

Supporting Information

Impact of anthropogenic emissions on biogenic secondary organic aerosol: Observation in the Pearl River Delta, South China

Yu-Qing Zhang^{1, *}, Duo-Hong Chen^{2, *}, Xiang Ding^{1, †}, Jun Li¹, Tao Zhang², Jun-Qi Wang¹, Qian Cheng^{1, 3}, Hao Jiang¹, Wei Song¹, Yu-Bo Ou², Peng-Lin Ye³, Gan Zhang¹, Xin-Ming Wang^{1, 4}

1 State Key Laboratory of Organic Geochemistry and Guangdong Provincial Key Laboratory of Environmental Protection and Resources Utilization, Guangzhou Institute of Geochemistry, Chinese Academy of Sciences, Guangzhou, 510640, China

2 State Environmental Protection Key Laboratory of Regional Air Quality Monitoring, Environmental Monitoring Center of Guangdong Province, Guangzhou, 510308, China

3 Aerodyne Research Inc., Billerica, Massachusetts 01821, United States

4 Center for Excellence in Regional Atmospheric Environment, Institute of Urban Environment, Chinese Academy of Sciences, Xiamen, 361021, China

* These authors contributed equally to this work.

† *Correspondence to:* Xiang Ding (xiangd@gig.ac.cn)

Contents of this file

Text S1

Figure S1 to S6

Table S1 to S5

Text S1 SOA-tracer method for source apportionment

The SOA-tracer method is developed by Kleindienst and co-workers. Based on chamber experiments, they determine the mass fractions of tracers in SOA (f_{SOA}) and SOC (f_{SOC}) for individual precursor:

$$f_{SOA} = \frac{\sum_i [tr_i]}{[SOA]}, \quad f_{SOC} = \frac{\sum_i [tr_i]}{[SOC]}$$

where $\sum_i [tr_i]$ is the sum of tracer concentrations for a precursor, and [SOA] and [SOC] are the measured SOA and SOC concentrations in chamber-generated SOA samples. The available f_{SOA} and f_{SOC} values were listed in Table S2. With these mass fractions in literatures and measured SOA tracers in the ambient air, SOA and SOC from different precursors have been estimated in different places of the world (Hu et al., 2008; Lewandowski et al., 2013; Stone et al., 2012; von Schneidmesser et al., 2009; Ding et al., 2014), with the assumption that the f_{SOA} and f_{SOC} values in the chamber samples are the same in the ambient air. In this study, the same set of SOA tracers reported by Kleindienst and co-workers were used for the SOC and SOA estimations (Table S2).

The uncertainty in the SOA-tracer method is induced from the analysis of organic tracers and the determination of conversion factors. The uncertainties in the tracers' analyses were estimated in the range of 15-157% (Table S2). The uncertainties in f_{SOA} were reported to be 25% for isoprene, 48% for monoterpenes, and 22% for β -caryophyllene (Kleindienst et al., 2007; Lewandowski et al., 2013). Considering these factors, the uncertainty of the estimating procedure was calculated through error propagation. The relative standard deviations (RSD) were 37% for SOA_I , 67% for SOA_M , and 158% for SOA_C . On average, the RSD of total BSOA (sum of the three BVOCs) was 59%.

