

1 **Highly resolved mapping of NO₂ vertical column densities from** 2 **GeoTASO measurements over a megacity and industrial area during** 3 **the KORUS-AQ campaign**

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12 **Abstract.** The Korea-United States Air Quality (KORUS-AQ) campaign is a joint study between the United States National
13 Aeronautics and Space Administration (NASA) and the South Korea National Institute of Environmental Research (NIER) to
14 monitor megacity and transboundary air pollution around the Korean Peninsula using airborne and ground-based
15 measurements. Here, tropospheric nitrogen dioxide (NO₂) slant column density (SCD) measurements were retrieved from
16 Geostationary Trace and Aerosol Sensor Optimization (GeoTASO) L1B data during the KORUS-AQ campaign (May 2 to
17 June 10, 2016). The retrieved SCDs were converted to tropospheric vertical column densities using the air mass factor (AMF)
18 obtained from a radiative transfer calculation with trace gas profiles and aerosol property inputs simulated with the Community
19 Multiscale Air Quality (CMAQ) model and surface reflectance data obtained from the Moderate Resolution Imaging
20 Spectroradiometer (MODIS). For the first time, we examine highly resolved (250 m × 250 m resolution) tropospheric NO₂
21 over the Seoul and Busan metropolitan regions, and the industrial regions of Anmyeon. We reveal that the maximum NO₂
22 VCDs were 4.94×10^{16} and 1.46×10^{17} molecules cm⁻² at 9 AM and 3 PM over Seoul, respectively, 6.86×10^{16} and $4.89 \times$
23 10^{16} molecules cm⁻² in the morning and afternoon over Busan, respectively, and 1.64×10^{16} molecules cm⁻² over Anmyeon.
24 The VCDs retrieved from the GeoTASO airborne instrument were correlated with those obtained from the Ozone Monitoring
25 Instrument (OMI) ($r = 0.48$), NASA's Pandora Spectrometer System ($r = 0.91$), and NO₂ mixing ratios obtained from in situ
26 measurements ($r = 0.07$ in the morning, $r = 0.26$ in the afternoon over the Seoul, and $r > 0.56$ over Busan). Based on our results,
27 GeoTASO is useful for identifying NO₂ hotspots and their spatial distribution in highly populated cities and industrial areas.

28 **1 Introduction**

29 Nitrogen dioxide (NO₂) is one of the most important atmospheric trace gases and plays a key role in aerosol production and
30 tropospheric ozone photochemistry (Boersma et al., 2004; Richter et al., 2005). Furthermore, high concentrations of NO₂ in
31 the atmosphere have adverse effects on human health, such as respiratory infections, and associated symptoms (Brauer et al.,
32 2002; Latza et al., 2009).

33 The main sources of NO₂ in the atmosphere are fossil fuel combustion from vehicles and thermal power plants, lightning, and
34 biogenic soil processes. Furthermore, NO₂ concentrations are highly correlated with population size (Lamsal et al., 2013). The
35 implementation of emission control technology and environmental regulation has led to a decrease in surface NO₂
36 concentrations in Western Europe, the United States, and Japan in the last few decades (Richter et al., 2005). The concentration
37 of NO₂ in major metropolitan cities in South Korea and China is over 3 times larger than over similarly sized cities in Europe
38 and United States, despite NO₂ concentration decreasing in China and South Korea (de Foy et al., 2016, Choo et al., 2020).

39 To date, several low-orbit space borne sensors, such as the Global Ozone Monitoring Experiment (GOME) (Burrows et al.,
40 1999), the Scanning Imaging Spectrometer for Atmospheric Cartography (SCIAMACHY) (Burrows et al., 1995), the Ozone

41 Monitoring Instrument (OMI) (Levelt et al., 2006), the GOME-2 (Callies et al., 2000), and the Tropospheric Monitoring
42 Instrument (TROPOMI) (Veefkind et al., 2012), have monitored atmospheric ozone and its precursors including NO₂ and
43 formaldehyde (HCHO) as a proxy for volatile organic compounds (VOCs). Furthermore, the Geostationary Environment
44 Monitoring Spectrometer (GEMS) (Choi et al., 2018; Kim et al., 2020), which was launched on February 18, 2020, will form
45 a constellation of geostationary satellites including the upcoming Tropospheric Emission: Monitoring of Pollution (TEMPO)
46 (Zoogman et al., 2017) and Sentinel-4 platforms, to continuously observe the air quality of the Northern Hemisphere during
47 the day.

48 NO₂ retrievals from space borne hyperspectral measurements are typically conducted using the differential optical absorption
49 spectroscopy (DOAS) method (Platt and Stutz, 2008) to first retrieve the view-dependent slant column density (SCD), and
50 then radiative transfer models are used to determine the vertical column density (VCD) using an air mass factor (AMF)
51 correction. Previous and ongoing space borne instruments use various radiative transfer codes and model input assumptions to
52 calculate NO₂ AMF values at coarse spatial resolution. Because AMF weighting has a large impact on NO₂ retrievals using
53 the DOAS method, it is important to use model input assumptions that most accurately match viewing and atmospheric
54 conditions. Several studies have demonstrated the sensitivity of AMF calculations to inaccurate model input parameters (e.g.,
55 *a priori* NO₂ vertical profile and aerosol properties) and *a priori* data (cloud information and surface reflectance) (Leitão et
56 al., 2010; Hong et al., 2017; Lorente et al., 2017; Boersma et al., 2018). NO₂ retrievals have also been consistently conducted
57 based on surface remote sensing measurements including the Multi-Axis DOAS (MAX-DOAS), Système D'Analyse par
58 Observations Zénithales (SAOZ) spectrometer (Pastel et al., 2014), and Pandora (Herman et al., 2009) systems. These ground-
59 based measurements can be used as validation references for both airborne and space borne measurements.

60 NO₂ retrievals from airborne remote sensing instruments, such as the Geostationary Coast and Air Pollution Event (GEO-
61 CAPE) Airborne Simulator (GCAS) (Kowalewski and Janz, 2014), the Heidelberg Airborne Imaging DOAS Instrument
62 (HAIDI) (General et al., 2014), the Geostationary Trace gas and Aerosol Sensor Optimization (GeoTASO) (Leitch et al., 2014),
63 the Airborne Prism Experiment (APEX; Popp et al., 2012), the Airborne Imaging DOAS instrument for Measurements of
64 Atmospheric Pollution (AirMAP; Meier et al., 2017; Schönhardt et al., 2015), the Small Whiskbroom Imager for atmospheric
65 composition monitorinG (SWING; Merlaud et al. 2018), and the Spectrolite Breadboard Instrument (SBI; Vlemmix et al.,
66 2017; Tack et al., 2019) have also been performed to identify local emission sources and obtain highly resolved horizontal
67 NO₂ distributions.

68 Observations using airborne measurements have an advantage as they enable the observation of horizontal distributions of
69 trace gases at resolutions higher than those of space-based satellites and provide data over a wider area than those of ground-
70 based observations. For example, Nowlan et al. (2018) retrieved tropospheric NO₂ VCDs over Houston, Texas, during the
71 Deriving Information on Surface Conditions from Column and Vertically Resolved Observations Relevant to Air Quality
72 (DISCOVER-AQ) campaign and identified a high correlation with data retrieved from Pandora. Popp et al. (2012) also
73 presented the morning and afternoon NO₂ spatial distribution in Zurich, Switzerland, using APEX. Tack et al. (2017) have
74 conducted high-resolution mapping of NO₂ over three Belgium cities (Antwerp, Brussels, and Liège) using APEX and Judd
75 et al. (2020) and Tack et al. (2021) compared NO₂ VCDs retrieved from GCAS/GeoTASO and APEX with those obtained
76 from TROPOMI over New York City and Antwerp and Brussels, respectively. Merlaud et al. (2013) observed NO₂ VCDs in
77 Turceni over Romania using SWING mounted on an unmanned aerial vehicle (UAV) during the Airborne Romanian
78 Measurements of Aerosols and Trace gases (AROMAT) campaign. These existing NO₂ retrievals, using airborne
79 measurements, have been useful in constraining regional air quality models due to the highly resolved source identification
80 and the ability to tie these results to ground-based observations.

81 This work focuses on airborne NO₂ retrievals from GeoTASO. This instrument was developed by Ball Aerospace to reduce
82 mission risk for UV-VIS air quality measurements from geostationary orbit for the GEMS and TEMPO missions (Leitch et
83 al., 2014). The retrieval of NO₂, SO₂, and HCHO observed from GeoTASO L1B data using DOAS and principal component

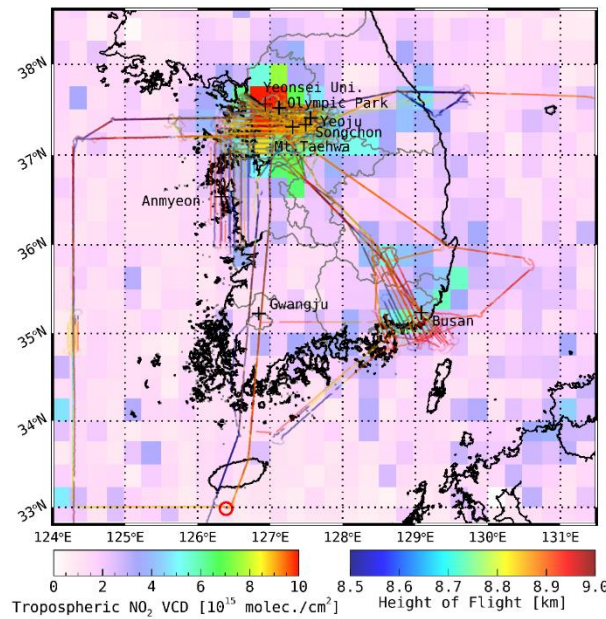
84 analysis (PCA) (Wold et al., 1987) was conducted through the DISCOVER-AQ and KORea-United States Air Quality
 85 (KORUS-AQ) campaigns (Nowlan et al., 2016; Judd et al., 2018; Choi et al., 2020; Chong et al., 2020). The KORUS-AQ
 86 campaign is a joint study between the National Institute of Environmental Research (NIER) and National Aeronautics and
 87 Space Administration (NASA) to monitor megacity air pollution and transboundary pollution, and to prepare for geostationary
 88 satellite (i.e., GEMS, TEMPO, and Sentinel-4) air quality observability (of trace gases and aerosols), organized from May to
 89 June 2016.

90 Although surface NO₂ concentrations in South Korea are the high due to high population density, high traffic volumes, and
 91 many industrial complexes and thermal power plants, and although NO₂ retrieval studies using airborne and ground
 92 measurements in North America, Europe, China, and Japan, data for South Korea remain limited. The specific objectives of
 93 this study are as follows:

- 94 (1) To retrieve tropospheric NO₂ vertical column data using GeoTASO measurements over polluted regions of the Seoul
 95 and Busan metropolitan areas and the Anmyeon industrial regions of the Korean Peninsula.
- 96 (2) To estimate NO₂ VCD uncertainties using error propagation accounting for spectral fitting errors and AMF
 97 uncertainties associated with input data errors, including aerosol optical depth (AOD), single scattering albedo (SSA),
 98 aerosol peak height (APH), and surface reflectance (SRF).
- 99 (3) To compare NO₂ VCDs retrieved from GeoTASO and those obtained from OMI and ground-based Pandora
 100 instruments, as well as surface in situ measurements.

