Interactive comment on "Anthropogenic and volcanic point source SO₂ emissions derived from TROPOMI onboard Sentinel 5 Precursor: first results" by Vitali Fioletov et al.

Anonymous Referee #1

The authors present in this paper first result for the estimation of SO2 emissions from point sources using TOPOMI data. They apply a well-documented methodology already applied in previous publications on other satellite data (e.g. OMI, OMPS). They demonstrate in this publication the impact of TROPOMI's high spatial resolution on the potential of the algorithm to estimate emissions from weaker sources as well as to finer separation of point sources. Issues relative to large scale biases are discussed and the results are compared with the ones estimated from OMI and OMPS data. The paper is well written and structured and should be accepted for final publication after considering my comments below.

We would like to thank the reviewer for the evaluation and comments that helped us improve the manuscript.

Page 5, Lime 11: The -3DU threshold is related to the spread of the distribution of the SO2 values? Please comment.

Apparently, large negative values are related to very rare specific conditions (probably due to extremally high HCHO amounts). To determine the threshold, the calculations were done for various cut-off limits from -1 DU to -5 DU. It was found that setting the limit below -3 DU changed the emission values to negative for rare sites where such values <-3DU occur. Rising the limit above -3DU was changing the emission estimates for most of the sites. Therefore the -3DU value was used. We added a sentence to address this issue:

"Lower than -3 DU threshold values produced negative emission values in some rare cases, while higher values affected the emission estimates themselves."

Page 5, lines 12-14: Probably the comma is misplaced, but generally this sentence needs rephrasing. It is not clear how this limit is set. Please elaborate here more. As it is now these limits seem arbitrary.

To make it clear, we changed the text to:
"To eliminate cases of transient volcanic SO2, days with high volcanic SO2 values were excluded from the analysis. If the highest 10% of SO2 values near the analyzed site were above a certain limit on a particular day, all data from the entire day were excluded. The limit depended on the emission strength and varies from 6 DU for sources emitting less than 100 kt per year to 15 DU for sources emitting >1000 kt per year (see Fioletov et al., (2016) for details)."

Page 6, Line1-9: The whole discussion on the AMF and the temperature dependence is confusing. Do the authors use a specially processed TROPOMI product for this paper (without temperature adjustment, but increased by 22%) and the official product is still the one where an AMF correction factor for the temperature dependence is applied?

To make it clear, we changed the text to:

"In this work, for consistency, we used TROPOMI SO₂ SCDs and converted them to VCDs using the same AMF as we utilized for OMI and OMPS (without any temperature adjustment). However, that meant that the obtained TROPOMI VCDs corresponded to 203 K as the original TROPOMI SCD were calculated for that temperature. To

remove the systematic difference with OMI/OMPS data caused by the difference in cross section temperature (203° K for TROPOMI vs. 293° K for OMI/OMPS), we increased the TROPOMI SO_2 VCDs by 22% (see Theys et al., (2017), their Figure 6, for justification)."

Page 6, line 24-27. Is there any possible explanation for this seasonality in standard deviation?

It likely has the same explanation as for the difference in the noise between low and high latitudes: low Sun means lower signals particularly due to a longer path through the ozone layer. We change the text to:

"Outside the tropical belt, there is also some seasonality in the standard deviation values with higher values occurring in winter and lower in summer (not shown) due to weaker signals at low Sun."

Page 7, line 5-10. Can the authors provide an explanation, why the standard deviations from TROPOMI are larger? Is it instrumental or a matter of spatial resolution differences?

A smaller size of TROPOMI pixels means that less photons can reach a single cell of the detector compared to OMI and OMIS. As mentioned, if TROPMI pixels are combined to match OMI/OMPS footprints, TROPOMI actually performs better (i.e., measurements have smaller uncertainties) than OMI/OMPS. We added that:

"As Figure 2 shows, the standard deviations (σ) for TROPOMI are roughly 1.5 time larger than OMI, and 3 times larger than OMPS since the TROPOMI footprint is smaller and each detector cell receives less photons than OMI and PMPS detector cells."

Page 10, lines 9-16: Is there any justification for the size of the rectangular where the fit is applied and its dependency on the source strength?

We added some additional explanation to the text:

"The parameter estimation was done using OMI pixels centered within a rectangular area that spreads ±L km across the wind direction, L km in the upwind direction and 3·L km in the downwind direction. As in the original study, the value of L was chosen to be 30km for small sources (under 100 kt yr^{-1}), 50km for medium sources (between 100 and 1000 kt yr^{-1}), and 90km for large sources (more than 1000 kt yr^{-1}). For small sources, different L values have little effect on the estimated parameters, but smaller values of L allow to separate individual sources where multiple sources are located in the same area. For larger sources, pixels with elevated SO2 values are located over larger areas and therefore the parameters estimated for higher L values have smaller uncertainties."

This manuscript provides the first SO2 emissions estimates for ~278 point sources using TROPOMI observations. The method applied has been used before on OMI and OMPS SO2 data. Results are compared with OMI and OMPS estimates, and specific issues related to the higher spatial resolution of TROPOMI and specific biases in TROPOMI are discussed. The paper is wel suited for this journal. I have the following comments.

We would like to thank the reviewer for the evaluation and comments that helped us improve the manuscript.

- p.5 l. 11-12 'To eliminate ..days then ..' somehow this sentence does not make sense to me.

It was a typo, it should be "when" instead of "then". Corrected.

- P.6 l. 8 what is meant with THIS systematic difference ?

We changed the text to:

"To remove the systematic difference with OMI/OMPS data caused by the difference in cross section temperature (203 K for TROPOMI vs. 293 K for OMI/OMPS), we increased the TROPOMI SO$_2$ VCDs by 22% (see Theys et al., (2017), their Figure 6, for justification). "

- P.6 l9 is the 22% the result of this different use of absorption cross sections at different temperatures ?

Yes. We edited that paragraph to make the cross-section temperature difference clear.

- P.6 l. 23 What do you mean with shorter exposure times at the edge of the swath. This is not correct I think. The SNR could be lower due to less binning across track, but that has nothing to do with shorter exp. times.

We agree with the reviewer and changed the text as suggested.

- P.8 if the biases are seasonal dependent what is its effect on the emissions calculated using annual means SO2 ?

The actual calculations were done for individual seasons. We added the explanation to page 10:

"To remove the local biases mentioned above, the average SO$_2$ VCD for the area located upwind from the source was calculated and then subtracted from the data. As the biases may be different from season to season, all calculation were done for 3-month periods (seasons) and then the annual emission rate was calculated by averaging seasonal emission rates. "

- Data availability: Should the locations (Fig. 9) and emissions (Fig. 10) not be made available through some database connected to the manuscript ? Otherwise other people can not use those.

We added a supplement with this information. Note that we also re-examined the set of emission sources and excluded four additional sources with the ratio of the estimated emission to its uncertainty below 4.

Anthropogenic and volcanic point source SO₂ emissions derived from TROPOMI onboard Sentinel 5 Precursor: first results

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Abstract.

The paper introduces the first TROPOMI-based sulfur dioxide (SO₂) emissions estimates for point sources. A total of about 500 continuously emitting point sources releasing from about 10 kT y^{-1} to more than 2000 kT y^{-1} of SO₂ per year, previously identified from Ozone Monitoring Instrument (OMI) observations were analysed using TROPOMI measurements for one full year, from April 2018 to March 2019. The annual emissions from these sources were estimated and compared to similar estimates from OMI and Ozone Mapping Profiling Suite (OMPS) measurements. Note that emissions from many of these 500 sources declined significantly since 2005 making their quantification more challenging. We were able to identify ~~278~~ 274 sources where annual emissions are significant and can be reliably estimated from TROPOMI. The standard deviations of TROPOMI vertical column density data, about 1 Dobson Unit (DU, where 1 DU = 2.69×10^{16} molecules/cm²) over tropics and 1.5 DU over high latitudes, are larger than those of OMI (0.6-1DU) and OMPS (0.3-0.4 DU). Due to its very high spatial resolution, TROPOMI produces 12-20 times more observations over a certain area than OMI and 96 times more than OMPS. Despite higher uncertainties of individual TROPOMI observations, TROPOMI data averaged over a large area have roughly two-three times lower uncertainties compared to OMI and OMPS data. Similarly, TROPOMI annual emissions can be estimated with uncertainties that are 1.5-2 times lower than the uncertainties of annual emissions estimates from OMI. While there are area biases in TROPOMI data over some regions that have to be removed for emission calculations, the absolute magnitude of these are modest, typically within ±0.25 DU, it can be comparable to SO₂ values over large sources.

1 Introduction

Sulfur dioxide (SO_2) is a major air pollutant that contributes to acid rain and aerosol formation, adversely affect the environment and human health, and impacts climate. Current and accurate information about SO_2 emissions is therefore required in modern air quality and climate models (e.g., Liu et al., 2018). The majority of SO_2 emissions are related to anthropogenic processes (e.g., combustion of sulfur-containing fuels, oil refining processes, metal ore smelting operations), although natural processes such as volcanic eruptions and degassing also play an important role. Information about emissions from SO_2 sources is not always available or up to date, and a sizable fraction of emission sources is even missing from conventional emission inventories (McLinden et al., 2016) with satellite measurements only now being used to fill this gap. Liu et al., (2018) demonstrated that merging such satellite-based emissions estimates with a conventional bottom-up inventory improves the agreement between the model and surface observations.

In the early 1980s, satellite measurements of backscattered radiation by the Total Ozone Mapping Spectrometer (TOMS) provided the first global estimates of SO_2 from large volcanic eruptions (Krueger, 1983). The TOMS instrument was capable of measuring backscattered solar ultraviolet (BUV) radiance at just several wavelengths. A hyperspectral instrument from the next generation (a UV-Visible imaging spectrometer), Global Ozone Monitoring Experiment (GOME) on the Earth Research Satellite 2 (ERS-2), launched in 1995, was able to detect major anthropogenic sources (Eisinger and Burrows, 1998; Khokhar et al., 2008). The launch of the Ozone Monitoring Instrument (OMI) aboard NASA's Earth Observing System Aura satellite (Levelt et al., 2017, 2006) with high spatial resolution, of up to 13 by 24 km^2 at nadir, but lower at the swath edges (de Graaf et al., 2016) started a new era in satellite air quality monitoring. Data from OMI as well as from Scanning Imaging Absorption Spectrometer for Atmospheric Chartography (SCIAMACHY), on the ENVISAT, the Global Ozone Monitoring Experiment-2 (GOME-2) on MetOp-A, -B, (Callies et al., 2000), and the Ozone Mapping and Profiler Suite (OMPS) aboard the NASA– NOAA Suomi National Polar-orbiting Partnership (Suomi NPP) were used to track SO_2 changes on the global and regional scales and estimate area- and point-source emissions (Carn et al., 2004, 2007; Fioletov et al., 2013; de Foy et al., 2009; Koukouli et al., 2016a, 2016b; Krotkov et al., 2016; Lee et al., 2009; Li et al., 2017; McLinden et al., 2012, 2014; Rix et al., 2012; Nowlan et al., 2011; Thomas et al., 2005; Zhang et al., 2017). Moreover, OMI measurements were used to evaluate the efficacy of cleantech solutions in reducing SO_2 emissions from industrial sources (Fioletov et al., 2013, 2016; Ialongo et al., 2018; Song and Yang, 2014).

