂ emission sources (Fioletov et al., 2016; McLinden et al., 2016).

TROPOMI, launched in October 2017 onboard the ESA and Copernicus Sentinel-5 Precursor (S-5P) platform, is an atmospheric mission with a dedicated focus on the tropospheric composition (Veefkind et al., 2012). With a spatial resolution as good as 3.5 x 5.5 km² per ground pixel (3.5 x 7 km² before August 2019), it is specifically designed to monitor atmospheric constituents from urban to global scales. The first observations of SO₂ by TROPOMI were focusing on relatively large volcanic sources and indeed revealed the great potential of the instrument to inform about global volcanic SO₂ degassing with high resolution and unprecedented sensitivity (Theys et al., 2019; Queiße et al. 2019). However, further investigation of anthropogenic and volcanic SO₂ sources using TROPOMI revealed problems with the current TROPOMI SO₂ retrievals for weak emission sources (Fioletov et al., 2020). In brief, large-scale and variable VCD biases on the order

of 0.25 Dobson Unit (DU; $1 \text{ DU} = 2.69 \times 10^{16} \text{ molecules} \times \text{cm}^{-2}$) are present in the data, which limits their use to medium to large SO_2 sources only.

The operational TROPOMI SO_2 algorithm is based on the Differential Optical Absorption Spectroscopy technique (DOAS; Platt and Stutz, 2008), and essentially works in three steps (details are given in Theys et al., 2017): a spectral analysis yielding SO_2 slant column densities (SCD), an empirical background correction of the SCDs and a radiative transfer calculation of air mass factors (AMF) to convert the corrected SCD into the VCD output ($\text{VCD} = \text{SCD}_{\text{cor}} / \text{AMF}$). As a matter of fact, the SO_2 SCD retrieval is subject to spectral misfits which can lead to systematic offsets. These SCD errors are difficult to correct and arise from imperfect DOAS forward modeling. Here, we propose an alternative spectral fitting approach, named COBRA, which strongly reduces the SO_2 SCD biases for the weak SO_2 columns and suppresses the need for the post-processing background correction. COBRA is akin to the PCA approach, which constitutes the basis of the OMI and OMPS SO_2 operational retrievals (Li et al., 2020b, 2020c). As demonstrated below, COBRA significantly improves the quality as compared with the current TROPOMI DOAS operational SO_2 product. The analysis of two and a half years of data oversampled at high resolution reveals many new SO_2 emission sources globally, highlighting the great performance of COBRA in terms of SO_2 detection.

The paper is structured as follows. Section 2 describes the algorithm and its practical implementation. In section 3, SO_2 retrievals from COBRA are evaluated against other satellite data sets, model results and ground-based observations. Section 4 presents long-term averaged global results. In Section 5, we apply an emission inversion scheme to the COBRA SO_2 data set and compare with previously estimated SO_2 emissions from the TROPOMI operational product. New SO_2 emission sources detected by the COBRA are discussed. Conclusions are given in Section 6.

2. METHODOLOGY

2.1 TROPOMI

In this study, we use observations from the TROPOMI instrument on the Sentinel-5 Precursor satellite. TROPOMI is a hyperspectral nadir sensor measuring solar radiation backscattered by the atmosphere and reflected by the Earth, in the ultraviolet, visible, near-infrared and shortwave infrared wavelength regions. TROPOMI delivers column amounts of minor atmospheric constituents, such as O_3 , NO_2 , SO_2 , HCHO , CO , CH_4 , as well as aerosol and cloud information

(Veefkind et al., 2012). The S-5P satellite is a polar orbiting platform crossing the equator at 13:30h local time. A nearly global coverage is achieved in one day owing to a 2600 km wide swath. The footprint on the ground of the satellite measurement depends mainly on the across-track position in the swath and on the spectral band. For SO₂, the ultraviolet spectral band 3 is used, and the swath is divided into 450 across-track positions (also referred to as ‘rows’). The spatial resolution for the center of the swath is approximately 3.5 x 7 km² (across-track x along-track) until 6 August 2019 when the sampling improved to 3.5 x 5.5 km².

For this work, we analyze data measured between 1 April 2018 and 31 December 2020, and solar zenith angles (SZA) less than 60°.

2.2 Algorithm description

As mentioned above, the operational TROPOMI SO₂ algorithm is based on the DOAS technique, the most widely used method to derive atmospheric trace gas constituents in the UV-visible spectral range. The inverse problem can be expressed (employing the notation of Rodgers, 2000):

$$y = K \cdot x + \epsilon \quad (1)$$

where $y = -\log(I/I_o)$ is the optical depth, i.e. the logarithmic ratio of the wavelength calibrated measured intensity (I) and the reference intensity spectrum (I_o) over a given wavelength range, x is the state vector including SCDs of relevant trace gases and closure fit parameters (e.g. for broadband effects), K is the forward model matrix with absorption cross-sections and other spectra, and ϵ is the measurement noise. The solution can be approximated by least-square fitting:

$$\hat{x} = (K^T S_\epsilon^{-1} K)^{-1} K^T S_\epsilon^{-1} y \quad (2)$$

where S_ϵ is the measurement error covariance matrix. The latter matrix is most often taken diagonal (no error correlations) or proportional to unity (unweighted least-square). Eqs. (1) and (2) describe the simplest DOAS approach and are given here for illustration purpose only. In practice, the DOAS problem is fundamentally non-linear in many aspects and DOAS software packages, such as QDOAS (Danckaert et al., 2017), support different non-linear retrieval options (e.g. for wavelength shift and squeeze, or intensity offset), with the aim to improve the quality of the retrievals.

For weakly absorbing tropospheric species, retrieval artefacts are frequent with DOAS (notably for satellite nadir geometry), and are attributed to spectral interferences, imperfect forward model and incomplete treatment of instrumental effects (e.g., polarization sensitivity). For UV nadir SO₂

1 retrievals in particular, biases in the data arise mainly from strong ozone absorption and imperfect
2 treatment of the non-elastic Rotational Raman Scattering (Ring) effect. It is generally difficult to
3 completely remove these offsets even after applying post-processing background corrections
4 (Theys et al., 2017; Fioletov et al., 2020).

5 The Covariance-Based Retrieval Algorithm (COBRA) presented here, and illustrated for
6 TROPOMI measurements, aims at correcting most of the artefacts in the DOAS SO₂ SCDs by
7 optimally retrieving a single parameter: the SO₂ SCD.

8 First introduced by von Clarmann et al. (2001), the retrieval approach was developed by Walker
9 et al. (2011) for nadir observations of SO₂ and NH₃ from the Infrared Atmospheric Sounding
10 Interferometer (IASI). Then, the technique, also known as Hyperspectral Range Index (HRI), has
11 been further refined and successfully applied to other trace gases and aerosols (e.g., Van Damme
12 et al., 2014; Franco et al., 2018; Clarisse et al., 2019). The method proved to be very sensitive and
13 led to superior data quality both in terms of precision and accuracy. Surprisingly, this technique
14 has, to our knowledge, never been applied in the UV-visible spectral range.

15 Starting from Eq. 1, we assume the measurement vector can be linearized around a background
16 SO₂-free spectrum $\bar{y}$:

$$y = \bar{y} + k.SCD + \epsilon_{bg} + \epsilon \quad (3)$$

18 with ϵ_{bg} being the [deviation of the SO₂-free component of the spectrum relative to the mean](#)
19 [spectrum](#) $\bar{y}$, and ϵ is the measurement noise. The SO₂ contribution to the measured spectral optical
20 depth is approximated by the product of the instrument slit convolved absorption cross-section
21 vector k (expressed in cm²/molecule) and the SO₂ slant column density SCD (in molecules/cm²).
22 Here, we use as input of the retrieval the same SO₂ absorption cross-section data (Bogumil et al.,
23 2003) [and the same approach for the wavelength calibration of the spectra](#), as for the operational
24 TROPOMI SO₂ retrievals (see Theys et al., 2017). The only difference is the wavelength interval
25 of 310.5 – 326 nm (see discussion below).

26 The basic principle of the method is to consider all contributions to the difference $(y - \bar{y})$ other
27 than SO₂ as an error term $(\epsilon_{bg} + \epsilon)$ with a Gaussian distribution. If one can define an ensemble Y
28 of N measured [SO₂-free](#) spectra, representative of the total $(\epsilon_{bg} + \epsilon)$ variability, and characterized
29 by a mean measurement vector $\bar{y}$ and a covariance matrix S :

$$S = \frac{1}{N-1} \cdot \sum_{i=1}^N (y_i - \bar{y})(y_i - \bar{y})^T \quad (4)$$

then the solution of the problem writes as:

$$\widehat{SCD} = \overline{SCD} + (k^T S^{-1} k)^{-1} k^T S^{-1} (y - \bar{y}) \quad (5)$$

where $\overline{SCD}$ is the mean SO₂ SCD of the ensemble ($\overline{SCD} = 0$ by definition). The error on the retrieved SCD is given by [the square root of the error covariance of the solution \(Rodgers, 2000\)](#):

$$\widehat{SCD}_{err} = \sqrt{(k^T S^{-1} k)^{-1}} \quad (6)$$

Fundamentally, COBRA generalizes the measurement error covariance matrix of Eq. 2 by incorporating geophysical background spectral variability (including all cross-correlations), variability from the atmosphere or induced by instrumental changes.

For spectra where no enhancements of SO₂ can be detected, the linearization (Eq. 3) simplifies to

$$y - \bar{y} = \epsilon_{bg} + \epsilon \quad (7)$$

Both sides of the equation have therefore the same probability distribution, and it follows that the covariance matrix associated with $\epsilon_{bg} + \epsilon$ can readily be constructed by applying Eq. 4 on a representative set of SO₂-free spectra. The key is to define the ensemble Y such that $y - \bar{y}$ cancels much of the systematic components of ϵ_{bg} .

A remarkable feature of COBRA is its simplicity. The SO₂ SCD retrieval in Eq. 5 reduces to a simple dot product between the $y - \bar{y}$ residue and $k^T S^{-1}$ (skipping the normalization factor $(k^T S^{-1} k)^{-1}$). The vector $k^T S^{-1}$ essentially contains the weights of each wavelength to the retrieved target column amount; the strength of the method relies in the fact that these weights are optimally determined by the measurements themselves. This is in contrast to the DOAS approach which mostly considers all wavelengths equal. Furthermore, DOAS also allows for cross-talks between the state vector elements, which can lead to an increase of the SCD data scatter (in particular for weak absorbers). This is obviously not the case for COBRA, as only a single parameter is retrieved, the SO₂ slant column. COBRA has other great advantages that we briefly outline here:

- [Except for the wavelength calibration step](#), the algorithm does not need a reference spectrum (I_o). Indeed, equations 4 and 5 involve differences of logarithmic intensity ratios

and thus I_o cancels out. Following the same logic, any constant spectral feature multiplicative to the radiance and shared by the ensemble Y will have no influence on the retrieved SCDs.