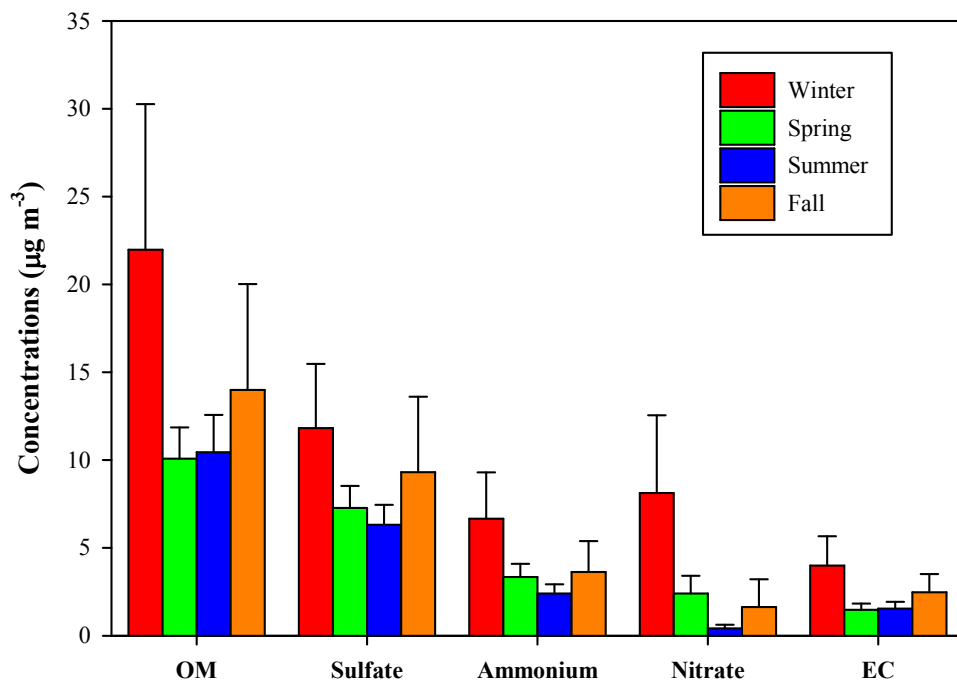


Figure S1 Seasonal variation of major components in PM_{2.5}. All the major components increased in winter and fall.

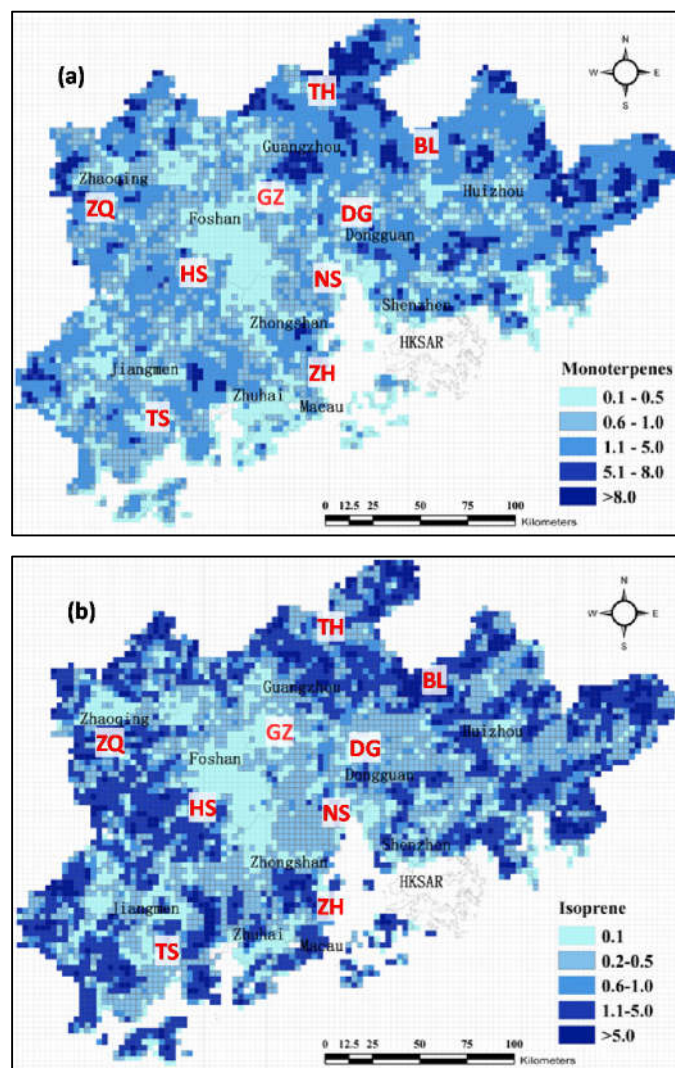


Figure S2 Spatial distribution of monoterpenes (a), and isoprene (b) emissions in the PRD (Zheng et al., 2010). The sampling sites are labeled.

