

## Ozone sensitivity analysis with the MM5-CMAQ modeling system for Shanghai

Li Li<sup>1,2</sup>, Changhong Chen<sup>2,\*</sup>, Cheng Huang<sup>2</sup>, Haiying Huang<sup>2</sup>, Gangfeng Zhang<sup>2</sup>, Yangjun Wang<sup>1</sup>,  
Minghua Chen<sup>2</sup>, Hongli Wang<sup>2</sup>, Yiran Chen<sup>2</sup>, D. G. Streets<sup>3</sup>, Jiamo Fu<sup>1</sup>

1. Institute of Environmental Pollution and Health, School of Environmental and Chemical Engineering, Shanghai University,  
Shanghai 200444, China. E-mail: [lili@saes.sh.cn](mailto:lili@saes.sh.cn)

2. Shanghai Academy of Environmental Sciences, Shanghai 200233, China

3. Decision and Information Sciences Division, Argonne National Laboratory, Argonne, IL 60439, USA

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### Abstract

Ozone has become one of the most important air pollution issues around the world. This article applied both  $O_3/(NO_y-NO_x)$  and  $H_2O_2/HNO_3$  indicators to analyze the ozone sensitivity in urban and rural areas of Shanghai, with implementation of the MM5-CMAQ modeling system in July, 2007. The meteorological parameters were obtained by using the MM5 model. A regional emission inventory with spatial and temporal allocation based on the statistical data has been developed to provide input emission data to the MM5-CMAQ modeling system. Results showed that the ozone concentrations in Shanghai show clear regional differences. The ozone concentration in rural areas was much higher than that in the urban area. Two indicators showed that ozone was more sensitive to VOCs in urban areas, while it tended to be  $NO_x$  sensitive in rural areas of Shanghai.

**Key words:** ozone sensitivity; indicators; Shanghai

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### Introduction

Shanghai is the most important economic center in China. It hosts around 19 million residents and occupies 6340 km<sup>2</sup>. The GDP and per capita GDP of Shanghai reached 1505 billion CNY and 78,989 CNY, respectively, in 2009 (Shanghai Statistical Yearbook, 2010). This represents an increase of almost 2.5 times during the period 2000–2009 (Shanghai Statistical Yearbook, 2001). Such a fast pace of economic growth drives high growth in energy consumption, and thus huge air pollutant emissions (Zhang et al., 2009), which causes the secondary air pollution becoming worse. Regional visibility is decreasing and ozone concentration is increasing, especially in summer. Ozone is considered to be one of the most serious air pollutants of concern in the United States, as well as in most metropolitan areas around the world (Streets et al., 2007), and it has attracted much interest for its impact on people's health. Many observations (Shan et al., 2008; Wang et al., 2001, 2005) show that high ozone concentrations have begun to appear in eastern China. Wang et al. (2003) found that the occurrence frequency of ozone concentration higher than 160  $\mu\text{g}/\text{m}^3$  reached 20% in some sites of the Yangtze River Delta (YRD). Tang et al. (2008)

found that 167 high ozone days with daily one hour ozone concentrations higher than 104  $\mu\text{g}/\text{m}^3$  occurred in 2006 in Shanghai.

Ozone is formed through a series of complex chemical reactions from the mixture of reactive volatile organic compounds (VOCs) and nitrogen oxides ( $NO_x$ ). The ratio of these two ingredients determines how much ozone is formed. Ozone increases rapidly between 9:00 am and 3:00 pm related to the average lifetime of the most reactive VOC compounds, which dominate the reactivity of hydroxyl radical ( $OH\cdot$ ), and start to decrease in the late afternoon, when its loss processes catch up with formation. During the nighttime, continuous nitrogen monoxide ( $NO$ ) emissions gradually titrate ozone, forming high  $NO_2$  mixing ratios. Under significant ozone and  $NO_2$  abundances, these species slowly react to form  $NO_3\cdot$  radicals, which serves as equivalent to daytime  $OH\cdot$  at night, oxidizing hydrocarbons. In this way,  $NO_x$  can be oxidized to  $NO_z$  at night. Once the sun comes up,  $NO_3$  is rapidly photolyzed. Thus, the control of  $O_3$  is a complicated problem due to the nature of the non-linear formation of  $O_3$  (Seinfeld and Pandis, 1998). Due to different emission characteristics of the precursors, the control of ozone may differ with regions. Many studies on the complex formation processes of ozone have been made (Geng et al., 2007; Ran et al., 2011; Tang, 2004; Tie et al., 2006; Wang and Li, 2002; Xu et al., 2006;