- The analysis of individual spectra with COBRA does not require the fit of a wavelength shift and squeeze, a common (and often time-consuming) practice in DOAS. Beirle et al (2013) have shown that the effects of spectral shift and squeeze can be linearized and represented by pseudo-absorbers. Therefore, their contributions (and variability) to the optical depth are accounted for by the covariance matrix.
- The COBRA results display low noise. This is a direct result of the COBRA approach in that the wavelengths with the largest background radiance variability will have the lowest weights on the retrieved SCD (Eq. 5).
- Very small biases are observed in the COBRA data (see next section). As a consequence, an empirical SCD background correction is not needed.
- The approach works in principle for any wavelength range. This allows flexibility in case of lower instrumental performance for certain wavelength regions.
- The covariance matrix S and mean measurement vector $\bar{y}$ can be pre-calculated and the implementation of COBRA then becomes very efficient in terms of processing time (about an order of magnitude faster than DOAS non-linear schemes).

However, the practical implementation for COBRA require some caution. The main difficulty lies in the definition of the ensemble Y used to construct S (and $\bar{y}$). The sample of N spectra should be highly representative of the measurement conditions under consideration, otherwise offsets in the SCDs will likely occur. Also, in principle, the spectra should be uncontaminated by absorption of the trace gas of interest. Finally, N should be large enough to insure statistically meaningful covariance results.

It should be stressed that COBRA is close in concept to the PCA SO_2 algorithm of Li et al. (2013, 2020a). In brief, the PCA scheme characterizes the background radiance variability using a number of leading PC spectra (typically 20-30), instead of a covariance matrix. The SO_2 column is then retrieved from the measured spectrum along with the PCs fitted parameters. In comparison, COBRA removes the need of having many parameters to fit. Only the SO_2 slant column density is determined and the background radiance variability is fully described by the covariance matrix. In a sense, COBRA can be considered as a generalization of the PCA scheme. It is therefore of great

interest to compare the two methods (see section 3.1). Having this perspective in mind, the parameters of COBRA for the retrieval of SO₂ from TROPOMI have been largely aligned with the PCA algorithm, to facilitate the comparison.

The input spectra for the covariance matrix calculation are analyzed separately for each TROPOMI row, to consider the row-dependent characteristics of the instrument. We also treat each orbit individually to account best for the orbit-to-orbit variability. The data are first screened for solar zenith angles larger than 60°, and to cope with the latitudinal dependence of the total ozone absorption and of the Ring effect, the data are divided into 6 equal and non-overlapping along-track segments. For each segment, an initial covariance matrix S is derived and initial estimates of SO₂ SCDs are inverted through equation 5. In a second step, improved estimates of S and SO₂ SCDs are obtained iteratively by removing SO₂ contaminated spectra from the ensemble Y . To do this, we use the ratio of the SO₂ SCD to its retrieval uncertainty (Eqs. 5 and 6), referred to as the signal to noise ratio (SNR):

$$SNR = \frac{k^T S^{-1} (y - \bar{y})}{\sqrt{k^T S^{-1} k}} \quad (8)$$

A fixed SNR upper value of 1.5 is used for the filtering. This is a rather strict value but tests over pristine regions indicate that this choice does not introduce biases in the SCDs data. The number of iterations is set to 4 but in general we found small changes in the results already after 2 iterations.

A lower limit on the number N of SO₂-free spectra is set to 50. If this limit is reached, because of a major volcanic eruption for example, the SO₂ SCD retrieval is entirely skipped for the corresponding row-segment pair. This is a limitation of the current algorithm version, but in the future a better handling of this problem would be possible, e.g. by using a covariance matrix fallback constructed from previously processed orbits. However, the amount of data skipped is small, on the order of 0.025% in total.

To help the comparison with the PCA SO₂ algorithm, we have used a spectral window from 310.5 to 326 nm (instead of 312-326 nm for the TROPOMI operational DOAS product), which includes the same strong SO₂ absorption bands as in the spectral range 310.5-340 nm used by Li et al. (2013). This choice is also motivated by the inclusion of the intense absorption band at 310.8 nm which leads to a further reduction of the noise on the SO₂ column by about 25%. Note that initial tests with the TROPOMI operational algorithm using the 310.5-326 nm window were actually not very successful (large SO₂ SCD offsets). On the contrary, with COBRA, we tested both

wavelength ranges (310.5-326 nm and 312-326 nm) and found only small differences between the retrieved SO₂ column patterns (Fig. S1).

In the following sections, SO₂ vertical columns will be presented. For the SCD to VCD conversion, we have used air mass factors from the operational product. Note that doing so is not strictly valid because one should expect lower AMFs due to the change in fitting window (from 310.5 to 326 nm to 312-326 nm). To account for this, we have applied a constant scaling factor of 1.15 to the retrieved SO₂ VCDs. Based on radiative transfer calculations, we found this to be a good first order correction. However, in the future, AMFs shall be recalculated properly. For the cloud filtering and AMF cloud correction, the operational cloud product OCRA/ROCINN CRB is used (Loyola et al., 2018; Compennolle et al., 2020).

As a final note, it should be reminded that the operational TROPOMI algorithm also handles the retrieval of large SO₂ VCDs, by making use of multiple fitting windows (as described in Theys et al., 2017). In this study, we have not applied COBRA on the alternative fitting windows. While there is no fundamental limitation to do so, COBRA is relevant mostly for low SO₂ columns. All the results presented in the next sections are for situations where the SO₂ VCDs are below 5 DU.

3. VERIFICATION OF THE RETRIEVALS

3.1 Comparison to satellite observations and CAMS

In order to evaluate the SO₂ data from COBRA, it is interesting to first investigate the bias and data scatter over a clean region and compare with the operational product (hereafter referred to as ‘DOAS’). In Figure 1, the mean and standard deviation of SO₂ slant columns over an equatorial Pacific region are shown for one particular orbit, as a function of the TROPOMI row. As can be seen from Fig. 1a, the DOAS data suffer from SCD offsets in the range of ± 0.25 DU, despite the background correction applied. These offsets have a low-frequency dependence component with the across-track position [which is not well understood](#), but also vary sharply from one row to the next (leading to stripes in the SO₂ maps). [Given that the background correction is applied separately for each row, this behavior points to limitations in the correction approach.](#) In contrast, the COBRA results have very small SCD biases (mostly below ± 0.025 DU) and no noticeable

across-track dependence. It follows that COBRA is a very powerful bias self-correction and destriping scheme. In Fig. 1b, the standard deviations of the SO₂ SCD values are shown for both algorithms. Compared to DOAS, it is clear that the data scatter is significantly improved with COBRA, by a factor of 2. It is understood that part of this noise reduction is due to the change in fitting window (section 2.2), but most of the improvement (~75%) is from the COBRA approach. From Fig. 1, it is clear that the combined reduction of bias and data scatter provided by COBRA over the DOAS results is very significant. From a practical point of view, a factor of 2 improvement of the data scatter means 4 times less pixels to average to reach a certain noise level. In Fig. 1b, we note also a distinct increase in data scatter for the outermost rows, for both DOAS and COBRA. This feature is due to difference in detector signal binning at the swath edges which leads to an increase in radiance shot noise. To keep the data of the best quality, we will not use the 50 outermost rows in the following of the paper.

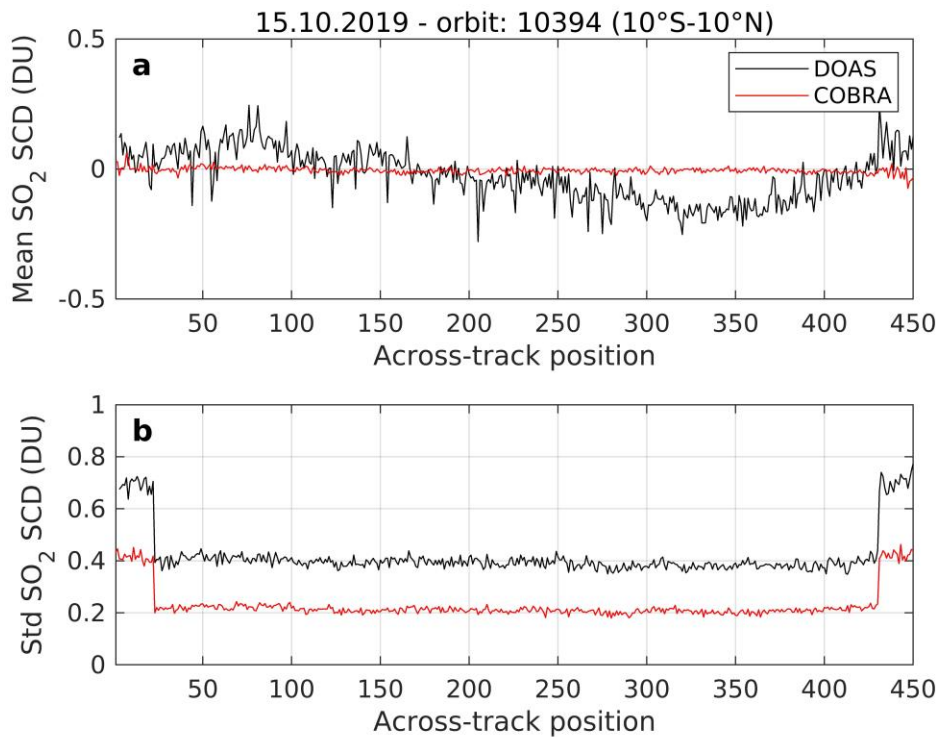


Figure 1. (a) Mean SO₂ slant columns from (black) DOAS (background corrected) and (red) COBRA for one orbit (10394 on 15 October 2019) over the equatorial Pacific region (10°S-10°N), as a function of the across-track position of TROPOMI, (b) same as (a) for the SO₂ SCD standard deviation.

Figure 2 compares the DOAS and COBRA seasonal averaged SO₂ VCD maps from September to November 2019. The data are gridded at a resolution of 0.1° x 0.1° and smoothed by a 2-dimensional 5-points box car function. Both DOAS and COBRA results are extracted using identical pixel selection criteria: SZA less than 60°, radiometric cloud fraction lower than 30% and TROPOMI rows 26-424. From Fig. 2, several artefacts are evident in the DOAS product. Negative values are found in the tropics and a large scale positive bias at mid-latitudes. In comparison, COBRA remarkably solves all the systematic biases found in the operational product whereas the signal from major SO₂ sources (e.g. in China, India, Middle-East, South Africa, Central and South America) is nicely preserved. Note that for individual pixels with unambiguous detection of SO₂ (typically SO₂ VCDs larger than 2 DU), the agreement between DOAS and COBRA is excellent (see e.g., Fig. S2). In Fig. 2, a closer look at the COBRA SO₂ map still reveals some negative values for specific locations. For instance, the Garabogazköl Basin near the Caspian Sea is particularly visible. It is characterized by a salt flat with a high albedo. This surface effect is apparently poorly represented in the radiance covariance, and leads to the negative values observed in the data.

