- 2 Sensitivity of the WRF-Chem v4.4 ozone, formaldehyde, and their precursors
- 3 simulations to multiple bottom-up emission inventories over East Asia during the
- 4 KORUS-AQ 2016 field campaign

- 6 Kyoung-Min Kim¹, Si-Wan Kim^{2*}, Seunghwan Seo¹, Donald R. Blake³, Seogju Cho⁴,
- 7 James H. Crawford⁵, Louisa Emmons⁶, Alan Fried⁷, Jay R. Herman^{8,9}, Jinkyu Hong¹,
- 8 Jinsang Jung¹⁰ Gabriele Pfister⁶, Andrew J. Weinheimer⁶, Jung-Hun Woo¹¹, and Qiang
- 9 Zhang¹²

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- ¹Department of Atmospheric Sciences, Yonsei University, Seoul, South Korea
- ²Irreversible Climate Change Research Center, Yonsei University, Seoul, South Korea
- ³Department of Chemistry, University of California at Irvine, Irvine, CA, US
- 14 ⁴Seoul Metropolitan Government Research Institute of Public Health and
- 15 Environment, Gyeonggi-do, South Korea
- ⁵NASA Langley Research Center, Hampton, VA, US
- ⁶National Center for Atmospheric Research, Boulder, CO, US
- ⁷Institute of Arctic and Alpine Research, University of Colorado, Boulder, CO, US
- 19 ⁸NASA Goddard Space Flight Center, Greenbelt, MD, US
- ⁹University of Maryland Baltimore County, Baltimore, MD, USA
- 21 ¹⁰Korea Research Institute of Standards and Science, Daejeon, South Korea
- 22 ¹¹Department of Advanced Technology Fusion, Konkuk University, Seoul, South
- 23 Korea
- ¹²Department of Earth System Science, Tsinghua University, Beijing, China

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*To whom correspondence should be addressed. E-mail: siwan.kim@yonsei.ac.kr

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1 Abstract

2 In this study, the WRF-Chem v4.4 model was utilized to evaluate the sensitivity of O₃ 3 simulations with different three bottom-up emission inventories (EDGAR-HTAP v2, v3, and KORUS v5) using surface and aircraft data in East Asia during the Korea-4 United States Air Quality (KORUS-AQ) campaign period in 2016. All emission 5 6 inventories were found to reproduce the diurnal variations of O₃ and its main precursor 7 NO₂, which is one of the major precursors of O₃, as compared to the surface monitor data. However, the spatial distributions of the daily maximum 8-hour average (MDA8) 8 9 O₃ in the model do not completely align with the observations. The model MDA8 O₃ had a negative (positive) bias north (south) of 30°N over China. All simulations 10 underestimated the observed CO by 50-60% over China and South Korea. In the Seoul 11 Metropolitan Area (SMA), EDGAR-HTAP v2, v3, and KORUS v5 simulated the 12 13 vertical shapes and diurnal patterns of O₃ and other precursors effectively, but the model underestimated the observed O₃, CO and HCHO concentrations. Notably, the model 14 15 aromatic VOCs were significantly underestimated with the three bottom-up emission 16 inventories, although the KORUS v5 shows improvements. The model isoprene (ISO) estimations had a positive bias relative to the observations, suggesting that the Model 17 18 of Emissions of Gases and Aerosols from Nature (MEGAN) version 2.04 overestimated 19 isoprene isoprene ISO emissions. Additional model simulations were conducted by 20 doubling CO and VOC emissions over China and South Korea to investigate the causes of the model O₃ biases and the effects of the long-range transport on the O₃ over South 21 22 Korea. The doubled CO and VOC emission simulations improved the model O₃ simulations for the local emission dominant case, but led to the model O₃ 23

- 1 overestimations for the transport dominant case, which emphasizes the need for
- 2 accurate representations of the local VOC emissions over South Korea.

1. Introduction

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2 Air pollutants not only harm human health but also affect radiative balance, resulting 3 in climate change (Anenberg et al., 2018; Franklin et al., 2015; Lee et al., 2014; Manning and von Tiedemann, 1995; Rosenzweig et al., 2008; Wild et al., 2001). 4 Anthropogenic activities are the primary source of air pollutant emissions, which have 5 6 significant temporal and spatial variability. Chemical transport models (CTMs) use 7 bottom-up emission data to simulate ambient concentrations of air pollutants. CTMs then process these emissions, tracking their impact through chemistry, transport, and 8 9 loss through deposition (Zhong et al., 2016). Therefore, sensitivity evaluations of CTMs 10 to anthropogenic emission data are an essential part of atmospheric modeling research. 11 Several bottom-up emission inventories are available for chemical modeling of 12 Asia, including the Multi-resolution Emission Inventory for China (MEIC), Regional Emission inventory in Asia (REAS), and Emissions Database for Global Atmospheric 13 14 Research-Hemispheric Transport of Air Pollution (EDGAR-HTAP). Since 2010, Tsinghua University has developed the high-resolution MEIC emission inventory for 15 China and updated the data to the v1.3, providing anthropogenic emissions by sector 16 and species from 2008 to 2017 (Zheng et al., 2018). REAS provides emission data in 17 Asia from 1950 to 2015 (Kurokawa and Ohara, 2020). In Europe, EDGAR-HTAP has 18 been developed and widely used for CTM simulations from global to regional scale 19 20 (Kim et al., 2021; Sharma et al., 2017; Sicard et al., 2021). Recently, EDGAR-HTAP v3 has been published, covering 19 years from 2000 to 2018 compared to only two 21 years (2008 and 2010) in the version 2 data (Crippa et al., 2023). Zhong et al. (2016) 22 23 compared REAS with EDGAR in July, 2007 over China, while Saikawa et al. (2017) compared 5 emission inventories including REAS, EDGAR, MEIC in China, without 24

- 1 validation. As bottom-up emission inventories are continuously updated for recent years,
- 2 there is an ongoing need to evaluate new emissions data.
- 3 The Ministry of Environment (MOE) in South Korea and National Aeronautics and Space Administration (NASA) in the U.S. conducted the Korea-United States Air 4 5 Quality (KORUS-AQ) campaign in May-June 2016. The campaign provided a variety 6 of data sets, including ground-based and airborne observations, useful for the validation 7 of model simulations. The KORUS emissions, developed by Konkuk University, were used by many modeling teams to simulate the air pollutant concentrations during the 8 9 campaign period. Numerous modeling studies wereare conducted forduring this period including validations of CTM results with diverse observation datasets. Miyazaki et al. 10 (2019) adjusted emission inventories using various satellite data sets and Model for 11 Interdisciplinary Research on Climate with chemistry (MIROC-Chem), resulting 12 13 improved simulations of tropospheric O₃. Goldberg et al. (2019) reported underestimations of NOx emissions in South Korea, particularly in Seoul. Souri et al. 14 (2020) also revealed the same issue in South Korea and conducted analysis of the 15 sensitivity of O₃ formation to adjustments in NOx and volatile organic compound (VOC) 16 17 emission derived from inverse modeling. Tang et al. (2019) revealed negative biases of simulated CO concentrations in East Asia by utilizing satellite data and the Community 18 19 Atmosphere Model with Chemistry (CAM-Chem). Choi et al. (2022) modified 20 anthropogenic VOC emissions through the inverse modeling using satellite HCHO observations with the Goddard Earth Observing System with Chemistry (GEOS-Chem), 21 22 which reduced O₃ and HCHO biases. 23 NowRecently, the most updated version of bottom-up emission inventories and

CTMs have become available for the air pollution modeling studies in East Asia. Earlier

research pointed out the prevailing issues associated with diverse bottom-up emission 1 2 inventories. In this study, we selected the EDGAR-HTAP versions 2 and +3, and 3 KORUS version 5 emission data and used the Weather Research and Forecasting model coupled with Chemistry (WRF-Chem) version 4.4 for intercomparison of the three 4 emissions data sets, aiming to elucidate the potential issues understand the status of -of 5 6 precursor emissions from bottom-up emission inventories and their uncertainties, which 7 that may could impact the O₃ formations in the model. Our analysis focused on O₃ and 8 its major precursors were selected were selected for model evaluation and the model 9 results were evaludated with surface observation data in China and South 10 Korea and aircraft data acquired over the South Korean peninsula and surrounding 11 waters. The manuscript is organized as follows. The data and methods section introduces 12 13 emission inventories, the numerical model, and meteorological and chemical 14 observations. The results section evaluates the model's meteorology and chemistry using routine surface observations over China and South Korea. Subsequently, the 15 model results employing three bottom-up emission inventories are compared with 16 sophisticated chemical observations obtained during the KORUS-AQ field campaign, 17 primarily over South Korea. This comparison summarized the model's performance 18 19 with each emission inventory. In the discussion section, strategies to enhance surface O₃ simulations, along with accurate precursor simulations, are proposed based on 20 various emission scenarios for urban and regional areas over China and South Korea. 21 The summary and conclusion section follow, providing overview of the key findings 22 23 and conclusions drawn from the study.

2. Data and Methods

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2.1. WRF-Chem model configurations

Oceanic and Atmospheric Administration (NOAA) and National Center for 4 Atmospheric Research (NCAR), to simulate meteorological variables and chemical 5 species in the atmosphere (Grell et al., 2005). The WRF-Chem v4.4 includes N₂O₅ 6 7 heterogeneous chemistry that consists of several chemical reactions related with ClNO₂ 8 and N₂O₅ reactions, resulting in nitrate aerosol. The reactions are incorporated in Secondary Organic Aerosol-Volatility Basis Set (SOA-VBS) with Regional 9 Atmospheric Chemistry Mechanism (RACM) chemistry option (chem = 108) in WRF-10 11 Chem (Li et al., 2016). 12 We set 59 vertically customized eta (η) levels as vertical layers. The model's first layer height is approximately 40 m above ground level for the entire domain. The 13 model's vertical layers are designed to include about 17 layers under 1.5 km to simulate 14 planetary boundary layer chemistry and near surface vertical distribution in detail. The 15 horizontal resolution is 28 x 28 km². The simulations in this study start at 12 UTC on 16 April 24 and end at 12 UTC on June 11. The model meteorology restarts every 12 UTC 17 (9 PM local time in South Korea) to minimize numerical errors. After the first 7 days 18 19 of model initiation (spin-up), we analyzed the model results from May 1 to June 10. We used China standard time (+8 UTC) and Korea standard time (+9 UTC) for evaluations 20 with observations. The model physics, chemistry, and aerosol schemes are summarized 21 22 in Table S1 with corresponding references. The Global Forecast System (GFS) Final 23 (FNL) analysis data are used for meteorological input and boundary conditions. The Community Atmosphere Model with Chemistry (CAM-Chem) output is used for 24

In this study, we utilized the WRF-Chem v4.4, which was developed by the National

- 1 chemical boundary conditions (https://www.acom.ucar.edu/cam-chem/cam-
- 2 <u>chem.html</u>https://rda.ucar.edu/datasets/ds313.7/) (Buchholz et al., 2019; Emmons et al.,
- 3 2020). We used the Model of Emissions of Gases and Aerosols from Nature (MEGAN)
- 4 v2.04 to calculate biogenic emissions (Guenther et al., 2006). We did not account for
- 5 fire emissions because of small impact on air quality simulations during the KORUS-
- 6 AQ campaign period (Park et al., 2021). In our sensitivity simulation with the Fire
- 7 INventory from NCAR (FINN) v2.5 fire emissions (Wiedinmyer et al., 2022), a
- 8 marginal increase in the simulated averaged daily maximum 8-hour average (MDA8)
- 9 O₃ of approximately 1 ppbv (1.6 %) was noted in China (Park et al., 2021)(Supporting
- 10 <u>information, Figure S1</u>).