101 2 KORUS-AQ campaign area, measurements, and model simulation

102 2.1 Campaign area



103
 104 **Figure 1. Flight paths of the NASA LaRC B200 aircraft carrying GeoTASO and the average tropospheric NO₂ VCDs**
 105 **obtained from OMI gridded to a 0.25° × 0.25° horizontal grid during the KORUS-AQ campaign period. The line color**
 106 **represents flight height. In this period, the GeoTASO observations focused on megacities (Seoul and Busan) and**
 107 **industrial complex area (Anmyeon) with high tropospheric NO₂ concentrations. The reference spectrum for spectral**
 108 **fitting is obtained from the radiation data over the Jeju Island (marked with red circle).**
 109

110 The Korean Peninsula, located on the Asia-Pacific coast, has a complex atmospheric environment by local emissions and long-
 111 range transport under appropriate weather conditions (Jeong et al., 2017; NIER and NASA, 2020; Choo et al., 2021). Seoul,
 112 the capital of South Korea, and the metropolitan area are densely populated, and power plants and industrial activities on the

113 northwest coast are conducted, which emit relatively large amounts of pollutants. The KORUS-AQ campaign conducted three-
 114 dimensional observations, including ground-based remote, aircraft, satellite observation, and air quality modeling, to
 115 understand the complex air quality and interpret the observations of GEMS launched in 2020. The KORUS-AQ campaign
 116 period was from May 2 to June 10, 2016. During the KORUS-AQ campaign, air pollutants were conducted using the GeoTASO
 117 on board the NASA Langley Research Center B200 aircraft to monitor air quality and long-range transport of pollutants over
 118 the Korean Peninsula (NIER and NASA, 2020). The GeoTASO observations were conducted 30 times in 23 d out of 40 d.
 119 Most observations were made once or twice a day. Each flight was planned and conducted on a day when the weather
 120 conditions were fine and flight hours were approximately 2-4 h. We show the average values of GeoTASO flight information
 121 such as flight time, altitude, speed, solar zenith angle (SZA), and viewing zenith angle (VZA) for the dates retrieved for NO₂
 122 VCD, aerosol properties (AOD, SSA) extracted from CMAQ, and cloud fraction and surface reflectance extracted from the
 123 Moderate Resolution Imaging Spectroradiometer (MODIS) in Table 1. Flight information on the date of aircraft observation
 124 can be found at http://www-air.larc.nasa.gov/missions/korus-aq/docs/KORUS-AQ_Flight_Summaries_ID122.pdf. Figure 1
 125 indicates the flight routes of B200 and the tropospheric NO₂ VCD obtained from the OMI during the campaign period. The
 126 observations were concentrated in the metropolitan areas of Seoul and Busan and the industrial areas of Anmyeon, with an
 127 average flight altitude of ~8.5 km during KORUS-AQ.

128

129 **Table 1. Summary of information on the dates when NO₂ VCD was retrieved during the KORUS-AQ period (LT = UTC + 9**
 130 **h). The average values of GeoTASO data sets for flight characteristics, aerosol properties, geometric information and cloud**
 131 **information.**

Date	Jun 5	Jun 9 AM	Jun 9 PM	Jun 10 AM	Jun 10 PM
ROI	Anmyeon	Seoul metropolitan		Busan metropolitan	
Flight time (LT)	13:11–17:20	7:48–12:00	13:46–17:52	8:02–11:38	13:05–15:19
Flight altitude (km)	8.6	8.4	8.5	8.6	8.5
Flight speed (km hr ⁻¹)	117.0	116.2	117.6	117.2	117.1
SZA (°)	39.2	36.1	45.3	35.9	33.0
VZA (°)	11.9	12.6	12.8	12.1	11.8
AOD	0.27	0.40	0.21	0.13	0.09
SSA	0.966	0.980	0.949	0.981	0.968
Surface reflectance	0.07	0.09	0.09	0.06	0.06

132

133 As shown in Fig. 1, GeoTASO observations were conducted focusing on highly NO₂-polluted regions in the Seoul and Busan
134 metropolitan areas and the Anmyeon region during the KORUS-AQ campaign. The Seoul metropolitan area (Seoul Special
135 City, Gyeonggi Province, and Incheon City) is one of the most densely populated areas worldwide, with a population of
136 approximately 20 million in 2016. Busan is the second-largest city in South Korea, with a population of approximately 3.4
137 million in 2016. Anmyeon is located southwest of Seoul, with petrochemical complexes, steel mills, and thermal power stations
138 in this area. The background color in Fig. 1 represents the average NO₂ VCD obtained from the OMI during the KORUS-AQ
139 campaign period, showing over 1×10^{16} molecules cm⁻² over the Seoul metropolitan area. The OMI data were obtained with
140 the Level 2.0 OMNO2 version 3.0 and downloaded from the NASA Earthdata search (<http://search.earthdata.nasa.gov/search/>).
141 We calculated the arithmetic means of tropospheric NO₂ VCDs, like Choo et al. (2020), to obtain the grid data (0.25° × 0.25°)
142 during the KORUS-AQ period. The average tropospheric NO₂ VCD data were excluded from May 30 2016 to Jun 9 2016,
143 when the OMI data did not exist during the campaign period.

144 2.2 Pandora

145 NO₂ VCDs retrieved from the GeoTASO were validated using those from NASA's Pandora Spectrometer system. The Pandora
146 spectrometer is a hyper-spectrometer that can provide direct sun measurements of UV-Vis spectra (280–525 nm with a full
147 width at half maximum (FWHM) of 0.6 nm) for observing atmospheric trace gases. During the KORUS-AQ, eight Pandora
148 instruments monitored NO₂ and ozone (O₃) VCD as depicted by plus symbols in Fig. 1. The retrieved data are available on the
149 KORUS-AQ pages of NASA's Goddard Space Flight Center website
150 (<https://avdc.gsfc.nasa.gov/pub/DSCOVER/Pandora/DATA/KORUS-AQ/>). We compared NO₂ VCDs obtained from five
151 Pandora measurements (Busan university: 35.24 °N, 129.08 °E; Olympic park: 37.52 °N, 127.13 °E; Songchon: 37.41 °N,
152 127.56 °E; Yeosu: 37.34 °N, 127.49 °E; Yonsei University: 37.56 °N, 126.93 °E) within 0.05° and 30 min with those from
153 GeoTASO. Because NO₂ has a short atmospheric lifetime, especially during the summer (Shah et al., 2020), its spatial and
154 temporal distributions vary notably. A detailed description of Pandora's operation during the KORUS-AQ campaign has
155 previously been reported (Herman et al., 2018; Spinei et al., 2018).

156 2.3 Ground-based in situ NO₂ measurement

157 Although the basic physical quantity of VCD and the surface mixing ratio from in situ measurements are different, comparison
158 of their spatiotemporal variations provides useful information for deriving surface air quality from airborne instruments (e.g.,
159 Jeong and Hong, 2021a; 2021b). In this study, we compare the NO₂ VCDs (molecules cm⁻²) retrieved from GeoTASO to
160 surface mixing ratios measured by ground-based in-situ monitoring network over South Korea (i.e., Air-Korea, a national real-
161 time air quality network; <https://www.airkorea.or.kr/>). The instruments use the chemiluminescence method (Kley and
162 McFarland, 1980), and approximately 400 air quality monitoring sites in Korea are registered in the system, providing hourly
163 surface NO₂ concentrations. We compared NO₂ VCDs retrieved from GeoTASO within 0.5 km and 30 min with NO₂
164 concentrations obtained from Air-Korea.

165 2.4 GeoTASO measurements

166 NO₂ VCDs were retrieved from the L1B radiance dataset (version: V02y) obtained using GeoTASO during the KORUS-AQ
167 campaign. The NASA Goddard Space Flight Center conducted the L1B radiance calibration, which included offset and smear
168 correction, gain matching, amplifier cross-talk correction, dark rate correction, integration normalization, sensitivity derivation,

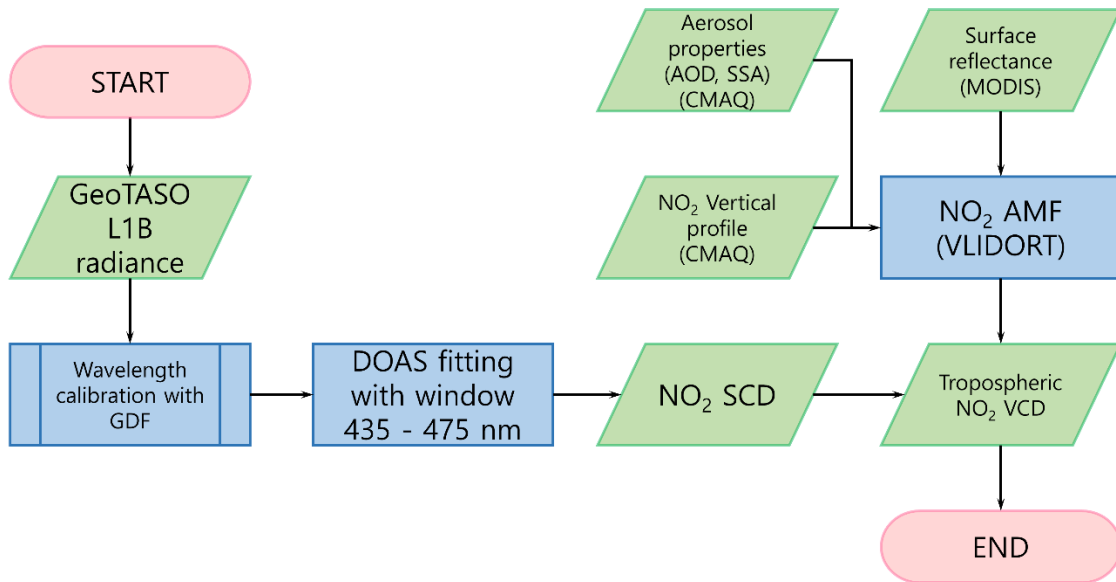
169 wavelength registration, geo-registration, non-linearity correction, and ground pixel geolocation (Kowalewski et al., 2017;
 170 Chong et al., 2020). The detailed specifications of GeoTASO are listed in Table 2 (Nowlan et al., 2016).