Retrieval results using the new COBRA are also evaluated in Fig. 2 against a scientific TROPOMI SO₂ product generated using the PCA approach. The settings of the experimental TROPOMI PCA SO₂ algorithm, including the spectral range and number of iterations, are identical to the operational OMI algorithm with the following exceptions: 1) TROPOMI pixels from each row are grouped into sectors of 20-degree latitude bands, instead of three sectors as in the OMI algorithm; 2) a third degree polynomial is removed from each Sun-normalized radiance spectrum before PCA analysis; 3) at maximum 20 PCs are used in the fitting instead of 30 in the current OMI algorithm (Li et al., 2020a); and 4) no attempts were made to reduce TROPOMI retrieval noise over the SAA affected areas. For this exercise, the PCA scheme uses as input the same SO₂ absorption cross-section data (Bogumil et al., 2003) as for the DOAS and COBRA retrievals, and the same selection of pixels. Figure 2 also compares the TROPOMI SO₂ columns (from DOAS, COBRA and PCA) to the operational OMPS SO₂ PCA retrievals NMSO₂_PCA_L2 V2 (Zhang et al., 2017; Li et al., 2020c). Although OMPS has a coarser resolution (50 x 50 km²) than TROPOMI, it provides nonetheless a useful reference data because it operates on the Suomi National Polar-orbiting Partnership (SNPP) satellite which flies in loose formation with S-5P (i.e. 3-5 minutes difference of overpass time). To allow a meaningful comparison, the OMPS pixels were selected similarly as

1 TROPOMI, i.e. with cloud radiance fraction lower than 30% and OMPS across-track positions 3-
2 34. Note finally that to avoid discrepancies due to different a-priori profiles in the TROPOMI and
3 OMPS retrievals, a fixed AMF of 0.4 was used for all four data sets. As can be seen from Fig.2,
4 an overall excellent agreement is found between COBRA and PCA retrievals, the observed SO₂
5 spatial distributions being essentially the same. However, the OMPS SO₂ data set has different
6 patterns over China (possibly due to sampling differences), and also appears noisier than the
7 TROPOMI results (as expected from the smaller number of pixels). When comparing the
8 TROPOMI COBRA and PCA maps, very consistent results are found. Yet, the quality of COBRA
9 seems slightly better than the PCA retrievals. In particular, COBRA is much less sensitive to the
10 South Atlantic Anomaly than PCA data, which exhibit many outliers in the corresponding region.
11 At mid-latitudes, there is also a slight positive bias (of about +0.1 DU on average) and higher noise
12 in the PCA results compared to COBRA.

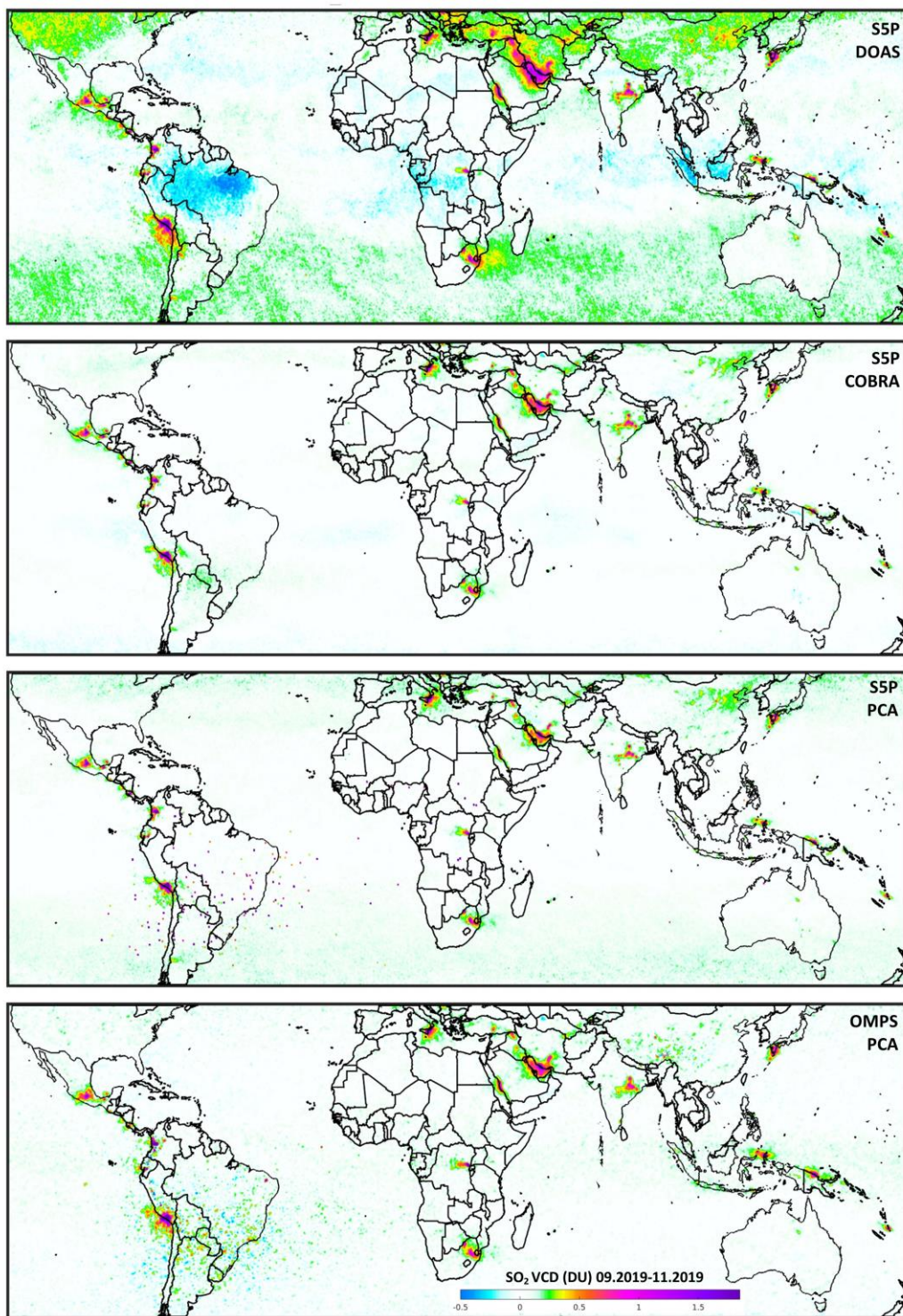


Figure 2. Comparison of seasonal mean SO_2 columns for September to November 2019 retrieved from TROPOMI DOAS, COBRA, PCA and OMPS PCA algorithms (from top to bottom). Consistent pixel selection criteria, gridding and retrieval settings are applied (see text). For all four data sets, a fixed AMF of 0.4 is applied.

We have estimated the data scatter for the three TROPOMI data sets, based on measurements from the same orbit over the Pacific as Fig. 1. Results are shown in Figure 3, as a function of latitude. We find that COBRA has a SCD noise level 20-25% lower than the PCA retrievals, and twice better than DOAS (as in Fig. 1). Translating the numbers of Figure 3 in terms of vertical columns for a typical pollution scenario, we estimate the retrieval precision for individual pixels typically to be 0.5 - 1 DU for COBRA.

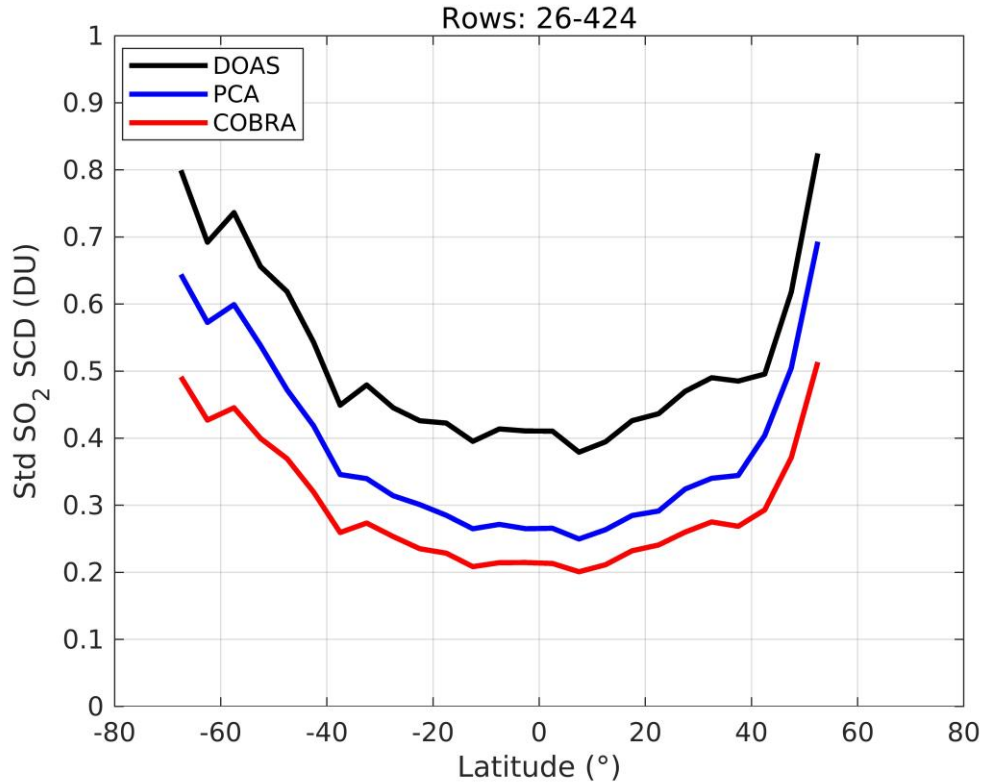


Figure 3. Standard deviation of the SO₂ slant columns as retrieved from DOAS (black), PCA (blue) and COBRA (red) for one TROPOMI orbit (10394 on 15 October 2019, same as Fig. 1) for rows 26-424, as a function of latitude (for 5° bins).

To further evaluate the overall quality of the COBRA retrievals, the SO₂ VCDs can also be compared to model data. Here, we have used the output of the Copernicus Atmosphere Monitoring Service (CAMS; <https://atmosphere.copernicus.eu/>) regional model, for September to November 2019. The CAMS regional air quality production is based on an ensemble of 9 European air quality models that are run at a resolution of 0.1° and produce 4-day, daily forecasts of the main atmospheric pollutants, including SO₂. The forecasts and analyses from all 9 models are combined

1 in calculating the median value of the individual outputs, which is designated as the ENSEMBLE
2 output and is the field used in this study. The CAMS regional ensemble data was obtained from
3 the Copernicus Atmosphere Data Store (ADS, <https://atmosphere.copernicus.eu/data>). More
4 information about the CAMS regional system can be found on the ECMWF website
5 ([https://confluence.ecmwf.int/display/CKB/CAMS+Regional:+European+air+quality+analysis+a
6 nd+forecast+data+documentation](https://confluence.ecmwf.int/display/CKB/CAMS+Regional:+European+air+quality+analysis+and+forecast+data+documentation)). The CAMS regional system used the CAMS-REG-
7 AP_v2_2_1 emissions (reference year: 2015) between June 2019 and February 2020, and the
8 updated CAMS-REG-AP_v3_1 emissions dataset (reference year: 2016) since February 2020.

9 In Figure 4, seasonal regional maps of S-5P SO₂ VCDs over Eastern Europe from the DOAS and
10 COBRA schemes are compared to the output of the CAMS regional model, for September to
11 November 2019. From the maps, it is clear that the COBRA results are in much better agreement
12 with the CAMS analysis than the DOAS data. Owing to the quasi-absence of bias and the low
13 noise level, the COBRA data allows better isolation of the emission sources. The agreement
14 between COBRA and CAMS is however not perfect and there are several explanations for this.
15 Most of the SO₂ emissions in this region are from coal-fired power plants and the emission
16 inventory used by CAMS is likely reflecting neither the actual activity nor the emission mitigation
17 solution (e.g. SO₂ scrubbers) at each power plant. Noteworthy is also the absence of SO₂ emissions
18 from Mt. Etna in CAMS. Secondly, the AMFs used here are calculated with SO₂ profiles from
19 TM5, a different model with a coarser resolution (1° x 1°) than CAMS regional. Therefore the
20 COBRA and CAMS SO₂ columns cannot be strictly compared. Nevertheless, the comparison in
21 Fig. 4 is encouraging. In the future, the COBRA SO₂ retrievals together with the corresponding
22 column averaging kernels (Eskes and Boersma, 2003) could be ingested in the CAMS assimilation
23 system to better constrain the model SO₂ output and emission estimates.