There are two major types of UV-visible SO_2 retrieval algorithms for nadir viewing instruments. The traditional Differential Optical Absorption Spectroscopy (DOAS) scheme is based on the approach where absorption cross sections of relevant atmospheric gases are adjusted by a nonlinear least squares fit procedure to the log ratio of a measured earthshine spectrum and a reference spectrum in a given wavelength interval (Theys et al., 2015). The DOAS algorithm requires information about the absorption spectra of all trace gases, non-elastic Rotational Raman scattering (Ring effect) and information about the instrument characteristics. The uncertainties of the DOAS algorithm arise from inaccurate modeling of the various physical processes in solar light absorption and scattering (e.g., Ring effect, surface properties) as well as

artifacts in the radiance measurements (e.g., stray light, wavelength shift). An alternative approach is used in the principal component analysis (PCA) algorithm. Instead of attempting to model all various factors other than SO_2, the PCA algorithm replaces them with characteristic features derived directly from the measurements over locations where no SO_2 is expected (Li et al., 2013, 2017, 2019). When applied to OMI measurements, both DOAS and PCA algorithms produce similar results, however the PCA algorithm-based data show reduced data scattering and smaller biases comparerd to the DOAS algorithm-based data (Fioletov et al., 2016).

The launch of TROPOspheric Monitoring Instrument (TROPOMI) on board Copernicus Sentinel-5 Precursor in October 2017 made it possible to monitor atmospheric pollutants with an unprecedented spatial resolution, 3.5 by 7 km^2 (Veefkind et al., 2012), that is at least 12 times better that resolution of OMI. Since August 6th 2019 the spatial resolution was further reduced in the flight direction, the TROPOMI ground pixel size is now 3.5 by 5.5 km^2. It has already been demonstrated that TROPOMI can successfully monitor trace gases such as ozone (Garane et al., 2019), NO_2 (Griffin et al., 2019), HCHO (de Smedt et al., 2018), CO (Borsdorff et al., 2019), CH_4 (Hu et al., 2018), and even BrO (Seo et al., 2019) as well as cloud properties (Loyola et al., 2018). The operational TROPOMI SO_2 retrieval algorithm utilizes the DOAS approach (Theys et al., 2017) and early observations demonstrated the benefits of high spatial resolution for monitoring volcanic plumes (Hedelt et al., 2019; Theys et al., 2019; Queißer et al., 2019). However, these first studies were focused on relatively high volcanic SO_2 levels. In this study, we perform analysis of TROPOMI SO_2 observations that include smaller anthropogenic and volcanic degassing sources. We applied a previously developed technique (Fioletov et al., 2015) to estimate SO_2 emissions from TROPOMI observations. About 500 SO_2 sources, previously identified using OMI 2005-2015 data (Fioletov et al., 2016), were examined and their emissions were estimated using TROPOMI data and then compared to emission estimates from OMI and OMPS.

2 Data Sets

2.1 Satellite SO_2 VCD data

The TROPOMI instrument onboard of the Sentinel-5 Precursor (S5P) satellite was launched on October 13, 2017. TROPOMI has the smallest spatial footprint of 3.5 by 7 km^2 (3.5 by 5.5 km^2 after August 2019) among the instruments of its class (Veefkind et al., 2012). TROPOMI measures spectra of backscattered solar light at 450 cross-track positions (or pixels) and provides daily global coverage. TROPOMI SO_2 Level 2 (/PRODUCT/sulfurdioxide_total_vertical_column) data processed with the S5P operational processing system UPAS version 01.01.05 (Theys et al., 2017) were used in this study. In the first step of the algorithm, SO_2 slant column densities (SCDs), representing the effective optical-path integral of SO_2 concentration, were retrieved using the DOAS method. An additional background correction was applied to remove possible biases in SCDs after the spectral retrieval step. The spectral fitting was done using the 312–326 nm window, although two other spectral windows (325–335 nm and 360–390 nm) were used for retrievals in cases of very high volcanic SO_2. The final product, the SO_2 vertical column densities (VCDs) were calculated from SCDs using conversion factors (air mass factors).

VCDs represent the number of SO_2 molecules (or total mass) in an atmospheric column per unit area. VCDs are commonly reported in Dobson Units (DU) where 1 DU = 2.69×10^{16} molecules/cm². The standard TROPOMI SO_2 data product additionally includes VCDs calculated for three volcanic scenarios: when a 1 km thick plume is located at ground level, at 7 km and 15 km. In this study we are focussed on anthropogenic and degassing volcanic emissions and used only data corresponding to the ground level plume.

OMI, a Dutch-Finnish UV-Visible wide field of view nadir-viewing spectrometer onboard NASA's Aura satellite was launched on July 15, 2004 (Schoeberl et al., 2006). Originally, it was able to provide daily global coverage with the resolution up to 13 by 24 km^2 at nadir (de Graaf et al., 2016; Levelt et al., 2006), but now about a half of its pixels are affected by a field-of–view blockage and stray light (the so-called "row anomaly") and SO_2 cannot be retrieved successfully from those pixels. The OMI detector has 60 cross-track positions. In our previous studies (Fioletov et al., 2016; McLinden et al., 2016), we excluded data from the first 10 and last 10 cross-track positions from the analysis to limit the across-track pixel width from 24 km to about 40 km. However, due to row anomaly, this currently limits the number of available pixels to 15-20. We found that excluding only the first and the last 5 cross-track positions does not change the emission estimates noticeable but reduces their uncertainties, so only the first and last 5 pixels were excluded from the current analysis. NASA operational Planetary Boundary Layer (PBL) SO_2 Level 2 data product was used in this study (OMSO2; Li et al., 2019a). This data product is produced with the Principal Component Analysis (PCA) algorithm (Li et al., 2013; 2017). The 310.5–340 nm spectral window was used for SO_2 retrievals. Detailed information on the OMI PCA SO_2 data sets and its characteristics is available elsewhere (Krotkov et al., 2016; McLinden et al., 2015). It should be noted that OMI DOAS algorithm-based data product is also available (Theys et al., 2015). While the results of the two algorithms are somewhat different, particularly in large-scale biases, emission estimates from the two algorithms demonstrate very similar results (Fioletov et al., 2016).

OMPS Nadir Mapper on board the Suomi National Polar-orbiting Partnership (or Suomi NPP) satellite operated by NASA and NOAA was launched in October 2011. The standard NASA OMPS SO_2 data product (NMSO2-PCA-L2) is based on the same PCA algorithm as the NASA OMI data product (Li et al., 2019b; Zhang et al., 2017). OMPS has a lower spatial resolution than OMI, 50 km by 50 km, but better signal-to-noise characteristics. OMPS SO_2 VCD data are retrieved for 35 cross-track positions. Similar to OMI data analysis, large OMPS pixels at the edges of the swath (rows < 2 or > 33) were excluded. Both OMI and OMPS SO_2 data are retrieved with the same PCA algorithm, and emissions estimates for the two satellite instruments are similar, although OMPS tends to miss or underestimate emissions from small sources (Zhang et al., 2017).

Suomi NPP and S5P are on the same orbit 3.5 minutes apart and cross the Equator at about 13:30 local time. Aura is on a similar polar orbit and cross the Equator at about 13:45 local time. Therefore, we can assume that there is no difference in the measurements of the three satellite instruments related to diurnal variations of SO_2. TROPOMI operational SO_2 data record starts in April 2018. In order to have one full year of data, we analyzed TROPOMI, OMI, and OMPS data for the period from April 2018 to March 2019.

For emission estimates, we examined SO$_2$ values within a 300 km radius from each emission source listed in the SO$_2$ point source catalogue (Fioletov et al., 2016). There are about 500 sources in the catalogue, however many sources emitting SO$_2$ in the first years of OMI operation, were below the OMI sensitivity level in 2018: either closed or now produce substantially reduced emissions due to scrubber installation. The most recent version of the SO$_2$ emission catalogue is available from NASA public archive (Fioletov et al., 2019) and at https://so2.gsfc.nasa.gov/measures.html.

2.2 Air mass factors and data filtering

Data filtering was applied to OMI, OMPS, and TROPOMI SO$_2$ data before the analysis. The current retrieval algorithms are optimized for low (0.05) surface albedo, therefore pixels that corresponds to snow-covered high albedo surfaces were excluded from the analysis. Measurements taken at high solar zenith angles (more than 70°) were also excluded. Only clear-sky data, defined as having a cloud radiance fraction (across each pixel) less than 20% were used. Negative SO$_2$ values that are less than -3 DU were also excluded. Lower than -3 DU threshold values produced negative emission values in some rare cases, while higher values affected the emission estimates themselves. To eliminate cases of transient volcanic SO2, days with high volcanic SO2 values were excluded from the analysis. If the highest 10% of SO2 values near the analyzed site were above a certain limit on a particular day, all data from the entire day were excluded. The limit depended on the emission strength and varies from 6 DU for sources emitting less than 100 kt per year to 15 DU for sources emitting >1000 kt per year (see Fioletov et al., (2016) for details). ~~To eliminate cases of transient volcanic SO$_2$, periods when high SO$_2$ values were caused by eruptions, days then the highest 10% of SO$_2$ values were above a limit near the analysed site were excluded from the analysis. The limit depends on the emission strength and varies from 6 DU for sources emitting less than 100 kt per year to 15 DU for sources emitting >1000 kt per year.~~

Information on airmass factors (AMFs) is required to convert TROPOMI SCDs to VCDs. AMFs depend on SO$_2$ vertical profile shape, solar zenith angle, observation geometry, total ozone absorption, clouds, and surface reflectivity. In the operational TROPOMI dataset, TM5 model calculations were used to obtain a-priori SO$_2$ vertical profile to calculate AMF for each TROPOMI pixel. The model estimates rely on "bottom-up" emission inventories derived from economic activity data and SO$_2$ emission factors for known sources, so that in the case of a missing source in the inventory, the model SO$_2$ profile shape would be representative of clean background areas, causing calculated AMFs to be biased high and VCDs being underestimated over that source.

