Supplementary of

- **Significant influence of oxygenated volatile organic compounds on atmospheric chemistry analysis: A case study in a typical industrial city in China**
- **Jingwen Dai et al.**
- *Correspondence*: Li Li (Lily@shu.edu.cn) and Kun Zhang (zk1231@shu.edu.cn)

Text S1 Adjustments of model parameter

 The base parameter settings of the OBM were shown in Sec 2.2. As formal studies have reported, PKU-Mo as a catalytic converter for NO² measurement can cause interferences from other nitrogen– 10 oxygen compounds (e.g., PAN, HNO₃), which can lead to an overestimation NO₂ by 30%~50% (Kim et al., 2015; Tan et al., 2017, 2019; Xu et al., 2013). In addition, strong anthropogenic emissions (such as vehicle emissions) around these sites, the model might not be able to reach the 13 steady state resulting in positive deviation (Li et al., 2014). Therefore, the observed $NO₂$ concentrations among those 5 sites were reduced by 30%~40% to compensate for interferences from catalytic converter (Xu et al., 2013), and NO steady-state approximations (NOss) was calculated (Del Negro et al., 1999), which were then used for OBM inputs. Simulation constrained by all 17 observed parameters including OVOCs and adjusted NO₂ and NO serves as the Base scenario. In order to investigate the impacts of model with OVOCs observationally constrained, scenario Free (Table S3) was conducted on top of the Base scenario by excluding observation constraints of OVOCs so that they were mainly generated by the oxidation of precursor VOCs in OBM.

Text S2 Model validation

22 The time series plot of modeled and observed O_3 [\(Figure S6\)](#page-8-0) and the model validation parameters 23 were shown in [\(Table S4\)](#page-18-0). The box model could be able to capture the diurnal variations of O_3 , but 24 the scenario M0 generally overestimated the O_3 peaks, which may be related to the overestimation 25 of NO₂. After adjusting for the observed NO₂ concentrations by cutting by 30%~40% (40% for ZL, 26 CQ and XD, 30% for CD and TZ) in scenario M1, the peak of O_3 in M1 was significantly decreased. It showed that the observed NO concentrations were significantly higher than the modeled NO in scenario M0 and M1. As discussed before, there was strong vehicle emissions nearby those sites especially during August 8-9, which may result in failing to reach steady stat. By further adjusting the input of NO to the approximate steady state concentration (NOss) in scenario M2 for each site, 31 the peak of O_3 was further reduced that counteracted the reduction of titration of O_3 by NO, and 32 closer to the observations overall [\(Figure S6](#page-8-0) (b)). The modeled O_3 remain underestimate or overestimate at some times during the daytime, and significantly underestimate at night [\(Figure](#page-12-0) [S11\)](#page-12-0). This discrepancy may be due to the limitations of the 0-D model to express the transport processes in the atmosphere due to its simplification of atmospheric physical processes. The simplification of physical process is acceptable for modeling of in situ photochemistry.

Text S3 Simulations of OVOCs in the Free scenario

 Similar to Base scenario but without OVOCs observationally constraints in the Free scenario, the hourly average concentration of OVOCs at five sites was 24.7±7.2 ppb, with a 59.1% overpredict of observations (15.5±11.3 ppb). OVOCs at CD (18.7±4.1 ppb), CQ (26.3±6.6 ppb), XD (24.7±7.0 ppb), and ZL (32.1±6.2 ppb) sites were overestimated in Free scenario by 81.4%, 88.4%, 42.1%, and 126.5%, respectively. The OVOCs concentrations in the atmosphere are subject to a combination of emission, secondary generation, and removal processes. Given that direct emissions of OVOCs are not considered in the Free scenario, the OVOCs concentrations in the model are determined by the generation and removal processes. In terms of the production process, it can be influenced by the emission of precursor VOCs indirectly. It has shown that in the presence of strong emission sources of VOCs, the model might not be able to reach an steady state, leading to a significant overestimation (Li et al., 2014). The observed OVOCs at TZ during August 8 were unusually high due to transient emissions [\(Figure S6](#page-8-0) (c)), pulling up the average levels. However, during the later days, the modeled OVOCs (15.5±10.7 ppb) were also higher (15.3%) than the 51 observed concentrations (13.4±11.5 ppb) consisting with the other sites.