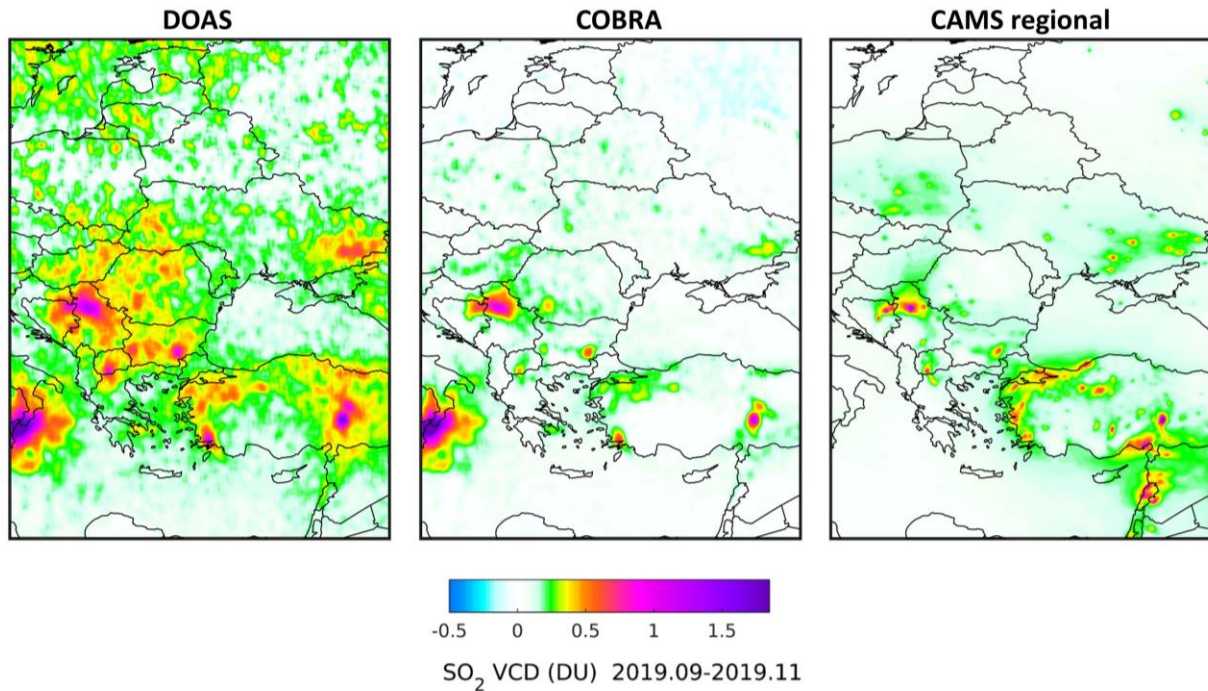


Figure 4. Seasonal mean SO₂ columns for September to November 2019 from (left-center) TROPOMI DOAS and COBRA retrievals, and (right) simulated by the CAMS regional model. The CAMS data are displayed at the 0.1° x 0.1° native resolution.

3.2 Comparison to ground-based MAX-DOAS observations

The Multi-Axis DOAS (MAX-DOAS) measurement technique is an established method to retrieve tropospheric trace gas columns and vertical profiles from a sequence of spectral observations performed at various elevation angles above the horizon (Hönninger and Platt, 2002; Tirpitz et al., 2021). MAX-DOAS measurements leverage the fact that low elevation measurements have enhanced sensitivity to atmospheric pollutants in the boundary layer and that the combination of the different elevations carries information on the vertical distribution of the trace gas of interest as well as aerosols. The simplest estimation of the tropospheric VCD from MAX-DOAS measurements is obtained by scaling the differential SCD at a given elevation angle (often 15° or 30°) with an AMF assuming a geometrical light path through the trace gas layer. Recently, more sophisticated approaches have been developed to retrieve the concentration profile in the troposphere using multiple elevation measurements.

Here we compare our TROPOMI SO₂ VCD data to MAX-DOAS observations at two sites, both characterized by relatively low SO₂ columns: Xianghe and Mohali (Table 1). In general, the

different MAX-DOAS instruments and SO₂ retrievals share common characteristics, practices and approaches, and the reader is referred to the publications listed in Table 1 for more detailed information.

For the comparison, we have used a common set of selection criteria for the satellite data. For each day, we selected the TROPOMI pixels within a 25 km radius circle around the station of interest, a strict radiometric cloud fraction threshold of 20%, SZA lower than 60° and AMF larger than 0.2. If the number of retained pixels is larger than 10 then the mean SO₂ VCD is calculated and compared to the averaged SO₂ column for the MAX-DOAS measurements within ±1 hour of the S-5P overpass time.

Regarding the ground-based retrievals, the SO₂ VCDs are estimated, for Mohali, using the 15° elevation SO₂ SCDs and geometrical AMFs. Conversely, the retrieved data for Xianghe consist of SO₂ profiles. These are integrated along the vertical to provide the VCDs. Moreover, to make the comparison between MAX-DOAS and TROPOMI more consistent, we have rescaled the TROPOMI VCDs using the satellite averaging kernels (Eskes and Boersma, 2003) and the MAX-DOAS SO₂ profiles at Xianghe.

Table 1. Summary of SO₂ VCDs comparison

Station	Reference	MAX-DOAS VCD calculation	Period	Mean SO ₂ VCD (DU) for SZA<50° (50°<SZA<60°)		
				MAX- DOAS	S-5P COBRA	S-5P DOAS
Xianghe, China 39.77°N 117°E	Wang et al. (2014)	Integrated profile	01.2020-10.2020	0.2 (0.28)	0.21 (0.34)	0.34 (0.77)
Mohali, India 30.67°N 76.74°E	Kumar et al. (2020)	15° elevation (geometrical)	05.2019-10.2020	0.21 (0.18)	0.18 (0.26)	0.16 (0.50)

The comparison results between COBRA and MAX-DOAS measurements are shown in Figure 5a and 5b, for Xianghe and Mohali, respectively. Overall, the agreement between COBRA and MAX-DOAS data is very good, keeping in mind that the levels of SO₂ columns are quite low. The slopes of the regression lines are close to unity. In Table 1, the mean SO₂ columns from MAX-DOAS, TROPOMI COBRA and DOAS retrievals are given at each station and for different SZAs. A striking feature of the comparison is that the COBRA results show similar good agreements over

a wide range of SZA. It further supports the idea that COBRA yields unbiased results over varying observation conditions. This is in contrast to the DOAS product which is clearly biased high for high SZA. For completeness, the comparison results between TROPOMI DOAS and MAX-DOAS measurements are shown in Fig. S3, for both Xianghe and Mohali stations. The agreement is clearly not as good as for the COBRA vs MAX-DOAS comparison, both in terms of the correlation coefficients and slopes of the regression lines.

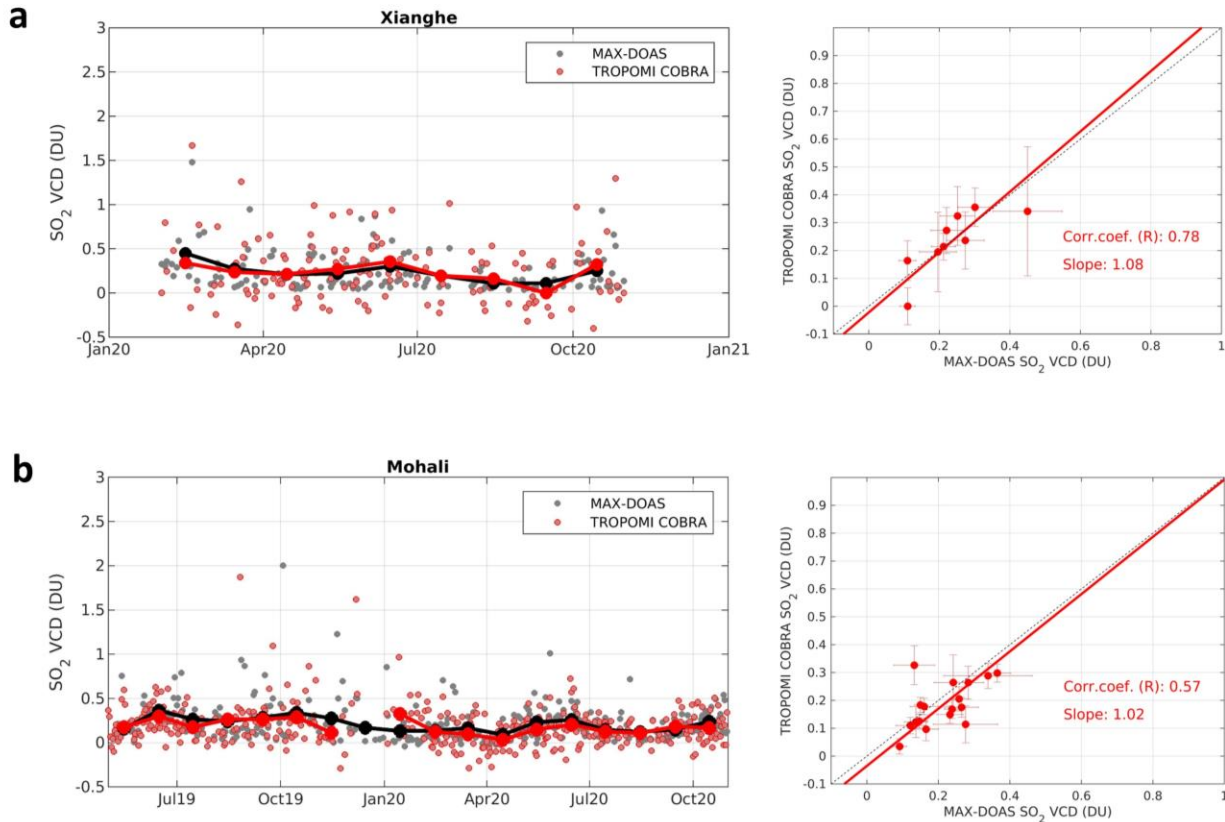


Figure 5. (left) Comparison of monthly mean SO₂ columns from MAX-DOAS and TROPOMI COBRA for (a) Xianghe, and (b) Mohali. The grey and pale red dots correspond to the individual days. (right) Scatter plots of monthly mean SO₂ columns of TROPOMI COBRA vs MAX-DOAS observations. Error bars are the standard errors on the monthly average SO₂ columns. The correlation coefficient and slope of the regression line are given as an inset for each plot.

