



Sulfur dioxide retrievals from TROPOMI onboard Sentinel-5 Precursor: algorithm theoretical basis

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Abstract. The TROPospheric Monitoring Instrument (TROPOMI) onboard the Copernicus Sentinel-5 Precursor (S-5P) platform will measure ultraviolet earthshine radiances at high spectral and improved spatial resolution (pixel size of $7\text{ km} \times 3.5\text{ km}$ at nadir) compared to its predecessors OMI and GOME-2. This paper presents the sulfur dioxide (SO_2) vertical column retrieval algorithm implemented in the S-5P operational processor UPAS (Universal Processor for UV/VIS Atmospheric Spectrometers) and comprehensively describes its various retrieval steps. The spectral fitting is performed using the differential optical absorption spectroscopy (DOAS) method including multiple fitting windows to cope with the large range of atmospheric SO_2 columns encountered. It is followed by a slant column background correction scheme to reduce possible biases or across-track-dependent artifacts in the data. The SO_2 vertical columns are obtained by applying air mass factors (AMFs) calculated for a set of representative a priori profiles and accounting for various parameters influencing the retrieval sensitivity to SO_2 . Finally, the algorithm includes an error analysis module which is fully described here. We also discuss verification results (as part of the algorithm development) and future validation needs of the TROPOMI SO_2 algorithm.

1 Introduction

Sulfur dioxide enters the Earth's atmosphere via both natural and anthropogenic processes. Through the formation of sulfate aerosols and sulfuric acid, it plays an important role on the chemistry at local and global scales and its impact ranges from short-term pollution to climate forcing. While about one-third of the global sulfur emissions originate from natural sources (volcanoes and biogenic dimethyl sulfide), the main contributor to the total budget is from anthropogenic emissions mainly from the combustion of fossil fuels (coal and oil) and from smelting. Over the last decades, a host of satellite-based UV–visible instruments have been used for the monitoring of anthropogenic and volcanic SO_2 emissions. Total vertical column density (VCD) of SO_2 has been retrieved with the sensors TOMS (Krueger, 1983), GOME (Eisinger and Burrows, 1998; Thomas et al., 2005; Khokar et al., 2005), SCIAMACHY (Afe et al., 2004), OMI (Krotkov et al., 2006; Yang et al., 2007, 2010; Li et al., 2013; Theys et al., 2015), GOME-2 (Richter et al., 2009; Bobrowski et al., 2010; Nowlan et al., 2011; Rix et al., 2012; Hörmann et al., 2013) and OMPS (Yang et al., 2013). In particular, the Ozone Monitoring Instrument (OMI) has largely demonstrated the value of satellite UV–visible remote sensing (1) in monitoring volcanic plumes in near-real time (Brenot et al., 2014) and changes in volcanic degassing at the global scale (Carn et al., 2016, and references therein) and (2) in detecting and quantifying large anthropogenic SO_2 emissions, weak or

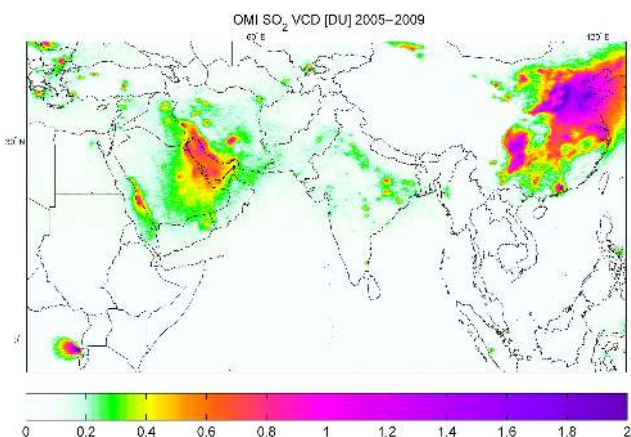


Figure 1. Map of averaged SO₂ columns from OMI clear-sky pixels for the 2005–2009 period.

unreported emission sources worldwide (Theys et al., 2015; Fioletov et al., 2016; McLinden et al., 2016) as well as investigating their long-term changes (Krotkov et al., 2016; van der A et al., 2016; He et al., 2016). An example map of OMI SO₂ columns (Theys et al., 2015) averaged over the 2005–2009 period is shown in Fig. 1, illustrating typical anthropogenic emission hotspots (China, eastern Europe, India and the Middle East) and signals from volcanic activity (e.g., from the volcanoes in DR Congo).

The 7-year-lifetime Sentinel-5p sensor TROPOMI (Veefkind et al., 2012) will fly on a polar low-Earth orbit with a wide swath of 2600 km. The TROPOMI instrument is a push-broom imaging spectrometer similar in concept to OMI. It has eight spectral bands covering UV to SWIR wavelengths. The SO₂ retrieval algorithm exploits measurements from band 3 (310–405 nm), with typical spectral resolution of 0.54 nm, signal-to-noise ratio of about 1000 and pixel size as good as 7 km × 3.5 km.

TROPOMI will continue and improve the measurement time series of OMI SO₂ and other UV sensors. Owing to similar performance to OMI in terms of signal-to-noise ratio and unprecedented spatial resolution, TROPOMI will arguably discern very fine details in the SO₂ distribution and will be able to detect point sources with annual SO₂ emissions of about 10 kT yr⁻¹ or lower (using oversampling techniques).

This paper gives a thorough description of the operational TROPOMI SO₂ algorithm and reflects the S-5P SO₂ L2 Algorithm Theoretical Basis Document v1.0. In Sect. 2, we first present the product requirements and briefly discuss the expected product performance in terms of precision and accuracy. It is then followed by the SO₂ column retrieval algorithm description. An error analysis of the retrieval method is presented in Sect. 3. Results from algorithm verification exercise using an independent retrieval scheme is given in Sect. 4. The possibilities for future validation of the retrieved SO₂ data product can be found in Sect. 5. Conclusions are

given in Sect. 6. Additional information on data product and auxiliary data, as well as a list of acronyms, is provided in the Appendix.

2 TROPOMI SO₂ algorithm

2.1 Product requirements

While UV measurements are highly sensitive to SO₂ at high altitudes (upper troposphere–lower stratosphere), the sensitivity to SO₂ concentration in the boundary layer is intrinsically limited from space due to the combined effect of scattering (Rayleigh and Mie) and ozone absorption that hamper the penetration of solar radiation into the lowest atmospheric layers. Furthermore, the SO₂ absorption signature suffers from the interference with the ozone absorption spectrum.

The retrieval precision (or random uncertainty) is driven by the signal-to-noise ratio of the recorded spectra and by the retrieval wavelength interval used, the accuracy (or systematic uncertainty) is limited by the knowledge on the auxiliary parameters needed in the different retrieval steps. Among these are the treatment of other chemical interfering species, clouds and aerosol, the representation of vertical profiles (gas, temperature, pressure), and uncertainties on data from external sources (e.g., surface reflectance).

Requirements on the accuracy and precision for the data products derived from the TROPOMI measurements are specified in the GMES Sentinels 4 and 5 and 5p Mission Requirements Document MRD (Langen et al., 2011), the Report of The Review Of User Requirements for Sentinels-4/5 (Bovensmann et al., 2011) and the Science Requirements Document for TROPOMI (van Weele et al., 2008). These requirements derive from the Composition of the Atmosphere: Progress to Applications in the user Community (CAPACITY) study (Kelder et al., 2005) and have been fine-tuned by the Composition of the Atmospheric Mission concEpts and Sentinel Observation Techniques (CAMELOT; Levelt et al., 2009) and Original and New Tropospheric composition and Air Quality measurements (ONTRAQ; Zweers et al., 2010) studies. The CAPACITY study has defined three main themes: the ozone layer (A), air quality (B), and climate (C), with further division into sub-themes. Requirements for SO₂ have been specified for a number of these sub-themes. In the following paragraphs, we discuss these requirements and the expected performances of the SO₂ retrieval algorithm (summary is given in Table 1).

2.1.1 Theme A3 – ozone layer assessment

This theme addresses the importance of measurements in the case of enhanced SO₂ concentrations in the stratosphere due to severe volcanic events. The long-term presence (up to several months) of SO₂ in the stratosphere contributes to the stratospheric aerosol loading and hence affects the cli-

Table 1. Requirements on SO₂ vertical column products as derived from the MRD. Left- and right-hand numbers in ranges denote accuracy and precision, respectively.

	Horizontal resolution (km)	Required uncertainty	Achievable uncertainty	Theme (table in MRTD)
Enhanced stratospheric column	50–200	30 % for VCD > 0.5 DU	Met for VCD > 0.5 DU	A3
Tropospheric column	5–20	30–60 % or 1.3×10^{15} molecules cm ⁻² (least stringent)	50 %/ $3\text{--}6 \times 10^{16}$ molecules cm ⁻²	B1, B2, B3
Total column	5–20	30–60 % or 1.3×10^{15} molecules cm ⁻² (least stringent)	50 %/ $3\text{--}6 \times 10^{16}$ molecules cm ⁻²	B1, B2, B3

mate and the stratospheric ozone budget. For such scenarios, the requirements state that the stratospheric vertical column should be monitored with a total uncertainty of 30 %. Although powerful volcanic events generally produce large amounts of SO₂, monitoring such a plume over extended periods of time also requires the detection of the plume after it has diluted during the weeks after the eruption.

From an error analysis of the proposed SO₂ algorithm (Sect. 3), we have assessed the major sources of uncertainty in the retrieved SO₂ column. One of the main contributors to the total uncertainty is instrumental noise. This source of error alone limits the precision to vertical columns above about 0.25 DU (1 DU = 2.69×10^{16} molec cm⁻²). For SO₂ in the stratosphere, the summing-up of the various uncertainties (Sect. 3) is believed to be around the required uncertainty of 30 % for diluted SO₂ plumes, provided that the vertical column is larger than 0.5 DU. Explosive volcanic eruptions capable of injecting SO₂ into the stratosphere regularly show stratospheric SO₂ columns of a few DU to several hundreds of DU or more, as was the case, for example, for the eruptions of Mt. Kasatochi (Yang et al., 2010) and Sarychev Peak (Carn et al., 2011). For very large SO₂ concentrations, the dynamical use of different fitting windows (see Sect. 2.2) enables the 30 % uncertainty level to be reached (see Sect. 4).

2.1.2 Theme B – air quality

This theme includes three sub-themes:

B1 Protocol monitoring: this involves the monitoring of abundances and concentrations of atmospheric constituents, driven by several agreements, such as the Gothenburg protocol, National Emission Ceilings, and EU Air Quality regulations.

B2 Near-real-time (NRT) data requirements: this comprises the relatively fast (~ 30 min) prediction and determination of surface concentrations in relation to health and safety warnings.

B3 Assessment: this sub-theme aims at answering several air-quality-related scientific questions, such as the effect on air quality of spatial and temporal variations in oxidizing capacity and long-range transport of atmospheric constituents.

A more detailed description of the air quality sub-themes can be found in Langen et al. (2011).

The user requirements on SO₂ products are equal for all three sub-themes. For the total vertical column and the tropospheric vertical column of SO₂, the user requirements state an absolute maximum uncertainty of 1.3×10^{15} molecules cm⁻² or 0.05 DU. This number derives from the ESA CAPACITY study, where the number was expressed as 0.4 ppbv for a 1.5 km thick boundary layer reaching up to 850 hPa. From the uncertainty due to instrument noise only, it is clear that the 0.05 DU requirement cannot be met on a single-measurement basis. This limitation was already found in the ESA CAMELOT study (Levelt et al., 2009).

For anthropogenic SO₂ typically confined in the planetary boundary layer (PBL), calculations performed within the CAMELOT study showed that the smallest vertical column that can be detected in the PBL is of about 1–3 DU (for a signal-to-noise ratio (S/N) of 1000). Although pollution hotspots can be better identified by spatial or temporal averaging, several uncertainties (e.g., due to varying surface albedo or SO₂ vertical profile shape) are not averaging out and directly limit the product accuracy to about 50 % or more. Though the difference between the MRD requirements and the expected TROPOMI performance is rather large, one could argue that the required threshold should not be a strict criterion in all circumstances. The user requirement of 0.05 DU represents the maximum uncertainty to distinguish (anthropogenic) pollution sources from background concentrations. Bovensmann et al. (2011) reviewed the MRD user requirements and motivated a relaxation of certain user requirements for specific conditions. For measurements in the PBL, the document proposes a relative

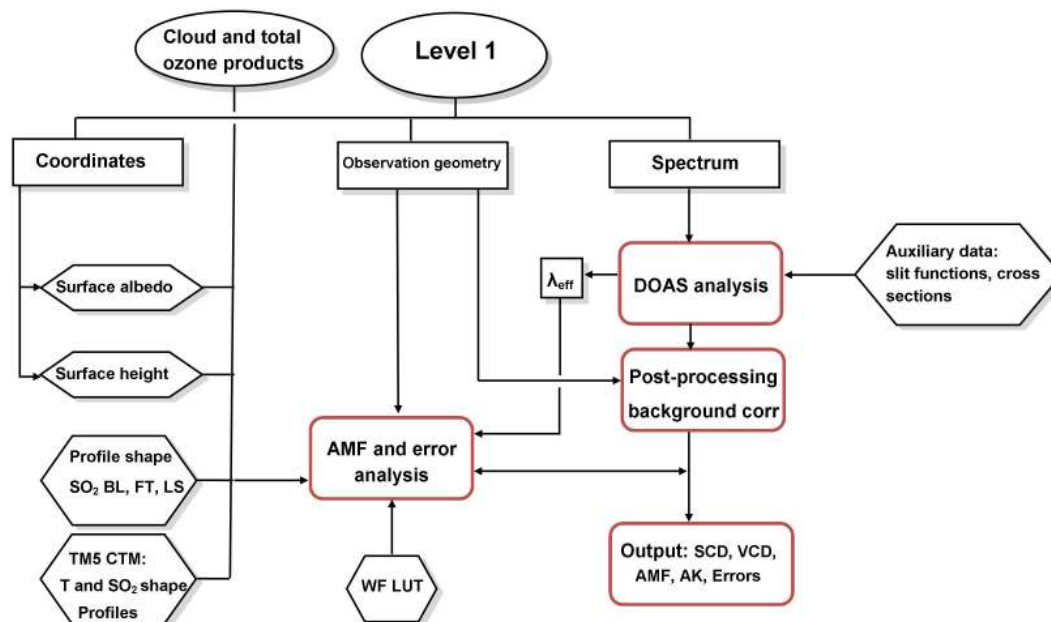


Figure 2. Flow Diagram of the TROPOMI DOAS retrieval algorithm for SO₂.

requirement of 30–60 % in order to discriminate between enhanced (> 1.5 ppbv), moderate (0.5–1.5 ppbv), and background concentrations (< 0.5 ppbv). It is expected that it will be possible to discriminate these three levels by averaging (spatiotemporally) TROPOMI data.

For volcanic SO₂ plumes in the free troposphere, a better measurement sensitivity is expected for TROPOMI. The expected precision is about 0.5 DU on the vertical column. The accuracy on the SO₂ vertical column will be strongly limited by the SO₂ plume height and the cloud conditions. As these parameters are highly variable in practice, it is difficult to ascertain the product accuracy for these conditions.

2.2 Algorithm description

The first algorithm to retrieve SO₂ columns from space-borne UV measurements was developed based on a few wavelength pairs (for TOMS) and has been subsequently applied and refined for OMI measurements (e.g., Krotkov et al., 2006; Yang et al., 2007, and references therein). Current algorithms exploit back-scattered radiance measurements in a wide spectral range using a direct fitting approach (Yang et al., 2010; Nowlan et al., 2011), a principal component analysis (PCA) method (Li et al., 2013) or (some form of) differential optical absorption spectroscopy (DOAS; Platt and Stutz, 2008); see, e.g., Richter et al. (2009), Hörmann et al. (2013), or Theys et al. (2015).

