

Global retrieval of TROPOMI tropospheric HCHO and NO₂ columns with improved consistency based on updated Peking University OMI NO₂ algorithm

Yuhang Zhang^{1,2}, Huan Yu², Isabelle De Smedt², Jintai Lin¹, Nicolas Theys², Michel Van Roozendael², Gaia Pinardi², Steven Compernelle², Ruijing Ni³, Fangxuan Ren¹, Sijie Wang¹, Lulu Chen¹, Jos Van Geffen⁴, Mengyao Liu⁴, Alexander M. Cede^{5,6}, Martin Tiefengraber⁶, Alexis Merlaud², Martina M. Friedrich², Andreas Richter⁷, Ankie Pitters⁴, Vinod Kumar^{3,8}, Vinayak Sinha⁸, Thomas Wagner³, Yongjoo Choi⁹, Hisahiro Takashima¹⁰, Yugo Kanaya¹¹, Hitoshi Irie¹², Robert Spurr¹³, Wenfu Sun², Lorenzo Fabris²

¹Laboratory for Climate and Ocean-Atmosphere Studies, Department of Atmospheric and Oceanic Sciences, School of Physics, Peking University, Beijing 100871, China

²Royal Belgian Institute for Space Aeronomy, BIRA-IASB, Ringlaan 3, 1180 Uccle, Belgium

³Max Planck Institute for Chemistry, Mainz 55128, Germany

⁴Royal Netherlands Meteorological Institute (KNMI), Utrechtseweg 297, 3730 AE De Bilt, the Netherlands

⁵NASA Goddard Space Flight Center, Greenbelt, MD 20771, USA

⁶Luftblick, Innsbruck, Austria

⁷Institute of Environmental Physics (IUP), University of Bremen, Otto-Hahn-Allee 1, Bremen 28359, Germany

⁸Department of Earth and Environmental Sciences, Indian Institute of Science Education and Research, Mohali, Punjab, India

⁹Hankuk University of Foreign Studies, Yongin, Republic of Korea

¹⁰Fukuoka University, Fukuoka, Japan

¹¹Research Institute for Global Change, Japan Agency for Marine-Earth Science and Technology (JAMSTEC), Yokohama, Japan

¹²Center for Environmental Remote Sensing, Chiba University (Chiba U), Chiba, Japan

¹³RT Solutions Inc., Cambridge, MA 02138, USA

Correspondence to: Jintai Lin (linjt@pku.edu.cn), Michel Van Roozendael (michel.vanroozendael@aeronomie.be)

Abstract. The TROPOspheric Monitoring Instrument (TROPOMI), onboard the Sentinel-5 Precursor (S5P) satellite launched in October 2017, is dedicated to monitoring the atmospheric composition associated with air quality and climate change. This paper presents the global retrieval of TROPOMI tropospheric formaldehyde (HCHO) and nitrogen dioxide (NO₂) vertical columns using an updated version of the Peking University OMI NO₂ (POMINO) algorithm, which focuses on improving the calculation of air mass factors (AMFs). The algorithm features explicit corrections for the surface reflectance anisotropy and aerosol optical effects, and uses daily high-resolution (0.25°×0.25°) a priori HCHO and NO₂ profiles from the Global Earth Observing System Composition Forecast (GEOS-CF) dataset. For cloud correction, a consistent approach is used for both HCHO and NO₂ retrievals, where (1) the cloud fraction is re-calculated at 440 nm using the same ancillary parameters as those used in the NO₂ AMF calculation, and (2) the cloud top pressure is taken from the operational FRESCO-S cloud product.

The comparison between POMINO and reprocessed (RPRO) operational products in April, July, October 2021 and January 2022 exhibits high spatial agreement, but RPRO tropospheric HCHO and NO₂ columns are lower by 10% to 20% over polluted regions. Sensitivity tests with POMINO show that the HCHO retrieval differences are mainly caused by different aerosol correction methods (implicit versus explicit), prior information of vertical profile shapes and background corrections; while the NO₂ retrieval discrepancies result from different aerosol corrections, surface reflectances and a priori vertical profile shapes as well as their non-linear interactions. With

44 explicit aerosol corrections, the HCHO structural uncertainty due to the cloud correction using different cloud
45 parameters is within $\pm 20\%$, mainly caused by cloud height differences. Validation against ground-based
46 measurements from global Multi-Axis Differential Optical Absorption Spectroscopy (MAX-DOAS) observations
47 and the Pandonia Global Network (PGN) shows that in April, July, October 2021 and January 2022, POMINO
48 retrievals present a comparable day-to-day correlation but a reduced bias compared to the RPRO products (HCHO:
49 $R = 0.62$, NMB = -30.8% versus $R = 0.68$, NMB = -35.0% ; NO_2 : $R = 0.84$, NMB = -9.5% versus $R = 0.85$,
50 NMB = -19.4%). An improved agreement of HCHO/ NO_2 ratio (FNR) with MAX-DOAS and PGN measurements
51 based on POMINO retrievals is also found (NMB: -14.8% versus -21.1%). Our POMINO retrieval provides a
52 useful source of information particularly for studies combining HCHO and NO_2 .

53 **1 Introduction**

54 Formaldehyde (HCHO) and nitrogen dioxide (NO_2) are important trace gases in the troposphere. They play a
55 critical role in the processes of tropospheric ozone (O_3) and aerosol formation, and have significant influences on
56 air quality, climate and human health (Beelen et al., 2014; Crutzen, 1970; Shindell et al., 2009). Methods to
57 retrieve tropospheric HCHO and NO_2 vertical column densities (VCDs), respectively in the ultraviolet (UV) and
58 visible (VIS) spectral ranges, have rapidly developed in the last decades, based on sensors mounted on both sun-
59 synchronous and geostationary satellites such as the Global Ozone Monitoring Experiment (GOME; Burrows et
60 al., 1999), SCanning Imaging Absorption spectroMeter for Atmospheric CHartographY (SCIAMACHY;
61 Bovensmann et al., 1999), Ozone Monitoring Instrument (OMI; Levelt et al., 2006), Global Ozone Monitoring
62 Experiment-2 (GOME-2; Callies et al., 2000), Ozone Mapping and Profiling Suite Nadier Mapper (OMPS-NM;
63 Dittman et al., 2002), TROPOspheric Monitoring Instrument (TROPOMI; Veefkind et al., 2012), Environmental
64 Trace Gases Monitoring Instrument (EMI; Zhang et al., 2020), Geostationary Environment Monitoring
65 Spectrometer (GEMS; Kim et al., 2020) and Tropospheric Emissions: Monitoring of Pollution (TEMPO;
66 Zoogman et al., 2017). Such satellite observations have been extensively used in studies related to long-term trend
67 and variabilities (De Smedt et al., 2010; Jiang et al., 2022; Richter et al., 2005), estimation of surface-level
68 concentrations (Cooper et al., 2022; Wei et al., 2022), constraining emissions of non-methane volatile organic
69 compounds (NMVOCs) and nitrogen oxides ($\text{NO}_x \equiv \text{NO} + \text{NO}_2$) (Kong et al., 2022; Lin, 2012; Stavrou et al.,
70 2018), non-linear ozone chemistry (Jin et al., 2017, 2023; Jin and Holloway, 2015) and impacts on the environment
71 and human health (Chen et al., 2022; Li et al., 2023).

72 The retrieval algorithms of tropospheric HCHO and NO_2 VCDs based on observations from spaceborne
73 instruments share many retrieval concepts. First, the slant column density (SCD) representing the trace gas
74 concentration integrated along the average light path is obtained by performing a spectral fit from backscattered
75 radiance and irradiance spectra. Second the SCD is converted to a VCD using air mass factors (AMFs) obtained
76 from radiative transfer (RT) calculations, which are a function of the observation geometry, cloud information,
77 aerosol properties, surface conditions and the shape of a priori vertical profiles. The main intrinsic differences
78 between HCHO and NO_2 retrievals are that (1) different wavelength ranges are used for each retrieval, and (2) the
79 final tropospheric HCHO VCDs are determined with additional background correction based on modelled HCHO
80 columns in the reference region in the Field of Regard (FOR) of satellite instruments, while for NO_2 a stratosphere-
81 troposphere separation is performed in order to obtain tropospheric columns.

82 Many studies have focused on improving or developing retrieval algorithms to generate scientific HCHO or NO₂
83 products for comparison with operational products and for applications. For example, Liu et al. (2021) present an
84 improved tropospheric NO₂ retrieval algorithm from TROPOMI measurements over Europe, which employs a
85 new stratosphere-troposphere separation and updated auxiliary parameters, including a more realistic cloud
86 treatment, for AMF calculation. Over East Asia, Liu et al. (2020) release a new TROPOMI product for
87 tropospheric NO₂ columns that features explicit aerosol corrections in the AMF calculation, and Su et al. (2020)
88 improve the TROPOMI tropospheric HCHO retrieval by optimizing the spectral fit and using a priori profiles
89 from a higher resolution regional chemistry transport model.

90 However, little attention has been paid to fixing the systematic differences in ancillary parameters between HCHO
91 and NO₂ AMF calculations. For instance, the TROPOMI reprocessed (RPRO) HCHO version 2.4.1 and NO₂
92 version 2.4.0 operational products make use of cloud information from different sources: the Optical Cloud
93 Recognition Algorithm/Retrieval of Cloud information using Neural Networks (OCRA/ROCINN) - Cloud as
94 Reflecting Boundaries (CRB) product is used for HCHO, while the Fast Retrieval Scheme for Clouds from
95 Oxygen absorptions bands - Sentinels (FRESCO-S) product is used for NO₂. Besides, the surface albedo used in
96 the current HCHO retrieval is the OMI-based monthly minimum Lambertian-equivalent reflectivity (MLER) at
97 340 nm with a spatial resolution of $0.5^\circ \times 0.5^\circ$ (lat. \times long.), whereas the one used in the NO₂ retrieval has been
98 updated with the KNMI TROPOMI directionally dependent Lambertian-equivalent reflectivity (DLER) v1.0
99 database at 440 nm with a spatial resolution of $0.125^\circ \times 0.125^\circ$. Finally, the radiative transfer model used for
100 HCHO AMF calculation is the linearized pseudo-spherical scalar and vector discrete ordinate radiative transfer
101 code (VLIDORT) version 2.6, whereas that used for NO₂ AMF calculation is the Double-Adding KNMI (DAK)
102 polarized radiative transfer code version 3.2. Such inconsistencies are an important limitation for studies
103 combining satellite HCHO and NO₂ products, such as analysis of ozone chemistry and wildfires (Jin et al., 2020,
104 2023). Therefore, there is a need for consistent retrievals of tropospheric HCHO and NO₂ VCDs. Moreover, the
105 TROPOMI operational HCHO and NO₂ products do not explicitly account for the optical effect of aerosols, and
106 use a priori profile shapes from the massively parallel version of the Tracer Model 5 (TM5-MP; Williams et al.,
107 2017) with a relatively coarse spatial resolution ($1^\circ \times 1^\circ$).

108 The Peking University OMI NO₂ (POMINO) algorithm offers a potential tool to address these limitations.
109 Founded by Lin et al. (2014), POMINO has been continuously developed and applied to the OMI, TROPOMI and
110 GEMS instruments (Lin et al., 2014, 2015; Liu et al., 2019, 2020; Zhang et al., 2023). POMINO features an
111 explicit treatment of aerosol optical effects and surface reflectance anisotropy, as well as a re-calculation of cloud
112 information using ancillary parameters consistent with those used for NO₂ AMF calculation. A smaller bias of
113 POMINO NO₂ data than the operational products has been reported from validation against independent ground-
114 based measurements (Liu et al., 2019, 2020; Zhang et al., 2023). However, the previous POMINO-TROPOMI
115 algorithm was limited to Asia, and its potential for HCHO retrieval remained unexplored.

116 In this paper, we present the global retrieval of TROPOMI tropospheric HCHO and NO₂ VCDs with much
117 improved consistency, based on an updated version of the POMINO algorithm. After describing the methods and
118 data in Section 2, we present the quantitative comparison of tropospheric HCHO and NO₂ columns between
119 POMINO and RPRO products (Sect. 3). We then discuss the structural uncertainty of HCHO and NO₂ retrieval
120 based on the POMINO algorithm, by conducting a series of sensitivity tests on cloud correction, aerosol correction,
121 surface reflectance and a priori profile shapes (Sect. 4). Tentative estimates of POMINO retrieval uncertainty are

122 given in Sect. 5. Finally, we use independent ground-based measurements from a global network of Multi-Axis
 123 Differential Optical Absorption Spectroscopy (MAX-DOAS) instruments and the Pandonia Global Network
 124 (PGN) to validate the tropospheric HCHO and NO₂ columns from the POMINO and RPRO products (Sect. 6).

125 2 Method and data

126 2.1 TROPOMI instrument and operational algorithms for HCHO and NO₂ retrieval

127 TROPOMI is an imaging spectrometer onboard the European Space Agency (ESA) Copernicus Sentinel-5
 128 Precursor (S5P) satellite launched on 13th October 2017, crossing the equator at around 13:30 local time (LT)
 129 (Veefkind et al., 2012). Its wide spectral range includes the ultraviolet (UV), visible (VIS), near-infrared (NIR)
 130 and shortwave infrared (SWIR), allowing monitoring of atmospheric trace gases, aerosols, clouds and surface
 131 properties. The original spatial resolution of about 7 km × 3.5 km (along-track × across-track) at nadir was refined
 132 to about 5.5 km × 3.5 km on the 6th of August 2019 by means of a reduction of the along-track integration time.
 133 The wide swath of about 2600 km in the across-track direction enables global coverage on a daily basis, except
 134 for narrow strips between orbits of about 0.5° wide at the equator.

135 The TROPOMI operational HCHO and NO₂ retrieval algorithms have been fully described in De Smedt (2022)
 136 and Van Geffen et al. (2022b), respectively. The first common step is to derive slant columns by performing a
 137 spectral fit using the Differential Optical Absorption Spectroscopy (DOAS) method. Specifics for the SCD
 138 retrieval are provided in Table S1. After the DOAS spectral fitting, a two-step normalization of the HCHO slant
 139 columns is performed to remove any remaining global offset and possible stripes. Then the corrected differential
 140 SCDs (dSCDs) are converted to vertical columns using AMFs at 340 nm. The AMFs are derived from a pre-
 141 calculated look-up table (LUT) storing altitude-dependent AMFs calculated with the VLIDORT v2.6 radiative
 142 transfer model. This approach implements implicit aerosol corrections by assuming that aerosols can be simply
 143 treated as “effective clouds”, and uses the OMI-based monthly MLER dataset for surface reflectance. The HCHO
 144 vertical profile shape is specified from TM5-MP daily analyses. For pixels with partly cloudy scenes, a cloud
 145 correction is applied based on the independent pixel approximation (IPA) (Martin et al., 2002), using cloud
 146 fraction (CF), cloud top pressure (CP) and cloud albedo information from the OCRA/ROCINN-CRB product:

$$147 \quad M = w \cdot M_{\text{cld}} + (1 - w) \cdot M_{\text{clr}} \quad (1)$$

148 In Eq. (1), w is the cloud radiance fraction (CRF), M_{cld} the cloudy-sky AMF and M_{clr} the clear-sky AMF. In the
 149 OCRA/ROCINN-CRB cloud retrieval, OCRA first computes the cloud fraction using a broad-band UV/VIS color-
 150 space approach with two colors: Green (405–495 nm) and Blue (350–395 nm); then ROCINN-CRB calculates the
 151 cloud height and cloud albedo using in and around the oxygen (O₂) A-band (~760 nm). In the final step, TM5-MP
 152 HCHO vertical columns in the reference region are added as the compensation for the background HCHO from
 153 methane (CH₄) oxidation in the equatorial Pacific. The final tropospheric HCHO VCD, N_V , can be written as
 154 follows:

$$155 \quad N_V = \frac{N_S - N_{S,0}}{M} + \frac{M_{\text{clear},0}}{M} N_{V,0}^{\text{TM5-MP}} \quad (2)$$

156 with $(N_S - N_{S,0})$ being the corrected HCHO differential slant column, M the HCHO AMF, $M_{\text{clear},0}$ the HCHO clear-
 157 sky AMF in the reference region ([90°S, 90°N], [180°W, 120°W]), and $N_{V,0}^{\text{TM5-MP}}$ the HCHO vertical column from

158 a daily latitude-dependent polynomial, which is fitted through 5° latitude bin means of TM5-MP HCHO vertical
159 columns in the reference region (De Smedt, 2022).