4. GLOBAL RESULTS

In this section, we present long-term global results from COBRA, based on two and a half years (April 2018 – December 2020) of cloud-free TROPOMI data (radiometric cloud fraction less than 30%). Using an oversampling technique, a global average SO₂ column map at 0.015° x 0.015° resolution was obtained and smoothed by a 2-dimensional 10-points box car function. Figure 6 shows the resulting SO₂ distribution for specific regions, over East China, India, the Middle East, South America, South Africa, US and Europe (the global map is also available in the form of a Google Earth /geotiff file, as supplementary material). Figure 6 also shows the locations of the SO₂ sources based on the latest OMI 2005-2019 catalogue (Fioletov et al., 2016), with a total of 588 sites, including power plants, smelters, oil and gas industry sources, and volcanoes. As can be seen, many of the sources of the catalogue are easily identified as SO₂ hotspots on the map. Conversely, there is also a significant number of sources in the inventory with no detectable SO₂ in the TROPOMI data, but one should keep in mind that the catalogue gathers emission sources since the beginning of the OMI data record in 2005 and several of these sources have ceased operations or decreased drastically their emissions since then (e.g. due to the operation of SO₂ scrubbers in coal power plants). To help identifying those sources, Figure 6 shows with a different marker the sources detected by OMI for 2018-2019 (i.e. the sources with emissions above the detection limit, as in Fioletov et al. (2016) and McLinden et al. (2016); see also Section 5.2). This is also helpful to highlight the differences in sensitivity, as many sources are detected by TROPOMI but not by OMI.

Generally speaking, the SO₂ maps of Figure 6 are very detailed. Biases over clean regions are remarkably low, and emissions-related patterns with SO₂ VCDs less than 0.25 DU are clearly visible in many places. By scrutinizing the SO₂ distributions, one can identify numerous sources from the current catalogue but also several potentially new source regions. However, some care must be taken in attributing new sources and relate this to the improved sensitivity of TROPOMI COBRA. First, the catalogue is arguably not resolving well the individual sources for dense regions (e.g., in East China and India) and, as a matter of fact, typically reports a total emission estimate of a point-like source for what is in reality a cluster of sources. While the algorithm to handle such clusters of sources exists (Fioletov et al., 2017), it has not been implemented in the catalogue yet. Second, the SO₂ catalogue is being populated on a best-effort basis, and a number of emission sources might be missing, in particular for emerging countries where industrial

infrastructures are built quickly. Third, SO₂ outflow from the strong sources - or clusters of sources - can lead to variations in the map and thus fictitious emission sources. Finally, retrieval artefacts, measurement noise or sampling related issues can also lead to false sources identification. Note that a comprehensive identification and classification of new sources from the COBRA SO₂ data is not the scope of the present study. Here, we aim at discussing plausible new SO₂ sources (i.e. not in the OMI catalogue). In Section 5.2, we will further demonstrate the excellent performance of COBRA to detect very weak emissions, for a limited number of sources.

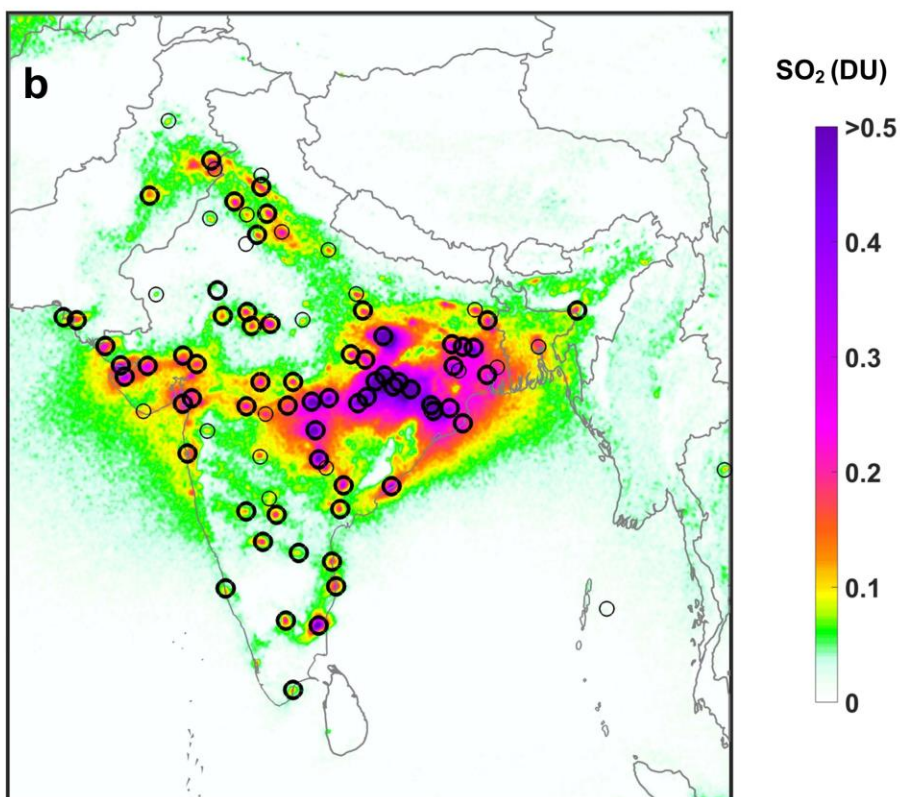
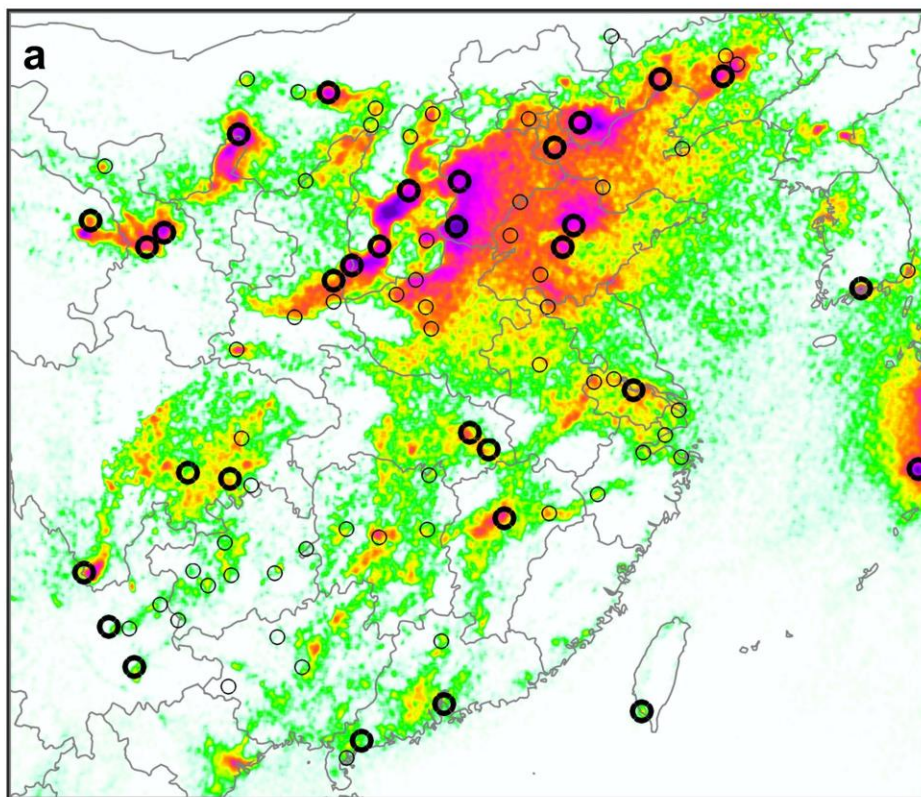
The new identified sources are characterized by low SO₂ column levels in the range 0.05 - 0.2 DU. For instance, in Fig. 6a we observe hotspots of SO₂ from power plants (mostly coal but also likely gas) in North- and South-Korea, northern Vietnam (near Haiphong), several Chinese provinces (e.g., Hubei, Guangxi, Guangdong) and along the coast of China. In Fig. 6b, several weak emission sources can be isolated in India (e.g., over the western coast and the Indo-Gangetic plain), Pakistan, Bangladesh and Sri Lanka (near the city of Colombo). In the Middle East (Fig. 6c), most of the SO₂ emissions are from oil and gas related industries, like power plants, gas flaring and refineries. Examples of weak SO₂ emissions can be found in Saudi Arabia, Oman, Egypt, Syria (near the city of Damascus) and Iran. In South America (Fig. 6d), new emission sources are popping up, notably, in Brazil (near Rio de Janeiro, São Paulo and Porto Alegre), and on both sides of the Andes (in Chile and Argentina). In South Africa (Fig. 6e), in addition to the strong emissions from the coal power plants of the Highveld, a clear SO₂ signal is detected over Cape town. Interestingly, the measured SO₂ distribution nicely matches the orography setting. In the US (Fig. 6f), the most striking emission region is the state of California, with enhanced SO₂ over the Central Valley and the city of Los Angeles. Over the central and eastern parts of the US, the emissions from power plants have declined dramatically over the last 15 years (Krotkov et al., 2016). However, the data still show enhanced SO₂ over some of them. In Europe (Fig. 6g), most of the observed enhanced SO₂ correspond to sources already in the catalogue. Still, a number of small spots are found e.g., in eastern Europe (Romania, Serbia, Kosovo, Hungary), Germany (near Leipzig), Turkey and Tunisia (Gulf of Gabes). Interestingly, enhanced SO₂ is also observed over the Gibraltar strait and Red sea which might result from shipping emissions.

Overall, the SO₂ maps of Fig. 6 nicely illustrate the great ability of TROPOMI to detect weak SO₂ point emissions sources when analyzed using a sensitive approach as COBRA. Using Google Earth imagery and information on industrial facilities location, we were able to confirm that many