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12 **2.2.** The model simulations using different anthropogenic emissions

2.2.1. Bottom-up emission data

- 14 EDGAR-HTAP v2, v3, and KORUS v5 anthropogenic bottom-up emissions inventories
- are compared with respect to their spatial distribution and total amount in Figure 1 and
- Table S23. We applied the same diurnal factor for all three emissions data by species,
- 17 following the diurnal patterns for the Los Angeles Basin as in Kim et al. (2016) (also
- 18 see Figure S_{24}).
- 19 EDGAR-HTAP v2 provides 2-dimensional emissions of CH₄, CO, SO₂, NOx (NO
- + NO₂), total non-methane volatile organic compound (NMVOC), NH₃, PM₁₀, PM_{2.5},
- BC, and OC in 2008 and 2010 with a horizontal resolution of 0.1° x 0.1°. We used 2010
- data since it is the most recent data available. The data are partitioned by each sector
- 23 and its sources such as air, ships, energy, industry, transport, residential, and agriculture

(https://edgar.jrc.ec.europa.eu/dataset htap v2). For East Asia, it included data from 1 the Model Inter-Comparison Study for Asia (MICS-Asia) and REAS v2.1. In South 2 Korea, it adopted data from the Clean Air Policy Support System (CAPSS) (Janssens-3 Maenhout et al., 2015), and the underlying emission data had an original horizontal 4 resolution of 0.25° x 0.25° over East Asia, which is resampled to 0.1° x 0.1° resolution 5 by raster resampling and aggregation. The specifically mapped speciated EDGAR-6 7 v2VOC data were obtained through the WRF-Chem (https://www.acom.ucar.edu/wrf-chem/download.shtml) in the anthro emiss program 8 with the Model for Ozone and Related chemical Tracers (MOZART) species (Supporting Information, Table S3). The anthro emiss program converts the EDGAR-10 11 HTAP v2 data into 28 x 28 km² grid by the RACM chemical species (Supporting 12 <u>Information, Table S4</u>). It mapped the MOZART volatile organic compounds (VOC) 13 species into the RACM VOC species (See the detailed equations in Supporting Information, Table S54) (Li et al., 2014; Emmons et al., 2010). 14

The EDGAR-HTAP v3 is extended to much longer time scale than the previous version EDGAR-HTAP (v2). The EDGAR-HTAP v3 covers 2000 to 2018 with a more detailed horizontal resolution (https://edgar.jrc.ec.europa.eu/dataset_htap_v3) (Crippa et al. 2023). While EDGAR-HTAP v2 uses MICS-Asia, only the REAS data are used in China and India in the EDGAR-HTAP v3. It adopts the CAPSS-Konkuk University (CAPSS-KU) data for South Korea and emission data provided by the Japanese government for Japan. We chose the data for 2016, according to the KORUS-AQ campaign period. Because the original EDGAR-HTAP v3 data provide VOC as total NMVOC with the unit of ton/month, we distributed the total NMVOC to MOZART VOC species with the ratio of each VOC species to total NMVOC from EDGAR-HTAP

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- 1 v2 in anthro emiss program. Then, the assigned EDGAR-HTAP v3 data were again
- 2 converted to the RACM.
- The KORUS v5 emission data represent 2016 in China and 2015 in other regions.
- 4 The Comprehensive Regional Emissions Inventory for Atmospheric Transport
- 5 Experiment (CREATE) v2.3 data from 2015 were used and the ship emissions from
- 6 CAPSS were added near the coastal region in South Korea (Jang et al., 2020; Woo et
- 7 al., 2012). The CREATE is originally developed by combining REAS, MEIC, Japan
- 8 Auto-Oil Program emission inventory (JATOP), and Korean Clear Air Policy Support
- 9 System (CAPSS). The NMVOC species from KORUS v5 were mapped following the
- 10 Statewide Air Pollution Research Center (SAPRC-99) mechanism, and we also
- assigned the SAPRC-99 species to RACM (Carter, 2000) (Supporting information,
- 12 Table S<u>5-6</u>4).
- Figure 1 shows the spatial distribution of NO, CO, and <u>TOL</u> (toluene <u>+ less reactive</u>
- 14 aromatics defined in RACM, see Table S4) emissions in May for each inventory. The
- NO_X emissions were assumed to be emitted as NO. The major cities in China and South
- 16 Korea had relatively high NO, CO, and TOL (toluene and less aromatics defined in
- 17 RACM, see Table S2) emissions, which are major precursors affecting O₃ formation.
- 18 We define three boxes representing Eastern China, South Korea, and the Seoul
- metropolitan area (SMA) and calculated the emissions (see Table S23). In South Korea
- 20 including SMA, EDGAR-HTAP v3 had the largest NO_X emission among the emission
- 21 inventories. The KORUS v5 has lower NO_X emissions in Eastern China by 46% and
- 22 39% compared to EDGAR-HTAP v2 and v3, respectively. The CO emission was the
- lowest in EDGAR-HTAP v2 in South Korea, being 56% (69%) lower than that in
- 24 KORUS v5 (EDGAR-HTAP v3). KORUS v5 showed the highest CO emissions in SMA

- 1 though EDGAR-HTAP v3 showed more CO emissions in South Korea. However,
- 2 KORUS v5 had the smallest CO emissions in China, being 7% (9%) lower than that in
- 3 EDGAR-HTAP v2 (v3). The TOL emission from KORUS v5 is higher than those from
- 4 EDGAR-HTAP v2 (EDGAR-HTAP v3) by 176% (98%) in China. The relative
- 5 difference of TOL between KORUS v5 and EDGAR-HTAP v2 (EDGAR-HTAP v3) is
- 6 larger in South Korea by 263%. On the other hand, EDGAR-HTAP v3 have the largest
- 7 total NMVOC emissions over China than EDGAR-HTAP v2 and KORUS v5 by 38 and
- 8 27 %, respectively. These discrepancies of VOC emissions may lead to a change in the
- 9 NOx/VOC-sensitive regime and O₃ production efficiency. The sensitivity of O₃
- 10 formation to NOx emission has discrepancies by its regime, which will be further
- 11 discussed in section 3.2.

13 **2.2.2. The model experiments**

- 14 The model experiments are summarized in Table 1. The simulations using EDGAR-
- 15 HTAP v2, v3, and KORUS v5 emissions are named as EDV2, EDV3, and KOV5,
- respectively. In this study, we found consistent underestimation of CO and VOC,
- 17 HCHO, TOL, and XYL for all emissions by -40% (\pm 2%) and, -25% (\pm 1%) (HCHO),
- $\frac{-67\% (\pm 21\%), -53\% (\pm 18\%), respectively, compared to DC-8 in South Korea. Here}{}$
- 19 TOL and XYL are lumped species including toluene and xylene, respectively. This is
- 20 in line with the results reported by Park et al. (2021), who found that almost every
- 21 model underestimated CO. Underestimation of CO in East Asia is a well-known feature
- revealed by many studies. For example, Gaubert et al. (2020) mentioned that CAM-
- 23 Chem underestimates CO during the KORUS-AQ campaign period and presented a CO

- 1 compensation method utilizing data assimilation with CO observations. Wada et al.
- 2 (2012) pointed out that EDGAR v4.1 underestimates anthropogenic CO emissions in
- 3 China by 45% compared to observation-based estimations of CO emissions. Moreover,
- 4 underestimation of VOC is also found for all anthropogenic emission inventories.
- 5 Kwon et al. (2021) estimated top-down emissions of anthropogenic VOCs utilizing
- 6 Geostationary Trace gas and Aerosol Sensor Optimization spectrometer (GeoTASO).
- 7 They found that top-down VOC emissions were up to 6.9 times higher than bottom-up
- 8 emissions (KORUS v5).
- 9 WithFor all emission inventories, O₃ is underestimated at most ground-based observation sites in South Korea. Therefore, tTo figure out the potential issue that may 10 affect causes of negative biases of O₃ in South Korea, we conducted threewo additional 11 model simulations using EDGAR-HTAP v3 that shows the lowest bias of O3 12 13 concentrations compared to DC-8 (-14.2 ppb) than EDGAR-HTAP v2 (-16.9 ppb) and 14 KORUS v5 (-18.1 ppb) over the SMA; the mean biases are -16.9, -14.2, and -18.1 ppb with EDV2, EDV3, and KOV5, respectively. : oOneTwo simulations are is with twice 15 16 the anthropogenic CO and VOC emissions in China (EDV3 Ch2) and South Korea 17 (EDV3 Ko2), respectively, and the third simulation other simulation uses double CO 18 and VOC emissions in both China and South Korea (EDV3 ChKo2) to investigate 19 possible improvements in the simulated O₃ and CO from these emission changes. To propose the strategies to improve surface O₃ simulations over China and South Korea, 20 21 we incorporated 4 additional emission scenarios involving the reduction of NO_x and/or VOC emissions over China. Specifically, we considered the cases with a 50% reduction 22 in NO_x emissions only, a 50% reduction in VOC emissions only, a simultaneous 50% 23

- 1 reduction in both NO_x and VOC emissions, and a 75% reduction in NO_x emissions only.
- 2 For more details, refer to Section 4 (Discussion).

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2.3. Observations

2.3.1. Meteorological data

- 6 The meteorological field that WRF-Chem reproduced is <u>evaluated</u> with the
- 7 surface synoptic observation (SYNOP) data operated by the World Meteorological
- 8 Organization (WMO) (http://www.meteomanz.com). Surface temperature, relative
- 9 humidity, and surface wind speed are adopted for model validation. As the SYNOP data
- are provided every 3 or 6-hourly, we selected model data when the observation data are
- available. There were 271 sites in China-Taiwan-Hongkong and 48 sites in South Korea.

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2.3.2. Ground-based observations

- 14 The surface observation network used in this study was obtained from Airkorea in South
- 15 Korea and the China Ministry of Ecology and Environment (MEE) in China. The
- Airkorea observation network provides 1-hourly measurements of NO₂, SO₂, CO, O₃,
- 17 PM₁₀, and PM_{2.5} at suburban, background, roadside, city, and port sites
- 18 (www.airkorea.or.kr). The concentrations of NO₂, CO, and O₃ are measured using the
- 19 chemiluminescent, non-dispersive infrared, and ultraviolet photometric methods,
- 20 respectively. In South Korea, there are indications of positive biases in NO₂ surface
- observations, potentially resulting in overestimations of 2~308.9%, particularly at
- 22 <u>suburban sites in spring (Jung et al., 2017).</u> The model data with 28 x 28 km² horizontal

- 1 resolution were linearly interpolated to the 365 sites in South Korea, and we selected
- 2 NO₂, O₃, and CO for model validation.—
- 3 The Chinese observations were provided by MEE through the website
- 4 (beijingair.sinaapp.com). Surface NO₂ over China was measured using a molybdenum
- 5 converter, which has the potential for positive biases due to other NO₂-related oxidation
- 6 products (Dunlea et al., 2007). In South Korea, the positive biases exist regarding NO₂
- 7 surface observations, which could to overestimations of 28.9% at suburban sites in
- 8 spring (Jung et al., 2017). CO was measured using infrared absorption (Zhang and Cao.,
- 9 2015), and there were 1454 stations in China during the campaign period.

Research Institute of Public Health and Environment (SIHE).

For validation of NO₂ and HCHO vertical column density, data from the Pandora spectrometer were used, which the model reproduced with emission inventories at the Olympic Park site (37.5232°N, 127.126°E). The HCHO data from Pandora is corrected because of internal off-gasing to avoid positive biases (Spinei et al., 2021). At the same observation site, surface NO₂ was also measured by a KENTEK NOx analyzer with photolytic method, and surface O₃ was measured using the same instrument. Ground-based HCHO was measured using Aerodyne QCL. We compared the observed diurnal cycle of vertical column and surface concentrations of NO₂ and HCHO with the model simulations utilizing EDV2, EDV3, and KOV5. We also used ground-based VOC data from gas chromatography flame ionization detector (GC-FID) operated by the Seoul

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2.3.3. Aircraft data

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The DC-8 research aircraft, operated by NASA, performed multiple flight 2 3 measurements with a variety of measuring instruments. We utilized 1 minute interval merged data of O₃, NO₂, CO, HCHO, and VOC along the 20 flight paths (Figure 27). 4 The nearest WRF-Chem grid is selected and then temporally and vertically interpolated 5 6 to the aircraft data using linear interpolation method to fully utilize the observations. 7 Atmospheric NO₂ and O₃ concentrations were measured using a 4-channel chemiluminescence instrument, with an uncertainty of 100 pptv + 30% and 5 ppbv + 8 9 10%, respectively. CO concentrations were observed using a diode laser spectrometer, with an uncertainty of 2% or 2 ppbv. The Compact Atmospheric Multi-species 10 Spectrometer (CAMS) was used to measure HCHO concentration, with a possible 3% 11 systematic error (Richter et al., 2015). We also utilized data from the Whole Air Sampler 12 (WAS) to analyze VOC species from different emission inventories (Colman et al., 13 14 2001). In this study, we focused on DC-8 observations below a height of 2 km to concentrate on planetary boundary layer (PBL) chemistry. The observation height was 15 determined by GPS altitude above ground level. 16

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3. Results

19 3.1. The model meteorology simulations

The model temperature and relative humidity were compared with surface observations in China and South Korea. The model-simulated temperature had a slight negative mean bias of -0.91 °C (correlation coefficient R = 0.90) in China, with the largest negative bias in southwestern China. In South Korea, the mean bias was -1.71 °C (R = 0.88). The

simulated relative humidity showed a negative bias of -20 to -10% in the North China 1 2 Plain (NCP) area and a positive bias of 10 to 20% in southwestern China. There was a negative bias of relative humidity over the west coastal area and a positive bias of 10 to 3 20% at most observation stations in South Korea. The correlation coefficients between 4 5 the model relative humidity and observations were 0.85 and 0.76 for China and South Korea, respectively. Overall, the comparisons showed decent model simulations of 6 7 meteorology. A negative temperature bias could result in a reduction of isoprene 8 emissions, as illustrated in Figure S3 of the Supporting Information, compared to the 9 estimates based on accurately simulated temperature (Figure S3). Even if there is negative bias of temperature, leading to a -3.3 % bias in actually emitted emissions, the 10 averaged impact on emissions in East Asia is negligible by 0.4 % (Supporting 11 12 Information, Figure S3), During the KORUS-AQ campaign period, WRF-Chem accurately simulated the 13 daytime PBL height from a laser ceilometer (CL-31, Vaisala Inc., Finland) observed at 14 Yonsei University in Seoul, South Korea (Lee et al., 2019). But, Travis et al. (2022) has 15 indicated the possibility of PBL height underestimations by CTM. Furthermore, due to 16 17 limitations of the instrument, the ceilometer has potential to inadequately estimate nighttime PBL height. It is primarily attributed to the method based on aerosol gradients 18 19 (Jordan et al., 2020). Therefore, the interpretation of simulated nighttime concentrations 20 of air pollutants should be approached with caution. More analysis of meteorological

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and Figure S42-S53).

fields, including PBL height, can be found in the Supporting Information (Table S75)