 Figure S1 (a) Wind rose and (b) O³ pollution rose diagram of each site during the observation period.

 Figure S2 Time series of (a) meteorological parameters and (b) major pollutant mixing ratios at five sites in Zibo.

Figure S4 (a) Comparison and (b) spatial distribution of VOCs components among five sites.

 Figure S5 OVOCs accumulation and contributions from local net photochemical production and emissions/transport, and winds at (a) CD, (b) CQ, (c) TZ, (d) XD, and (e) ZL sites, respectively, and (f) time variations of REmis&Trans for all sites. RNetProd, REmis&Trans, RDep and RMeas in the legend represent local net O3 photochemical production, emissions and regional transport, deposition and observed OVOCs formation rates, respectively.

 Figure S6 Time series of O3, NO*^x* **from observations (Obs), simulations (Base and Free scenarios) and NO steady state (NOss), and that of OVOCs only including input species from observation and Free scenario.**

 Figure S7 (a) Simulated average daytime variation of RO*x* **(RO2, HO2, and OH) and NO³ radicals at five sites, and (b) the effects of OVOCs observationally constrains on radical concentrations, calculated by (Free – Base).**

 Figure S8 Average diurnal profiles of sources and sinks of (a) RO*^x* **and (b) O³ in the Base scenario.**

Figure S9 Daytime (8:00-18:00 LT) average budgets of RO*^x* **radicals (in ppb h-1) at each site in Base scenario and the difference between Free and Base scenario. The first values were the rates of Base, followed by the difference between Free and Base, where '-' means that the rate of Free scenario is lower than that of Base (in green), and conversely '+' means that the rate of Free is higher than that of Base (in orange). Primary RO***^x* **sources and sinks are in** 96 **red and blue, respectively, and the black lines represent the processes in** $\mathbb{R}O_\mathbf{x}$ **and** $\mathbb{N}O_\mathbf{x}$ **recycling.**

 Figure S10 Comparison of daytime atmospheric oxidation capacity (AOC) between Base and Free scenario.

 Figure S11 Heat map of O³ concentration difference (ΔO³ = Sim - Obs) between simulated and observed of Base and Free scenario

106 **Table S2 Comparison of VOC concentrations and compositions in this study with former studies.**

City	Site	Type	Period	Species	TVOCs	Alkanes	Alkenes	Aromatics	Acetylene	OVOCs	Halocarbons	References
Zibo	CD	Downwind	August 8-13, 2021	74	35.3	13.4	3.4	4.1	4.0	10.4		
	CQ	Upwind			42.6	16.9	3.9	7.5	0.5	13.9		
	TZ	Suburban			55.1	29.4	3.8	2.1	0.0	19.7		This study
	XD	Industrial			47.0	22.3	3.4	2.9	1.6	16.8		
	${\rm ZL}$	Urban			41.3	14.3	5.8	$6.2\,$	0.0	14.9		
	$\ensuremath{\mathsf{T}}\ensuremath{\mathsf{Z}}$	Suburban	High-O ₃ episodes in July 2019	56	58.1	43.8	3.7	5.5	3.1			
	${\bf BJ}$	Urban			23.8	13.8	3.2	3.7	2.4			(Li et al., 2021)
	XD	Suburban			38.1	22.5	7.8	3.4	$3.2\,$			
Qingdao		Rural	October 5 to	106	7.6	4.7	1.6	$0.6\,$	0.2	$0.4\,$		(Liu et al., 2021a)
			November 10, 2018									
Rizhao		Urban	December 2021 to	111	19.7	8.6	2.1	1.4	0.7	4.0	3.6	(Zhang et al., 2023)
			October 2022									
Jinan		Downtown	June 2010 to May 2012	55	25.3	14.3	$7.0\,$	$4.0\,$				(Wang et al., 2016)
Shanxi	LL		2019-2020	115	44.4	19.4	5.3	4.5	1.8	10.8	2.7	
	$\rm LF$	Urban			45.7	14.3	9.1	3.2	2.9	13.2	2.6	(Liu et al., 2023)
	YC				37.5	13.9	5.9	2.4	3.1	9.6	2.7	
Beijing		Urban	2018	99	29.1	12.4	2.9	2.1	2.1	6.4	$3.0\,$	(Li et al., 2022)
Tianjing		Suburban	November 1, 2018 to March 15, 2019	54	30.6	17.3	6.5	3.9	2.9			(Gu et al., 2020)
Xianghe			August 7-25, 2018		28.1	13.5	3.1	$6.0\,$		5.5		
	Suburban		December 1, 2018 to January 5, 2019	65	58.0	28.6	9.8	8.3		11.3		(Yang et al., 2021)