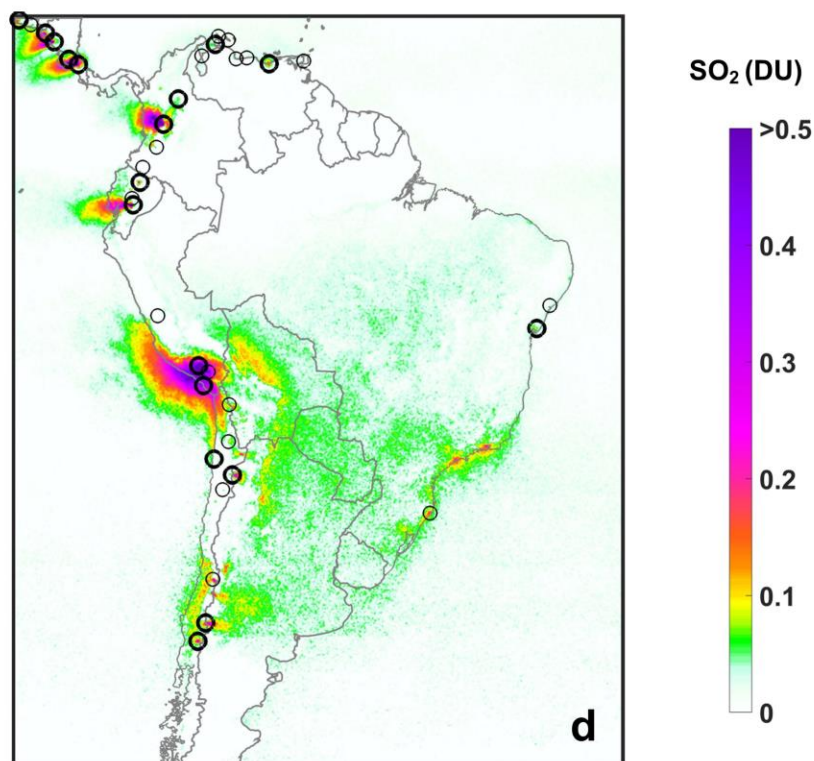
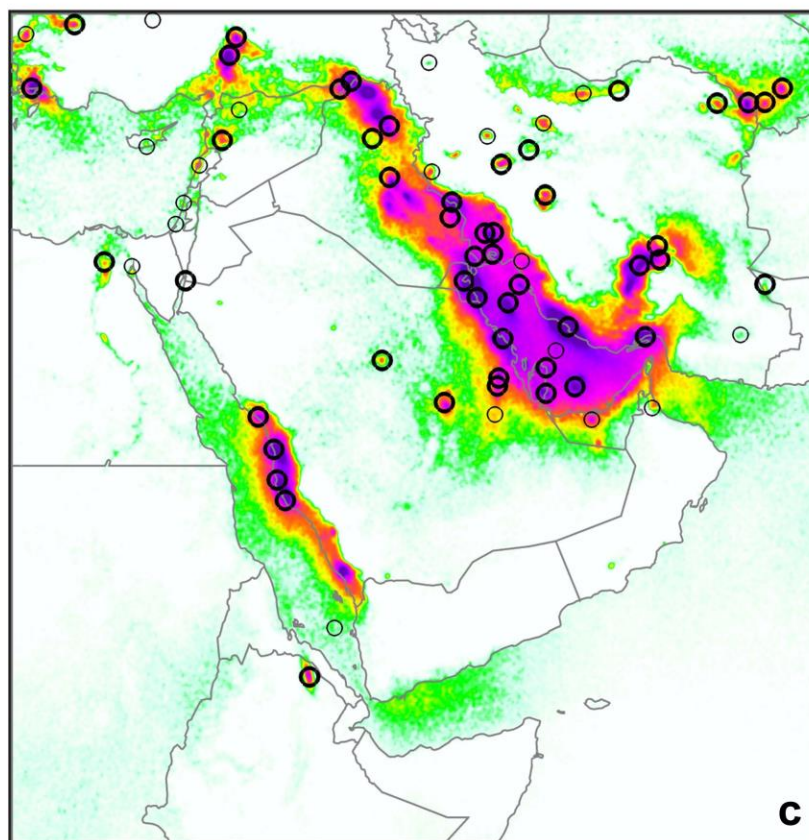
features in the SO₂ map are real sources. For this, we have also compared our SO₂ data to tropospheric NO₂ column maps from TROPOMI. An example of comparison is shown in Fig. S4 for a region over Central Asia. There, the SO₂ emissions sources in the catalogue are mostly from coal power plants and smelters, in the Xinjiang province (China) and east Kazakhstan. As can be seen in Fig. S4, several other SO₂ emission hotspots are detected (notably in the Xinjiang province) which clearly coincide with locations with enhanced tropospheric NO₂.

Nevertheless, several patterns in the SO₂ map (Fig. 6) are hard to relate to point source emissions. In particular, the SO₂ signal observed over Cape town (Fig. 6e) and Los Angeles (Fig. 6f) could be due to area sources rather than point emissions. Over South America (Fig. 6d) and eastern US (Fig. 6f), the apparent SO₂ background is intriguing. It is unclear whether this could be due to real SO₂ emissions or not. We also identify several artefacts in the data. Unsurprisingly, biases in the data occur for specific conditions which are under-sampled or not optimally represented by the covariance matrix. These are most often surface-related effects (due to peculiar albedo or elevated terrain). One illustration of this problem is given in Fig. 6c, over the Nile Valley. Although some real SO₂ emissions are found in the area, with SO₂ VCDs larger than ~0.1 DU, there are also unexpected enhancements in the SO₂ column that follow the Nile River. These are probably due to the very dark surfaces there. Similarly, elevated values are also found further South in Sudan and Ethiopia, over vegetated scenes. However, the resulting SO₂ VCD biases are overall very small, typically less than 0.04 DU ($\sim 1 \times 10^{15}$ molecules/cm²), and can be suppressed by a local bias correction or more sophisticated approaches.

As mentioned above, the attribution of new sources based on SO₂ maps is not straightforward. Efficient space-based techniques do exist, to isolate sources and estimate their emissions (Fioletov et al., 2015; McLinden et al., 2016; Clarisse et al., 2019). However, applying such methods systematically to the TROPOMI COBRA SO₂ data goes beyond the scope of this paper. Instead, in the next section, we will estimate the SO₂ emissions for the known largest sources, and demonstrate the potential of COBRA for the retrieval of weak emissions, for a limited number of new sites.



1 Figure 6. Averaged SO₂ column (in DU) for April 2018 to December 2020 over (a) East China, (b)
2 India, (c) the Middle East, (d) South America, (e) South Africa, (f) US and (g) Europe. The black
3 circles mark the locations of SO₂ sources detected by OMI (in bold for the 2018-2019 period, see
4 text). Due to the massive eruption of Raikoke on 21 June 2019, all data in the northern hemisphere,
5 for the 3 months period after the eruption, are filtered out.



1
2 Figure 6. Continued.

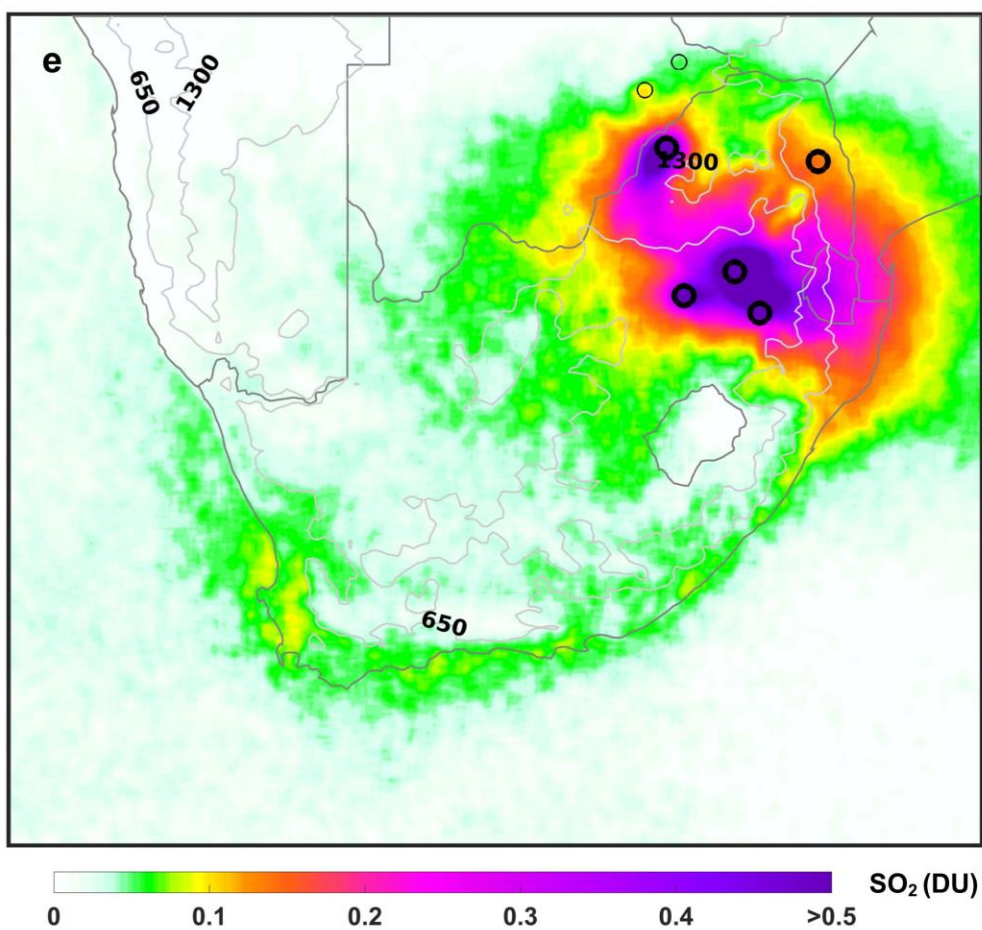
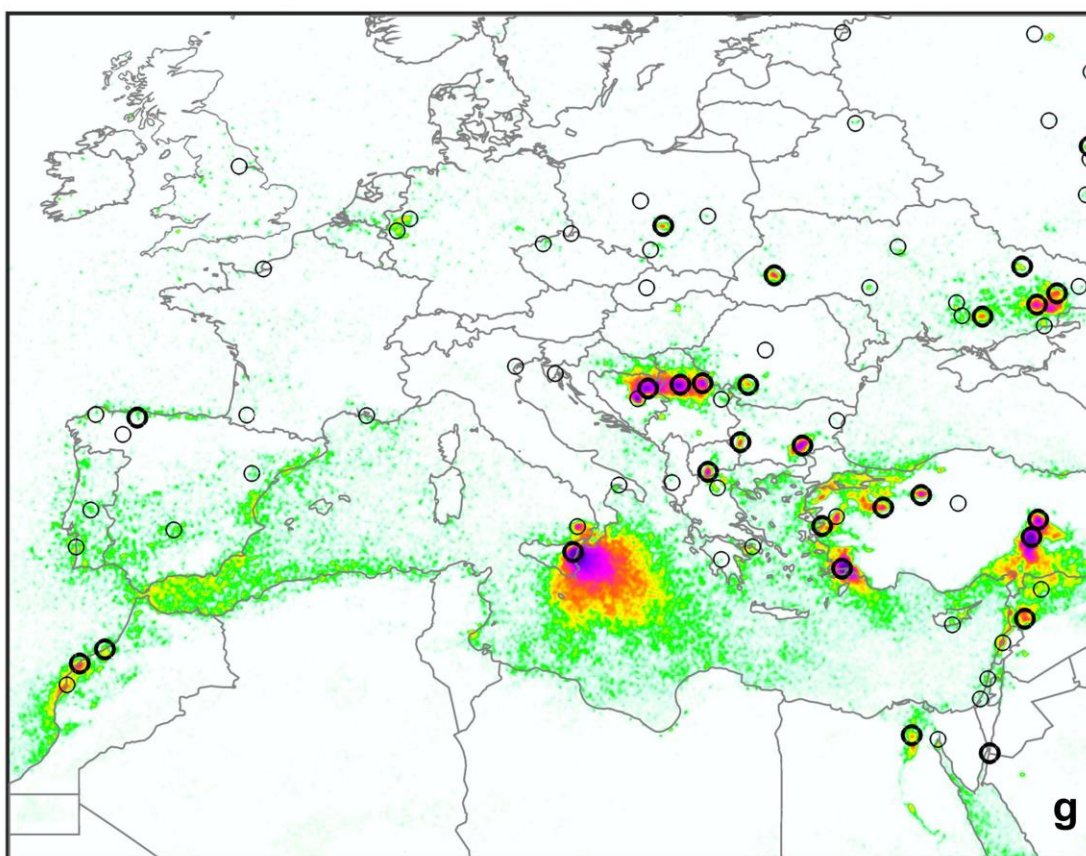
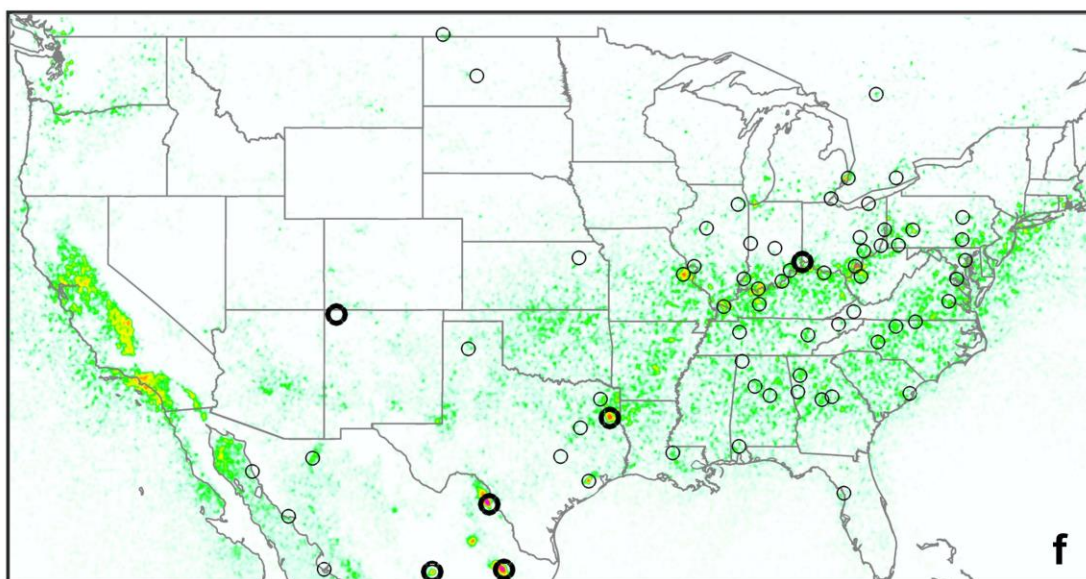