3.2. Evaluations with routine surface chemical observational data

- 2 The study compared simulated concentrations of O₃, NO₂, and CO with data from
- 3 routine surface observational networks (Table 2 and Figure 32-76). First, the diurnal
- 4 variations of the model O₃ using different emissions inventories were compared with
- observations for each subregion (Table 2 and Figure 32). Overall, all emissions
- 6 inventories successfully reproduced diurnal variations and absolute values of O₃ for
- 7 most regions, but there were notable discrepancies in several regions.
- 8 —In the North China Plain (NCP) region, EDV2 led to a negative model O₃ bias (-
- 9 12 ppb) with R=0.65, while EDV3 and KOV5 simulated O₃ better with reduced biases
- and increased correlations (R=0.68-0.71). The high NOx emissions are higher
- than relative to the VOC emissions in NCP, leadingled –to a low formaldehyde to NO2
- 12 ratio (FNR) (<10.7), suggesting that the NCP area is in a VOC-limited regimeegion
- withfor all emission inventories (Table 3). Due to the elevated reactive VOC emissions
- in EDV3 and KOV5 compared to EDV2, both EDV3 and KOV5 show improved O₃
- simulations. Similarly, EDV2 had a negative O₃ bias (-17 ppb) with R=0.62 in the
- 16 Yangtze River Delta (YRD) area, but EDV3 and KOV5 much improved the simulations,
- which was also observed in the Northeastern China (NEC) area. However, the model
- 18 O₃ concentrations based on the three emission inventories were overestimated in the
- 19 Sichuan-Chongqing-Guizhou (SCG) and Southeastern China (SEC) area. In SCG and
- 20 SEC, the WRF-Chem simulated higher biogenic isoprene emissions compared to
- 21 anthropogenic TOL and XYL emissions by up to a factor of 10, leading to a high FNR
- 22 (>1). -In Pearl River Delta (PRD), EDV2 showed the lowest bias (-0.3 ppb) compared
- 23 to EDV3 and KOV5 because EDV3 and KOV5 have elevated anthropogenic VOC
- 24 emissions as well as enhanced biogenic isoprene emissions under a VOC-limited

- 1 regime (Table 3). In the suburban area of Northern China (NOC), all emission
- 2 inventories reasonably simulated hourly O₃ concentrations.
- 3 Averaged O₃ was well simulated in South Korea (KOR) with low biases (-1 to 0.7
- 4 ppb), but a negative bias appears over the Seoul metropolitan area (SMA) with all
- 5 emissions (-5.5 to -3.5 ppb) (Table 2). The WRF-Chem simulations indicated SMA as
- 6 a highly NOx-saturated region (FNR < 0.2), resulting in being extremely VOC-sensitive
- 7 chemistry regime for O₃ production. The uThis nderestimated model O₃ levels in this
- 8 region suggests implies the possibility of deficient insufficient anthropogenic VOC
- 9 emissions in SMA across for all emission inventories (Table 3). A detailed This will be
- 10 <u>more analyzed</u>discussion will be provided <u>-in section 3.3.</u>

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The study also analyzed the mean values of daily maximum 8 hour average (MDA8) O₃ concentration at each site and their spatial distributions for the entire campaign period (Figure 43). The spatial distributions of the model MDA8 O₃ were not well correlated with those of the observations. But, notable disparities were observed in simulating MDA8 O₃ when the different emissions were used. For the north and eastern part of China including Beijing and Shanghai, large negative biases disappear when using EDV3 and KOV5. KOV5 only shows a significant correlation with the surface MDA8 O₃ observations (including 929 sites) than EDV2 and EDV3 in China (0.43 versus 0.01, 0.20). The correlations between the time series of the model MDA8 O₃ and observations varied at each site, with about 40-60% of sites (depending on the emission inventories) showing a correlation coefficient greater than 0.6 (see Supporting Information, Figure S46), and the locations of these sites were scattered. The correlation slightly improved with hourly O₃ concentrations instead of MDA8 O₃, with about 50-60% of sites having a correlation coefficient greater than 0.6 (Supporting Information,

Figure S64). For this metric, high correlations occurred in pollution hot spots north of 30°N and the South Coast of China, in which the ratio of HCHO to NO₂ (FNR) was much less than 1, suggesting VOC-limited/NOx-saturated chemical regime (Supporting Information, Figure S75). The model MDA8 O₃ were underestimated for the pollution hot spots with a low HCHO to NO₂ ratio located north of 30°N, suggesting a possibility of model underestimations of anthropogenic VOC emissions causing model MDA8 O₃ biases at these sites. In contrast, the simulated MDA8 O₃ was generally overestimated for sites south of 30°N in which HCHO concentrations were high (Supporting Information, Figure S75). Zhang et al. (2020) reported that simulated biogenic isoprene (ISO) from MEGAN was overestimated compared to observation sites under 35°N in China.

The EDV2 and EDV3 showed a positive NO₂ bias over the YRD, NCP, and PRD regions, which include large cities in China (Table 2 and Figure 54-65). On the other hand, EDV2 and EDV3 had smalllow negative NO₂ biases in the NEC and NOC regions. during the daytime, which implies underestimation of NOx emissions in background sites (Figure 54). While EDV2 simulates NEC and NOC as NOx saturated regime. EDV3 and KOV5 shows higher FNR indicating transition and VOC saturated regime. respectively. As the lower NOx change the chemical regime from NOx saturated to VOC saturated with KOV5, higher VOC emissions resulted in the lowest O₃ biases in NEC and NOC. However, there are still some underestimations of O₃ even though EDV3 and KOV5 are used. Chen et al. (2022) reported that soil NOx could increase O₃ concentrations in NEC and NOC about 2-10 ppb. All models demonstrated reasonable NO₂ model performance in the SCG region, where MDA8 O₃ was overestimated

(Figure 32 and 54). In the YRD region, there were large positive NO₂ biases with EDV2, 1 EDV3, and KOV5 (ranging from 6.4 to 22.7 ppb). Liu et al. (2021) reported that YRD 2 is in a VOC-limited regime when using EDV2 The findings indicated that a reduction 3 in NOx emissions led to an increase in O₃ concentrations, while a reduction in VOC 4 emissions resulted in lower O₃ concentrations. The lower bias of reason why O₃ is well 5 simulated in YRD, even though NO2 is highly overestimated in this region, can be 6 7 attributed to the combined influence of higher anthropogenic NOx emissions and VOC originated from both anthropogenic and biogenic sources (Figure S75). In contrast, 8 9 KOV5 underestimated NO2 in the NCP region, while EDV2 and EDV3 did not, which implies that KOV5 overestimates O₃ in NCP due to the overpredicted NOx titration 10 effect on O₃. All emissions showed significant discrepancies compared to NO₂ 11 12 observations in the SEC area, with a low correlation coefficient (0.19 to 0.26). EDV3 13 showed the lowest smallest bias of -1.9 ppb (-0.8 ppb) compared to EDV2 and KOV5 in South Korea (SMA) showing most reasonable NOx emissions in this region. The 14 15 daily averaged NO₂ exhibited spatial distributions similar to MDA8 O₃ and CO (Figure 65). The slopes of regression between the three model simulations and observations 16 were 1.31, 1.03, and 0.8 for EDV2, EDV3, and KOV5, respectively, in China. The 17 correlation coefficients between the simulated NO₂ utilizing EDV2, EDV3, and KOV5 18 and surface data were around 0.6 in China. EDV2, EDV3, and KOV5 demonstrated 19 20 good correlations with observations in South Korea (R = 0.69-0.74). Correlation coefficient (R) was the highest with KOV5 in South Korea (R=0.74). 21 22 Likewise, the diurnal patterns of Ox (= $NO_2 + O_3$) are well simulated with all 23 emission inventories (Supporting Information, Figure S8). The spatial distribution and 24 diurnal patterns of Ox are similar to O₃ except YRD (Supporting Information, Figure

1 S9). In YRD, the overestimations of Ox with all emission inventories reveals that same

2 <u>issue of NO₂ overestimations in Figure 5. Even though O₃ is well simulated in this</u>

3 region, the negative impact of NOx titration to O₃ formation is compensated with the

overestimated anthropogenic and biogenic VOC emissions as mentioned above.-

5 The simulated CO was averaged at each site and compared with observations

6 during the KORUS-AQ campaign period (Figure 76). The three model results showed

7 similar spatial distributions to observations, indicating higher CO concentrations in the

8 NCP, YRD, and PRD regions than their surrounding areas. However, all simulations

failed to reproduce the abundance of CO, indicating large negative biases throughout

10 the country. The bias was larger in South Korea than in China.

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3.3. Evaluations with the airborne and special surface chemical observations during KORUS-AQ

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3.3.1. The aircraft observations

Figure 27 shows the flight paths flown by the DC-8 during the KORUS-AQ campaign 16 17 period. In Table 43, we compare the model results for O₃, NO₂, CO, HCHO, TOL, XYL, 18 ETE (Ethene or OL2), and ISO with the corresponding observed values for all flight 19 tracks under 2 km height in South Korea (Table 3). On average, the model 20 underestimated O₃ by 15-18 ppb, with EDV3 exhibiting the lowest O₃ bias (-15.1 ppb) compared to EDV2 and KOV5 (-16.8 and -17.5 ppb, respectively). All emissions 21 showed positive biases for NO₂ (0.64 to 1.72 ppb), ETE (0.08 to 0.14 ppb), and ISO 22 23 (0.1 to 0.11 ppb). However, the model significantly underestimated CO, HCHO, TOL, and XYL for all three emissions. Given the large spatial variability of air pollutants in 24 South Korea, we also sampled aircraft data from six regions (see Figure 27) and 25

- 1 compared the three model results with the aircraft observations under 2 km height 2 (Figure 8).
- 3 The flight tracks that surveyed large power plants and factories in the Chungnam region on a daily basis are shown in Figure S106 in the Supporting Information. The 4 largest negative model O₃ bias was observed over the Chungnam region, with a 5 6 difference of 38-41 ppb. Emission estimation uncertainties can be significant over this region, where there are large point sources such as coal-burning power plants and 7 petrochemical industries. The model NO₂ agreed with the aircraft observations in SMA, 8 9 but it tends to overestimate the measurements in the other areas. There were substantial model overestimations of NO2 with EDV3 over the Chungnam and Busan areas, while 10 KOV5 showed the most reasonable model NO2 simulations. The model CO near the 11 surface was underestimated in the entire domain, resulting in high negative model CO 12 biases relative to the aircraft observations across the six regions (Figure 8). We 13 14 additionally conducted sensitivity test to investigate the contribution of CO to O₃ concentrations in SMA (Supporting information, Figure S11). Doubling CO emissions 15 in China did notn't significantly change O₃ concentrations at all levels under 2 km. Only 16 17 1.4 ppb of O₃ concentrations are changed on average during all flight observations-. AdditionallyIn addition, Wwe also evaluaidated the model HCHO, which can be 18 19 formed by oxidation of other VOCs but also directly emitted by anthropogenic sources, 20 to investigate potential issueuncertainties inof anthropogenic VOC emissions. *tThe*
- model HCHO was underestimated by all emission inventories for all subregions, with
- 22 negative biases being evident in the SMA, Yellow Sea, and Chungnam regions.
- Other model VOC species, such as TOL, XYL, ETE, and ISO, were also analyzed.
- 24 These VOC species are classified by their chemical structures and reactivities in the

RACM (Stockwell et al., 1997) (Table S42). For example, TOL includes toluene and 1 relatively less reactive aromatics, while XYL includes xylene and more reactive 2 aromatics. The WAS data from DC-8 were lumped into RACM (Supporting 3 Information Table S86, Lu et al., 2013) and were compared with aircraft observations. 4 5 When the model TOL or XYL was compared with the observed toluene and xylene, the model using KOV5 reasonably reproduced the observed concentrations (light gray bars 6 7 in Figure 8). However, the model TOL (even using KOV5) underestimated the observed lumped TOL for most of the regions except for Busan (bars including the dark gray part 8 9 in Figure 8). The model using KOV5 reasonably reproduced the observed xylene or XYL, except for the Chungnam and Busan regions. The observed ethene (or ETE) 10 concentrations were low (< 0.5 ppb), except for the Chungnam region, where the 11 12 average of measurements was 2.1 ppb. The model ethene concentration was higher than 13 the observations for the SMA, Kyungbuk, and Busan regions, while it had a large negative bias $(-1.6 \sim -1.3 \text{ ppb})$ for the Chungnam region. Regarding isoprene (ISO), one 14 15 of the most important biogenic VOCs, the model values were larger than the observations by a factor of 2. 16

In summary, underestimated CO and aromatic VOCs are the main features, along with underestimated ozone and HCHO. The largest discrepancies occur over the Chungnam region, where large point sources are located on the west coast of South Korea. The detailed statistics over the SMA and Chungnam area can be obtained from the Supporting Information (Table S97-S108).

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Figure 9 displays the vertical distributions of observed and simulated O₃ and related species over SMA. The shapes of the simulated profile were in agreement with the observations. Particularly, the model accurately reproduced the observed NO₂ profiles

though the surface NO₂ is underestimated by -4.2 to -0.8 ppb in SMA (Table 2 and Figure 9b). The underestimation of simulated surface NO2 is explained by the overestimation of molybdenum converter method; surface concentrations of NO₂ from molybdenum converter is larger than photolytic converter by 13.6% on average and 64% at 4 pm (Figure 10). Although diurnal pattern of surface NO₂ at 12-20 LT is explained by the overestimation of molybdenum converter, there are still some other possible reasons; 1) the emission factor used in this study is fromwas developed for the Los Angeles Basin, which may need to be adjusted for that might be not adequate to SMA, 2) the uncertainty of HOx and ROx radicals from other sources can affect the NO₂ concentrations.

However, the simulated O₃ and HCHO had negative biases of 16.4 ppb and 0.73 ppb, respectively, persisting from the surface to 2 km. Additionally, the simulated CO underestimated the observations by 40% throughout the vertical layer. While the model TOL and XYL, utilizing KOV5, agreed well with the observations below 1 kmat surface level and had the lowest bias of -0.88 and -0.12 ppb under 2 km, the results using EDV2 and EDV3 substantially underestimated the observations throughout the layer (Supporting information, Table S9). On the other hand, the model simulated ETE and ISO overestimated the observations below 1 km over SMA.

