

# 1 **Highly resolved mapping of NO<sub>2</sub> 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 (NO<sub>2</sub>) slant column density (SCD) measurements were retrieved from  
16 Geostationary Trace and Aerosol Sensor Optimization (GeoTASO) L1B data during the KORUS-AQ campaign (May 1 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 NO<sub>2</sub>  
21 over the Seoul and Busan metropolitan regions, and the industrial regions of Anmyeon. We reveal that the maximum NO<sub>2</sub>  
22 VCDs were  $4.94 \times 10^{16}$  and  $1.46 \times 10^{17}$  molecules cm<sup>-2</sup> at 9 AM and 3 PM over Seoul, respectively,  $6.86 \times 10^{16}$  and  $4.89 \times$   
23  $10^{16}$  molecules cm<sup>-2</sup> in the morning and afternoon over Busan, respectively, and  $1.64 \times 10^{16}$  molecules cm<sup>-2</sup> over Anmyeon.  
24 The VCDs retrieved from the GeoTASO airborne instrument were well correlated with those obtained from the Ozone  
25 Monitoring Instrument (OMI) ( $r = 0.70$ ), NASA's Pandora Spectrometer System ( $r = 0.79$ ), and NO<sub>2</sub> mixing ratios obtained  
26 from in situ measurements ( $r = 0.45$  in the morning,  $r = 0.81$  in the afternoon over the Seoul, and  $r > 0.78$  over Busan). Based  
27 on our results, GeoTASO is useful for identifying hotspots of NO<sub>2</sub> and its spatial distribution in highly populated cities and  
28 industrial areas.

## 29 1 Introduction

30 Nitrogen dioxide (NO<sub>2</sub>) is one of the most important atmospheric trace gases and plays a key role in aerosol production and  
31 tropospheric ozone photochemistry (Boersma et al., 2004; Richter et al., 2005). Furthermore, high NO<sub>2</sub> concentrations in the  
32 atmosphere have adverse effects on human health, such as respiratory infections, domestic heating, and associated symptoms  
33 (Brauer et al., 2002; Latza et al., 2009).

34 The major sources of NO<sub>2</sub> in the atmosphere are from fossil fuel combustion from vehicles and thermal power plants, lightning,  
35 flash production, and biogenic soil processes. In addition, it has been found that NO<sub>2</sub> concentrations are highly correlated with  
36 population size (Lamsal et al., 2013). The implementation of emission control technology and environmental regulation has  
37 led to a decrease in surface NO<sub>2</sub> concentrations in Western Europe, the United States, and Japan in the last few decades (Richter  
38 et al., 2005). The NO<sub>2</sub> concentration over major metropolitan cities in South Korea and China are over 3 times larger than over  
39 similarly sized cities in Europe and United States, despite NO<sub>2</sub> concentration decreasing in China and South Korea (de Foy et  
40 al., 2016, Choo et al., 2020).

41 To date, several low-orbit space borne sensors, such as the Global Ozone Monitoring Experiment (GOME) (Burrows et al.,  
42 1999), the Scanning Imaging Spectrometer for Atmospheric Cartography (SCIAMACHY) (Burrows et al., 1995), the Ozone  
43 Monitoring Instrument (OMI) (Levelt et al., 2006), the GOME-2 (Callies et al., 2000), and the Tropospheric Monitoring  
44 Instrument (TROPOMI) (Veefkind et al., 2012), have monitored atmospheric ozone and its precursors including NO<sub>2</sub> and  
45 formaldehyde (HCHO) as a proxy for volatile organic compounds (VOCs). Furthermore, the Geostationary Environment  
46 Monitoring Spectrometer (GEMS) (Choi et al., 2018; Kim et al., 2020), which was launched on February 18, 2020, will form  
47 a constellation of geostationary satellites including the upcoming Tropospheric Emission: Monitoring of Pollution (TEMPO)  
48 (Zoogman et al., 2017) and Sentinel-4 platforms, to continuously observe the air quality of the Northern Hemisphere during  
49 the daytime.

50 NO<sub>2</sub> retrievals from space borne hyperspectral measurements are typically conducted using the differential optical absorption  
51 spectroscopy (DOAS) method (Platt and Stutz, 2008) to first retrieve the view-dependent slant column density (SCD), and  
52 then radiative transfer models are used to determine the vertical column density (VCD) using an air mass factor (AMF)  
53 correction. Previous and ongoing space borne instruments use various radiative transfer codes and model input assumptions to  
54 calculate NO<sub>2</sub> AMF values at fairly coarse spatial resolution. Because the AMF weighting has a large impact on NO<sub>2</sub> retrievals  
55 using the DOAS method, it is important to use model input assumptions that most accurately match the viewing and  
56 atmospheric conditions. Several studies have demonstrated the sensitivity of AMF calculations to inaccurate model input  
57 parameters (e.g., *a priori* NO<sub>2</sub> vertical profile and aerosol properties) and *a priori* data (cloud information and surface  
58 reflectance) (Leitão et al., 2010; Hong et al., 2017; Lorente et al., 2017; Boersma et al., 2018). NO<sub>2</sub> retrievals have also been  
59 consistently conducted based on surface remote sensing measurements including the Multi-Axis DOAS (MAX-DOAS),  
60 Système D'Analyse par Observations Zénithales (SAOZ) spectrometer (Pastel et al., 2014), and Pandora (Herman et al., 2009)

61 systems. These ground-based measurements can be used as validation references for both airborne and space borne  
62 measurements.

63 Furthermore, NO<sub>2</sub> retrievals from airborne remote sensing instruments, such as the Geostationary Coast and Air Pollution  
64 Event (GEO-CAPE) Airborne Simulator (GCAS) (Kowalewski and Janz, 2014), the Heidelberg Airborne Imaging DOAS  
65 Instrument (HAIDI) (General et al., 2014), the Geostationary Trace gas and Aerosol Sensor Optimization (GeoTASO) (Leitch  
66 et al., 2014), the Airborne Prism Experiment (APEX; Popp et al., 2012), the Airborne Imaging DOAS instrument for  
67 Measurements of Atmospheric Pollution (AirMAP; Meier et al., 2017; Schönhardt et al., 2015), the Small Whiskbroom Imager  
68 for atmospheric composition monitorinG (SWING; Merlaud et al. 2018), and the Spectrolite Breadboard Instrument (SBI;  
69 Vlemmix et al., 2017; Tack et al., 2019) have also been performed to identify local emission sources and obtain highly resolved  
70 horizontal NO<sub>2</sub> distributions.

71 Observations using airborne measurements have an advantage as they enable the observation of horizontal distributions of  
72 trace gases at resolutions higher than space-based satellites and provide data over a wider area than ground-based observations.  
73 For example, Nowlan et al. (2018) retrieved tropospheric NO<sub>2</sub> VCDs over Houston, Texas, during the Deriving Information  
74 on Surface Conditions from Column and Vertically Resolved Observations Relevant to Air Quality (DISCOVER-AQ)  
75 campaign and identified a high correlation with data retrieved from Pandora. Popp et al. (2012) also presented the morning  
76 and afternoon NO<sub>2</sub> spatial distribution in Zurich, Switzerland, using APEX. Tack et al. (2017) have conducted high-resolution  
77 mapping of NO<sub>2</sub> over three Belgium cities (Antwerp, Brussels, and Liège) using APEX and Judd et al. (2020) and Tack et al.  
78 (2021) compared NO<sub>2</sub> VCDs retrieved from GCAS/GeoTASO and APEX with those obtained from TROPOMI over New  
79 York City and Antwerp and Brussels, respectively. Merlaud et al. (2013) observed NO<sub>2</sub> VCDs in Turceni over Romania using  
80 SWING mounted on an unmanned aerial vehicle (UAV) during the Airborne Romanian Measurements of Aerosols and Trace  
81 gases (AROMAT) campaign. These existing NO<sub>2</sub> retrievals, using airborne measurements, have been useful for constraining  
82 regional air quality models due to the highly resolved source identification and the ability to tie these results to the ground-  
83 based observations.

84 This work focuses on airborne NO<sub>2</sub> retrievals from GeoTASO. This instrument was developed by Ball Aerospace to reduce  
85 mission risk for the UV-VIS air quality measurements from geostationary orbit for the GEMS and TEMPO missions (Leitch  
86 et al., 2014). The retrieval of NO<sub>2</sub>, SO<sub>2</sub>, and HCHO observed from GeoTASO L1B data using DOAS and principal component  
87 analysis (PCA) (Wold et al., 1987) was conducted through the DISCOVER-AQ and KOREa-United States Air Quality  
88 (KORUS-AQ) campaigns (Nowlan et al., 2016; Judd et al., 2018; Choi et al., 2020; Chong et al., 2020). The KORUS-AQ  
89 campaign is a joint study between the National Institute of Environmental Research (NIER) and National Aeronautics and  
90 Space Administration (NASA) to monitor megacity air pollution and transboundary pollution, and to prepare for geostationary  
91 satellite (i.e., GEMS, TEMPO, and Sentinel-4) air quality observability (of trace gases and aerosols), organized from May to  
92 June 2016.

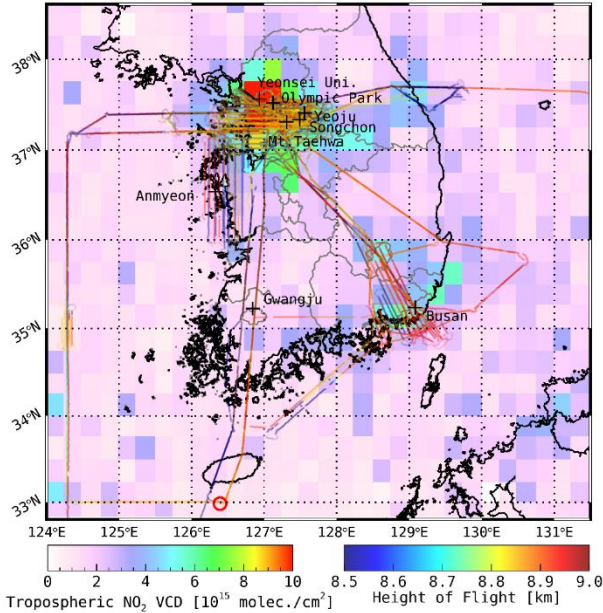
93 Although surface NO<sub>2</sub> concentrations in South Korea are high due to high population density, high traffic volumes, and many  
94 industrial complexes and thermal power plants, and whereas NO<sub>2</sub> retrieval studies using airborne and ground measurements

over North America, Europe, China, and Japan have been conducted, data for South Korea remain limited. The specific aims of this study are as follows:

- (1) To retrieve tropospheric NO<sub>2</sub> vertical column data using GeoTASO measurements over polluted regions of the Seoul and Busan metropolitan areas and the Anmyeon industrial regions of the Korean Peninsula.
- (2) To estimate NO<sub>2</sub> VCD uncertainties using error propagation accounting for spectral fitting errors and AMF uncertainties associated with input data errors, including aerosol optical depth (AOD), single scattering albedo (SSA), aerosol loading height (ALH), and surface reflectance.
- (3) To compare NO<sub>2</sub> VCDs retrieved from GeoTASO and those obtained from OMI and ground-based Pandora instruments, as well as surface in situ measurements.