Direct fitting schemes in which on-the-fly radiative transfer simulations are made for all concerned wavelengths and resulting simulated spectra are adjusted to the spectral observations, are in principle the most accurate. They are able to cope with very large SO₂ columns (such as those occurring

during explosive volcanic eruptions), i.e., conditions typically leading to a strongly nonlinear relation between the SO₂ signal and the VCD. However, the main disadvantage of direct fitting algorithms with respect to DOAS (or PCA) is that they are computationally expensive and are out of reach for TROPOMI operational near-real-time processing, for which the level 1b data flow is expected to be massive and deliver around 1.5 million spectral measurements per orbit (~ 15 orbits daily) for band 3 (with a corresponding data size of 6 GB). To reach the product accuracy and processing performance requirements, the approach adopted here applies DOAS in three different fitting windows (within the 310–390 nm spectral range) that are still sensitive enough to SO₂ but less affected by nonlinear effects (Bobrowski et al., 2010; Hörmann et al., 2013).

Figure 2 shows the full flow diagram of the SO₂ retrieval algorithm including the dependencies on auxiliary data and other L2 products. The algorithm and its application to OMI data are also described in Theys et al. (2015), although there are differences in some settings. The baseline operation flow of the scheme is based on a DOAS retrieval algorithm and is identical to that implemented in the retrieval algorithm for HCHO (also developed by BIRA-IASB; see De Smedt et al., 2016). The main output parameters of the algorithm are SO₂ vertical column density, slant column density, air mass factor, averaging kernels (AKs) and error estimates. Here, we will first briefly discuss the principle of the DOAS VCD retrieval before discussing the individual steps of the process in more detail.

First, the radiance and irradiance data are read from an S-5P L1b file, along with geolocation data such as pixel coor-

dinates and observation geometry (sun and viewing angles). At this stage cloud cover information (see Table A3 in Appendix A) is also obtained from the S-5P cloud L2 data, as required for the calculation of the AMF, later in the scheme. Then relevant absorption cross section data, as well as characteristics of the instrument (e.g., slit functions), are used as input for the determination of SO₂ slant column density. As a baseline, the slant column fit is done in a sensitive window from 312 to 326 nm. For pixels with a strong SO₂ signal, results from alternative windows where the SO₂ absorption is weaker can be used instead. An empirical offset correction (dependent on the fitting window used) is then applied to the SCD. The latter correction accounts for systematic biases in the SCDs. Following the SCD determination, the AMF is estimated based on a pre-calculated weighting functions (or box AMFs) look-up table (LUT). This look-up table is generated using the LLinearized Discrete Ordinate Radiative Transfer (LIDORT) code (Spurr, 2008) and has several entries: cloud cover data, topographic information, observation geometry, surface albedo, effective wavelength (representative of the fitting window used), total ozone column and the shape of the vertical SO₂ profile. The algorithm also includes an error calculation and retrieval characterization module (Sect. 3) that computes the averaging kernels (Eskes and Boersma, 2003), which characterize the vertical sensitivity of the measurement and which are required for comparison with other types of data (Veefkind et al., 2012).

The final SO₂ vertical column is obtained by

$$N_v = \frac{N_s - N_s^{\text{back}}}{M}, \quad (1)$$

where the main quantities are the vertical column (N_v), the slant column density (N_s) and the values used for the background correction (N_s^{back}). M is the air mass factor.

2.2.1 Slant column retrieval

The backscattered radiance spectrum recorded by the space instrument differs from the solar spectrum because of the interactions of the photons with the Earth's atmosphere and surface reflection. Hence, the reflectance spectra contains spectral features that can be related to the various absorbing species and their amounts in the atmosphere. The DOAS method aims at the separation of the highly structured trace gas absorption spectra and broadband spectral structures. The technique relies on a number of assumptions that can be summarized as follows:

- The spectral analysis and atmospheric radiative transfer computations are treated separately by considering one averaged atmospheric light path of the photons traveling from the sun to the instrument.
- The absorption cross sections are not strongly dependent on pressure and temperature. Additionally, the averaged light path should be weakly dependent on the

wavelength – for the fitting window used – which enables defining an effective absorption (slant) column density. It should be noted that this is not strictly valid for the SO₂ DOAS retrieval because of strong absorption by ozone and in some cases SO₂ itself (for large SO₂ amounts).

- Spectrally smoothed structures due broadband absorption, scattering and reflection processes can be well reproduced by a low-order polynomial as a function of wavelength.

Photons collected by the satellite instrument may have followed very different light paths through the atmosphere depending on their scattering history. However, a single effective light path is assumed, which represents an average of the complex paths of all reflected and scattered solar photons reaching the instrument within the spectral interval used for the retrieval. This simplification is valid if the effective light path is reasonably constant over the considered wavelength range. The spectral analysis can be described by the following equation:

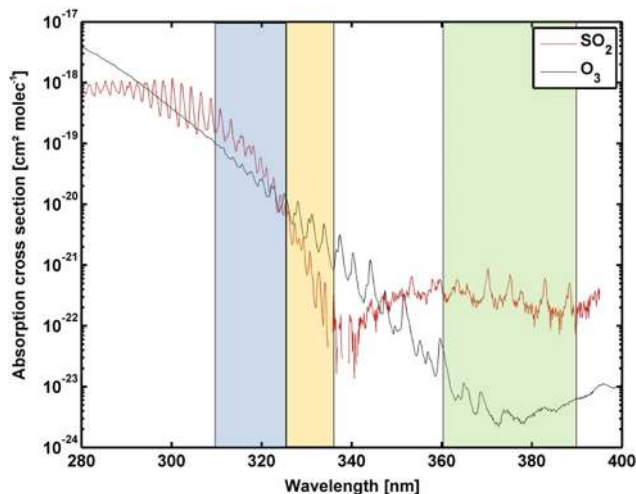
$$\ln \frac{\pi I(\lambda)}{\mu_0 E_0(\lambda)} = - \sum_j \sigma_j(\lambda) N_{s_j} + \sum_p c_p \lambda^p, \quad (2)$$

where $I(\lambda)$ is the observed backscattered earthshine radiance ($\text{W m}^{-2} \text{nm}^{-1} \text{sr}^{-1}$), E_0 is the solar irradiance ($\text{W m}^{-2} \text{nm}^{-1}$) and $\mu_0 = \cos \theta_0$. The first term on the right-hand side indicates all relevant absorbing species with absorption cross sections σ_j ($\text{cm}^2 \text{molec}^{-1}$). Integration of the number densities of these species along the effective light path gives the slant column density N_{s_j} (molec cm^{-2}). Equation (2) can be solved by least-squares fitting techniques (Platt and Stutz, 2008) for the slant column values. The final term in Eq. (2) is the polynomial representing broadband absorption and (Rayleigh and Mie) scattering structures in the observed spectrum and also accounts for possible errors such as uncorrected instrument degradation effects, uncertainties in the radiometric calibration or possible residual (smooth) polarization response effects not accounted for in the level 0–1 processing.

Apart from the cross sections for the trace gases of interest, additional fit parameters need to be introduced to account for the effect of several physical phenomena on the fit result. For SO₂ fitting, these are the filling-in of Fraunhofer lines (Ring effect) and the need for an intensity offset correction. In the above, we have assumed that for the ensemble of observed photons a single effective light path can be assumed over the adopted wavelength fitting interval. For the observation of (generally small) SO₂ concentrations at large solar zenith angles (SZAs) this is not necessarily the case. For such long light paths, the large contribution of O₃ absorption may lead to negative SO₂ retrievals. This may be mitigated by taking the wavelength dependence of the O₃ SCD over the

Table 2. DOAS settings used to retrieve SO₂ slant columns.

Fitting intervals 1 and 2	312–326 nm (w1), 325–335 nm (w2)
Cross sections	SO ₂ : 203K (Bogumil et al., 2003) O ₃ : 228 and 243 K with Io correction (Brion et al., 1998) Pseudo-O ₃ cross sections ($\lambda\sigma_{O_3}$, $\sigma_{O_3}^2$; Pukite et al., 2010) Ring effect: two eigenvectors (Vountas et al., 1998) generated for 20 and 87° solar zenith angles using LIDORT-RRS (Spurr et al., 2008)
Polynomial	Fifth order
Fitting interval 3	360–390 nm (w3)
Cross sections	SO ₂ : Hermans et al. (2009) extrapolated at 203K NO ₂ : 220 K (Vandaele et al., 1998) O ₂ -O ₂ : Greenblatt et al., 1990 Ring effect: single spectrum (Chance and Spurr, 1997)
Polynomial	Fourth order
Intensity offset correction	Linear offset
Spectrum shift and stretch	Fitted
Spectral spikes removal procedure	Richter et al. (2011)
Reference spectrum	Baseline: daily solar irradiance Foreseen update: daily averaged earthshine spectrum in Pacific region (10° S–10° N, 160° E–120° W); separate spectrum for each detector row. NRT: averaged spectra of the last available day; offline: averaged spectra of the current day

**Figure 3.** Absorption cross sections of SO₂ and O₃. The blue, yellow and green boxes delimit the three SO₂ fitting windows 312–326, 325–335 and 360–390 nm, respectively.

fitting window into account, as will be described in the next section.

The different parts of the DOAS retrieval are detailed in the next subsections and Table 2 gives a summary of settings used to invert SO₂ slant columns. Note that, in Eq. (2), the daily solar irradiance is used as a baseline for the reference spectrum. As a better option, it is generally preferred to use daily averaged radiances, selected for each across-track position, in the equatorial Pacific. In the NRT algorithm, the last valid day can be used to derive the reference spectra,

while in the offline version of the algorithm, the current day should be used. Based on OMI experience, it would allow, for example, for better handling of instrumental artifacts and degradation of the recorded spectra for each detector. At the time of writing, it is planned to test this option during the S-5P commissioning phase.

Wavelength fitting windows

DOAS measurements are in principle applicable to all gases having suitable narrow absorption bands in the UV, visible, or near-IR regions. However, the generally low concentrations of these compounds in the atmosphere, and the limited signal-to-noise ratio of the spectrometers, restrict the number of trace gases that can be detected. Many spectral regions contain several interfering absorbers and correlations between absorber cross sections can sometimes lead to systematic biases in the retrieved slant columns. In general, the correlation between cross sections decreases if the wavelength interval is extended, but then the assumption of a single effective light path defined for the entire wavelength interval may not be fully satisfied, leading to systematic misfit effects that may also introduce biases in the retrieved slant columns (e.g., Pukite et al., 2010). To optimize DOAS retrieval settings, a trade-off has to be found between these effects. In the UV–visible spectral region, the cross-section spectrum of SO₂ has its strongest bands in the 280–320 nm range (Fig. 3). For the short wavelengths in this range, the SO₂ signal, however, suffers from a strong increase in Rayleigh scattering and ozone absorption. In practice, this leads to a very small SO₂ signal in the satellite spectra compared to ozone absorp-

tion, especially for tropospheric SO₂. Consequently, SO₂ is traditionally retrieved (for GOME, SCIAMACHY, GOME-2, OMI) using sensitive windows in the 310–326 nm range. Note that even in this range the SO₂ absorption can be 3 orders of magnitude lower than that of ozone.

The TROPOMI SO₂ algorithm is using a multiple-window approach:

- 312–326 nm: classical fitting window, ideal for small columns. This window is used as baseline. If nonlinear effects due to high SO₂ amounts are encountered, one of the two following windows will be used instead.
- 325–335 nm: in this window, differential SO₂ spectral features are 1 order of magnitude smaller than in the classical window. It allows the retrieval of moderate SO₂ columns, an approach similar to the one described by Hörmann et al. (2013).
- 360–390 nm: SO₂ absorption bands are 2–3 orders of magnitude weaker than in the classical window and are best suited for the retrieval of extremely high SO₂ columns (Bobrowski et al., 2010)

Note that in the 325–335 and 360–390 nm windows the Rayleigh scattering and ozone absorption are less important than in the baseline 312–326 nm window (see also Fig. 3).

Specifically, in the first two intervals, absorption cross sections of O₃ at 228 and 243 K are included in the fit and, to better cope with the strong (nonlinear) ozone absorption at short wavelengths, the retrieval also includes two pseudo-cross sections following the approach of Pukite et al. (2010): $\lambda\sigma_{\text{O}_3}$ and $\sigma_{\text{O}_3}^2$ calculated from the O₃ cross-section spectrum at 228 K. The correction for the Ring effect is based on the technique outlined by Vountas et al. (1998). This technique involves a PCA of a set of Ring spectra, calculated for a range of solar zenith angles. The first two of the resulting eigenvectors appear to accurately describe the Ring spectra, with the first eigenvector representing the filling-in of Fraunhofer lines and the second mostly representing the filling-in of gas absorption features. In the retrieval algorithm, these vectors are determined by orthogonalizing two Ring spectra, calculated by LIDORT-RRS (Spurr et al., 2008), a version of LIDORT accounting for rotational Raman scattering, for a low SZA (20°) and a high SZA (87°), respectively.

Wavelength calibration and convolution to TROPOMI resolution

The quality of a DOAS fit critically depends on the accuracy of the alignment between the earthshine radiance spectrum, the reference spectrum and the cross sections. Although the level 1b will contain a spectral assignment, an additional spectral calibration is part of the SO₂ algorithm. Moreover, the DOAS spectral analysis also includes the fit of shift and stretch of radiance spectra because the TROPOMI spectral registration will differ from one ground pixel to another, e.g.,

due to thermal variations over the orbit as well as due to inhomogeneous filling of the slit in flight direction.

The wavelength registration of the reference spectrum can be fine-tuned by means of a calibration procedure making use of the solar Fraunhofer lines. To this end, a reference solar atlas, E_s , accurate in absolute vacuum wavelength to better than 0.001 nm (Chance and Kurucz, 2010) is degraded at the resolution of the instrument, through convolution by the TROPOMI instrumental slit function.

Using a nonlinear least-squares approach, the shift (Δ_i) between the reference solar atlas and the TROPOMI irradiance is determined in a set of equally spaced sub-intervals covering a spectral range large enough to encompass all relevant fitting intervals. The shift is derived according to the following equation:

$$E_0(\lambda) = E_s(\lambda - \Delta_i), \quad (3)$$

where E_s is the solar spectrum convolved at the resolution of the instrument and Δ_i is the shift in sub-interval i . A polynomial is then fitted through the individual points in order to reconstruct an accurate wavelength calibration $\Delta(\lambda)$ for the complete analysis interval. Note that this approach allows one to compensate for stretch and shift errors in the original wavelength assignment.

In the case of TROPOMI, the procedure is complicated by the fact that such calibrations must be performed (and stored) for each separate spectral field on the CCD detector array. Indeed, due to the imperfect characteristics of the imaging optics, each row of the TROPOMI instrument must be considered a separate spectrometer for analysis purposes.

In a subsequent step of the processing, the absorption cross sections of the different trace gases must be convolved with the instrumental slit function. The baseline approach is to use slit functions determined as part of the TROPOMI key data. Slit functions are delivered for each binned spectrum and as a function of wavelength. Note that an additional feature of the prototype algorithm allows for an effective slit function of known line shape to be dynamically fitted (e.g., asymmetric Gaussian). This can be used for verification and monitoring purpose during commissioning and later on during the mission.

More specifically, wavelength calibrations are made for each TROPOMI orbit as follows:

1. The TROPOMI irradiances (one for each row of the CCD) are calibrated in wavelength over the 310–390 nm wavelength range, using 10 sub-windows.
2. The earthshine radiances and the absorption cross sections are interpolated (cubic spline interpolation) on the calibrated wavelength grid, prior to the DOAS analysis.
3. During spectral fitting, shift and stretch parameters are further derived to align radiance and irradiance spectra.