3.3.2. The ground-based observations

During the KORUS-AQ campaign, Pandora and surface measurements were co-located at the Olympic Park. Figure 10 compares the observed diurnal cycle of Pandora vertical columns and surface concentrations of NO₂ and HCHO with the model simulations. The photolytic converter was used to measure surface NO₂ to minimize positive bias

- 1 from the molybdenum converter. All emissions reasonably simulated the diurnal
- 2 patterns of vertical column and surface NO₂ and HCHO concentrations.
- The surface NO₂ peak appeared at 07 LT in the model and 08 LT in the observations,
- 4 associated with the increase of traffic and the under-developed convective boundary
- 5 layer. On the other hand, the Pandora NO₂ column amount increased from 06 LT to 12
- 6 LT and stayed at that value throughout the afternoon, indicating the increase of NO_x
- 7 emissions from morning to afternoon. The model-simulated NO₂ columns agreed with
- 8 those from Pandora in terms of absolute values and diurnal variations. The opposite
- 9 patterns between surface and column NO₂ were attributed to the change of boundary
- 10 layer height; NO₂ is concentrated near the surface layer as mixed layer is shallow in the
- morning and vertically well mixed during the daytime resulting in low surface NO₂
- 12 <u>concentrations also shown in (Crawford et al., (2020). On the other hand, vertical</u>
- 13 <u>column NO₂ concentrations show large values in the afternoon due to the consistent</u>
- 14 emission of NOx. The simulated and observed HCHO show similar diurnal variations,
 - but a

- All three emissions inventories underestimated both column and surface HCHO
- values by up to -8.5×10^{15} molecules·cm⁻² (-46%) at 7 LT and -0.9 ppbv (-26%) at the
- surface on average. The underestimations of the model HCHO relative to the Pandora
- and surface observations are similar to findings from comparisons of the model results
- 20 with the aircraft data (Figure 9). Therefore, the model VOC performance needs to be
- 21 investigated at the Olympic Park.
- The diurnal variations of the model O₃, CO, TOL, and XYL were evaluated against
- 23 the surface observations at the Olympic Park acquired during the KORUS-AQ
- campaign (Figure 11). The diurnal pattern and hourly averaged mixing ratio of O₃ were

- 1 well simulated with the three emission inventories with slight model negative biases.
- 2 The observed CO was 2.7 times higher than the model on average. Considering the
- diurnal profile of observed TOL and XYL, KOV5 reduced the model exhibited smaller
- 4 negative biases fromthan EDV2 and EDV3, but it still showed negative biases. The
- 5 model TOL and XYL showed peak concentrations at 08 LT, but the observation had a
- 6 maximum value at 06 LT. The model biases of XYL (-3.7 to -0.6 ppb, -89 to -20%)
- 7 were much larger than those in TOL at the surface. Our study demonstrates that the
- 8 improvement of VOC emission/chemistry representations in the model is necessary for
- 9 better simulations of air quality over SMA and South Korea.

- 3.4. The model performances over South Korea for the Local and Transport Cases
- 12 Previous studies have used meteorological conditions to classify synoptic patterns that
- affect air pollutant concentrations (Park et al. 2021; Peterson et al. 2019). In contrast,
- we defined the Transport and Local cases by comparing model results that used the
- 15 EDV3 base emission and the EDV3 zero-out-Chinese emission (see Figure 12). The
- 16 Local case comprises May 4, May 20, June 2, and June 3 (Supporting Information,
- 17 Figure S127), while the Transport case includes May 25, May 26, and May 31
- 18 (Supporting Information, Figure S<u>13</u>8). The Local (Transport) case in this study
- 19 generally aligns with the Stagnant and Blocking (Transport) cases in Peterson et al.
- 20 (2019); Stagnant and Blocking is the period that large anticyclone is located over South
- 21 Korea, and Transport case is the period that South Korea is largely affected by long-
- 22 range transport of air pollutants by westerly wind. The Local case has a Chinese
- 23 contribution to O₃ of under 11%, whereas the Transport case has a Chinese contribution
- 24 to O₃ of over 46%. EDV3 performed better in simulating O₃ for the Transport case

- 1 compared to EDV2 and KOV5, with a bias of only 2.7 ppb in comparison with the DC-
- 8 airborne observations. In contrast, for the Local case, all emissions had a negative
- bias ranging from 15.5-18.2 ppb., which implies that the insufficient local emissions
- 4 of O₃ precursors in the emission inventories are much important that the Chinese
- 5 <u>emissions.</u> See the Table S<u>11 and S12</u>9 in Supporting Information to obtain detailed
- 6 information of model performances against DC-8 measurements for different cases.
- 7 Surface concentrations of O₃ at Olympic Park also exhibited enhanced contributions
- 8 from Chinese anthropogenic emissions for Transport case (Figure S149). This section
- 9 focuses on the model simulations using EDV3 and its modified versions, EDV3_Ch2.
- 10 EDV3 Ko2 –and EDV3 ChKo2 (doubling Chinese and South Korean CO and VOC
- 11 emissions).
- Figure 13 illustrates the biases in the model O₃, CO, and HCHO using EDV3 and
- its variants relative to DC-8 observations over SMA. The plot highlights differences in
- biases for the Local and Transport cases. The model O₃ biases were negative, and the
- absolute values of biases were larger in the Local case than in the Transport case (-20%)
- versus -6%). The model CO biases were also negative, and not only the absolute values
- of biases but also relative differences were larger in the Transport case than in the Local
- 18 case despite the higher CO concentrations in Transport case, which implies that the
- 19 <u>background CO emissions are much biased in East Asia (Table S11)</u>. The model HCHO
- 20 biases were negative and similar for the two cases, except for a larger discrepancy
- between model and observation in the Local case than in the Transport case
- 22 representing deficiency of reactive VOC emissions from local sources in SMA.
- Doubling Chinese CO and VOC emissions (EDV3 Ch2) only slightly reduced
- biases in the Local case, whereas doubling South Korean CO and VOC emissions

(EDV Ko2) reduced biases more compared to the EDV3 Ch2 case. Doubling South 1 Korean CO and VOC emissions, as well as Chinese CO and VOC emissions (EDV3 2 3 ChKo2) led to the best results in O₃ and CO, were necessary to substantially reduce the model biases for the Local case. For the Transport case, doubling Chinese CO and VOC 4 5 emissions reduced biases to almost zero for CO and HCHO, but the model O₃ was much overestimated, with 14% positive biases (from an original bias of -6%). Doubling South 6 7 Korean COo and VOC emissions reduced the biases in O3 and CO a bit, but overestimated HCHO. The overestimation of O₃ in Transport case in the EDV3 Ch2 8 9 and EDV3 ChKo2 cases can be explained by not only excessive ISO but also overpredicted background O3 from doubled CO and VOC emissions in China 10 (Supporting information, Table S9-S13). Doubled CO and VOC emissions 11 overestimated O₃ concentrations over the Yellow Sea, which implies that the enhanced 12 background O₃ increase can enhanceincrease the O₃ level in SMA (Supporting 13 Information, Figure S15) (Kim et al., 2023). 14 15 Further increasing South Korean CO and VOC emissions in addition to the increase of Chinese CO and VOC emissions led to overestimations of O₃ (20%) and HCHO 16 17 (33%). Even though negative bias of HCHO is compensated by doubling CO and VOC emissions in China, the additional increase of South Korea CO and VOC emissions 18 19 (EDV3 ChKo2) still simulates low concentrations of TOL and XYL in the model but high concentrations of ETE, which implies that overestimations of other reactive VOCs 20 21 in SMA. The VOC speciation map can be another uncertainty that could affect O₃ 22 simulations in East Asia. These sensitivity tests modifying EDV3 indicate that increases 23 in CO and VOC emissions over South Korea improve the model O₃, CO, and VOC 24 simulations. However, increasing Chinese VOC (and CO) emissions may overestimate 1 the model O₃ for the studied period.

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3 4. Discussion: strategy for accurate surface O₃ simulations over urban and regional areas in China and South Korea 4 5 Due to unprecedentedly rich observational data sets acquired during KORUS-AQ, we investigated the status of O₃ simulations and outlined directions for their improvements 6 7 in SMA and South Korea. In this section, strategies for the enhanced accuracy of surface O₃ simulations over urban and regional areas in China and South Korea are discussed. 8 9 The discussion is based on the model simulations incorporating various emission 10 scenarios derived from EDV3. In Figures 14 and 15, diverse emission cases are labeled 11 from C1 to C7. Specifically C1, C2, and C3 correspond to EDV3 ChKo2, EDV3 Ch2, and EDV3 Ko2, respectively. Meanwhile, C4, C5, C6, and C7 represent scenarios 12 13 involving a 50% reduction in Chinese NO_x emissions, a 50% reduction in Chinese VOC emissions, a simultaneous 50% reduction in both Chinese NO_x and VOC emissions, 14 and a 75% reduction of Chinese NO_x emission, respectively, as discussed in Kim et al. 15 (2023). Examining various options involving the increase and decrease of NO_x and 16 17 VOC emissions from C1 to C7 sheds light on the direction for improving O₃ simulations. 18 Figure 14 illustrates the model O₃ and NO₂ biases (%) in each region for all cases based on EDV3 (Supporting Information, Table S14-S15 for detailed). EDV3 19 20 demonstrated good performance in simulating O₃ and NO₂ for the NCP, KOR, NEC, and NOC region. The most substantial model O₃ biases were observed in SCG and SEC, 21 22 with minimal model NO₂ biases. Conversely, the largest model NO₂ biases were found 23 in YRD and PRD, accompanied by modest model O₃ biases. Improvements are needed 24 for model O₃ in SCG, SEC, YRD, and PRD with reasonable NO₂ simulations. For SCG

1	and SEC, the C5 case (50% VOC emission reduction only) exhibited the lowest O ₃
2	biases. Doubled Chinese VOC emission case (C1 and C2) in SCG and SEC resulted in
3	increased O ₃ biases to ~100%, compared to 66% in the EDV3 case. In this study, the
4	anthropogenic VOC emissions were reduced. Further reductions of biogenic VOC
5	emissions as well as anthropogenic emissions need to be explored in the future. For
6	SCG and SEC, a reduction in NO _x emissions also led to a slight decrease in O ₃ biases.
7	FNR values for the two regions are about 1.3, which turned out to be still VOC-limited
8	or in a transitional state. For the YRD and PRD regions, first, NO _x emissions need to
9	be reduced to improve NO ₂ biases in the model. The case C6 (50% reductions in both
10	NOx and VOC emissions) yielded the most favourable O3 and NO2 simulations. Solely
11	reducing NO _x emissions (as in case C4) increase O ₃ biases by 25-36% relative to EDV3.
12	The FNR values for YRD and PRD are 0.32 and 0.52, respectively, placing them in the
13	VOC-limited regime (FNR < 1). In general, an increase in Chinese VOC emissions (as
14	observed in cases C1 and C2) resulted in elevated surface ozone levels for all regions,
15	including KOR. For NCP, KOR, NEC, and NOC where the model O ₃ and NO ₂ agree
16	with the observations, reducing VOC proves to be an effective strategy for decreasing
17	surface O ₃ .
18	In Figure 15, the model O ₃ and NO ₂ biases (%) in the 12 mega cities in China and
19	South Korea are illustrated for all cases. Refer to Supporting Information Table S16 and
20	S17 for specific values. EDV3 showed effective performance in simulating O ₃ and NO ₂
21	for cities such as Beijing, Tianjin, Hangzhou, SMA, and Xian. The most substantial
22	model O ₃ biases were observed in Chengdu and Chongqing, with minimal model NO ₂
23	biases. In contrast, the notable model NO ₂ biases were identified in Shanghai, Nanjing,
24	Guangzhou, Shenzhen, and Wuhan, accompanied by modest model O ₃ biases. For

1 Chengdu and Chongqing, situated roughly in SCG, the C5 case (50% VOC emission

2 reduction only) results in the lowest O₃ biases with decent NO₂ simulations. For

3 Shanghai, Nanjing, Guangzhou, Shenzhen, and Wuhan, case C6 (50% reductions in

4 <u>both NOx and VOC emissions) produced the most favorable O₃ and NO₂ simulations.</u>

Simply reducing NO_x emissions (as in case C4) increase O₃ biases in these cities.

Overall, the increase in Chinese VOC emissions (cases C1 and C2) resulted in elevated

7 surface ozone levels for all cities, including SMA with an increase in biases, biases,

except for Shanghai. Reduction of only VOC emissions (C5) led to the lowest surface

9 O₃ levels for all cities.

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54. Summary and conclusions

We conducted sensitivity tests using WRF-Chem with three different bottom-up emission inventories (EDGAR-HTAP v2, v3, and KORUS v5) to investigate the impacts of different emissions on the simulation of O3 and precursors in East Asia. This study is the first to use EDGAR-HTAP v3 with WRF-Chem v4.4 and extends the validation domain to the whole of China during the KORUS-AQ campaign period. We extensively evaluidated these emission inventories using both ground and aircraft observations in East Asia.