Figure 6. Continued. The gray lines are the topography isolines (in meter). Note that to further reduce the data scatter, the SO_2 map was smoothed by a 2-dimensional 20-points box car function (instead of 10-points function for the other sub-figures).



1
2
3 Figure 6. Continued.

5. EMISSIONS ESTIMATES

Satellite observations are being increasingly used to estimate SO₂ emissions. In particular, new methods have been very successful in deriving reliable emission rates, and even detecting missing sources, by combining satellite SO₂ columns and wind information, without the need of atmospheric chemistry transport models (e.g., Beirle et al., 2014; Fioletov et al., 2016, 2017, 2020; McLinden et al., 2016; Carn et al., 2017). These techniques have been used to derive global SO₂ emissions inventory from OMI observations (Liu et al., 2018). Recently, Fioletov et al. (2020) presented an analysis using the TROPOMI operational SO₂ product and found overall consistent results with the OMI emissions estimates. The TROPOMI-based emissions uncertainties were found a factor of 1.5 - 2 lower than the ones from OMI. In this section, we repeat the same analysis using the COBRA SO₂ retrievals and investigate the added value of COBRA for the estimation of SO₂ emissions. The details of the inversion technique can be found in Fioletov et al. (2015) and references above.

In brief, the method considers a potential point source and apply a wind rotation of the satellite measured SO₂ VCDs around this location. This first step enables to align all plume dispersion patterns along a fixed direction and leads to an improved SO₂ detection limit. By contrasting the upwind and downwind averaged SO₂ columns, the wind rotation procedure allows to confirm whether the test location is a real emission source and also to correct for a possible bias in the data. Note that for this first step, the retrieved SO₂ VCDs are rescaled using site-specific AMFs so that realistic SO₂ emission profile shapes (based on the elevation of the site and climatological boundary-layer height) are used for all analyzed sources.

The second part of the retrieval method deals with the emission estimate itself. The averaged downwind SO₂ field is modelled by an exponential modified Gaussian function which accounts for the SO₂ total mass, e-folding time and plume width. From the fitted parameters, the average SO₂ emission rate can be derived directly. Here the baseline inversion is however not to fit all three parameters but rather to prescribe the e-folding time and plume width, and therefore the only parameter derived from the fit is the SO₂ total mass which is directly proportional to the SO₂ emission rate.

5.1 SO₂ emissions for large sources

The method was applied to the SO₂ data from COBRA for 274 large emissions sources, including power plants, volcanoes, oil and gas sources and smelters, distributed worldwide. In Fig. 7a, the results are compared to the analysis of Fioletov et al. (2020) using the TROPOMI DOAS product, for the period from April 2018 to March 2019.

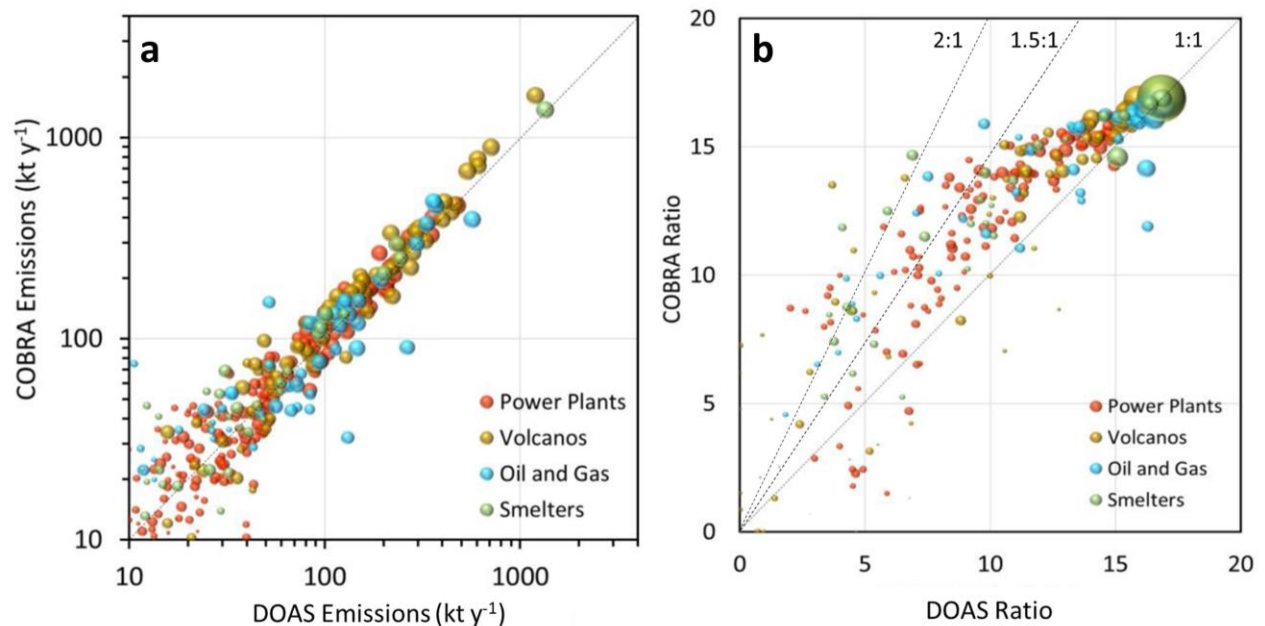


Figure 7. (a) Estimated SO₂ emissions from TROPOMI, based on the COBRA and DOAS algorithms analyzed for power plants, volcanoes, oil and gas industries, smelters sources. (b) Ratios of the estimated emissions and the corresponding uncertainties. The size of the marker is proportional to the average of COBRA and DOAS (a) ratios between emission value to its corresponding uncertainty, (b) estimated SO₂ emissions.

In general, the emission estimates from COBRA and DOAS are fairly consistent for all four source types. For two-thirds of all sources, the differences between DOAS and COBRA emission estimates are within ± 3 times the standard deviations of the DOAS-based emissions. However, it was found that the local bias in the satellite data (as derived from the upwind SO₂ columns) are much higher with DOAS (~ 0.25 DU) than with COBRA (~ 0.05 DU), and that the large differences between DOAS and COBRA emission estimates for some sources, are related to problems with the DOAS algorithm. Also, as result of the large improvement in the noise level, the estimated

emissions uncertainties are significantly improved with COBRA compared to DOAS, by 20-50% on average (see Fig. 7b).

It should be emphasized though that the improvement of emission uncertainties depends on the emission level. The sources considered here are relatively large sources that have been previously detected by OMI. The TROPOMI COBRA SO₂ data set presented in this study combines the advantages of high spatial resolution, low noise level and almost no bias. It has therefore the potential to detect weaker sources (as shown in Sect. 4).

5.2. Detection of weak emissions

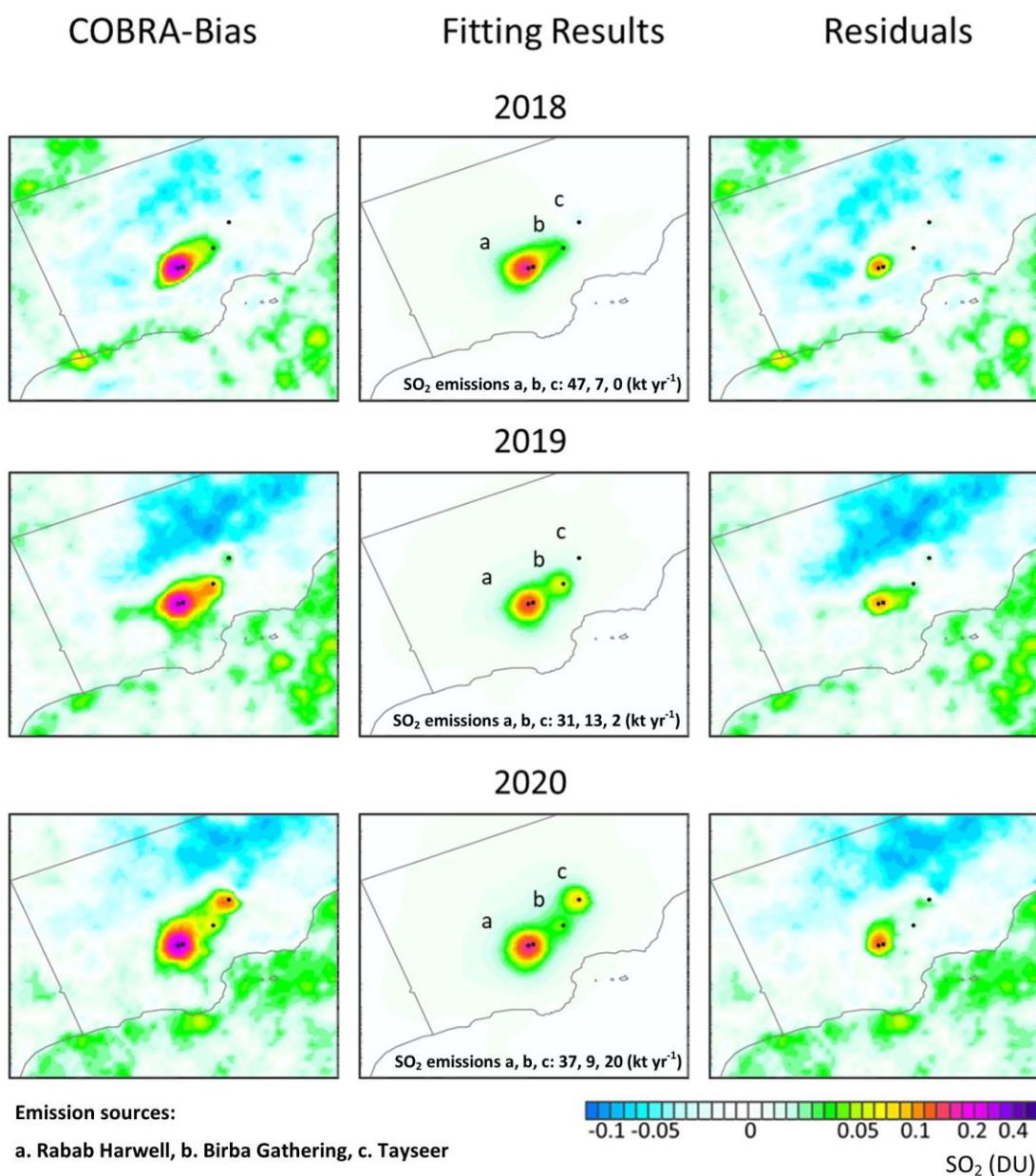
It is enlightening to estimate the lowest level of SO₂ emission detectable by COBRA. Clearly, it is expected to be dependent on the observation conditions and generally speaking the best detection limit is obtained for sites with low noise on the SO₂ SCDs and the highest measurement sensitivity (i.e. high AMFs). These sites are found at low latitudes and in particular at high elevations or for high albedos. To estimate the emission detection limit, we define the statistical significance of an emission signal as three times its standard error. Based on the global sources presented above (section 5.1), we performed statistics using this metric. To avoid biases by the strongest sources, we only considered the sources with estimated emissions less than 50 kt y⁻¹. The resulting detection limit values are found in the range between 4 and 11 kt y⁻¹ depending on the AMF, with a mean value of 8 kt y⁻¹. It is important to realize that this limit of detection is remarkably low, at least twice better than using TROPOMI DOAS data. It is also a factor of 4 smaller than the detection limit of 30-40 kt y⁻¹ offered by OMI for the first years of operation (Fioletov et al., 2016; McLinden et al., 2016). This suggests that the TROPOMI COBRA implementation is [excellent](#) in exploiting the gain in spatial resolution of TROPOMI compared to OMI (~16 times smaller pixel sizes). This finding is supported by the fact that the noise levels for individual pixels for TROPOMI COBRA and OMI PCA VCDs are similar (not shown; see also section 3 of Fioletov et al., 2020).