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

### 2.1 Campaign area



**Figure 1. Flight paths of the NASA LaRC B200 aircraft carrying GeoTASO and the average tropospheric NO<sub>2</sub> VCDs obtained from OMI gridded to a 0.25°×0.25° horizontal grid during the KORUS-AQ campaign period. The line colour represents flight height. In this period, the GeoTASO observations focused on megacities (Seoul and Busan) and industrial complex area (Anmyeon) with high tropospheric NO<sub>2</sub> concentrations. The reference spectrum for spectral fitting is obtained from the radiation data under the Jeju Island (marked with red circle).**

The Korean Peninsula, located on the Asia-Pacific coast, has a complex atmospheric environment by local emissions and long-range transport under appropriate weather conditions (Jeong et al., 2017; NIER and NASA, 2020; Choo et al., 2021). In

115 particular, Seoul, the capital of South Korea, and the metropolitan area are densely populated, and power plants and industrial  
 116 activities on the northwest coast are carried out, which emits relatively large amounts of pollutants. The KORUS-AQ campaign  
 117 conducted three-dimensional observations, including ground-based remote, aircraft, satellite observation, and air quality  
 118 modelling, to understand the complex air quality and interpret the observations of GEMS launched in 2020. The KORUS-AQ  
 119 campaign period was from May 2 to June 10, 2016. During the KORUS-AQ campaign, measurements of air pollutants were  
 120 carried out by using the GeoTASO on board the NASA Langley Research Center B200 aircraft to monitor air quality and long-  
 121 range transport of pollutants over the Korean Peninsula (NIER and NASA, 2020). The GeoTASO observations conducted a  
 122 total of 30 times over 23 days out of 40 days. Most observations were made once or twice a day. Each flight were planned and  
 123 conducted on a day when weather conditions were fine and flight hours were approximately 2-4 hours. Flight information on  
 124 the date of aircraft observation can be found in detail at [http://www-air.larc.nasa.gov/missions/korus-aq/docs/KORUS-](http://www-air.larc.nasa.gov/missions/korus-aq/docs/KORUS-AQ_Flight_Summaries_ID122.pdf)  
 125 [AQ\\_Flight\\_Summaries\\_ID122.pdf](http://www-air.larc.nasa.gov/missions/korus-aq/docs/KORUS-AQ_Flight_Summaries_ID122.pdf). Figure 1 shows the flight routes of B200 and the tropospheric NO<sub>2</sub> VCDs obtained from  
 126 the OMI during the campaign period. The observations were concentrated in the metropolitan areas of Seoul and Busan and  
 127 the industrial areas of Anmyeon, with a flight altitude of 8,000–9,000 m.  
 128 As shown in Fig. 1, GeoTASO observations were conducted focusing on highly NO<sub>2</sub>-polluted regions in the Seoul and Busan  
 129 metropolitan areas and the Anmyeon region during the KORUS-AQ campaign. The Seoul metropolitan area (Seoul Special  
 130 City, Gyeonggi Province, and Incheon City) is one of the most densely populated areas worldwide, with a population of  
 131 approximately 20 million in 2016. Busan is the second-largest city in South Korea, with a population of approximately 3.4  
 132 million in 2016. Anmyeon is located southwest of Seoul with petrochemical complexes, steel mill works, and thermal power  
 133 stations in this area. The background colour in Fig. 1 represents the average NO<sub>2</sub> VCD obtained from the OMI during the  
 134 KORUS-AQ campaign period, showing over  $1 \times 10^{16}$  molecules cm<sup>-2</sup> over the Seoul metropolitan area. The OMI data obtained  
 135 by the Level 2.0 OMNO2 version 3.0 and downloaded from the NASA's Earthdata search  
 136 (<http://search.earthdata.nasa.gov/search/>). We calculated the arithmetic means of the tropospheric NO<sub>2</sub> VCDs, similar to Choo  
 137 et al. (2020), to obtain the grid data (0.25° × 0.25°) during KORUS-AQ period. The average tropospheric NO<sub>2</sub> VCD data were  
 138 excluded from 30 May 2016 to 9 Jun 2016, when the OMI data did not exist during the campaign period.

## 139 **2.2 Pandora**

140 NO<sub>2</sub> VCDs retrieved from the GeoTASO were validated using those from NASA's Pandora Spectrometer system. The Pandora  
 141 spectrometer is a hyper-spectrometer that can provide direct sun measurements of UV/Vis spectra (280–525 nm with a full  
 142 width at half maximum (FWHM) of 0.6 nm) for observing atmospheric trace gases. During the KORUS-AQ, eight Pandora  
 143 instruments monitored NO<sub>2</sub> and ozone (O<sub>3</sub>) VCD as depicted by plus symbols in Fig. 1. The retrieved data are available on the  
 144 KORUS-AQ pages of NASA's Goddard Space Flight Center website  
 145 (<https://avdc.gsfc.nasa.gov/pub/DSCOVER/Pandora/DATA/KORUS-AQ/>). We compared NO<sub>2</sub> VCDs obtained from five  
 146 Pandora measurement (Busan university: 35.24 °N, 129.08 °E; Olympic park: 37.52 °N, 127.13 °E; Songchon: 37.41 °N,

127.56 °E; Yeosu: 37.34 °N, 127.49 °E; Yonsei University: 37.56 °N, 126.93 °E) within 0.05 degree and 30 min with those from GeoTASO. Because NO<sub>2</sub> has a short atmospheric lifetime, especially during the summer (Shah et al., 2020), its spatial and temporal distributions vary notably. A detailed description of Pandora’s operation during the KORUS-AQ campaign has been previously reported (Herman et al., 2018; Spinei et al., 2018).

**2.3 Ground-based in situ NO<sub>2</sub> measurement**

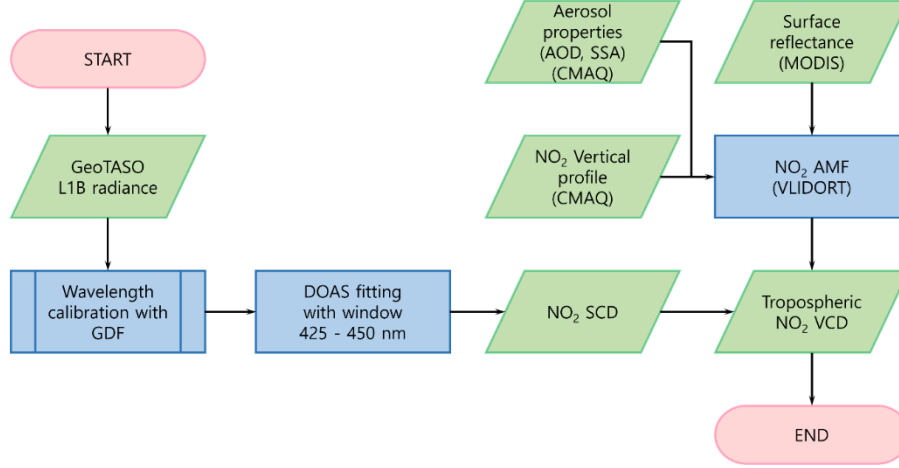
Although the basic physical quantity of VCD and surface mixing ratio from in-situ measurements are different, comparison of their spatiotemporal variations provides useful information for deriving surface air quality from airborne instruments (e.g., Jeong and Hong, 2021a; 2021b *and references therein*). In this study, we compare the NO<sub>2</sub> VCDs (molecules cm<sup>-2</sup>) retrieved from GeoTASO to surface mixing ratios measured by ground-based in-situ monitoring network over South Korea (i.e., Air-Korea, a national real-time air quality network; <https://www.airkorea.or.kr/>). The instruments utilize the chemiluminescence method (Kley and McFarland, 1980), and approximately 400 air quality monitoring sites in Korea are registered in the system, providing hourly surface NO<sub>2</sub> concentrations. We compared NO<sub>2</sub> VCDs retrieved from GeoTASO within 0.5 km and 30 min with NO<sub>2</sub> concentrations obtained from Air-Korea.

**2.4 GeoTASO measurements**

NO<sub>2</sub> VCDs were retrieved from the L1B radiance dataset (version: V02y) obtained using GeoTASO during the KORUS-AQ campaign. The NASA Goddard Space Flight Center conducted the L1B radiance calibration, which included offset and smear correction, gain matching, amplifier cross-talk correction, dark rate correction, integration normalisation, sensitivity derivation, wavelength registration, geo-registration, non-linearity correction, and ground pixel geolocation (Kowalewski et al., 2017; Chong et al., 2020). The detailed specifications of GeoTASO are listed in Table 1.

**Table 1. Summary for GeoTASO instrument and optical specification.**

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



**Figure 2. Flowchart of the algorithm for retrieving tropospheric NO<sub>2</sub> data from GeoTASO.**

#### 2.4.1 NO<sub>2</sub> slant column density retrieval

Figure 2 shows the flowchart for retrieving the tropospheric NO<sub>2</sub> VCD from the GeoTASO. We first retrieved NO<sub>2</sub> SCDs using the DOAS method (Platt, 1994). Nonlinear least square minimisation was used to retrieve the NO<sub>2</sub> SCDs which minimize the difference between the measured optical depth and the modelled value in QDOAS software (Eq. (1); Danckaert et al., 2012).