The three emission inventories accurately reproduced the diurnal profiles and absolute values of surface O₃ for most subregions in China, except for the SCG and SEC areas. However, discrepancies were observed in the model performance for the MDA8 O₃ concentrations, with poor correlations observed over regions with high

- 1 HCHO concentrations (south of 30°N) and relatively low ratios of FNR (north of 30°N).
- 2 The emission inventories reasonably reproduced the spatial distribution of daily surface
- 3 NO₂ concentrations. However, we found that CO was considerably underestimated by
- 4 the emission inventories over both China and South Korea.
- 5 We evaluated the model simulations against vertical profile measurements of O₃,
- 6 NO₂, CO, HCHO, TOL, XYL, ETE, and ISO from the DC-8 aircraft, as well as surface
- 7 observations over South Korea. The simulated vertical shapes of O₃, NO₂, CO, HCHO,
- 8 TOL, XYL, ETE, and ISO agreed well with the DC-8 measurements in the SMA,
- 9 although negative biases were observed for O₃, CO, TOL, XYL, and HCHO, with the
- 10 largest discrepancy between the model results and observations in the Chungnam area.
- 11 When we compared the simulations with the surface in-situ measurements and
- 12 PANDORA observations at the Olympic Park in Seoul, the model accurately
- 13 reproduced the diurnal patterns of surface and vertical columns of NO₂ and HCHO.
- However, we found that the model underestimated TOL and XYL. This underestimation
- of TOL and XYL is one of the reasons why the model underestimates O₃ concentrations,
- as VOCs contribute to NO to NO2 conversions resulting in O3 production via
- 17 photochemistry.
- We also classified the flight tracks into two categories: Local and Transport cases.
- 19 We found that the negative bias of O₃ was much larger under the Local case than the
- 20 Transport case. When the increment of CO and VOC emissions in South Korea is taken
- 21 into account, the biases of O₃ are significantly reduced, indicating the need for local
- 22 emission adjustments to decrease O₃ bias in South Korea.
- To improve surface O₃ simulations over China and South Korea using EDV3,
- lowering VOC emissions are advantageous for SCG and SEC including urban areas

- 1 like Chengdu and Chongqing. Meanwhile, for YRD and PRD regions, as well as cities
- 2 such as Shanghai, Nanjing, Guangzhou, Shenzhen, and Wuhan, both NOx and VOC
- 3 emissions should be reduced to enhance model performances. Increase in VOC
- 4 emissions adversely affected the model's accuracy in simulating O₃ in China, leading
- 5 to increased biases.
- 6 Our study revealed a consistent overestimation of isoprene over SMA. The
- 7 uncertainty of biogenic VOC emissions from MEGAN can affect the model O₃
- 8 performance. Therefore, to achieve more accurate simulations of O₃ in East Asia, it is
- 9 essential to explore precise representations of both anthropogenic and biogenic VOC
- 10 emissions.

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Code and data availability

- 13 WRF-Chem source codes are distributed by NCAR
- 14 (https://doi:10.5065/D6MK6B4K). WRF-Chem v4.4 is available in the GitHub (wrf-
- model, 2022). The exact version of WRF-Chem codes and configuration files are
- archived at https://doi.org/10.5281/zenodo.8260026 (Kim et al., 2023). National
- 17 Centers for Environmental Prediction (NCEP) FNL data can be accessed from Research
- Data Archive (RDA) (NCEP, 2019). The CAM-Chem data for boundary conditions is
- 19 also obtained from RDA (ACOM, 2019). The EDGAR-HTAP v2 data can be
- 20 downloaded in the website (https://edgar.jrc.ec.europa.eu/dataset htap v2). The
- 21 EDGAR-HTAP v3 is archived on Zenodo (Crippa, 2023). The KORUS-AQ data are
- 22 available in the website (https://www-air.larc.nasa.gov/cgi-bin/ArcView/korusaq)
- 23 (doi:10.5067/Suborbital/KORUSAQ/DATA01). The EDGAR-HTAP v2, v3, and

- 1 KORUS v5 data including emission processing programs are available at
- 2 https://doi.org/10.5281/zenodo.8260026 (Kim et al., 2023).

4 Author contribution

- 5 KMK conducted simulations, analysis and wrote the paper. SWK designed this study,
- 6 secured funding, performed analysis and wrote the paper. SS supported model set-up
- 7 and contributed to refining the paper. DRB measured VOC data from DC-8. SC
- 8 acquired ground-based in-situ VOC data at Olympic Park. JHC performed analysis and
- 9 wrote the paper. LE and GF assisted in setting up the model emissions and discussed
- 10 about the model performance. AF measured HCHO data from DC-8. JRH measured
- Pandora data (NO₂, HCHO). JH retrieved PBL height and discussed about the results.
- 12 JJ acquired NO₂ data at Olympic Park with different methods. AJW acquired NO₂ and
- 13 O₃ data from DC-8. JHW and QZ provided emissions inventories and related
- information. All authors reviewed the manuscript.

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Competing interests

- 17 At least one of the (co-)authors is a member of the editorial board of Geoscientific
- 18 Model Development.

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1 **Table List** 2 3 Table 1. The model experiments with different emissions. 4 5 Table 2. Comparison of the ground-based hourly O₃, NO₂, and CO observations with the simulations utilizing EDGAR-HTAP v2 (EDV2) and v3 (EDV3) and KORUS v5 6 (KOV5) in each regional box (unit = ppb). N is the number of samples. R is correlation 7 coefficient. 8 9 Table 3. Comparison of total NOx, TOL, XYL, biogenic isoprene emissions, and 10 formaldehyde to NO₂ ratio (FNR) for different emission data sets in each regional box. 11 The MEGAN biogenic isoprene emissions are equally applied to all simulations using 12 different emission data. (unit = mol/s for emissions) 13 14 15 Table 43. Comparison of aircraft-based 1-minuite-interval O₃, NO₂, CO, HCHO, TOL, XYL, ETE, and ISO observations with EDV2, EDV3, and KOV5 for all flight cases 16 under 2 km height (unit = ppb). N is the number of samples. R is correlation coefficient. 17

Figure List

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3 Figure 1. The averaged spatial distribution map of the NO, CO, and TOL (toluene + less reactive aromatics) emissions from EDGAR-HTAP v2, v3, and KORUS v5 in May. The 4 5 boxes represent Northern China (NOC, 38-42°N/106-110°E), Sichuan-Chongqing-6 Guizhou (SCG, 27-33°N/103-109°E), Pearl River Delta (PRD, 21.5-24°N/112-115.5°E), 7 Southeastern China (SEC, 24-28°N/116-120°E), Yangtze River Delta (YRD, 30-33°N/119-122°E), South Korea (KOR, 34.5-38°N/126-130°E), North China Plain (NCP, 8 9 34-41°N/113-119°E), and Northeastern China (NEC, 43-47°N/124-130°E). NOC, NEC, 10 and SEC are denoted by blue boxes (non-urban). NCP, SCG, PRD, YRD, and KOR are denoted by red boxes (urban). 11

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Figure 72. The DC-8 flight paths during the KORUS-AQ campaign period (black) and 6 regional boxes (1: Seoul Metropolitan Area (SMA); 2: Yellow Sea; 3: Chungnam; 4: Kyungbuk; 5: Gwangju; 6: Busan) (red).

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17 Figure 23. Averaged O3 concentrations from ground-based observations and model simulations over the areas that distinguish urban (red box) and non-urban (green box) 18 region (central plot). Box-averaged diurnal cycle (solid lines) of O₃ and 1/4 of standard 19 20 deviations (filled area) from observations (black), EDV2 (greensky blue), EDV3 (blue), 21 and KOV5 (red) by local time are shown. The results are shown for Northern China 22 (NOC, 38-42°N/106-110°E), Sichuan-Chongqing-Guizhou (SCG, 27-33°N/103-109°E), Pearl River Delta (PRD, 21.5-24°N/112-115.5°E), Southeastern China (SEC, 24-23 28°N/116-120°E), Yangtze River Delta (YRD, 30-33°N/119-122°E), South Korea 24 25 (KOR, 34.5-38°N/126-130°E), North China Plain (NCP, 34-41°N/113-119°E), and Northeastern China (NEC, 43-47°N/124-130°E). 26

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Figure 43. Comparison of (a) the campaign averaged ground-based maximum daily average of 8-hour O₃ (MDA8 O₃) (unit: ppb) observations and WRF-Chem simulations with (d) EDGAR-HTAP v2 (EDV2), (e) v3 (EDV3), (f) KORUS v5 (KOV5) and (g, h, i) the differences between the observations and model results. The sub-regions are presented with red (urban) and green (non-urban) boxes. The scatter plots comparing averaged observations and the three-emission-based WRF-Chem simulations (greensky blue; EDV2, blue; EDV3, red; KOV5) are shown in (b) and (c) for Eastern China and South Korea, respectively. (a, d-e) Color-filled circles in (a), (d), (e), and (f) represent the averaged MDA8 O₃ for the whole campaign period (1st May to 10th June).

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Figure 45. The same as Figure 23 except NO₂.

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Figure \frac{56}{10}. The same as Figure \frac{34}{10} except daily NO<sub>2</sub> (unit: ppb).
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      Figure 67. The same as Figure 34 except daily CO (unit: ppm).
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      Figure 7. The DC-8 flight paths during the KORUS-AQ campaign period (black) and 6
      regional boxes (1: Seoul Metropolitan Area (SMA); 2: Yellow Sea; 3: Chungnam; 4:
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      Kyungbuk; 5: Gwangju; 6: Busan) (red).
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      Figure 8. The mean (bars) and 1/4 of standard deviations (whiskers) of (a) O<sub>3</sub>, (b) NO<sub>2</sub>,
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      (c) CO, (d) HCHO, (e) TOL, (f) XYL, (g) ethene (ETE), and (h) isoprene (ISO) (unit =
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      ppb) from DC-8 (dark grey), EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) for
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      each box are shown, respectively. TOL and XYL are calculated based on Table S68
      (Supporting Information). The contribution of toluene to TOL and m/p-Xylene + o-
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      Xylene to XYL is represented with light grey bars (e, f). The sampling numbers are
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      represented with magenta color above the plots.
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      ETE, and (h) ISO from DC-8 (black), EDV2 (greensky blue), EDV3 (blue), and KOV5
      (red) in SMA under 2 km height above ground level. The 1/2 of standard deviations are
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      represented with black whiskers in each 200m layer. The sample number is presented
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      with magenta color on the right side of the plots.
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      Figure 10. The diurnal cycles of vertical columns and surface concentrations of (a) NO<sub>2</sub>
      and (b) HCHO from Pandora spectrometer (column), and ground-based instruments
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      (TEI 42i NOx analyzer and Aerodyne QCL) at the Olympic Park site (37.5232°N,
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      127.126°E). EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) are compared with
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      observations. The WRF-Chem vertical column concentrations are produced by
      summing all vertical layers.
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      Figure 11. Diurnal cycles of surface (a) O<sub>3</sub>, (b) CO, (c) TOL, and (d) XYL at the
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      Olympic Park site. EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) are compared
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      with the observations. 1/4 of standard deviations are represented with grey shades. The
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      average period is from the 11th May to the 10th June.
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Figure 12. Averaged O₃ (bars) and 1/4 of standard deviations (whiskers) (unit: ppbv)

for the 20 DC8 flights (under 2 km height). The observations (grey) are compared with

the model results utilizing EDV2 (greensky blue), EDV3 (blue), and KOV5 (red).

- 1 White hatch-filled bars over blue bars are the contribution of Chinese emissions to O₃
- 2 concentrations obtained from the default and sensitivity model runs with/without
- 3 Chinese anthropogenic emissions. The Local (5/4,20 and 6/2,3) and Transport
- 4 (5/25,26,31) cases are shaded with light blue and orange, respectively.

- 6 Figure 13. The biases in (a) the model O₃, (b) CO, and (c) HCHO concentrations (bars)
- 7 relative to the DC-8 observations under 2 km height over SMA (dark gray: EDV3, red:
- 8 EDV3 Ch2, blue: EDV3 ChKo2): (left panel) Local and (right panel) Transport case.
- 9 Fractional differences (%) are shown in the white boxes.

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- 11 Figure 14. Comparison of relative biases ((Model-Observation)/Observation, unit=%)
- of daily O₃ and NO₂ at surface observation sites during the KORUS-AQ campaign
- period from sensitivity simulation (C1-7) with EDV3 in each region (NCP, SCG, YRD,
- 14 PRD, KOR, NEC, NOC, and SEC). C1; EDGAR-HTAP v3 with double CO and VOC
- emission in China and South Korea, C2; EDGAR-HTAP v3 with double CO and VOC
- 16 emission in China, C3; EDGAR-HTAP v3 with double CO and VOC emission in South
- 17 Korea, C4; EDGAR-HTAP v3 with 50% NOx reduction in China, C5; EDGAR-HTAP
- 10 Roled, C+, LDOAR-IITAI V5 WILL 50/0 TOX ICCCCION III CHINA, C5, LDOAR-IITAI
- 18 <u>v3 with 50% VOC reduction in China, C6; EDGAR-HTAP v3 with 50% NOx and VOC</u>
- reduction in China, C7; EDGAR-HTAP v3 with 75% NOx reduction in China.

- 21 Figure 15. Same as Figure 14 except that the region is changed to cities; Beijing (39.4-
- 22 41.1N, 115.4-117.5E), Tianjin (38.55-40.25N, 116.7-118.1E), Chengdu (30.05-31.5N,
- 23 <u>103-105E</u>), Chongqing (28.15-32.25N, 105.3-110.2E), Shanghai (30.7-31.5N, 120.85-
- 24 122E), Hangzhou (29.2-30.6N, 118.3-120.9E), Nanjing (31.2-32.65N, 118.35-119.25E),
- 25 Guangzhou (22.55-24N, 112.9-114.05E), Shenzhen (22.4-22.9N, 113.7-114.65E),
- 26 SMA (37.2-37.8N, 126.5-127.3E), Wuhan (29.95-31.4N, 113.65-115.1E), and Xian
- 27 (33.65-34.75N, 107.65-109.9E).-

Table 1. The model experiments with different emissions

Experiments	Emissions
EDV2	EDGAR HTAP v2
EDV3	EDGAR HTAP v3
KOV5	KORUS v5
EDV3_Ch2	EDGAR HTAP v3 with double CO, VOC emission in China
EDV3_ChKo2	EDGAR HTAP v3 with double CO, VOC emission in China & South Korea

Table 1. The model experiments with different emissions.