\* Corresponding author. E-mail: [chench@saes.sh.cn](mailto:chench@saes.sh.cn)

Xu and Zhang, 2006; Zhao et al., 2004;). However, since Shanghai occupies a large area, the emissions rates, emission characteristics of the ozone precursors vary greatly. This means that the ozone formation mechanism differs with areas in Shanghai. Studies on the ozone sensitivity at different sites in Shanghai are quite limited up to now. With development of the numerical air quality models, using indicators to study the ozone sensitivity in different regions becomes more possible. In this study, we chose two sites representing the rural and urban areas of Shanghai respectively, and applied  $\text{H}_2\text{O}_2/\text{HNO}_3$  and  $\text{O}_3/(\text{NO}_y\text{-NO}_x)$  ratios to study the ozone sensitivity characteristics. The results are helpful for guiding the development of ozone control strategies to be made in Shanghai.

## 1 Materials and methods

The methodology used in this article is to simulate the atmospheric processes over both Shanghai and its surrounding areas using the Models-3/Community Multi-scale Air Quality (CMAQ) modeling system (Version 4.4), developed by the US EPA (Byun and Ching, 1999). The CMAQ model is used for regional- and urban-scale air quality simulations, integrating a number of air quality issues (particulate matter, ozone, acid deposition, visibility, etc.) into a so-called “one-atmosphere” approach. The driving meteorological inputs for this work are provided by the fifth-generation NCAR/Penn State Mesoscale Model (MM5), version 3.6.2, and the meteorology-chemistry interface processor (MCIP) was used to transfer MM5 output into gridded meteorological field data as the input to CMAQ. The Carbon Bond – IV chemical mechanism (CB-IV) was used in the CMAQ model, which consists of 36 chemical species, 93 chemical reactions, and 11 photochemical reactions (Lamb, 1982). Geographic Infor-

mation System (GIS) technology is applied in gridding the regional emission inventory to the model domain.

### 1.1 Model domain and simulation episodes

The model domain is based on a Lambert Conformal map projection, using a one-way nested mode with 81 km (covering all China, Japan, Korea, parts of India and Southeast Asia); 27 km (covering eastern China); 9 km (covering major city-clusters including Shandong Province, the Yangtze River Delta (YRD) and the Pearl River Delta (PRD)) and 3 km (covering most of the YRD area, with Shanghai in the center) grid resolutions respectively. The large domain is centered at (118°E, 32°N). The YRD domain has  $99 \times 102$  horizontal grid cells. The model domain is shown in Fig. 1.

The pollution episode is from July 21–30, 2007, when the photochemical reaction is very active and regional air pollution is significant. The initial conditions were prepared by running the model five days ahead of the period starting with a clean initial condition. The model employs 14 vertical layers of varying thickness with denser layers in the lower atmosphere to better resolve the mixing height.

### 1.2 Regional emission inventory

Since one-way nested modeling is used in this study, emission inventories for different simulation domains need to be prepared. In this work, the anthropogenic emissions in Shanghai in 2007 are based on the pollution source survey launched in China in 2007, which was a national scale anthropogenic pollution source investigation, including major air pollution sources like power plants, industry, vehicles and residential sources. The regional emission inventory in the YRD is built using emission factors and local statistics (Jiangsu Statistical Yearbook, 2008;