The PCA algorithm uses spectrally dependent SO$_2$ Jacobians instead of AMFs. To make it consistent to the previous operational OMI Band Residual Difference (BRD) algorithm, the present PCA algorithm assumes the same fixed conditions that correspond to a typical summertime conditions in the eastern USA and PCA retrievals can therefore be interpreted as having an effective AMF of 0.36 as in the BRD algorithm (Krotkov et al., 2006). However, a constant AMF does not represent conditions such as high elevations or enhanced aerosol loading. As in our previous studies (Fioletov et al., 2016; McLinden et al., 2016) a single site-specific AMF was calculated for each source (McLinden et al., 2014) and applied to both OMI/OMPS and TROPOMI estimated emissions.

As one of the main goals of this study is to compare TROPOMI SO$_2$ data and emissions estimates to those from OMI and OMPS, we used a constant AMF of 0.36 for illustration maps while for the emissions estimates, we converted TROPOMI SO$_2$ SCDs to VCDs using the same site-specific AMFs thereby removing it as a potential source of variability. It should be also noted that the spectral fitting window used in the TROPOMI algorithm is different from the window in the PCA algorithm. However, we estimated that that effect is small (under 10%) compared to other sources of uncertainties.

The SO$_2$ absorption cross section has a moderate temperature dependence, with absorption increasing for higher temperatures, and there is a difference in how this dependence was handled in TROPOMI and OMI/OMPS retrievals. In the TROPOMI spectral fit, an SO$_2$ cross section at 203°K was used and then the retrieved VCDs were adjusted by applying an AMF correction factor using temperature from the European Centre for Medium-Range Weather Forecast (ECMWF) operational model (Theys et al., 2017). The OMI/OMPS retrieval algorithm uses the SO$_2$ cross section at 293°K (Krotkov et al., 2006) without any adjustment. In this work, for consistency, we ~~converted~~ used TROPOMI SO$_2$ SCDs and converted them to VCDs using the same AMFs ~~approach~~ as we utilized for OMI and OMPS (without any temperature adjustment). However, that meant that~~, meaning~~ the obtained TROPOMI VCDs ~~now~~ corresponded to 203° K as the original TROPOMI SCD were calculated for that temperature. To remove ~~this~~ the systematic difference with OMI/OMPS data caused by the difference in cross section temperature (203° K for TROPOMI vs. 293° K for OMI/OMPS), we increased the TROPOMI SO$_2$ VCDs by 22% (see Theys et al., (2017), their Figure 6, for justification).

2.5 Wind and snow data

The emission estimation algorithm requires wind data. As in several previous studies (Fioletov et al., 2015; McLinden et al., 2016), European Centre for Medium-Range Weather Forecasts (ECMWF) reanalysis data (Dee et al., 2011) (http://apps.ecmwf.int/datasets/) were extracted for every satellite pixel. Wind profiles are available every 6 hours on a 0.75° horizontal grid and are interpolated in time and space to the location of each satellite pixel center. U- and V- (west-east and south-north, respectively) wind-speed components were averaged for 1-km thick layers and the winds for the layer that corresponds to the site altitude was used. The Interactive Multisensor Snow and Ice (IMS) Mapping System data (Helfrich et al., 2007) were used to screen out pixels over snow-covered surface with high albedo.

3 TROPOMI SO$_2$

For brevity, from this point we refer to "SO$_2$ VCD" as simply to "SO$_2$". It can be expected that a smaller pixel size of TROPOMI would yield a lower signal to noise level. Figure 1 shows the standard deviation of SO$_2$ values at four sites, each located at different latitudes, as a function of the TROPOMI cross-track position. The selected sites have relatively small SO$_2$ emissions, so the standard deviations are determined by the instrumental noise and possible retrieval uncertainties. The standard deviations at the 20 cross-track positions at the edges of the swath are particularly high due to less binning across track ~~shorter exposure time~~, which motivated our decision (in addition to a larger footprint) to exclude them from the

analysis. There is also a clear increase in the noise from low to high latitudes with the noise standard deviations at a sub-polar site nearly double compared to tropical sites. Outside the tropical belt, there is also some seasonality in the standard deviation values with higher values occurring in winter and lower in summer (not shown) due to weaker signals at low Sun.

The standard deviation of SO_2 retrievals for the three satellite instruments as a function of latitude is shown in Figure 2 for the period from April 2018 to March 2019. The plot is based on satellite measurements over clean areas (150 km-300 km distance from the catalogue source locations) and represent background noise levels of SO_2. Large sources with annual SO_2 emissions above 1000 kt per year where the high standard deviations are likely to be influenced by the SO_2 variability itself were excluded from this analysis. Sources inside the South Atlantic Anomaly (SAA) region were also excluded. The standard deviations of TROPOMI data (about 1 DU over tropics and 1.5 DU over high latitudes) are larger than those of OMI (0.6-1DU) and OMPS (0.3-0.4 DU) data. The standard deviations are particularly large (1.6-2.2 DU) for the first and the last 20 pixels in the TROPOMI 450 pixels-wide swath, which were excluded from further analysis.

As Figure 2 shows, the standard deviations (σ) for TROPOMI are roughly 1.5 time larger than OMI, and 3 times larger than OMPS since the TROPOMI footprint is smaller and each detector cell receives less photons than OMI and PMPS detector cells. However, the pixel size for TROPOMI is much smaller, and so the number of observations (n) over the same area for TROPOMI is 12 and 96 times that of OMI and OMPS, respectively. Considering these two factors, and assuming the standard error is proportional to $\sigma/\sqrt{n}$ (assuming that the errors of individual pixels are not correlated), then the uncertainty of a TROPOMI average will be roughly a factor of 2 smaller than OMI and a factor of 3 smaller than OMPS In fact, due to the OMI row anomaly, the number of TROPOMI pixels over the same area is now a factor of 20 larger.

The global distribution of mean SO_2 from TROPOMI (smoothed using oversampling technique or pixel averaging technique with a 30 km radius, see e.g., Fioletov et al., (2011), Sun et al., (2018)) is very similar to that from OMI and OMPS (Figure 3). All three instruments clearly show elevated values over the Persian Gulf, China, Mexico, and India, as well as many anthropogenic "hotspots" such as Norilsk (Bauduin et al., 2014; Khokhar et al., 2008), a cluster of power plants in South Africa, and large volcanic sources such as Kilauea, Hawaii, and Ambrym, Vanuatu. All three satellite data sets shown in Figure 3 do not demonstrate large biases seen in the older versions of OMI, GOME-2, and SCIAMACHY data (see Fioletov et al., (2013), their Figure 1). Except for the hotspot-affected areas, SO_2 values from all three instruments are typically within the ±0.25 DU range. It is also interesting to note that the South Atlantic Anomaly (SAA), an area of increased flux of energetic "solar wind" particles that may intercept instruments in low-Earth orbits such as these, significantly increases the uncertainties of OMI and OMPS data (as well as data from GOME-2 and SCIAMACHY), but has little effect on TROPOMI data.

There are, however, still some differences in the absolute values between OMI, OMPS, and TROPOMI over some regions. Zoomed-in plots of mean SO_2 over four regions of elevated SO_2 values, Northern China, India, Mexico, and Iran are shown in Figure 4. TROPOMI SO_2 means are, in general, higher than OMI and OMPS values over these regions suggesting possible biases in TROPOMI data. The spatial scale of these biases (thousands of km) is larger than the scale of elevated SO_2 values from a typical industrial source (one-two hundreds of km), so we will call them "large-scale biases". Note that the

biases are very small, only 0.1-0.2 DU, however even such small biases could affect emission estimates since the SO_2 enhancements from many sources are really tiny, a few tenths of a DU. These large-scale biases are common in satellite SO_2 retrievals. Their magnitude often depends on the retrieval algorithm and the same satellite measurements (i.e., calibrated level 1B data) processed with different SO_2 algorithms produce different biases. For example, GOME-2 data processed with

5   the original operational algorithm (Valks and Loyola, 2008) had larger biases than the SO_2 data product based on a direct fitting method developed by the Harvard-Smithsonian Center for Astrophysics, Cambridge, Massachusetts, (Nowlan et al., 2011): see Figure 1 in Fioletov et al., (2013). The origin of such biases is not always known although an imperfect removal of the very strong ozone absorption, that itself depends on stratospheric temperature and shape of the ozone profile, could be one of the contributing factors.

10   OMI data processed with a DOAS algorithm (Theys et al., 2015) that is similar to the present TROPOMI algorithm, also had larger biases over some areas than seen in the PCA-based data (Fioletov et al., 2016). However, as was also noted by Fioletov et al., (2016), both algorithms produce very similar results if the large-scale biases are removed, for example, by comparing up-wind and down-wind values around an SO_2 emission source. In case of large-scale biases in the area with multiple sources, the bias can be accounted for by introducing functions that change slowly with latitude and longitude as

15   suggested by Fioletov et al., (2017). This multi-source algorithm accounts for the bias using Legendre polynomials of latitude and longitude and their products and the emissions using functions that represent plumes from individual sources. As an example, Figure 5 shows original data from TROPOMI, OMI and OMPS over Europe and the same data with the local bias removed using 6th degree polynomials (see Fioletov et al., (2017) for details). As Figure 5 suggests, large-scale biases seen in the original TROPOMI data are removed by this statistical fitting procedure. Note that OMPS data also show some

20   large-scale biases over that region. The maps with the large-scale bias removed look very similar for all three satellite data sources and all the major SO_2 hotspots are clearly seen. Note that there is practically no bias in OMI data over southern Europe hence OMI data with and without bias removed appear very similar.

    The problem of the bias in TROPOMI data as well as in data from other satellites requires further investigation and probably improvements of the SO_2 algorithms. In case of TROPOMI, we saw such biases over many major areas of interest:

25   China, India, Europe, the Persian Gulf. The biases are often larger than the signals from emission sources that creates an impression that TROPOMI values over such sources are larger than those from OMI. It also appears that the biases are larger in winter-fall than in summer and are also larger over water. It will be possible to investigate time dependence of these biases as more TROPOMI data become available.