171

172 **Table 2. Summary of the GeoTASO instrument and optical specification.**

L1B version	V02y
Full cross-track field of view	45°
Single-pixel cross-track field of view	0.046°
Wavelength	UV: 290–400 nm
	VIS: 415–695 nm
Spectral resolution	UV: ~0.39 nm
(full width at half maximum, FWHM)	VIS: ~0.88 nm
CCD	1,056 (wavelength) × 1,033 (cross-track)
Spatial resolution before binning	~35 m (along-track) × 7 m (cross-track)
Spatial resolution after binning	~250 m (along-track) × 250 m (cross-track)

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174

175 **Figure 2. Flowchart of the algorithm for retrieving tropospheric NO₂ data from GeoTASO.**

176

177 2.4.1 NO₂ slant column density retrieval

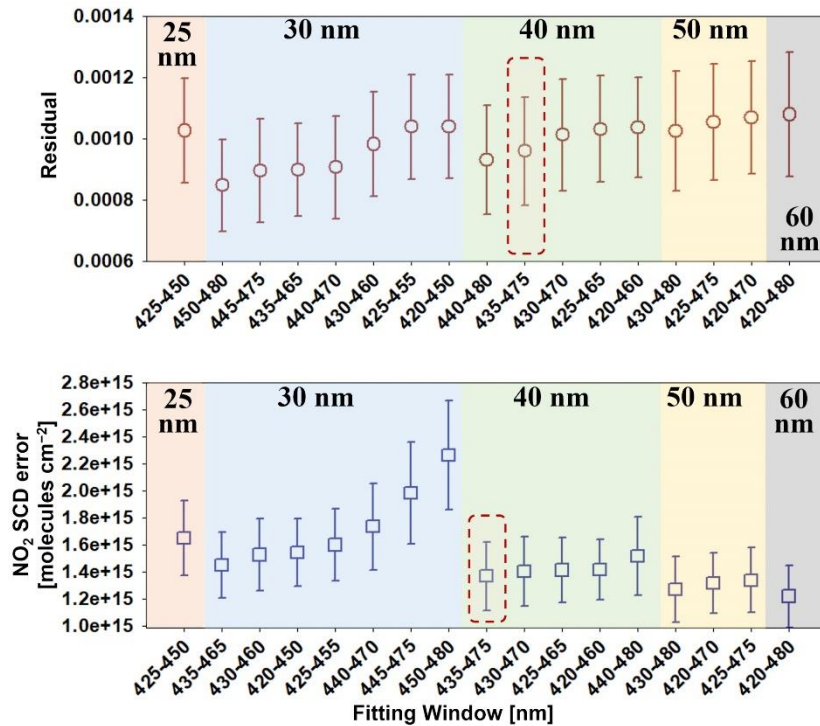
178 Figure 2 indicates the flowchart for retrieving the tropospheric NO₂ VCD from the GeoTASO. We first retrieved NO₂ SCDs
 179 using the DOAS method (Platt, 1994). Nonlinear least square minimization was used to retrieve the NO₂ SCDs which
 180 minimizes the difference between the measured optical depth and the modeled value in QDOAS software (Eq. (1); Danckaert
 181 et al., 2012).

$$182 \frac{\ln I(\lambda)}{\ln I_0(\lambda)} = -(\sum_{j=1}^m \rho_j \times \sigma'_j(\lambda) + B(\lambda) + R(\lambda) + A(\lambda) + N(\lambda)), \quad (1)$$

183

184 Where $I(\lambda)$ is the measured earthshine radiance at wavelength λ ; I_0 is the reference radiance from the reference sector (southern
 185 ocean of the Jeju Island denoted as the red circle in Fig. 1; 32.983°N, 126.392°E) at 9 AM on May 1 2016. The Community
 186 Multiscale Air Quality (CMAQ) modeling system data indicated that the NO₂ VCD from the surface to 50 hPa over this
 187 reference sector on this day was 6.75×10^{15} molecules cm⁻², and the mean of total NO₂ VCD obtained from the OMI during
 188 the KOURS-AQ period was 4.77×10^{15} molecules cm⁻² with a standard deviation of 1.33×10^{15} molecules cm⁻². We also
 189 confirmed the stability of NO₂ distribution over this area using the TROPOMI offline data from 2019 to 2020. In this period,
 190 the NO₂ VCD from the TROPOMI was 4.81×10^{15} molecules cm⁻² with a standard deviation of 0.43×10^{15} molecules cm⁻².
 191 The NO₂ VCD used as a reference sector obtained from CMAQ was mainly dominated by stratospheric NO₂ VCD. However,
 192 stratospheric NO₂ VCD has a relatively lower than tropospheric NO₂ VCD. The ρ_j represents the SCD of each species j ; $\sigma_j(\lambda)$
 193 represents the differential gas phase absorption cross-section convolved with the Gaussian distribution function (GDF) with
 194 GeoTASO FWHM (the UV and VIS range were 0.34–0.49 nm and 0.70–1.00 nm, respectively (Nowlan et al., 2016)) at
 195 wavelength λ of species j , respectively.

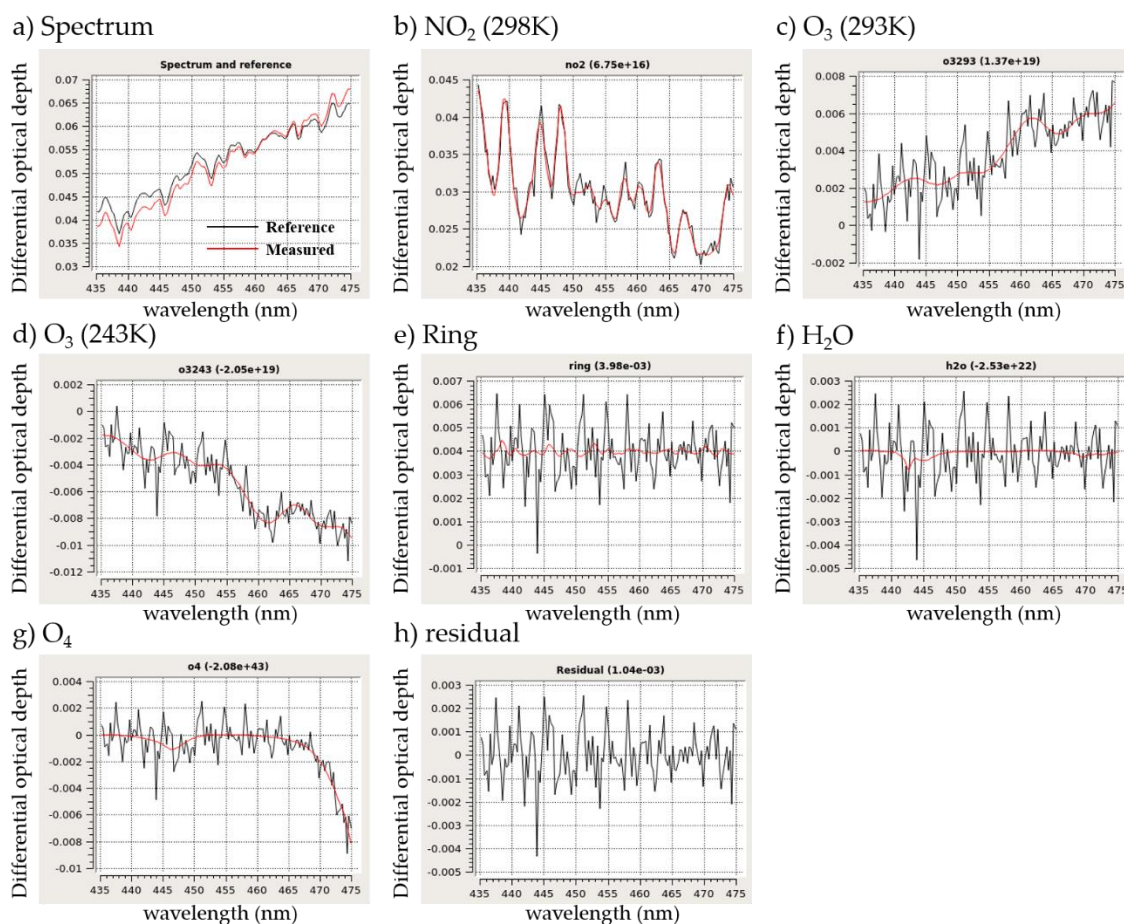
196 We used the measured radiances at the reference sector to calculate differential slant column density (dSCD) over the whole
 197 domain of the GetoTASO measurements. CMAQ calculation over the reference sector (i.e., 6.75×10^{15} molecules cm⁻²) was
 198 adopted as the reference SCD (SC_0), which is added to all dSCD values to convert to the SCD. The reference sector is known
 199 as a background area but is occasionally affected by the long-range transport of NO₂ from upwind areas. Considering the
 200 standard deviation of the OMI measurements accounts for such effects during the measurement period, we estimate the
 201 maximum uncertainties of the SC_0 can be calculated from this value (i.e., 1.33×10^{15} molecules cm⁻²) in addition to the
 202 difference of the mean values between the CMAQ and OMI (i.e., 1.98×10^{15} molecules cm⁻²). Therefore, our best estimate of
 203 the uncertainty of the SC_0 is the root of the sum of squares of these values (i.e., 2.38×10^{15} molecules cm⁻²).



205
 206 **Figure 3. Residuals and NO₂ SCD errors of 17 spectral fitting window candidates (May 17, 2016, across track number: 15).**

207
 208 The spectral fitting window was selected based on the sensitivity test with 17 fitting window candidates from 420 to 480
 209 with the length of the fitting window from 25 to 60 nm. Spectral fitting residuals and NO₂ SCD errors have been investigated
 210 for 17 spectral fitting window candidates (Fig. 3).

211 In terms of the residual, when the NO₂ fitting window includes a wavelength region less than 430 nm, it has a larger residual
 212 compared to the case where it does not. The higher residual can include the more noise signals that cannot be calculated
 213 mathematically, which can become an uncertainty for the NO₂ SCD retrievals. Therefore, we excluded the fitting window
 214 which includes wavelengths less than 430 nm for the GeoTASO NO₂ retrievals during the KORUS-AQ campaign. In the case
 215 of the NO₂ SCD error, it was confirmed that the longer the fitting window length, the lower the NO₂ SCD error appeared
 216 regardless of including the wavelength region less than 430 nm. Therefore, for the stable NO₂ SCD retrieval, an appropriate
 217 spectral fitting window needs to be selected which can minimize the residual with a moderate length of the fitting window. To
 218 find the optimal fitting window, we set the threshold value based on the above results: residual < 0.001, NO₂ SCD error < 1.4
 219 $\times 10^{15}$ molecules cm⁻², the length of fitting window > 30 nm. Then, the fitting window of 435–475 nm was selected for the
 220 GeoTASO NO₂ retrievals during the KORUS-AQ campaign. To determine the wavelength registration more accurately in the
 221 narrow fitting window, additional wavelength calibration of the spectra for each of the 33 across track pixels was performed
 222 using a high-resolution solar reference spectrum (Kurucz solar spectrum) (Chance and Kurucz, 2010) with the GDF. The
 223 absorption cross-sections of NO₂ (Vandaele et al., 1998), O₃ (Bogumil et al., 2000), H₂O (Rothman et al., 2010), and the Ring
 224 effect as pseudo-absorbers (Chance and Spurr, 1997) were used to construct the model equation; and B(λ), R(λ), A(λ), and
 225 N(λ) are the broad absorption of trace gases, extinction by Mie and Rayleigh scattering, variation in the spectral sensitivity of
 226 the detector or spectrograph, and noise, respectively, which were accounted for by an 8th order polynomial. An example of the
 227 spectral fitting results is presented in Fig. 4.
 228



229
 230 **Figure 4.** An example of the spectral fitting results of NO₂ retrievals from GeoTASO during the KORUS-AQ campaign (at Gangnam,
 231 Seoul on June 9, 2016). Red and black line in the panel (a) represent measured and reference spectrum, respectively. The panels of
 232 (b) to (h) depict examples of spectral fitting results of (b) NO₂, (c) O₃ (293 K), (d) O₃ (243 K), (e) ring, (f) H₂O, (g) O₄, where red and
 233 black lines are the absorption cross section of target species and the fitting residual plus the absorption of the target species,
 234 respectively. The panel (h) indicates the fitting residual of this example.