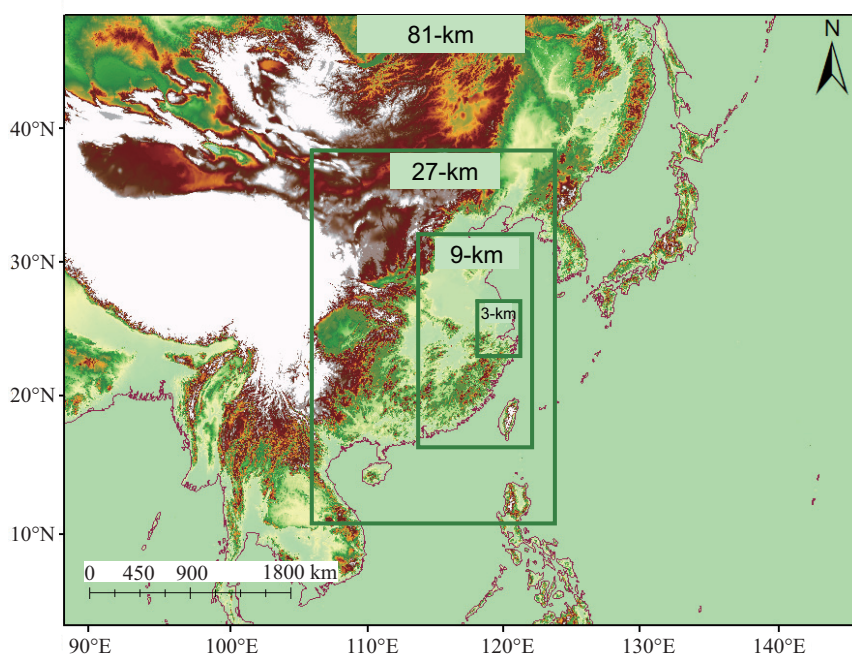


Fig. 1 One-way nested model domain.

Zhejiang Statistical Yearbook, 2008). Supporting data on emission factors and related activity data were assembled from related studies (Streets et al., 2001, 2003a, 2003b; US EPA, 2006; MEP, 1996). The main sources considered in the anthropogenic emission inventory in the YRD in 2007 include industry, transportation, residential and agriculture sectors. The emission sources of the industry sector include the emissions from fuel combustion processes of power plants, boilers, furnaces, kilns, and non-combustion processes such as iron and steel manufacturing, oil refining, cement producing. Transport emission sources mainly consist of vehicle exhaust and road dust emissions. The residential emission sources cover residential fuel combustion emissions, domestic paint and solvent use, gas evaporation in the service stations, etc. The agriculture emission sources include the emissions from livestock feeding, fertilizer application and biomass burning. Huang et al. (2011) has reported detailed information about development of the emission inventory in the Yangtze River Delta.

Emissions of the area outside the YRD are taken from the INTEX-B emission inventory (Zhang et al., 2009). This emission inventory is the latest one that can be found for China. The energy consumption has increased from 2006 to 2007, and thus emissions may also be different between the two years. However, we do not have an updated emission inventory that could be used. The previous version of the INTEX-B emission inventory has been described and demonstrated to be reliable for China in previous studies (Carmichael et al., 2003; Streets et al., 2003a, 2003b).

Table 1 summarizes the emissions used in the modeling for 16 major cities in the YRD. The total anthropogenic VOCs emissions in the YRD in 2007 were around 2767 kilo tons, which mainly comes from non-combustion sources, including oil refining, chemical producing, and fugitive emissions from paint and solvent use in industry, sharing 9%, 30%, and 20% of the total, respectively. In addition, vehicle emission and fugitive emissions from domestic paint and solvent use contribute 15% and 14% of total VOCs emission, respectively. For biogenic VOCs

emissions, this paper used the natural VOCs emission inventory of GEIA Global Emissions Inventory Activity 1990 (<http://geiacenter.org>). In July, total biogenic VOCs emissions in Shanghai were 3286 tons, taking a share of 23% of the total biogenic emissions in the YRD.

### 1.3 Volatile organic compounds (VOCs) speciation

The VOCs speciation is one of the most important factors that influence the ozone simulation results. The VOCs sources include refinery process, chemical process, industrial use of paint and solvent, vehicle exhaust, domestic use of paint and solvent, gas evaporation, biomass burning and biogenic emissions. Some local experimental results of coking industry and vehicle exhaust were adopted to determine the source profiles, and parts of the studies have been published previously (Jia et al., 2009; Lu et al., 2010). The VOCs speciation for other sources without local measurement in the modeling work is mainly compiled based on a literature survey (Liu et al., 2008; Wang et al., 2008; Yuan et al., 2010). The VOCs species are then inserted into the CB-IV mechanism in the CMAQ model.

### 1.4 Model verification

Figure 2 shows comparisons between observed and modeled meteorological parameters including temperature, wind speed, wind direction and relative humidity during the period of July 21–30, 2007. The figure gives average data of six meteorology monitoring sites, including Xujiahui, Nanhui, Qingpu, Baoshan, Jinshan and Minhang. The average bias of temperature, humidity, wind speed and wind direction are 0%, –12%, 15% and –5% respectively. Hourly comparisons of the meteorological parameters show that MM5 can reflect the variation trends of the major meteorological conditions. The selected parameters adopted in MM5 can be used in the pollutant concentration simulation.