    As mentioned, the uncertainties of TROPOMI data averaged over a certain area are 2-3 times smaller than those for

30   OMI and OMPS. Due to its very high spatial resolution, a single year of TROPOMI can provide as much information on the SO_2 distribution around hotspots as compared with several years of OMI or OMPS. Figure 6 (top) shows the mean SO_2 over Bosnia & Herzegovina and Serbia in 2018 from OMI and TROPOMI and over the 2014-2018 period from OMI, using oversampling technique (see Sun et al. (2018) and references therein). In these countries, SO_2 emissions were not under the same strict emission cutting regulations as in the EU countries. Emissions from the power plants shown in Figure 6 remained

nearly constant in 2014-2018. A simple version of the oversampling technique was applied where a geographical grid is established around the source and the mean value of all satellite pixels centred within a 30 km radius from each grid point is calculated. As the mean is calculated, the standard error of the mean can be also calculated and used to evaluate the significance of that mean value by analysing the ratio of the mean value to its standard error. Figure 6 shows both, the mean values (the top row) and the ratios (the bottom row). Although individual TROPOMI SO_2 values are noisier than for OMI, the much larger volume of TROPOMI data contributing to the mean makes it appear less noisy than a one-year OMI map and only a five-year OMI average demonstrates a TROPOMI-like level of noise. This is further confirmed by the ratio maps (Figure 6, bottom): TROPOMI one-year ratios are as high as 25, while OMI one-year ratios are under 10 and only five-year ratios are close to those for one-year TROPOMI values.

Although averaging of multiple years of OMI data can produce the same or even higher signal to noise ratio as one year of TROPOMI data, OMI cannot provide the same level of detail as TROPOMI due to the difference in the instrument spatial resolutions. The high spatial resolution of TROPOMI also makes it possible to see individual sources in areas where multiple sources are in a close proximity. As an example, Figure 7 shows the mean SO_2 over a cluster of power plants in South Africa using one year of TROPOMI data and the entire (2005-2019) available record of OMI. For this plot, the pixel averaging with a 10 km radius was used (smaller radiuses make the OMI map too noisy to see individual sources). Although we used a very small radius for averaging, it is hard to distinguish individual sources in the OMI map, while on the one-year TROPOMI map they appear as local maxima or "hotspots".

High TROPOMI spatial resolution makes it possible in some cases to resolve an individual, persistent SO_2 plume. As an example, the mean SO_2 over Hawaii for the period from April 2018 to March 2019 is shown in Figure 8. The source, Kilauea volcano, is located at 1200 m above sea level, while the mountains north and north-west of the volcano are as high as 4000 m. The area is dominated by easterly winds. TROPOMI data demonstrate that, on average, elevated SO_2 values are not observed above the volcano peak. This means that the symmetrical modified Gaussian plume model used for emissions calculations may not describe the actual plume very well in this particular case. OMI data with their lower spatial resolution do not really show these features of the SO_2 distribution.

4 Emissions estimates

A method developed to estimate emissions from point sources from OMI data (Fioletov et al., 2015) was applied here to TROPOMI, OMI, and OMPS data. The method is based on a fit of satellite data to an empirical plume model developed to describe the SO_2 spatial distribution near emission point sources. First, satellite measurements are merged with wind data and the rotation technique applied (Pommier et al., 2013; Valin et al., 2013), so the satellite data can be analysed assuming that the wind always has the same direction. Then, emissions and lifetimes were estimated using the exponentially modified Gaussian fit (Beirle et al., 2014; Fioletov et al., 2015; de Foy et al., 2015) appropriate for a near-point source. The fitted plume model depends on three parameters, total mass (α) near the source, the lifetime or, more accurately, decay time (τ),

and the plume width (σ). Finally, the emission strength (E) is calculated from τ by $E=\alpha/\tau$. For each source, all three parameters can be derived from a fit using a nonlinear regression model, but when doing so the uncertainties in the non-linear parameters (τ and σ) are often large. To minimize this uncertainty, all emissions were derived using a mean τ and σ determined by averaging over values obtained from the non-linear fits. Thus, only one parameter (α) is derived from the fit

5 that turns the algorithm into a simple linear regression model (Fioletov et al., 2016).

The three parameters for each of the three satellite instruments were estimated using April 2018 – March 2019 data. It can be expected that that the lifetime τ that characterizes the plume decay is the same for all three instruments. Indeed, we found that the average value of τ is about 6 hours for all three of them. The plume width σ depends on the instrument pixel size and is expected to be different. We estimated that, as in the previous study (Fioletov et al., 2016), σ is about 20 km for

10 OMI. For OMPS with its larger pixels, the average σ value is about 25 km. For TROPOMI, the average value of σ is about 15 km. However, many SO_2 sources are not really "point sources". Industrial sources are often comprised of several individual facilities located a few km apart. For example, in Norilsk, there are three major smelting factories located 8-10 km apart. For relatively large OMI pixels, this typically does not affect σ calculations. For TROPOMI, however, we can see that for real "point" sources σ is smaller, about 10 km, than for sources with multiple facilities. Our sensitivity study

15 suggests that a change of sigma from 15 km to 10 km reduces the emission estimates by about 20%. A better characterization of the emission sources will be required in the future in order to improve emission estimates for sources with multiple facilities.

The calculations were performed in the same manner as the original study for OMI data (Fioletov et al., 2016). The parameter estimation was done using OMI pixels centered within a rectangular area that spreads $\pm L$ km across the wind

20 direction, L km in the upwind direction and $3 \cdot L$ km in the downwind direction. As in the original study, the value of L was chosen to be 30km for small sources (under 100 kt yr^{-1}), 50km for medium sources (between 100 and 1000 kt yr^{-1}), and 90km for large sources (more than 1000 kt yr^{-1}). For small sources, different L values have little effect on the estimated parameters, but smaller values of L allow to separate individual sources where multiple sources are located in the same area. For larger sources, pixels with elevated SO_2 values are located over larger areas and therefore the parameters estimated for

25 higher L values have smaller uncertainties. ~~To avoid impacts of local biases on emission estimates, the mean SO_2 upwind from the source was subtracted from the data. To minimize interference from other sources, the fitting was done for a rectangular area that for small sources (<100 kt SO_2 yr^{-1}) spreads ±30 km across the wind direction, 30 km in the upwind direction and 90 km in the downwind direction. For large sources (>1000 kt SO_2 yr^{-1}), these numbers are 90 km, 90 km and 270 km respectively (.~~ Only pixels with associated wind speeds between 0.5 and 45 km h^{-1} were used for the fitting. The

30 overall uncertainty of the method is about 50%. There are several factors that contribute to the emission estimate uncertainty, however the major contributors, uncertainties in A~~F~~Ms and τ, appear as scaling factors that affect TROPOMI, OMI, and OMPS-based estimates the same way. To remove the local biases mentioned above, the average SO_2 VCD for the area located upwind from the source was calculated and then subtracted from the data. As the biases may be different from season

to season, all calculation were done for 3-month periods (seasons) and then the annual emission rate was calculated by averaging seasonal emission rates. Additional information about the algorithm and uncertainly analysis can be found in (Fioletov et al., 2016).

We examined all sources listed in the catalogue (Fioletov et al., 2016) and calculated emissions for the period from

5   April 2018 to March 2019 and their uncertainties using data for the three satellite instruments. It should be mentioned that although the catalogue contains about 500 sources, many were either closed or their emissions declined significantly due to several possible factors such as the installation of scrubbers and reduction in coal consumption. This includes most of the sources in the USA, European Union and many sources in China. Volcanic degassing emissions also vary with time (Carn et al., 2017) and some of the volcanoes that were active at the beginning of OMI operations did not emit high amounts of SO_2

10 in 2018-2019. Therefore, a decline in the number of catalogue sources detectable by TROPOMI is not entirely unexpected. The map of catalogue sources that are detectable from one year of TROPOMI data is shown in Figure 9 and their coordinates and emission levels are given in the Supplement. The following criteria were used to identify a source as detectable: (a) the sources should have an emission to uncertainty ratio exceeding 5 or (b) being between 4 3.6 and 5, but with a clear "hotspot" at the source with a downwind tail. There are only 20 sites in the (b) category, and we examined them on the case-by-case

15 basis. A total of 278 274 sites including 150 147 Power Plants, 19 Smelters, 41 40 Oil and Gas industry-relates sources, and 68 volcanoes with annual emissions from 10 to 2000 kt SO_2 yr^{-1} that satisfy these conditions were detected.

Scatter plots of TROPOMI, OMI, and OMPS-based emission estimates for all SO_2 catalogue sites are shown in Figure 10 a and b. Emissions estimates from OMI are on the horizontal axis of the both panels. Both OMPS and TROPOMI emissions estimates show a good agreement with OMI estimates for sources with estimated emissions above 50-60 kt SO_2

20 yr^{-1} (calculated as an average of emission estimates from the three instruments). For them, the correlation coefficients are about 0.97 for both instruments. However, the correlation coefficient is only 0.3 if calculated just for sources that emit less than 60 kt SO_2 yr^{-1}. There is practically no systematic biases between estimates from the three instruments.

Not surprisingly, statistical uncertainties of the emissions estimates from the three satellite instruments are also highly correlated (Figure 10 c and d). In general, the OMPS-based emission uncertainties are slightly larger than those based

25 on OMI data. The OMI-based emission uncertainties are almost always larger than those from TROPOMI data.

The relative TROPOMI emission uncertainties are lower than those from OMI. To illustrate that, Figure 10 e and f show scatter plots of the ratios of their signal to uncertainty ratios. For very large sources (>1000 kt yr^{-1}), the emission to uncertainty ratio is dominated by SO_2 variability, not by the noise in satellite data. For example, SO_2 emissions from volcanic sources could be very different from day to day. Even if the emissions are fairly constant, different weather

30 conditions (e.g., dry conditions vs. rain) affect the SO_2 dispersion patterns observed by satellites. For these very large sources, such SO_2 variability is larger than instrumental errors and the emission to uncertainty ratio is nearly the same for all three instruments. For smaller sources, however, measurement uncertainties play a bigger role. For OMPS, the ratios are mostly below the 1:1 line meaning that the uncertainties of OMPS-based emission estimates are higher than those based on OMI data. It is opposite for TROPOMI, where the ratios are mostly above the 1:1 line. Moreover, for medium-size and small

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sources, the ratios group around the 1.5:1 and 2:1 lines meaning that the TROPOMI emission estimate uncertainties are 1.5-2 times lower than those for OMI.

As all three satellite data sets can provide relatively independent emission estimates, the present satellite based SO_2 emission inventory could be further improved by combining emission estimates from the three sources. Due to its high resolution, and hence lower detection limit, TROPOMI can potentially identify many more sources than OMI and OMPS and then obtain emission estimates for them. An exhaustive analysis of this is beyond the scope of this paper. However, as an example of the sizable advantage offered by TROPOMI, Figure 11 shows the mean TROPOMI SO_2 distribution (from April 2018 to March 2019) at the border between Iran and Turkmenistan. The biggest source is the Khangiran gas refinery (1), an Iranian source that is included in the Catalogue. The second largest source is located near Mary, Turkmenistan (2), is related to gas exploration. The LANDSAT satellite images show that the source was build in 2012-2014. While one year of OMI data shows a signal from that region, they hardly can point to the source location. TROPOMI data clearly show a hotspot in both mean SO_2, as shown, and the high signal-to-noise ratio (not shown), right at the source location. Moreover, there are two other sources that can be resolved by TROPOMI. One of them, located east of Khangiran, could be related to two power plants (Toos and Ferdosi (3)) that are 1 km apart. This source can also be used as an illustration of the difference in emission uncertainties between TROPOMI and OMI/OMPS. TROPOMI-based emission estimates for this source is 14 kt SO_2 yr^{-1} with the standard error of 2.8 kt yr^{-1}, five times lower than the emission strength itself. The standard errors of OMI and OMPS-based emission estimates are 6.1 kt yr^{-1} and 7.1 kt yr^{-1} respectively, 2-3 times that of TROPOMI.