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Esx periments	<u>Emissions</u>
EDV2	EDGAR-HTAP v2
EDV3	EDGAR-HTAP v3
KOV5	KORUS v5
EDV3_Ch2	EDGAR-HTAP v3 with double CO, VOC emission in China
EDV3_Ko2	EDGAR-HTAP v3 with double CO, VOC emission in South Korea
EDV3_ChKo2	EDGAR-HTAP v3 with double CO, VOC emission in China & South Korea

Table 2. Comparison of the ground-based hourly O₃, NO₂, and CO observations with the simulations utilizing EDGAR-HTAP v2 (EDV2) and v3 (EDV3) and KORUS v5 (KOV5) in each regional box (unit = ppb). N is the number of samples. R is correlation coefficient.

	Region		1) NCP	1),a) SCG	1) YRD	1) PRD	1),b) KOR (SMA)	^{2),c)} NEC	2),d) NOC	^{2),e)} SEC
	N		190	104	93	68	358 (125)	45	28	43
O ₃	OBS	Mean	44.5	34.6	38.2	27.9	41.5 (36.6)	40.9	44.3	26.1
		Mean	32.2	53.5	21.6	27.6	40.5 (31.1)	28.6	39.4	40.8
	EDV2	Bias	-12.3	18.9	-16.6	-0.3	-1.0 (-5.5)	-12.3	-4.9	14.7
		R	0.65	0.53	0.62	0.61	0.59 (0.60)	0.48	0.63	0.52
		Mean	43.4	57.5	35.7	34.7	41.0 (32.6)	35.2	43.7	45.5
	EDV3	Bias	-1.1	23.0	-2.5	6.8	-0.5 (-4.0)	-5.7	-0.6	19.4
		R	0.68	0.55	0.66	0.65	0.56 (0.57)	0.63	0.67	0.55
		Mean	49.0	55.3	41.1	35.7	42.2 (33.1)	37.1	43.8	42.4
	KOV5	Bias	4.5	20.7	2.8	7.8	0.7 (-3.5)	-3.8	-0.5	16.3
		R	0.71	0.53	0.65	0.70	0.62 (0.64)	0.62	0.67	0.54
NO ₂	OBS	Mean	17.5	13.8	17.1	12.9	23.2 (32.5)	13.5	11.9	9.6
		Mean	25.8	12.7	39.8	22.0	18.8 (29.6)	13.7	12.9	11.0
	EDV2	Bias	8.3	-1.0	22.7	9.1	-4.3 (-3.0)	0.2	1.0	1.5
		R	0.45	0.37	0.38	0.54	0.51 (0.34)	0.49	0.47	0.19
		Mean	21.8	12.2	30.4	21.0	21.3 (31.8)	11.2	10.3	11.3
	EDV3	Bias	4.3	-1.6	13.3	8.1	-1.9 (-0.8)	-2.3	-1.6	1.7
		R	0.44	0.34	0.36	0.52	0.49 (0.31)	0.49	0.52	0.22
		Mean	13.9	7.5	23.5	13.3	17.7 (28.3)	7.0	7.7	7.7
	KOV5	Bias	-3.6	-6.3	6.4	0.3	-5.5 (-4.2)	-6.5	-4.2	-1.9
		R	0.44	0.37	0.41	0.52	0.51 (0.39)	0.49	0.51	0.26
	OBS	Mean	835	597	694	636	443 (493)	527	579	655
		Mean	373	389	455	282	175 (210)	206	162	258
	EDV2	Bias	-462	-208	-239	-354	-267 (-283)	-321	-417	-397
		R	0.24	0.20	0.42	0.30	0.31 (0.30)	0.21	0.09	0.18
СО		Mean	374	359	535	282	196 (208)	221	162	256
	EDV3	Bias	-461	-238	-159	-354	-247 (-285)	-306	-417	-398
		R	0.22	0.19	0.35	0.31	0.26 (0.33)	0.24	0.10	0.25
		Mean	355	358	475	305	190 (217)	231	176	266
	KOV5	Bias	-480	-239	-219	-331	-253 (-276)	-296	-404	-388
		R	0.27	0.21	0.48	0.29	0.35 (0.36)	0.15	0.10	0.13

^{5 1)} Urban area, 2) Non-urban area

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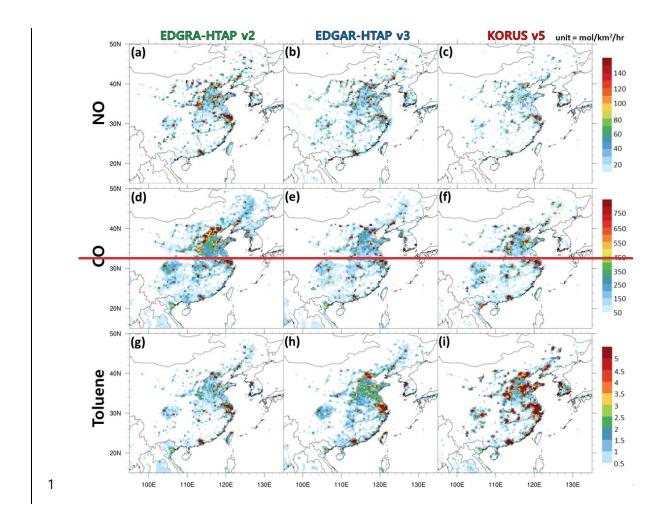
a) Sichuan-Chongqing-Guizhou, b) South Korea, c) Northeastern China, d) Northern China, e) Southeastern China

Table 3. Comparison of total NOx, TOL, XYL, biogenic isoprene emissions, and formaldehyde to NO₂ ratio (FNR) for different emission data sets in each regional box. The MEGAN biogenic isoprene emissions are equally applied to all simulations using different emission data. (unit = mol/s for emissions)

Type	emissions	<u>NCP</u>	<u>SCG</u>	YRD	PRD	KOR(SMA)	<u>NEC</u>	NOC	SEC
NOx emission	EDV2	<u>5967</u>	<u>1500</u>	<u>2366</u>	<u>1178</u>	990(196)	<u>987</u>	<u>688</u>	<u>590</u>
	EDV3	<u>5202</u>	<u>1654</u>	<u>1642</u>	<u>1091</u>	1191(214)	<u>876</u>	<u>597</u>	<u>662</u>
	KOV5	3237	902	1166	607	886(191)	<u>513</u>	<u>373</u>	410
	EDV2	<u>140</u>	<u>56</u>	<u>84</u>	<u>47</u>	<u>27(6)</u>	<u>26</u>	<u>8</u>	<u>20</u>
TOL emission	EDV3	<u>220</u>	<u>77</u>	<u>99</u>	<u>68</u>	<u>27(8)</u>	<u>40</u>	<u>9</u>	<u>36</u>
	KOV5	<u>403</u>	<u>106</u>	<u>234</u>	<u>155</u>	98(26)	<u>68</u>	<u>21</u>	<u>79</u>
	EDV2	<u>84</u>	<u>34</u>	<u>51</u>	<u>28</u>	<u>15(4)</u>	<u>15</u>	<u>4</u>	<u>12</u>
XYL emission	EDV3	<u>132</u>	<u>46</u>	<u>60</u>	<u>41</u>	<u>16(4)</u>	<u>24</u>	<u>6</u>	<u>22</u>
	KOV5	133	<u>35</u>	<u>79</u>	<u>52</u>	<u>41(9)</u>	<u>21</u>	<u>7</u>	<u>26</u>
	genic e emission	<u>132</u>	<u>364</u>	<u>43</u>	<u>127</u>	135(6)	<u>106</u>	<u>23</u>	<u>310</u>
ENID	EDV2	<u>0.252</u>	<u>1.3122</u>	0.19	0.52	0.532(0.197)	<u>0.6857</u>	0.76	<u>1.1805</u>
<u>FNR</u> (14- 16LT)	EDV3	<u>0.440</u>	<u>1.3023</u>	<u>0.323</u>	<u>0.524</u>	<u>0.432(0.186)</u>	<u>0.9380</u>	0.941	<u>1.3321</u>
10L1)	KOV5	<u>0.7268</u>	2.323	0.48	<u>10.0095</u>	0.71(0.2219)	1.4425	1.49	1.9173

Table 34. Comparison of aircraft-based 1-minuite-interval O₃, NO₂, CO, HCHO, TOL, XYL, ETE, and ISO observations with EDV2, EDV3, and KOV5 for all flight cases under 2 km height (unit = ppb). N is the number of samples. R is correlation coefficient.

Species	Type	N	Mean	Bias	σ	R
O ₃	OBS		84.4		19.9	
	EDV2	£101	67.5	-16.8	16.7	0.44
	EDV3	5191	69.3	-15.1	17.8	0.43
	KOV5		66.9	-17.5	15.8	0.50
	OBS		2.19		4.49	
NO	EDV2	5047	3.06	0.87	4.60	0.71
NO_2	EDV3	5047	3.91	1.72	5.34	0.67
	KOV5		2.83	0.64	4.73	0.73
	OBS		253		100	
CO	EDV2	5575	148	-105	48	0.60
CO	EDV3	5575	156	-97	47	0.59
	KOV5		146	-107	43	0.62
	OBS		2.37		1.64	
исио	EDV2	5265	1.75	-0.62	1.01	0.69
НСНО	EDV3	5365	1.78	-0.59	1.02	0.67
	KOV5		1.80	-0.57	1.10	0.71
	OBS	730	2.60		2.02	
TOL	EDV2		0.47	-2.13	0.38	0.39
TOL	EDV3	/30	0.55	-2.05	0.48	0.38
	KOV5		1.58	-1.01	1.30	0.37
	OBS		0.73		0.65	
VVI	EDV2	200	0.23	-0.50	0.23	0.30
XYL	EDV3	289	0.30	-0.43	0.31	0.30
	KOV5		0.49	-0.24	0.47	0.27
	OBS		0.42		1.59	
ETE	EDV2	2573	0.51	0.09	0.65	0.14
ETE	EDV3	23/3	0.56	0.14	0.76	0.15
	KOV5		0.51	0.08	0.58	0.20
ISO	OBS		0.08		0.09	
	EDV2	1204	0.18	0.10	0.21	0.41
	EDV3	1294	0.19	0.11	0.20	0.41
	KOV5		0.17	0.10	0.20	0.42



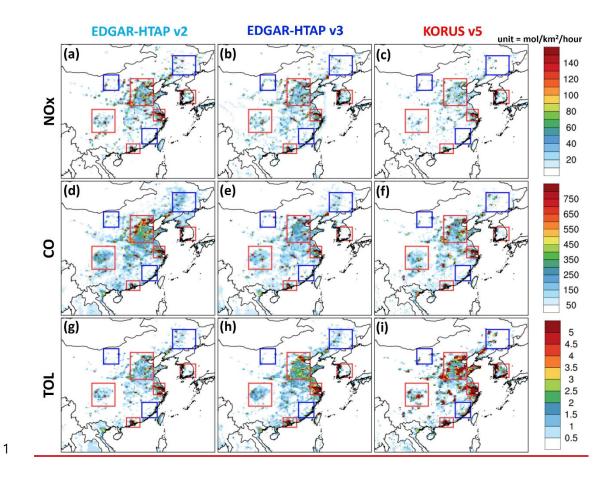
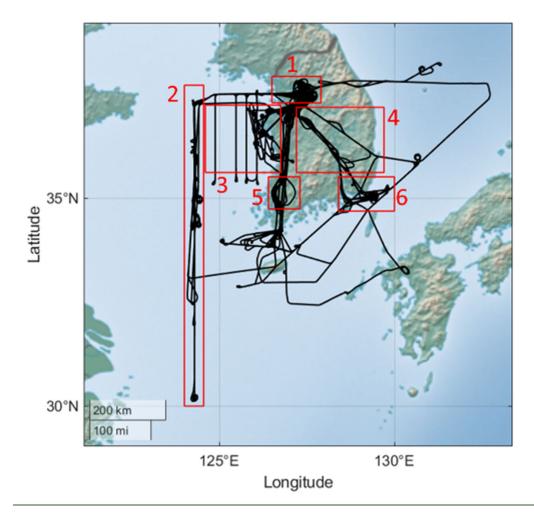


Figure 1. The averaged spatial distribution map of the NO, CO, and TOL (toluene + less reactive aromatics) emissions from EDGAR-HTAP v2, v3, and KORUS v5 in May. The boxes represent Northern China (NOC, 38-42°N/106-110°E), Sichuan-Chongqing-Guizhou (SCG, 27-33°N/103-109°E), Pearl River Delta (PRD, 21.5-24°N/112-115.5°E), Southeastern China (SEC, 24-28°N/116-120°E), Yangtze River Delta (YRD, 30-33°N/119-122°E), South Korea (KOR, 34.5-38°N/126-130°E), North China Plain (NCP, 34-41°N/113-119°E), and Northeastern China (NEC, 43-47°N/124-130°E). NOC, NEC, and SEC are denoted by blue boxes (non-urban). NCP, SCG, PRD, YRD, and KOR are denoted by red boxes (urban).



<u>Figure 72</u>. The DC-8 flight paths during the KORUS-AQ campaign period (black) and 6 regional boxes (1: Seoul Metropolitan Area (SMA); 2: Yellow Sea; 3: Chungnam; 4: Kyungbuk; 5: Gwangju; 6: Busan) (red).