Figure 3 shows a comparison between observed and modeled NO<sub>2</sub> concentrations at four national observational sites including Putuo, Xuhui, Luwan and Weifang

**Table 1** Emissions of major anthropogenic species in the YRD in 2007

Province	City	Area (km <sup>2</sup> )	Total anthropogenic emissions (kilo ton)				
			SO <sub>2</sub>	NO <sub>x</sub>	PM <sub>10</sub>	PM <sub>2.5</sub>	VOCs
Jiangsu	Suzhou	1650	174.0	368.6	389.2	208.1	483.3
	Nantong	355	73.0	90.5	111.6	56.8	87.7
	Wuxi	1623	217.0	183.7	347.5	167.8	191.7
	Changzhou	1872	108.0	90.7	211.6	98.8	73.2
	Taizhou	444	20.0	52.5	64.9	31.2	62.8
	Zhenjiang	1082	60.0	103.3	208.3	96.4	45.9
	Yangzhou	980	32.0	110.4	81.8	48.3	58.1
	Nanjing	4723	148.0	144.2	255.9	129.7	153.2
	Zhejiang	16 cities in the YRD	74321	1676.0	2293.0	3115.9	1510.6

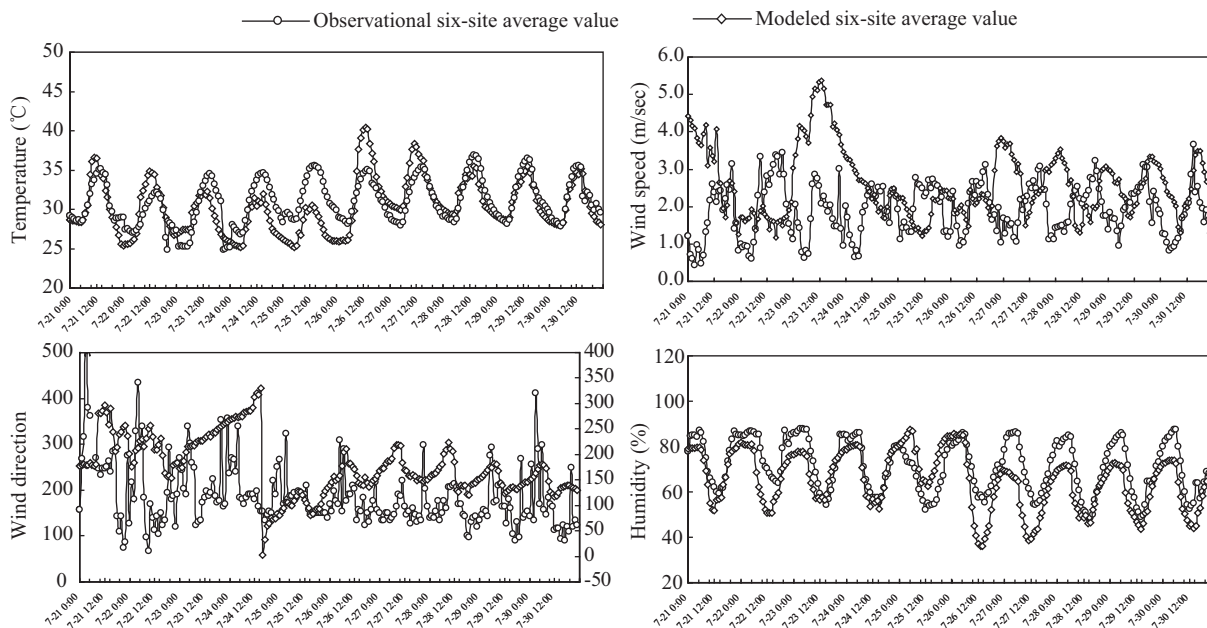


Fig. 2 Comparisons of meteorological parameters between model and observational data during July 21–30, 2007.