5 Summary and discussion

The first analysis of TROPOMI near-surface SO_2 for the period from April 2018 to March 2019 reveals global distributions and features very similar to those seen from OMI and OMPS: elevated values over the Persian Gulf, India, China; major "hotspots" over Norilsk, Russia; South Africa; major volcanic persistent sources such as Kilauea, Hawaii, and Ambrym, Vanuatu. Outside the areas affected by these "hotspots" all three instruments typically demonstrate low background SO_2 values within ±0.25 DU.

Over clean areas the spatial standard deviations of TROPOMI data (about 1 DU over tropics and 1.5 DU over high latitudes) are larger than those of OMI (0.6-1 DU) and OMPS (0.3-0.4 DU) data. However, despite higher uncertainties of individual TROPOMI pixels, spatially averaged TROPOMI data over respective field-of-views have uncertainties that are 2-3 times smaller than that from OMI and OMPS data. As a result, annual mean SO_2 maps smoothed by a spatial filtering appears less noisy than corresponding OMI maps. In terms of the signal to noise ratio, TROPOMI smoothed one-year mean values have the same uncertainties as 4-5 years of smoothed mean values based on OMI data.

We tested about 500 SO_2 sources previously detected from OMI data in 2005-2015, however, many of these sources emitted much less SO_2 in 2018-2019 than in the beginning of OMI operation. That includes, for example, almost all US

sources, many sources in Europe and China. We were able to identify 278 274 sources where annual (from April 2018 to March 2019) emissions can be estimated from TROPOMI data. Their emissions are in the range from 10 to 2000 kt SO_2 yr^{-1}.

Currently TROPOMI is able to provide point-source SO_2 emission estimates that have 1.5-2 times lower uncertainties than those from OMI, but it is less than expected taking into account that the number of useful TROPOMI pixels is about 20 times higher than that of OMI. If the statistical uncertainties are inversely proportional to the square root of the number of averaged pixels and their uncertainties are the same, then it is expected that the standard errors for TROPOMI emissions estimates should be about 4-5 times lower than those for OMI. Taking into account that the SO_2 uncertainties of individual TROPOMI pixels are 1.5-2 times larger than those for OMI one can expect that TROPOMI emission uncertainties would be 2.5-3.3 times lower than those for OMI, higher than the values of 1.5-2 that we derived directly from emission estimates. This may suggest that errors in individual TROPOMI pixels are correlated, for example, due to large-scale biases.

There are larger scale spatial biases in TROPOMI data over some areas that appear to be larger than similar biases in OMI and OMPS data. While the absolute magnitude of the biases is not very large, 0.1-0.2 DU, it can be comparable to the SO_2 enhancements over large sources. Due to these biases, SO_2 values over some sources may appear larger in TROPOMI data than in OMI and OMPS data. If, however, such biases are removed by, for example, a statistical fitting procedure, TROPOMI annual mean SO_2 maps are very similar to OMI and OMPS. It also appears that these TROPOMI biases have a larger amplitude in winter and fall, although it is hard to say that this is a repeatable seasonal effect based on just one year of data.

Biases are very common in early versions of all satellite SO_2 products and currently their origin is still not completely clear. The very small SO_2 absorption signal in the UV needs to be detected against a large contribution from the ozone absorption. The latter is a strong function of stratospheric temperature (and hence ozone profile). Any imperfection in any of these parameters may yield a bias in retrieved SO_2. This, however, does not explain biases in the tropical region where ozone variability is low. Development of a PCA-type algorithm for TROPOMI could help reducing these biases and improve the overall quality of the data. An improved version of the TROPOMI processing algorithm that includes a PCA component may produce a data product with smaller biases and lower noise than the present operational algorithm. Such improved algorithm is now under development. Preliminary test TROPOMI SO_2 retrievals applying a PCA-based algorithm have shown some promises, and work is underway to better understand algorithmic differences between DOAS and PCA.

6 Data availability

OMI and OMPS PCA SO_2 data used in this study have been publicly released as part of the Aura OMI Sulfur Dioxide Data Product (OMSO2) and can be obtained free of charge from the Goddard Earth Sciences (GES) Data and Information Services Center (Li et al., 2019 a, b). TROPOMI data are freely available from the European Union Copernicus Sentinel 5p data hub (https://s5phub.copernicus.eu; http://www.tropomi.eu).

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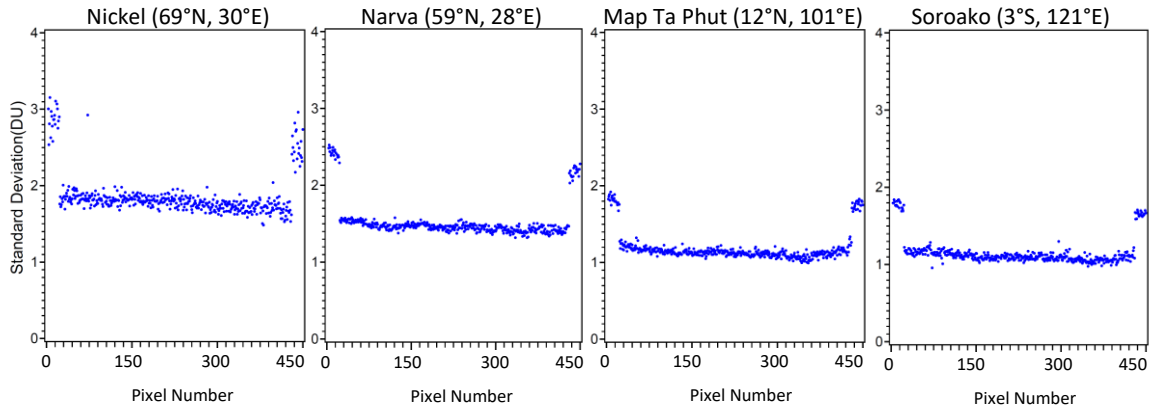
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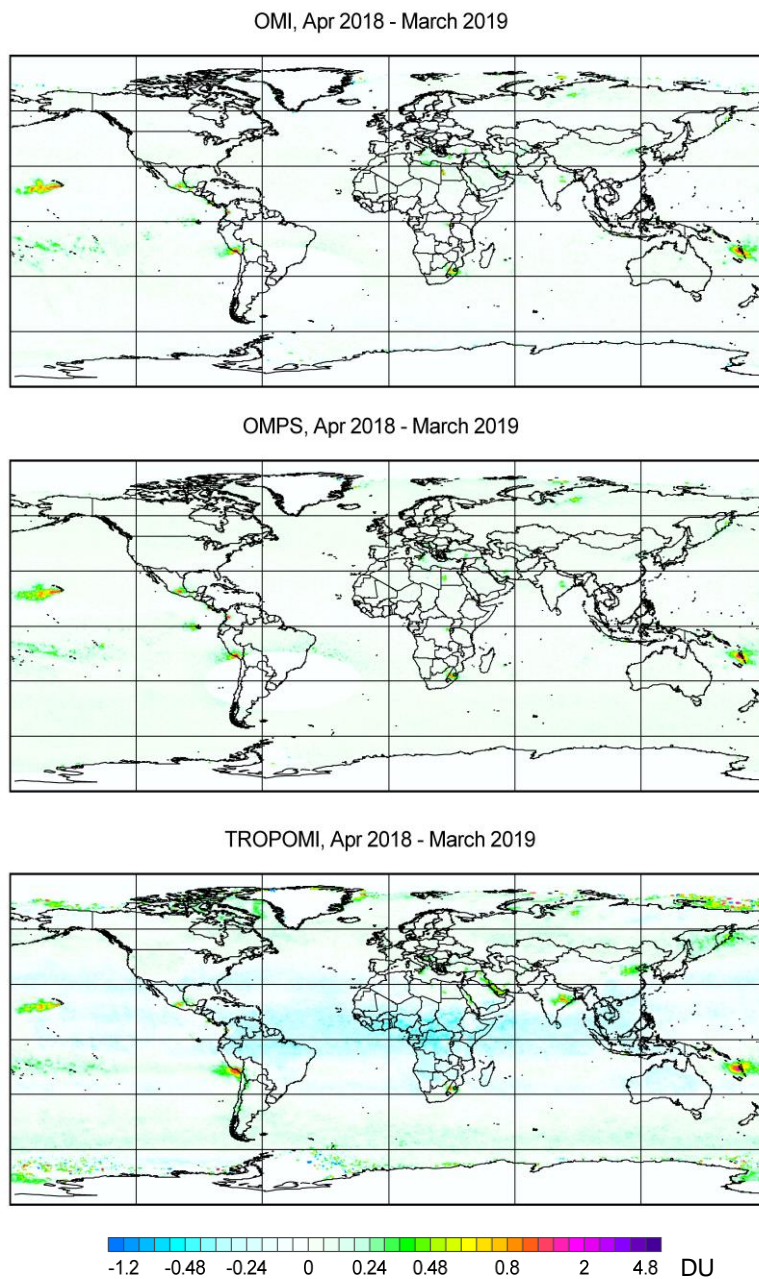
Figure 1. The SO$_2$ standard deviation as a function of the TROPOMI cross-track position (pixel number) at four sites illustrate a decline from high to low latitudes for the period from April 2018 to March 2019. The selected four sites represent sources with very low SO$_2$ emissions and therefore the standard deviations represent the measurement uncertainties. The SO$_2$ values retrieved at the first and last 20 pixels have noticeably higher standard deviations and are excluded from the analysis.
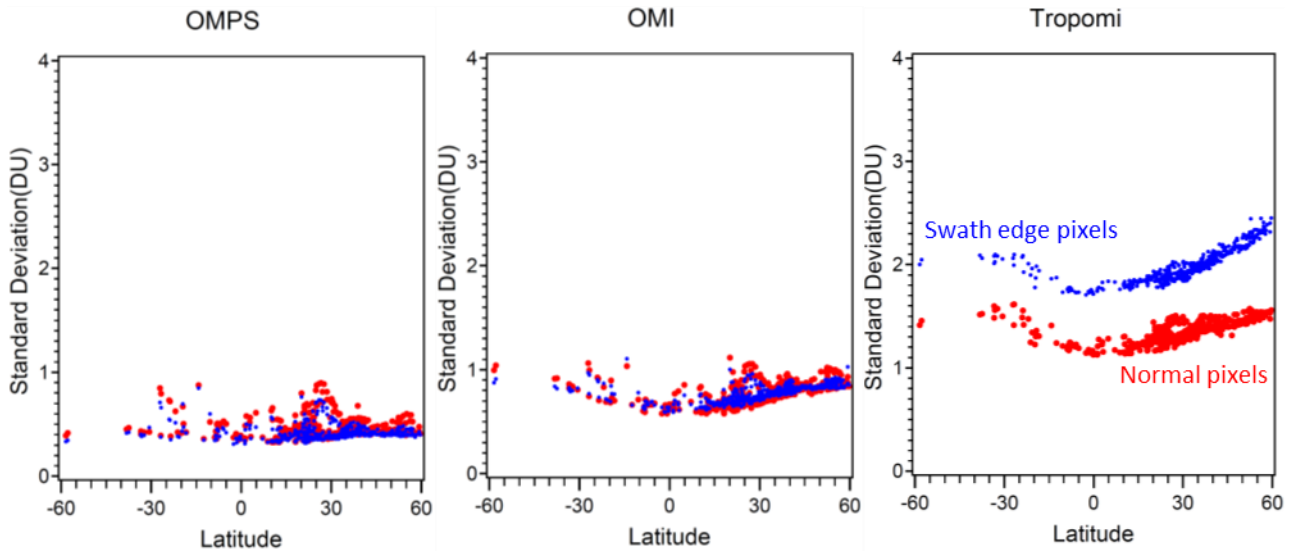
Figure 2. The OMPS, OMI, and TROPOMI SO$_2$ standard deviation vs. latitude for "normal" (red) and "swath edge" (blue) pixels. "Swath edge" pixels were defined as the first and the last 3, 5, and 20 pixels for OMPS, OMI, and TROPOMI respectively. All other pixels were considered as "normal". The plot is based on satellite measurements centred between 150 km and 300 km around the sources from the catalogue. The sources under the South Atlantic Anomaly (SAA) are excluded.