In the following, we demonstrate the potential of the TROPOMI COBRA SO₂ data set to detect and quantify weak emissions. For this, we use a slightly adapted version of the inversion technique of section 5.1, and illustrate the method on a selection of new emission sources.

The region of interest is the Dhofar governorate in southern Oman. There, the exploitation of oil and gas fields is growing fast, with a number of rapidly evolving projects of exploration and

production. In Figure 8 (left column), the yearly averaged TROPOMI SO₂ maps over South Oman are shown for 2018-2020. One can clearly identify and isolate 3 main emission locations, namely the Rabab Harwell integrated plant (18.03°N, 54.64°E), the Birba Gathering station (18.32°N, 55.10°E) and the Tayseer gas field (18.71°N, 55.34°E). We note that these emission sources are not listed in any emission inventory and the actual locations of the sources are approximated from available visible imagery (e.g. Google Earth). A noticeable feature in Figure 8 is the very low observed SO₂ column level, in particular over Birba Gathering and Tayseer with SO₂ VCDs of 0.03 – 0.1 DU, reflecting again the great sensitivity of COBRA. To estimate the SO₂ emissions from the TROPOMI data, the source method used in section 5.1 has been refined and tuned for this particular case study. A multi-source SO₂ emission retrieval was applied as in Fioletov et al. (2017) with one modification: a regression term proportional to the elevation was added to the fit to adjust for a small altitude-related bias in retrieved SO₂ (the values were slightly lower over the mountains near the Arabian Sea coast). This multi-source method is motivated by the fact that the sources are close to each other (~ 50-100 km distance) and the emissions cannot be fitted separately. Here, the approach basically allows for overlaps of the modelled SO₂ spatial distributions: the emissions from the individual sources are then adjusted so that the total SO₂ modelled field fits best the observed SO₂ VCD distribution. In Figure 8, the results of the fit are shown (center column), as well as the residuals of the fit (right column). The estimated annual SO₂ emissions for the 3 sources are given in the inset of Figure 8. Note that for this particular case, the emission detection limit (as defined above) is typically of about 6 kt yr⁻¹. For the Rabab Harwell site, the algorithm retrieves rather high and stable emissions over the years, with an average value of about 40 kt yr⁻¹, which is well above the estimated detection limit. Interestingly, the Rabab Harwell site has large residuals of ~0.1 DU for all years. This suggests that the point source representation used here is likely not sufficient to explain the observations and it is possible that there are many small contributing sources in the area. For the Birba Gathering site, the estimated emissions are much smaller and lie in the range of 7-13 kt yr⁻¹. Yet, there is a good confidence that these emissions are real, given that the estimates are a factor of 1-2 larger than the limit of detection. However, it is clear that the uncertainty of the emission estimates are quite large. For the Tayseer site, an SO₂ signal could be detected only recently. In 2019, the estimated emissions are of 2 kt yr⁻¹, i.e. below the detection limit, and in 2020, the SO₂ emissions strongly increased to about 20 kt yr⁻¹, probably as a result of a change in operation at the production facility. Finally, note that no significant residuals could be

1 found neither for Tayseer nor Birba Gathering site, and this suggests a point source behavior, at
 2 both sites.



3
 4 Figure 8. (Left) Yearly mean TROPOMI SO₂ columns retrieved from COBRA over South Oman
 5 for 2018 (April to December), 2019, 2020 (from top to bottom), after bias correction for the effect
 6 of elevation (see text). Three distinct SO₂ spots are discernable from the maps and are the results
 7 of emissions from oil and gas fields, referred as Rabab Harwell, Birba Gathering and Tayseer,
 8 (center) results of the fitting of TROPOMI SO₂ data. The estimated annual SO₂ emissions (given

in the inset for the three sources) are used to reconstruct the SO₂ column field, (right) residuals of the fit or the difference between TROPOMI and fitting results.

In summary, the analysis over South Oman of Figure 8 nicely illustrates the strength of a highly sensitive scheme such as COBRA when applied to a high spatial resolution instrument as TROPOMI. The fact that such low SO₂ emissions can be tracked and quantified with that level of detail is remarkable. Although not shown, the emission inversion scheme was successfully applied not only to the South Oman sources but also to other test sites, as they were found in the global SO₂ map (Figure 6). The wind-rotation technique when applied to TROPOMI COBRA SO₂ data is arguably a promising tool to monitor weak SO₂ emissions and track the activity from rapidly emerging production facilities worldwide. However, applying the inversion scheme at the global scale is a significant effort, as it also requires some level of manual intervention and testing. For instance, the information on source type, location, etc. is typically lacking, and the supporting visible imagery - useful for identifying industrial facilities - is often outdated.

6. CONCLUSIONS

A new spectral fitting method for the retrieval of sulfur dioxide columns in the UV was presented and demonstrated for TROPOMI. Based on a dynamical total measurement error covariance, the method, called COBRA, allows reducing considerably the noise level (by a factor of 2) and biases present in the current TROPOMI DOAS SO₂ operational product. COBRA provides greater sensitivity to low SO₂ columns, and this conclusion is supported by MAX-DOAS observations. Preliminary comparison of COBRA to PCA retrievals suggests similar and even better algorithm performance. The SO₂ vertical column precision for individual pixel is in the range 0.5 - 1 DU. The main limitation of the method relates to the set of spectra chosen to build the covariance matrix, to what extent it is uncontaminated by SO₂ and its distribution representative of the observations in the absence of SO₂. In particular, very bright or very dark scenes may be poorly represented by the covariance matrix. These conditions (i.e. intensity outliers) can lead to retrieval artefacts. However, the systematic VCD uncertainty (contribution from the COBRA spectral fit only) is very small, typically less than 0.04 DU.

The benefit of COBRA is clearly demonstrated in this work using long-term oversampled averages. Owing to the excellent quality of the data (in terms of precision and accuracy), the high

1 spatial resolution of TROPOMI can be better exploited. Zoomed SO₂ maps reveal new emission
2 sources worldwide, with low SO₂ columns of 0.05 - 0.2 DU, or even lower.

3 By using the COBRA SO₂ data over large emission sources, we have recalculated the SO₂
4 emissions obtained by Fioletov et al. (2020) that were based on the TROPOMI operational SO₂
5 product. While the derived emission rates agree generally well, we found that the uncertainties on
6 the emissions are significantly lower (up to 50%) using COBRA than with the operational product.
7 This opens the possibility to retrieve SO₂ emissions for weakly emitting sources, and we present a
8 number of examples that demonstrate the potential of the COBRA data in this direction.

9 With an estimated annual emission detection limit of about 8 kt yr⁻¹, the TROPOMI COBRA SO₂
10 data provides unique access to weak anthropogenic and volcanic point sources, and can help
11 completing current SO₂ emission inventories. It can also be used to track more accurately weak or
12 rapid changes in SO₂ levels, e.g., due to COVID-19 lockdown measures (Levelt et al., 2021) as
13 well as estimate seasonal and even monthly emissions. Finally, COBRA data would be particularly
14 relevant for the CAMS assimilation system as well.

15 COBRA is a good candidate for an implementation in the TROPOMI operational processor, with
16 limited computational resources and without the need for a separate background correction processor.
17 COBRA is also adaptable to other satellite instruments, including from geostationary platforms.
18 In particular, the European Sentinel-4 mission would likely benefit from a COBRA approach for
19 the retrieval of SO₂ columns, as the atmosphere will be sounded under unfavorable large
20 observation angles.

21 Future work could also be dedicated to the application of COBRA to historical sensors, in order to
22 produce a consistent long-term SO₂ data record, but also to the retrieval of other molecules.

CODE AND DATA AVAILABILITY

The TROPOMI COBRA SO₂ dataset is available from the corresponding author on request. The TROPOMI DOAS SO₂ product is publically available on the Copernicus Sentinel-5P data hub (<https://s5phub.copernicus.eu>). The TROPOMI PCA SO₂ dataset is available from Dr. Can Li on request. The OMPS PCA SO₂ is publically available from Goddard Earth Sciences (GES) Data Information Service Center (DISC) (https://daac.gsfc.nasa.gov/datasets/OMPS_NPP_NMSO2_PCA_L2_2/summary).

The CAMS regional data are available from the Copernicus Atmosphere Data Store (<https://atmosphere.copernicus.eu/data/>). The SO₂ emissions estimates can be obtained from Dr. Vitali Fioletov on request. The MAX-DOAS measurements used to validate the satellite SO₂ data are available on request from Drs. François Hendrick (Xianghe), Thomas Wagner, Vinod Kumar (Mohali).

AUTHOR CONTRIBUTIONS

N.T. prepared the manuscript and figures with contributions from all the coauthors. N.T., I.D.S., C.Le., L.C., J.V., H.B., M.V.R. contributed to the development of the COBRA algorithm, processing of the data and satellite comparison. V.F., C.McL., D.G. estimated the SO₂ emissions. C.Li, N.K. developed the TROPOMI and OMPS PCA algorithms and provided data for the comparison. P.H. and D.L. contributed to the development of the TROPOMI DOAS algorithm, processing of the data and satellite comparison. A.I. and R.R. provided CAMS SO₂ data. T. W., V.K., F.H., M.V.R. analyzed and provided MAX-DOAS data. All authors contributed to the interpretation of the results and improvement of the manuscript.

COMPETING INTERESTS,

The authors declare that they have no conflict of interest.

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