Table 3. Criteria for selecting alternative fitting windows.

Window number	w1	w2	w3
Wavelength range	312–326 nm	325–335 nm	360–390 nm
Derived slant column	S1	S2	S3
Application	Baseline for every pixel	S1 > 15 DU and S2 > S1	S2 > 250 DU and S3 > S2

The reference wavelength grid used in the DOAS procedure is the (optimized) grid of the TROPOMI solar irradiance.

Spike removal algorithm

A method to remove individual hot pixels or detector pixels affected by the South Atlantic Anomaly has been presented for NO₂ retrievals in Richter et al. (2011). Often only a few individual detector pixels are affected, and in these cases, it is possible to identify and remove the noisy points from the fit. However, as the amplitude of the distortion is usually only of the order of a few percent or less, it cannot always be found in the highly structured spectra themselves. Higher sensitivity for spikes can be achieved by analyzing the residual of the fit where the contribution of the Fraunhofer lines, scattering, and absorption is already removed.

When the residual for a single detector pixel exceeds the average residual of all detector pixels by a chosen threshold ratio (the tolerance factor), the pixel is excluded from the analysis, in an iterative process. This procedure is repeated until no further outliers are identified, or until the maximum number of iterations is reached (here fixed to 3). This is especially important to handle the degradation of 2-D detector arrays such as OMI or TROPOMI. However, this improvement of the algorithm has a non-negligible impact on the time of processing. At the time of writing, the exact values for the tolerance factor and maximum number of iterations of the spike removal procedure are difficult to ascertain and will only be known during operations. To assess the impact on the processing time, test retrievals have been done on OMI spectra using a tolerance factor of 5 and a limit of three iterations (this could be relaxed), leading to an increase in processing time by a factor of 1.5.

Fitting window selection

The implementation of the multiple-fitting-window retrieval requires selection criteria for the transition from one window to another. These criteria are based on the measured SO₂ slant columns. As a baseline, the SO₂ SCD in the 312–326 nm window will be retrieved for each satellite pixel. When the resulting value exceeds a certain criterion, the slant column retrieval is taken from an alternative window. As part of the algorithm development and during the verification exercise (Sect. 4), closed-loop retrievals have been performed

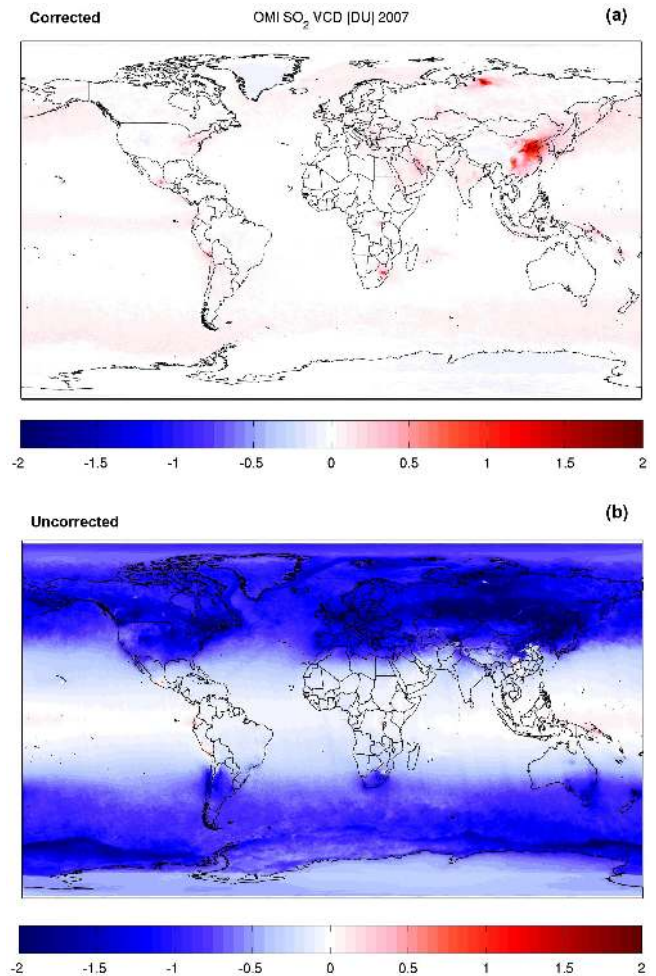


Figure 4. OMI SO₂ vertical columns (DU) averaged for the year 2007 (top) with and (bottom) without background correction. Only clear-sky pixels (cloud fraction lower than 30 %) have been kept. AMFs calculated from SO₂ profiles from the IMAGES global model are applied to the slant columns (Theys et al., 2015).

and application of the algorithm to real data from the GOME-2 and OMI instruments lead to threshold values and criteria as given in Table 3.

2.2.2 Offset correction

When applying the algorithm to OMI and GOME-2 data, across-track/viewing-angle-dependent residuals of SO₂ were found over clean areas and negative SO₂ SCDs are found at high SZA which need to be corrected (note that this is a common problem of most algorithms to retrieve SO₂ from space UV sensors). A background correction scheme was found mostly necessary for the SO₂ slant columns retrieved in the baseline fitting window. The adopted correction scheme depends on across-track position and measured O₃ slant column as described below.

The correction is based on a parameterization of the background values that are then subtracted from the measurements. The scheme first removes pixels with high SZA ($> 70^\circ$) or SCDs larger than 1.5 DU (measurements with presumably real SO₂) and then calculates the offset correction by averaging the SO₂ data on an ozone slant column grid (bins of 75 DU). This is done independently for each across-track position and hemisphere, and the correction makes use of measurements averaged over a time period of 2 weeks preceding the measurement of interest (to improve the statistics and minimize the impact of a possible extended volcanic SO₂ plume on the averaged values).

It should be noted that the O₃ slant column is dependent on the wavelength when applying the approach of Pukite et al. (2010):

$$\text{SCD}(\lambda) = \text{SCD}_{T1} + \text{SCD}_{T2} + \lambda \text{SCD}_\lambda + \sigma_s(\lambda) \text{SCD}_s. \quad (4)$$

SCD_{T1} and SCD_{T2} are the retrieved ozone slant columns corresponding to the ozone cross sections at two temperatures ($T1$, $T2$) included in the fit. SCD_λ and SCD_s are the retrieved parameters for the two pseudo-cross sections $\lambda \cdot \sigma_s$ and σ_s^2 (σ_s being the O₃ cross section at $T1$). In order to apply the background correction, the O₃ slant column expression (Eq. 4) is evaluated at 313 nm (read below).

An example of the effect of the background correction is shown in Fig. 4 for OMI. One can see that after correction (top panel) the retrievals show smooth/unstriped results and values close to zero outside the polluted areas. In some regions (in particular at high latitudes), residual columns can be found, but are generally lower than 0.2 DU.

For the two additional fitting windows, residual SO₂ levels are relatively small in comparison to the column amounts expected to be retrieved in these windows. However, simplified background corrections are also applied to the alternative windows: the offset corrections use parameterizations of the background slant columns based on latitude (bins of 5°), cross-track position and time (2-week moving averages as for the baseline window). To avoid contamination by strong volcanic eruptions, only the pixels are kept with SCD less than 50 and 250 DU for the fitting windows 325–335 and 360–390 nm, respectively.

It should be noted that the background corrections do not imply saving 2 weeks of SO₂ L2 data in intermediate products, but only the averaged values ($\sum_{i=1} N \text{SCD}_i / N$) over the predefined working grids (note: the numerators $\sum_{i=1} N \text{SCD}_i$ and denominators N are stored separately).

This background correction is well suited for the case of a 2-D-detector array such as TROPOMI, for which across-track striping can possibly arise due to imperfect cross-calibration and different dead/hot pixel masks for the CCD detector regions. This instrumental effect can also be found for scanning spectrometers, but since these instruments only have one single detector, such errors do not appear as stripes. These different retrieval artifacts can be compensated for (up

to a certain extent) using background corrections which depend on the across-track position. All of these corrections are also meant to handle the time-dependent degradation of the instrument. Note that experiences with OMI show that the most efficient method to avoid across-track stripes in the retrievals is to use row-dependent mean radiances as control spectrum in the DOAS fit.

2.2.3 Air mass factors

The DOAS method assumes that the retrieved slant column (after appropriate background correction) can be converted into a vertical column using a single air mass factor M (representative of the fitting interval):

$$M = \frac{N_s}{N_v}, \quad (5)$$

which is determined by radiative transfer calculations with LIDORT version 3.3 (Spurr, 2008). The AMF calculation is based on the formulation of Palmer et al. (2001):

$$M = \int m'(p) \cdot s(p) dp, \quad (6)$$

with $m' = m(p)/C_{\text{temp}}(p)$, where $m(p)$ is the so-called weighting function (WF) or pressure-dependent air mass factor, C_{temp} is a temperature correction (see Sect. 2.2.3.7) and s is the SO₂ normalized a priori mixing ratio profile, as a function of pressure (p).

The AMF calculation assumes Lambertian reflectors for the ground and the clouds and makes use of pre-calculated WF LUTs at 313, 326 and 375 nm (depending on the fitting window used). Calculating the AMF at these three wavelengths was found to give the best results using closed-loop retrievals (see Auxiliary material of Theys et al., 2015). The WF depends on observation geometry (solar zenith angle: SZA; line-of-sight angle: LOS; relative azimuth angle: RAA), total ozone column (TO3), scene albedo (alb), surface pressure (p_s), cloud top pressure (p_{cloud}) and effective cloud fraction (f_{eff}).

Examples of SO₂ weighting functions are displayed in Fig. 5 (as a function of height for illustration purpose) and show the typical variations in the measurement sensitivity as a function of height, wavelength and surface albedo.

The generation of the WF LUT has been done for a large range of physical parameters, listed in Table 4. In practice, the WF for each pixel is computed by linear interpolation of the WF LUT at the a priori profile pressure grid and using the auxiliary data sets described in the following subsections. Linear interpolations are performed along the cosine of solar and viewing angles, relative azimuth angle and surface albedo, while a nearest-neighbor interpolation is performed in surface pressure. In particular, the grid of surface pressure

Table 4. Physical parameters that define the WF look-up table.

Parameter	Number of grid points	Grid values	Symbol
Atmospheric pressure (hPa)	64	1056.77, 1044.17, 1031.72, 1019.41, 1007.26, 995.25, 983.38, 971.66, 960.07, 948.62, 937.31, 926.14, 915.09, 904.18, 887.87, 866.35, 845.39, 824.87, 804.88, 785.15, 765.68, 746.70, 728.18, 710.12, 692.31, 674.73, 657.60, 640.90, 624.63, 608.58, 592.75, 577.34, 562.32, 547.70, 522.83, 488.67, 456.36, 425.80, 396.93, 369.66, 343.94, 319.68, 296.84, 275.34, 245.99, 210.49, 179.89, 153.74, 131.40, 104.80, 76.59, 55.98, 40.98, 30.08, 18.73, 8.86, 4.31, 2.18, 1.14, 0.51, 0.14, 0.03, 0.01, 0.001	p_l
Altitude corresponding to the atmospheric pressure, using a US standard atmosphere (km)	64	-0.35, -0.25, -0.15, -0.05, 0.05, 0.15, 0.25, 0.35, 0.45, 0.55, 0.65, 0.75, 0.85, 0.95, 1.10, 1.30, 1.50, 1.70, 1.90, 2.10, 2.30, 2.50, 2.70, 2.90, 3.10, 3.30, 3.50, 3.70, 3.90, 4.10, 4.30, 4.50, 4.70, 4.90, 5.25, 5.75, 6.25, 6.75, 7.25, 7.75, 8.25, 8.75, 9.25, 9.75, 10.50, 11.50, 12.50, 13.50, 14.50, 16.00, 18.00, 20.00, 22.00, 24.00, 27.50, 32.50, 37.50, 42.50, 47.50, 55.00, 65.00, 75.00, 85.00, 95.00	z_l
Solar zenith angle (°)	17	0, 10, 20, 30, 40, 45, 50, 55, 60, 65, 70, 72, 74, 76, 78, 80, 85	θ_0
Line-of-sight angle (°)	10	0, 10, 20, 30, 40, 50, 60, 65, 70, 75	θ
Relative azimuth angle (°)	5	0, 45, 90, 135, 180	φ
Total ozone column (DU)	4	205, 295, 385, 505	TO3
Surface albedo	14	0, 0.01, 0.025, 0.05, 0.075, 0.1, 0.15, 0.2, 0.25, 0.3, 0.4, 0.6, 0.8, 1.0	A_s
Surface/cloud top pressure (hPa)	17	1063.10, 1037.90, 1013.30, 989.28, 965.83, 920.58, 876.98, 834.99, 795.01, 701.21, 616.60, 540.48, 411.05, 308.00, 226.99, 165.79, 121.11	p_s
AMF wavelength	3	313, 326, 375	

is very thin near the ground in order to minimize interpolation errors caused by the generally low albedo of ground surfaces. Furthermore, the LUT and model pressures are scaled to the respective surface pressures in order to avoid extrapolations outside the LUT range.

Observation geometry

The LUT covers the full range of values for solar zenith angles, line-of-sight angles and relative azimuth angles that can be encountered in the TROPOMI measurements. The observation geometry is readily present in the L1b data for each satellite pixel.

Total ozone column

The measurement sensitivity at 313 nm is dependent on the total ozone absorption. The LUT covers a range of ozone

column values from 200 to 500 DU for a set of typical ozone profiles. The total ozone column is directly available from the operational processing of the S-5P total ozone column product.

Surface albedo

The albedo value is very important for PBL anthropogenic SO₂ but less critical for volcanic SO₂ when it is higher in the atmosphere. For the surface albedo dimension, we use the climatological monthly minimum Lambertian equivalent reflector (minLER) data from Kleipool et al. (2008) at 328 nm for w1 and w2, and 376 nm for w3. This database is based on OMI measurements and has a spatial resolution of $0.5^\circ \times 0.5^\circ$. Note that other surface reflectance databases with improved spatial resolution (more appropriate for TROPOMI) will likely become available and these data sets will be considered for next algorithmic versions.

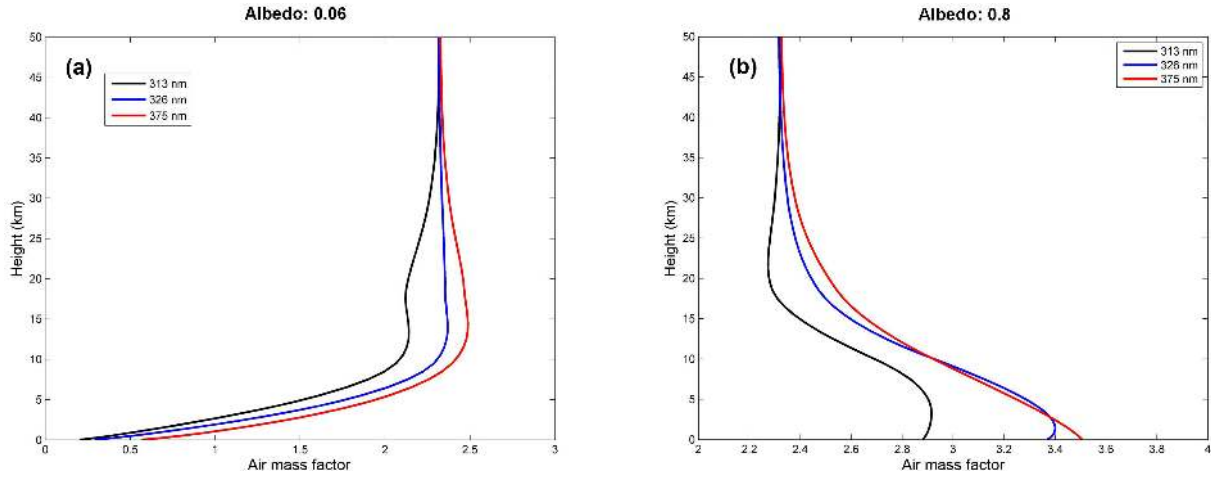


Figure 5. SO₂ box AMFs at 313, 326 and 375nm for albedo of (a) 0.06 and (b) 0.8. SZA: 40°; LOS: 10°; RAA: 0°; surface height: 0 km.