OMI, Apr 2018 - March 2019

OMPS, Apr 2018 - March 2019

TROPOMI, Apr 2018 - March 2019

-1.2 -0.48 -0.24 0 0.24 0.48 0.8 2 4.8 DU

Figure 3. Mean SO$_2$ (DU) over the globe from TROPOMI, OMI, and OMPS for the period from April 2018 to March 2019. Data are smoothed by oversampling technique with radius R=30 km. The area of the South Atlantic Anomaly (SAA) is left blank on the OMI and OMPS maps. The SAA largely increases uncertainties of OMI and OMPS SO$_2$ data but has a much smaller effect on TROPOMI SO$_2$ data.
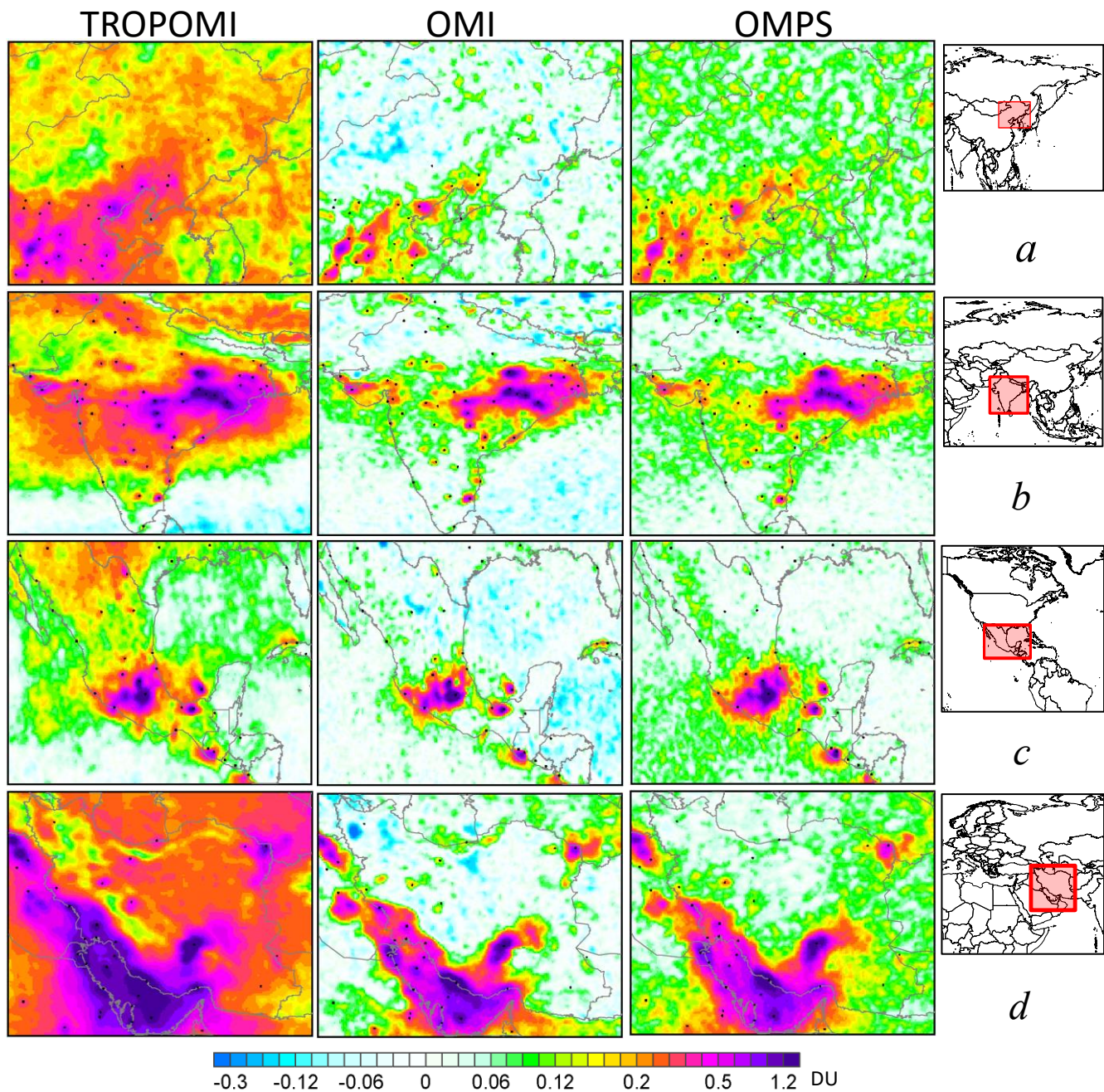
Figure 4. Mean SO$_2$ (DU) from TROPOMI, OMI, and OMPS over northern China (a), India (b), Mexico (c), and Iran (d) in the period from April 2018 to March 2019. Large-scale biases make it difficult to interpret Tropomi SO$_2$ data and compare them to OMI and OMPS directly. Data are smoothed by oversampling technique with radius R=30 km. The black dots indicate the SO$_2$ sources. Note that the color scale is different from the scale in Figure 3.
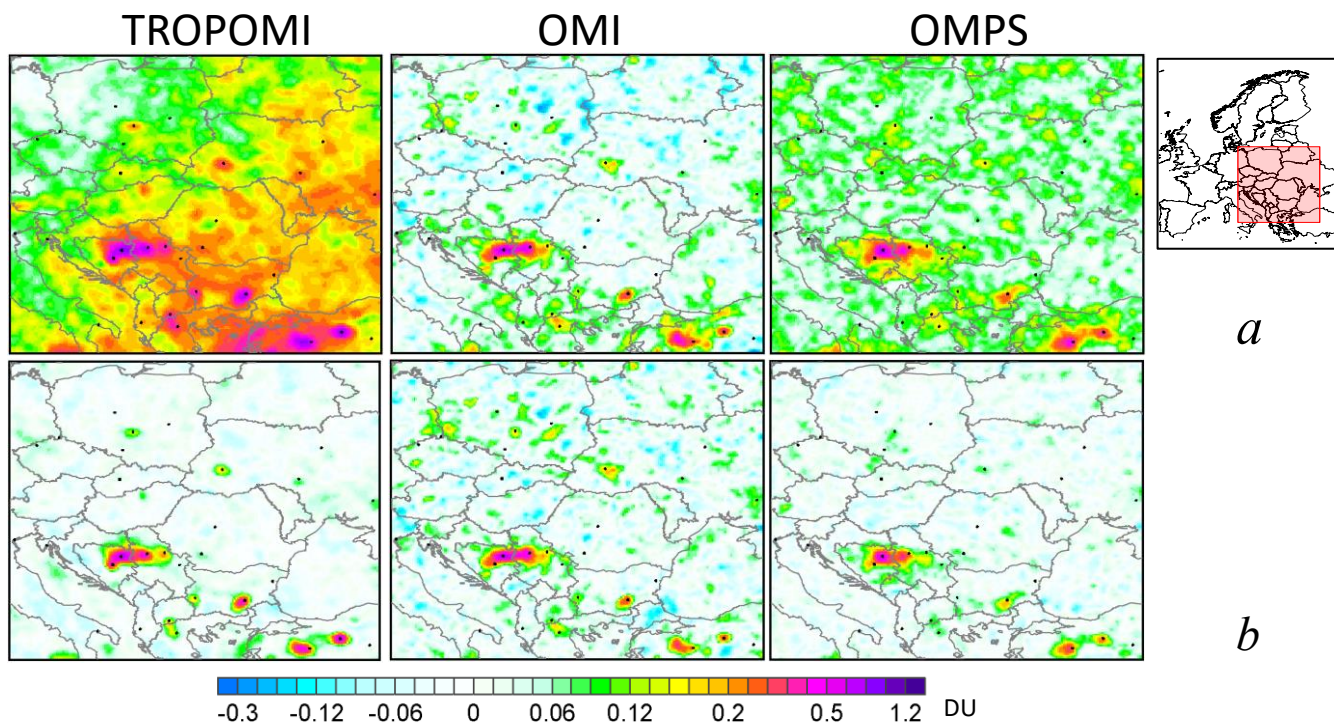
Figure 5. TROPOMI, OMI and OMPS mean SO$_2$ over Eastern and Southern Europe in the period from April 2018 to March 2019 (a). The same data, but with large-scale bias removed (b). Data are smoothed by oversampling pixel averaging with R=30 km radius; the black dots indicate the SO$_2$ emission sources.