235

236 2.4.2 NO₂ AMF calculation

237 AMF, the ratio of SCD to VCD, can be calculated using the scattering weight (ω) and shape factor (S) (Palmer et al., 2001) in
238 Eq. (2)–(5).

$$239 \quad AMF = \frac{SCD}{VCD} \quad (2)$$

$$240 \quad AMF = AMF_G \int_{z_1}^{z_2} \omega(z)S(z)dz, \quad (3)$$

$$241 \quad \omega(z) = -\frac{1}{AMF_G} \frac{\partial \ln I_B}{\partial \tau}, \quad (4)$$

$$242 \quad S(z) = \frac{\alpha(z)n(z)}{\int_{z_1}^{z_2} \alpha(z)n(z)dz'} \quad (5)$$

243

244 Where AMF_G represents the geometric AMF, I_B is the earthshine radiance, τ is the optical depth, α is the absorption cross-
245 section, and n is the number density of the absorber. NO₂ AMF was calculated using a linearized pseudo-spherical scalar and
246 vector discrete ordinate radiative transfer model (VLIDORT, version 2.6; Spurr and Christi, 2014). Aerosol properties, such
247 as AOD, SSA, APH, and *a priori* NO₂ vertical profile information, were simulated using the CMAQ, and surface reflectivity
248 was obtained from MODIS (Collection 6). The surface reflectance products, MCD43A3, available at a 500 m spatial resolution,
249 provide an estimate of the surface spectral reflectance including MODIS bands 1 through 7. Here, MODIS band 3 (459–479
250 nm) was used, because this band is the closest the wavelength (455 nm) used in the calculation of AMF in this study. APH
251 was assumed to be the peak height of the aerosol extinction coefficient simulated in CMAQ, and the aerosol profile applied
252 GDF based on APH (Hong et al., 2017). For pixels without reflectance information, AMF was not calculated. The products
253 were corrected for atmospheric conditions, such as aerosol, gases, and Rayleigh scattering. In previous studies (Lamsal et al.,
254 2017; Nowlan et al., 2018; Judd et al., 2019; Chong et al., 2020), an AMF was described for both above and below aircraft
255 altitude is used to convert NO₂ SCDs to VCDs using Eq. (6)–(8).

$$256 \quad AMF \uparrow = AMF_G \int_{z_A}^{z_{TOA}} \omega(z)S(z)dz, \quad (6)$$

$$257 \quad AMF \downarrow = AMF_G \int_{z_0}^{z_A} \omega(z)S(z)dz, \quad (7)$$

$$258 \quad NO_2 \text{ VCD} \downarrow = \frac{NO_2 \text{ SCD} - AMF \uparrow \cdot NO_2 \text{ VCD} \uparrow}{AMF \downarrow}, \quad (8)$$

259 Where $AMF \uparrow$ and $AMF \downarrow$ are AMF above and below aircraft, respectively, and $NO_2 \text{ VCD} \uparrow$ represents NO₂ VCD above the
260 aircraft obtained from a chemical transport model (CTM). However, here we calculated $NO_2 \text{ VCD} \downarrow$ by dividing NO₂ SCDs
261 by $AMF \downarrow$ as the CMAQ only simulates the troposphere (surface to 50 hPa). However, as the stratospheric and free tropospheric
262 NO₂ ($NO_2 \text{ VCD} \uparrow$) column densities over megacities and industrial areas are much lower than tropospheric NO₂ column
263 densities, (Valks et al., 2011), we assume that the uncertainties in the AMF without considering the upper atmosphere are
264 negligible in this study.

265

266 2.5 Chemical model description

267 Vertical profiles from CMAQ (Byun and Ching, 1999; Byun and Schere, 2006), a CTM, were used to calculate AMFs. The
268 CMAQ simulations were conducted with a horizontal resolution of 15 × 15 km and had 27 vertical layers from the surface to
269 50 hPa. The meteorological fields were prepared using the advanced research Weather Research and Forecasting (WRF)
270 Advanced Research WRF (ARW) Model (Skamarock et al., 2008). Anthropogenic emissions were generated based on the
271 KORUS v5.0 model (Woo et al., 2012), and biogenic emissions were simulated using the Model of Emissions of Gases and
272 Aerosols from Nature (MEGAN v2.1; Guenther et al., 2006; 2012). Besides anthropogenic and biogenic emissions, the Fire
273 Inventory from NCAR (FINN; Wiedinmyer et al., 2006, 2011) was used to update the pyrogenic emission fields.

274 The CMAQ AOD was calculated by integrating the aerosol extinction coefficient (Q_{ext}), which is the sum of scattering (Q_{sca})
 275 and absorption (Q_{abs}) coefficients, over all vertical layers (z) as follows:

$$276 \quad AOD = \int Q_{ext}(z) dz = \int \{Q_{sca}(z) + Q_{abs}(z)\} dz, \quad (9)$$

$$277 \quad Q_{abs}[Mm^{-1}] = \sum_i \sum_j \{ (1 - \omega_{ij}) \cdot \beta_{ij} \cdot f_{ij}(RH) \cdot [C]_{ij} \}, \quad (10)$$

$$278 \quad Q_{sca}[Mm^{-1}] = \sum_i \sum_j \{ \omega_{ij} \cdot \beta_{ij} \cdot f_{ij}(RH) \cdot [C]_{ij} \}, \quad (11)$$

279 Here, ω_{ij} indicates SSA of particulate species i for the particulate mode (or size bin) j , β_{ij} denotes the mass extinction
 280 efficiency, $f_{ij}(RH)$ is the hygroscopicity factor according to the relative humidity (RH), and $[C]_{ij}$ is the concentration of
 281 particulate species. CMAQ SSA is defined as the ratio of the integrated Q_{sca} to AOD, and NO_2 vertical profiles were obtained
 282 from NO_2 concentrations at each vertical layers by conducting CMAQ simulations. Details of the model descriptions and
 283 calculations of optical properties are given by Lee et al. (2020) and Malm and Hand (2007).

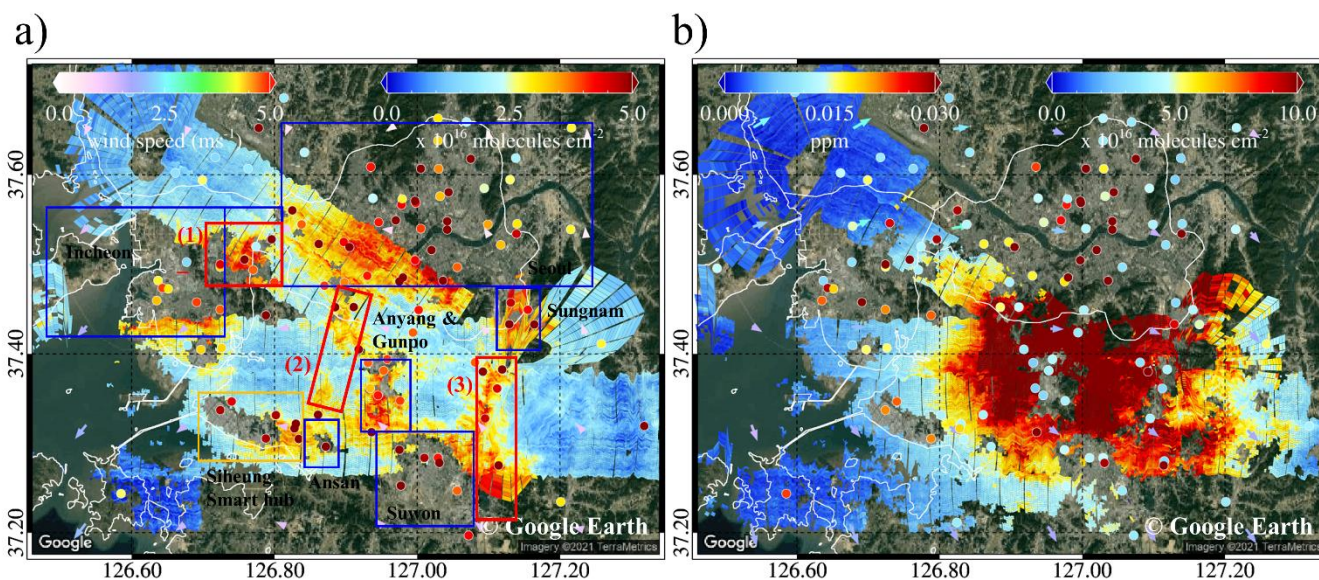
284 3 Results and discussion

285 3.1 NO_2 VCD retrieval

286 3.1.1 Seoul metropolitan region

287 We show the final NO_2 VCDs from 250 m spatial resolution. Because of NO_2 VCD, we selected the dates observed in both
 288 the morning and afternoon during the KORUS-AQ period over the Seoul metropolitan area, Busan, and Anmyeon. The
 289 retrieved dates for NO_2 VCDs were Jun 5, 9, and 10, 2016.

290 The population of the Seoul metropolitan region is approximately 20 million, which is approximately 40% of the total
 291 population of South Korea. It is rare to obtain high-resolution horizontal NO_2 VCD distributions using airborne measurements
 292 in the morning and afternoon, especially in Asian megacities. Figure 5 indicates tropospheric NO_2 VCDs over Seoul on June
 293 9 2016, at 9 AM and 3 PM local time (LT). Because of an issue with imaging systems, enlarged views (Fig. 5-Fig. 8) present
 294 a slightly stripy appearance from the GeoTASO observation (Nowlan et al., 2016; Chong et al., 2020).



295
 296 **Figure 5. Tropospheric NO_2 VCD, in the Seoul metropolitan region on June 9, 2016, retrieved from GeoTASO: a) at 9 AM and b) at**
 297 **3 PM. The red boxes represent expressways (counterclockwise from left to right, (1) Gyeongin Expressway, (2) Seohaean Expressway,**
 298 **and (3) Gyeongbu Expressway), the orange box indicates the industrial complex, and the blue boxes indicate the major cities (Seoul,**
 299 **Incheon, Suwon, Bucheon, Anyang, Gunpo, Sungnam, and Ansan) of the Seoul metropolitan region. Colors of the circles depict the**
 300 **NO_2 surface mixing ratio obtained from Air-Korea. The color arrows indicate the wind direction and speed at 1000 hPa over Seoul**
 301 **metropolitan region, obtained via the Unified Model (UM) simulations (background RGB image is from Google Earth;**
 302 **<https://www.google.com/maps/>).**

303

304 In the morning, NO₂ VCDs retrieved from GeoTASO were highly correlated with expressways (red boxes in Fig. 5), such as
 305 the Gyeongin, Seohaean, and Gyeongbu Expressways, and over major cities with heavy traffic, such as Seoul, Bucheon, Ansan,
 306 Anyang, and Suwon. GeoTASO observed NO₂ VCD values three-times higher ($>3 \times 10^{16}$ molecules cm⁻²) in these areas
 307 compared to the surrounding rural areas. High NO₂ VCD values above 6×10^{16} molecules cm⁻² were observed above the
 308 Gyeongin Expressway, which has very heavy traffic in a relatively short section, and the Gunpo Complex Logistics zone,
 309 where diesel vehicle traffic is also high. The main NO₂ source regions and the regions where high NO₂ VCD values were
 310 observed were highly consistent at 9 AM because the wind speed at this time—as obtained from the unified model (UM) based
 311 Regional Data Assimilation and Prediction System (RDAPS) of the Korea Meteorological Administration (KMA)—was as
 312 low as 0.1 ms⁻¹ and the average wind direction was 84.7° at 1000 hPa over Seoul metropolitan region. The average daily traffic
 313 volume of these expressways exceeds 150,000 vehicles, and the total number of vehicles registered in these major cities is >
 314 6,000,000, with an average daily mileage per car per day of over 38 km. Detailed information on these cities and expressways
 315 is listed in Table 3 and 4. Based on the level of vehicular traffic, combustion using gasoline and diesel engines leads to high
 316 overall emissions of NO₂ in the Seoul metropolitan region (Kendrick et al., 2015).