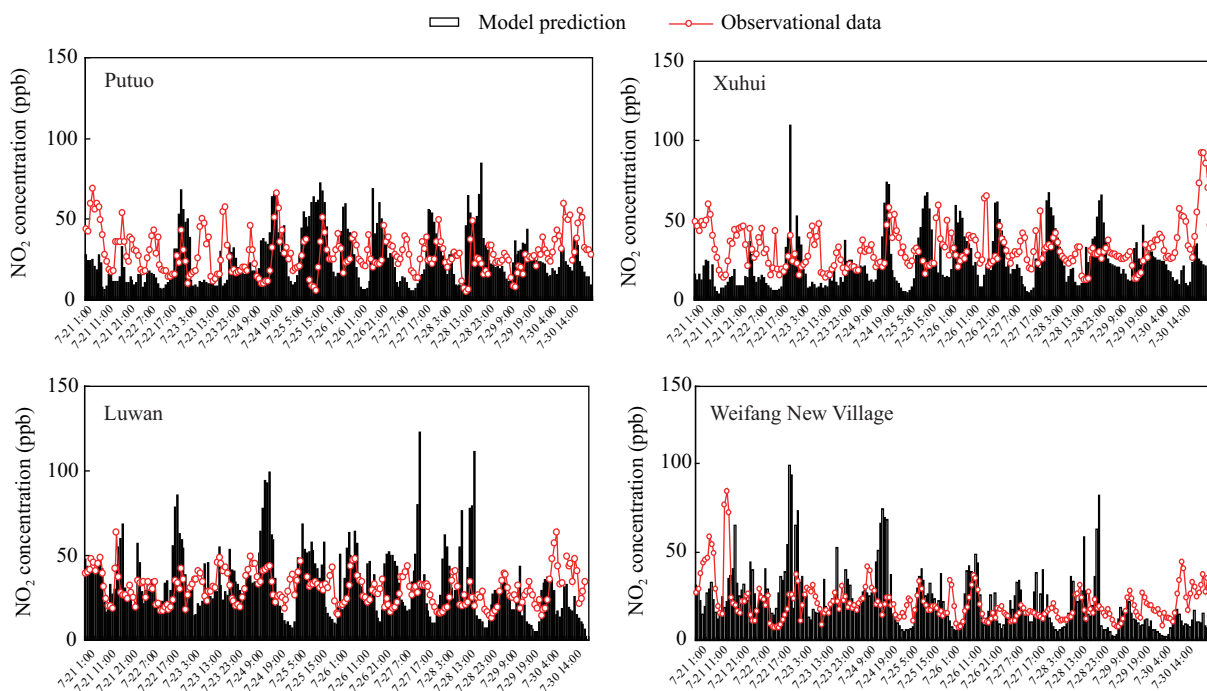


Fig. 3 Comparison of CMAQ model simulations for NO<sub>2</sub> concentrations against observations.

New Village Sites during July 21–30 in 2007. Generally speaking, the modeled data of CMAQ are similar to the observational pollutant trends. The emission inventory of NO<sub>2</sub> in the YRD in 2007, the meteorological field and the modeling results of CMAQ could reflect the NO<sub>2</sub> pollution situation in the YRD.

Figure 4 shows the comparisons between the calculated and observed data of O<sub>3</sub> hourly concentrations at the four national observational sites during July 21–30 in 2007. The results show that the model can well reflect the daily changes of O<sub>3</sub> concentration. With increase of radiation, O<sub>3</sub> concentration rises; while in the afternoon, with decrease of radiation, O<sub>3</sub> concentration gradually

declines. The O<sub>3</sub> concentration gradually decreases and even disappears in the evening. As is shown in the figure, there is still discrepancy between the observed and computed values of ozone concentrations, which might be caused by the out-of-dated biogenic VOCs emissions. As is known, biogenic VOCs contribute significantly to the ozone formation, since the ISOP and TERPB are very active in atmospheric chemistry reactions. However, the biogenic VOCs emissions are based on GEIA, which uses emission data for 1990, possibly quite different from the current situation. The update of biogenic VOCs emissions will be done in a future study. In addition, the calculated O<sub>3</sub> concentrations at midnight are not as low as expected,