108 **Table S3 Summary of main meteorological parameters and average levels of pollutants** 109 **during the observation period.**

Parameter	CD	CQ	TZ	XD	${\rm ZL}$
WS (m s ⁻¹)	$2.1 + 1.2$	2.0 ± 1.0	1.9 ± 1.0	1.9 ± 1.1	1.5 ± 1.0
$T (^{\circ}C)$	27.3 ± 2.9	28.2 ± 2.9	27.1 ± 3.1	27.4 ± 3.3	27.3 ± 3.4
RH (%)	69.1 ± 11.8	70.4 ± 13.1	85.4 ± 17	74.6 ± 13.9	70.9±14.9
P(hPa)	984.6 ± 2.5	1005.2 ± 2.6	1005.6 ± 2.6	$1001.6{\pm}2.6$	996.1 ± 2.5
BLH(m)	421.2±512.2	451.5 ± 510.1	421.3 ± 465.1	463.9±541.2	443.8±528.3
SSR (10^5 J m^{-2})	6.7 ± 8.3	6.5 ± 8.1	6.3 ± 7.7	$6.6 + 8.2$	$6.6 + 8.2$
NO (ppb)	$3.9 + 4.7$	4.5 ± 7.8	$1.9 + 4.1$	$1.1 + 1.1$	2.6 ± 2.9
$NO2$ (ppb)	$10.8 + 5.1$	12.7 ± 8.1	10.4 ± 6.7	11.4 ± 6.7	14.8 ± 6.6
$SO2$ (ppb)	3.4 ± 1.3	$3.0 + 0.5$	2.2 ± 1.6	1.4 ± 1.3	3.9 ± 1.5
CO (ppb)	508 ± 173.6	1176.4±578.4	674.3 ± 190.9	1261.4 ± 1174.1	868 ± 258.3
O_3 (ppb)	58.6 ± 30.0	56.4 ± 34.2	51.0 ± 27.8	56.1 ± 29.4	57.4 ± 32.2
TVOCs (ppb)	35.7 ± 12.5	42.3 ± 15.4	58.5±35.0	49.6 ± 19.0	40.6 ± 10.3
Alkanes (ppb)	13.2 ± 6.2	16.5 ± 8.5	$30.2 + 21$	23.3 ± 11.2	13.5 ± 5.6
Alkenes (ppb)	3.3 ± 1.8	3.3 ± 1.6	2.9 ± 1.7	$2.8 + 1.3$	$5.6 + 3.0$
Aromatics (ppb)	4.0 ± 1.7	$7 + 3.6$	2.2 ± 1.2	3.1 ± 1.5	$6.3 + 4.7$
OVOCs (ppb)	$10.7 + 5$	14.5 ± 6.7	22.1 ± 22.5	17.9 ± 8.5	$14.6 + 4.8$
Isoprene (ppb)	$0.2 + 0.2$	$0.6 + 0.6$	1.1 ± 0.8	$0.7 + 0.5$	$0.6 + 0.7$
Alkyne (ppb)	$4.4 + 4.1$	$0.4 + 0.7$	0 ± 0	$1.9 + 1.6$	0 ± 0

		Base	Free		
Site	IOA	\mathbb{R}	IOA	\mathbb{R}	
CD	0.80	0.88	0.90	0.88	
CQ	0.90	0.87	0.86	0.87	
TZ	0.88	0.88	0.85	0.88	
XD	0.86	0.88	0.83	0.89	
ZL	0.88	0.89	0.88	0.87	

111 **Table S4 Modeled O³ assessment of Base and Free scenario.**

112 **Table S5 Comparison concentrations of the Base and Free scenario modeling parameters,**

113 **including OVOCs, O3, RO2, HO2, and OH) at the five sites.**

Note: Concentrations of OVOCs and O₃ in ppb, RO₂, HO₂ and OH in molecules cm⁻³.; $|\Delta O_3|$ = |Sim – Obs|.