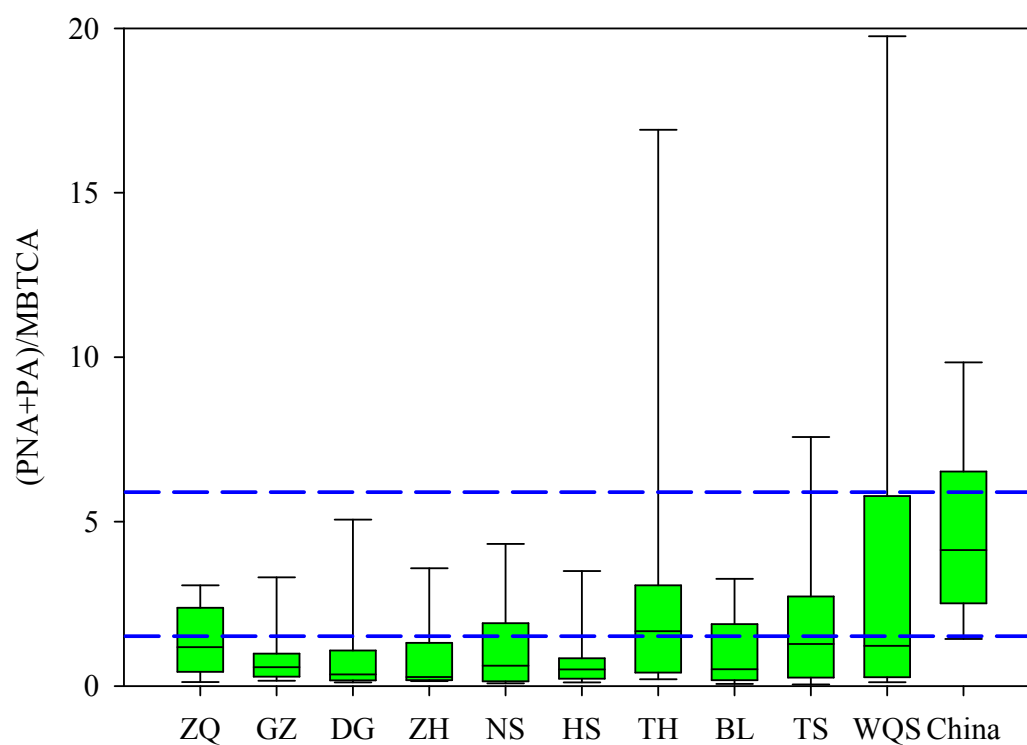


Figure S3 Spatial distribution of (PNA+PA)/MBTCA ratios at 9 sites in the PRD. The (PNA+PA)/MBTCA ratios between two blue dash lines (1.51–5.91) indicate fresh SOA_M from chamber studies (Eddingsaas et al., 2012; Offenberg et al., 2007). Box with error bars represent 10th, 25th, 75th, 90th percentiles at each site. The line in the box is the median at each site. The data at WQS site during 2008 in the PRD (Ding et al., 2012) and at 12 sites during 2012-2013 in China (Ding et al., 2016) were reported in our previous studies.