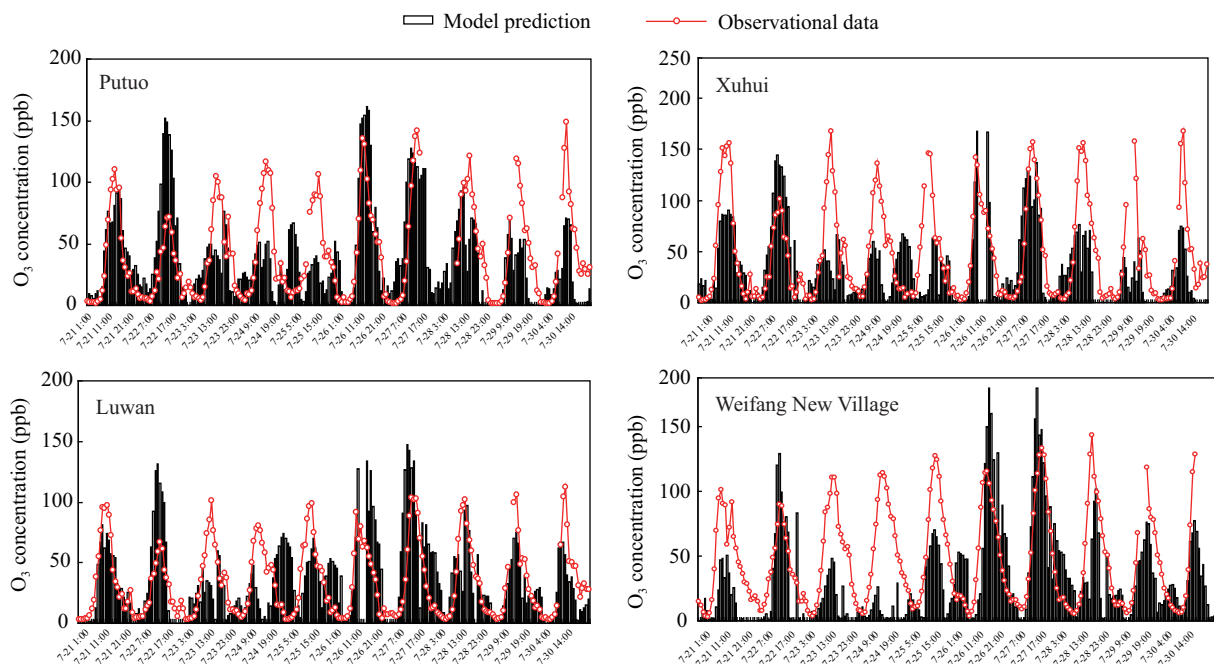


Fig. 4 Comparison of CMAQ model simulations for  $O_3$  concentrations against observations.

which is due to the PBL height simulation by MM5. In this work, the MRF parameters are adopted in MM5 modeling, which gives high PBL height in daytime and low PBL at night. This systematic error causes that the ozone concentration at daytime to be too low while is higher than the observations at night time.

### 1.5 Ozone sensitivity analysis

The photochemical reactions producing ozone are propagated by cycling of  $HO_x$  radicals. The pathway for  $HO_x$  loss is determined by the relative abundance of  $HO_x$  and  $NO_x$ . Ozone production tends to be either  $NO_x$ -sensitive if  $HO_x$ -loss occurs primarily by self reaction of peroxy radicals, or  $NO_x$ -saturated if the primary  $HO_x$ -loss pathway is via reaction of  $NO$  and  $OH$  (Sillman et al., 1990). Three indicators for ozone formation sensitivity, including  $O_3/(NO_y-NO_x)$ ,  $HCHO/NO_y$  and  $H_2O_2/HNO_3$  have been developed (Sillman and Samson, 1995; Sillman and He, 2002). Related ozone sensitivity studies show that the ratios are good indicators for regions that are sensitive to either  $NO_x$  or VOCs emissions changes (Chen, 2002). In addition, the ratios indicate the degree of ozone response to changes in precursor emissions.

To understand the ozone formation mechanism in different areas of Shanghai, this study applied  $O_3/(NO_y-NO_x)$  and  $H_2O_2/HNO_3$  indicators to study the sensitivity of ozone. Regions with a higher ratio have a higher degree of  $NO_x$  emission sensitivity, while regions with a lower ratio have a higher degree of VOCs emission sensitivity. In buffer areas, the  $O_3/(NO_y-NO_x)$  is between 8–10 and  $H_2O_2/HNO_3$  is between 0.35–0.6. To obtain the difference of ozone sensitivity at urban and rural sites in Shanghai, this study chooses Dianshan Lake (DSL) as a representative site of rural areas and Jiang'an (JA) as a representative site of urban areas. Locations of the two representative sites are shown in Fig. 5.