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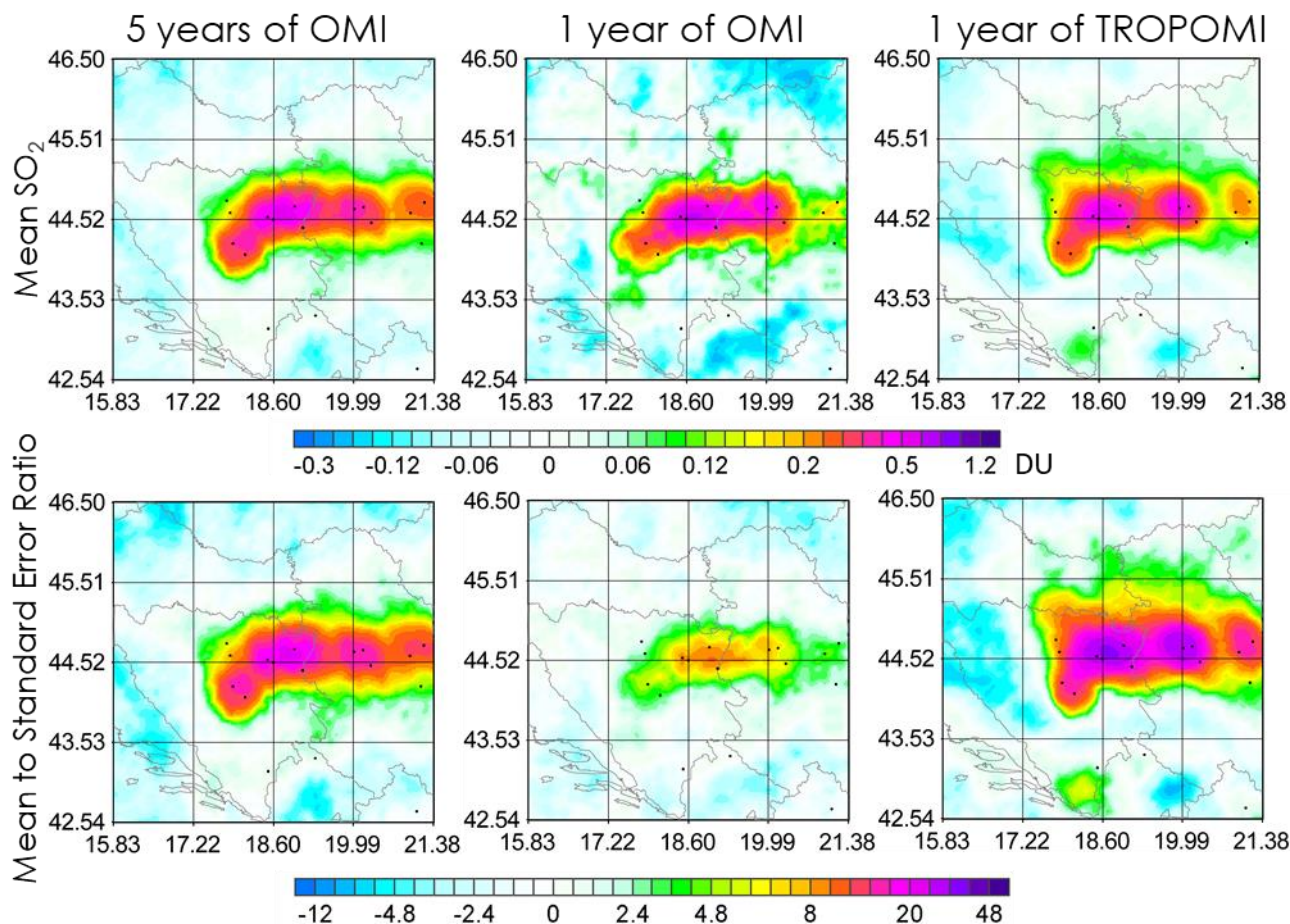
5 **Figure 6.** SO$_2$ hotspot over Serbia and Bosnia and Hercegovina: mean OMI values over a 5-year period (2014-2018) and mean OMI and TROPOMI values over 1-year period (from April 2018 to March 2019). Data are smoothed by oversampling/pixel averaging with 30 km radius; constant bias is removed; the black dots indicate the emission sources. The top panels show mean SO$_2$, the bottom panel show the ratios of the mean SO$_2$ value to the standard error of the mean.
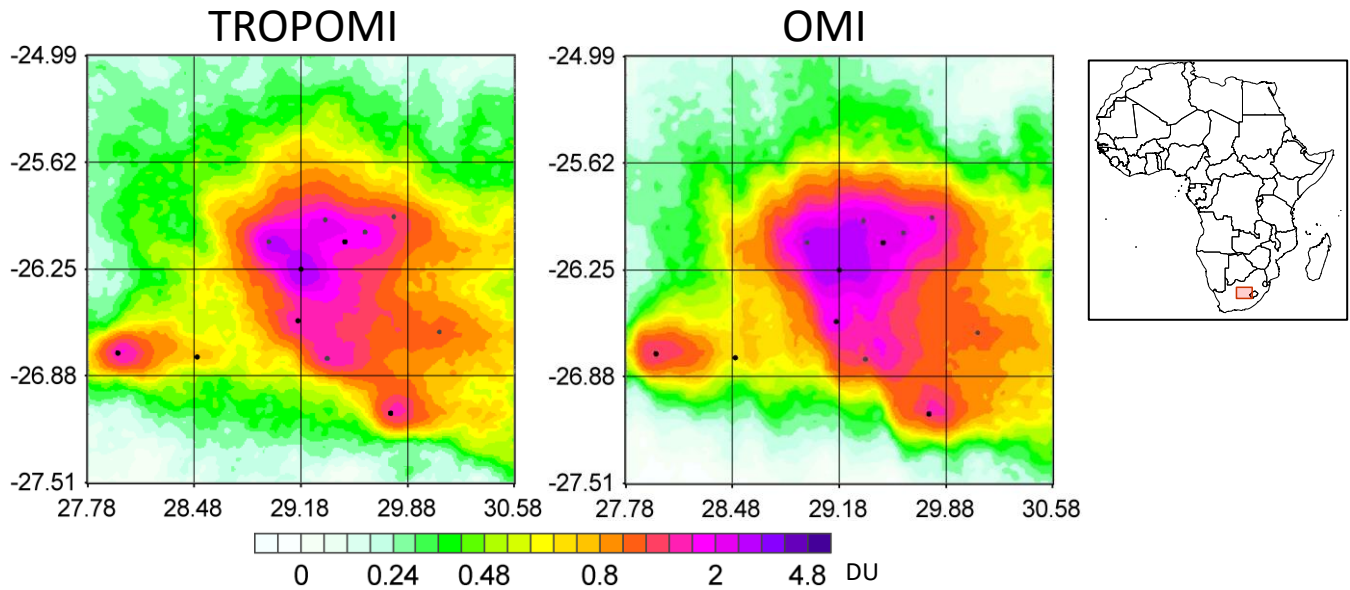
Figure 7. The mean TROPOMI SO_2 over a cluster of power plants in South Africa in the period from April 2018 to March 2019 and the mean 2005-2019 OMI SO_2 over the same region. Data are smoothed by oversampling/pixel averaging with 10 km radius. The black dots indicate the emission sources. Note that the color scale is different from the scale in the previous figures.
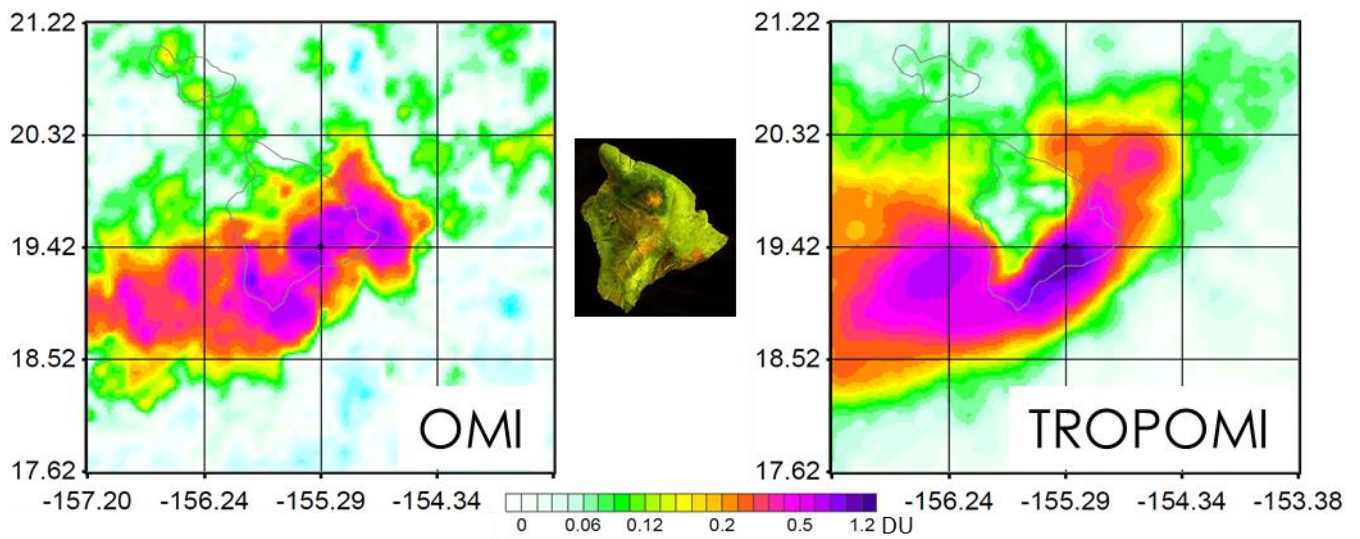
Figure 8. Mean SO$_2$ in DU over Kilauea Volcano, Hawaii, from OMI and TROPOMI data for the period from April 2018 to March 2019. The volcano is in the centre of the map. The influence of orography on the SO$_2$ distribution is clearly visible due to the high spatial resolution of Tropomi. A Sentinel-1 image from 23 May 2018 that illustrates the island's orography is shown in the middle of the figure.
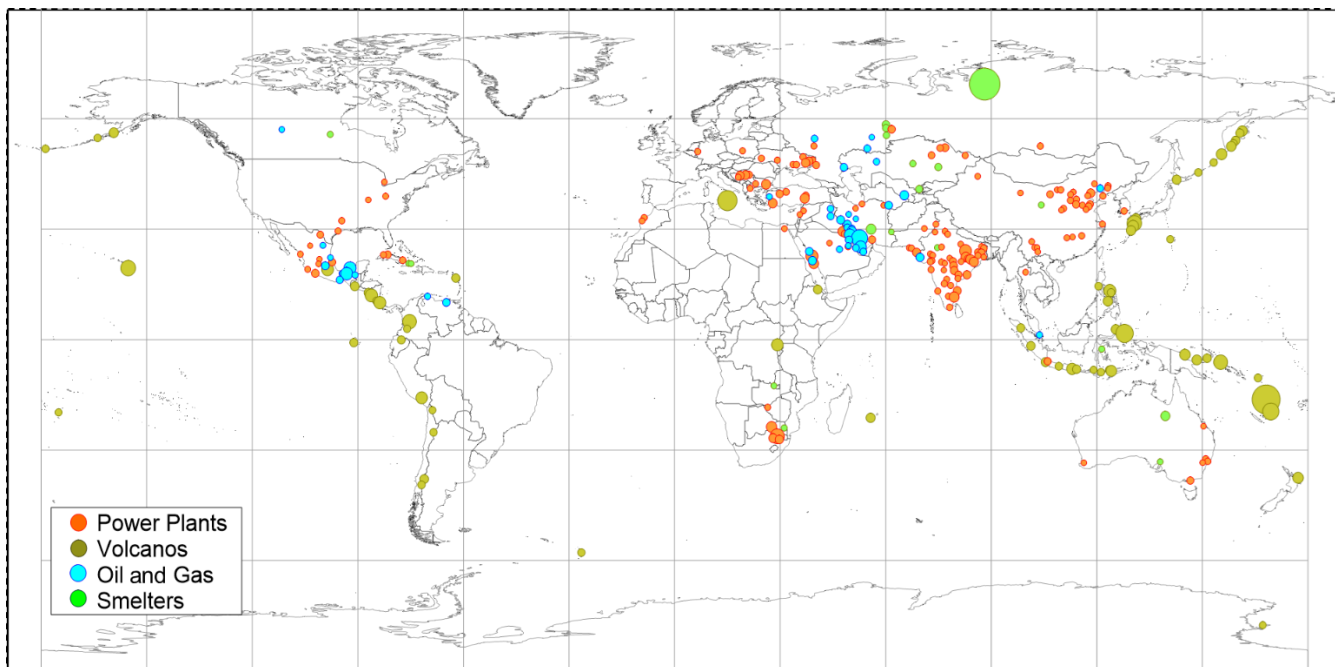