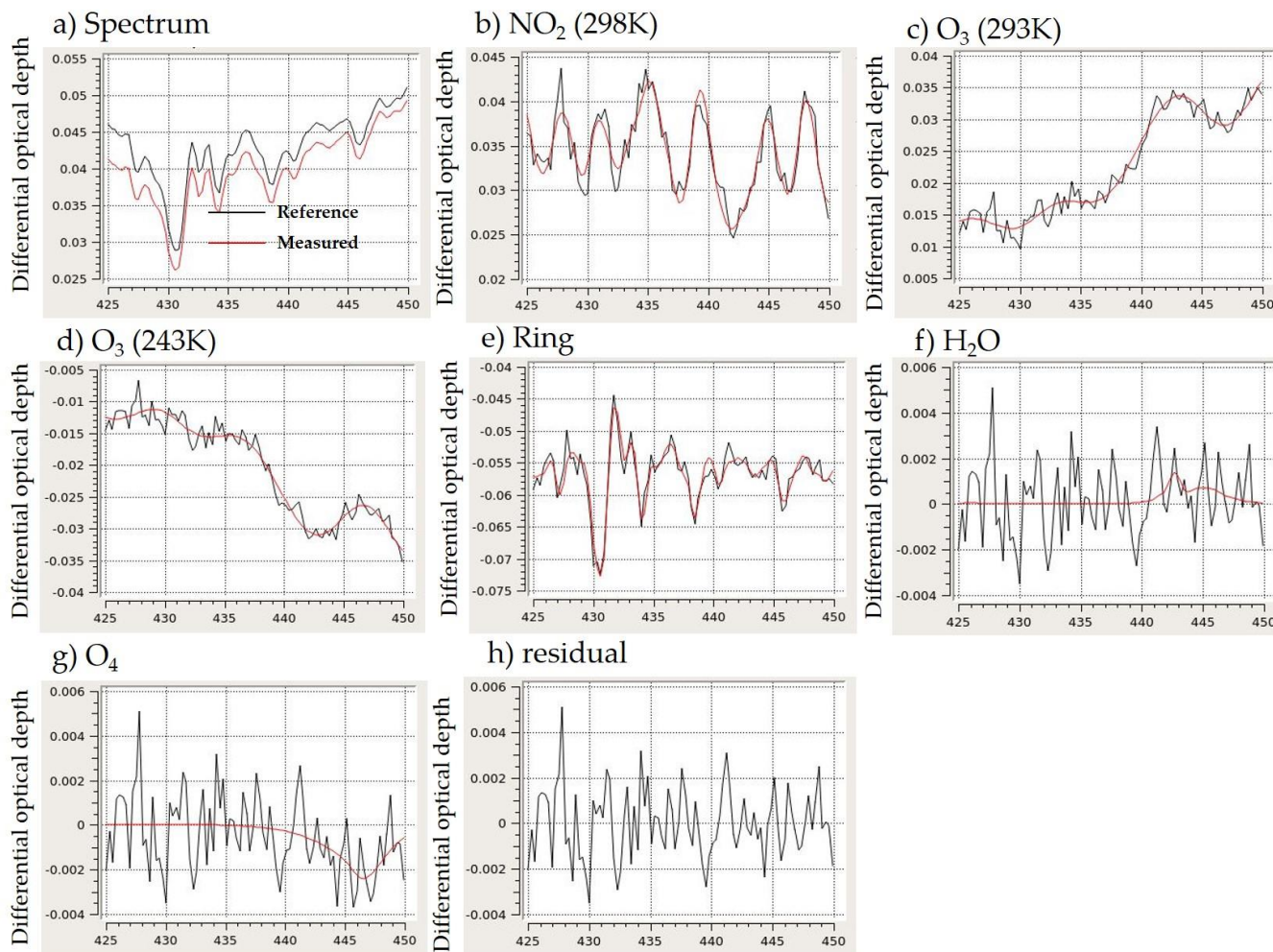
$$\frac{\ln I(\lambda)}{\ln I_0(\lambda)} = -(\sum_{j=1}^m \rho_j \times \sigma'_j(\lambda) + B(\lambda) + R(\lambda) + A(\lambda) + N(\lambda)) \quad (1)$$

Where  $I(\lambda)$  is the measured earthshine radiance at wavelength  $\lambda$ ;  $I_0$  is the reference radiance from the sea surface south of Jeju Island (red circle in Fig. 1, 32.983°N, 126.392°E) on 09 AM in 1 May 2016. The Community Multiscale Air Quality (CMAQ) modelling system data indicated that the NO<sub>2</sub> VCD from the surface to 50 hPa at this point on this day was  $6.75 \times 10^{15}$  molecules cm<sup>-2</sup> (averaged NO<sub>2</sub> VCD obtained from OMI available during KOURS-AQ period is  $4.77 \times 10^{15}$  molecules cm<sup>-2</sup> and standard deviation of  $1.33 \times 10^{15}$  molecules cm<sup>-2</sup>, respectively);  $\rho_j$  represents the SCD of each species  $j$ ;  $\sigma'_j(\lambda)$  represents the differential gas phase absorption cross-section convolved with the Gaussian distribution function (GDF) with 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 wavelength  $\lambda$  of

187 species  $j$ , respectively. The spectral fitting window was selected from 425 to 450 nm. To determine the wavelength registration  
 188 more accurately in the narrow fitting window, additional wavelength calibration of the spectra for each of the 33 across track  
 189 pixels was performed using a high-resolution solar reference spectrum (Kurucz solar spectrum) (Chance and Kurucz, 2010)  
 190 with the GDF. The absorption cross-sections of  $\text{NO}_2$  (Vandaele et al., 1998),  $\text{O}_3$  (Bogumil et al., 2000),  $\text{H}_2\text{O}$ , and the Ring  
 191 effect as pseudo-absorbers (Chance and Spurr, 1997) were used to construct the model equation; and  $B(\lambda)$ ,  $R(\lambda)$ ,  $A(\lambda)$ , and  
 192  $N(\lambda)$  are the broad absorption of the trace gases, extinction by Mie and Rayleigh scattering, variation in the spectral sensitivity  
 193 of the detector or spectrograph, and noise, respectively, which were accounted by an 8<sup>th</sup> order polynomial. An example of the  
 194 spectral fitting results is presented in Fig. 3.

195

196



197

198 **Figure 3.** An example of the spectral fitting results of  $\text{NO}_2$  retrievals from GeoTASO during the KORUS-AQ campaign (at Gangnam,  
 199 Seoul on 9 June, 2016). Red and black line in the panel (a) represent measured and reference spectrum, respectively. The panels



from (b) to (h) depict examples of spectral fitting results of (b) NO<sub>2</sub>, (c) O<sub>3</sub>(293K), (d) O<sub>3</sub>(243K), (e) ring, (f)H<sub>2</sub>O, (g) O<sub>4</sub> where red and black lines are absorption cross section of target species and the fitting residual plus the absorption of the target species, respectively. The panel (h) shows fitting residual of this example.

203

#### 204 2.4.2 NO<sub>2</sub> AMF calculation

205 AMF, the ratio of SCD to VCD, can be calculated using the scattering weight ( $\omega$ ) and shape factor ( $S$ ) (Palmer et al., 2001) in  
206 Eq. (2)–(5).

$$207 \text{ AMF} = \frac{SCD}{VCD} \quad (2)$$

$$208 \text{ AMF} = \text{AMF}_G \int_{z_1}^{z_2} \omega(z) S(z) dz \quad (3)$$

$$209 \omega(z) = - \frac{1}{\text{AMF}_G} \frac{\partial \ln I_B}{\partial \tau} \quad (4)$$

$$210 S(z) = \frac{\alpha(z)n(z)}{\int_{z_1}^{z_2} \alpha(z)n(z) dz} \quad (5)$$

211 Where  $\text{AMF}_G$  represents the geometric AMF,  $I_B$  is the earthshine radiance,  $\tau$  is the optical depth,  $\alpha$  is the absorption cross-  
212 section, and  $n$  is the number density of the absorber. NO<sub>2</sub> AMF was calculated using a linearised pseudo-spherical scalar and  
213 vector discrete ordinate radiative transfer model (VLIDORT, version 2.6; Spurr and Christi, 2014). Aerosol properties, such  
214 as AOD, SSA, and *a priori* NO<sub>2</sub> vertical profile information, were simulated using the CMAQ, and surface reflectivity was  
215 obtained from the Moderate Resolution Imaging Spectroradiometer (MODIS) (Collection 6). The surface reflectance products,  
216 MOD09CMG and MYD09CMG, available at a 0.05 degree (~5.6 km) spatial resolution, provide an estimate of the surface  
217 spectral reflectance including MODIS bands 1 through 7. The products were corrected for atmospheric conditions such as  
218 aerosol, gasses, Rayleigh scattering. In previous studies (Lamsal et al., 2017; Nowlan et al., 2018; Judd et al., 2019; Chong et  
219 al., 2020), an AMF were described for both above and below aircraft altitude is used to convert NO<sub>2</sub> SCDs to VCDs using Eq.  
220 (6)–(8).

$$221 \text{ AMF} \uparrow = \text{AMF}_G \int_{z_A}^{z_{TOA}} \omega(z) S(z) dz \quad (6)$$

$$222 \text{ AMF} \downarrow = \text{AMF}_G \int_{z_0}^{z_A} \omega(z) S(z) dz \quad (7)$$

$$223 \text{ NO}_2 \text{ VCD} \downarrow = \frac{\text{NO}_2 \text{ SCD} - \text{AMF} \uparrow \cdot \text{NO}_2 \text{ VCD} \uparrow}{\text{AMF} \downarrow} \quad (8)$$

224 Where  $\text{AMF} \uparrow$  and  $\text{AMF} \downarrow$  are AMF above and below aircraft, respectively, and  $\text{NO}_2 \text{ VCD} \uparrow$  represents NO<sub>2</sub> VCD above the  
225 aircraft obtained from a chemical transport model (CTM). However, here we calculated NO<sub>2</sub> VCD $\downarrow$  by dividing NO<sub>2</sub> SCDs  
226 by  $\text{AMF} \downarrow$  because stratospheric and free tropospheric NO<sub>2</sub> ( $\text{NO}_2 \text{ VCD} \uparrow$ ) column densities are much lower than tropospheric  
227 NO<sub>2</sub> column densities, especially in megacities and industrial areas (Valks et al., 2011).

228

## 229 2.5 Chemical model description

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

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

$$239 \text{ AOD} = \int Q_{sca}(z) dz = \int \{Q_{sca}(z) + Q_{abs}(z)\} dz \quad (9)$$

$$240 Q_{abs}[\text{Mm}^{-1}] = \sum_i \sum_j \{ (1 - \omega_{ij}) \cdot \beta_{ij} \cdot f_{ij}(RH) \cdot [C]_{ij} \} \quad (10)$$

$$241 Q_{sca}[\text{Mm}^{-1}] = \sum_i \sum_j \{ \omega_{ij} \cdot \beta_{ij} \cdot f_{ij}(RH) \cdot [C]_{ij} \} \quad (11)$$

242 Here,  $\omega_{ij}$  indicates SSA of particulate species  $i$  for the particulate mode (or size bin)  $j$ ,  $\beta_{ij}$  denotes the mass extinction  
243 efficiency,  $f_{ij}(RH)$  is the hygroscopicity factor according to the relative humidity (RH), and  $[C]_{ij}$  is the concentration of  
244 particulate species. CMAQ SSA is defined as the ratio of the integrated  $Q_{sca}$  to AOD, and  $\text{NO}_2$  vertical profiles were obtained  
245 from  $\text{NO}_2$  concentrations at each vertical layers by conducting CMAQ simulations. Details of the model descriptions and  
246 calculations of optical properties are given in Lee et al. (2020) and Malm and Hand (2007).