Clouds

The AMF calculations for TROPOMI partly cloudy scenes use the cloud parameters (cloud fraction f_c , cloud albedo A_c , cloud top pressure ctp) supplied by the nominal S-5P cloud algorithm OCRA/ROCINN in its Clouds as Reflecting Boundaries (CRB) implementation (Loyola et al., 2016). The cloud surface is considered to be a Lambertian reflecting surface and the treatment of clouds is achieved through the independent pixel approximation (IPA; Martin et al., 2002), which considers an inhomogeneous satellite pixel as being composed (as for the radiance intensity) of two independent homogeneous scenes, one completely clear and the other completely cloudy. The weighting function is expressed as

$$m(p) = \Phi m_{\text{cloud}}(p) + (1 - \Phi) m_{\text{clear}}(p), \quad (7)$$

where Φ is the intensity-weighted cloud fraction or cloud radiance fraction:

$$\Phi = \frac{f_{\text{eff}} I_{\text{cloud}}}{f_{\text{eff}} I_{\text{cloud}} + (1 - f_{\text{eff}}) I_{\text{clear}}}. \quad (8)$$

The suffixes “clear” and “cloudy” refer to the WF and intensity calculation corresponding to a fully clear or cloudy pixel, respectively. The WF LUT is therefore accompanied by an intensity LUT with the same input grids. Both LUTs have been generated for a range of cloud cover fractions and cloud top pressures.

Note that the variations in the cloud albedo are directly related to the cloud optical thickness. Strictly speaking, in a Lambertian (reflective) cloud model approach, only thick clouds can be represented. An effective cloud fraction corresponding to an effective cloud albedo of 0.8 ($f_{\text{eff}} \cong f_c \frac{A_c}{0.8}$) can be defined in order to transform optically thin clouds into equivalent optically thick clouds of reduced extent. Note that

in some cases (thick clouds with $A_c > 0.8$) the effective cloud fraction can be larger than one and the algorithm assumes $f_{\text{eff}} = 1$. In such altitude-dependent air mass factor calculations, a single cloud top pressure is assumed within a given viewing scene. For low effective cloud fractions (f_{eff} lower than 10%), the current cloud top pressure output is highly unstable and it is therefore reasonable to consider the observation a clear-sky pixel (i.e., the cloud fraction is set to 0 in Eq. 8) in order to avoid unnecessary error propagation through the retrievals, which can be as high as 100%. Moreover, it has been shown recently by Wang et al. (2016) using multi-axis DOAS (MAX-DOAS) observations to validate satellite data that, in the case of elevated aerosol loadings in the PBL (typically leading to apparent f_{eff} up to 10%), it is recommended to apply clear-sky AMFs rather than total AMFs (based on cloud parameters) that presumably correct implicitly for the aerosol effect on the measurement sensitivity.

It should be noted that the formulation of the pressure-dependent air mass factor for a partly cloudy pixel implicitly includes a correction for the SO₂ column lying below the cloud and therefore not seen by the satellite, the so-called ghost column. Indeed, the total AMF calculation as expressed by Eqs. (6) and (7) assumes the same shape factor and implies an integration of the a priori profile from the top of the atmosphere to the ground, for each fraction of the scene. The ghost column information is thus coming from the a priori profile shapes. For this reason, only observations with moderate cloud fractions (f_{eff} lower than 30%) are used, unless it can be assumed that the cloud cover is mostly situated below the SO₂ layer, i.e., a typical situation for volcanic plumes injected into the upper troposphere or lower stratosphere.

Surface height

The surface height (z_s) is determined for each pixel by interpolating the values of a high-resolution digital elevation map, GMTED2010 (Danielson and Gesch, 2011).

Profile shapes

It is generally not possible to know at the time of observation what the SO₂ vertical profile is and whether the observed SO₂ is of volcanic origin or from pollution (or both). Therefore, the algorithm computes four vertical columns for different hypothetical SO₂ profiles.

Three box profiles of 1 km thickness, located in the boundary layer, upper troposphere and lower stratosphere, are used. The first box profile stands for typical conditions of well mixed SO₂ (from volcanic or anthropogenic emissions) in the boundary layer, while the upper-troposphere and lower-stratosphere box profiles are representative of volcanic SO₂ plumes from effusive and explosive eruptions, respectively.

In order to have more realistic SO₂ profiles for polluted scenes, daily forecasts calculated with the global TM5 chemical transport model (Huijnen et al., 2010) will also be used. TM5 will be operated with a spatial resolution of 1° × 1° in latitude and longitude, and with 34 sigma pressure levels up to 0.1 hPa in the vertical direction. TM5 will use 3 h meteorological fields from the European Centre for Medium-Range Weather Forecast (ECMWF) operational model (ERA-Interim reanalysis data for reprocessing, and the operational archive for real time applications and forecasts). These fields include global distributions of wind, temperature, surface pressure, humidity, (liquid and ice) water content, and precipitation. A more detailed description of the TM5 model is given at <http://tm.knmi.nl/> and by van Geffen et al. (2016).

For the calculation of the air mass factors, the profiles are linearly interpolated in space and time, at the pixel center and S-5P local overpass time, through a model time step of 30 min. For NRT processing, the daily forecast of the TM5 model (located at KNMI) will be ingested by the UPAS operational processor.

To reduce the errors associated to topography and the lower spatial resolution of the model compared to the TROPOMI 7 km × 3.5 km spatial resolution, the a priori profiles need to be rescaled to effective surface elevation of the satellite pixel. The TM5 surface pressure is converted by applying the hypsometric equation and the assumption that temperature changes linearly with height (Zhou et al., 2009):

$$p_s = p_{\text{TM5}} \left(\frac{T_{\text{TM5}}}{T_{\text{TM5}} + \Gamma(z_{\text{TM5}} - z_s)} \right)^{-\frac{g}{R\Gamma}} \quad (9)$$

where p_{TM5} and T_{TM5} are the TM5 surface pressure and temperature, $\Gamma = 6.5 \text{ K km}^{-1}$ the lapse rate, z_{TM5} the TM5 terrain height, and z_s surface elevation for the satellite ground

pixel. The TM5 SO₂ profile is shifted to start at p_s and scaled so that volume mixing ratios are preserved (see Zhou et al., 2009).

Temperature correction

The SO₂ absorption cross sections of Bogumil et al. (2003) show a clear temperature dependence which has an impact on the retrieved SO₂ SCDs depending on the fitting window used. However, only one temperature (203 K) is used for the DOAS fit, therefore a temperature correction needs to be applied: $\text{SCD}' = C_{\text{temp}} \cdot \text{SCD}$. While the SO₂ algorithm provides vertical column results for a set of a priori profiles, applying this correction to the slant column is not simple and as a workaround it is preferred to apply the correction directly to the AMFs (or box AMFs to be precise) while keeping the (retrieved) SCD unchanged: $\text{AMF}' = \text{AMF}/C_{\text{temp}}$. This formulation implicitly assumes that the AMF is not strongly affected by temperature, which is a reasonable approximation (optically thin atmosphere). The correction to be applied requires a temperature profile for each pixel (which is obtained from the TM5 model):

$$C_{\text{temp}} = 1/[1 - \alpha \cdot (T[\text{K}] - 203)], \quad (10)$$

where α equals 0.002, 0.0038 and 0 for the fitting windows 312–326, 325–335 and 360–390 nm, respectively. The parameter α has been determined empirically by fitting Eq. (10) through a set of data points (Fig. 6), for each fitting window. Each value in Fig. 6 is the slope of the fitting line between the SO₂ differential cross sections at 203 K vs. the cross section at a given temperature. In the fitting window 360–390 nm, no temperature correction is applied ($\alpha = 0$) because the cross sections are quite uncertain. Moreover, the 360–390 nm wavelength range is meant for extreme cases (strong volcanic eruptions) for SO₂ plumes in the lower stratosphere, where a temperature of 203 K is a good baseline.

Aerosols

The presence of aerosol in the observed scene (likely when observing anthropogenic pollution or volcanic events) may affect the quality of the SO₂ retrieval (e.g., Yang et al., 2010). No explicit treatment of aerosols (absorbing or not) is foreseen in the algorithm as there is no general and easy way to treat the aerosols effect on the retrieval. At processing time, the aerosol parameters (e.g., extinction profile or single-scattering albedo) are unknown. However, the information on the S-5P UV absorbing aerosol index (AAI) by Zweers (2016) will be included in the L2 SO₂ files as it gives information to the users on the presence of aerosols for both anthropogenic and volcanic SO₂. Nevertheless, the AAI data should be used/interpreted with care. In an offline future version of the SO₂ product, absorbing aerosols might be included in the forward model, if reliable information on

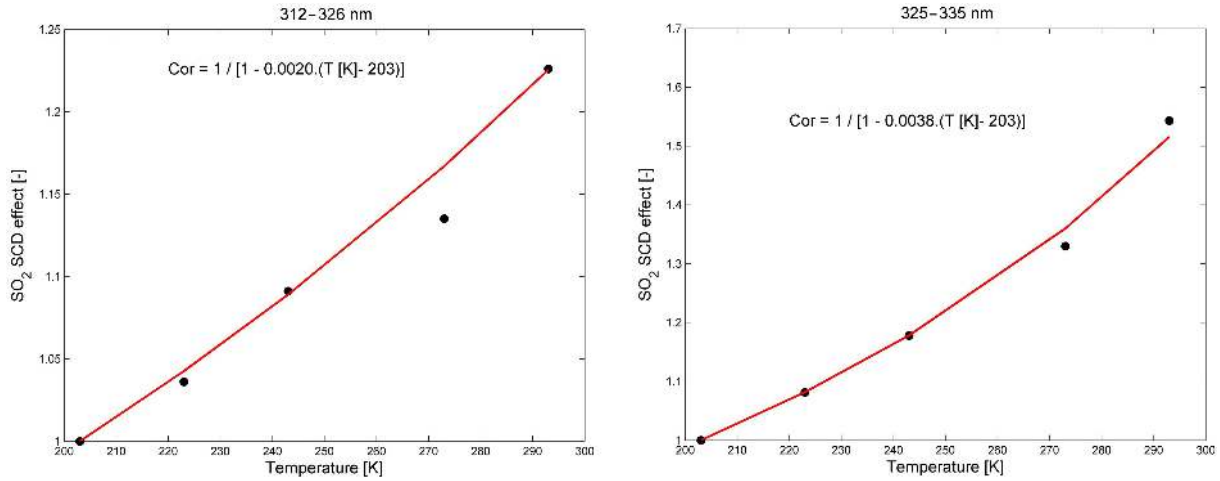


Figure 6. Effect of temperature (relative to 203 K) on SO₂ retrieved SCD for fitting windows 312–326 (left) and 325–335 nm (right). The red lines show the adopted formulation of C_{temp} (Eq. 10). Note that, for the 312–326 nm window, the result at 273K has been discarded from the fit as it seems rather inconsistent with the dependence at other temperatures.

absorbing aerosol can be obtained from the AAI and the S-5P aerosol height product (Sanders and de Haan, 2016).

3 Error analysis

3.1 Introduction

The total uncertainty (accuracy and precision) on the SO₂ columns produced by the algorithm presented in Sect. 2 is composed of many sources of error (see also, e.g., Lee et al., 2009). Several of them are related to the instrument, such as uncertainties due to noise or knowledge of the slit function. These instrumental errors propagate into the uncertainty on the slant column. Other types of error can be considered model errors and are related to the representation of the physics in the algorithm. Examples of model errors are uncertainties on the trace gas absorption cross sections and the treatment of clouds. Model errors can affect the slant column results or the air mass factors.

The total retrieval uncertainty on the SO₂ vertical columns can be derived by error propagation, starting from Eq. (1) and if one assumes uncorrelated retrieval steps (Boersma et al., 2004; De Smedt et al., 2008):

$$\sigma_{N_v}^2 = \left(\frac{\sigma_{N_S}}{M}\right)^2 + \left(\frac{\sigma_{N_S^{back}}}{M}\right)^2 + \left(\frac{(N_S - N_S^{back})\sigma_M}{M^2}\right)^2, \quad (11)$$

where σ_{N_S} and $\sigma_{N_S^{back}}$ are the errors on the slant column N_S and on the background correction N_S^{back} , respectively.

The error analysis is complemented by the total column averaging kernel (AK) as described in Eskes and Boersma (2003):

$$AK(p) = \frac{m'(p)}{M} \quad (12)$$

(m' is the weighting function, Eq. 6), which is often used to characterize the sensitivity of the retrieved column to a change in the true profile.

3.2 Error components

The following sections describe and characterize 20 error contributions to the total SO₂ vertical column uncertainty. These different error components and corresponding typical values are summarized in Tables 5 and 6. Note that, at the time of writing, the precise effect of several S-5P-specific error sources are unknown and will be estimated during operations.

A difficulty in the error formulation presented above comes from the fact that it assumes the different error sources/steps of the algorithm to be independent and uncorrelated, which is not strictly valid. For example, the background correction is designed to overcome systematic features/deficiencies of the DOAS slant column fitting, and these two steps cannot be considered independent. Hence, summing up all the corresponding error estimates would lead to overestimated error bars. Therefore, several error sources will be discussed in the following subsections without giving actual values at this point. Their impact is included and described in later subsections.

Another important point to note is that one should also (be able to) discriminate systematic and random components of

Table 5. Systematic and random error components contributing to the total uncertainty on the SO₂ slant column.

#	Error source	Type*	Parameter uncertainty	Typical uncertainty on SO ₂ SCD
1	SO ₂ absorption cross section	S	6 % (window 1) 6 % (window 2) unknown (window 3)	6 %
2	SO ₂ and O ₃ absorption	S & R		Errors 9 & 10
3	Other atmospheric absorption or interference	S & R		Error 9
4	Radiance shot noise	R	$S/N = 800\text{--}1000$	0.3–0.5 DU (window 1) 5 DU (window 2) 60 DU (window 3)
5	DOAS settings	S	1 nm, polynomial order	< 11 % (window 1) < 6 % (window 2) < 8 % (window 3)
6	Wavelength and radiometric calibration	S	Wavelength calibration Radiometric calibration Additive errors should remain below 2 %	Wavelength calibration and spectral shifts can be corrected by the algorithm to less than 5 % effect on the slant column Intensity offset correction in principle treats (small) radiometric calibration errors
7	Spectral response function		TBD	TROPOMI-specific Expected uncertainty: 10 %
8	Other spectral features		Strongly dependent on interfering signal	–
9	Background correction	S & R		0.2 DU

* R: random; S: systematic.

a given error source V :

$$\sigma_V^2 = \frac{\sigma_{V(\text{rand})}^2}{n} + \sigma_{V(\text{syst})}^2, \quad (13)$$

here n is the number of pixels considered. However, they are hard to separate in practice. Therefore, each of the 20 error contributions are (tentatively) classified as either “random” or “systematic” errors, depending on their tendencies to average out in space/time or not.

3.2.1 Errors on the slant column

Error sources that contribute to the total uncertainty on the slant column originate both from instrument characteristics and uncertainties/limitations on the representation of the physics in the DOAS slant column fitting algorithm. For the systematic errors on the slant column, the numbers provided in Table 5 have been determined based on sensitivity tests (using the QDOAS software).