160 For NO₂, a de-stripping is also applied to the fitted slant columns even though the systematic across-track features
161 are very small (Van Geffen et al., 2020). The second step is the stratosphere-troposphere separation, where TM5-
162 MP is used to assimilate TROPOMI total NO₂ SCDs, determine the stratospheric NO₂ SCDs and, by subtraction,
163 infer the tropospheric NO₂ SCDs. To calculate tropospheric NO₂ AMFs, the operational algorithm applies implicit
164 aerosol corrections, uses NO₂ a priori profile shapes from TM5-MP daily analyses, and adopts a DLER at 440 nm
165 from the KNMI TROPOMI DLER v1.0 surface reflectance database. For the cloud correction, it takes the cloud
166 top pressure from the FRESCO-S product (using the O₂ A-band at ~760 nm) and retrieves an effective cloud
167 fraction (ECF) by fitting the observed continuum reflectance to a simulated reflectance at 440 nm, assuming an
168 optically thick Lambertian cloud with a fixed cloud albedo of 0.8. The tropospheric NO₂ VCD, N_V^{trop} , can be
169 written as follows:

$$170 \quad N_V^{\text{trop}} = \frac{N_S^{\text{total}} - N_S^{\text{strat}}}{M} \quad (3)$$

171 with $(N_S^{\text{total}} - N_S^{\text{strat}})$ the tropospheric NO₂ slant column and M the tropospheric NO₂ AMF.

172 **2.2 Improved POMINO-TROPOMI algorithm for global HCHO and NO₂ AMF calculations**

173 Focusing on the improvement of global HCHO and NO₂ AMF calculations as well as their consistency, we use an
174 updated POMINO-TROPOMI parallelized AMFv6 package (Figure S1) driven by the LLinearized Discrete
175 Ordinate Radiative Transfer code (LIDORT) version 3.6 inherited from previous POMINO products (Liu et al.,
176 2020). The DOAS spectral fit, HCHO dSCD background correction and NO₂ stratosphere-troposphere separation
177 are not included in this study, so corrected HCHO dSCDs and tropospheric NO₂ SCDs are directly taken from the
178 RPRO HCHO v2.4.1 product and RPRO NO₂ v2.4.0 product, respectively. Compared to the previous HCHO
179 v2.3.0 processor, HCHO v2.4.1 processor uses new improved Level 1b v2.1.0 data products as input, and has been
180 applied for a full mission reprocessing starting from 7th May 2018. For NO₂, the improvements of the v2.4.0
181 processor include the use of a DLER climatology derived from TROPOMI observations and new improved Level
182 1b v2.1.0 data products as input, which has also been used for a full mission reprocessing from 1st May 2018.
183 Detailed information of S5P TROPOMI L2 HCHO and NO₂ processing baseline, including the processor version,
184 in-operation period and relevant improvements can be found at [https://sentiwiki.copernicus.eu/web/s5p-](https://sentiwiki.copernicus.eu/web/s5p-processing)
185 [processing](https://sentiwiki.copernicus.eu/web/s5p-processing).

186 Table1 lists the main improvements in the POMINO AMF algorithm compared to the RPRO algorithms. POMINO
187 calculates the AMFs with online pixel-by-pixel RT simulations rather than using the LUT. Explicit aerosol
188 corrections are implemented at the corresponding wavelengths of HCHO and NO₂, respectively, based on the
189 aerosol information from Global Earth Observing System Composition Forecast (GEOS-CF; Keller et al., 2021)
190 v1.0 and Moderate Resolution Imaging Spectroradiometer (MODIS) satellite data. We convert GEOS-CF vertical
191 volume mixing ratio profiles to optical depth profiles for each aerosol type, i.e., dust, sulfate-nitrate-ammonium
192 (SNA), organic carbon (OC), black carbon (BC) and sea salt, by using high-spectral-resolution aerosol optical
193 parameters from the GEOS-Chem website
194 (https://ftp.as.harvard.edu/gcgrid/data/aerosol_optics/hi_spectral_res/v9-02/, last access: 23 July 2024). We then
195 convert component-specific aerosol information to vertical profiles of aerosol extinction coefficient, single

196 scattering albedo and phase function. We further use monthly aerosol optical depth (AOD) data from
197 MODIS/Aqua Collection 6.1 MYD04_L2 dataset, with spatial and temporal interpolation for missing values, to
198 constrain the model AOD (Lin et al., 2014). Daily a priori HCHO and NO₂ profile shapes at TROPOMI overpass
199 time are also obtained from GEOS-CF v1.0 at the spatial resolution of 0.25° × 0.25°. Detailed comparison of the
200 specifications between GEOS-CF and TM5-MP is provided in Table S2.

201 In NO₂ AMF calculations, to account for the surface reflectance anisotropy over lands and coastal ocean regions,
202 we use bidirectional reflectance distribution function (BRDF) coefficients around 470 nm (band 3; bandwidth:
203 459 – 479 nm) from the MODIS MCD43C2.061 dataset. The reason for the choice of MODIS BRDF over KNMI
204 TROPOMI DLER is that the operational MODIS BRDF algorithm fully characterizes the dependence of surface
205 reflectance on the solar zenith angle (SZA), viewing zenith angle (VZA) and relative azimuth angle (RAA) by a
206 linear combination of an isotropic parameter plus the volumetric and geometric scattering kernels (Roujean et al.,
207 1992; Zhou et al., 2010), while the DLER model only considers the satellite viewing angle (Tilstra et al., 2024).
208 For HCHO, given that the UV spectral band is not included in the MODIS instrument, we decided to use the
209 climatological DLER at 340 nm from the KNMI TROPOMI DLER v2.0 database.

210 To allow a consistent cloud correction, we use the same cloud information for both HCHO and NO₂ AMF
211 calculation. For each pixel, we acquire the cloud parameters by (1) taking the cloud top pressure from the
212 FRESCO-S cloud product, and (2) re-calculating the cloud fraction at 440 nm in a similar way as used in the
213 operational NO₂ algorithm. To simulate the TOA reflectance at 440 nm to derive cloud fraction, we use the
214 ancillary parameters fully consistent with those used in NO₂ AMF calculation, i.e., a surface reflectance derived
215 from MODIS BRDF coefficients and explicit aerosol information. Previous studies have demonstrated that in
216 most cases, explicit aerosol corrections lead to reduced cloud (radiance) fractions, especially over regions with
217 heavy aerosol loads such as the North China Plain in winter (Lin et al., 2015); while over regions where frequent
218 aerosol-cloud overlap occurs such as Southeast China in spring, the explicit corrections for absorbing aerosols
219 overlying the cloud deck lead to increased cloud fraction (Jethva et al., 2018). Such differences are because the
220 optical effects of aerosols are separated from those of clouds.

221 Based on the POMINO structure, we implemented a series of sensitivity tests to assess the importance of structural
222 uncertainties that arise when different ancillary parameters or methodologies are applied to the same data. For
223 HCHO, we first conducted the test “Fst_ORcp” (Case F1) by (1) re-calculating the cloud fraction at 340 nm based
224 on the reflectance derived using TROPOMI L1B radiance dataset version 2.1 in TROPOMI spectral band 3 (305-
225 400 nm), and irradiance dataset version 2.1 for the Ultra-violet, Visible and Near-Infrared (UVN) module post-
226 processed by BIRA-IASB, and (2) using the cloud top pressure from OCRA/ROCINN-CRB product. Therefore,
227 the differences between POMINO HCHO columns (Case F0) and those of the test “Fst_ORcp” represent the
228 structural uncertainty from the cloud correction using different cloud products. Based on the test “Fst_ORcp”, we
229 separately evaluate the effect of aerosol correction, surface reflectance and a priori profile shapes by conducting
230 the tests “Fst_imaer” (Case F2), “Fst_mler” (Case F3) and “Fst_tm5” (Case F4), respectively. Note that in all
231 sensitivity tests, only HCHO AMFs are changed accordingly, while we keep using GEOS-CF HCHO columns for
232 background correction.

233 Similarly, for NO₂ AMF calculations, based on POMINO NO₂ retrievals as the reference (Case N0), tests
234 “Nst_imaer” (Case N1), “Nst_dler” (Case N2) and “Nst_tm5” (Case N3) are used to quantify the individual effect
235 of aerosol correction, surface reflectance and a priori profile shapes. However, we noticed that the NO₂ differences

236 between POMINO and RPRO products can hardly be explained by the linear combination of the individual effect
 237 of each ancillary parameter as in the HCHO analysis. Therefore, we further conducted an additional test “Nst_joint”
 238 (Case N4) to “mimic” the AMF calculation in the RPRO algorithm, quantifying the joint effect of implicit aerosol
 239 corrections, KNMI TROPOMI DLER and TM5-MP a priori NO₂ profile shapes.

240 **Table 1.** Comparison of ancillary parameters between POMINO and RPRO operational products, and sensitivity tests on the
 241 corresponding ancillary parameters (“S.A.P.” means “Same as POMINO”).

Species	Product or sensitivity test case	RT model	Aerosol correction	Surface reflectance	Cloud correction	A priori profiles
HCHO	RPRO v2.4.1	VLIDORT v2.6 (LUT)	Implicit	OMI-based monthly MLER at 340 nm	CF and CP: OCRA/ROCINN-CRB	TM5-MP (1° × 1°)
	POMINO (Case F0)	LIDORT v3.6 (online)	Explicit	KNMI TROPOMI v2.0 DLER at 340 nm ⁽¹⁾	CF and CP: same as POMINO NO ₂	GEOS-CF (0.25° × 0.25°)
	Fst_ORcp (Case F1)	S.A.P.	S.A.P.	S.A.P.	CF: calculated at 340 nm CP: OCRA/ROCINN-CRB	S.A.P.
	Fst_imaer (Case F2)	S.A.P.	Implicit	S.A.P.	CF: re-calculated at 340 nm ⁽²⁾ CP: OCRA/ROCINN-CRB	S.A.P.
	Fst_mlcr (Case F3)	S.A.P.	S.A.P.	KNMI TROPOMI v2.0 MLER at 340 nm ⁽¹⁾	CF: re-calculated at 340 nm ⁽³⁾ CP: OCRA/ROCINN-CRB	S.A.P.
	Fst_tm5 (Case F4)	S.A.P.	S.A.P.	S.A.P.	CF: calculated at 340 nm CP: OCRA/ROCINN-CRB	TM5-MP (1° × 1°)
(1) KNMI TROPOMI v2.0 DLER at 340 nm over lands and coastal ocean regions, and MLER at 340 nm over open oceans.						
(2) Fst_imaer (Case F2) cloud fraction is re-calculated with implicit aerosol corrections and different from that of Case F1.						
(3) Fst_mlcr (Case F3) cloud fraction is re-calculated with KNMI TROPOMI v2.0 MLER and different from that of Case F1.						
NO ₂	RPRO v2.4.0	DAK v3.2 (LUT)	Implicit	KNMI TROPOMI v1.0 DLER at 440 nm	CF: calculated at 440 nm CP: FRESCO-S	TM5-MP (1° × 1°)
	POMINO (Case N0)	LIDORT v3.6 (online)	Explicit	MODIS MCD43C2.061 BRDF around 470 nm ⁽⁴⁾	CF: re-calculated at 440 nm CP: FRESCO-S	GEOS-CF (0.25° × 0.25°)
	Nst_imaer (Case N1)	S.A.P.	Implicit	S.A.P.	CF: re-calculated at 440 nm ⁽⁶⁾ CP: FRESCO-S	S.A.P.
	Nst_dler (Case N2)	S.A.P.	S.A.P.	KNMI TROPOMI v2.0 DLER at 440 nm ⁽⁵⁾	CF: re-calculated at 440 nm ⁽⁷⁾ CP: FRESCO-S	S.A.P.
	Nst_tm5 (Case N3)	S.A.P.	S.A.P.	S.A.P.	S.A.P.	TM5-MP (1° × 1°)
	Nst_joint (Case N4)	S.A.P.	Implicit	KNMI TROPOMI v2.0 DLER at 440 nm ⁽⁵⁾	CF: re-calculated at 440 nm ⁽⁸⁾ CP: FRESCO-S	TM5-MP (1° × 1°)

(4) MODIS MCD43C2.061 BRDF around 470 nm over lands and coastal ocean regions, and KNMI TROPOMI v2.0 MLER at 440 nm over open oceans.

(5) KNMI TROPOMI v2.0 DLER at 440 nm over lands and coastal ocean regions, and MLER at 440 nm over open oceans.

(6) Nst_imaer (Case N1) cloud fraction is re-calculated with implicit aerosol corrections and different from that of Case N0.

(7) Nst_dler (Case N2) cloud fraction is re-calculated with KNMI TROPOMI v2.0 DLER and different from that of Case N0.

(8) Nst_joint (Case N4) cloud fraction is re-calculated with implicit aerosol corrections and KNMI TROPOMI v2.0 DLER, and different from that of Case N0.

242

243 2.3 Ground-based MAX-DOAS datasets

244 Ground-based MAX-DOAS instruments can provide vertical columns and profiles of trace gases from the surface
 245 up to the lower free troposphere (around 4 km). The measurement sensitivity is the highest near the surface and
 246 decreases at higher altitudes. Information on ground-based MAX-DOAS measurements used in this study is
 247 summarized in Table 2 with locations specified in Figure S2. For each site, we use Fiducial Reference
 248 Measurements for Ground-based DOAS Air-Quality Observations (FRM₄DOAS; <https://frm4doas.aeronomie.be/>,
 249 Van Roozendaal et al., 2024) version 01.01 harmonized HCHO and NO₂ data if available, otherwise we use data
 250 generated by principal investigators of each instrument using non-harmonized retrieval settings. The aim of the
 251 FRM₄DOAS project is to minimize inhomogeneities in the current MAX-DOAS network to provide reference

252 datasets for satellite data validation. So far, many MAX-DOAS sites have been used for validation (De Smedt et
 253 al., 2021; Pinardi et al., 2020; Verhoelst et al., 2021; Yombo Phaka et al., 2023), but this is the starting point of
 254 the FRM₄DOAS project and much more sites will join the centralized processing facility.
 255 According to previous studies, the total estimated uncertainty of ground-based MAX-DOAS measurements in
 256 polluted conditions is about 30% for HCHO and NO₂ VCDs (De Smedt et al., 2021; Verhoelst et al., 2021). The
 257 mean bias is due mainly to systematic uncertainties related to AMF calculations. The uncertainty may also vary
 258 when different report strategies are used. Routine validation results show an overall bias of -37% for HCHO and
 259 -28% for NO₂ in the operational TROPOMI products compared to MAX-DOAS measurements in the validation
 260 report (available at <https://mpc-vdaf.tropomi.eu/>).

261 **Table 2.** MAX-DOAS datasets used for the validation. The sites are listed in alphabetical order based on the first letter of the
 262 site name.

Station, country (lat/long)	Species	Owner/group	Retrieval type	Reference
Athens, Greece (38.05°N, 23.86°E)	NO ₂	IUPB ⁽¹⁾	FRM ₄ DOAS 01.01	https://frm4doas.aeronomie.be/ Van Roozendaal et al. (2024)
Bremen, German (53.10°N, 8.85°E)	HCHO and NO ₂	IUPB	FRM ₄ DOAS 01.01	https://frm4doas.aeronomie.be/ Van Roozendaal et al. (2024)
Cabauw, the Netherlands (51.97°N, 4.93°E)	HCHO and NO ₂	KNMI ⁽²⁾	FRM ₄ DOAS 01.01	https://frm4doas.aeronomie.be/ Van Roozendaal et al. (2024)
Cape Hedo, Japan (26.87°N, 128.25°E)	NO ₂	JAMSTEC ⁽³⁾	Parameterized profiling (PP)	Kanaya et al. (2014)
Chiba, Japan (35.63°N, 140.10°E)	NO ₂	ChibaU ⁽⁴⁾	Parameterized profiling (PP)	Irie et al. (2011, 2012, 2015)
De Bilt, the Netherlands (52.10°N, 5.18°E)	HCHO and NO ₂	KNMI	FRM ₄ DOAS 01.01	https://frm4doas.aeronomie.be/ Van Roozendaal et al. (2024)
Fukue, Japan (32.75°N, 128.68°E)	NO ₂	JAMSTEC	Parameterized profiling (PP)	Kanaya et al. (2014)
Kinshasa, Democratic Republic of Congo (4.3°S, 15.30°E)	HCHO and NO ₂	BIRA-IASB ⁽⁵⁾	FRM ₄ DOAS 01.01	https://frm4doas.aeronomie.be/ Van Roozendaal et al. (2024)
Mohali, India (30.67°N, 76.74°E)	HCHO and NO ₂	IISER ⁽⁶⁾ /MPIC ⁽⁷⁾	QA4ECV harmonization procedure	De Smedt et al. (2021); Kumar et al. (2020)
Xianghe, China (39.75°N, 116.96°E)	HCHO and NO ₂	BIRA-IASB	FRM ₄ DOAS 01.01	https://frm4doas.aeronomie.be/ Van Roozendaal et al. (2024)
Yokosuka, Japan (35.32°N, 139.65°E)	NO ₂	JAMSTEC	Parameterized profiling (PP)	Kanaya et al. (2014)

(1) Institute of Environmental Physics, University of Bremen

(2) Royal Netherlands Meteorological Institute

(3) Japan Agency for Marine-Earth Science and Technology

(4) Chiba University

(5) Royal Belgian Institute for Space Aeronomy

(6) Indian Institute of Science Education and Research

(7) Max Planck Institute for Chemistry

263

264 2.4 PGN/Pandora datasets

265 The Pandonia Global Network (PGN) is a large-scale global network providing ground-based observations of
 266 multiple atmospheric reactive trace gases, including HCHO and NO₂, and associated uncertainty values for
 267 satellite validation and other scientific activities. It is based on ground-based passive spectrometer systems called
 268 “Pandora” that can perform sun, moon and sky observations. The datasets have been widely used to validate
 269 HCHO and NO₂ measurements from satellite instruments and field campaigns (Herman et al., 2019; Kai-
 270 Sikhakhane et al., 2024; Li et al., 2021; Liu et al., 2024a; Verhoelst et al., 2021).