Reference

 Chen, G., Liu, T., Chen, J., Xu, L., Hu, B., Yang, C., Fan, X., Li, M., Hong, Y., Ji, X., Chen, J., and 117 Zhang, F.: Atmospheric oxidation capacity and O₃ formation in a coastal city of southeast China: Results from simulation based on four-season observation, J. Environ. Sci., 136, 68–80, https://doi.org/10.1016/j.jes.2022.11.015, 2024.

 Del Negro, L. A., Fahey, D. W., Gao, R. S., Donnelly, S. G., Keim, E. R., Neuman, J. A., Cohen, R. C., Perkins, K. K., Koch, L. C., Salawitch, R. J., Lloyd, S. A., Proffitt, M. H., Margitan, J. J., Stimpfle, R. M., Bonne, G. P., Voss, P. B., Wennberg, P. O., McElroy, C. T., Swartz, W. H., Kusterer, T. L., Anderson, D. E., Lait, L. R., and Bui, T. P.: Comparison of modeled and observed values of NO² and JNO² during the Photochemistry of Ozone Loss in the Arctic Region in Summer (POLARIS) mission, J. Geophys. Res.-Atmos., 104, 26687–26703, https://doi.org/10.1029/1999JD900246, 1999.

 Gu, Y., Liu, B., Li, Y., Zhang, Y., Bi, X., Wu, J., Song, C., Dai, Q., Han, Y., Ren, G., and Feng, Y.: Multi-scale volatile organic compound (VOC) source apportionment in Tianjin, China, using a receptor model coupled with 1-hr resolution data, Environ. Pollut., 265, 115023, https://doi.org/10.1016/j.envpol.2020.115023, 2020.

 Han, Y., Huang, X., Wang, C., Zhu, B., and He, L.: Characterizing oxygenated volatile organic compounds and their sources in rural atmospheres in China, J. Environ. Sci., 81, 148–155, https://doi.org/10.1016/j.jes.2019.01.017, 2019.

 Hong, Z., Li, M., Wang, H., Xu, L., Hong, Y., Chen, J., Chen, J., Zhang, H., Zhang, Y., Wu, X., Hu, B., and Li, M.: Characteristics of atmospheric volatile organic compounds (VOCs) at a mountainous forest site and two urban sites in the southeast of China, Sci. Total Environ., 657, 1491–1500, https://doi.org/10.1016/j.scitotenv.2018.12.132, 2019.

 Huang, X.-F., Wang, C., Zhu, B., Lin, L.-L., and He, L.-Y.: Exploration of sources of OVOCs in various atmospheres in southern China, Environ. Pollut., 249, 831–842, https://doi.org/10.1016/j.envpol.2019.03.106, 2019.

 Hui, L., Liu, X., Tan, Q., Feng, M., An, J., Qu, Y., Zhang, Y., and Jiang, M.: Characteristics, source apportionment and contribution of VOCs to ozone formation in Wuhan, Central China, Atmos. Environ., 192, 55–71, https://doi.org/10.1016/j.atmosenv.2018.08.042, 2018.

 Kim, S., Kim, S.-Y., Lee, M., Shim, H., Wolfe, G. M., Guenther, A. B., He, A., Hong, Y., and Han, J.: Impact of isoprene and HONO chemistry on ozone and OVOC formation in a semirural South Korean forest, Atmos. Chem. Phys., 15, 4357–4371, https://doi.org/10.5194/acp-15-4357-2015, 2015.

 Li, C., Liu, Y., Cheng, B., Zhang, Y., Liu, X., Qu, Y., An, J., Kong, L., Zhang, Y., Zhang, C., Tan, Q., and Feng, M.: A comprehensive investigation on volatile organic compounds (VOCs) in 2018 150 in Beijing, China: Characteristics, sources and behaviours in response to O_3 formation, Sci. Total Environ., 806, 150247, https://doi.org/10.1016/j.scitotenv.2021.150247, 2022.

 Li, J., Zhai, C., Yu, J., Liu, R., Li, Y., Zeng, L., and Xie, S.: Spatiotemporal variations of ambient volatile organic compounds and their sources in Chongqing, a mountainous megacity in China, Sci. Total Environ., 627, 1442–1452, https://doi.org/10.1016/j.scitotenv.2018.02.010, 2018.