## 247 3 Results and discussion

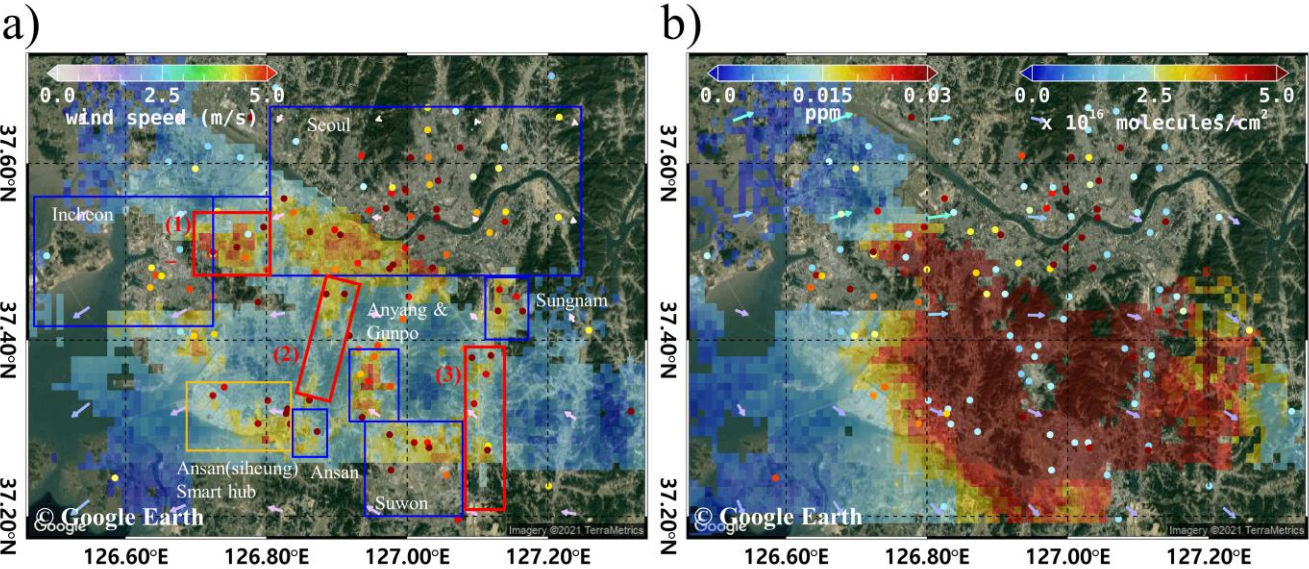
### 248 3.1 $\text{NO}_2$ VCD retrieval

249 We showed the finally  $\text{NO}_2$  VCDs by binning them with  $0.01^\circ \times 0.01^\circ$  from 250 m spatial resolution. Although the spatial  
250 binning  $\text{NO}_2$  VCDs were compared to those at native resolution, we noted that the spatiotemporal variability was still able to  
251 be clearly distinguished from the background at  $0.01^\circ$  binning resolution. Chong et al. (2020) showed that larger VCDs at 250  
252 m resolutions do not necessarily lead to larger VCDs at wider resolutions. As the results of  $\text{NO}_2$  VCD, we selected the dates  
253 observed in both the morning and afternoon during the KORUS-AQ period over Seoul metropolitan area, Busan, and Anmyeon.  
254 The retrieved dates for  $\text{NO}_2$  VCDs were 5, 9, and 10 Jun, 2016.

#### 255 3.1.1 Seoul metropolitan region

256 The population of the Seoul metropolitan region is approximately 20 million, which is approximately 40% of the total  
257 population of South Korea. It is very rare to obtain high-resolution horizontal  $\text{NO}_2$  VCD distributions using airborne

258 measurements in the morning and afternoon, especially in Asian megacities. Fig. 4 shows tropospheric NO<sub>2</sub> VCDs over Seoul  
 259 on 9 June 2016, at 9 AM and 3 PM local time (LT). According to the Terra/Aqua CLDMASK data (Ackerman et al., 1998),  
 260 on this day, the cloud fraction was less than 0.3 over the entire domain of Fig. 4.



261  
 262 **Figure 4. Tropospheric NO<sub>2</sub> VCD, binned to a 0.01°×0.01° horizontal grid, in the Seoul metropolitan region on 9, June 2016 retrieved**  
 263 **from GeoTASO: a) at 9 AM and b) at 3 PM. The red boxes represent expressways (counterclockwise from left to right, (1) Gyeongin**  
 264 **expressway, (2) Seohaean expressway, and (3) Gyeongbu expressway), the orange box indicates the industrial complex, and the blue**  
 265 **boxes indicate the major cities (Seoul, Incheon, Suwon, Bucheon, Anyang, Gunpo, Sungnam, and Ansan) of the Seoul metropolitan**  
 266 **region. Colours of the circles depict the NO<sub>2</sub> surface mixing ratio obtained from Air-Korea. The colour arrows show the wind**  
 267 **direction and speed at 1000 hPa over Seoul metropolitan region, obtained via the Unified Model (UM) simulations (background**  
 268 **RGB image is from Google Earth; <https://www.google.com/maps/>).**

269  
 270 In the morning, NO<sub>2</sub> VCDs retrieved from GeoTASO were highly correlated with expressways (red boxes in Fig. 4), such as  
 271 the Gyeongin, Seohaean, and Gyeongbu Expressways, and over major cities with heavy traffic, such as Seoul, Bucheon, Ansan,  
 272 Anyang, and Suwon. GeoTASO observed NO<sub>2</sub> VCD values three-times higher ( $>3 \times 10^{16}$  molecules cm<sup>-2</sup>) in these areas  
 273 compared to the surrounding rural areas. In particular, high NO<sub>2</sub> VCD values above  $6 \times 10^{16}$  molecules cm<sup>-2</sup> were observed  
 274 above the Gyeongin Expressway, which has very heavy traffic in a relatively short section, and the Gunpo Complex Logistics  
 275 zone, where diesel vehicle traffic is also high. The major NO<sub>2</sub> source regions and the regions where high NO<sub>2</sub> VCD values  
 276 were observed were highly consistent at 9 AM because the wind speed at this time—as obtained from the unified model (UM)  
 277 based Regional Data Assimilation and Prediction System (RDAPS) of the Korea Meteorological Administration (KMA)—  
 278 was as low as 0.1 ms<sup>-1</sup> and the average wind direction was 84.7° at 1000 hPa over Seoul metropolitan region. The average  
 279 daily traffic volume of these expressways exceeds 150,000 vehicles, and the total number of vehicles registered in these major  
 280 cities is  $> 6,000,000$ , with an average daily mileage per car per day of over 38 km. Detailed information on these cities and

expressways is listed in Table 2 and Table 3. Based on the level of vehicular traffic, combustion using gasoline and diesel engines leads to high overall emissions of NO<sub>2</sub> in the Seoul metropolitan region (Kendrick et al., 2015).

**Table 2. The population, number of registered vehicles, and average mileage per car per day of major cities in the Seoul metropolitan region obtained from the Korean Statistical Information Service (<https://kosis.kr/eng>).**

City	Population (millions)	Vehicle registration number (thousands)	Average mileage (km/car/day)
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

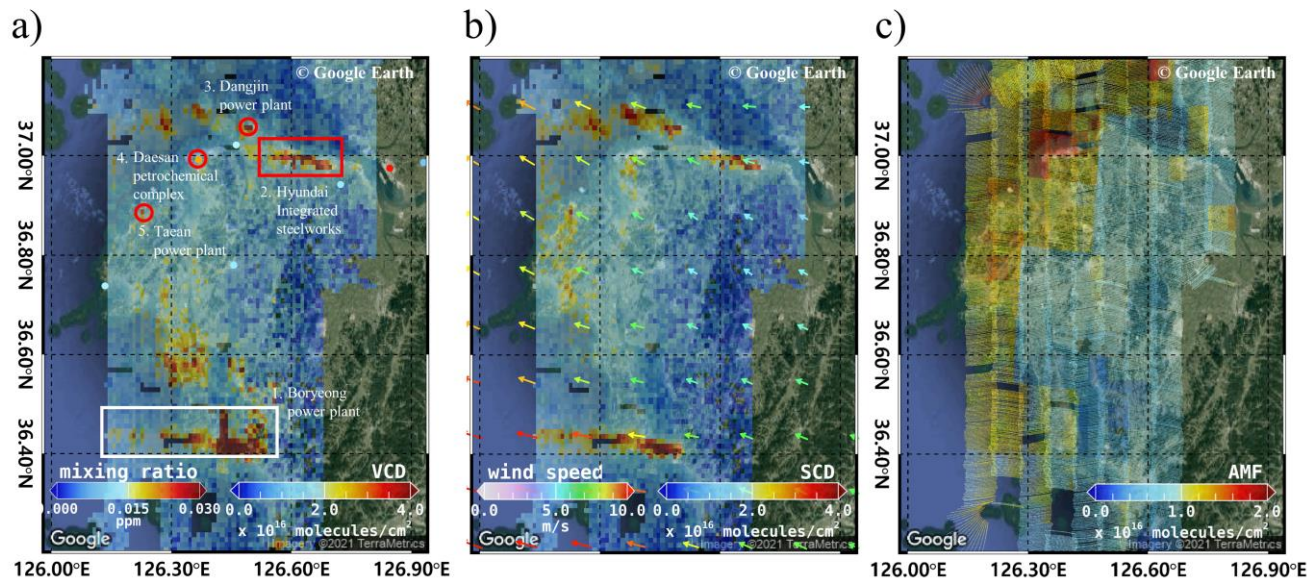
**Table 3. Daily average traffic volume on the Gyeongin, Gyeongbu, and Seohaean Expressways obtained using the Traffic Monitoring System (<https://www.road.re.kr>).**

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

Compared the data from the morning, the average wind speed and wind direction were 1.7 m/s and 284.5° at 1000 hPa in the afternoon and the afternoon had extremely high tropospheric NO<sub>2</sub> VCD values (exceeding  $5 \times 10^{16}$  molecules cm<sup>-2</sup>) in most of the Seoul metropolitan regions including rural areas, whereas the NO<sub>2</sub> mixing ratio (MR) obtained from Air-Korea decreases in the afternoon. According to Tzortziou et al. (2018), similar results were retrieved from the Pandora site in Seoul, with higher afternoon NO<sub>2</sub> VCDs than in the morning. This result is presumed to be due to the reason that the amount of NO<sub>2</sub> produced by chemical conversion of nitric oxide (NO) by O<sub>3</sub> and VOCs in the atmosphere, along with NO<sub>x</sub> generated by regional emissions (traffic) in the Seoul metropolitan region, is greater than the amount lost by photolysis and transport to nearby areas (Herman

et al., 2018). In addition, the increase in tropospheric NO<sub>2</sub> VCD in the afternoon is presumed to be due to the accumulation and dispersion of NO<sub>2</sub> according to the change in the planetary boundary layer height (Ma et al., 2013).