271 Herman et al. (2009) reported that the nominal estimated uncertainty of total NO₂ columns is 0.27×10^{15}
272 molec.cm⁻² for the random part and 2.7×10^{15} molec.cm⁻² for the systematic part, and an uncertainty of 20% is
273 reported by comparisons with in-situ measurements (Verhoelst et al., 2021). However, the newer PGN NO₂
274 rnv3p1-8 data, which are employed in this study, have considerably lower uncertainties due to changes in (1) the
275 optical setup, (2) the gas-calibration approach and (3) a more accurate NO₂ effective temperature estimation. As
276 reported in the PGN data products Readme ([https://publications.pandonia-global-](https://publications.pandonia-global-network.org/manuals/PGN_DataProducts_Readme.pdf)
277 [network.org/manuals/PGN_DataProducts_Readme.pdf](https://publications.pandonia-global-network.org/manuals/PGN_DataProducts_Readme.pdf)), the combined uncertainty increases with decreasing
278 SZA, reaching $\sim 0.45 \times 10^{15}$ molec.cm⁻² for NO₂ rnv3p1-8 data and $\sim 1.2 \times 10^{15}$ molec.cm⁻² for HCHO rfus5p1-8
279 data at SZA=10° (median uncertainty over 137 data sets). The report uncertainty does not yet include the impact
280 of spectral fitting quality and is therefore a lower limit. This uncertainty component will be included in a future
281 PGN release; at Izana site, it is estimated to increase the reported uncertainty at SZA=10° to 1.0×10^{15} molec.cm⁻²
282 for NO₂ and 3.0×10^{15} molec.cm⁻² for HCHO.

283 In this work, we use HCHO rfus5p1-8 and NO₂ rnv3p1-8 direct sun total column measurements only from the
284 ESA Validation Data Centre (EVDC) (<https://evdc.esa.int>, last access: 17 July 2024), because the PGN sub-dataset
285 submitted to EVDC undergoes a more thorough quality check, in which the issues in PGN HCHO retrievals are
286 mostly mitigated. A total of 22 sites across the globe have valid measurements for HCHO and NO₂ validation in
287 the period of study (Figure S2).

288 **2.5 Data use and validation statistics**

289 For comparison between satellite HCHO data, we filter out the retrieved data based on the following criteria: we
290 exclude pixels with RPRO quality assurance values (QA) ≤ 0.5 , which includes SZA or VZA $> 70^\circ$ or activated
291 snow/ice flag. We also exclude pixels with POMINO-derived CRFs at 440 nm greater than 0.5, to minimize the
292 impact of cloud contamination. The same criteria are applied to the NO₂ comparison as well. To examine the
293 spatial distribution, gridded tropospheric HCHO and NO₂ VCDs in April, July, October 2021, and January 2022
294 at a resolution of $0.25^\circ \times 0.25^\circ$ are calculated using an area-weighted oversampling technique (Zhang et al., 2023).
295 For comparisons between satellite and ground-based HCHO data, we take two successive steps for data processing.
296 First, we calculate the daily average HCHO columns from ground-based MAX-DOAS and PGN measurements
297 within the time window between 11:00 and 16:00 LT. For PGN data, we only use those with the flag “assured
298 high quality” (data quality flag of 0) or “not-assured high quality” (data quality flag of 10)
299 ([https://www.pandonia-global-network.org/wp-content/uploads/2024/11/PGN_DataProducts_Readme_v1-8-](https://www.pandonia-global-network.org/wp-content/uploads/2024/11/PGN_DataProducts_Readme_v1-8-9.pdf)
300 [9.pdf](https://www.pandonia-global-network.org/wp-content/uploads/2024/11/PGN_DataProducts_Readme_v1-8-9.pdf)). Then we calculate daily average satellite HCHO columns based on pixels selected using the cloud
301 information from POMINO retrieval, with the pixel center located within a radius of 20 km to the instruments.
302 The daily collocated data pair is considered valid only if 10 satellite pixels or more are used for calculation. The
303 processing for NO₂ data is different from that of HCHO in three aspects: (1) the time window for NO₂ is between
304 13:00 to 14:00 LT, as the diurnal variation of NO₂ is much stronger than that of HCHO; (2) the radius between
305 the satellite pixel center and the instrument is 5 km, considering the much larger spatial gradient of the NO₂
306 distribution and less noise in the NO₂ retrieval; (3) we derive PGN tropospheric NO₂ columns each day by
307 subtracting stratospheric NO₂ columns from the RPRO NO₂ v2.4.0 L2 product over the instrument from the total
308 NO₂ columns, in order to make them comparable with satellite tropospheric NO₂ columns (Pinardi et al., 2020).
309 Based on collocated HCHO and NO₂ columns, we further compare the daily tropospheric column ratio of

310 formaldehyde to nitrogen dioxide (FNR) derived from satellite products and ground-based MAX-DOAS and PGN
311 measurements.

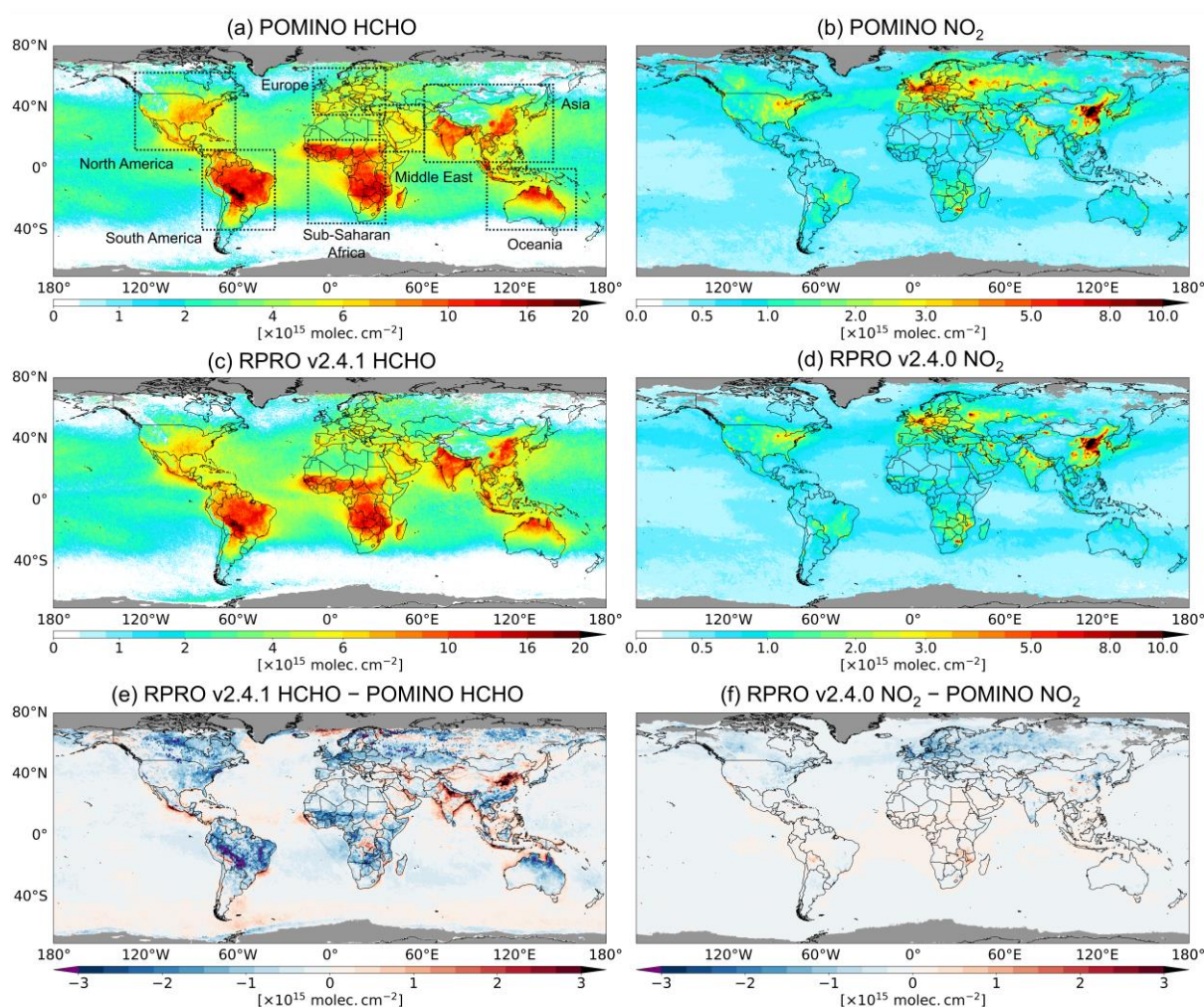
312 To quantify the performance of satellite products relative to ground-based measurements, we derive slope, offset
313 and correlation of the linear regression using the robust Theil-Sen estimator (Sen, 1968), which is insensitive to
314 occasional outliers. In a relative sense, we use normalized mean bias (NMB) to quantify the deviation between
315 satellite and ground-based measurements:

$$316 \quad \text{NMB} = \frac{\overline{\Omega^{\text{SAT}}} - \overline{\Omega^{\text{ground-based}}}}{\overline{\Omega^{\text{ground-based}}}} \times 100\% \quad (4)$$

317 with Ω being the HCHO or NO₂ vertical column in Sects. 6.1 and 6.2, and FNR in Sect. 6.3.

318 **3 Comparison of HCHO and NO₂ columns between POMINO and RPRO products**

319 Figures 1a and c illustrate the global distribution of tropospheric HCHO VCDs averaged over April, July, October
320 2021 and January 2022 from POMINO and RPRO retrieval, respectively. High levels of tropospheric HCHO
321 columns ($> 10 \times 10^{15}$ molec.cm⁻²) are evident over the Amazonia Rainforest, Sub-Saharan Africa, South and East
322 Asia as well as North Australia. Enhanced HCHO concentrations are also noticeable in the southeastern United
323 States of America (USA) and Mexico, while localized hotspots with lower magnitudes are evident in the Middle
324 East and Europe. Over the remote background regions, HCHO is primarily from CH₄ oxidation, and the abundance
325 is about 3×10^{15} molec.cm⁻² at maximum. Similarly, Figs. 1b and d show the POMINO and RPRO tropospheric
326 NO₂ VCDs in April, July, October 2021 and January 2022. High NO₂ columns are visible over three well-known
327 polluted regions, i.e., North China Plain, West Europe, and East USA, with strong hotspot signals over megacities
328 and metropolitan areas across the globe. Low NO₂ content in the remote atmosphere comes from aviation and
329 ship emissions, natural biogenic emissions, lightning and oxidation of long-lifetime species such as peroxyacetyl
330 nitrate (PAN).

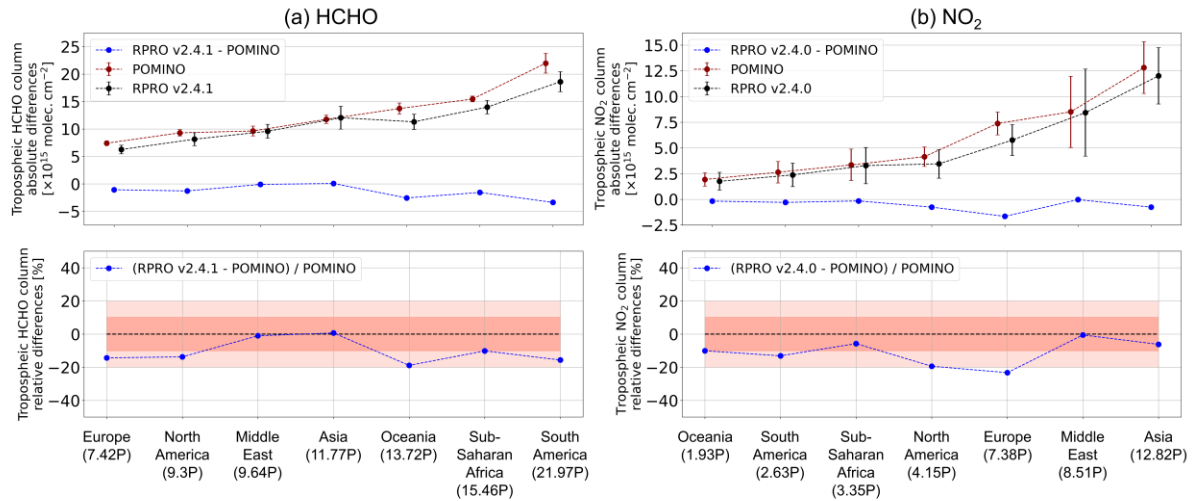


331

332 **Figure 1.** Spatial distribution of POMINO tropospheric HCHO and NO₂ VCDs (**a** and **b**), RPRO tropospheric HCHO and
 333 NO₂ VCDs (**c** and **d**), and respective absolute differences (**e** and **f**) at a spatial resolution of 0.25° × 0.25° averaged in April,
 334 July, October 2021, and January 2022. The black dashed rectangles illustrate the spatial range of the regions used for
 335 comparison. The regions in gray mean that there are no valid observations.

336 A high qualitative agreement is observed for both HCHO and NO₂ VCDs between RPRO and POMINO retrievals,
 337 as the same HCHO dSCDs and tropospheric NO₂ SCDs are used. However, as shown in Fig. 1e, RPRO HCHO
 338 tropospheric columns are lower by 2×10^{15} molec.cm⁻² or more over almost all regions with elevated HCHO
 339 columns except North India and North China Plain; RPRO NO₂ columns are also lower than those of POMINO
 340 over most East China, India, Europe, and North America by up to about 20% in a relative sense, despite the
 341 positive differences over Sub-Saharan Africa and some cities such as Xi'an, Teheran, and Los Angeles (Fig. 1f).
 342 We further make the comparison in seven specific regions (bounded by black rectangles in Fig. 1a): North America
 343 (125°W-60°W, 10°N-65°N), South America (85°W-35°W, 40°S-10°N), Europe (10°W-35°E, 35°N-60°N), Sub-
 344 Saharan Africa (15°W-35°E, 35°S-20°N), Middle East (30°E-60°E, 10°N-40°N), Asia (60°E-145°E, 5°N-55°N), and
 345 Oceania (100°E-160°E, 40°S-0°). Figure 2 shows the comparison results over the most polluted areas in each
 346 region, defined as where the POMINO tropospheric HCHO or NO₂ VCDs averaged over April, July, October
 347 2021 and January 2022 exceed their 99 percentiles; results for regional mean comparisons are shown in Figure
 348 S3. For HCHO, RPRO data are consistently lower than POMINO by around 15% over polluted areas in five
 349 regions, although the difference is small over the Middle East and Asia because of the cancellation between

350 positive and negative differences on the finer spatial scale. For NO_2 , RPRO is smaller than POMINO by -19.4%
 351 for North America and -23.3% for Europe. Detailed comparisons for each month are shown in Figure S4 and S5.
 352 Overall, POMINO and RPRO HCHO and NO_2 retrievals show excellent agreement in a qualitative sense, but the
 353 column values differ by 10% to 20% on average over polluted areas around the world. Such differences result
 354 from the different cloud correction, aerosol correction, surface reflectance and vertical profile shapes used in AMF
 355 calculations, which will be further discussed in Sect. 4.



356
 357 **Figure 2.** Absolute and relative differences between POMINO and RPRO (a) HCHO and (b) NO_2 tropospheric columns
 358 averaged in April, July, October 2021, and January 2022 over polluted areas (defined as where POMINO mean HCHO or NO_2
 359 columns exceed their 99 percentiles) in seven regions. Regions are sorted as a function of POMINO mean HCHO or NO_2
 360 columns, with values (in the unit of “P” as $\text{Pmolec.cm}^{-2} = 1 \times 10^{15} \text{ molec.cm}^{-2}$) shown in the brackets in the bottom axis. Mean
 361 POMINO (red) and RPRO (black) columns are also plotted with the absolute differences in the upper panel. Error bars
 362 represent the standard deviations of the columns. Pink areas indicate 10% and 20% relative differences.