With all effects summed in quadrature, the various contributions are estimated to account for a systematic error of

about 20 % + 0.2 DU of the background-corrected slant column ($\sigma_{N_s, \text{syst}} = 0.2 \times (N_s - N_s^{\text{back}}) + 0.2 \text{ DU}$).

For the random component of the slant column errors, the error on the slant columns provided by the DOAS fit is considered (hereafter referred to as SCDE) as it is assumed to be dominated by and representative of the different random sources of error.

Error source 1: SO₂ cross section

Systematic errors on slant columns due to SO₂ cross-section uncertainties are estimated to be around 6 % (Vandaele et al., 2009) in window 1 (312–326 nm) and window 2 (325–335 nm) and unknown in window 3 (360–390 nm). In addition, the effect of the temperature on the SO₂ cross sections has to be considered as well. Refer to Sect. 3.2.2 for a discussion of this source of error.

Table 6. Systematic and random error components contributing to the total uncertainty on the SO₂ air mass factor.

#	Error	Type ¹	Parameter uncertainty	Typical uncertainty on the AMF
10	AMF wavelength dependence	S		10 %
11	Model atmosphere	S	O ₃ profile P, T profiles	~ 5–10 % small
12	Forward model	S	< 5 %	< 5 %
13	Surface albedo ²	S	0.02	15 % (PBL) 5 % (FT) 1 % (LS)
14	Cloud fraction ²	R	0.05	5 % (PBL) 15 % (FT) 1 % (LS)
15	Cloud top pressure ²	R	50 hPa	50 % (PBL) 50 % (FT) 1 % (LS)
16	Cloud correction	R		< 5 % on yearly averaged data
17	Cloud model		TBD	
18	SO ₂ profile shape	S		anthropogenic SO ₂ 20–35 % volcanic SO ₂ large (low albedo), < 50 % (high albedo)
19	Aerosol	S & R		Anthropogenic SO ₂ ≈ 15 % (Nowlan et al., 2011). Volcanic SO ₂ (aerosols: ash/sulfate): ~ 20 % (Yang et al., 2010)
20	Temperature correction	R		~ 5 %

¹ R: random; S: systematic. ² Effect on the AMF estimated from Fig. 6.

Error source 2: O₃ and SO₂ absorption

Nonlinear effects due to O₃ absorption are to a large extent accounted for using the Taylor expansion of the O₃ optical depth (Pukite et al., 2010). Remaining systematic biases are then removed using the background correction; hence, residual systematic features are believed to be small (please read also the discussion on errors 9 and 10). The random component of the slant column error contributes to SCDE.

Nonlinear effects due to SO₂ absorption itself (mostly for volcanic plumes) are largely handled by the triple windows retrievals, but – as will be discussed in Sect. 4 – the transition between the different fitting windows is a compromise and there are cases where saturation can still lead to rather

large uncertainties. However, those are difficult to assess on a pixel-to-pixel basis.

Error source 3: other atmospheric absorption/interferences

In some geographical regions, several systematic features in the slant columns remain after the background correction procedure (see discussion on error 9: background correction error) and are attributed to spectral interferences not fully accounted for in the DOAS analysis, such as incomplete treatment of the Ring effect. This effect also has a random component and contributes to the retrieved SCD error (SCDE).

Error source 4: radiance shot noise

It has a major contribution to the SCDE and it can be estimated from typical S/N values of S-5P in UV band 3 (800–1000, according to Veeffkind et al., 2012). This translates to typical SCD random errors of about 0.3–0.5, 5 and 60 DU for window 1, 2 and 3, respectively. Note that real measurements are needed to consolidate these numbers.

Error source 5: DOAS settings

Tests on the effect of changing the lower and upper limits of the fitting windows by 1 nm and the order of the closure polynomial (4 instead of 5) have been performed. Based on a selection of orbits for the Kasatochi eruption (wide range of measured SCDs), the corresponding SCD errors are less than 11, 6 and 8 % for window 1, 2 and 3, respectively.

Error source 6: wavelength and radiometric calibration

Tests on the effect of uncertainties in the wavelength calibration have been performed in the ESA CAMELOT study. The numbers are for a shift of 1/20th of the spectral sampling in the solar spectrum and 1/100th of the spectral sampling in the earthshine spectrum. The shift can be corrected for, but interpolation errors can still lead to a remaining uncertainty of a few percent.

Regarding radiometric calibration, the retrieval result is in principle insensitive to flat (spectrally constant) offsets on the measured radiance because the algorithm includes an intensity offset correction. From the ESA ONTRAQ study it was found that additive error signals should remain within 2 % of the measured spectrum.

Error source 7: spectral response function

Uncertainties in the S-5P instrumental slit functions can lead to systematic errors on the retrieved SO₂ slant columns (to be determined).

Error source 8: other spectral features

Unknown or untreated instrumental characteristics such as stray light and polarization sensitivity can introduce spectral features that may lead to bias in the retrieved slant column data. To a certain extent these can be prevented by the DOAS polynomial and the intensity offset correction settings, as long as the perturbing signals are a smooth function of wavelength. Conversely, high-frequency spectral structures can have potentially a large impact on SO₂ retrievals depending on their amplitude and whether they interfere with SO₂ absorption structures. At the time of writing, it is hard to evaluate these measurement errors. Once the instrument will be operating, such measurement errors will be characterized and correct for, either as part of L1b processor or in the form of pseudo-absorption cross sections in the DOAS analysis.

In the ONTRAQ study, testing sinusoidal perturbation signals showed that the effect of spectral features on the retrieval result depends strongly on the frequency of the signal. Additive signals with an amplitude of 0.05 % of the measurement affect the retrieved SO₂ slant column by up to 30 %. The effect scales more or less linearly with the signal amplitude.

Error source 9: background/dstriping correction

This error source is mostly systematic and important for anthropogenic SO₂ or for monitoring degassing volcanoes. Based on OMI and GOME-2 test retrievals, the uncertainty on the background correction is estimated to be <0.2 DU. This value accounts for limitations of the background correction and is compatible with residual slant columns values typically found (after correction) in some clean areas (e.g., above the Sahara), or for a possible contamination by volcanic SO₂, after a strong eruption.

3.2.2 Errors on the air mass factor

The error estimates on the AMF are listed in Table 6 and are based on simulations and closed-loop tests using the radiative transfer code LIDORT. One can identify two sources of errors on the AMF. First, the adopted LUT approach has limitations in reproducing the radiative transfer in the atmosphere (forward model errors). Secondly, the error on the AMF depends on input parameter uncertainties. This contribution can be broken down into a squared sum of terms (Boersma et al., 2004):

$$\sigma_M^2 = \left(\frac{\partial M}{\partial \text{alb}} \cdot \sigma_{\text{alb}} \right)^2 + \left(\frac{\partial M}{\partial \text{ctp}} \cdot \sigma_{\text{ctp}} \right)^2 + \left(\frac{\partial M}{\partial \text{feff}} \cdot \sigma_{\text{feff}} \right)^2 + \left(\frac{\partial M}{\partial s} \cdot \sigma_s \right)^2, \quad (14)$$

where σ_{alb} , σ_{ctp} , σ_{feff} , σ_s are typical uncertainties on the albedo, cloud top pressure, cloud fraction and profile shape, respectively.

The contribution of each parameter to the total air mass factor error depends on the observation conditions. The air mass factor sensitivities ($\frac{\partial M}{\partial \text{parameter}}$), i.e., the air mass factor derivatives with respect to the different input parameters, can be derived for any particular condition of observation using the altitude-dependent AMF LUT, created with LIDORTv3.3, and using the a priori profile shapes. In practice, a LUT of AMF sensitivities has been created using reduced grids from the AMF LUT and a parameterization of the profile shapes based on the profile shape height.

Error source 10: AMF wavelength dependence

Because of strong atmospheric absorbers (mostly ozone) and scattering processes, the SO₂ AMF shows a wavelength dependence. We have conducted sensitivity tests to determine

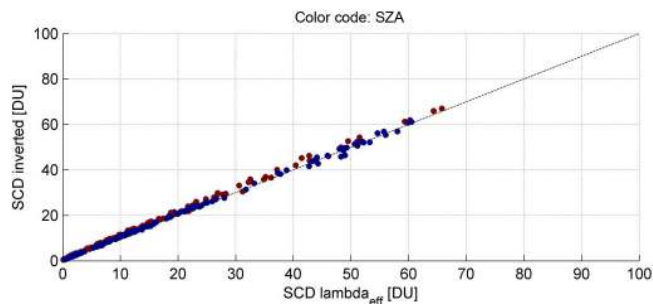


Figure 7. Retrieved SO₂ slant columns versus simulated SCDs at a wavelength of 313 nm from synthetic spectra (SZA: 30, 70°) in the spectral range 312–326 nm and for SO₂ layers in the boundary layer, upper troposphere and lower stratosphere. The different points correspond to different values for the line-of-sight angle (0, 45°), surface albedo (0.06, 0.8), surface height (0, 5 km) and total ozone column (350, 500 DU). SO₂ vertical columns as input of the RT simulations are a maximum of 25 DU.

the optimal wavelengths for AMF calculations representative of each of the three fitting windows. To do so, synthetic radiances and SO₂ SCDs have been generated using LIDORT for typical observations scenarios and at spectral resolution and sampling compatible with S-5P. The spectra have been analyzed by DOAS and the retrieved SCDs have been compared to the calculated SCDs at different wavelengths. It appears from this exercise that 313, 326 and 375 nm provide the best results, for window 1, 2 and 3, respectively. Figure 7 shows an illustration of these sensitivity tests in the baseline window; an excellent correlation and slope close to 1 is found for the scatter plot of retrieved versus simulated slant columns using an effective wavelength of 313 nm for the AMF. Overall, for low solar zenith angles, the deviations from the truth are less than 5 % in most cases, except for boundary layer (BL) SO₂ at a 1 DU column level and for low-albedo scenes (deviations up to 20 %). For high solar zenith angles deviations are less than 10 % in most cases, except for BL SO₂ at a 1 DU column level and for low-albedo scenes (underestimation by up to a factor of 2).

Error source 11: model atmosphere

This error relates to uncertainties in the atmospheric profiles used as input of LIDORT for the weighting function look-up-table calculations.

Although the effect of O₃ absorption on the AMF is treated in the algorithm, the O₃ profiles used as input of LIDORT are not fully representative of the real profiles and typical errors (including error due to interpolation) of 5–10 % can occur.

A test has been performed by replacing the US standard atmosphere pressure and temperature profiles by high latitude winter profiles and the impact on the results is found to be small.

Error source 12: radiative transfer model

This error source is believed to be small, less than 5 % (Hendrick et al., 2006; Wagner et al., 2007).

Error source 13: surface albedo

A typical uncertainty on the albedo is 0.02 (Kleipool et al., 2008). This translates to an error on the air mass factor after multiplication by the slope of the air mass factor as a function of the albedo (Eq. 14) and can be evaluated for each satellite pixel. As an illustration, Fig. 8 shows the expected dependence of the AMF with albedo and also with the cloud conditions. From Fig. 8a, one concludes that the retrievals of SO₂ in the BL are much more sensitive to the exact albedo value than for SO₂ higher up in the atmosphere, for this particular example.

More substantial errors can be introduced if the real albedo differs considerably from what is expected, for example in the case of the sudden snowfall or ice cover. The snow/ice cover flag in the L2 file will therefore be useful for such cases.

Error source 14: cloud fraction

An uncertainty on the cloud fraction of 0.05 is considered. The corresponding AMF error can be estimated through Eq. (14; see Fig. 8b) or by analytic derivation from Eqs. (6)–(8).

Error source 15: cloud top pressure

An uncertainty on the cloud top height of 0.5 km (~ 50 hPa) is assumed. The corresponding AMF error can be estimated through Eq. (14). Figure 8c illustrates the typical behavior of signal amplification/shielding for a cloud below/above the SO₂ layer. One can see that the error (slope) dramatically increases when the cloud is at a height similar to the SO₂ bulk altitude.

Error source 16: cloud correction

Sensitivity tests showed that applying the independent pixel approximation or assuming cloud-free pixels makes a difference of only 5 % on yearly averaged data (for anthropogenic BL SO₂ VC with cloud fractions less than 40 %).

Error source 17: cloud model

Cloud as layer (CAL) is the baseline of the S-5P cloud algorithm, but a Lambertian equivalent reflector (LER) implementation will be used for NO₂, SO₂ and HCHO retrievals. The error due to the choice of the cloud model will be evaluated during the operational phase.

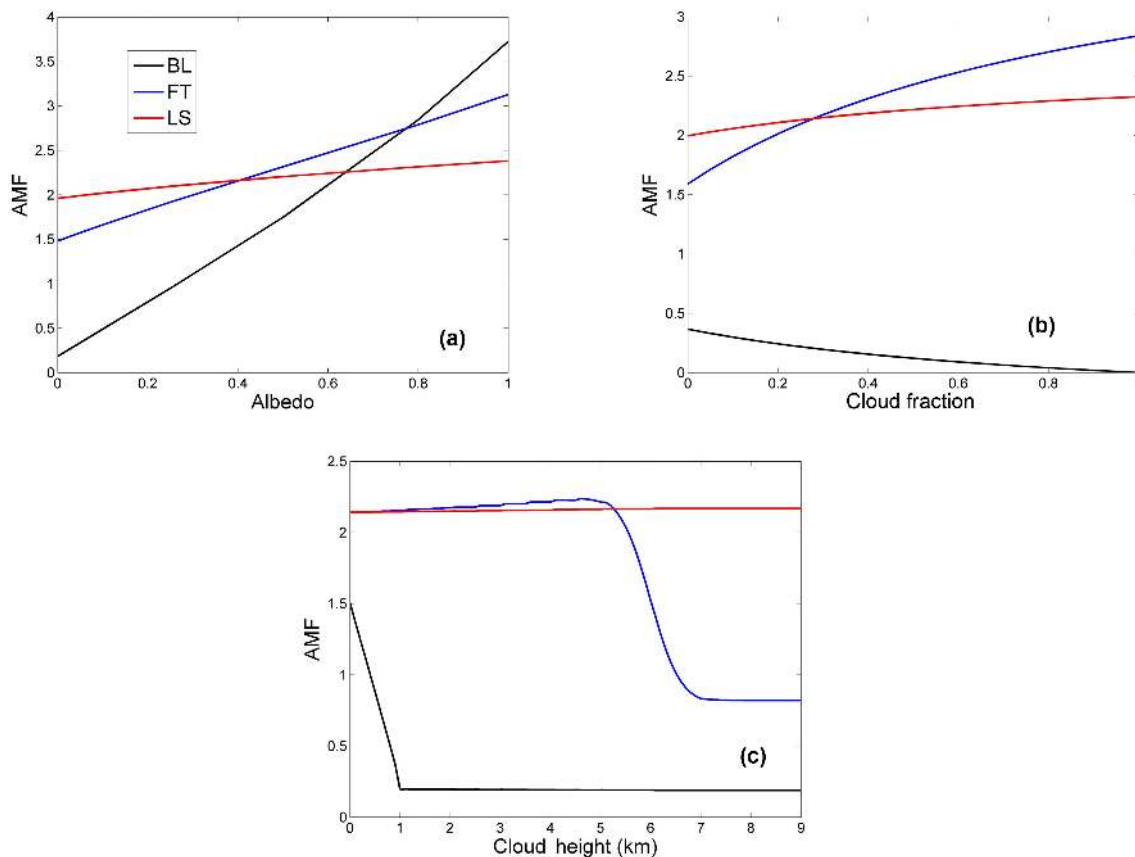


Figure 8. Air mass factors at 313 nm for SO₂ in the boundary layer (BL: 0–1 km), free troposphere and lower stratosphere (FT, LS: Gaussian profiles with maximum height at 6, 15 km; FWHM: 1 km). Calculations are for SZA = 40°, Los = 10°, RAA = 0° and surface height = 0 km. AMFs are displayed as a function of the (a) albedo for clear-sky conditions, (b) cloud fraction for albedo = 0.06, cloud albedo = 0.8 and cloud top height = 2 km and (c) cloud top height for albedo = 0.06, cloud albedo = 0.8 and cloud fraction = 0.3.