317

318 **Table 3. The population, number of registered vehicles, and average mileage per car per day of the major cities in the Seoul**
 319 **and Busan metropolitan region obtained from the Korean Statistical Information Service (<https://kosis.kr/eng>).**

City	Population (millions)	Vehicle registration number (thousands)	Average mileage (km car ⁻¹ day ⁻¹)
Seoul	9.776	3,083	37.1
Incheon	2.914	1,402	41.7
Bucheon	0.848	284	37.2
Ansan	0.744	289	40.8
Anyang	0.596	206	39.6
Gunpo	0.286	87	38.8
Suwon	1.241	467	38.1
Sungnam	0.994	358	36.3
Busan	3.389	1,295	40.1
Daegu	2.450	1,121	37.1
Changwon	1.080	551	37.5
Kimhae	0.529	250	38.0

320

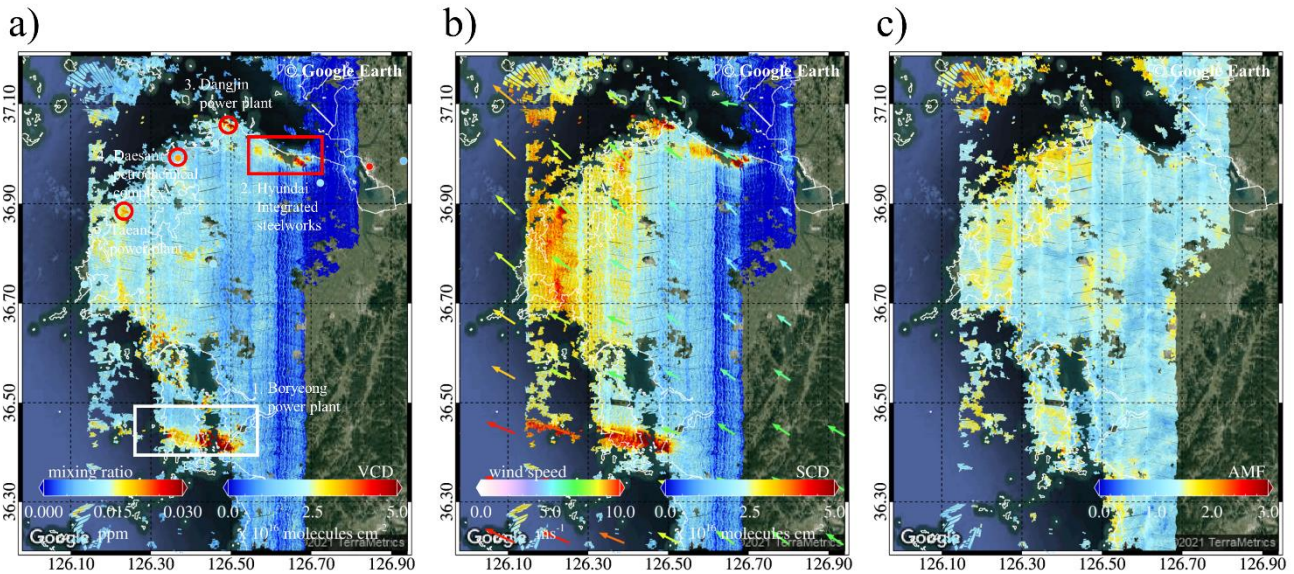
321 **Table 4. Daily average traffic volume on the Gyeongin, Gyeongbu, and Seohaean Expressways obtained using the Traffic**
 322 **Monitoring System (<https://www.road.re.kr>).**

Expressway	Daily average traffic volume
Gyeongin Expressway	162,369
Gyeongbu Expressway	173,413
Seohaean Expressway	150,298

323

324 Compared to the data of the morning, the average wind speed and wind direction were 1.7 ms^{-1} and 284.5° at 1000 hPa in the
 325 afternoon and the afternoon had extremely high tropospheric NO_2 VCD values (exceeding 5×10^{16} molecules cm^{-2}) in most of
 326 the Seoul metropolitan regions including rural areas, whereas the NO_2 mixing ratio (MR) obtained from Air-Korea decreases
 327 in the afternoon. According to Tzortziou et al. (2018), similar results were retrieved from the Pandora site in Seoul, with higher
 328 afternoon NO_2 VCDs than in the morning. This result is because the amount of NO_2 produced by chemical conversion of nitric
 329 oxide (NO) by O_3 and VOCs in the atmosphere, along with NO_x generated by regional emissions (traffic) in the Seoul
 330 metropolitan region, is greater than the amount lost by photolysis and transport to nearby areas (Herman et al., 2018).
 331 Furthermore, the increase in tropospheric NO_2 VCD in the afternoon is likely due to the accumulation and dispersion of NO_2
 332 according to the height of the change in the planetary boundary layer (Ma et al., 2013).

333 3.1.2 Industrial and power plant regions in Anmyeon



334
 335 **Figure 6. a) Tropospheric NO_2 VCD and b) NO_2 SCD retrieved from GeoTASO, and c) NO_2 AMF, native resolution (250 m)**
 336 **calculated using VLIDORT over Anmyeon in South Korea on 5 June 2016. The colored arrows indicate wind speed and wind**
 337 **direction at 850 hPa from the Unified Model (UM) simulations. The red circles and rectangle in panel (a) represent the major NO_2**
 338 **emission sources, such as steelworks and power plants (background RGB image is from Google Earth;**
 339 **<https://www.google.com/maps/>).**

340
 341 The high spatial resolution of the tropospheric NO_2 VCD from GeoTASO over the Anmyeon industrial region, where many
 342 industrial facilities and several power plants are distributed, is shown in Fig. 6. Panels a and b of this figure indicate the binned
 343 tropospheric NO_2 VCD and NO_2 SCD retrieved from GeoTASO L1B data, respectively, between 13:00 and 17:00 LT on June
 344 5 2016. Panel c depicts the calculated AMF of NO_2 from native resolution over the domain. GeoTASO observations detected
 345 moderate and strong NO_2 emission sources in this area: (1) Boryeong power plant, (2) Hyundai integrated steelworks, (3)
 346 Dangjin power plant, (4) Daesan Petrochemical Complex, and (5) Taean Power Plant. High NO_2 VCD values ($> 5 \times 10^{16}$
 347 molecules cm^{-2}) were observed over steel mill works, petrochemical complexes, and power plants, whereas values were
 348 comparatively low ($< 1 \times 10^{16}$ molecules cm^{-2}) over small cities including Seosan, Dangjin, and Boryeong with populations of
 349 less than 0.1 million, and the Sohaean Expressway. In 2016, the annual NO_x emissions from Hyundai steelworks and the
 350 Dangjin and Boryeong power plants were approximately 10.3, 11.9, and 16.8 kt year^{-1} , respectively. The NO_x emission rates
 351 of major industrial facilities in the Anmyeon region are shown in Table 5.

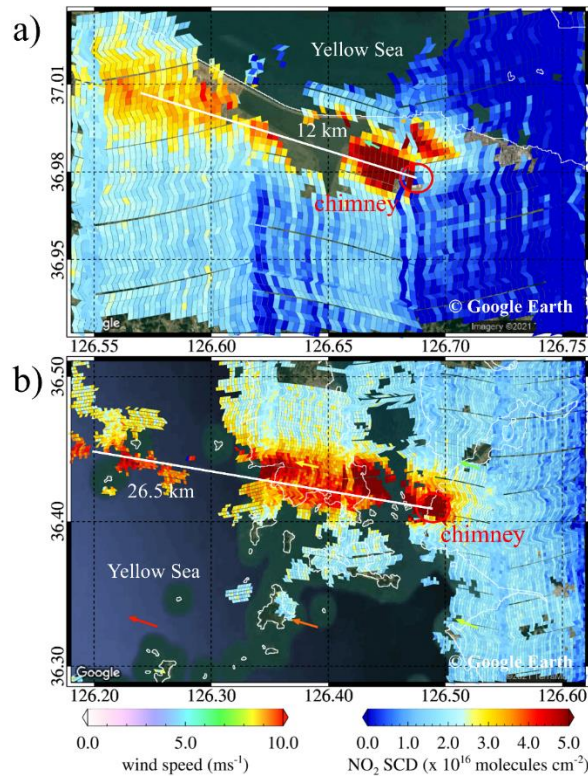
352

353 Table 5. NOx emission rates in 2016 from major industrial facilities in the Anmyeon region obtained from the Continuous
354 Emission Monitoring System of the Korea Environment Corporation (<https://www.stacknsky.or.kr/eng/index.html>).