363 4 Sensitivity tests on AMF input parameters

364 As listed in Table 1, we implement a series of sensitivity tests to quantify the structural uncertainty from either
 365 individual or joint effect of using different ancillary parameters in the HCHO and NO_2 AMF calculation. The time
 366 period selected for the sensitivity analysis is July 2021 and January 2022, representing the summer and winter
 367 time, respectively. Note that one of the most important features of the POMINO HCHO and NO_2 retrievals is that
 368 they use the same cloud parameters for consistent cloud correction. Therefore, besides discussing the effect of
 369 cloud correction based on POMINO cloud parameters, we also compare the differences between HCHO columns
 370 retrieved using different cloud parameters, especially the cloud top pressures. The influences of aerosol correction,
 371 surface reflectance, a priori profile shapes and their joint effect are discussed in the subsequent sub-sections.

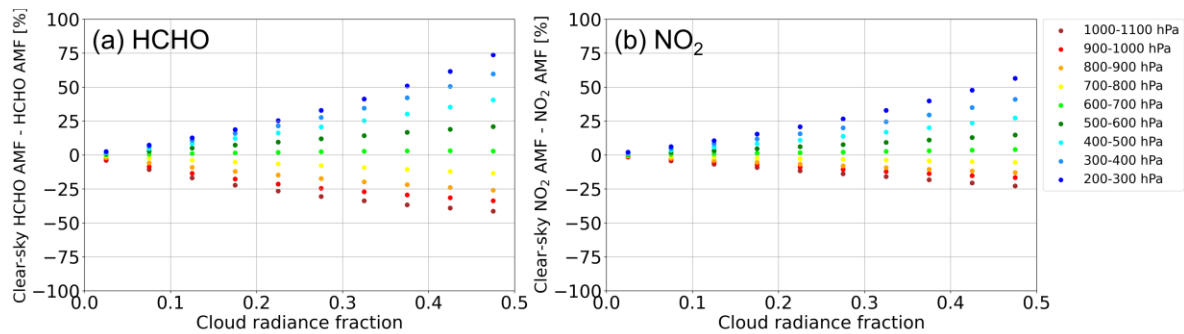
372 4.1 Cloud correction

373 4.1.1 Effect of cloud correction based on POMINO cloud parameters

374 When calculating tropospheric AMFs, it is important to account for the influence of clouds on the radiative transfer
 375 process in the atmosphere (Boersma et al., 2011; De Smedt et al., 2021; Lorente et al., 2017; Martin et al., 2002).
 376 Clouds can either enhance or reduce the sensitivity to the trace gas molecules depending on their height relative
 377 to the trace gas layers (the so-called “albedo” or “shielding” effect, respectively). Despite the relatively large
 378 uncertainty of retrieved cloud parameters in near-cloud-free scenario (defined here as $\text{CF} \leq 0.1$ or $\text{CRF} \leq 0.4$)

379 (Richter and Burrows, 2002), most HCHO and NO₂ AMF algorithms make use of the IPA method (Sect. 2.1) to
380 explicitly account for the cloud effect.

381 Figure 3 shows the differences between clear-sky AMF and total AMF of all pixels with HCHO or NO₂ QA > 0.5
382 in July 2021 and January 2022, based on the FRESCO-S cloud top pressures and POMINO re-calculated cloud
383 fractions at 440 nm with explicit aerosol corrections. For both HCHO and NO₂, the differences between clear-sky
384 AMF and total AMF are negative when cloud top pressures are higher than 700 hPa, and their magnitudes continue
385 to increase along with the cloud top pressures. The negative differences can be as large as -30% for HCHO and
386 -20% for NO₂ when the CRFs are in the interval of 0.45 to 0.5 and cloud top pressures are higher than 900 hPa.
387 This illustrates the “albedo” effect of low clouds by increasing the contribution of photons from near-surface
388 layers to the ensemble of photons received at the satellite instrument and thus leading to higher total AMF.



389

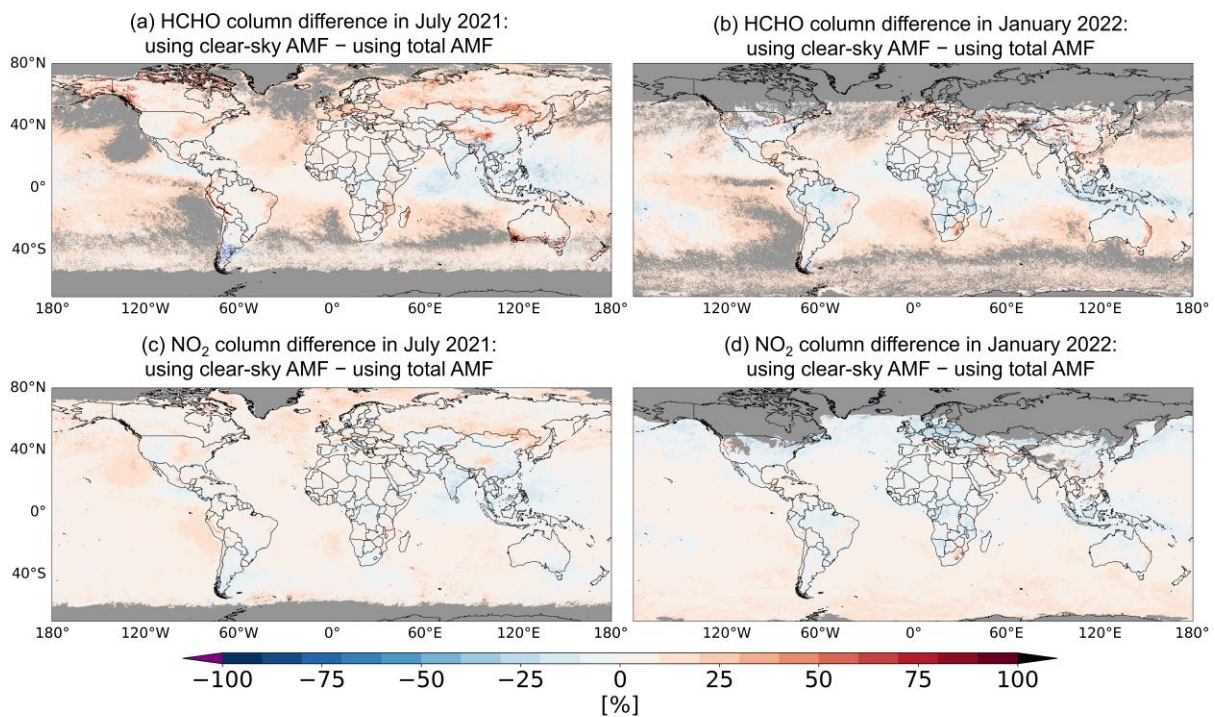
390 **Figure 3.** Differences of (a) HCHO and (b) NO₂ clear-sky AMF to total AMF for different cloud radiance fraction with an
391 interval of 0.05 in different cloud top pressure ranges (shown in different colors). All pixels with HCHO or NO₂ QA > 0.5 in
392 July 2021 and January 2022 are included.

393 On the contrary, clouds with cloud top pressure lower than 700 hPa reflect most photons back to the top of
394 atmosphere as a “shield” before they reach the HCHO or NO₂ abundant layers. As a result, positive differences of
395 clear-sky AMF to total AMF occur, and they increase as the cloud top pressures decrease, reaching 50% or more
396 when CRFs are in the interval of 0.4 to 0.5 and cloud top pressures are lower than 400 hPa. This result is also in
397 line with the previous study by Lorente et al. (2017).

398 In the global view (Figure 4), for both HCHO and NO₂ columns, the difference due to cloud correction (i.e., using
399 clear-sky AMF versus total AMF) is ±10% on average over high-value regions and can reach 40% over specific
400 areas. Note that all these comparisons are based on HCHO and NO₂ a priori profile shapes from GEOS-CF. The
401 signs and values of the differences might be different when using the profile shapes from another model, along
402 with the structural uncertainty discussed in Sect. 4.1.2.

403 One issue existing in the process of cloud correction in the POMINO retrieval is that only the cloud fraction is re-
404 calculated with explicit aerosol corrections, while the cloud top pressure is taken from the external dataset, i.e.,
405 the FRESCO-S cloud product, in which the aerosols are implicitly accounted for. As a result, this step introduces
406 presumably an aerosol overcorrection issue in the cloud top pressures of partly cloudy pixels, and therefore brings
407 in additional uncertainties in the AMF calculations. Lin et al. (2015) reported that excluding aerosols leads to an
408 increase of O₂-O₂-based cloud top pressures (from 700–900 hPa to 750–950 hPa) over eastern China, but it is
409 difficult to clarify the mechanism due to its complexity (Lin et al., 2014). Currently there is no direct way to
410 estimate the effect of aerosol correction on the FRESCO-S cloud height retrieval without doing O₂ A-band cloud
411 retrieval tests, which is beyond the scope of this study. However, below we give an estimation of the uncertainty
412 in POMINO HCHO and NO₂ vertical columns caused by this issue.

413 Given the fact that, in the retrieval algorithm, the cloud is assumed to be an optically thick Lambertian reflector
 414 with a high albedo of 0.8, the cloudy-sky AMF (and hence tropospheric AMF) is very sensitive to the accuracy of
 415 the cloud height when the cloud is low and vertically mixed with the aerosols and trace gases. In these cases, we
 416 can assume that the retrieved cloud height is primarily influenced by aerosols (Van Geffen et al., 2022a), therefore
 417 the aerosol overcorrection issue becomes non-negligible. Focusing on valid pixels for which the difference
 418 between the surface pressure and the FRESCO-S cloud top pressure is equal to 100 hPa or less ($\sim 17.5\%$ and $\sim 19.9\%$
 419 of total pixels in July 2021 and January 2022, respectively), the aerosol overcorrection uncertainty can be roughly
 420 estimated from the difference of HCHO and NO₂ vertical columns retrieved using either aerosol-corrected clear-
 421 sky AMFs (aerosol correction applied; cloud correction not applied) or aerosol-corrected total AMFs (both aerosol
 422 and cloud corrections applied). Based on the results shown in Figure S6, we tentatively estimate the uncertainty
 423 to be in the range from 10% to 15% for HCHO, and within 10% for NO₂. The estimated NO₂ uncertainty level is
 424 also supported by the sensitivity test results in Liu et al. (2020). They implemented a “semi-explicit” aerosol
 425 correction approach, in which aerosol optical effects are explicitly corrected for clear-sky AMFs, but are excluded
 426 for the cloudy-sky portion of partly cloudy pixels, and found the NO₂ differences due to the aerosol correction
 427 choice for cloudy-sky AMFs vary from 3.1% to 11.2% over eastern China in July 2018. The tentatively estimated
 428 uncertainty range above is comparable to or less than that from other ancillary parameters (Sect. 5), and only
 429 needs to be taken into account for partly cloudy pixels with low clouds.



430

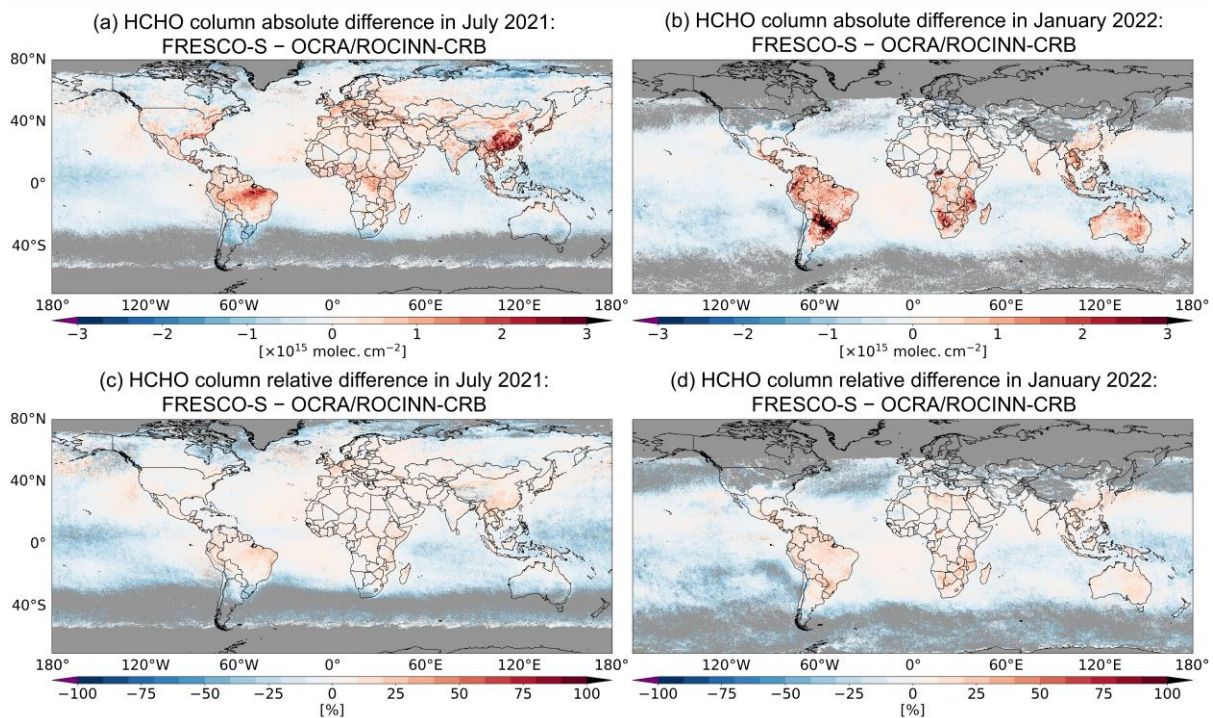
431 **Figure 4.** Relative differences of tropospheric HCHO (a and b) and NO₂ (c and d) columns derived using clear-sky POMINO
 432 AMF to those using total POMINO AMF in July 2021 and January 2022. The regions in gray mean that there are no valid
 433 observations.

434 4.1.2 Structural uncertainty of cloud correction based on different cloud parameters

435 The structural uncertainty of the cloud correction can be evaluated using cloud parameters from different cloud
 436 products. Lorente et al. (2017) have demonstrated that the systematic differences in cloud top pressure can lead
 437 to substantial differences in tropospheric NO₂ AMFs and VCDs. Focusing on HCHO in this section, we first

438 compare the effective cloud fractions and cloud top pressures either calculated in different ways or from different
 439 products. As shown in the left column of Figure S7, POMINO-based ECF calculated at 440 nm and 340 nm as
 440 well as OCRA/ROCINN-CRB ECF show similar global patterns in July 2021. Despite the differences over certain
 441 areas, great agreement is exhibited between OCRA/ROCINN-CRB ECF and POMINO-based ECF calculated at
 442 440 nm (linear regression slope of 0.92, offset of 0.02 and correlation coefficient of 0.80), and between POMINO-
 443 based ECF calculated at 340 nm and 440 nm (linear regression slope of 0.93, offset of 0.01 and correlation
 444 coefficient of 0.93). However, the OCRA/ROCINN-CRB cloud top pressures are significantly higher than those
 445 of the FRESCO-S product over the Amazonia Rainforest, Equatorial Africa and East China by 100-300 hPa, while
 446 the FRESCO-S cloud top pressures tend to be higher over many other places such as the Intertropical Convergence
 447 Zone (ITCZ) over the oceans (Fig. S6f). The comparison results over China are also qualitatively consistent with
 448 the findings by Latsch et al. (2022), in which the ROCINN CRB cloud heights differ significantly from those of
 449 FRESCO-S when considering low cloud fraction and lowest cloud height values that are critical for tropospheric
 450 trace gas retrievals. Such differences are systematic and are caused by different methodologies and ancillary
 451 parameters used in each cloud retrieval (Loyola et al., 2018; Van Geffen et al., 2022a), which are also reported in
 452 recent validation exercises using independent cloud measurements (Compernelle et al., 2021).

453 As shown in Fig. 5, by comparing the result of POMINO to the test “Fst_ORcp” (Case F1, using the
 454 OCRA/ROCINN-CRB cloud top pressures and the POMINO-based ECFs calculated at 340 nm), we find
 455 differences of HCHO columns by up to 20% on average over highly polluted regions, as well as a positive
 456 increment over South America. Over remote background regions such as the Pacific Ocean, however, negative
 457 differences are found of $0.5-1 \times 10^{15}$ molec.cm⁻². We attribute these differences to different OCRA/ROCINN-
 458 CRB and FRESCO-S cloud top pressures, as ECFs in POMINO and Case “Fst_ORcp” are very close. Note that
 459 this is a tentative estimate of HCHO column structural uncertainty from the choices of cloud parameters for cloud
 460 correction, because the results are dependent on the explicit aerosol corrections and HCHO priori profile shapes
 461 used in the tests.