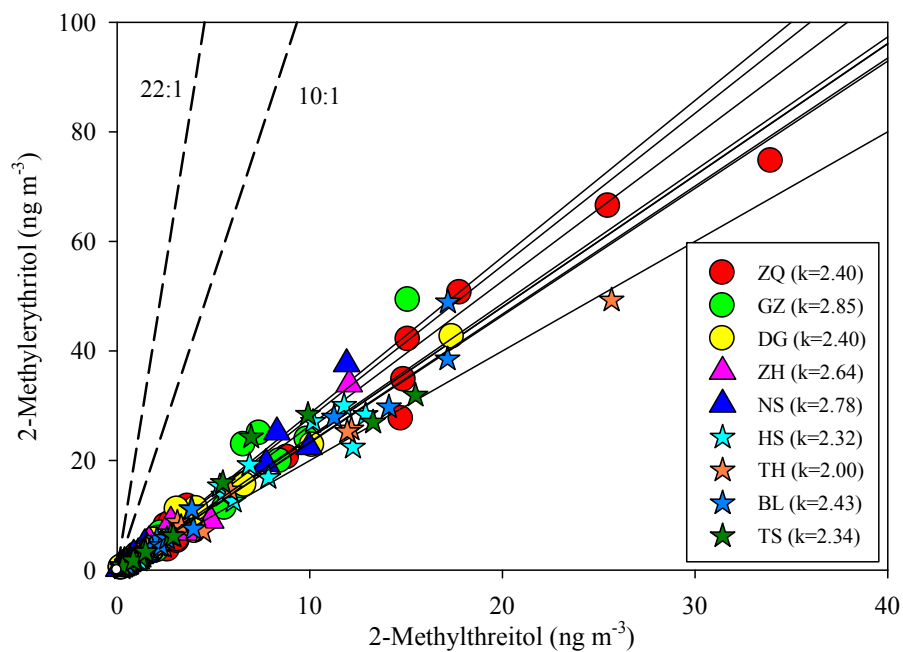


Figure S4 Significant correlations between 2-methyltetrol isomers at 9 sites in the PRD. K indicates the slope of each linear regression. The dash lines indicate the ratio range of 2-methyltetrol isomers in the SOA from isoprene ozonolysis (Riva et al., 2016).

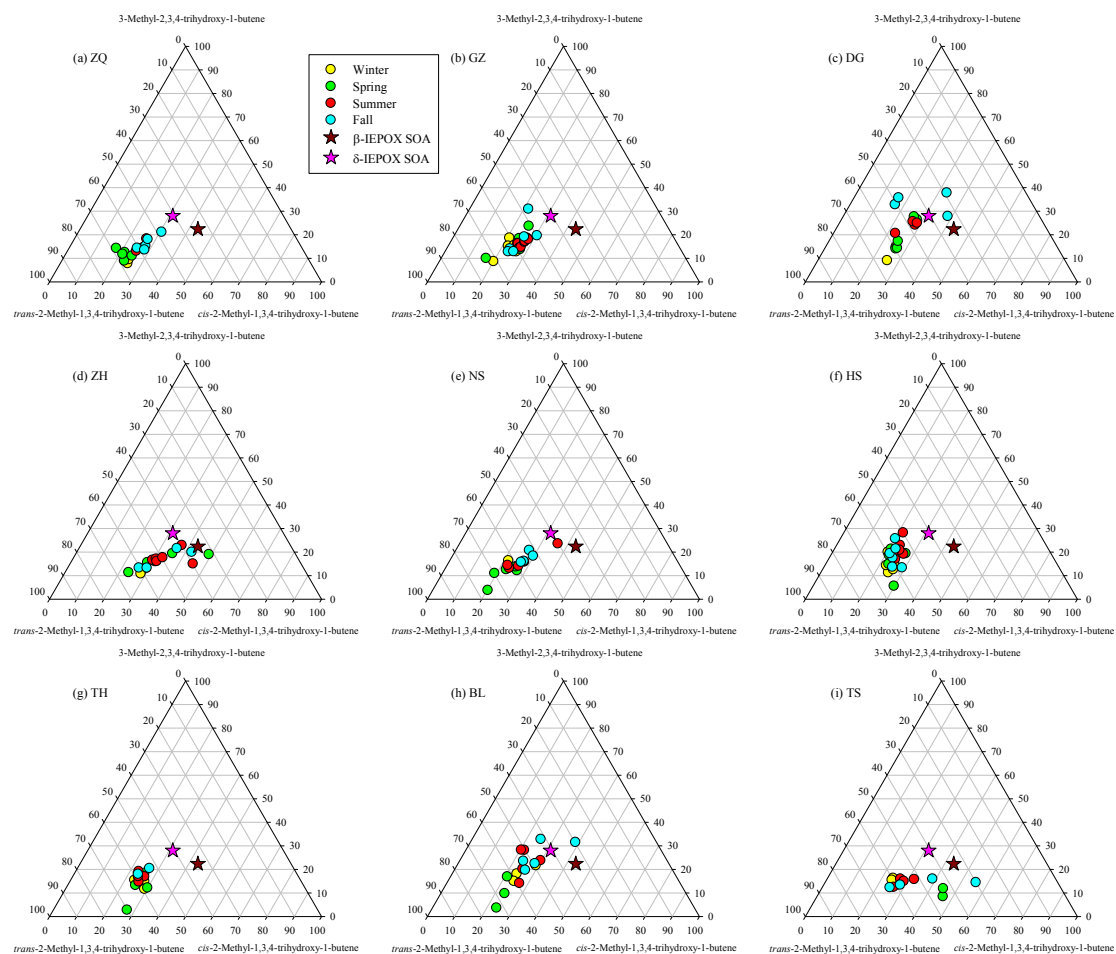


Figure S5 Intercomparison of C₅-alkene triols compositions at 9 sites and in β -IEPOX and δ -IEPOX derived SOA (Lin et al., 2012).