Industrial facilities	NOx emission rate (kg year ⁻¹)
Boryeong power plant	16,788,438
Hyundai integrated steelworks	10,271,075
Dangjin power plant	11,852,972
Daesan petrochemical complex	3,397,939
Taeon power plant	15,466,022

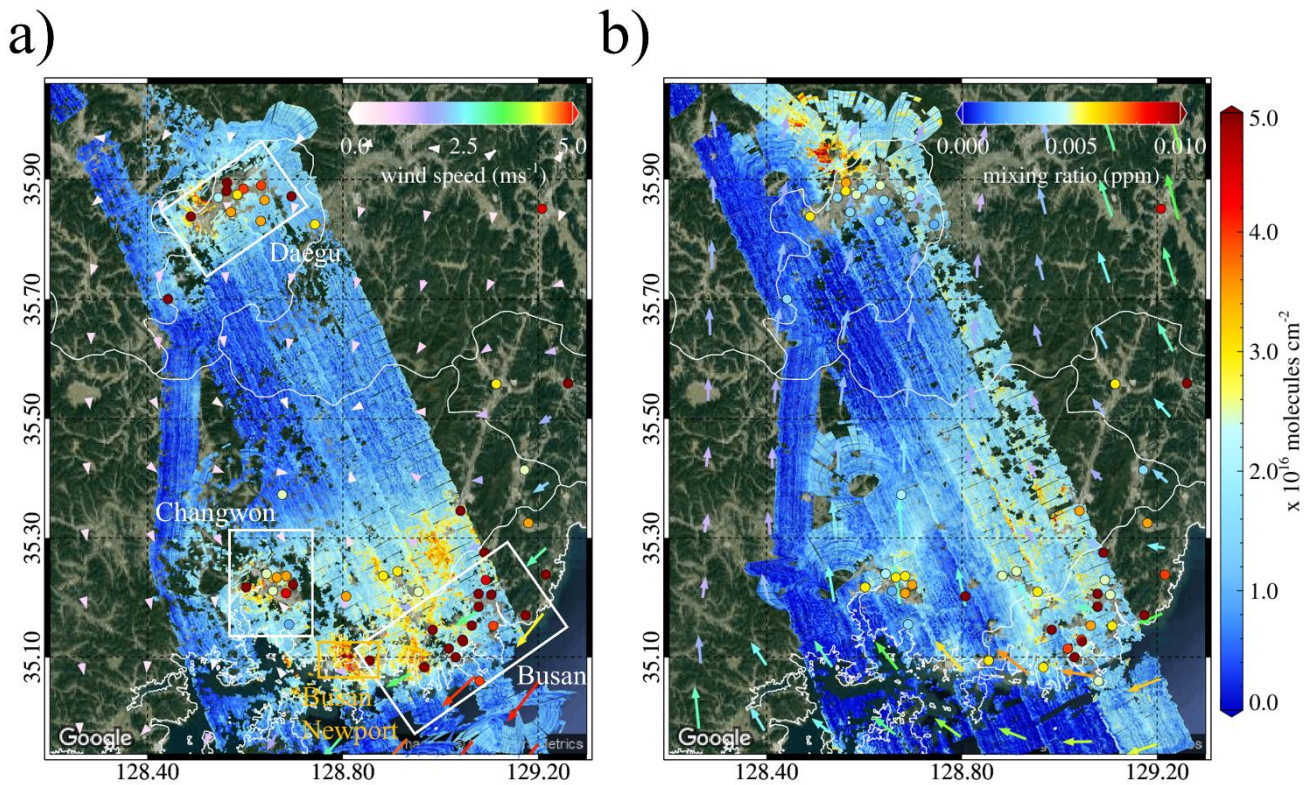
355

356 Figure 6 shows high NO₂ concentrations of the main industrial facilities in the Anmyeon region, where the combustion of
357 fossil fuel in factories and thermal power plants leads to high emissions (Prasad et al., 2012). Due to relatively sparse
358 distribution over rural areas, the Air-Korea measurements did not detect the major NO₂ plume as shown in Fig. 6a. Thus,
359 airborne remote sensing systems, such as GeoTASO, can effectively complement ground-based networks for monitoring minor
360 and major NO_x emissions, particularly over these remote industrial regions.



361
362 **Figure 7. Enlarged view of GeoTASO tropospheric NO₂ SCD observation over a) Hyundai steel works, indicated by the red box in**
363 **Figure 6, and b) the Boryeong power plant, indicated by the white box in Figure 6. The arrows represent the wind direction and**
364 **speed at 850 hPa from the Unified Model (UM) simulations, respectively (background RGB image is from Google Earth;**
365 **<https://www.google.com/maps/>).**

366
367 The GeoTASO data captured not only NO_x emissions from the chimneys of steelworks and power plants but also its transport
368 by the wind. Figure 7a and 7b show enlarged views of tropospheric NO₂ SCD retrieved using GeoTASO over the Hyundai
369 steelworks (red box in Fig. 6) and the Boryeong power plant (white box in Fig. 6). The arrows in Fig. 7 represent the prevailing
370 wind direction and speed from RDAPS. NO₂ emitted from the chimneys of these sites was transported to the Yellow Sea,
371 traveling distances of over 26.5 km at speeds of approximately 6 ms⁻¹. According to Chong et al. (2020), similar results were
372 found for SO₂ emitted and transported from these sites.



374

375 **Figure 8. Tropospheric NO₂ VCD in the Busan metropolitan region in the (a) morning and (b) afternoon of June 10, 2016.**
 376 **The wind speed (colors scale) and wind direction (arrows) at 1000 hPa pressure level were obtained from the Unified Model (UM)**
 377 **simulations. The white boxes represent major cities such as Busan, Daegu, and Changwon. The orange box represents Busan**
 378 **Newport (the background RGB image is from Google Earth; <https://www.google.com/maps/>).**

379

380 Figure 8a and 8b show tropospheric NO₂ VCD retrieved from the GeoTASO L1B data over the Busan metropolitan region on
 381 June 10 2016 in the morning (between 08:00 and 11:00 LT) and afternoon (between 13:00 and 16:00 LT), respectively. The
 382 arrows in Fig. 8 indicate the wind speed and wind direction of 1000 hPa obtained from the UM-RDAPS, with the average
 383 wind speed and wind direction of 0.9 ms⁻¹, and 55.4°, 1.9 ms⁻¹ and 147.0°, respectively, in the morning and afternoon. High
 384 NO₂ VCDs were observed above urban areas, port, industrial complexes, and the inter-city road between Busan and Changwon.
 385 Like the Seoul metropolitan regions, combustion using gasoline and diesel engines is estimated to contribute to the high NO_x
 386 emission. In the morning, NO₂ VCDs were high (approximately 3 × 10¹⁶ molecules cm⁻²) in the major cities and, especially,
 387 around Busan Newport, with values exceeding 7 × 10¹⁶ molecules cm⁻². In comparison, in the mountainous regions between
 388 Daegu and Busan, the NO₂ VCD values were less than 1 × 10¹⁶ molecules cm⁻² during the same period. The spatial distribution
 389 of tropospheric NO₂ VCDs was like that in the Seoul metropolitan regions, with high values over major cities and roads
 390 (compare Figs. 5 and 8). In Busan, fossil fuel combustion that uses both road vehicles and ships is likely to contribute to the
 391 NO_x emissions. In the afternoon, unlike the Seoul metropolitan region, tropospheric NO₂ VCD over Busan decreased by over
 392 3 × 10¹⁶ molecules cm⁻², which also corresponds with NO₂ MR data obtained from the Air-Korea sites. Detailed information
 393 on these cities is listed in Table 3.

394

395 3.2 Error estimation

396 The accuracy of the NO₂ VCD retrieval using the DOAS method depends on both the AMF calculation and the spectral fitting
 397 error of the SCD retrieval. Retrieval errors of the NO₂ VCD were estimated using error propagation analysis as expressed in
 398 Eq. (12).

$$399 \frac{\varepsilon_{VCD}}{VCD} = \sqrt{\left(\frac{\varepsilon_{SCD}}{SCD}\right)^2 + \left(\frac{\varepsilon_{AMF}}{AMF}\right)^2}, \quad (12)$$

400 Where ε_{VCD} is the total error of NO₂ VCD. The error of NO₂ SCD (ε_{SCD}) is obtained from the spectral fitting error of NO₂ SCD
 401 via the DOAS spectral fitting. ε_{AMF} indicates the error of NO₂ AMF caused by uncertainties in the model input parameters for
 402 AMF calculation. Uncertainties in aerosol properties (AOD, SSA, and APH) and surface reflectance for the RTM calculations
 403 are the major factors affecting NO₂ AMF accuracy (Boersma et al., 2004; Leitão et al., 2010; Hong et al., 2017). Therefore, in
 404 this present study, we quantified the NO₂ AMF errors (ε_{AMF}) due to uncertainties in the input parameters independent of each
 405 other using Eq. (13):

$$406 \varepsilon_{AMF} = \sqrt{\left(\frac{\partial AMF}{\partial AOD}\right)^2 \sigma AOD^2 + \left(\frac{\partial AMF}{\partial SSA}\right)^2 \sigma SSA^2 + \left(\frac{\partial AMF}{\partial ALH}\right)^2 \sigma ALH^2 + \left(\frac{\partial AMF}{\partial SFR}\right)^2 \sigma SFR^2} = \sqrt{\sum_{i=1}^4 \left(\frac{\partial AMF}{\partial \chi_i}\right)^2 \sigma_{\chi_i}^2}, \quad (13)$$

407 where $\frac{\partial AMF}{\partial \chi_i}$ are partial derivatives of NO₂ AMF regarding the input parameters (χ_i), σ_{χ_i} represents the uncertainty of the χ_i . The
 408 σ of AOD, SSA, surface reflectance, and APH are assumed to be 30% (Ahn et al., 2014), 0.04 (Jethva et al., 2014), 0.005 +
 409 0.05 × surface reflectance (EOS Land Validation; <https://landval.gsfc.nasa.gov>), and 1 km (Fishman et al., 2012), respectively,
 410 in this study. To derive $\left(\frac{\partial AMF}{\partial \chi_i}\right)^2$, the true χ_i is input to the RTM to simulate ‘true’ NO₂ AMF. For the AOD, SSA, APH, and
 411 surface reflectance (SFR), perturbed NO₂ AMF was simulated using RTM with $\chi_i \pm \sigma_{\chi_i}$. $\partial \chi_i$ denotes the difference between the
 412 ‘centre’ χ_i and $\chi_i \pm \sigma_{\chi_i}$, and ∂AMF is the difference between the ‘centre’ NO₂ AMF (AMF_{centre}) simulated with ‘centre’ input
 413 values and the perturbed NO₂ AMF ($AMF_{perturbed}$) simulated using the perturbed input parameters $\chi_i \pm \sigma_{\chi_i}$ (i.e. the original input
 414 parameters modified by the uncertainty). The simulation for calculating the ε_{AMF} was conducted using the input parameters
 415 on 9 June 2016.

416

417 **Table 6. Total NO₂ VCD caused by uncertainties in NO₂ SCD and NO₂ AMF (the average for the flight on June 9, 2016).**