### 3.1.2 Industrial and power plant regions in Anmyeon



**Figure 5. a) Tropospheric NO<sub>2</sub> VCD and b) NO<sub>2</sub> SCD retrieved from GeoTASO, and c) NO<sub>2</sub> AMF, native resolution (250 m) calculated using VLIDORT over Anmyeon in South Korea on 5 June 2016. The NO<sub>2</sub> VCD and SCD were gridded into to a spatial resolution of 0.01°×0.01°. The colored arrows indicate wind speed and wind direction at 850 hPa from the Unified Model (UM) simulations. The red circles and rectangle in panel (a) represent the major NO<sub>2</sub> emission sources, such as steelworks and power plants (background RGB image is from Google Earth; <https://www.google.com/maps/>).**

The high spatial resolution of tropospheric NO<sub>2</sub> VCD from GeoTASO over the Anmyeon industrial region, where many industrial facilities and several power plants are distributed, is shown in Fig. 5. The panels a and b of this figure show the binned tropospheric NO<sub>2</sub> VCD and NO<sub>2</sub> SCD retrieved from GeoTASO L1B data, respectively, between 13:00 and 17:00 LT on 5 June 2016. The panel c depicts the calculated AMF of NO<sub>2</sub> from native resolution over the domain. The GeoTASO observations clearly detected moderate and strong NO<sub>2</sub> emission sources over this area: (1) Boryeong power plant, (2) the Hyundai integrated steelworks, (3) Dangjin power plant, (4) the Daesan Petrochemical Complex, and (5) Taean power plant. High NO<sub>2</sub> VCD values ( $> 5 \times 10^{16}$  molecules cm<sup>-2</sup>) were observed over steel mill works, petrochemical complexes, and power plants, whereas values were comparatively low ( $< 1 \times 10^{16}$  molecules cm<sup>-2</sup>) over small cities including Seosan, Dangjin, and Boryeong with populations of less than 0.1 million, and the Seohaean Expressway. In 2016, the annual NO<sub>x</sub> emissions by the Hyundai steelworks and the Dangjin and Boryeong power plants were about 10.3, 11.9, and 16.8 kt year<sup>-1</sup>, respectively. NO<sub>2</sub> emission rates from major industrial facilities in the Anmyeon region are shown in Table 4.

318

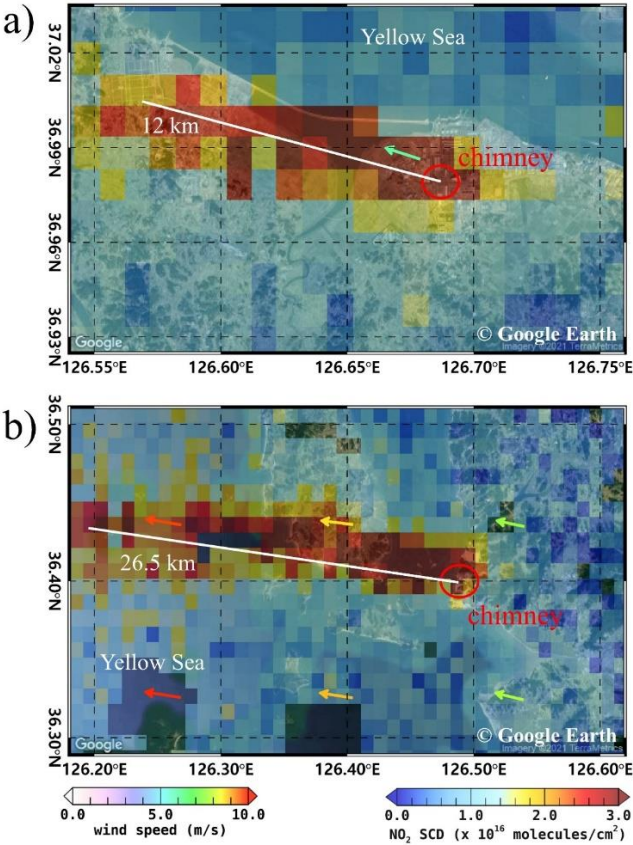
319 **Table 4. NO<sub>2</sub> emission rates from major industrial facilities in the Anmyeon region obtained from the Continuous Emission**  
320 **Monitoring System of the Korea Environment Corporation (<https://www.stacknsky.or.kr/eng/index.html>).**

Industrial facilities	NO <sub>2</sub> emission rate (2016) (kg/year)
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

321



Figure 5 shows high NO<sub>2</sub> concentrations of major industrial facilities in the Anmyeon region, where fossil fuel combustion in factories and thermal power plants leads to high emissions (Prasad et al., 2012). Due to relatively sparse distribution over rural areas, the Air-Korea measurements did not detect the major NO<sub>2</sub> plume as shown in Fig. 5a. Thus, airborne remote sensing systems, such as GeoTASO, can effectively compliment the ground-based networks for monitoring minor and major NO<sub>2</sub> emissions, particularly over these remote industrial regions.

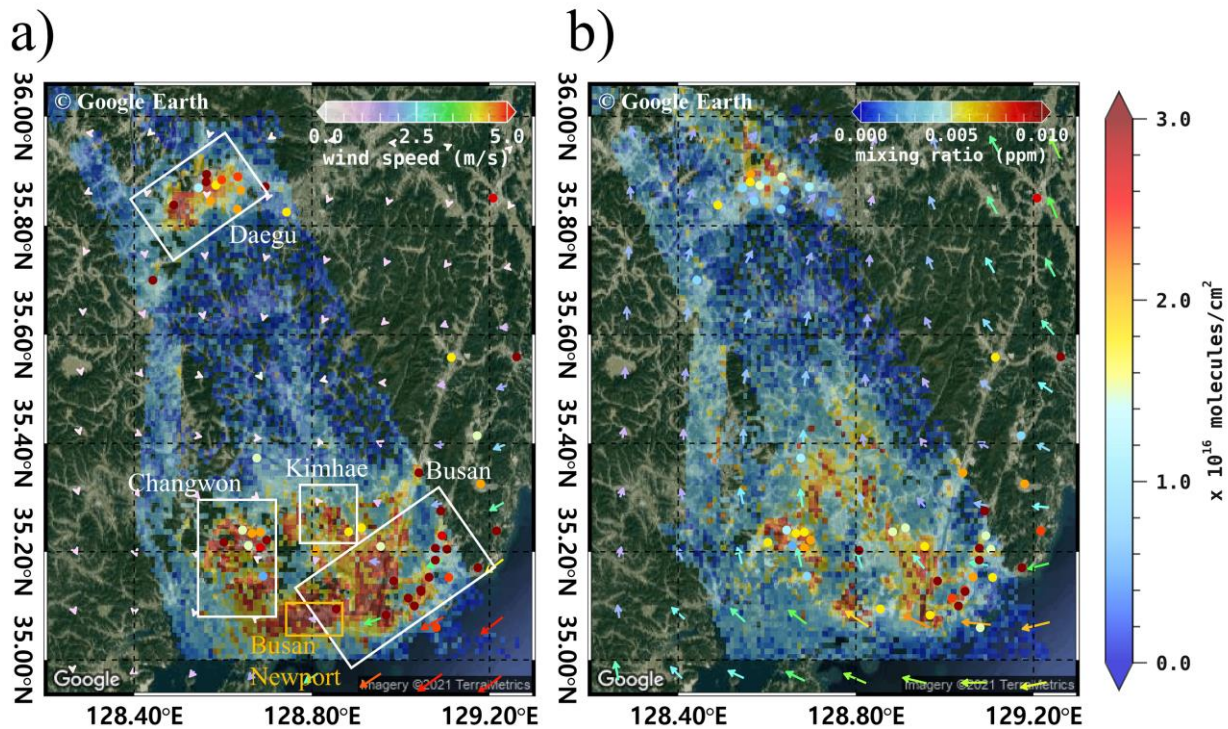


**Figure 6. Enlarged view of GeoTASO tropospheric NO<sub>2</sub> VCD observation over a) Hyundai steel works, indicated by the red box in Figure 5, and b) the Boryeong power plant, indicated by the white box in Figure 5. The data were gridded to a spatial resolution of 0.01°×0.01°. The arrows represent the wind direction and speed at 850 hPa from the Unified Model (UM) simulations, respectively (background RGB image is from Google Earth; <https://www.google.com/maps/>).**

The GeoTASO data captured not only NO<sub>2</sub> emissions from the chimneys of steelworks and power plants but also its transport by the wind. Fig. 6a and 6b show enlarged views of tropospheric NO<sub>2</sub> SCD retrieved using GeoTASO over the Hyundai steelworks (red box in Fig. 5) and the Boryeong power plant (white box in Fig. 5). The arrows in Fig. 6 represent the prevailing wind direction and speed from RDAPS. NO<sub>2</sub> emitted from the chimneys of these sites was transported to the Yellow Sea,

travelling distances of over 26 km at speeds of approximately  $6 \text{ ms}^{-1}$ . According to Chong et al. (2020), similar results were found for  $\text{SO}_2$  emitted and transported from these sites.