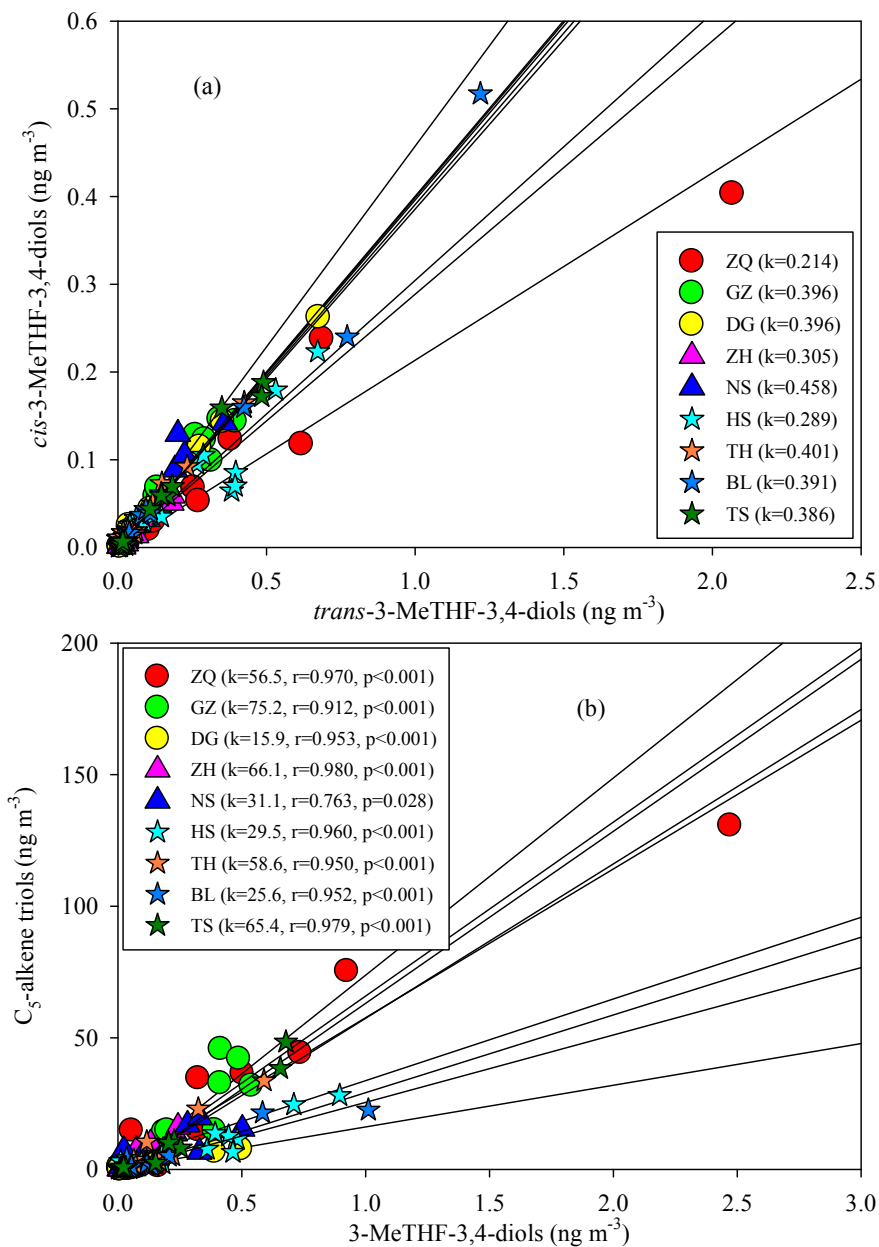


Figure S6 Significant correlations among the SOA_i tracers. K indicates the slope of each linear regression.