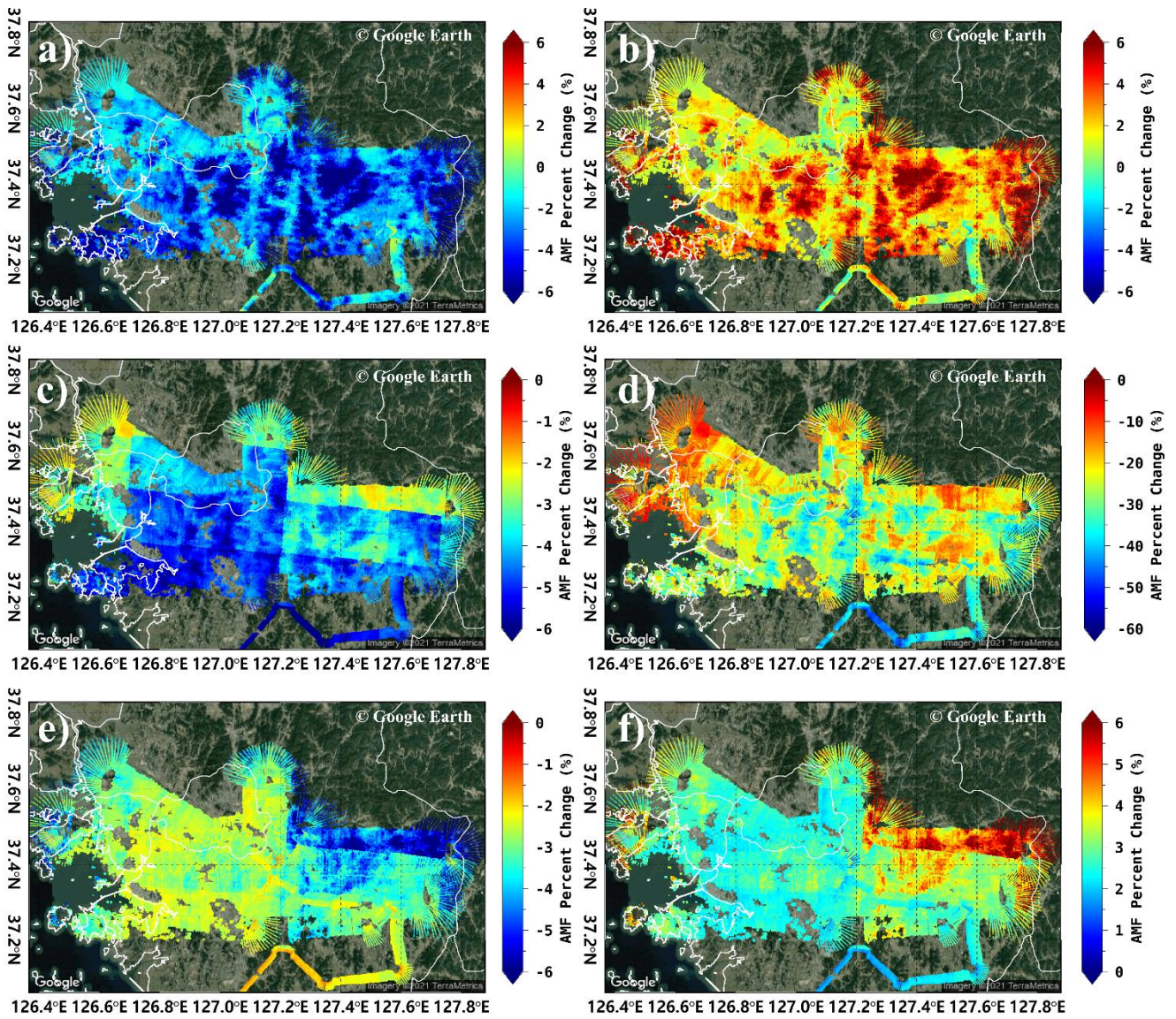
	AOD	2.8%
	SSA	4.1%
NO ₂ AMF errors	Aerosol peak height	22.3%
	Surface reflectance	2.8%
	Total NO₂ AMF error due to aerosol uncertainties	23.3%
	NO₂ SCD error	11.7%
	NO₂ VCD error	26.9%

418

419 Table 6 lists the estimated NO₂ VCD error on June 9 2016 for each source based on the error propagation method. The error
 420 estimation was conducted for the pixels where root mean square residual < 0.001 and NO₂ VCD > 5 × 10¹⁵ molecules cm⁻²
 421 since NO₂ SCD precision is reported to be highly decreased in low NO₂ conditions (Hong et al., 2017). The total NO₂ VCD
 422 error was 26.9% with a high portion of NO₂ AMF error. The NO₂ SCD error was calculated to be 11.7%, showing the

423 importance of accurate DOAS spectral fitting for deriving NO₂ SCD. The total AMF error due to uncertainties in the input
 424 parameters was calculated to be 23.3%. Among model input parameters, the effect of APH on NO₂ AMF becomes high (22.3%),
 425 indicating the importance of accurate aerosol profile information. APH sensitively affects NO₂ AMF because near the surface
 426 where trace gases and aerosols are well mixed, aerosols lead to multiple scattering effects and the light absorption of trace
 427 gases is due to increasing light path (Castellanos et al., 2015; Hong et al., 2017). Especially, APH can be the most important
 428 input parameter in the Asia region where high loadings of aerosol plumes persist throughout the year. The NO₂ AMF
 429 calculation errors due to uncertainties in SSA and AOD were 4.1% and 2.8%, respectively. The NO₂ AMF calculation error
 430 due to uncertainties in aerosol optical properties (SSA and AOD) appears to be smaller than those in a previous study (Leitão
 431 et al., 2010). The smaller effect of the aerosol properties can be explained by the moderate aerosol loading (AOD = 0.40) on
 432 the day of flight day. The NO₂ AMF errors become larger under high AOD conditions. The smallest effect of SRF was found
 433 on NO₂ AMF calculation error, which was calculated based on the uncertainty of the SRF of the satellite-based product
 434 (MODIS). Therefore, it may be an unrealistic number for the airborne NO₂ AMF calculation. Once the uncertainty of airborne-
 435 based SRF is provided, considering its measurement geometry and finer spatial resolution, more realistic airborne-based NO₂
 436 AMF calculation error due to uncertainties in SRF can be estimated. The can of the *a priori* NO₂ profile shape also be a factor
 437 to cause calculation error for NO₂ AMF, as reported in previous studies (Leitão et al., 2010, Meier et al., 2016, Hong et al.,
 438 2017). Therefore, it is necessary to calculate the contribution of the shape of the NO₂ profile *a priori* on the accuracy of NO₂
 439 AMF in the future. Moreover, the resulting uncertainties of input parameters of a GeoTASO ground pixel need to be considered
 440 by combining the initial uncertainties of CTM and satellite-based products, and by the variability of the parameters within the
 441 respective CTM (AOD, SSA, and APH) and satellite (SFR) grid box. If values such as SFR are assumed constant over larger
 442 areas, the fundamental spatial variability in this these data increases the uncertainty of the AMF and hence of the determined
 443 NO₂ VCD on the respective finer spatial scale. In addition, the uncertainty from the assumption on the SC₀ and the uncertainty
 444 from ignoring the NO₂ above the aircraft in the AMF calculations are needed to be considered in the error analysis. This
 445 analysis should be considered in further study.

$$446 \text{AMF}_{\text{percent_change}} = \frac{\text{AMF}_{\text{perturbed}} - \text{AMF}_{\text{centre}}}{\text{AMF}_{\text{centre}}} \times 100, \quad (14)$$



447

448 **Figure 9.** Percent change between AMF calculated using the CMAQ model simulation and those using a) 30% lower AOD,
 449 b) 30% higher AOD, c) 0.04 lower SSA, d) 1km higher APH, compared to the model outputs. The percentage change for AMF
 450 calculated using MODIS data and those using e) $0.005 + 0.05 \times \text{SFR}$ lower SFR, f) $0.005 + 0.05 \times \text{SFR}$ higher SFR (background
 451 RGB image is from Google Earth; <https://www.google.com/maps/>).

452

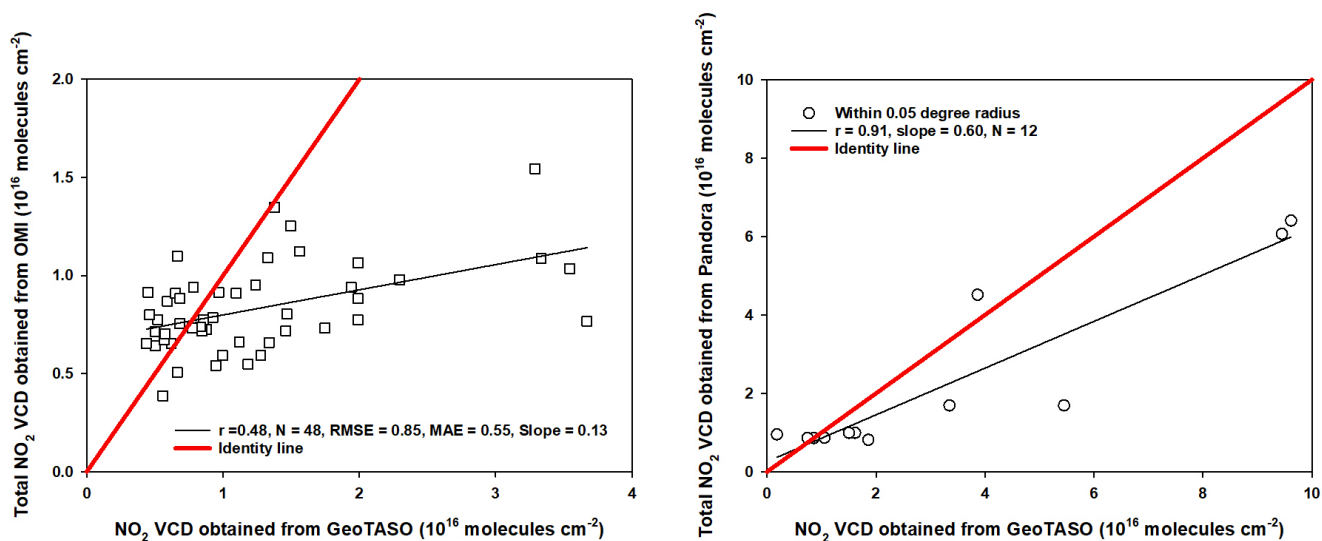
453 In this study, we also investigated the spatial distribution of AMF calculation errors associated with uncertainties in aerosol
 454 properties (AOD, SSA, and APH) and SFR. The percent change in NO_2 AMF ($\text{AMF}_{\text{percent_change}}$) was calculated on each spatial
 455 pixel using Eq. (14). Figure 9a and 9b indicate the percentage change error between the calculated AMFs using the CMAQ
 456 AOD data with 30% lower (Fig. 9a) and 30% higher (Fig. 9b) values, respectively. The AMF decreased and increased by up
 457 to 10% with decreasing and increasing AOD, respectively, in the Seoul metropolitan region. We estimated that, under low
 458 aerosol loading conditions, an increase in AOD near the surface leads to an increase in the scattering probability within the
 459 surface layer with high NO_2 concentrations. Figure 9c indicates the percent change error between the calculated AMFs using
 460 CMAQ SSA data with a 0.04 lower value. The AMF decreased with decreasing SSA because the absorption of light increased.
 461 APH was also found to highly affect the accuracy of the AMF calculations (Fig. 9d). The APH uncertainty of 1 km decreased
 462 the AMFs with an average $\text{AMF}_{\text{percent_change}}$ of -25% on the flight day. Especially, on the pixels where $\text{AOD} > 0.6$, the average
 463 $\text{AMF}_{\text{percent_change}}$ was found to be -26% whereas that was -27% on the pixels where $\text{AOD} < 0.4$, showing the combined effect
 464 of aerosol loading and aerosol profile shape on the NO_2 AMF calculations. Figure 9e and 9f indicate the percentage change

465 error between the calculated AMFs using the MODIS surface reflectance data with $0.005 + 0.05 \times \text{SFR}$ lower (Fig. 9e) and
 466 $0.005 + 0.05 \times \text{SFR}$ higher (Fig. 9f) values, respectively. The AMF decreased by approximately 3% when the SFR decreases,
 467 and vice versa when it increased.

468 3.3 Validation of NO₂ VCDs retrieved from GeoTASO

469 The tropospheric NO₂ VCDs retrieved from GeoTASO L1B data (NO_{2,G}) were compared with those obtained from OMI total
 470 NO₂ VCDs (NO_{2,O}) and Pandora (NO_{2,P}). The NO_{2,O} were only available for June 10 during the campaign period. Therefore,
 471 we compared only 48 NO_{2,G} and NO_{2,O} data points within a radius of 20 km and 30 min, which yielded a correlation coefficient
 472 of 0.48 with a slope of 0.13 (Fig. 10 a). To validate, All NO_{2,G} within a radius 20 km of the OMI center coordinate were
 473 averaged.