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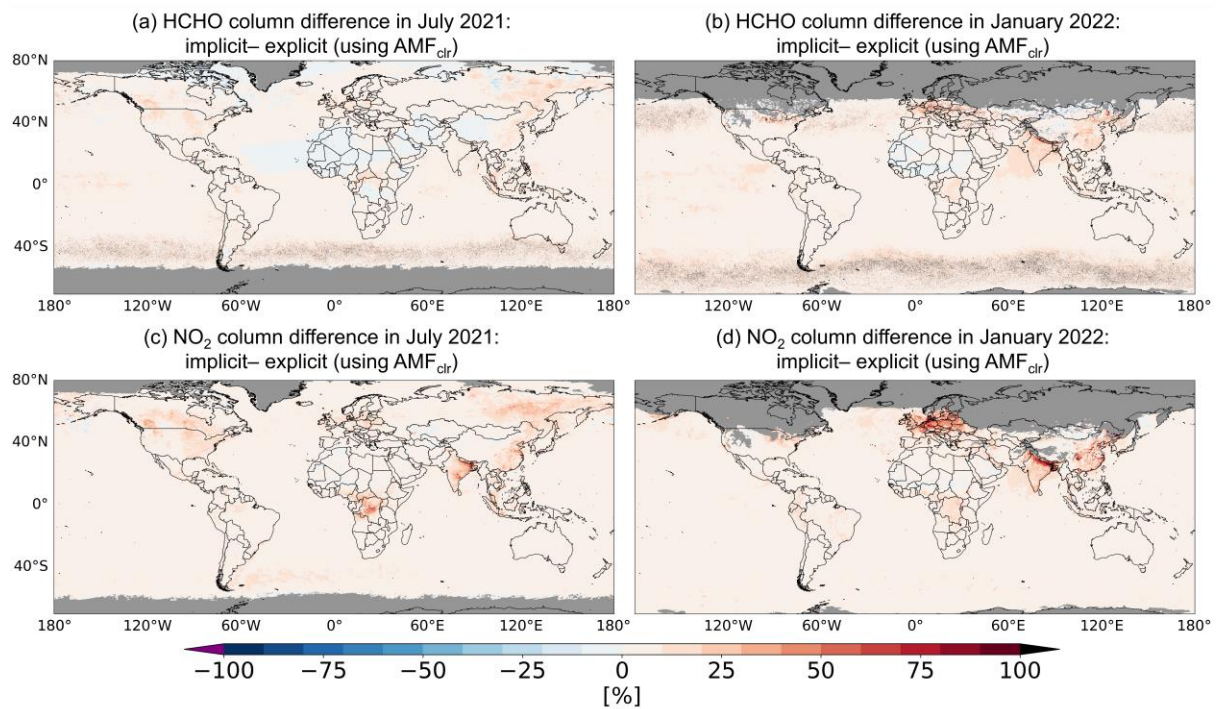
463 **Figure 5.** Absolute (first row) and relative differences (second row) of tropospheric HCHO columns of POMINO (using
464 FRESCO-S cloud top pressures) to those of the sensitivity test “Fst_ORcp” (using OCRA/ROCINN-CRB cloud top pressures)
465 in July 2021 and January 2022. Different cloud top pressures are emphasized in the title. The regions in gray mean that there
466 are no valid observations.

467 In summary, the implementation of the cloud correction in HCHO and NO₂ retrievals is necessary, and the
468 structural uncertainty due to different cloud parameters needs to be taken into consideration in product comparisons.
469 On the other hand, given the different spectral ranges used for trace gas retrievals (HCHO: 340 nm; NO₂: 440 nm)
470 and cloud retrievals (OCRA/ROCINN-CRB: O₂ A-band between 758 and 771 nm; FRESCO-S: O₂ A-band around
471 760 nm), cloud parameters should always be used with caution, especially for low-cloud-fraction conditions. For
472 example, in the ROCINN-CRB model, priori OCRA cloud fractions smaller than 0.05 are set to zero, and the
473 ROCINN retrieval is not activated under such “clear-sky” conditions. Instead of the NIR spectral range, the O₂-
474 O₂ cloud algorithm uses the O₂-O₂ absorption window around 477 nm, but it is more sensitive to low clouds and
475 aerosols. Therefore, further work is still needed to address such discrepancies.

476 **4.2 Aerosol correction**

477 The influence of aerosols on AMF calculations is very complicated because they depend on the type of aerosols
478 (scattering or absorbing) and their height relative to the trace gases. The AMFs are generally increased when non-
479 absorbing aerosols are vertically collocated with or lower than the trace gases, while an opposite effect arises
480 when the non-absorbing aerosols reside vertically higher than the trace gases; On the other hand, absorbing
481 aerosols (e.g., black carbon) always reduce the sensitivity of the satellite instruments to the trace gases (Leitão et
482 al., 2010; Lin et al., 2014, 2015; Liu et al., 2024b). Figure S8 shows a global map of AOD at 340 nm and 440 nm
483 used in POMINO retrievals. Areas with heavy aerosol loads in July 2021 include North America, Equatorial Africa,
484 Middle East, India and East China due to biomass burning and/or anthropogenic activities; while in January 2022,
485 the aerosol content is significant in Equatorial Africa, North India and North China Plain. Different aerosol
486 corrections can directly change the clear-sky AMF, affect the retrieval of cloud information (cloud fraction in
487 particular) and modulate the AMF in the cloudy portion of the pixel. The latter two effects influence the total AMF
488 in an indirect way, and the impact on cloud information is often more significant than on cloudy-sky AMF
489 (Vasilkov et al., 2021).

490 Figure 6 shows that when using clear-sky AMFs to derive vertical columns, implicit aerosol corrections lead to
491 higher HCHO columns by 10% to 20 % over North America in July 2021, and the differences exceed 20% over
492 North India and East China in January 2022. A similar pattern is shown in the NO₂ comparison. This is because
493 when aerosols that reside vertically lower than or are mixed with HCHO and NO₂ molecules are excluded (i.e., in
494 the case of implicit corrections), the calculated AMFs are lower than those with explicit aerosol corrections. On
495 the other hand, for scenarios with strong anthropogenic emissions or biomass burning, where most HCHO and
496 NO₂ molecules are near the surface while aerosols reside above these trace gases, implicit aerosol corrections
497 neglect the strong “shielding” effect of the scattering aerosols and the strong absorption of photons by the
498 absorbing aerosols (e.g., BC), which leads to higher AMFs and lower vertical columns. The negative differences
499 of HCHO columns over the Democratic Republic of Congo in July 2021 (Fig. 6a) can be explained by the second
500 case.



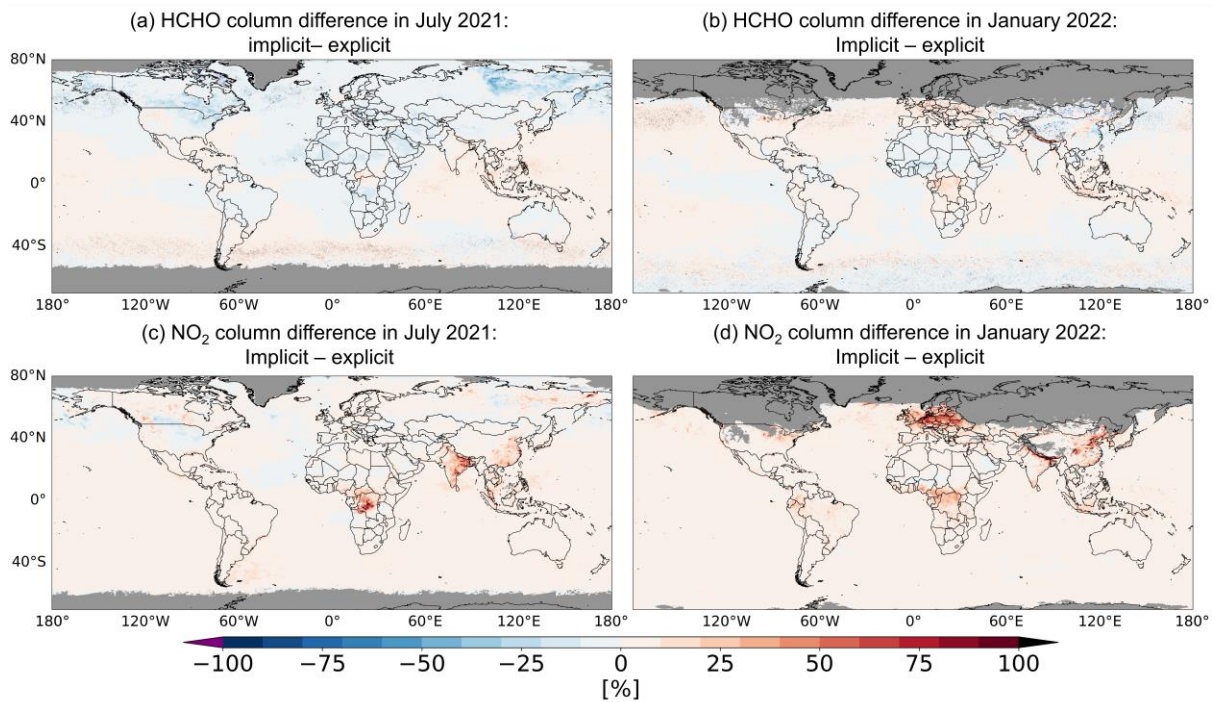
501

502 **Figure 6.** Relative differences of tropospheric HCHO (a and b) and NO₂ (c and d) columns retrieved using clear-sky AMF
 503 with implicit aerosol corrections to those with explicit aerosol corrections in July 2021 and January 2022. The regions in gray
 504 mean that there are no valid observations.

505 For cloudy-sky AMF, the impact of non-absorbing aerosols above a cloud is negligible since we assume the cloud
 506 to be an optically thick Lambertian reflectivity with a high albedo of 0.8 (Vasilkov et al., 2021). For absorbing
 507 aerosols above the clouds, they can reduce the backscattered radiance and hence affect the cloudy-sky AMF.
 508 However, Jethva et al. (2018) show that the occurrence of above-cloud absorbing aerosols is most frequent over
 509 coastal and oceanic regions because of the long-range transport of aerosols and low-level stratocumulus clouds.
 510 Over Southeast Asia during the springtime, the cloudy-sky frequency of occurrence of above-cloud absorbing
 511 aerosols is 20% to 40%, probably caused by biomass burning activities. Retrievals under these conditions are
 512 mostly discarded because the cloud fractions are too high to meet the filtering criteria for valid pixels (Sect. 2.5).
 513 Therefore, the overall influence of implicit aerosol corrections on the cloud-sky AMF can be neglected and the
 514 influence on the retrieval of cloud information, especially cloud fraction, is much more significant.

515 As explained in Sect. 2.2, explicit aerosol corrections affect the retrieved cloud (radiance) fraction due to the
 516 inclusion of aerosol radiative contribution. This is also confirmed in Figure S9 that compares retrieved cloud
 517 radiance fractions for the implicit versus explicit aerosol correction settings, in both UV and visible bands. As
 518 shown in Figure 7, when using cloud-corrected AMFs to consider both direct and indirect aerosol optical effects
 519 on the retrieval, the sign of HCHO relative differences over many regions is reversed from positive to negative
 520 compared to Figs. 6a and b, such as North and South America. This reflects the enhanced cloud “albedo” effect
 521 that increases the calculated HCHO scattering weights over the areas where cloud layers are vertically near or
 522 below the HCHO layers. As for NO₂, similar results due to enhanced cloud “albedo” effect are found over North
 523 America and East Russia in July 2021 (Fig. 7c), but the overall pattern in January 2022 remains the same as that
 524 in Fig. 6d. Over the polluted regions in Asia and Europe, implicit aerosol corrections increase the retrieved NO₂
 525 columns by 20% to 40% on average. This is because most NO₂ molecules over these polluted areas reside within
 526 1 km above the ground and below the FRESCO-S cloud layers during wintertime, so the increased cloud fractions

527 due to implicit aerosol corrections enhance the “shielding” effect on tropospheric NO₂ AMF calculation and hence
 528 higher NO₂ columns. The signs of the HCHO and NO₂ differences over North China Plain are not the same,
 529 probably because of the differences between HCHO and NO₂ vertical profile shapes.

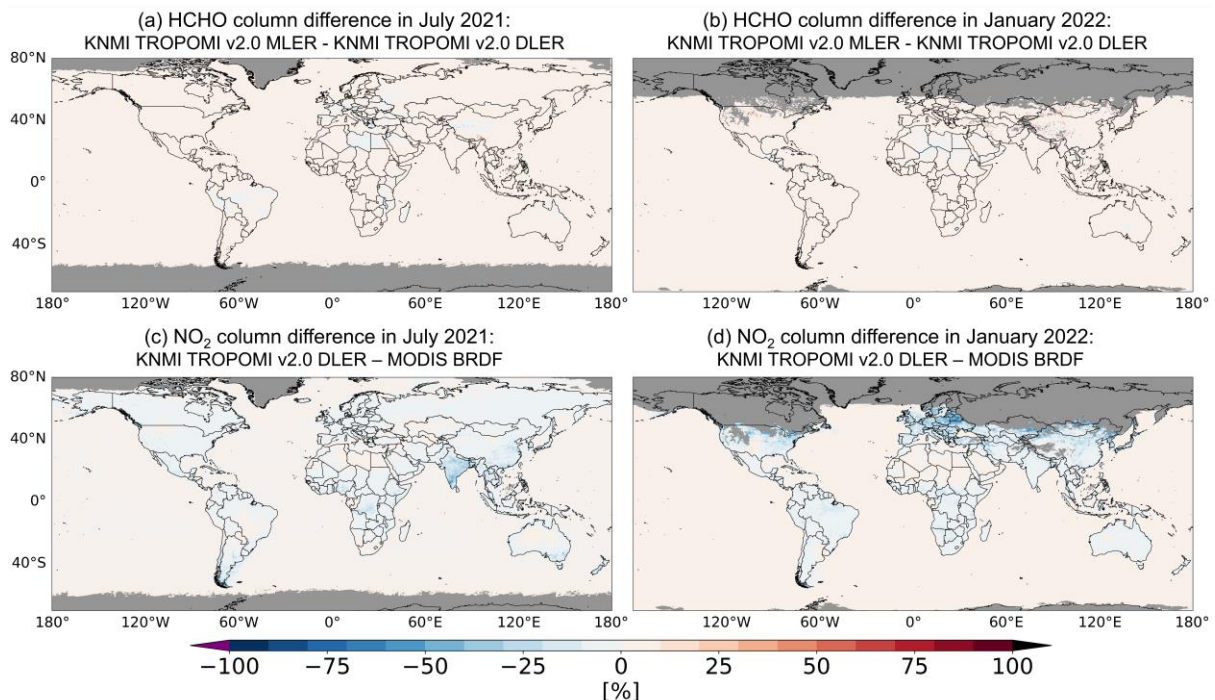


530
 531 **Figure 7.** Relative differences of tropospheric HCHO (a and b) and NO₂ (c and d) columns retrieved using cloud-corrected
 532 total AMF with implicit aerosol corrections (Cases “Fst_imaer” and “Nst_imaer”) to those with explicit aerosol corrections
 533 (Case “Fst_ORcp” and POMINO NO₂) in July 2021 and January 2022. The regions in gray mean that there are no valid
 534 observations.

535 4.3 Surface reflectance

536 Compared to the LER model, which simply assumes the surface to be a Lambertian reflector, DLER partly
 537 accounts for the anisotropy of the surface reflectance by building a certain relationship between the reflectance
 538 and the satellite VZA, but its dependence on the SZA and RAA is still not included. The BRDF model fully
 539 considers the surface optical property as a function of SZA, VZA, RAA and wavelength. At 340 nm, the
 540 directionality of the surface reflectance is small over most regions (Kleipool et al., 2008). Figure S10 compares
 541 the MODIS BRDF-derived blue-sky albedo (BSA, Schaepman-Strub et al., 2006) around 470 nm and KNMI
 542 TROPOMI DLER at 440 nm over lands and coastal ocean regions. In both months, DLER shows higher values
 543 than MODIS BSA except over desert and mountain regions, and the positive differences are larger than 0.1 over
 544 India in July 2021 and East Europe in January 2022. Reasons for these differences are not clear yet, but they are
 545 likely associated with different parameters and corrections for aerosols and snow/ice cover in the algorithm. The
 546 accuracy of the MODIS operational BRDF/albedo product (MCD43) is estimated by 5% to 10% of the field data
 547 at most validation sites studied so far (<https://modis-land.gsfc.nasa.gov/ValStatus.php?ProductID=MOD43>).
 548 Chong et al. (2024) also provide an estimation of random uncertainties in MODIS MCD43C1 surface reflectances
 549 for various surface types, which vary in the range of 0.01 to 0.03 for most cases.
 550 Figures 8a and b present the influence of surface reflectance on HCHO retrievals. As it is well known that the
 551 directionality of surface reflectance plays a marginal role in the retrieval based on the UV band, nearly no
 552 difference is shown between HCHO columns retrieved using KNMI TROPOMI DLER and MLER at 340 nm.