Error source 18: profile shape

A major source of systematic uncertainty for most SO₂ scenes is the shape of the vertical SO₂ distribution. The corresponding AMF error can be estimated through Eq. (14) and estimation of uncertainty on the profile shape. Note that vertical columns are provided with their averaging kernels, so that column data might be improved for particular locations by using more accurate SO₂ profile shapes based on input from models or observations.

For anthropogenic SO₂ under clear-sky conditions, sensitivity tests using a box profile from 0 to 1 ± 0.5 km above ground level, or using the different profiles from the CAMELOT study (Levelt et al., 2009), give differences in AMFs in the range of 20–35%. Note that for particular conditions SO₂ may also be uplifted above the top of the boundary layer and sometimes reach upper-tropospheric levels (e.g., Clarisse et al., 2011). SO₂ weighting functions displayed in Fig. 5 show that the measurement sensitivity is then increased by up to a factor of 3 and therefore constitutes a major source of error.

In the SO₂ algorithm, the uncertainty on the profile shape is estimated using one parameter describing the shape of the TM5 profile: the profile height, i.e., the altitude (pressure) below which resides 75% of the integrated SO₂ profile. $\frac{\partial M}{\partial s}$ is approached by $\frac{\partial M}{\partial s_h}$, where s_h is half of the profile height. Relatively small variations in this parameter have a strong impact on the total air mass factors for low-albedo scenes, because altitude-resolved air mass factors decrease strongly in the lower troposphere, where the SO₂ profiles peak (see, e.g., Fig. 5).

For volcanic SO₂, the effect of the profile shape uncertainty depends on the surface or cloud albedo. For low-albedo scenes (Fig. 5a), if no external information on the SO₂ plume height is available, it is a major source of error at all wavelengths. Vertical columns may vary by up to a factor of 5. For high-albedo scenes (Fig. 5b), the error is less than 50%. It should be noted that these conditions are often encountered for strong eruptions injecting SO₂ well above the cloud deck (high reflectivity). Further uncertainty on the retrieved SO₂ column may arise if the vertical distribution shows dis-

tinct layers at different altitudes, due to the different nature of successive phases of the eruption.

In the SO₂ algorithm, three 1 km thick box profiles are used in the AMF calculations, mostly to represent typical volcanic SO₂ profiles. The error due to the profile shape uncertainty is estimated by varying the box center levels by 100 hPa.

Error source 19: aerosols

The effect of aerosols on the air mass factors are not explicitly considered in the SO₂ retrieval algorithm. To some extent, however, the effect of the non-absorbing part of the aerosol extinction is implicitly included in the cloud correction (Boersma et al., 2004). Indeed, in the presence of aerosols, the cloud detection algorithm is expected to overestimate the cloud fraction, resulting partly in a compensation effect for cases where aerosols and clouds are at similar heights. Absorbing aerosols have a different effect on the air mass factors and can lead to significant errors for high aerosol optical depths (AODs). In the TROPOMI SO₂ product, the absorbing aerosol index field can be used to identify observations with elevated absorbing aerosols.

Generally speaking, the effect of aerosols on AMF is highly variable and strongly depends on aerosol properties (AOD, height and size distribution, single-scattering albedo, scattering phase function, etc.). Typical AMFs uncertainties due to aerosols found in the literature are given in Table 6. As aerosols affect cloud fraction, cloud top height and to some extent the albedo database used, correlations between uncertainties on these parameters are to be expected.

Error source 20: temperature correction

The DOAS scheme uses an SO₂ cross section at only one temperature (Bogumil et al., 2003, at 203 K), which is in general not representative of the effective temperature corresponding to the SO₂ vertical profile. This effect is in principle accounted for by the temperature correction (which is applied in practice to the AMFs; see Sect. 2.2.3.7) but with a certain associated error of ~ 5 %.

4 Verification

The SO₂ retrieval algorithm presented in Sect. 2, and hereafter referred as “prototype algorithm”, has been applied to OMI and GOME-2 spectra. The results have been extensively verified and validated against different satellite and ground-based data sets (e.g., Theys et al., 2015; Fioletov et al., 2016; Wang et al., 2016). Here we report on further scientific verification activities that took place during the ESA S-5P L2WG project.

In addition to the prototype algorithm, a scientific algorithm (referred as “verification algorithm”) has been developed in parallel. Both algorithms have been applied to syn-

thetic and real (OMI) spectra and results were compared. In this study, we only present and discuss a selection of results (for OMI).

4.1 Verification algorithm

The S-5P TROPOMI verification algorithm was developed in close cooperation between the Max Planck Institute for Chemistry (MPIC) in Mainz (Germany) and the Institut für Methodik und Fernerkundung as part of the Deutsches Institut für Luft- und Raumfahrt Oberpfaffenhofen (DLR-IMF). Like the prototype algorithm (PA), the verification algorithm (VA) uses a multiple fitting window DOAS approach to avoid nonlinear effects during the SCD retrieval in the case of high SO₂ concentrations in volcanic plumes. However, especially the alternatively used fitting windows differ strongly from the ones used for the PA and are entirely located in the lower UV range:

- 312.1–324 nm (standard retrieval – SR): similar to baseline PA fitting window, ideal for small columns
- 318.6–335.1 nm (medium retrieval – MR): this fitting window is essentially located in between the first and second fitting window of the PA and was mainly introduced to guarantee a smoother transition between the baseline window and the one used for high SO₂ concentrations. The differential SO₂ spectral features are still about 1 order of magnitude smaller than in the baseline window.
- 323.1–335.1 nm (alternative retrieval – AR): similar to the intermediate fitting window of the PA. This fitting window is used in the case of high SO₂ concentrations. Although it is expected that volcanic events with extreme SO₂ absorption are still affected by nonlinear absorption in this window, the wavelength range is sufficient for most volcanic events.

Furthermore, the VA selection criteria for the transition from one window to another are not just based on fixed SO₂ SCD thresholds. The algorithm allows for a slow and smooth transition between different fit ranges by linearly decreasing the weight of the former fitting window and at the same time increasing the weight of the following fitting window:

1. for SO₂ SCD ≤ 4 × 10¹⁷ molec cm⁻² (≈ 15 DU):

$$\text{SO}_2 \text{ SCD} = \text{SR};$$

2. for 4 × 10¹⁷ molec cm⁻² < SO₂ SCD < 9 × 10¹⁷ molec cm⁻²:

$$\text{SO}_2 \text{ SCD} = \text{SR} \times \left[1 - \frac{\text{SR}}{9 \times 10^{17} \text{ molec cm}^{-2}} \right] + \text{MR} \times \left[\frac{\text{SR}}{9 \times 10^{17} \text{ molec cm}^{-2}} \right];$$

3. for SO₂ SCD $\geq 9 \times 10^{17}$ molec cm⁻² (≈ 33 DU):

$$\text{SO}_2 \text{ SCD} = \text{MR};$$

4. for 9×10^{17} molec cm⁻² < SO₂ SCD < 4.6×10^{18} molec cm⁻²:

$$\text{SO}_2 \text{ SCD} = \text{MR} \times \left[1 - \frac{\text{MR}}{4.6 \times 10^{18} \text{ molec cm}^{-2}} \right] + \text{AR} \times \left[\frac{\text{AR}}{4.6 \times 10^{18} \text{ molec cm}^{-2}} \right];$$

5. for SO₂ SCD $\geq 4.6 \times 10^{18}$ molec cm⁻² (≈ 171 DU):

$$\text{SO}_2 \text{ SCD} = \text{AR}.$$

To convert the final SO₂ SCDs into vertical column densities, a single-wavelength AMF for each of the three fitting windows (SO₂ SR, MR and AR) is calculated using the LIDORT LRRS v2.3 (Spurr et al., 2008). The AMF depends on the viewing angles and illumination, surface and cloud conditions as well as on the O₃ total column, which is taken from the O₃ total column retrieval. A cloudy and clear-sky AMF is calculated using temperature-dependent cross sections for SO₂ (Bogumil et al., 2003) and O₃ (Brion et al.,

1984): $\text{AMF}(\lambda) = \frac{\ln\left(\frac{I_{+\text{SO}_2}}{I_{-\text{SO}_2}}\right)}{\tau_{\text{SO}_2}}$ with ($I_{+\text{SO}_2}$) and ($I_{-\text{SO}_2}$) being simulated earthshine spectra with and without including SO₂ as a trace gas, respectively. Both AMFs are combined using the cloud fraction information. Like the PA, the VA is calculated for different a priori SO₂ profiles (center of mass at 2.5, 6 and 15 km) and a temperature correction is applied (see Sect. 2.2.3.7). In contrast to the PA the VA uses Gaussian-shaped SO₂ profiles with a FWHM of 2.5 km rather than box profiles as in the PA. This choice, however, has only a minor influence on the AMF.

For further details on the VA, the reader is referred to the S-5P Science Verification Report (available at https://earth.esa.int/web/sentinel/user-guides/sentinel-5p-tropomi/document-library/-/asset_publisher/w9Mnd6VPjXlc/content/sentinel-5p-tropomi-science-verification-report) for more detailed description and results.

4.2 Verification results

For the intercomparison, the prototype algorithm and verification algorithm were applied to OMI data for three different SO₂ emission scenarios: moderate volcanic SO₂ VCDs on 1 May 2005, caused by the eruption of the Anatahan volcano, elevated anthropogenic SO₂ VCDs, on 1 May 2005, from the Norilsk copper smelter (Russia), and strongly enhanced SO₂ VCDs, on 8 August 2008, after the massive eruption of Mt. Kasatochi.

In the following, both algorithms use the same assumption of an SO₂ plume located at 15 km altitude for the AMF calculation. Even if this choice is not realistic for some of the presented scenarios, it minimizes the influence of differences in

the a priori settings. Main deviations between prototype and verification algorithm are therefore expected to be caused by the usage of different fit windows (determining their sensitivity and fit error) and especially the corresponding transition criteria.

Figure 9 shows the resulting maps of the SO₂ VCD for the VA (upper panels) and PA (lower panels) for the three selected test cases. As can be seen, both algorithms result in similar SO₂ VCDs; however, a closer look reveals some differences, such as the maximum VCDs, which are not necessarily appearing at the same locations. For the Anatahan case for instance, the maximum VCD is seen closer to the volcano at the eastern end of the plume for the PA, while it appears to be further downwind for the VA. This effect can be explained by the corresponding fit windows used for both algorithms, which may result in deviating SO₂ VCDs, especially for SO₂ scenarios where the best choice is difficult to assess. This is illustrated in Fig. 10, showing scatter plots of VA versus PA SO₂ VCDs for the three test cases (Anatahan, Norilsk and Kasatochi) color-coded differently depending on the fitting window used for VA (left) and PA (right), respectively. While the PA uses strictly separated results from the individual fit windows, the VA allows a smooth transition whenever resulting SO₂ SCDs are found to be located in between subsequent fit ranges.

For all three test cases, it appears that the PA is less affected by data scattering for low SO₂ or SO₂ free measurements than the VA. For the shortest UV fit windows, both algorithms mainly agree, but VA VCDs tend to be higher by 10–15 % than the PA VCDs for the Anatahan and Kasatochi measurements but interestingly not for the Norilsk case. For SO₂ VCDs around 7 DU the PA seems to be slightly affected by saturation effects in 312–326 nm window, while VA already makes use of a combined SR/MR SCD. For larger SO₂ VCDs (> 10 DU), data sets from both algorithms show an increased scattering, essentially resulting from the more intensive use of fitting windows at longer wavelengths (for which the SO₂ absorption is weaker). While it is difficult to conclude which algorithm is closer to the actual SO₂ VCDs, the combined fit windows of the VA probably are better suited (in some SO₂ column ranges) for such scenarios as the SO₂ cross section is generally stronger for lower wavelength (< 325 nm) when compared to the intermediate fit window of the PA.

For extremely high SO₂ loadings, i.e., for the Kasatochi plume on 8 August 2008, the DOAS retrievals from PA and VA require all three fit windows to prevent systematic underestimation of the resulting SO₂ SCDs due to nonlinear absorption caused by very high SO₂ concentrations within the volcanic plume. Figure 9 (right panel) shows that the SO₂ distribution is similar for both algorithms, including the location of the maximum SO₂ VCD.

From Fig. 10 (lowest panel), it can be seen that the VA shows higher values for SO₂ VCDs < 100 DU, for all three fit windows. For very high SO₂ VCDs, it seems that the ver-

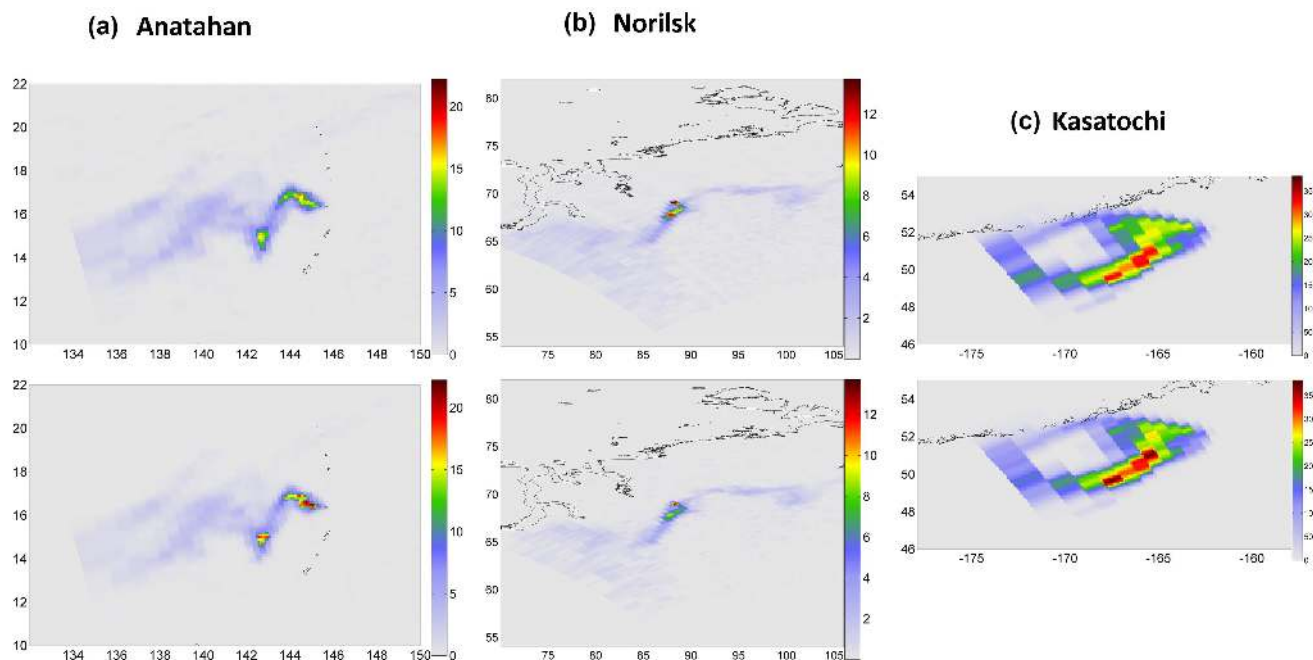


Figure 9. OMI SO₂ VCD (expressed in DU) for the verification (upper panels) and prototype algorithms (lower panels) for the three selected scenarios: during the Anatahan eruption (left), over the Norilsk copper smelter area (center) and for the volcanic eruption of Kasatochi (right). Note that, for each case, the color bar has been scaled to the maximum SO₂ VCD from both algorithms.

ification algorithm is already slightly affected by an underestimation of the SO₂ VCD caused by nonlinear radiative transfer effects in the SO₂ AR fit window, while the PA retrievals in the 360–390 nm fit range are insensitive to saturation effects. We note, however, that the Kasatochi plume also contained significant amounts of volcanic ash and we cannot rule out a possible retrieval effect of volcanic ash on the observed differences between PA and VA SO₂ results. Finally, we have also investigated other cases with extreme concentrations of SO₂, and contrasting results were found compared to the Kasatochi case. For example, on 4 September 2014, PA retrieved up to 260 DU of SO₂ during the Icelandic Bárðarbunga fissure eruption, while VA only found 150 DU (not shown). Compared to Kasatochi, we note that this specific scenario is very different as for the plume height (the SO₂ plume was typically in the lowermost troposphere \sim 3 km a.s.l.) and it is likely to play a role in the discrepancy between PA and VA results.