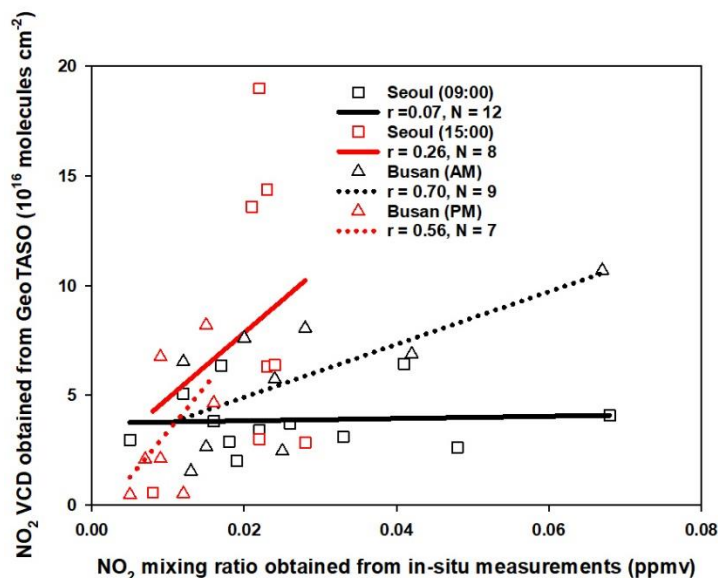
474 The NO₂ values are relatively low, as GeoTASO observation is conducted in a region with low NO₂ compared to the Seoul
 475 metropolitan and the overpass time of OMI is approximately 13:30 LT when NO₂ decreased. The low slope value is because
 476 the OMI with low spatial resolution does not reflect the spatial NO₂ inhomogeneity in the pixel.



477
 478 **Figure 10. Scatter plots of a) NO₂ VCD retrieved from GeoTASO and total NO₂ VCD obtained from OMI and b) total NO₂ VCD**
 479 **obtained from Pandora and NO₂ VCD retrieved from GeoTASO, respectively.**

480
 481 To compare NO_{2,G} data, we made a comparison with total NO₂ VCD obtained from the Pandora system (NO_{2,P}) during the
 482 KORUS-AQ campaign period. NO_{2,P} obtained from Busan University, Olympic Park, Songchon, Yeosu, and Yonsei University
 483 Pandora sites on June 5, 9, and 10 were used for the GeoTASO validation (Fig. 1). NO_{2,G} and NO_{2,P} columns at these sites are
 484 compared in Fig. 11. To compare NO_{2,G} and NO_{2,P}, we used averaged NO_{2,G} retrieved from 16 across tracks with the smallest
 485 viewing zenith angle and averaged 30 min NO₂ obtained from pandora measurement within a radius of approximately 0.05°.
 486 NO_{2,G} and NO_{2,P} were correlated (R = 0.91, with a slope of 0.60), however, when NO_{2,P} was lower than 1×10^{16} molecules
 487 cm⁻², the correlation coefficient between NO_{2,G} and NO_{2,P} was < 0.1. The weak correlation at low NO₂ levels most likely
 488 reflects differences in viewing geometries and the horizontal inhomogeneity of the measured NO₂ between Pandora and
 489 GeoTASO. Furthermore, Pandora and GeoTASO can be used for the NO₂ validation of geostationary satellites, such as GEMS.
 490 However, because the number of pandora is limited in this campaign, we difficulty validating NO₂ retrieved from GeoTASO
 491 under various conditions. Many ground-based remote sensing measurements are needed to validate GEMS under various
 492 conditions.

493
 494
 495



497

498 **Figure 11.** Scatter plot of the NO₂ VCDs retrieved from GeoTASO, and NO₂ surface mixing ratio obtained from Air-Korea.
 499 The black and red squares represent the NO₂ data at 9 AM and 3 PM (local time) in the Seoul metropolitan region,
 500 respectively. The black and red triangles represent those in the morning and afternoon, over Busan, respectively.

501

502 To compare the spatiotemporal distribution of NO₂ VCDs retrieved from GeoTASO, NO_{2,G} compared with surface spatial
 503 patterns, NO_{2,G} was compared with NO_{2,A} for GeoTASO data within a radius of approximately 0.05 km and 30 min (Fig. 11).

504 To compare NO_{2,G} and NO_{2,A}, we used averaged NO_{2,G} retrieved from 16 across tracks and averaged 30 min within a radius of
 505 0.05°. Because in situ measurements provide NO₂ VMR (NO_{2,A})(ppmv) once per hour, NO_{2,A} of the nearest time is used to
 506 compare with NO_{2,G}. The correlation coefficient (R) between NO_{2,G} (molecules cm⁻²) and NO_{2,A} at 9 AM and 3 PM LT in the
 507 Seoul metropolitan region was 0.07 and 0.26, respectively. When using only roadside station data from Air-Korea, the R-value
 508 for the morning increased to 0.72, which implies GeoTASO is more sensitive to emissions from NO₂ source areas, such as
 509 roadsides (Fig. 5). Because the comparison, there were large differences in the morning and afternoon. These results were
 510 identified because synoptic meteorology played an important role from June 1 to June 10, 2016 (Choi et al., 2019). As described
 511 by Judd et al. (2018), the spatial distribution for NO₂ VCDs appears to reflect the emission source in local industrialized regions
 512 and transportations in the morning with relatively weak winds. NO₂ concentration often increases in the late morning,
 513 indicating that the emission process proceeds faster than the NO₂ removal process. As the planetary boundary layer heights
 514 (PBLH) in early afternoon increase and surface NO₂ is mixed through a deeper PBLH, the NO₂ VCDs distribution showed a
 515 wider increase in most of the Seoul metropolitan area and the column amounts continue to increase (Judd et al., 2018).

516 When comparing NO₂ VCDs with surface NO₂ concentrations, it should be highlighted that it is a nonlinear relationship
 517 between NO_{2,G} and NO_{2,A}. Although it may vary depending on weather conditions, high NO₂ VCDs from airborne observations
 518 can sometimes be detected with low surface NO₂ concentrations. When exhaust gases emitted from industrial facilities occur
 519 at a certain altitude (stacks/chimneys), NO_{2,G} show high NO₂ VCDs, but NO_{2,A} may be observed to have a low concentration.
 520 Unfortunately, in the Anmyeon industrial region, NO_{2,G} and NO_{2,A} could not be compared due to spatial restrictions because
 521 the distribution of ground observation stations is concentrated in metropolitan areas.

522 In the Busan metropolitan area, the R-value of the NO_{2,G} and NO_{2,A} data had a correlation coefficient greater than 0.56. This
 523 reflects the more even horizontal distribution of NO₂ in the afternoon, when diffusion from the source areas had occurred.
 524 However, for a more accurate comparison, NO₂ VCD data should be converted to NO₂ MR based on mixing layer height,
 525 temperature, and pressure profile data (Kim et al., 2017; Qin et al., 2017; Jeong and Hong, 2021a). However, because the
 526 number of pandora and satellite data is limited in this campaign, we had difficulties in validating NO₂ retrieved from GeoTASO

527 under various conditions. Because ground-based, airborne and space borne remote sensing measurements have their own
528 advantages and disadvantages, it is recommended a comprehensive observation campaign involving all of ground-based,
529 airborne and space-borne measurements should be conducted continuously for the upcoming new era of geostationary
530 environmental satellites.

531 **4 Conclusions**

532 For the first time, we have retrieved NO₂ VCD data using airborne GeoTASO observations over the Seoul metropolitan
533 region—one of the most populous cities worldwide, the Busan metropolitan region—the second-largest city in South Korea,
534 and Anmyeon, with thermal power plants and industrial complexes. By retrieving NO₂ data using GeoTASO L1B radiance, it
535 was possible to observe the spatial distribution of NO₂ in these metropolitan and industrial regions. In the morning,
536 tropospheric NO₂ VCD in Seoul showed a strong horizontal gradient between rural and urban areas. In urban areas,
537 tropospheric NO₂ VCD was high, with values exceeding 3×10^{16} molecules cm⁻²; in rural areas, values were typically below
538 1×10^{16} molecules cm⁻². Extremely high values over 10×10^{16} molecules cm⁻² were also observed in both rural and urban
539 areas. In Anmyeon, GeoTASO observations showed that NO₂ is mainly emitted from the chimneys of industrial complexes
540 and thermal power plants, and subsequently transported by wind approximately 30 km to the Yellow Sea on the west coast of
541 the Korean Peninsula. In the Busan metropolitan region, in the morning, tropospheric NO₂ VCDs showed a pattern similar to
542 the Seoul metropolitan region, with high values above the inter-city road. However, unlike Seoul, tropospheric NO₂ VCDs in
543 Busan decreased in the afternoon due to local different weather conditions locally.

544 To compare the data retrieved from the GeoTASO system, we compared NO_{2,G} with NO_{2,O} obtained from the OMI, NO_{2,A}
545 obtained from Air-Korea, and NO_{2,P} obtained from the Pandora observation system. When the distance between two
546 observations was below 20 km or 0.05° within 30 min, the correlation coefficients were relatively high (R = 0.48, and 0.91,
547 respectively). However, the correlation between NO_{2,G} and NO_{2,A} over the Seoul metropolitan region was extremely weak (R
548 = 0.07) in the morning because of the more pronounced NO₂ horizontal gradient.

549 The GeoTASO system successfully observed NO₂ VCDs with high horizontal spatial resolution for both metropolitan and
550 industrial regions. This demonstrates that airborne remote sensing measurements from GeoTASO, similar to GCAS, APEX
551 and others, can be an effective tool for the validation of trace gases retrieved from environmental satellites, including the OMI,
552 TROPOMI, and GOME-2; these systems can obtain high-resolution measurements over relatively wide areas. However, to
553 validate geostationary environmental satellites with higher spatiotemporal resolutions, such as the GEMS, TEMPO, and
554 sentinel-4, additional validation strategies are needed. Based on error estimation, it can be concluded that aerosol properties
555 are relevant and should be determined and NO₂ vertical profile retrieval performed using, for example, LIDAR, MAX-DOAS,
556 and sondes. This is important because the accuracy of aerosol properties, surface reflectance and the NO₂ vertical profiles
557 affects the accuracy of AMF calculations (Leitão et al., 2010; Hong et al., 2017; Lorente et al., 2017; Boersma et al., 2018).
558 Furthermore, as we observed in the Seoul metropolitan area, closer spaced observations using ground-based remote sensing
559 systems and in situ measurements are needed as NO₂ displays large horizontal gradients, especially in the morning.

560 **Author contributions**

561 **GH** and **HH** designed and implemented the research. **KL** provided the CTM data. **GH** developed the code for model running
562 and performed the RTM simulations. **HH** and **UJ** contributed to the analysis of ground-based data. **GH** and **WC** conducted
563 the sensitivity test. **GH**, **KL**, **HH**, **UJ**, **WC**, and **JJS** revised and edited the paper. **HH**, **UJ**, and **WC** provided constructive
564 comments. All authors contributed to this works.

565 **Competing interests**

566 The authors declare that they have no conflict of interest.

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