### 3.1.3 Busan metropolitan region



**Figure 7. Tropospheric  $\text{NO}_2$  VCD in the Busan metropolitan region in the (a) morning and (b) afternoon of 10 June 2016. The data were gridded into to a spatial resolution of  $0.01^\circ \times 0.01^\circ$ . The wind speed (colours scale) and wind direction (arrows) at 1000 hPa pressure level were obtained from the Unified Model (UM) simulations. The white boxes represent major cities such as Busan, Daegu, Changwon, and Kimhae. The orange box represents Busan Newport (background RGB image is from Google Earth; <https://www.google.com/maps/>).**

Fig. 7a and 7b show tropospheric  $\text{NO}_2$  VCD retrieved from the GeoTASO L1B data over the Busan metropolitan region on 10 June 2016 in the morning (between 08:00 and 11:00 LT) and afternoon (between 13:00 and 16:00 LT), respectively. The arrows in Fig. 7 show the wind speed and wind direction of 1000 hPa obtained from the UM-RDAPS, with the average wind speed and wind direction of 0.9 m/s and  $55.4^\circ$ , 1.9 m/s and  $147.0^\circ$ , respectively, in the morning and afternoon. High  $\text{NO}_2$  VCDs were observed above urban areas, the port, industrial complexes, and the inter-city road between Busan and Changwon. Similar to the Seoul metropolitan regions, it is estimated that combustion using gasoline and diesel engines contributes to the high  $\text{NO}_2$  emission. In the morning,  $\text{NO}_2$  VCDs were high (approximately  $3 \times 10^{16}$  molecules  $\text{cm}^{-2}$ ) in the major cities and, especially, around Busan Newport, with values exceeding  $7 \times 10^{16}$  molecules  $\text{cm}^{-2}$ . In comparison, in the mountainous regions



355 between Daegu and Busan, NO<sub>2</sub> VCD values were less than  $1 \times 10^{16}$  molecules cm<sup>-2</sup> during the same period. The spatial  
 356 distribution of tropospheric NO<sub>2</sub> VCDs was similar to that in the Seoul metropolitan regions, with high values over major cities  
 357 and roads (compare Figs. 4 and 7). In Busan, fossil fuel combustion using both road vehicles and ships likely contributes to  
 358 the NO<sub>2</sub> emissions. In the afternoon, unlike Seoul metropolitan region, tropospheric NO<sub>2</sub> VCD over Busan decreased by over  
 359  $3 \times 10^{16}$  molecules cm<sup>-2</sup>, which also corresponds with NO<sub>2</sub> MR data obtained from the Air-Korea sites. Detailed information  
 360 on these cities is listed in Table 5.

361

362 **Table 5. The population, number of registered vehicles, and average mileage per car per day of major cities in the Busan**  
 363 **metropolitan region obtained from the Korean Statistical Information Service (<https://kosis.kr/eng>).**

City	Population (millions)	Vehicle registration number (thousands)	Average mileage (km/car/day)
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

364

### 365 3.2 Error estimation

366 NO<sub>2</sub> VCD retrieval accuracy using the DOAS method depends on both the AMF calculation and spectral fitting error of the  
 367 SCD retrieval. Retrieval errors of the NO<sub>2</sub> VCD were estimated using error propagation analysis as expressed in Eq. (12).

$$368 \frac{\varepsilon_{VCD}}{VCD} = \sqrt{\left(\frac{\varepsilon_{SCD}}{SCD}\right)^2 + \left(\frac{\varepsilon_{AMF}}{AMF}\right)^2} \quad (12)$$

369 Where  $\varepsilon_{VCD}$  is the total error of NO<sub>2</sub> VCD. The error of NO<sub>2</sub> SCD ( $\varepsilon_{SCD}$ ) is obtained from the spectral fitting error of NO<sub>2</sub> SCD  
 370 via DOAS spectral fitting.  $\varepsilon_{AMF}$  indicates the error of NO<sub>2</sub> AMF caused by uncertainties in the model input parameters for  
 371 AMF calculation. Uncertainties in aerosol properties (AOD, SSA, and ALH) and surface reflectance for the RTM calculations  
 372 are known to be the major factors affecting NO<sub>2</sub> AMF accuracy (Boersma et al., 2004; Leitão et al., 2010; Hong et al., 2017).  
 373 Therefore, in this present study, we quantified the NO<sub>2</sub> AMF errors ( $\varepsilon_{AMF}$ ) due to uncertainties in the input parameters  
 374 independent of one another using Eq. (13):

$$375 \varepsilon_{AMF} = \sqrt{\left(\frac{\partial AMF}{\partial AOD}\right)^2 \sigma AOD^2 + \left(\frac{\partial AMF}{\partial SSA}\right)^2 \sigma SSA^2 + \left(\frac{\partial AMF}{\partial ALH}\right)^2 \sigma ALH^2 + \left(\frac{\partial AMF}{\partial SFR}\right)^2 \sigma SFR^2} = \sqrt{\sum_{i=1}^4 \left(\frac{\partial AMF}{\partial \chi_i}\right)^2 \sigma_{\chi_i}^2}, \quad (13)$$

376 where  $\frac{\partial AMF}{\partial \chi_i}$  are partial derivatives of NO<sub>2</sub> AMF with respect to the input parameters ( $\chi_i$ ),  $\sigma_{\chi_i}$  represents the uncertainty of the  
 377  $\chi_i$ . The  $\sigma$  of AOD, SSA, surface reflectance, and ALH are assumed as 30% (Ahn et al., 2014), 0.04 (Jethva et al., 2014),  
 378 (0.005+0.05×surface reflectance; EOS Land Validation; <https://landval.gsfc.nasa.gov>), and 1 km (Fishman et al., 2012),

379 respectively, in this study. To derive  $(\frac{\partial AMF}{\partial \chi_i})^2$ , the true  $\chi_i$  is input to the RTM to simulate ‘true’ NO<sub>2</sub> AMF. For the AOD, SSA,  
 380 ALH, and surface reflectance (SFR), perturbed NO<sub>2</sub> AMF was simulated using RTM with  $\chi_i \pm \sigma\chi_i$ .  $\partial\chi_i$  denotes the difference  
 381 between the ‘true’  $\chi_i$  and  $\chi_i \pm \sigma\chi_i$ , and  $\partial AMF$  is the difference between the ‘true’ NO<sub>2</sub> AMF (AMF<sub>true</sub>) simulated with ‘true’  
 382 input values and the new NO<sub>2</sub> AMF (AMF<sub>new</sub>) simulated using the perturbed input parameters  $\chi_i \pm \sigma\chi_i$  (i.e. the original input  
 383 parameters modified by the uncertainty). The simulation for calculating the  $\varepsilon_{AMF}$  was conducted using the input parameters  
 384 on 9 June 2016. On the flight day, average (standard deviation) values of AOD, SSA, ALH, and surface reflectance were 0.39  
 385 (0.10), 0.98 (0.001), 0.27 km (0.10 km), and 0.09 (0.04), respectively.

387 **Table 6. Total errors of NO<sub>2</sub> VCD caused by uncertainties in NO<sub>2</sub> SCD and NO<sub>2</sub> AMF (the average for the flight on 9 June**  
 388 **2016).**

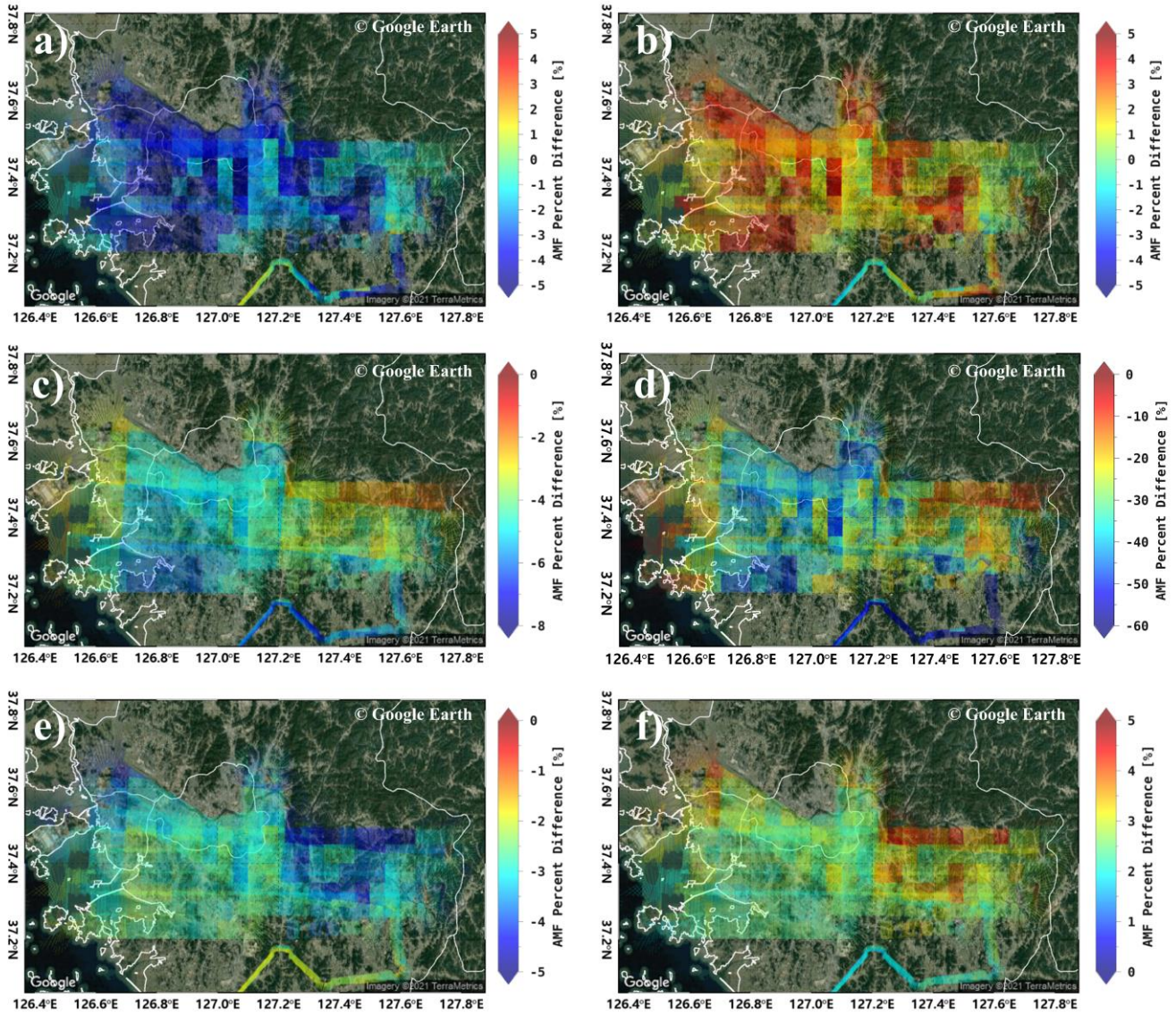
NO <sub>2</sub> AMF errors	AOD	3.0%
	SSA	4.2%
	aerosol loading height	26.4%
	surface reflectance	2.8%
	<b>total NO<sub>2</sub> AMF error due to aerosol uncertainties</b>	<b>27.8%</b>
NO <sub>2</sub> SCD error		11.9%
NO <sub>2</sub> VCD error		31.1%