 Li, K., Wang, X., Li, L., Wang, J., Liu, Y., Cheng, X., Xu, B., Wang, X., Yan, P., Li, S., Geng, C., 156 Yang, W., Azzi, M., and Bai, Z.: Large variability of O_3 -precursor relationship during severe ozone polluted period in an industry-driven cluster city (Zibo) of North China Plain, J. Clean Prod., 316, 128252, https://doi.org/10.1016/j.jclepro.2021.128252, 2021.

 Li, X., Rohrer, F., Brauers, T., Hofzumahaus, A., Lu, K., Shao, M., Zhang, Y. H., and Wahner, A.: Modeling of HCHO and CHOCHO at a semi-rural site in southern China during the PRIDE-

 PRD2006 campaign, Atmos. Chem. Phys., 14, 12291–12305, https://doi.org/10.5194/acp-14- 12291-2014, 2014.

 Li, Y., Yin, S., Yu, S., Yuan, M., Dong, Z., Zhang, D., Yang, L., and Zhang, R.: Characteristics, 164 source apportionment and health risks of ambient VOCs during high ozone period at an urban site
165 in central plain, China, Chemosphere, 250, 126283. in central plain, China, Chemosphere, 250, 126283, https://doi.org/10.1016/j.chemosphere.2020.126283, 2020.