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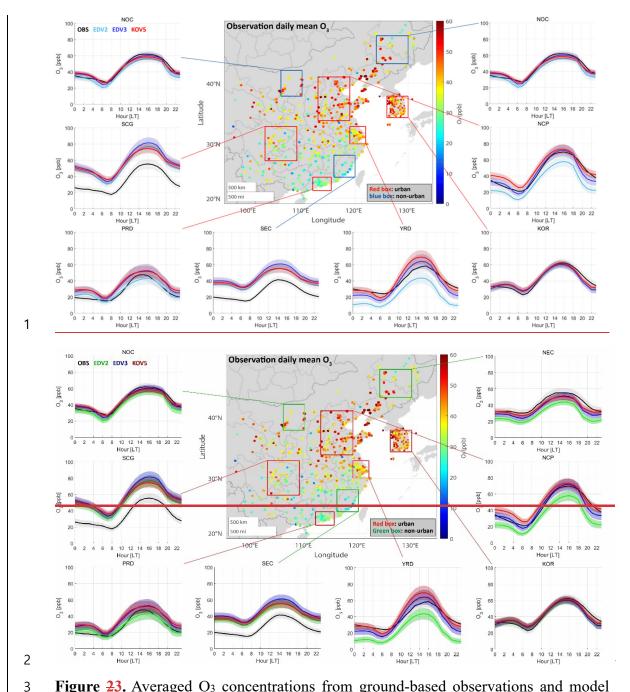


Figure 23. Averaged O₃ concentrations from ground-based observations and model simulations over the areas that distinguish urban (red box) and non-urban (green box) region (central plot). Box-averaged diurnal cycle (solid lines) of O₃ and 1/4 of standard deviations (filled area) from observations (black), EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) by local time are shown. The results are shown for Northern China (NOC, 38-42°N/106-110°E), Sichuan-Chongqing-Guizhou (SCG, 27-33°N/103-109°E), Pearl River Delta (PRD, 21.5-24°N/112-115.5°E), Southeastern China (SEC, 24-28°N/116-120°E), Yangtze River Delta (YRD, 30-33°N/119-122°E), South Korea (KOR, 34.5-38°N/126-130°E), North China Plain (NCP, 34-41°N/113-119°E), and Northeastern China (NEC, 43-47°N/124-130°E).

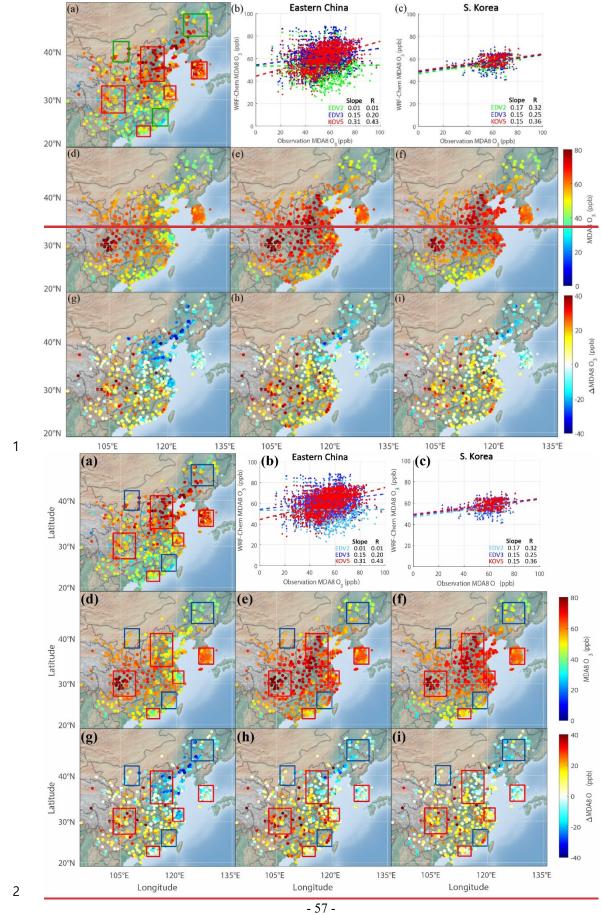


Figure 34. Comparison of (a) the campaign averaged ground-based maximum daily average of 8-hour O₃ (MDA8 O₃) (unit: ppb) observations and WRF-Chem simulations with (d) EDGAR-HTAP v2 (EDV2), (e) v3 (EDV3), (f) KORUS v5 (KOV5) and (g, h, i) the differences between the observations and model results. The sub-regions are presented with red (urban) and green (non-urban) boxes. The scatter plots comparing averaged observations and the three-emission-based WRF-Chem simulations (greensky blue; EDV2, blue; EDV3, red; KOV5) are shown in (b) and (c) for Eastern China and South Korea, respectively. (a, d-e) Color-filled circles in (a), (d), (e), and (f) represent the averaged MDA8 O₃ for the whole campaign period (1st May to 10th June).

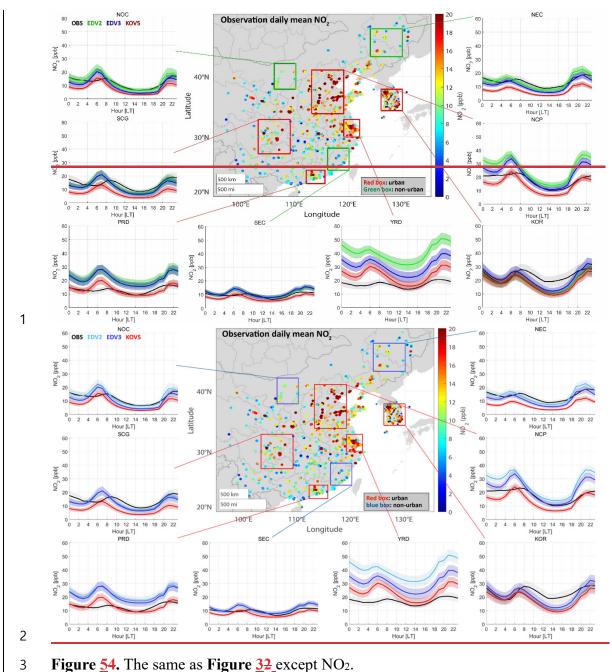
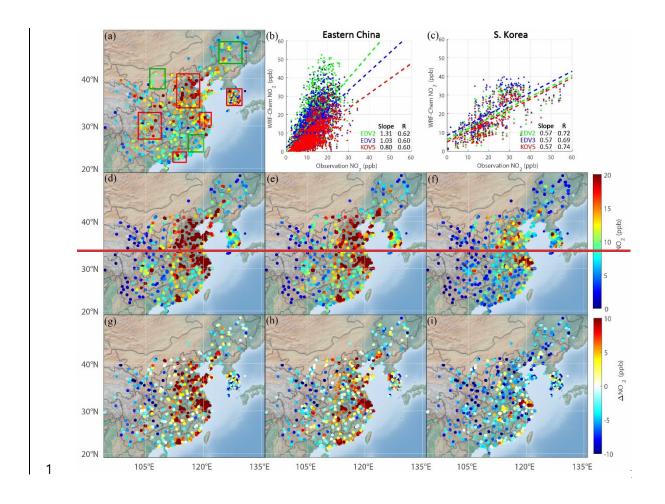


Figure 54. The same as Figure 32 except NO2.



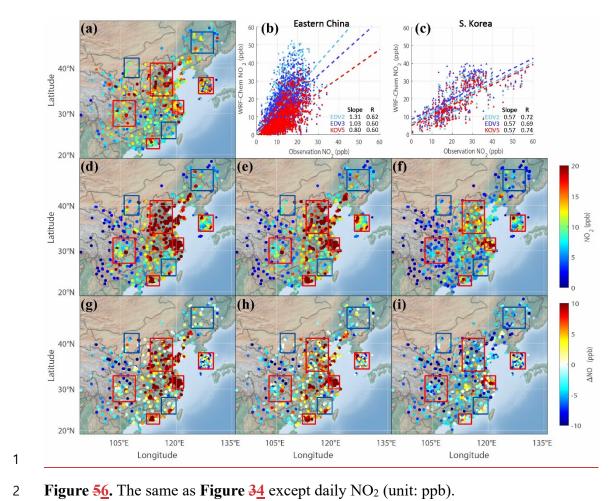
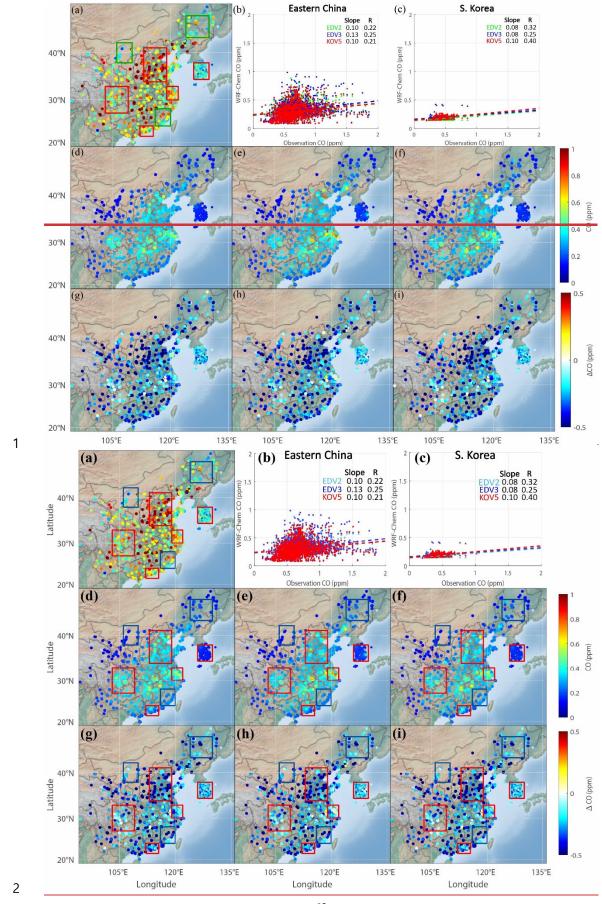


Figure 56. The same as Figure 34 except daily NO₂ (unit: ppb).



1 **Figure 67.** The same as **Figure 34** except daily CO (unit: ppm).

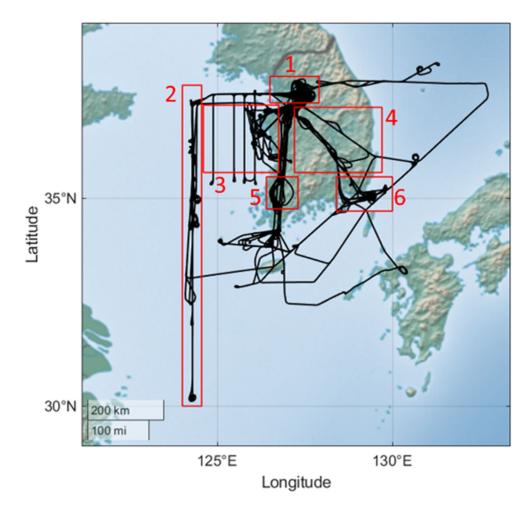
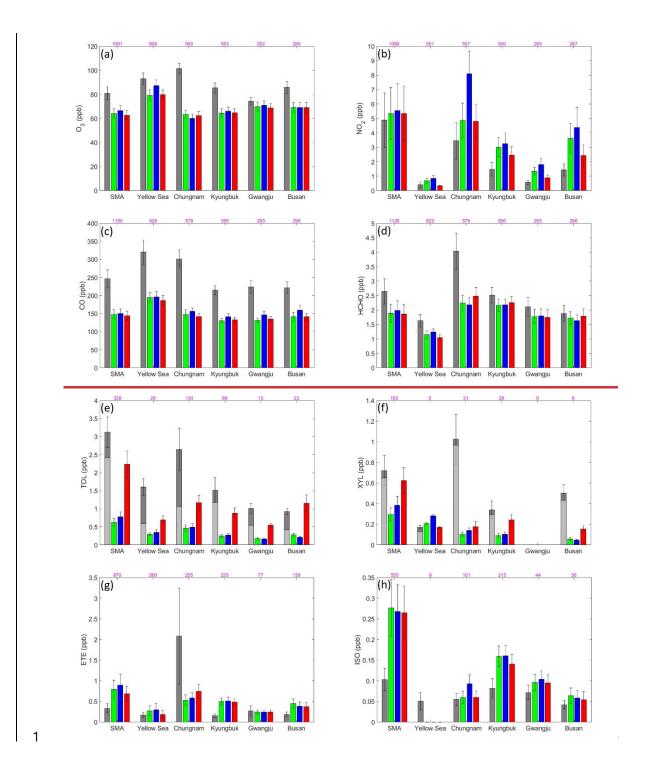


Figure 7. The DC-8 flight paths during the KORUS-AQ campaign period (black) and 6 regional boxes (1: Scoul Metropolitan Area (SMA); 2: Yellow Sea; 3: Chungnam; 4: Kyungbuk; 5: Gwangju; 6: Busan) (red).