553 However, the systematic differences between different MLER products are a more important source of the
 554 structural uncertainty in HCHO AMFs. For example, KNMI TROPOMI MLER albedo at 340 nm is found to be
 555 consistently lower than OMI climatology monthly MLER albedo used in the RPRO product by 0.01–0.05
 556 (Kleipool et al., 2008; Tilstra et al., 2024).



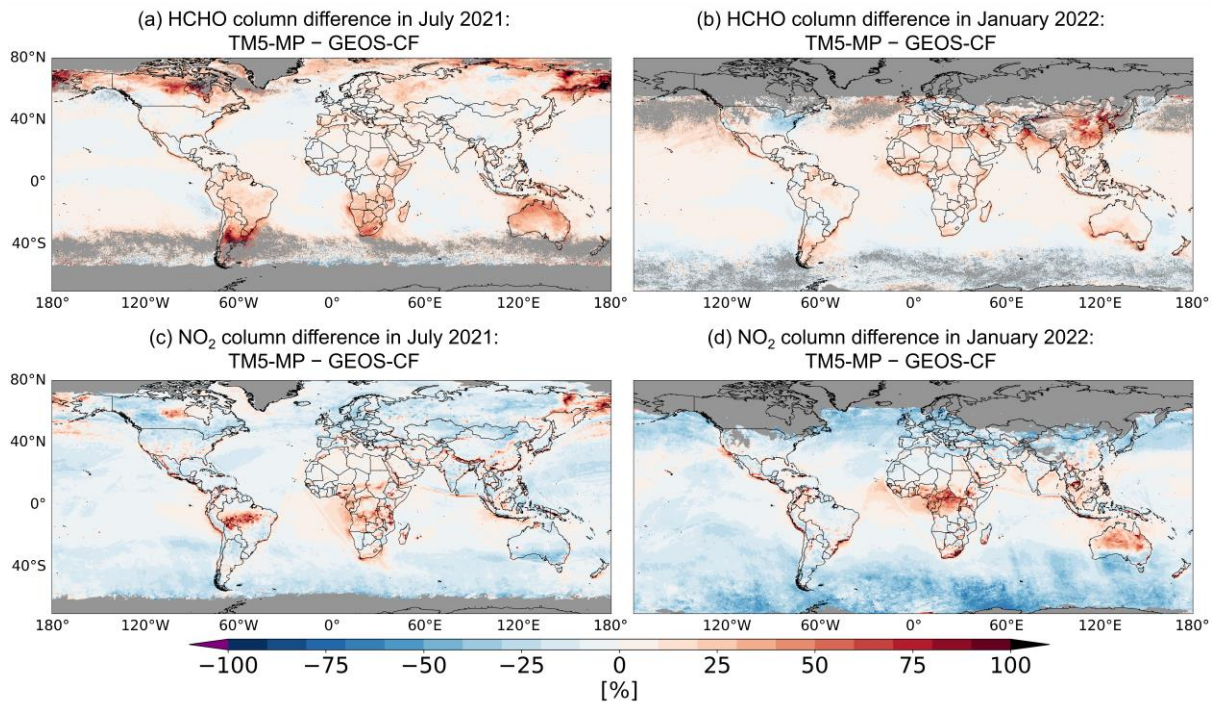
557
 558 **Figure 8.** Relative differences of tropospheric HCHO columns retrieved using KNMI TROPOMI v2.0 MLER at 340 nm (Case
 559 “Fst_mler”) to those using KNMI TROPOMI v2.0 DLER at 340 nm (Case “Fst_ORcp”) (a and b), and relative differences of
 560 tropospheric NO₂ columns retrieved using KNMI TROPOMI v2.0 DLER at 440 nm (Case “Nst_dler”) to those using MODIS
 561 BRDF at 440 nm (POMINO NO₂) (c and d) in July 2021 and January 2022. The regions in gray mean that there are no valid
 562 observations.

563 As for NO₂, Figs. 8c and d show significantly lower tropospheric NO₂ VCDs in the test “Nst_dler” (Case N2)
 564 than those in the reference POMINO retrieval (Case N0) over most land areas. In January 2022, the NO₂ columns
 565 retrieved using KNMI TROPOMI DLER are lower by 30% on average over the polluted regions with NO₂
 566 columns larger than 10×10^{15} molec.cm⁻² in Europe and North America. Like aerosols, the influence of surface
 567 reflectance on AMFs is also a combination of the direct effect on clear-sky AMF and the indirect effect through
 568 cloud correction (Boersma et al., 2011). As discussed by Tilstra (2024), DLER should not be considered as the
 569 optimal replacement for the BRDF in the VIS wavelength. If the directional surface reflection can be modelled in
 570 the RT calculation, it is better to use BRDF to derive surface reflectance for tropospheric NO₂ AMF calculation.

571 4.4 A priori profiles

572 In POMINO, we consistently use GEOS-CF HCHO and NO₂ vertical profile shapes as the prior information for
 573 AMF calculations. Compared with TM5-MP model of which the spatial resolution is $1^\circ \times 1^\circ$, GEOS-CF features
 574 a much finer spatial resolution ($0.25^\circ \times 0.25^\circ$). The horizontal distributions of GEOS-CF and TM5-MP
 575 tropospheric HCHO and NO₂ VCDs are shown in Figure S11, and comparisons of monthly mean HCHO and NO₂
 576 vertical profile shapes between the models and the ground-based MAX-DOAS measurements are shown in Figure
 577 S12. The collocation of model profiles and MAX-DOAS profiles follows the same methodology as described in
 578 Sect. 2.5. The differences between GEOS-CF, TM5-MP and MAX-DOAS profiles reflect the imperfections in

579 these data yet to be fully characterized (Keller et al., 2021; Williams et al., 2017), and they are also an important
 580 source of structural uncertainty in HCHO and NO₂ retrievals.
 581 Figure 9 shows the differences in retrieved HCHO and NO₂ VCDs caused by using different a priori vertical
 582 profile shapes. The HCHO and NO₂ columns retrieved with TM5-MP prior information are obtained using AMFs
 583 re-calculated by combining interpolated POMINO averaging kernels (AK) and TM5-MP a priori profile shapes.
 584 As shown in Figs. 9a and b, the spatial patterns of HCHO relative differences are variable over different places
 585 and in different months, and are generally more significant than the individual effects of clouds, aerosols and
 586 surface reflectance changes (Figs. 4, 7 and 8). At the regional level, the HCHO structural uncertainty from a priori
 587 profile shapes is 20% to 30% over the background clean areas, and 10% to 20% over the polluted areas. In contrast,
 588 the NO₂ differences caused by different a priori profile shapes are around 10% over the clean areas and reach 30%
 589 or more over the polluted areas. Over East China, India and the Middle East, localized differences over cities and
 590 polluted regions are obvious (Figs. 9c and d), reflecting the significant differences between TM5-MP and GEOS-
 591 CF NO₂ profile shapes. Besides, distinctive patterns along the coastal lines are visible, especially in the HCHO
 592 relative differences. This is caused by the relatively coarse horizontal resolution of TM5-MP, in which the large
 593 heterogeneity of HCHO vertical distribution is smoothed in the 1° × 1° grid.

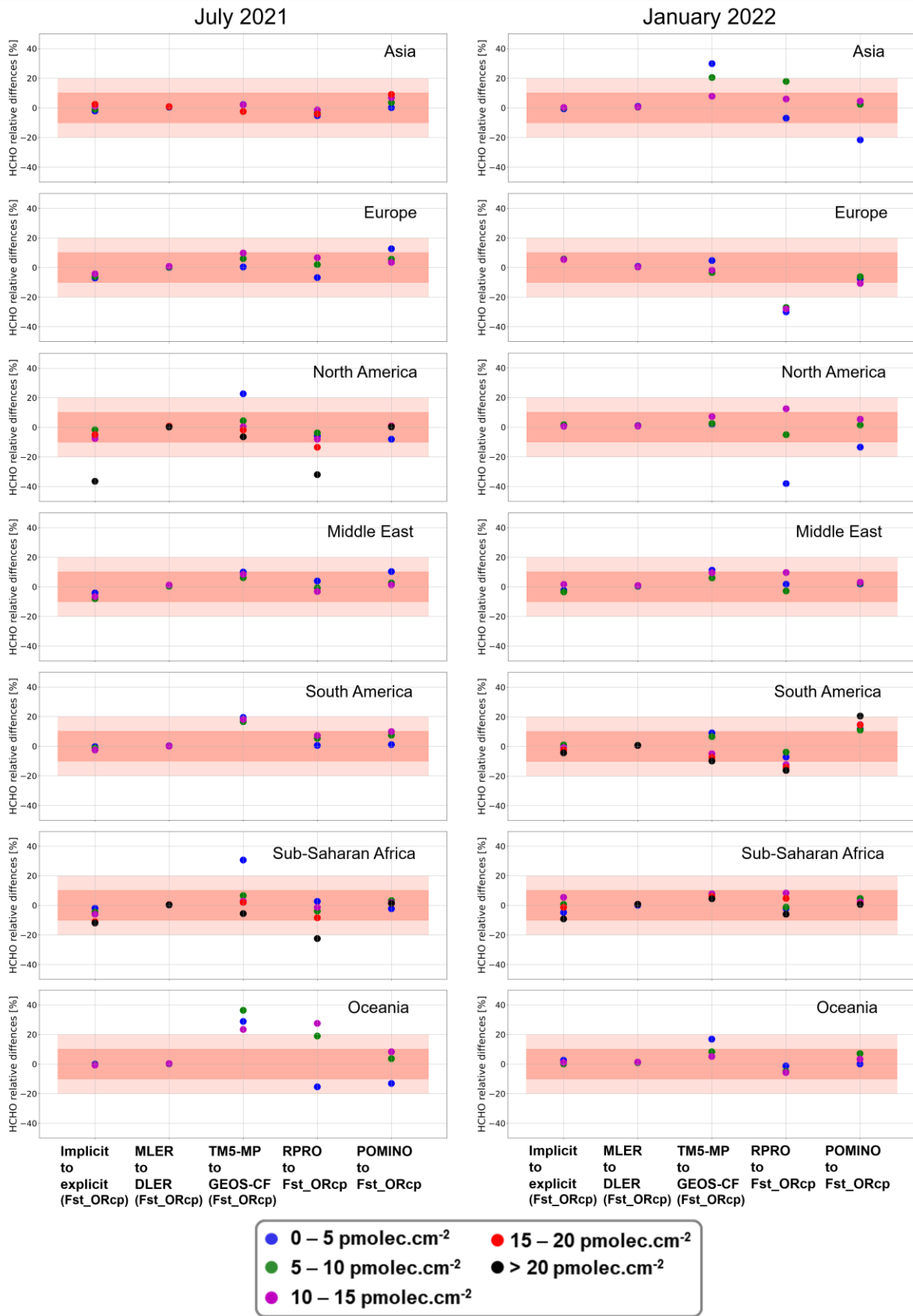


594
 595 **Figure 9.** Relative differences of tropospheric HCHO (a and b) and NO₂ (c and d) columns retrieved with TM5-MP priori
 596 profiles (Cases “Fst_tm5” and “Nst_tm5”) to those with GEOS-CF priori profiles (Case “Fst_ORcp” and POMINO NO₂) in
 597 July 2021 and January 2022. The regions in gray mean that there are no valid observations.

598 4.5 Summarizing the impacts of input parameters

599 As shown in each sub-figure of Figure 10, the first three columns summarize the structural uncertainty of aerosol
 600 correction, surface reflectance and a priori profile shapes on the HCHO retrieval in the corresponding region and
 601 month. As noted in Sect. 2.2, we consistently use GEOS-CF HCHO columns for background correction in every
 602 HCHO sensitivity test case. The TM5-MP HCHO columns over background regions are systematically lower than
 603 those of GEOS-CF by about 0.5×10^{15} molec.cm⁻² on average (Fig. S11), which strongly affects the comparisons
 604 over the low-HCHO regions.

605 Over clean areas ($\text{HCHO columns} < 5 \times 10^{15} \text{ molec.cm}^{-2}$), a priori profile shapes are the primary source of the
606 HCHO structural uncertainty (third column in Fig. 10). However, the differences between “Fst_tm5” and the
607 reference case “Fst_ORcp” are not in alignment with those of RPRO to the reference case, as manifested in the
608 consistent drop of the blue line from the third (“Fst_tm5” – reference) to the fourth column (RPRO – reference).
609 This drop can be attributed to the systematic issue in the background correction. Over most areas with HCHO
610 columns larger than $5 \times 10^{15} \text{ molec.cm}^{-2}$, relative to the same reference case, the HCHO differences caused by
611 using implicit aerosol corrections and TM5-MP priori profile shapes match well with those of RPRO product (the
612 fourth column). However, the lower values of RPRO than the reference case in Europe in January 2022 do not
613 agree with the combined results of tests “Fst_imaer” and “Fst_tm5”. This indicates that the higher OMI-based
614 climatology monthly MLER used in RPRO retrieval is probably the dominant factor. Furthermore, the influence
615 of cloud correction using different cloud parameters, especially the cloud top pressures, varies from -20% to 20%
616 depending on the specific regions and seasons. This is also an important factor for the HCHO differences between
617 POMINO and RPRO retrievals.



618

619

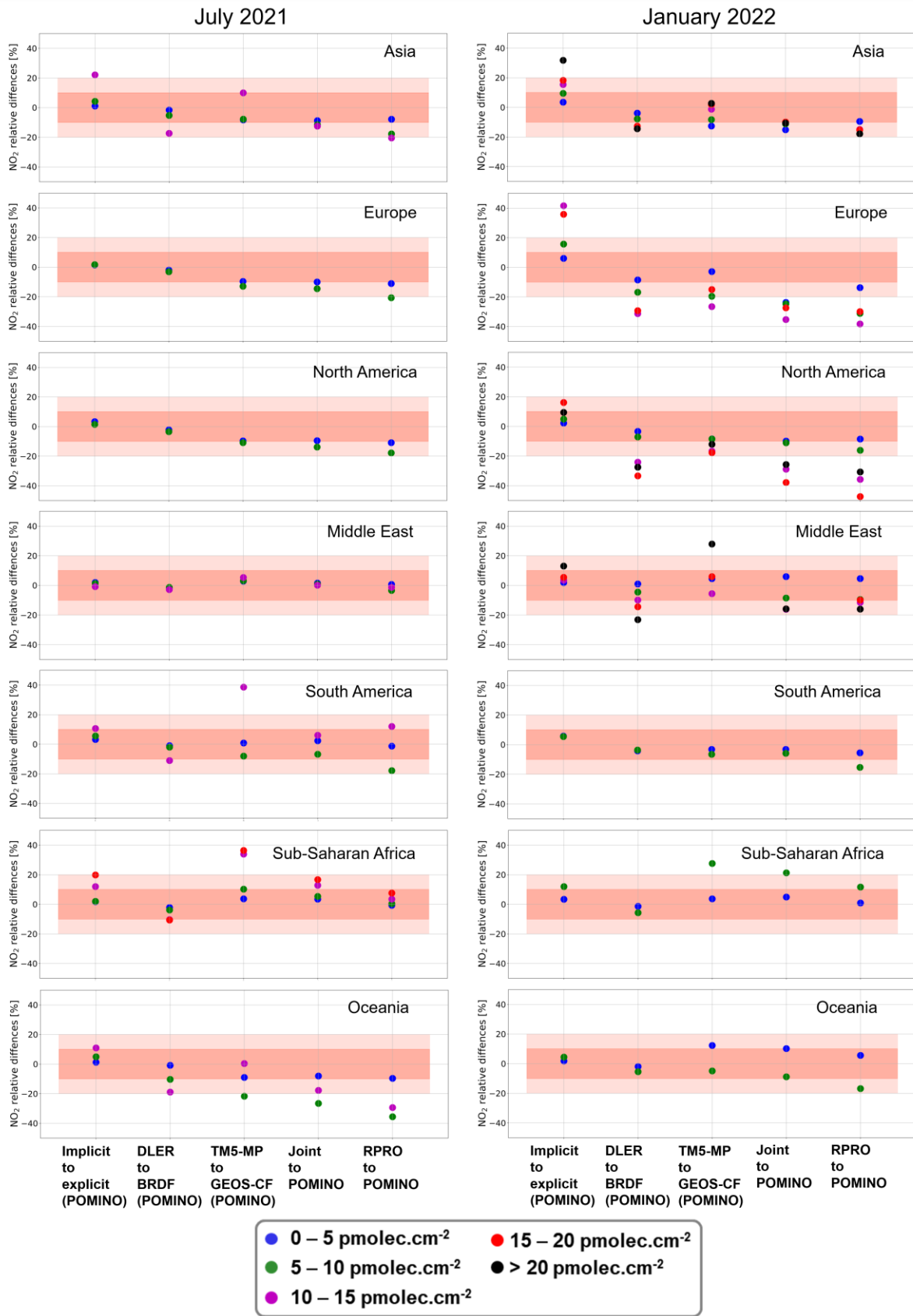
620

621

Figure 10. HCHO relative differences of the sensitivity test “Fst_imaer” (Case F2, first column), “Fst_mler” (Case F3, second column), “Fst_tm5” (Case F4, third column), RPRO product (fourth column) and POMINO product (fifth column) to the reference “Fst_ORcp” (Case F1) over seven regions in July 2021 and January 2022.