- Liu, Y., Shen, H., Mu, J., Li, H., Chen, T., Yang, J., Jiang, Y., Zhu, Y., Meng, H., Dong, C., Wang, W., and Xue, L.: Formation of peroxyacetyl nitrate (PAN) and its impact on ozone production in the coastal atmosphere of Qingdao, North China, Sci. Total Environ., 778, 146265, https://doi.org/10.1016/j.scitotenv.2021.146265, 2021a.
- Liu, Y., Wang, H., Jing, S., Peng, Y., Gao, Y., Yan, R., Wang, Q., Lou, S., Cheng, T., and Huang, C.: Strong regional transport of volatile organic compounds (VOCs) during wintertime in Shanghai megacity of China, Atmos. Environ., 244, 117940, https://doi.org/10.1016/j.atmosenv.2020.117940, 2021b.
- Liu, Y., Qiu, P., Xu, K., Li, C., Yin, S., Zhang, Y., Ding, Y., Zhang, C., Wang, Z., Zhai, R., Deng, Y., Yan, F., Zhang, W., Xue, Z., Sun, Y., Ji, D., Li, J., Chen, J., Tian, H., Liu, X., and Zhang, Y.:
- 177 Analysis of VOC emissions and O₃ control strategies in the Fenhe Plain cities, China, J. Environ.
- Manage, 325, 116534, https://doi.org/10.1016/j.jenvman.2022.116534, 2023.
- Ma, Z., Liu, C., Zhang, C., Liu, P., Ye, C., Xue, C., Zhao, D., Sun, J., Du, Y., Chai, F., and Mu, Y.: The levels, sources and reactivity of volatile organic compounds in a typical urban area of Northeast China, J. Environ. Sci., 79, 121–134, https://doi.org/10.1016/j.jes.2018.11.015, 2019.
- Simayi, M., Shi, Y., Xi, Z., Li, J., Yu, X., Liu, H., Tan, Q., Song, D., Zeng, L., Lu, S., and Xie, S.: Understanding the sources and spatiotemporal characteristics of VOCs in the Chengdu Plain, China, through measurement and emission inventory, Sci. Total Environ., 714, 136692, https://doi.org/10.1016/j.scitotenv.2020.136692, 2020.
- Song, M., Tan, Q., Feng, M., Qu, Y., Liu, X., An, J., and Zhang, Y.: Source Apportionment and Secondary Transformation of Atmospheric Nonmethane Hydrocarbons in Chengdu, Southwest China, J. Geophys. Res.-Atmos., 123, 9741–9763, https://doi.org/10.1029/2018JD028479, 2018.
- Song, M., Li, X., Yang, S., Yu, X., Zhou, S., Yang, Y., Chen, S., Dong, H., Liao, K., Chen, Q., Lu, K., Zhang, N., Cao, J., Zeng, L., and Zhang, Y.: Spatiotemporal variation, sources, and secondary transformation potential of volatile organic compounds in Xi'an, China, Atmos. Chem. Phys., 21, 4939–4958, https://doi.org/10.5194/acp-21-4939-2021, 2021.
- Tan, Z., Fuchs, H., Lu, K., Hofzumahaus, A., Bohn, B., Broch, S., Dong, H., Gomm, S., Haeseler, R., He, L., Holland, F., Li, X., Liu, Y., Lu, S., Rohrer, F., Shao, M., Wang, B., Wang, M., Wu, Y., Zeng, L., Zhang, Y., Wahner, A., and Zhang, Y.: Radical chemistry at a rural site (Wangdu) in the 196 North China Plain: observation and model calculations of OH, HO₂ and RO₂ radicals, Atmos. Chem.
197 Phys., 17, 663–690, https://doi.org/10.5194/acp-17-663-2017, 2017. Phys., 17, 663–690, https://doi.org/10.5194/acp-17-663-2017, 2017.
- Tan, Z., Lu, K., Jiang, M., Su, R., Wang, H., Lou, S., Fu, Q., Zhai, C., Tan, Q., Yue, D., Chen, D., Wang, Z., Xie, S., Zeng, L., and Zhang, Y.: Daytime atmospheric oxidation capacity in four Chinese 200 megacities during the photochemically polluted season: a case study based on box model simulation,
201 Mtmos. Chem. Phys., 19, 3493–3513, https://doi.org/10.5194/acp-19-3493-2019, 2019. Atmos. Chem. Phys., 19, 3493–3513, https://doi.org/10.5194/acp-19-3493-2019, 2019.
- Wang, N., Li, N., Liu, Z., and Evans, E.: Investigation of chemical reactivity and active components of ambient VOCs in Jinan, China, Air Qual Atmos Health, 9, 785–793, https://doi.org/10.1007/s11869-015-0380-1, 2016.
- 205 Wang, X., Yin, S., Zhang, R., Yuan, M., and Ying, Q.: Assessment of summertime O₃ formation and the O3-NO*x*-VOC sensitivity in Zhengzhou, China using an observation-based model, Sci. Total Environ., 813, 152449, https://doi.org/10.1016/j.scitotenv.2021.152449, 2022.
- Xu, K., Liu, Y., Li, C., Zhang, C., Liu, X., Li, Q., Xiong, M., Zhang, Y., Yin, S., and Ding, Y.: Enhanced secondary organic aerosol formation during dust episodes by photochemical reactions in the winter in Wuhan, J. Environ. Sci., 133, 70–82, https://doi.org/10.1016/j.jes.2022.04.0181001- 0742, 2023.
- Xu, Z., Wang, T., Xue, L. K., Louie, P. K. K., Luk, C. W. Y., Gao, J., Wang, S. L., Chai, F. H., and Wang, W. X.: Evaluating the uncertainties of thermal catalytic conversion in measuring atmospheric nitrogen dioxide at four differently polluted sites in China, Atmos. Environ., 76, 221–226, https://doi.org/10.1016/j.atmosenv.2012.09.043, 2013.
- Yang, Y., Shao, M., Keßel, S., Li, Y., Lu, K., Lu, S., Williams, J., Zhang, Y., Zeng, L., Nölscher, A. C., Wu, Y., Wang, X., and Zheng, J.: How the OH reactivity affects the ozone production efficiency: case studies in Beijing and Heshan, China, Atmos. Chem. Phys., 17, 7127–7142, https://doi.org/10.5194/acp-17-7127-2017, 2017.
- Yang, Y., Wang, Y., Huang, W., Yao, D., Zhao, S., Wang, Y., Ji, D., Zhang, R., and Wang, Y.: Parameterized atmospheric oxidation capacity and speciated OH reactivity over a suburban site in the North China Plain: A comparative study between summer and winter, Sci. Total Environ., 773, 145264, https://doi.org/10.1016/j.scitotenv.2021.145264, 2021.
- Zhang, Z., Sun, Y., and Li, J.: Characteristics and sources of VOCs in a coastal city in eastern China
- 225 and the implications in secondary organic aerosol and O_3 formation, Sci. Total Environ., 887, 164117, https://doi.org/10.1016/j.scitotenv.2023.164117, 2023.