Figure 9. SO$_2$ emission sources seen by TROPOMI in 2018. We checked ~500 locations where OMI detected SO$_2$ emissions in 2005-2014 (Fioletov et al., 2016). Note that some of them are not active now or have their emissions significantly reduced. Tropomi can "see" 278 sites including 150 Power Plants, 19 Smelters, 41 Oil and Gas industry-relates sources, and 68 Volcanoes with annual emissions from 10 to 2000 kt. The size of the symbols is proportional to the annual emission values.
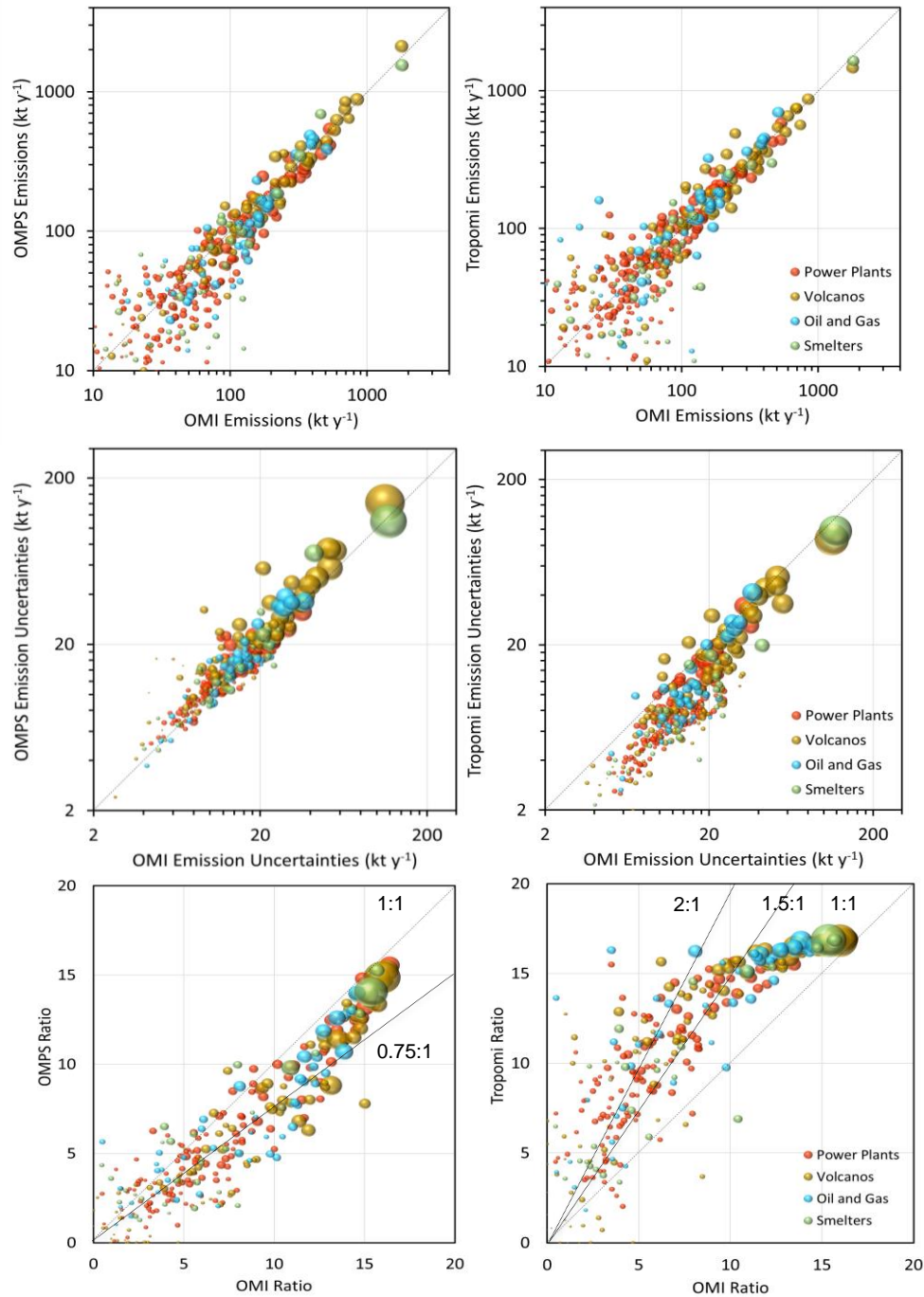
Figure 10. (a) OMI, OMPS, and TROPOMI-based estimated emissions in kt per year. The bubble area is proportional to the ratio of emission to uncertainty. The bigger the bubble, the more reliable the estimate. (b) Emission uncertainties in kt per year. The bubble area is proportional to the emission rate. The bigger the bubble the higher emissions. (c) Ratios between estimated emissions and their uncertainties. The bubble area is proportional to the emission rate.
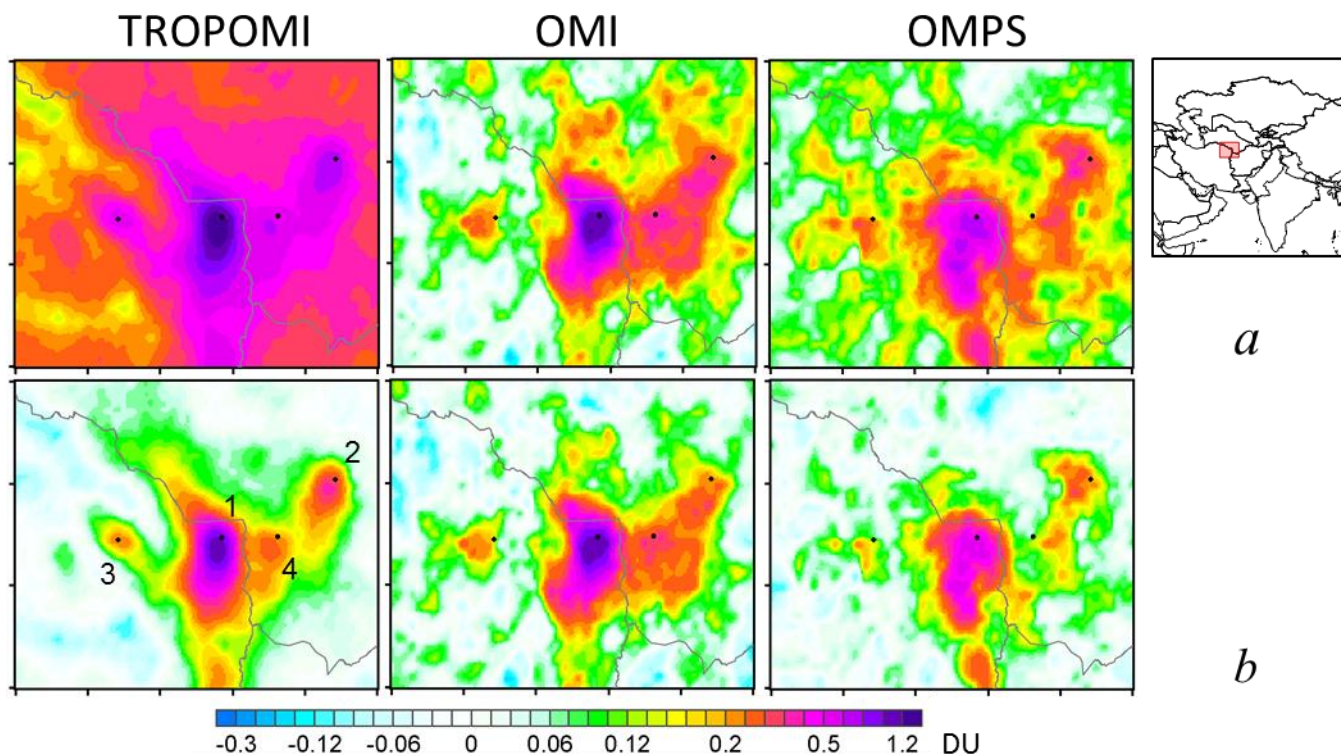
Figure 11. TROPOMI, OMI and OMPS mean SO_2 (DU) over the Khangiran region for the period from April 2018 to March 2019 (a). The same data, but with large-scale bias removed (b). Data are smoothed by oversampling/pixel averaging with R=20 km radius. The black dots indicate the SO_2 emission sources: gas refinery at Khangiran, Iran (1); gas exploration sources at Mary (2) and Sovetabad (4), Turkmenistan; Toos and Ferdosi power plants, Iran (3).

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