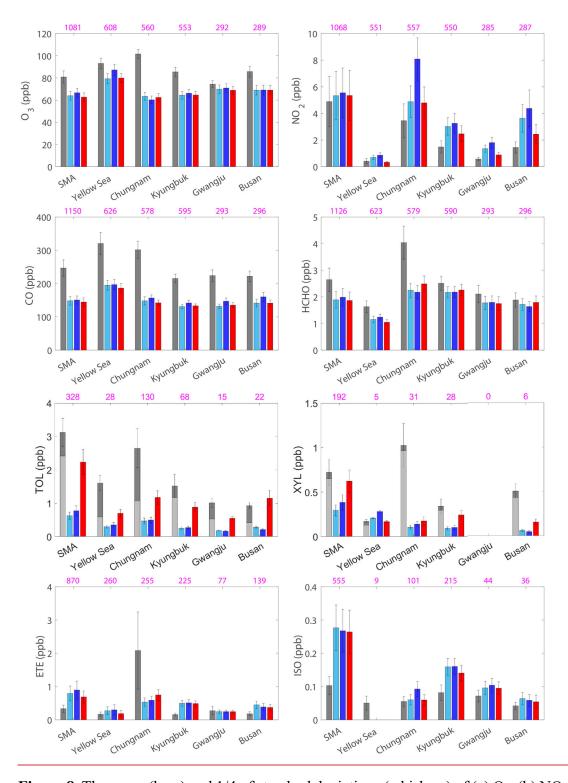


Figure 8. The mean (bars) and 1/4 of standard deviations (whiskers) of (a) O₃, (b) NO₂, (c) CO, (d) HCHO, (e) TOL, (f) XYL, (g) ethene (ETE), and (h) isoprene (ISO) (unit = ppb) from DC-8 (dark grey), EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) for each box are shown, respectively. TOL and XYL are calculated based on Table S68 (Supporting Information). The contribution of toluene to TOL and m/p-Xylene + o-Xylene to XYL is represented with light grey bars (e, f). The sampling numbers are represented with magenta color above the plots.

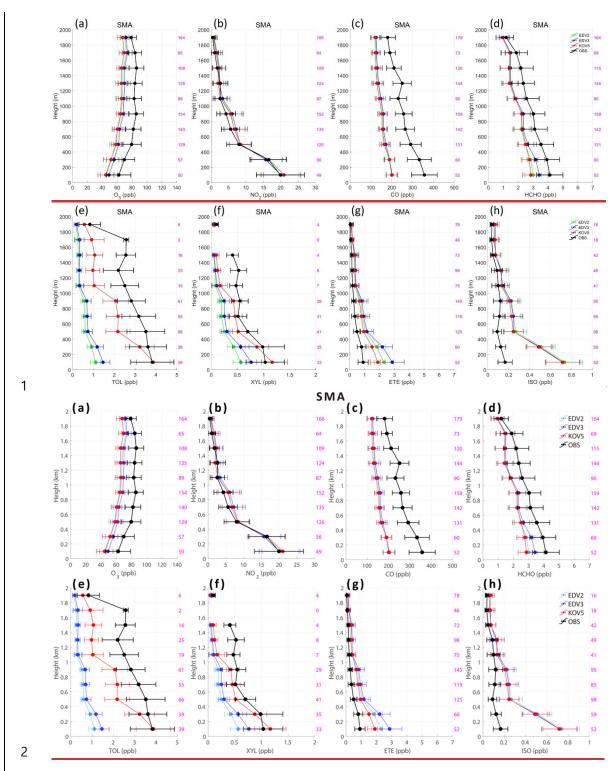


Figure 9. Vertically averaged (a) O₃, (b) NO₂, (c) CO, (d) HCHO, (e) TOL, (f) XYL, (g) ETE, and (h) ISO from DC-8 (black), EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) in SMA under 2 km height above ground level. The 1/2 of standard deviations are represented with black whiskers in each 200m layer. The sample number is presented with magenta color on the right side of the plots.

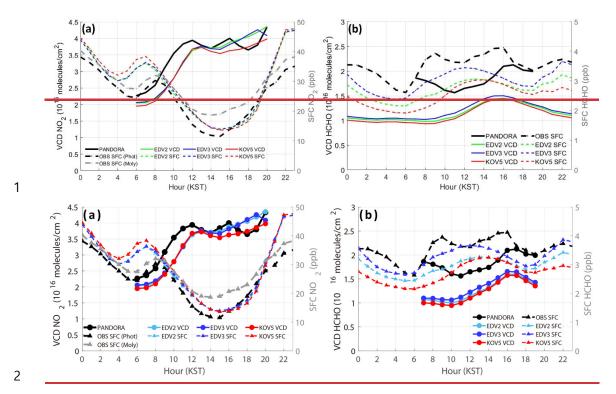


Figure 10. The diurnal cycles of vertical columns and surface concentrations of (a) NO₂ and (b) HCHO from Pandora spectrometer (column), and ground-based instruments (TEI 42i NO_x analyzer and Aerodyne QCL) at the Olympic Park site (37.5232°N, 127.126°E). Surface concentrations of NO₂ are obtained by the two methods: molybdenum converter and photolytic method. EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) are compared with observations. The WRF-Chem vertical column concentrations are produced by summing all vertical layers.

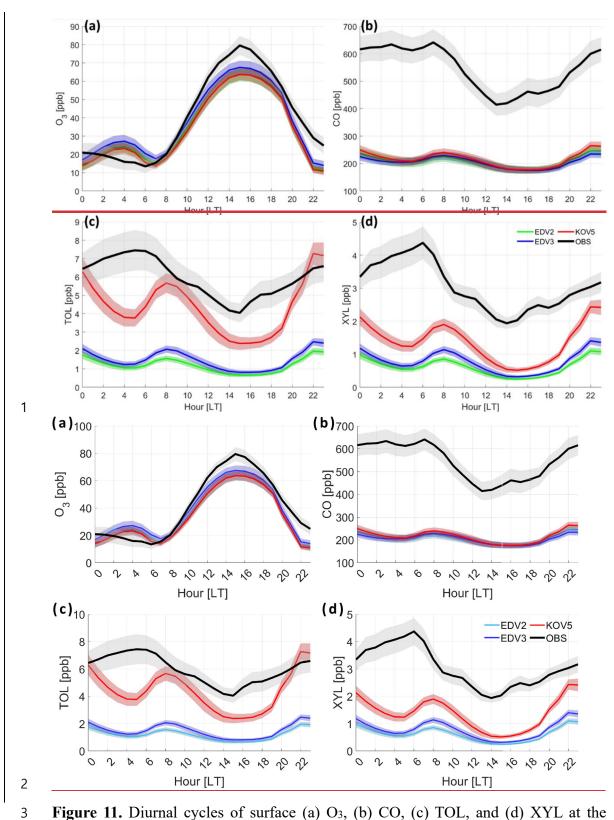


Figure 11. Diurnal cycles of surface (a) O₃, (b) CO, (c) TOL, and (d) XYL at the Olympic Park site. EDV2 (greensky blue), EDV3 (blue), and KOV5 (red) are compared with the observations. 1/4 of standard deviations are represented with grey shades. The average period is from the 11th May to the 10th June.

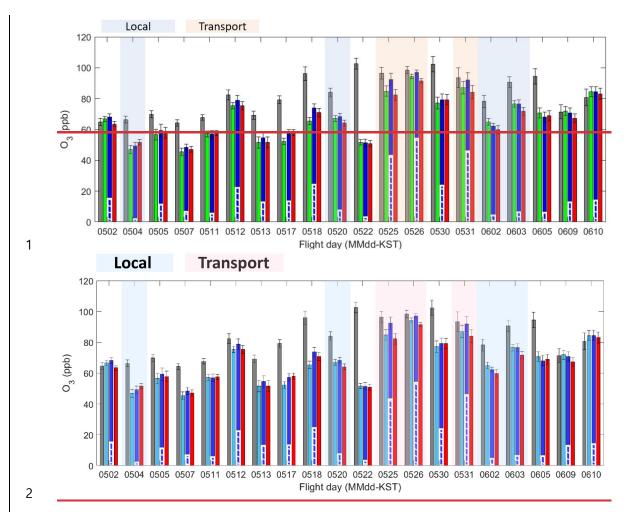
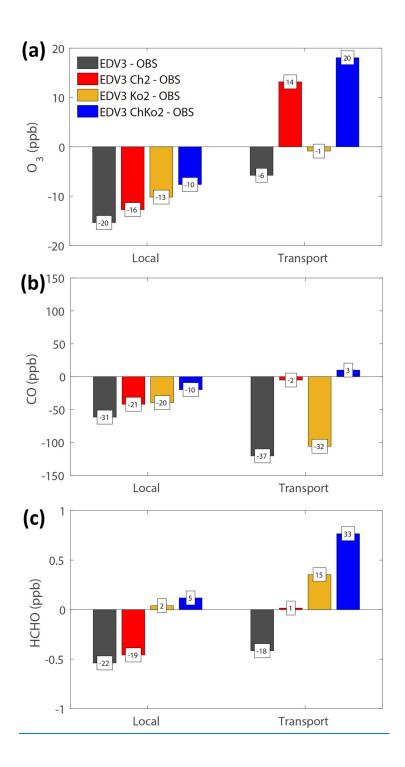


Figure 12. Averaged O₃ (bars) and 1/4 of standard deviations (whiskers) (unit: ppbv) for the 20 DC8 flights (under 2 km height). The observations (grey) are compared with the model results utilizing EDV2 (greensky blue), EDV3 (blue), and KOV5 (red). White hatch-filled bars over blue bars are the contribution of Chinese emissions to O₃ concentrations obtained from the default and sensitivity model runs with/without Chinese anthropogenic emissions. The Local (5/4,20 and 6/2,3) and Transport (5/25,26,31) cases are shaded with light blue and orange, respectively.



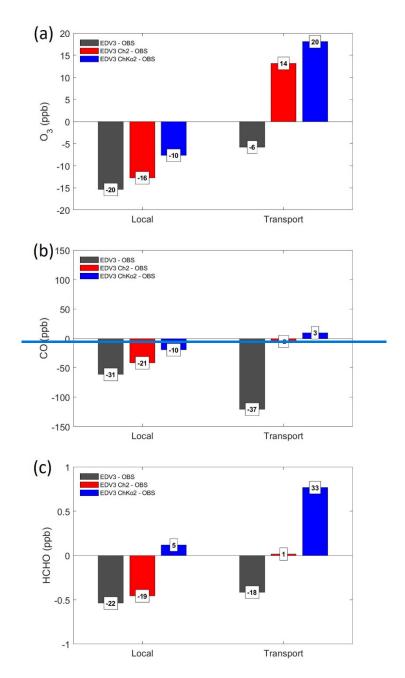


Figure 13. The biases in (a) the model O₃, (b) CO, and (c) HCHO concentrations (bars) relative to the DC-8 observations under 2 km height over SMA (dark gray: EDV3, red: EDV3 Ch2, orange: EDV3 Ko2, red: EDV3 ChKo2): (left panel) Local and (right panel) Transport case. Fractional differences (%) are shown in the white boxes.

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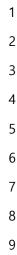
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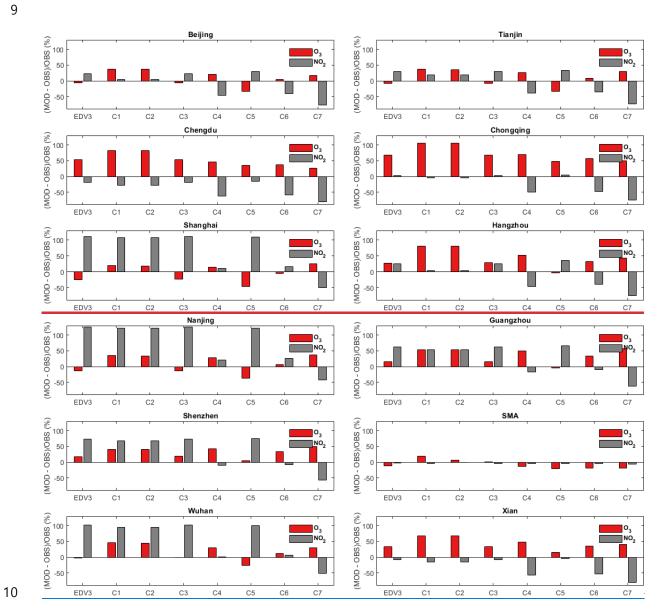
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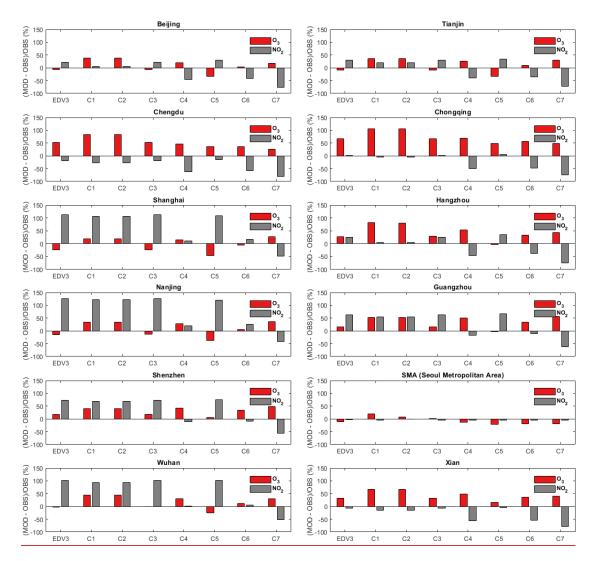


Figure 15. Same as Figure 14 except that the region is changed to cities; Beijing (39.4-41.1N, 115.4-117.5E), Tianjin (38.55-40.25N, 116.7-118.1E), Chengdu (30.05-31.5N, 103-105E), Chongqing (28.15-32.25N, 105.3-110.2E), Shanghai (30.7-31.5N, 120.85-122E), Hangzhou (29.2-30.6N, 118.3-120.9E), Nanjing (31.2-32.65N, 118.35-119.25E), Guangzhou (22.55-24N, 112.9-114.05E), Shenzhen (22.4-22.9N, 113.7-114.65E), SMA (37.2-37.8N, 126.5-127.3E), Wuhan (29.95-31.4N, 113.65-115.1E), and Xian (33.65-34.75N, 107.65-109.9E).