622

623 For NO₂, the first three columns in Figure 11 show the individual effect of each input parameter on the NO₂
624 retrieval in each region. Apparently, the relative differences between RPRO and POMINO (the fifth column) are
625 in discrepancy with the sum of the differences between each of the three cases (“Nst_imaer”, “Nst_dler” and
626 “Nst_tm5”) and the reference POMINO retrieval, especially over polluted areas in North America, Europe and
627 Asia in January 2022. However, the NO₂ columns of the test “Nst_joint” (Case N4) show high agreement with
628 those of the RPRO product when compared to the POMINO retrieval (fourth column in Fig. 11); a similar result
629 is shown for the spatial distribution in Figure S13. Therefore, the NO₂ differences between POMINO and RPRO
630 are the result of compensation effects between different aerosol corrections on one hand, and different surface
631 reflectances as well as vertical profile shapes on the other hand. These results demonstrate the non-linear joint
632 effects of aerosols, surface reflectance, clouds and a priori profiles in the AMF calculation, which are consistent
633 with the previous findings (Lin et al., 2015; Liu et al., 2020). The remaining differences between “Nst_joint” and
634 RPRO NO₂ columns are caused by their different ways to obtain tropospheric NO₂ AMFs, i.e., online pixel-
635 specific RT calculation versus LUT-based interpolation (Lin et al., 2014).



636

637 **Figure 11.** NO₂ relative differences of the sensitivity test “Nst_imaer” (Case N1, first column), “Nst_dler” (Case N2, second
 638 column), “Nst_tm5” (Case N3, third column), “Nst_joint” (Case N4, fourth column) and RPRO product (fifth column) to
 639 POMINO product as the reference (Case N0) over seven regions in July 2021 and January 2022.

640 5 Uncertainty estimates

641 The theoretical uncertainties of the POMINO retrievals can be analytically derived by uncertainty propagation
 642 based on the Eqs. 2 and 3 (Boersma et al., 2004). However, it is difficult to estimate the overall AMF uncertainty
 643 for each pixel, as one challenge is the amount of computational costs of sensitivity calculations with the online
 644 pixel-by-pixel RT simulations. Nonetheless, random uncertainties of the observations can be reduced by spatial
 645 and temporal averaging, although the systematic uncertainties from the main retrieval steps remain. There remains
 646 lack of information to separate random and systematic uncertainties accurately. Here we provide a preliminary
 647 estimate of the uncertainty budget for monthly averaged HCHO and NO₂ columns from POMINO retrievals
 648 (Tables 3 and 4), based on our sensitivity tests and validations as well as previous work.

649 For HCHO, the systematic differential slant column uncertainty is 25% for regions with low columns and 15%
 650 for regions with elevated columns (De Smedt, 2022; De Smedt et al., 2018). The background correction
 651 uncertainty is significant for low columns (around 40%), in which the systematic uncertainty from the dSCD
 652 normalization is estimated to be 0 to 4×10^{15} molec.cm⁻², and the uncertainty from the model background is 0 to
 653 2×10^{15} molec.cm⁻². The AMF uncertainty, which is the largest contributor to the vertical column uncertainty, is
 654 mainly dependent on the errors of the ancillary parameters tested in Sect. 4. The AMF uncertainty induced by the
 655 error of a priori profile shapes is the largest with 30% to 60% over clean regions and around 20% over polluted
 656 regions. The errors of cloud parameters and surface reflectance are assumed to contribute to the AMF uncertainty
 657 by 10% to 20%, and the errors in the aerosol parameters contribute to the AMF uncertainty by about 5% for
 658 regions with low columns and 10% for regions with elevated columns. In addition, the uncertainty due to the
 659 aerosol overcorrection issue for partly cloudy pixels with low cloud height is estimated 10% to 15% (Sect. 4.1.1).
 660 Overall, the HCHO AMF uncertainty is estimated to be about 70% for clean regions and 30% for polluted regions,
 661 respectively.

662 **Table 3.** Estimated uncertainty budget of POMINO HCHO vertical columns for monthly mean low and elevated columns
 663 (higher than 10×10^{15} molec.cm⁻²).

	Remote regions / low columns	Elevated column regions / periods
Differential slant column uncertainties (De Smedt, 2022)	25%	15%
Background correction uncertainties (De Smedt, 2022)	40%	10%
● dSCD normalization uncertainties		$0 - 4 \times 10^{15}$ molec.cm ⁻²
● model background uncertainties		$0 - 2 \times 10^{15}$ molec.cm ⁻²
AMF uncertainties	70%	30%
● from a priori profiles uncertainties	60%	20%
● from aerosol correction uncertainties	5%	10%
● from surface reflectance uncertainties	20%	10%
● from cloud correction uncertainties	20%	10%
● from aerosol overcorrection issue uncertainties (only for partly cloudy pixels with low clouds)	15%	10%
Tropospheric vertical column uncertainty	85%	35%

664
 665 For NO₂, the total SCD uncertainty is reported to be 0.5 to 0.6×10^{15} molec.cm⁻² and a constant value of $0.2 \times$
 666 10^{15} molec.cm⁻² is assigned to the uncertainty of the stratospheric SCDs (Van Geffen et al., 2022b). For
 667 tropospheric AMF, the uncertainty caused by aerosol-related errors is estimated to be 10% to 20%, and the errors
 668 in a priori NO₂ profile shapes is estimated to cause an AMF uncertainty of 10% on average based on the sensitivity
 669 test. The contribution from cloud parameters and surface reflectance to the NO₂ AMF uncertainty is estimated to

670 be on the same level as that of a priori profile shapes. For pixels partly covered by low clouds over both clean and
 671 polluted regions, the AMF uncertainty contributed from the aerosol overcorrection issue is within 10%. By adding
 672 these errors in quadrature, the overall NO₂ AMF uncertainty is 25% to 30%.

673 **Table 4.** Estimated uncertainty budget of monthly mean POMINO NO₂ vertical columns.

	All regions
Total slant column uncertainties (Van Geffen et al., 2022b)	$0.5 - 0.6 \times 10^{15} \text{ molec.cm}^{-2}$
Stratospheric slant column uncertainties (Van Geffen et al., 2022b)	$0.2 \times 10^{15} \text{ molec.cm}^{-2}$
AMF uncertainties	25% – 30%
● from a priori profiles uncertainties	10%
● from aerosol correction uncertainties	10% – 20%
● from surface reflectance uncertainties	10%
● from cloud correction uncertainties	10%
● from aerosol overcorrection issue uncertainties (only for partly cloudy pixels with low clouds)	10%
Tropospheric vertical column uncertainty	$0.3 \times 10^{15} \text{ molec.cm}^{-2} + [0.2 \text{ to } 0.4] \times \text{VCD}$

Note: the uncertainty in the total slant columns is mostly absorbed by the stratosphere-troposphere separation step, and may not propagate into the tropospheric slant columns. (Van Geffen et al., 2015)

674
 675 By wrapping up the estimated relative contributions to the vertical column uncertainty, the total uncertainty of
 676 POMINO HCHO VCDs is estimated to be 85% over regions with low columns, and 35% over regions with high
 677 columns. For the POMINO NO₂ retrieval, the overall uncertainty budget can be approximated as 0.3×10^{15}
 678 $\text{ molec.cm}^{-2} + [0.2 \text{ to } 0.4] \times \text{VCD}$. This tentative estimation of the POMINO retrieval uncertainties is in agreement
 679 with the error analysis by De Smedt (2022) and Van Geffen et al. (2022b), and is supported by the validation
 680 results against the independent ground-based measurements (Sect. 6.1). Quantification of the errors at an
 681 individual pixel level have been achieved in previous studies (Boersma et al., 2004; Chong et al., 2024; Van Geffen
 682 et al., 2022b). As an alternative option to the Gaussian error propagation method, artificial-intelligence-based
 683 methods are an appealing approach to be tried in our future work.

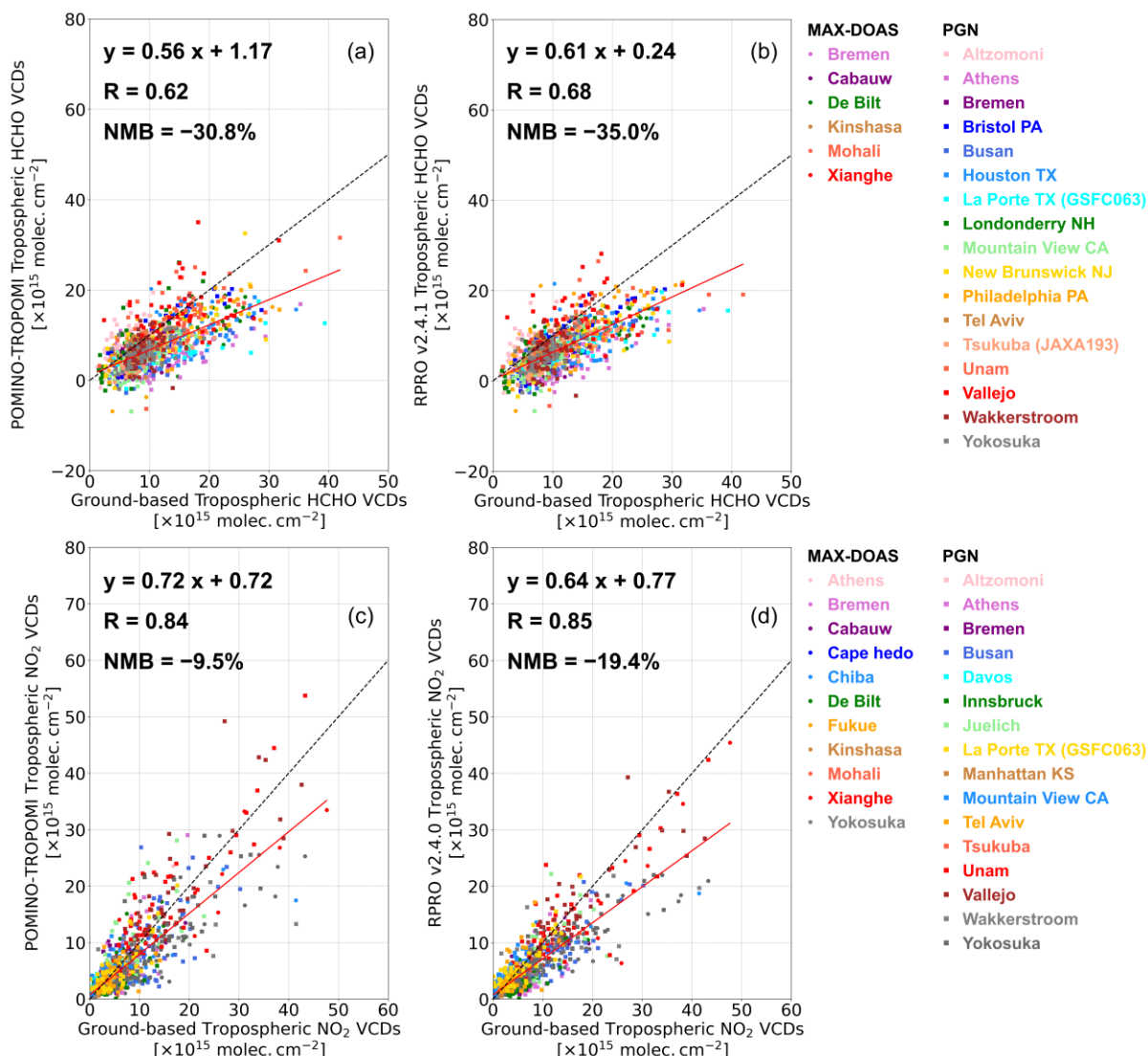
684 **6 Validation against global MAX-DOAS network and PGN measurements**

685 In this section, we present the validation results of POMINO and RPRO retrievals against independent ground-
 686 based measurements from the global MAX-DOAS network and PGN. Separate comparisons of tropospheric
 687 HCHO and NO₂ columns are given in Sect. 6.1, the effect of vertical smoothing is discussed in Sect. 6.2, and the
 688 satellite-based and ground-based FNRs are evaluated in Sect. 6.3.

689 **6.1 Validation of tropospheric HCHO and NO₂ columns**

690 Figures 12a and b present the scatterplots of daily satellite HCHO columns against ground-based measurements
 691 in April, July, October 2021 and January 2022. Each data point represents a day and site. There is a lower slope
 692 and higher positive offset for POMINO compared with those of RPRO product (slope: 0.56 versus 0.61; offset:
 693 1.17 versus 0.24). This is in line with the discussion in Sect. 4.5 that POMINO employs higher HCHO columns
 694 from GEOS-CF for background correction, which is the major component of HCHO columns over areas with low
 695 HCHO level. Furthermore, at 13 polluted ground-based sites where HCHO columns are higher than 10×10^{15}
 696 molec.cm^{-2} , POMINO HCHO columns show smaller bias at 8 sites (Figure S14). Overall, POMINO exhibits a

697 smaller negative NMB (−30.8%) than RPRO (−35.0%). Statistics of separate validation results against MAX-
 698 DOAS and PGN measurements are given in Table S3.



699
 700 **Figure 12.** Scatterplots of tropospheric HCHO (a and b) and NO₂ (c and d) columns between satellite products (POMINO and
 701 RPRO) and ground-based measurements in April, July, October 2021 and January 2022. The slope, offset and correlation from
 702 a linear regression using the robust Theil-Sen estimator and normalized mean bias (NMB) are given in each panel and plotted
 703 as the red line. The black dashed line is the 1:1 line. Each MAX-DOAS (marked by circles) and PGN site (marked by squares)
 704 is color-coded and listed on the right side.

705 For NO₂, a better agreement with ground-based measurements is found for POMINO tropospheric columns than
 706 for RPRO (slope: 0.72 versus 0.64; offset: 0.72 versus 0.77; NMB: −9.5% versus −19.4%). At remote MAX-
 707 DOAS sites where tropospheric NO₂ columns are around 1×10^{15} molec.cm^{−2} or less (Fig. S14), satellite
 708 tropospheric NO₂ columns are higher by $0.3\text{--}1 \times 10^{15}$ molec.cm^{−2}. This is in line with the previous validation
 709 studies (Kanaya et al., 2014; Pinardi et al., 2020; Verhoelst et al., 2021; Zhang et al., 2023), and is probably
 710 because that a majority of NO₂ molecules over remote regions are in the free troposphere, which are above the
 711 detection height of ground-based MAX-DOAS instruments but can be well observed by spaceborne instruments.
 712 At the six most-polluted sites with mean tropospheric NO₂ columns higher than 10×10^{15} molec.cm^{−2}, POMINO
 713 features a much-reduced bias of −14.5% compared with RPRO product (−22.0%). This is because of the explicit

714 correction for aerosol “shielding” effect over highly polluted sites and lower surface reflectance, which reduces
 715 the NO₂ scattering weights near the surface and hence increases the retrieved NO₂ columns.