389  
 390 Table 6 lists the estimated NO<sub>2</sub> VCD error on 9 June 2016 for each sources based on the error propagation method. The error  
 391 estimation was conducted for the pixels where root mean square residual < 0.001 and NO<sub>2</sub> VCD > 5 × 10<sup>15</sup> molecules cm<sup>-2</sup>  
 392 since NO<sub>2</sub> SCD precision is reported to be highly decreased in low NO<sub>2</sub> conditions (Hong et al., 2017). The total NO<sub>2</sub> VCD  
 393 error was 31.1% with a high portion of NO<sub>2</sub> AMF error. The NO<sub>2</sub> SCD error was calculated to be 11.9%, showing the  
 394 importance of accurate DOAS spectral fitting to derive NO<sub>2</sub> SCD. The total AMF error due to uncertainties in input parameters  
 395 was calculated to be 27.8%. Among model input parameters, the effect of ALH on NO<sub>2</sub> AMF become highest (26.4%),  
 396 indicating importance of accurate aerosol profile information. ALH sensitively affects NO<sub>2</sub> AMF because near the surface  
 397 where trace gases and aerosols are well mixed, aerosols lead to multiple scattering effects and the light absorption of trace  
 398 gases due to increasing light path (Castellanos et al., 2015; Hong et al., 2017). Especially, ALH can be the most important  
 399 input parameter in the Asia region where high loadings of aerosol plumes persist throughout the year. The NO<sub>2</sub> AMF

400 calculation errors due to uncertainties in SSA and AOD were 4.2% and 3.0%, respectively. The NO<sub>2</sub> AMF calculation error  
 401 due to uncertainties in aerosol optical properties (SSA and AOD) seems smaller than those in the previous study (Leitão et al.,  
 402 2010). The smaller effect of aerosol properties can be explained by the moderate aerosol loading (AOD = 0.39) on the flight  
 403 day. It is expected that NO<sub>2</sub> AMF errors become larger under high AOD condition. The smallest effect of SRF was found on  
 404 NO<sub>2</sub> AMF calculation error. A priori NO<sub>2</sub> profile shape also can be one of factors to cause calculation error for NO<sub>2</sub> AMF as  
 405 reported in the previous studies (Leitao et al., 2010, Meier et al., 2016). It is necessary to calculate the effect of a priori NO<sub>2</sub>  
 406 profile shape on airborne NO<sub>2</sub> AMF error in the future. Moreover, the resulting uncertainties of input parameters of a  
 407 GeoTASO ground pixel need to be considered by combining the initial uncertainties of CTM and satellite-based products, and  
 408 by the variability of the parameters within a grid box. This kind of analysis should be taken into account in further study.

$$\begin{aligned}
 409 \quad AMF_{percent\_diff} &= \frac{\partial AMF}{(AMF_{true} + AMF_{new}) \div 2} \times 100 & (14)
 \end{aligned}$$

410



411

412 **Figure 8.** Percentage difference between AMF calculated using the CMAQ model simulation and those using a) 30% lower  
 413 AOD, b) 30% higher AOD, c) 0.04 lower SSA, d) 1km higher ALH, compared to the model outputs. The percentage difference  
 414 for AMF calculated using MODIS data and those using e) 20%  $0.005 + 0.05 \times \text{SFR}$  lower SFR, f)  $0.005 + 0.05 \times \text{SFR}$  higher SFR  
 415 (background RGB image is from Google Earth; <https://www.google.com/maps/>).

416

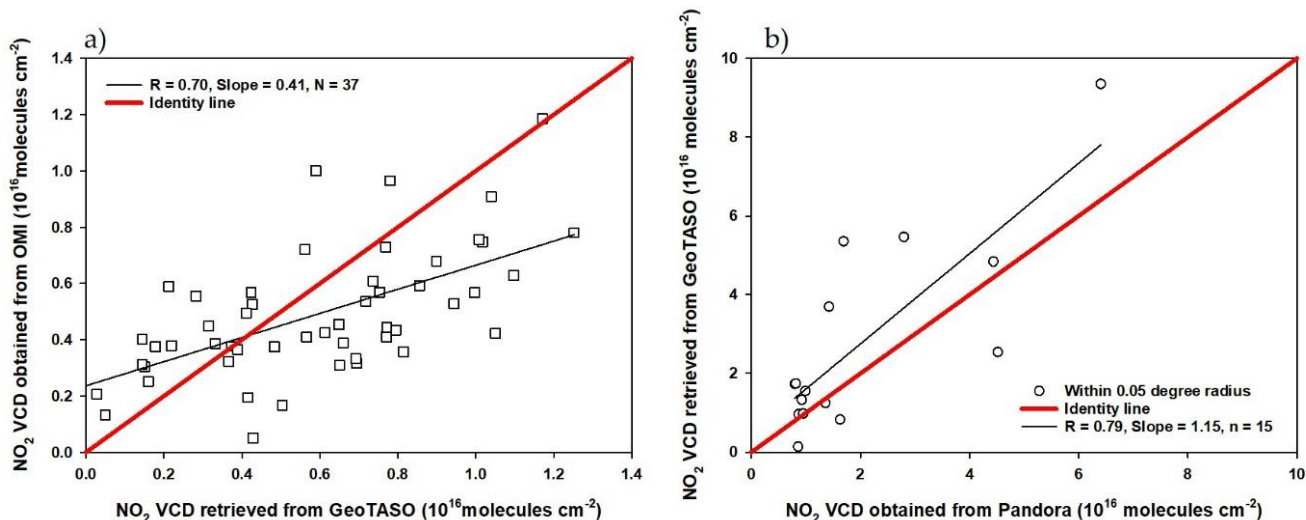
417 In this present study, we additionally investigated the spatial distribution of AMF calculation errors associated with  
 418 uncertainties in aerosol properties (AOD, SSA, ALH, and SFR). Percent difference of  $\text{NO}_2$  AMF ( $\text{AMF}_{\text{percent\_diff}}$ ) was calculated  
 419 on each spatial pixel using Eq. (14). Fig. 8a and 8b show the percent difference error between the calculated AMFs using the

420 CMAQ AOD data with 30% lower (Fig. 8a) and 30% higher (Fig. 8b) values, respectively. The AMF decreased and increase  
 421 by up to 7% with decreasing and increasing AOD, respectively, in the Seoul metropolitan region. We estimated that, under  
 422 low aerosol loading conditions, an increase in AOD near the surface leads to an increase in the scattering probability within  
 423 the surface layer with high NO<sub>2</sub> concentrations. Fig. 8c shows the percent difference error between the calculated AMFs using  
 424 CMAQ SSA data with a 0.04 lower value. The AMF decreased with decreasing SSA because the absorption of light increased.  
 425 The ALH was also found to highly affect the accuracy of the AMF calculations (Fig. 8d). The ALH uncertainty of 1 km  
 426 decreased AMFs with average AMF<sub>percent\_diff</sub> of -27% on the flight day. Especially, on the pixels where AOD > 0.6, the average  
 427 AMF<sub>percent\_diff</sub> was found to be -37% while that was -23% on the pixels where AOD < 0.4, showing the combined effect of  
 428 aerosol loading and aerosol profile shape on the NO<sub>2</sub> AMF calculations. Fig. 8e and 8f show the percentage difference error  
 429 between the calculated AMFs using the MODIS surface reflectance data with  $0.005 + 0.05 \times \text{SFR}$  lower (Fig. 8e) and  $0.005 +$   
 430  $0.05 \times \text{SFR}$  higher (Fig. 8f) values, respectively. The AMF decreased by about 6% when surface reflectance decrease, and vice  
 431 versa when it increased.

### 432 3.3 Validation of NO<sub>2</sub> VCDs retrieved from GeoTASO

433 Tropospheric NO<sub>2</sub> VCDs retrieved from GeoTASO L1B data (NO<sub>2,G</sub>) were compared with those obtained from OMI NO<sub>2</sub>  
 434 VCDs (NO<sub>2,O</sub>) and Pandora (NO<sub>2,P</sub>). The NO<sub>2,O</sub> were only available for 10 June during the campaign period. Therefore, we  
 435 only compared 37 NO<sub>2,G</sub> and NO<sub>2,O</sub> data points within a radius of 20 km and 30 min, which yielded a correlation coefficient  
 436 of 0.70 with a slope of 0.41 (Fig. 9 a)). In order to validate, All NO<sub>2,G</sub> within a radius 20 km of the OMI center coordinate were  
 437 averaged.

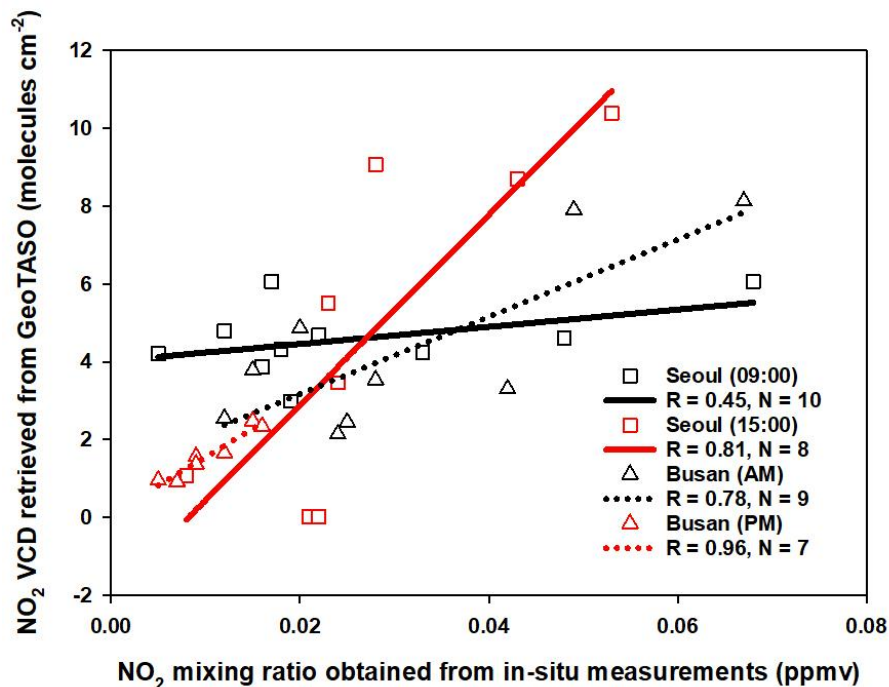
438 The NO<sub>2</sub> values are relatively low since GeoTAOS observation is carried out in a region with low NO<sub>2</sub> compared to Seoul  
 439 metropolitan and the overpass time of OMI is about 13:30 LT when NO<sub>2</sub> decreased. It is thought that the reson the low slope  
 440 value is because the OMI with low spatial resolution does not reflect the spatial NO<sub>2</sub> inhomogeneity in the pixel.