In summary, we found that the largest differences between prototype and verification algorithms are due to the fitting window transitions and differences of measurement sensitivity of the fitting windows used (all subject differently to nonlinear effects). Verification results have shown that the prototype algorithm produces reasonable results for all the expected scenarios, from modest to extreme SO₂ columns, and are therefore adequate for treating the TROPOMI data. In a future processor update, the method could, however, be refined.

5 Validation of TROPOMI SO₂ product

In this section, we give a brief summary of possibilities (and limitations) to validate the TROPOMI SO₂ product with independent measurements.

Generally speaking, the validation of a satellite SO₂ column product is a challenge for several reasons, on top of which is the representativeness of the correlative data when compared to the satellite retrievals. Another reason comes from the wide range of SO₂ columns in the atmosphere that vary from about 1 DU level for anthropogenic SO₂ and low-level volcanic degassing to 10–1000 DU for medium to extreme volcanic explosive eruptions.

The space-borne measurement of anthropogenic SO₂ is difficult because of the low column amount and reduced measurement sensitivity close to the surface. The SO₂ signal is covered by the competing O₃ absorption and the column accuracy is directly affected by the quality of the background correction applied. Among the many parameters of the SO₂ retrieval algorithm that affect the results, the SO₂ vertical profile shape is of utmost importance for any comparison with correlative data. The SO₂ column product accuracy is also directly impacted by the surface albedo used as input for the AMF calculation, the cloud correction/filtering and aerosols. In principle, all these effects will have to be addressed in future validation efforts.

The measurement of volcanic SO₂ is facilitated by SO₂ columns often larger than for anthropogenic SO₂. However,

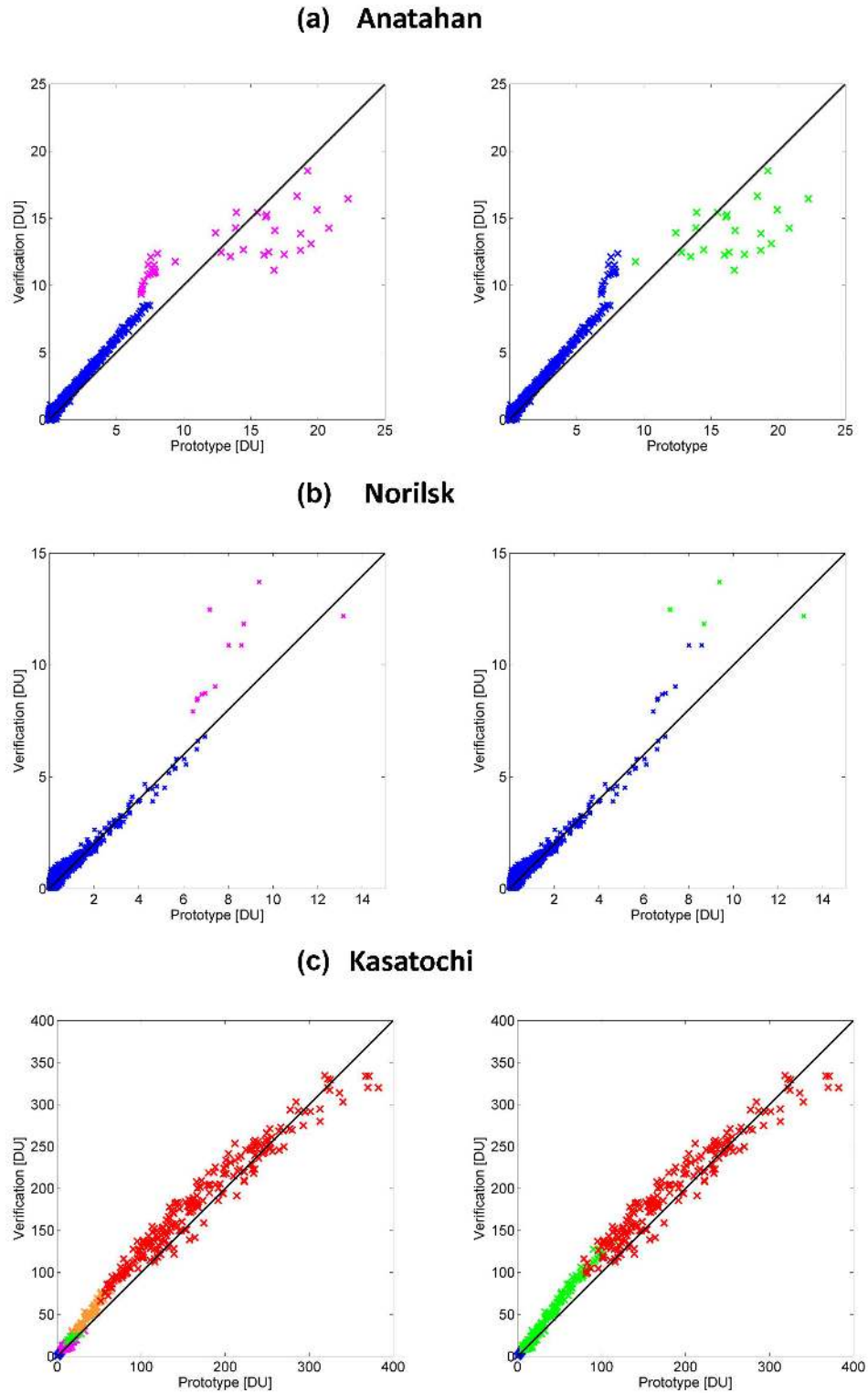


Figure 10. OMI SO₂ VCD (DU) scatter plots for PA (x axis) and VA (y axis) for the three test cases: the Anatahan eruption, Norilsk anthropogenic emissions and the Kasatochi eruption (from top to bottom). The different fit windows used for both algorithms are color-coded: VA on left panels (blue: SR; purple: SR/MR; green: MR; orange: MR/AR; red: AR), PA on right panels (blue: 312–326 nm; green: 325–335 nm; red: 360–390 nm). For the three scenarios, the prototype and verification algorithms agree fairly well with $r^2 \sim 0.9$.

the total SO₂ column is strongly dependent on the height of the SO₂ plume, which is highly variable and usually unknown. For most volcanoes, there is no ground-based equipment to measure SO₂ during an appreciable eruption and even if it is the case, the data are generally difficult to use for validation. For strong eruptions, volcanic plumes are transported over long distances and can be measured by ground-based and aircraft devices, but generally there are only a handful of data sets available and the number of coincidences is rather small.

For both anthropogenic and volcanic SO₂ measurements, the vertical distribution of SO₂ is a key parameter limiting the product accuracy. If reliable (external) information on the SO₂ profile (or profile shape) is available, it is recommended to recalculate the SO₂ vertical columns by using this piece of information and the column averaging kernels that can be found in the TROPOMI SO₂ L2 files.

5.1 Ground-based measurements

When considering the application of ground-based instruments for the validation of satellite SO₂ observations, several types of instruments are to be considered.

Brewer instruments have the advantage to operate as part of a network (<http://www.woudc.org>), but the retrieved SO₂ columns are generally found inaccurate for the validation of anthropogenic SO₂. Yet in some cases they might be used for coincidences with volcanic clouds, typically for SO₂ VCDs larger than 5–10 DU. Multi-axis DOAS (MAX-DOAS) or direct-sun DOAS measurements (e.g., from Pandora instruments) can be used to validate satellite SO₂ columns from anthropogenic emissions (e.g., Theys et al., 2015; Jin et al., 2016; Wang et al., 2016), but caution must be exercised in the interpretation of the results because realistic SO₂ profile shapes must be used by the satellite retrieval scheme. While direct-sun DOAS retrievals are independent of the SO₂ profile shape, MAX-DOAS observations carry information on the SO₂ vertical distribution, but it is not obvious that the technique is directly applicable to the validation of satellite SO₂ retrievals, because the technique is not able to retrieve the full SO₂ profile. Another important limitation comes from the fact that ground-based DOAS and satellite instruments have very different fields of view and are therefore probing different air masses. This can cause a large discrepancy between ground-based and satellite measurements in the case of strong horizontal gradients of the SO₂ column field. DOAS instruments scanning through volcanic plumes are now routinely measuring volcanic SO₂ emissions, as part of the Network for Observation of Volcanic and Atmospheric Change (NOVAC; Galle et al., 2010), for an increasing number of degassing volcanoes. Ongoing research focusses on calculating SO₂ fluxes from those measurements and accounting for nontrivial radiative transfer effects (e.g., light dilution; see Kern et al., 2009). NOVAC flux data could be used for comparison with TROPOMI SO₂ data,

but this requires techniques to convert satellite SO₂ vertical column into mass fluxes (see, e.g., Theys et al., 2013, and references therein; Beirle et al., 2014). Similarly, fast-sampling UV cameras are becoming increasingly used to measure and invert SO₂ fluxes and are also relevant to validate TROPOMI SO₂ data over volcanoes or anthropogenic point sources (e.g., power plants). It should be noted, however, that ground-based remote sensing instruments operating nearby SO₂ point sources are sensitive to newly emitted SO₂ plumes, while a satellite sensor like TROPOMI will measure aged plumes that have been significantly depleted in SO₂. While in some cases it is possible to compensate for this effect by estimating the SO₂ lifetime, e.g., directly from the space measurements (Beirle et al., 2014), the general situation is that the SO₂ loss rate is highly variable (especially in volcanic environments), and this can lead to strong discrepancies when comparing satellite and ground-based SO₂ fluxes.

In addition to optical devices, there are also in situ instruments measuring surface SO₂ mixing ratios. This type of instrument can only validate surface concentrations, and additional information on the SO₂ vertical profile (e.g., from model data) is required to make the link with the satellite retrieved column. However, in situ instruments are being operated for pollution monitoring in populated areas, and allow for extended and long-term comparisons with satellite data (see, e.g., Nowlan et al., 2011).

5.2 Aircraft and mobile measurements

Airborne and mobile instruments provide valuable and complementary data for satellite validation.

In the case of volcanic explosive eruptions, satisfactory validation results can be obtained by comparing satellite and fixed ground DOAS measurements of drifting SO₂ plumes, as shown by Spinei et al. (2010), but the comparison generally suffers from the small number of coincidences. Dedicated aircraft campaign flights (e.g., Schumann et al., 2011) can in principle improve the situation. Their trajectory can be planned with relative ease to cross sustained eruptive plumes. However, localized high SO₂ concentrations may be carried away too quickly to be captured by aircraft or have diluted below the threshold limit for satellite detection before an aircraft can respond. An important database of SO₂ aircraft measurements is provided by the CARIBIC/IAGOS project, which exploits automated scientific instruments operating long-distance commercial flights. Measurements of volcanic SO₂ during the eruptions of Mt. Kasatochi and Eyjafjallajökull and comparison with satellite data have been reported by Heue et al. (2010, 2011).

An attempt to validate satellite SO₂ measurements using a mobile DOAS instrument for a fast moving (stratospheric) volcanic SO₂ plume was presented by Carn and Lopez (2011). Although the agreement between both data sets was found reasonable, the comparison was complicated

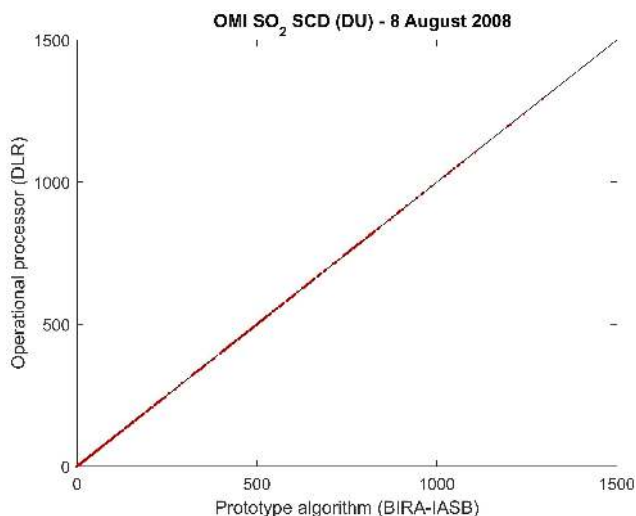


Figure 11. Comparison of SO₂ SCDs between prototype algorithm and operational processor for the OMI test data of 8 August 2008.

by the relatively fast displacement of the volcanic cloud with respect to the ground spectrometer and clear heterogeneity on scales smaller than a satellite pixel. For degassing volcanoes or new fissure eruptions, mobile DOAS traverse measurements under the plume offer unique opportunities to derive volcanic SO₂ fluxes that could be used to validate satellite measurements.

For polluted regions, measurements of anthropogenic SO₂ by airborne nadir-looking DOAS sensors are able to produce high-spatial-resolution mapping of the SO₂ column field (e.g., during the AROMAT campaigns, <http://uv-vis.aeronomie.be/aromat/>) that could be used to validate TROPOMI SO₂ product or give information on horizontal gradients of the SO₂ field (e.g., in combination with coincident mobile DOAS measurements) that would be particularly useful when comparing satellite and MAX-DOAS data (see discussion in Sect. 5.1). Equally important are also limb-DOAS or in situ instruments to provide information on the vertical distribution of SO₂ which is crucial for satellite validation (e.g., Krotkov et al., 2008).

5.3 Satellite measurements

Intercomparison of satellite SO₂ measurements generally provides a convenient and easy way to evaluate at a glance the quality of a satellite product, for instance by comparing SO₂ maps. Often, it also provides improved statistics and geographical representativeness, but it poses a number of problems because, when different satellite sensors are compared, they have also different overpass times, swaths, spatial resolutions and measurement sensitivities to SO₂.

For volcanic SO₂, satellite measurements often provide the only data available for the first hours to days after an eruption event and satellite intercomparison is thus the only prac-

tical way to assess the quality of the retrievals. To overcome sampling issues mentioned above, intercomparison of SO₂ masses integrated over the measured volcanic plume is often performed. For TROPOMI, current satellite instruments will be an important source of data for cross-comparisons. Although non-exhaustive, the following is a list of satellite sensors that could be used: OMI, OMPS, GOME-2 and IASI (MetOp-A, -B, and the forthcoming -C), AIRS, CrIS, VIIRS and MODIS. As mentioned above, the intercomparison of satellite SO₂ products is difficult and in this respect the plume altitude is a key factor of the satellite SO₂ data accuracy. Comparison of TROPOMI and other satellite SO₂ products will benefit not only from the advent of scientific algorithms for the retrieval of SO₂ plume heights but also from the use of volcanic plume height observations using space lidar instruments (e.g., CALIOP and the future EarthCare mission).