716 6.2 Effect of vertical smoothing for validation

717 To test the impact of different vertical sensitivity from the ground and space, MAX-DOAS FRM₄DOAS v01.01
 718 harmonized HCHO and NO₂ datasets were used. The data provides 20-layer-resolved (from surface to ~ 600 hPa)
 719 MAX-DOAS averaging kernels and vertical profiles (posterior and prior to the retrievals). Following the “vertical
 720 smoothing” technique (Rodgers and Connor, 2003) described in detail by Vigouroux et al. (2020), we first
 721 substituted the priori profile shapes used in MAX-DOAS retrieval with either GEOS-CF or TM5-MP profile
 722 shapes to get corrected MAX-DOAS retrieved profiles:

$$723 \quad \mathbf{x}'_{\text{MD}} = \mathbf{x}_{\text{MD}} + (\mathbf{A}_{\text{MD}} - \mathbf{I})(\mathbf{x}_{\text{MD},a} - \mathbf{x}_{\text{Sat},a}) \quad (5)$$

724 with \mathbf{x}'_{MD} denoting the corrected MAX-DOAS retrieved profile, \mathbf{x}_{MD} the original MAX-DOAS profile, \mathbf{A}_{MD} the
 725 MAX-DOAS averaging kernel matrix, \mathbf{I} the unit matrix, $\mathbf{x}_{\text{MD},a}$ the MAX-DOAS a priori profile and $\mathbf{x}_{\text{Sat},a}$ the
 726 satellite a priori profile (i.e., from GEOS-CF or TM5-MP) re-gridded to the MAX-DOAS retrieval resolution from
 727 the surface to 600 hPa. To account for the trace gas content in the free troposphere, especially for HCHO, we
 728 further extend the corrected MAX-DOAS profile to the tropopause with the satellite profile above 600 hPa that is
 729 scaled to ensure vertical continuity of the overall tropospheric profile. After that, we perform the smoothing
 730 process using either POMINO or RPRO averaging kernels:

$$731 \quad c_{\text{MD}}^{\text{smoothed}} = \mathbf{a}_{\text{Sat}} \cdot \mathbf{x}'_{\text{MD}} \quad (6)$$

732 with $c_{\text{MD}}^{\text{smoothed}}$ the smoothed MAX-DOAS column, \mathbf{a}_{Sat} the satellite averaging kernel vector and \mathbf{x}'_{MD} the
 733 corrected MAX-DOAS retrieved profile from Eq. (5). We compare the smoothed MAX-DOAS data with satellite
 734 retrievals and the statistics are summarized in Table 5.

735 For the five MAX-DOAS sites available (Table 2), we find that after smoothing, the linear regression slope gets
 736 improved for both HCHO products. The negative bias of POMINO is reduced by about 10% but that of RPRO
 737 product is increased by about 4%. This is because POMINO HCHO averaging kernels are smaller than those of
 738 RPRO between the surface to about 800 hPa, resulting in lower smoothed MAX-DOAS HCHO columns compared
 739 to those using RPRO HCHO averaging kernels. Smaller POMINO HCHO averaging kernels at low altitudes are
 740 due to enhanced “shielding” effect from explicit aerosol corrections and lower KNMI TROPOMI MLER than
 741 OMI-based climatological monthly MLER used in RPRO HCHO.

742 For NO₂, among the six sites (Table 2), after applying the vertical smoothing technique, the negative NMB
 743 increases from -7.3% to -15.7% for POMINO and decreases from -24.6% to -8.5% for RPRO, even though a
 744 better day-to-day correlation is found for both products. Again, such changes are caused by the different averaging
 745 kernels used in the two satellite products.

746 Due to the scarcity of the MAX-DOAS sites for analysis here (Tables 2 and 5) and the under-representativeness
 747 in their spatial distribution (Table 2), a general conclusion cannot be made on the overall impact of vertical
 748 smoothing now. Nevertheless, the comparison results indicate the importance of considering the different vertical
 749 sensitivity between spaceborne and ground-based MAX-DOAS instruments, and different a priori profile shapes
 750 used to derive the vertical columns during the validation practice (De Smedt et al., 2021; Dimitropoulou et al.,
 751 2022; Yombo Phaka et al., 2023).

752 **Table 5.** Effect of vertical smoothing on the comparisons of TROPOMI and MAX-DOAS data.

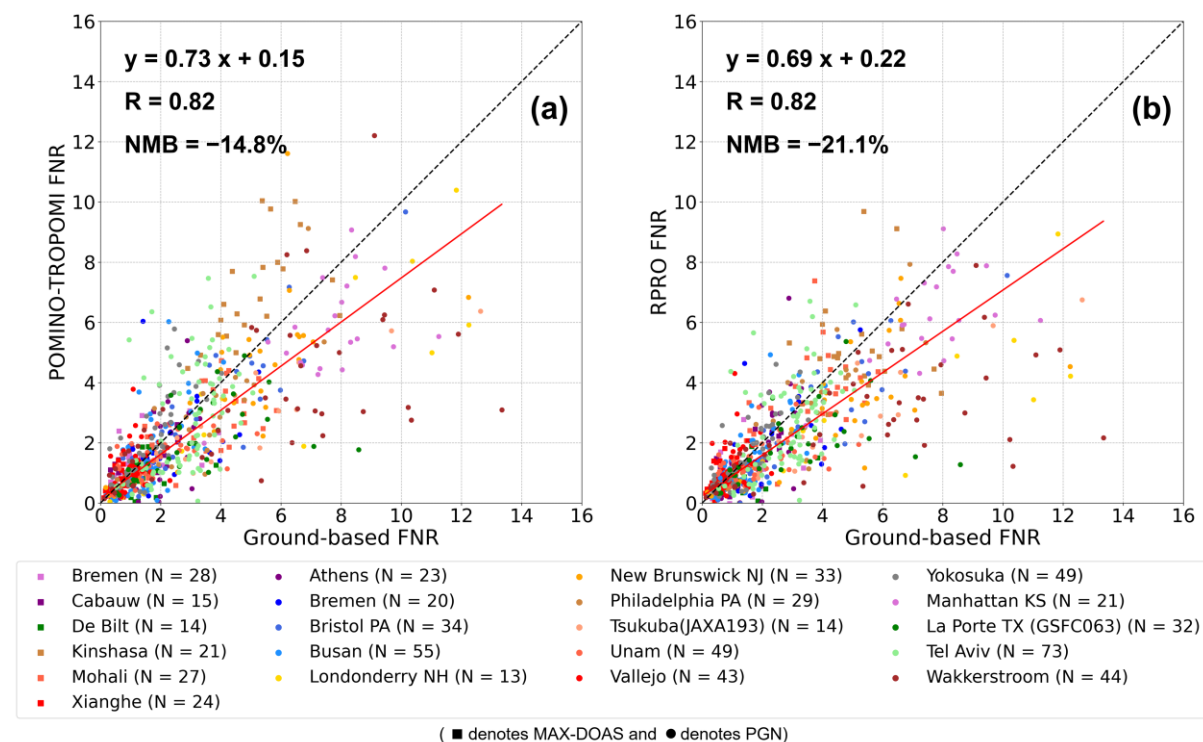
HCHO (five sites)	Direct comparisons		Vertical smoothing applied	
	POMINO	RPRO	POMINO	RPRO
Slope	0.56	0.65	1.08	0.72
Offset [10^{15} molec. cm^{-2}]	2.15	0.18	-1.58	-0.78
Correlation	0.63	0.66	0.66	0.73
NMB	-22.6%	-30.8%	-10.9%	-34.2%

NO ₂ (six sites)	Direct comparisons		Vertical smoothing applied	
	POMINO	RPRO	POMINO	RPRO
Slope	0.80	0.64	0.72	0.74
Offset [10^{15} molec. cm^{-2}]	0.38	0.46	0.74	0.98
Correlation	0.81	0.84	0.90	0.86
NMB	-7.3%	-24.6%	-15.7%	-8.5%

753

754 6.3 Comparisons of FNR

755 The FNR is an important space-based indicator of the ozone chemistry regimes and its sensitivity to precursor
756 emissions. Figure 13 shows the scatterplots of daily FNR derived from POMINO and RPRO products against
757 ground-based measurements, i.e., MAX-DOAS and PGN, in April, July, October 2021 and January 2022. A better
758 agreement is found between POMINO and ground-based FNR with improved linear regression statistics (slope:
759 0.73 versus 0.69; offset: 0.15 versus 0.22) and reduced NMB (-14.8% versus -21.1%) compared to those of
760 RPRO products. Moreover, the regression results are better in the comparisons for FNR than those in the
761 individual comparisons for either HCHO or NO₂ tropospheric VCDs (Sect. 6.1). This demonstrates the potential
762 of using POMINO HCHO and NO₂ retrievals to improve the studies on the ozone sensitivity analysis for NO_x as
763 well as VOC emission controls.



764

765 **Figure 13.** Scatterplots of daily tropospheric column ratio of formaldehyde to nitrogen dioxide (FNR) derived from satellite
766 products (a for POMINO and b for RPRO) and ground-based measurements in April, July, October 2021 and January 2022.
767 The slope, offset and correlation from a linear regression using the robust Theil-Sen estimator and normalized mean bias (NMB)
768 are given in each panel and plotted as the red line.

769 Note that most ground-based sites used here are in the North America, Europe, South Korea and Japan, but very
770 few or even no sites in other countries or continents (Figure S2). Thus, further validation with ground-based
771 measurements in combination with model simulations is needed over other regions, especially those where ozone
772 chemistry regimes change rapidly.

773 **7 Conclusions**

774 We developed an updated version of the POMINO algorithm providing HCHO and NO₂ AMF calculations, which
775 offers global tropospheric HCHO and NO₂ VCDs retrievals of TROPOMI with improved consistency compared
776 to current products. Compared to the independently developed RPRO HCHO and NO₂ operational algorithms
777 using different ancillary parameters, the POMINO algorithm includes: (1) the surface reflectance anisotropy by
778 using KNMI TROPOMI v2.0 DLER at 340 nm for HCHO and MODIS BRDF coefficients around 470 nm for
779 NO₂, (2) an explicit aerosol correction for both species based on GEOS-CF aerosol information and MODIS AOD
780 at corresponding wavelengths, (3) high-resolution (0.25° × 0.25°) a priori HCHO and NO₂ profile shapes from
781 GEOS-CF dataset and (4) a consistent cloud correction based on cloud top pressures taken from the FRESCO-S
782 cloud product and cloud fractions re-calculated at 440 nm using the same ancillary parameters as those used in
783 NO₂ AMF calculation.

784 High qualitative agreement of tropospheric HCHO and NO₂ columns is found between POMINO and RPRO
785 products in April, July, October 2021 and January 2022. However, RPRO HCHO columns are lower by 15% on
786 average than the POMINO HCHO columns over the polluted areas around the world, and the negative differences
787 of RPRO tropospheric NO₂ columns can reach -20% over specific areas.

788 To clarify the reasons for the differences between POMINO and RPRO columns and quantify the structural
789 uncertainty from ancillary parameters in the AMF calculation, we performed a series of sensitivity tests on the
790 cloud correction, aerosol correction, surface reflectance and a priori profile shapes. We find that based on
791 POMINO-recalculated cloud fraction at 440 nm and FRESCO-S cloud top pressures, differences between clear-
792 sky AMFs and total AMFs vary from -25% to more than 50% for both HCHO and NO₂, depending on the cloud
793 fraction and the relative height between clouds and trace gases. When using cloud top pressure data from
794 OCRA/ROCINN-CRB instead of FRESCO-S, a large decrease of tropospheric HCHO columns is found ($> 2 \times$
795 10^{15} molec.cm⁻²) over Amazonia Rainforest and southeast China, and the negative differences over polluted
796 regions are about 20% on average.

797 The influence of the implicit aerosol corrections used in operational products is within 10% on the HCHO retrieval,
798 while higher NO₂ columns by 20% to 40% over the polluted areas in January 2022 are found with implicit aerosol
799 corrections. Comparisons of retrieved NO₂ columns using clear-sky AMFs and total AMFs with implicit aerosol
800 corrections prove that the positive difference for NO₂ is dominated by the enhanced “shielding” effect of clouds
801 over NO₂ layers. The directionality of the surface reflectance has a very small impact on the HCHO retrieval in
802 the UV band, but the structural uncertainty of surface reflectance for NO₂ over polluted areas can reach 30%. The
803 HCHO structural uncertainty from a priori profile shapes is 20% to 30% over the background areas and 10% to
804 20% over the polluted areas. In contrast, the NO₂ differences due to different a priori profile shapes reach 30% or
805 more over the polluted areas. The additional test on the joint effect of these parameters shows notable non-linear
806 influences from aerosol correction, surface reflectance, cloud correction and a priori profile shapes in the RT
807 calculation.

808 Direct comparisons of tropospheric HCHO and NO₂ columns between satellite retrievals and ground-based
809 measurements from the global MAX-DOAS network and PGN show that both POMINO HCHO and NO₂
810 retrievals feature a reduced bias in comparison to RPRO products (HCHO: -30.8% versus -35.0%; NO₂: -9.5%
811 versus -19.4%), especially at the polluted sites. The effect of the vertical smoothing is significant and strongly
812 depends on the satellite averaging kernels. A better agreement of daily FNR with smaller bias is also found
813 between POMINO products and PGN measurements in comparison to results obtained with RPRO products
814 (NMB: -14.8% versus -21.1%).

815 Overall, we demonstrate the promising performance of TROPOMI-based POMINO algorithm for global HCHO
816 and NO₂ retrieval. However, there are still several limitations in our study. First, the aerosol overcorrection issue
817 for partly cloudy pixels exists in the current POMINO algorithm, which has been discussed in detail in Sect. 4.1.1.
818 The uncertainty due to this issue is estimated to be within 15% for HCHO and 10% for NO₂. Given that
819 TROPOMI-based O₂-O₂ cloud data have become available, we plan to improve the current POMINO algorithm
820 by performing O₂-O₂ cloud retrieval for both cloud fraction and cloud top pressure with explicit aerosol
821 corrections in the future, as has been done in the POMINO-OMI and POMINO-GEMS products (Lin et al., 2015;
822 Liu et al., 2019; Zhang et al., 2023).

823 Second, it should be noted that the indirect aerosol effect on HCHO and NO₂ retrievals through clouds is strongly
824 sensitive to the cloud top pressures and the trace gas profile shapes. Using OMI O₂-O₂ based cloud parameters or
825 FRESCO-S cloud top pressures stored in the operational NO₂ L2 product before version 1.4.0, previous studies
826 have shown lower NO₂ columns over polluted North China Plain when retrieved with implicit aerosol corrections
827 (Lin et al., 2015; Liu et al., 2020). This is because the cloud top pressures in those studies are higher, which result
828 in larger AMF values when implicit (instead of explicit) aerosol corrections are used. Besides, certain biases still
829 exist in the current FRESCO-S cloud top pressures, such as the overestimation over the ITCZ. The effect of a
830 priori profile shapes is also significant for both HCHO and NO₂ retrievals, and it deserves more attention in the
831 future analysis. Comprehensive evaluations of cloud retrievals and model performance with independent
832 measurements are needed in future studies.

833 Nevertheless, the POMINO algorithm that aims at improving the consistency in multi-gas retrieval shows great
834 potential and can be easily adapted to other satellite instruments, e.g. GEMS, TEMPO, as well as Sentinel-4 and
835 Sentinel-5 missions. The global tropospheric HCHO and NO₂ VCD retrievals presented in our study are also of
836 value for subsequent applications such as ozone chemistry analysis and emission controls.

837
838 *Data availability.* The POMINO HCHO and NO₂ datasets will be available soon on our website ([http://www.pku-](http://www.pku-atmos-acm.org/acmProduct.php)
839 [atmos-acm.org/acmProduct.php](http://www.pku-atmos-acm.org/acmProduct.php)). Before release, the data presented in the study are available from the
840 corresponding authors upon request. The S5p TROPOMI RPRO HCHO v2.4.1 L2 product and RPRO NO₂ v2.4.0
841 L2 product are available at Copernicus Data Space Ecosystem | Europe's eyes on Earth
842 (<https://dataspace.copernicus.eu/>, last access: 17 July 2024). The ground-based MAX-DOAS measurements can
843 be provided upon request to the corresponding authors. The PGN/Pandora direct sun measurements are available
844 at the ESA Validation Data Centre (EVDC, 2024) (<https://evdc.esa.int>, last access: 7 July 2024) and Pandonia
845 Global Network (2024) (<https://www.pandonia-global-network.org/>, last access: 17 July 2024).

846
847 *Supplement.*

848

849 *Author contributions.* YZ, JL, NT and MVR conceived this research. YZ, HY, IDS, JL, NT and MVR designed
850 the algorithm. YZ, HY, IDS, JL, MVR, GP, AM and SC designed the validation process together. YZ performed
851 all calculations. RS provided LIDORT model. RN, FR, SW, LC, JVG, ML, WS and LF provided data and technical
852 support for satellite retrievals. GP and SC provided methodological support for validation. AMC is the network
853 principal investigator (PI) for PGN instruments. GP, SC, AMC and MT provided the discussion for PGN
854 uncertainty estimation. MVR, GP, AM, MMF, AR, AP, VK, VS, TW, YC, HT, YK and HI provided ground-based
855 MAX-DOAS measurements. YZ wrote the paper with inputs from JL, NT, IDS and MVR. All co-authors revised
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857

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