**Figure 9. Scatter plots of a) NO<sub>2</sub> VCD retrieved from GeoTASO and those obtained from OMI and b) NO<sub>2</sub> VCD obtained from Pandora and those retrieved from GeoTASO, respectively.**

To validate the accuracy of NO<sub>2,G</sub> data, we made a comparison with NO<sub>2</sub> VCD obtained from the Pandora system (NO<sub>2,P</sub>) during the KORUS-AQ campaign period. NO<sub>2,P</sub> obtained from Busan University, Olympic Park, Songchon, Yeosu, and Yonsei University Pandora sites on June 5, 9, and 10 were used for the GeoTASO validation (Fig. 1). NO<sub>2,G</sub> and NO<sub>2,P</sub> columns at these sites are compared in Fig. 10. In order to compare NO<sub>2,G</sub> and NO<sub>2,P</sub>, we used averaged NO<sub>2,G</sub> retrieved from 16 across track with smallest viewing zenith angle and averaged 30 min NO<sub>2</sub> obtained from Pandora measurement within a radius of 0.05 degree. NO<sub>2,G</sub> and NO<sub>2,P</sub> were correlated ( $R = 0.79$ , with a slope of 1.15), however, when NO<sub>2,P</sub> was lower than  $1 \times 10^{16}$  molecules cm<sup>-2</sup>, the correlation coefficient between NO<sub>2,G</sub> and NO<sub>2,P</sub> was  $< 0.1$ . The weak correlation at low NO<sub>2</sub> levels are most likely to reflect the differences in viewing geometries and the horizontal inhomogeneity of the measured NO<sub>2</sub> between Pandora and GeoTASO. Also, from this result, it is thought that it can be used for NO<sub>2</sub> validation of geostationary satellite such as GEMS using Pandora and GeoTASO. However, since the number of Pandora is limited in this campaign, we had difficulties to validate NO<sub>2</sub> retrieved from GeoTASO under various conditions. I believe that many ground-based remote sensing measurements are needed to validate GEMS under various conditions.





**Figure 10.** Scatter plot of the  $\text{NO}_2$  VCDs retrieved from GeoTASO, and  $\text{NO}_2$  surface mixing ratio obtained from Air-Korea. The black and red squares represent the  $\text{NO}_2$  data at 9 AM and 3 PM (local time) over the Seoul metropolitan region, respectively. The black and red triangles represent those in the morning and afternoon, over Busan, respectively.

To evaluate the spatiotemporal distribution of  $\text{NO}_2$  VCDs retrieved from GeoTASO,  $\text{NO}_{2,G}$  in comparisons to surface spatial patterns,  $\text{NO}_{2,G}$  was compared with  $\text{NO}_{2,A}$  for GeoTASO data within a radius of approximately 0.05 km and 30 min (Fig. 9). In order to compare  $\text{NO}_{2,G}$  and  $\text{NO}_{2,A}$ , we used averaged  $\text{NO}_{2,G}$  retrieved from 16 across track and averaged 30 min within a radius of 0.05 degree. Since in-situ measurements provides  $\text{NO}_2$  VMR ( $\text{NO}_{2,A}$ )(ppmv) once per hour,  $\text{NO}_{2,A}$  of the nearest time is used to compare with  $\text{NO}_{2,G}$ . The correlation coefficient (R) between  $\text{NO}_{2,G}$  (molecules  $\text{cm}^{-2}$ ) and  $\text{NO}_{2,A}$  at 9 AM and 3 PM LT in the Seoul metropolitan region was 0.45 and 0.81, respectively. When using only roadside station data from Air-Korea, the R-value for the morning increased to 0.83, which implies GeoTASO is more sensitive to emissions from  $\text{NO}_2$  source areas, such as roadsides. As a result of the comparison, there were large differences in the morning and afternoon. These results were identified because synoptic meteorology played an important role from June 1 to June 10, 2016 (Choi et al., 2019). As described by Judd et al. (2018), the spatial distribution for  $\text{NO}_2$  VCDs appears that reflects the emission source in local industrialized regions and transportations in the morning with relatively weak winds. In general,  $\text{NO}_2$  concentration increases to late morning, indicating that the emissions process proceeds faster than the  $\text{NO}_2$  removal process. As the planetary boundary layer heights (PBLH) in early afternoon increase and surface  $\text{NO}_2$  is mixed through a deeper PBLH, the  $\text{NO}_2$  VCDs distribution showed a wider increase in most of the Seoul metropolitan area and the overall column amounts continue to increase (Judd et al., 2018).

480 In addition, when comparing NO<sub>2</sub> VCDs with surface NO<sub>2</sub> concentrations, it should be interpreted carefully that it is a non-  
481 linear relationship between NO<sub>2,G</sub> and NO<sub>2,A</sub>. Although it may vary depending on weather conditions, high NO<sub>2</sub> VCDs from  
482 airborne observations may sometimes be detected with low surface NO<sub>2</sub> concentrations. In particular, when exhaust gases  
483 emitted from industrial facilities are happen at a certain altitude (stacks/chimneys), NO<sub>2,G</sub> show high NO<sub>2</sub> VCDs, but NO<sub>2,A</sub>  
484 may be observed to have a low concentration. Unfortunately, in Anmyeon industrial region, NO<sub>2,G</sub> and NO<sub>2,A</sub> could not be  
485 compared due to spatial restrictions because the distribution of ground observation stations is concentrated in metropolitan  
486 areas.

487 In the Busan metropolitan area, the R-value of the NO<sub>2,G</sub> and NO<sub>2,A</sub> data had a correlation coefficient greater than 0.78. This  
488 reflects the more even horizontal distribution of NO<sub>2</sub> in the afternoon, when diffusion from the source areas had taken place.  
489 However, for a more accurate comparison, NO<sub>2</sub> VCD data should be converted to NO<sub>2</sub> MR based on mixing layer height,  
490 temperature, and pressure profile data (Kim et al., 2017; Qin et al., 2017; Jeong and Hong, 2021a). However, since the number  
491 of pandora and satellite data is limited in this campaign, we had difficulties to validate NO<sub>2</sub> retrieved from GeoTASO under  
492 various conditions. Since ground-based, airborne and space borne remote sensing measurements has their own advantage and  
493 disadvantage, I believe that a comprehensive observation campaign involving all of groud-based, airborne and space borne  
494 measurements should be carried out continuously for upcoming new era of geostationary environmental satellite.

#### 495 **4 Conclusions**

496 For the first time, we have retrieved NO<sub>2</sub> VCD data using airborne GeoTASO observations over the Seoul metropolitan  
497 region—one of the most populous cities worldwide, the Busan metropolitan region—the second-largest city in South Korea,  
498 and Anmyeon, with thermal power plants and industrial complexes. By retrieving NO<sub>2</sub> data using GeoTASO L1B radiance, it  
499 was possible to observe the spatial distribution of NO<sub>2</sub> over these metropolitan and industrial regions. In the morning,  
500 tropospheric NO<sub>2</sub> VCD over Seoul showed a strong horizontal gradient between rural and urban areas. In urban areas,  
501 tropospheric NO<sub>2</sub> VCD was high, with values exceeding  $3 \times 10^{16}$  molecules cm<sup>-2</sup>; in rural areas, values were typically below  
502  $1 \times 10^{16}$  molecules cm<sup>-2</sup>. Extremely high values over  $10 \times 10^{16}$  molecules cm<sup>-2</sup> were also observed in both rural and urban  
503 areas. In Anmyeon, GeoTASO observations showed NO<sub>2</sub> is mainly emitted from the chimneys of industrial complexes and  
504 thermal power plants, and subsequently transported by wind approximately 30 km to the Yellow Sea of the west coast of the  
505 Korean Peninsula. In the Busan metropolitan region, in the morning, tropospheric NO<sub>2</sub> VCDs showed a similar pattern to the  
506 Seoul metropolitan region, with high values above the inter-city road. However, in contrast to Seoul, tropospheric NO<sub>2</sub> VCDs  
507 in Busan decreased in the afternoon due to different weather conditions locally.

508 To compare the data retrieved from the GeoTASO system, we compared NO<sub>2,G</sub> with NO<sub>2,O</sub> obtained from the OMI, NO<sub>2,A</sub>  
509 obtained from Air-Korea, and NO<sub>2,P</sub> obtained from the Pandora observation system. When the distance between two  
510 observations was below 20 km or 0.05 degree within 30 min, the correlation coefficients were relatively high (R = 0.70, and



79, respectively). However, the correlation between  $\text{NO}_{2,G}$  and  $\text{NO}_{2,A}$  over the Seoul metropolitan region was weak ( $R = 0.45$ ) in the morning because of the more pronounced  $\text{NO}_2$  horizontal gradient. The GeoTASO system successfully observed  $\text{NO}_2$  VCDs with a high horizontal spatial resolution for both metropolitan and industrial regions. This demonstrates that airborne remote sensing measurements from GeoTASO, similar to GCAS, APEX and others, can be a very effective tool for the validation of trace gases retrieved from environmental satellites, including the OMI, TROPOMI, and GOME-2; these systems can obtain high-resolution measurements over relatively wide areas. However, to validate geostationary environmental satellites with higher spatiotemporal resolutions, such as the GEMS, TEMPO, and sentinel-4, additional validation strategies are needed. Based on error estimation, it can be concluded that aerosol properties are relevant and should be determined and  $\text{NO}_2$  vertical profile retrieval performed using, for example, LIDAR, MAX-DOAS, and sondes. This is important because the accuracy of aerosol properties and the  $\text{NO}_2$  vertical profiles affects the accuracy of AMF calculations (Leitão et al., 2010; Hong et al., 2017; Lorente et al., 2017; Boersma et al., 2018). Furthermore, as we observed in the Seoul metropolitan area, more closely spaced observations using ground-based remote sensing systems and in situ measurements are needed as  $\text{NO}_2$  displays large horizontal gradients, especially in the morning.

#### **Author contributions**

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

#### **Competing interests**

The authors declare that they have no conflict of interest.

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