For both anthropogenic SO₂ and volcanic degassing SO₂, the satellite UV sensors OMI, GOME-2 and OMPS can be compared to TROPOMI SO₂ data by averaging data over certain polluted regions. This procedure will give valuable information on the data quality, but, in some cases, the comparison will suffer from differences in spatial resolution. A more robust and in-depth comparison would be to use different TROPOMI SO₂ data sets generated by different retrieval algorithms and investigate the differences in the various retrieval steps (spectral fitting, corrections, radiative transfer simulations, error analysis).

6 Conclusions

Based on the heritage from GOME, SCIAMACHY, GOME-2 and OMI, a DOAS retrieval algorithm has been developed for the operational retrieval of SO₂ vertical columns from TROPOMI level 1b measurements in the UV spectral range. Here we describe its main features.

In addition to the traditionally used fitting window of 312–326 nm, the new algorithm allows for the selection of two additional fitting windows (325–335 and 360–390 nm), reducing the risk of saturation and ensuring accurate SO₂ column retrieval even for extreme SO₂ concentrations as observed for major volcanic events. The spectral fitting procedure also includes an advanced wavelength calibration scheme and a spectral spike removal algorithm.

After the slant column retrieval, the next step is a background correction, which is empirically based on the O₃ slant column (for the baseline fitting window) and across-track position and accounts for possible across-track dependencies and instrumental degradation.

The SO₂ slant columns are then converted into vertical columns by means of air mass factor calculations. The latter is based on weighting function look-up tables with dependencies on the viewing geometry, clouds, surface pressure, albedo and ozone and is applied to pre-defined box profiles

and TM5 CTM forecast profiles. In addition, the algorithm computes DOAS-type averaging kernels and a full error analysis of the retrieved columns.

In this paper we have also presented verification results using an independent algorithm for selected OMI scenes with enhanced SO₂ columns. Overall, the prototype algorithm agrees well with the verification algorithm, demonstrating its ability in retrieving accurately medium to very high SO₂ columns. We have discussed the advantages and limitations of both prototype and verification algorithms.

Based on the experience with GOME-2 and OMI, the TROPOMI SO₂ algorithm is expected to have a comparable level of accuracy. Due to its high signal-to-noise ratio, TROPOMI will be capable of at least achieving comparable retrieval precision as its predecessors but at a much finer spatial resolution of 7 km × 3.5 km at best. For single measurements, the user requirements for tropospheric SO₂ concentrations will not be met, but improved monitoring of strong pollution and volcanic events will be possible by spatial and temporal averaging the increased number of observations of TROPOMI. Nevertheless, it will require significant validation work and here we have discussed some of the inherent challenges for both volcanic and anthropogenic SO₂ retrievals. Correlative measurements from ground-based, aircraft/mobile, and satellite instruments will be needed over different regions and various emission scenarios to assess and characterize the quality of TROPOMI SO₂ retrievals.

The baseline algorithm presented here, including all its modules (slant column retrieval, background correction, air mass factor calculation and error analysis), has been fully implemented in the S-5P operational processor UPAS by the DLR team. Figure 11 illustrates the status of the implementation for one day of OMI test data, as an example for the slant columns retrievals. A nearly perfect agreement is found between SCD results over 4 orders of magnitude. A similar match between prototype algorithm and operational processor is found for all other retrieval modules.

For more information on the TROPOMI SO₂ L2 data files, the reader is referred to the S-5P SO₂ Product User Manual (Pedergrana et al., 2016).

7 Data availability

The TROPOMI SO₂ retrieval algorithm has been tested on OMI L1 and L2 operational data, publicly available from the NASA Goddard Earth Sciences (GES) Data and Information Services Center (<http://disc.sci.gsfc.nasa.gov/Aura/OMI/omso2.shtml>). The static auxiliary datasets used as input of the TROPOMI SO₂ retrieval algorithm are publicly available. The links to the data sets are in the references included in Table A2. Other underlying research data are available upon request from Nicolas Theys (theys@aeronomie.be).

Appendix A: Feasibility, information on data product and ancillary data

High-level data product description

In addition to the main product results, such as SO₂ slant column, vertical column and air mass factor, the level 2 data files will contain several additional parameters and diagnostic information. Table A1 gives a minimum set of data fields that will be present in the level 2 data. A one-orbit SO₂ column level 2 file will be about 640 MB. More details about the operational level 2 product based on the NetCDF data format and the CF metadata convention are provided in the SO₂ Product User Model (Pedernana et al., 2016).

It should be noted that the averaging kernels are given only for the a priori profiles from the TM5 CTM (to save space). The averaging kernels for the box profiles can be estimated by scaling the provided averaging kernel (corresponding to TM5 profiles): $AK_{\text{box}}(p) = AK(p)$. Following the AK formulation of Eskes and Boersma (2004), the scaling factor is given simply by AMF ratios: $AMF_{\text{TM5}} / AMF_{\text{box}}$.

Auxiliary information

The algorithm relies on several external data sets. These can be either static or dynamic. An overview is given in Tables A2 and A3.

Table A1. List of output fields in the TROPOMI SO₂ products. nAlong × nAcross corresponds to the number of pixels in an orbit along track and across track, respectively; n.u.: no unit.

Name/data	Symbol	Unit	Description	Data type	Number of entries per observation
Date		n.u.	Date and time of the measurement YYM-MDDHHMMSS.MS	characters	nAlong
Latitudes	lat	degree	Latitudes of the four pixel corners + center	float	5 × nAlong × nAcross
Longitudes	long	degree	Longitudes of the four pixel corners + center	float	5 × nAlong × nAcross
SZA	θ_0	degree	Solar zenith angle	float	nAlong × nAcross
VZA	θ	degree	Viewing zenith angle	float	nAlong × nAcross
RAA	φ	degree	Relative azimuth angle	float	nAlong × nAcross
SCD	N_s	mol m ⁻²	SO ₂ slant column density	float	nAlong × nAcross
SCDcorr	N_s^c	mol m ⁻²	SO ₂ slant column density background corrected	float	nAlong × nAcross
VCD	N_v	mol m ⁻²	SO ₂ vertical column density (4 values)	float	4 × nAlong × nAcross
Wdow flag	Wflag	n.u.	Flag for the fitting window used (1, 2, 3)	integer	nAlong × nAcross
AMF	M	n.u.	Air mass factor (4 values)	float	4 × nAlong × nAcross
Cloud-free AMF	M_{clear}	n.u.	Cloud-free air mass factor (4 values)	float	4 × nAlong × nAcross
Cloudy AMF	M_{cloud}	n.u.	Fully cloudy air mass factor (4 values)	float	4 × nAlong × nAcross
CF	f_c	n.u.	Cloud fraction	float	nAlong × nAcross
CRF	Φ	n.u.	Cloud radiance fraction	float	nAlong × nAcross
CP	p_{cloud}	Pa	Cloud top pressure	float	nAlong × nAcross
CH	z_{cloud}	m	Cloud top height	float	nAlong × nAcross
CA	A_{cloud}	n.u.	Cloud top albedo	float	nAlong × nAcross
Albedo	A_s	n.u.	Surface albedo	float	nAlong × nAcross
Aerosol index	AAI	n.u.	Absorbing aerosol index	float	nAlong × nAcross
Chi-squared	χ^2	n.u.	Chi-squared of the fit	float	nAlong × nAcross
VCD error	σN_v	mol m ⁻²	Total error on the vertical column (individual measurement)	float	4 × nAlong × nAcross
SCD random error	σN_{s_rand}	mol m ⁻²	Random error on the slant column	float	nAlong × nAcross
SCD systematic error	σN_{s_syst}	mol m ⁻²	Systematic error on the slant column	float	nAlong × nAcross
AMF random error	σM_{rand}	n.u.	Random error on the air mass factor (4 values)	float	4 × nAlong × nAcross
AMF systematic error	σM_{syst}	n.u.	Systematic error on the air mass factor (4 values)	float	4 × nAlong × nAcross
Averaging kernel	AK	n.u.	Total column averaging kernel (for a priori profile from CTM)	float	34 × nAlong × nAcross
Averaging kernel scalings for box profiles	scaling box	n.u.	Factors to apply to the averaging kernel function to obtain the corresponding averaging kernels for the three box profiles	float	3 × nAlong × nAcross
SO ₂ profile	n_a	n.u.	A priori profile from CTM (volume mixing ratio)	float	34 × nAlong × nAcross
Surface altitude	z_s	m	Digital elevation map	float	nAlong × nAcross
Surface pressure	p_s	Pa	Effective surface pressure of the satellite pixel	float	nAlong × nAcross
TM5 level coefficient a	A_i	Pa	TM5 pressure level coefficients that effectively define the mid-layer levels (from ECMWF)	float	24
TM5 level coefficient b	A_i	n.u.		float	24

Table A2. Static auxiliary data for the S-5P SO₂ algorithm.

Name/data	Symbol	Unit	Source	Pre-process needs	Comments
Absorption cross sections					
SO ₂	σ_{SO_2}	cm ² molec ⁻¹	Bogumil et al. (2003), 203, 223, 243, 293 K Hermans et al. (2009), all temperatures	Convolution at the instrumental spectral resolution using the provided slit function	
Ozone	σ_{o3218} σ_{o3243}	cm ² molec ⁻¹	Brion et al. (1998); 218 and 243 K		
BrO	σ_{BrO}	cm ² molec ⁻¹	Fleischmann et al. (2004), 223 K		
NO ₂	σ_{NO_2}	cm ² molec ⁻¹	Vandaele et al. (1998), 220 K		–
O ₄ (O ₂ -O ₂)	σ_{O_4}	cm ⁵ molec ⁻²	Greenblatt et al. (1990)		
High-resolution reference solar spectrum	E_s	W m ⁻² nm ⁻¹	Chance and Kurucz (2010)	–	–
Ring effect	$\sigma_{\text{ringev}1}$ $\sigma_{\text{ringev}2}$	cm ² molec ⁻¹	Two Ring cross sections generated internally	A high-resolution reference solar spectrum and the instrument slit function are needed to generate the data set	Calculated in an ozone containing atmosphere for low and high SZA, using LIDORT_RRS (Spurr et al., 2008) and a standard atmosphere (CAMELOT European Pollution atmospheric profile)
Nonlinear absorption effect	O ₃ σ_{o3l} σ_{o3sq}	nm cm ² molec ⁻¹ cm ⁴ molec ⁻²	Two pseudo-cross sections generated internally	The O ₃ cross section at 218 K is needed	Calculated from the Taylor expansion of the wavelength and the O ₃ optical depth (Pukite et al., 2010)
Instrument slit function	SF	n.u.	Slit function by wavelength/detector	–	Values between 300 and 400 nm
Surface albedo	A_s	n.u.	OMI-based monthly minimum LER (update of Kleipool et al., 2008)	–	
Digital elevation map	z_s	m	GMTED2010 (Danielson et al., 2011)		Average over the ground pixel area.

Table A2. Continued.

Name/data	Symbol	Unit	Source	Pre-process needs	Comments
SO ₂ profile	n_a	n.u.	One-kilometer-thick box profiles, with three different peak altitudes, representing different altitude regimes: boundary layer: from the surface altitude to 1 km above it; free troposphere: centered around 7 km altitude; lower stratosphere: centered around 15 km altitude Daily SO ₂ profiles forecast from TM5	–	TM5 profiles from the last available day if the TM5 profiles of the current day are not available
Look-up table of pressure-resolved AMFs	m	n.u.	Calculated internally with the LIDORTv3.3 RTM (Spurr, 2008)	–	For the different fitting windows (312–326, 325–335, 360–390 nm), the assumed vertical column is 5, 100, 500 DU, respectively
Temperature correction parameters	α	K ⁻¹	Bogumil et al. (2003)	–	–

Table A3. Dynamic auxiliary data for the S-5P SO₂ algorithm.

Name/data	Symbol	Unit	Source	Pre-process needs	Backup if not available
S-5P level 1B Earth radiance	I	$\text{mol s}^{-1} \text{m}^{-2} \text{nm}^{-1} \text{sr}^{-1}$	S-5P L1b product	–	No retrieval
S-5P level 1B sun irradiance	E_0	$\text{mol s}^{-1} \text{m}^{-2} \text{nm}^{-1}$	S-5P L1b product	Wavelength recalibrated using a high-resolution reference solar spectrum	Use previous measurement
S-5P cloud fraction	f_c	n.u.	S-5P operational cloud product	–	
S-5P cloud top pressure	p_{cloud}	Pa	based on a Lambertian cloud model (Loyola et al., 2016)		No retrieval
S-5P cloud top albedo	A_{cloud}	n.u.	UPAS processor		
SO ₂ profile	n_a	n.u.	Daily forecast from TM5 CTM run at KNMI.	–	Use TM5 CTM profile from last available day
Temperature profile	T	K	Daily forecast from TM5 CTM run at KNMI	–	Use TM5 CTM profile from last available day
S-5P absorbing aerosol index	AAI	n.u.	S-5P operational AAI product (Zweers, 2016) Used for flagging KNMI processor	–	Missing information flag
Snow/ice flag		n.u.	Near-real-time global Ice and Snow Extent (NISE) data from NASA	–	Use snow/ice climatology

Table A4. Acronyms and abbreviations.

AAI	Absorbing aerosol index
AK	Averaging kernel
AMF	Air mass factor
AOD	Aerosol optical depth
AR	Alternative retrieval
BrO	Bromine monoxide
CAL	Cloud as layer
CAMELOT	Composition of the Atmospheric Mission concepts and Sentinel Observation Techniques
CAPACITY	Composition of the Atmosphere: Progress to Applications in the user Community
CCD	Charge-coupled device
CRB	Clouds as Reflecting Boundaries
CTM	Chemical transport model
DOAS	Differential optical absorption spectroscopy
DU	Dobson unit (1 DU = 2.6867×10^{16} molecules cm ⁻²)
ECMWF	European Centre for Medium-Range Weather Forecast
ESA	European Space Agency
FT	Free troposphere
FWHM	Full width at half maximum
GMES	Global Monitoring for Environment and Security
GOME-2	Global Ozone Monitoring Experiment-2
HCHO	Formaldehyde
IPA	Independent pixel approximation
IR	Infrared
L2WG	Level-2 Working Group
LER	Lambertian equivalent reflector
LIDORT	Linearized Discrete Ordinate Radiative Transfer
LOS	Line-of-sight angle
LS	Lower stratosphere
LUT	Look-up table
MAX-DOAS	Multi-axis DOAS
MR	Medium retrieval
NO ₂	Nitrogen dioxide
NOVAC	Network for Observation of Volcanic and Atmospheric Change
NRT	Near-real time
OCRA	Optical Cloud Recognition Algorithm
O ₃	Ozone
OMI	Ozone Monitoring Instrument
OMPS	Ozone Mapping Profiler Suite
PA	Prototype algorithm
(P)BL	Planetary boundary layer
PCA	Principal component analysis
ROCINN	Retrieval Of Cloud Information using Neural Networks
RRS	Rotational Raman scattering
RTM	Radiative transfer model
RAA	Relative azimuth angle
S-5P	Sentinel-5 Precursor
SCIAMACHY	SCanning Imaging Absorption spectroMeter for Atmospheric ChartographY
SCD	Slant column density
SCDE	Slant column density error
S/N	Signal-to-noise ratio
SO ₂	Sulfur dioxide
SR	Standard retrieval
SWIR	Shortwave infrared
SZA	Solar zenith angle
TOMS	Total Ozone Mapping Spectrometer
TROPOMI	Tropospheric Monitoring Instrument
UPAS	Universal Processor for UV/VIS Atmospheric Spectrometers
UV	Ultraviolet
VA	Verification algorithm
VC(D)	Vertical column density
WF	Weighting function

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