Table S2 SOA tracers and f_{SOA} and f_{SOC} values for SOA estimation

	Monoterpenes ^a	Isoprene ^b	β -Caryophyllene ^b
SOA Tracers ^c	PNA (15%) ^d PA (34%) ^d MBTCA (62%) ^d HGA (96%) ^d HDMGA (67%) ^d	2-MTLs (41%) ^d 2-MGA (43%) ^d 3-MeTHF-3,4-diols (52%) C ₅ -alken triols (93%)	CA (157%) ^d
f_{SOA} ($\mu\text{g } \mu\text{g}^{-1}$)	0.044 (48%) ^e	0.063 (25%) ^e	0.0109 (22%) ^e
f_{SOC} ($\mu\text{g } \mu\text{gC}^{-1}$)	0.059	0.155	0.023

^a The f_{SOA} and f_{SOC} values for monoterpenes are calculated based on the data reported by Offenberg et al. (2007). ^b The f_{SOA} and f_{SOC} values for isoprene, and β -caryophyllene are reported by Kleindienst et al. (2007). ^c The numbers in brackets are uncertainties in tracer measurement. ^d These tracers are used to calculate f_{SOA} and estimate ambient SOA. ^e The numbers in brackets are the uncertainties of f_{SOA} values reported by Kleindienst et al. (2007).

Table S3 Correlation analysis of HO₂-channel SOA_I tracers with O₃

	Coefficient (r)	<i>p</i> -value
3-MeTHF-3,4-diols	0.343	<0.001
C ₅ -alkene triols	0.388	<0.001
2-Methyltetrols	0.386	<0.001
HO ₂ -chanle SOA _I tracers	0.409	<0.001

Table S4 Correlations among HO₂-channel SOA_I tracers

	3-MeTHF-3,4-diols	C ₅ -alkene triols	2-Methyltetrols
3-MeTHF-3,4-diols	1	0.789	0.792
C ₅ -alkene triols		1	0.787
2-Methyltetrols			1

All the correlations are significant ($p < 0.001$)

Table S5 Rate constants and lifetimes of SOA precursors

	α -Pinene	β -Pinene	Isoprene	β -Caryophyllene
	Rate constants at 298 K ($\text{cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$) ^a			
OH	5.25×10^{-11}	7.88×10^{-11}	9.99×10^{-11}	1.97×10^{-10}
O₃	9.01×10^{-17}	1.50×10^{-17}	1.28×10^{-17}	1.16×10^{-14}
	Lifetimes (hrs) ^b			
OH	0.53	0.35	0.28	0.14
O₃	3.64	21.9	25.7	0.03

^a Rate constants are provided by MCMv3.2 (<http://mcm.leeds.ac.uk/MCMv3.2>).

^b Lifetimes are estimated using summer average concentration of OH radical ($\sim 1 \times 10^7 \text{ molecules cm}^{-3}$) in the PRD (Hofzumahaus et al., 2009), and annual average O₃ concentration ($67.7 \mu\text{g m}^{-3}$) in Table S1.

References:

- Ding, X., Wang, X., Gao, B., Fu, X., He, Q., Zhao, X., Yu, J., and Zheng, M.: Tracer based estimation of secondary organic carbon in the Pearl River Delta, South China, *J. Geophys. Res-Atmos.*, 117, D05313, 10.1029/2011JD016596, 2012.
- Ding, X., He, Q. F., Shen, R. Q., Yu, Q. Q., and Wang, X. M.: Spatial distributions of secondary organic aerosols from isoprene, monoterpenes, β -caryophyllene, and aromatics over China during summer, *J. Geophys. Res-Atmos.*, 119, 11877-11891, 10.1002/2014JD021748, 2014.
- Ding, X., Zhang, Y. Q., He, Q. F., Yu, Q. Q., Shen, R. Q., Zhang, Y., Zhang, Z., Lyu, S. J., Hu, Q. H., Wang, Y. S., Li, L. F., Song, W., and Wang, X. M.: Spatial and seasonal variations of secondary organic aerosol from terpenoids over China, *J. Geophys. Res-Atmos.*, 121, 14661–14678, 10.1002/2016JD025467, 2016.
- Eddingsaas, N. C., Loza, C. L., Yee, L. D., Chan, M., Schilling, K. A., Chhabra, P. S., Seinfeld, J. H., and Wennberg, P. O.: α -pinene photooxidation under controlled chemical conditions – Part 2: SOA yield and composition in low- and high-NO_x environments, *Atmos. Chem. Phys.*, 12, 7413-7427, 10.5194/acp-12-7413-2012, 2012.
- Hofzumahaus, A., Rohrer, F., Lu, K., Bohn, B., Brauers, T., Chang, C.-C., Fuchs, H., Holland, F., Kita, K., Kondo, Y., Li, X., Lou, S., Shao, M., Zeng, L., Wahner, A., and Zhang, Y.: Amplified trace gas removal in the troposphere, *Science*, 324, 1702-1704, 10.1126/science.1164566, 2009.
- Hu, D., Bian, Q., Li, T. W. Y., Lau, A. K. H., and Yu, J. Z.: Contributions of isoprene, monoterpenes, β -caryophyllene, and toluene to secondary organic aerosols in Hong Kong during the summer of 2006, *J. Geophys. Res-Atmos.*, 113, D22206, 10.1029/2008JD010437, 2008.
- Kleindienst, T. E., Jaoui, M., Lewandowski, M., Offenberg, J. H., Lewis, C. W., Bhave, P. V., and Edney, E. O.: Estimates of the contributions of biogenic and anthropogenic hydrocarbons to secondary organic aerosol at a southeastern US location, *Atmos. Environ.*, 41, 8288-8300, 10.1016/j.atmosenv.2007.06.045, 2007.
- Lewandowski, M., Piletic, I. R., Kleindienst, T. E., Offenberg, J. H., Beaver, M. R., Jaoui, M., Docherty, K. S., and Edney, E. O.: Secondary organic aerosol characterisation at field sites across the United States during the spring–summer period, *Int. J. Environ. An. Ch.*, 93, 1084-1103, 10.1080/03067319.2013.803545, 2013.
- Lin, Y.-H., Zhang, Z., Docherty, K. S., Zhang, H., Budisulistiorini, S. H., Rubitschun, C. L., Shaw, S., Knipping, E., Edgerton, E. S., Kleindienst, T. E., Gold, A., and Surratt, J. D.: Isoprene epoxydiols as precursors to secondary organic aerosol formation: Acid-catalyzed reactive uptake studies with authentic standards, *Environ. Sci. Technol.*, 46, 189-195, 10.1021/es202554c, 2012.
- Offenberg, J. H., Lewis, C. W., Lewandowski, M., Jaoui, M., Kleindienst, T. E., and Edney, E. O.: Contributions of toluene and α -pinene to SOA formed in an irradiated toluene/ α -pinene/NO_x/ air mixture: Comparison of results using ¹⁴C content and SOA organic tracer methods, *Environ.Sci. Technol.*, 41, 3972-3976, 10.1021/es070089+, 2007.
- Riva, M., Budisulistiorini, S. H., Zhang, Z., Gold, A., and Surratt, J. D.: Chemical characterization of secondary organic aerosol constituents from isoprene ozonolysis in the presence of acidic aerosol, *Atmos. Environ.*, 130, 5-13, 10.1016/j.atmosenv.2015.06.027, 2016.

- Stone, E. A., Nguyen, T. T., Pradhan, B. B., and Man Dangol, P.: Assessment of biogenic secondary organic aerosol in the Himalayas, *Environ. Chem.*, 9, 263-272, 10.1071/EN12002, 2012.
- Von Schneidmesser, E., Zhou, J., Stone, E. A., Schauer, J. J., Shpund, J., Brenner, S., Qasrawi, R., Abdeen, Z., and Sarnat, J. A.: Spatial variability of carbonaceous aerosol concentrations in east and west Jerusalem, *Environ. Sci. Technol.*, 44, 1911-1917, 10.1021/es9014025, 2009.
- Zheng, J., Zheng, Z., Yu, Y., and Zhong, L.: Temporal, spatial characteristics and uncertainty of biogenic VOC emissions in the Pearl River Delta region, China, *Atmos. Environ.*, 44, 1960-1969, 10.1016/j.atmosenv.2010.03.001, 2010.