Fig. 5 Locations of the two representative sites in Shanghai.

## 2 Results and discussions

### 2.1 Difference of ozone concentrations at rural and urban sites

The modelling system indicates that the ozone concentration in Shanghai shows great difference between rural and urban areas. Figure 6 shows the modeled hourly ozone concentrations at Dianshan Lake and Jing'an sites during July 21–30, 2007. It is obvious that the ozone concentration at Dianshan Lake is higher than that at Jing'an. The average ozone hourly concentrations at Dianshan Lake and Jing'an during the modelling episode are 65.25 and 35.40 ppb, respectively. The average ozone hourly concentration at Dianshan Lake is 46% higher than that at Jing'an. The average concentration of ozone at 14:00 on each day

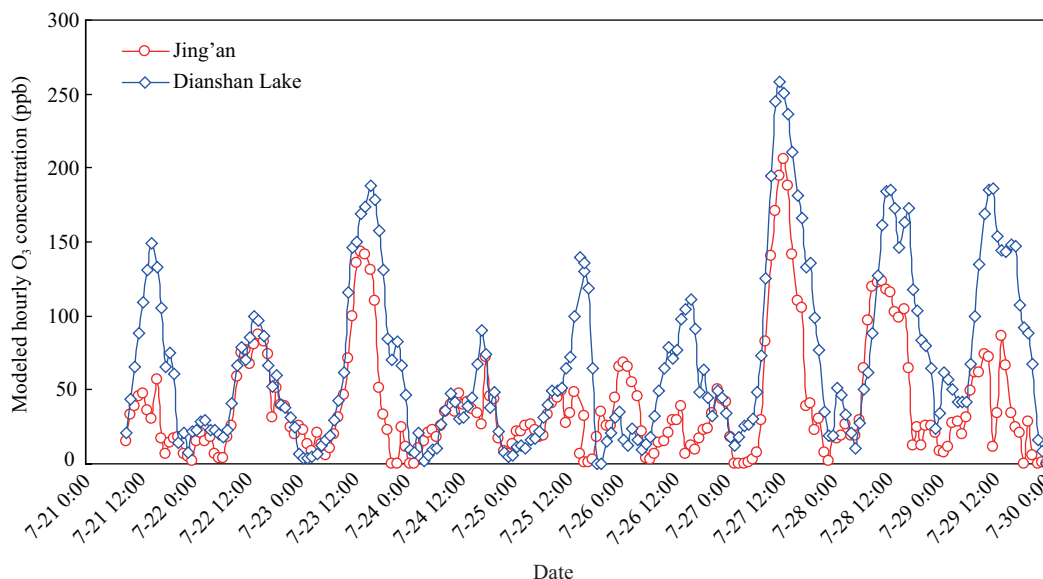


Fig. 6 Modeled hourly ozone concentrations at two sites in Shanghai during Jul 21–30, 2007.

during July 21–30, 2007 at Dianshan Lake is 129.72 ppb, which is 28% higher than the Grade II National Ambient Air Quality Standard. The average concentration of ozone at 14:00 on each day during July 21–30, 2007 at Jing'an is 79.58 ppb, which is 39% lower than that at Dianshan Lake. Results show that the ozone concentrations in rural areas are generally higher than those in urban areas, and the exceedance rate of ozone concentration in rural areas should be paid more attention to. The main reason causing this ozone distribution characteristic is that the ozone is a secondary air pollutant, which is produced through photochemical reactions of NO<sub>x</sub>, VOCs, and CO in the presence of sunlight. The formation of ozone is influenced by the emissions of precursors like NO<sub>x</sub>, VOCs, and the solar radiation intensity. In the urban area of Shanghai, the emissions of NO<sub>x</sub> and VOCs are significant, which produces high ozone concentration in daytime and it is then transported to the rural area. During the night time, the high NO emissions in the urban area gradually titrate ozone, which causes the ozone concentration becoming lower. However, the NO emissions in the rural area are much lower, which preserves the high ozone concentration.

2.2 Ozone sensitivity distribution in Shanghai areas

Figure 7 shows distributions of the average value of O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> at 14:00 during July 21–30, 2007 in Shanghai, modeled by MM5-CMAQ modeling system. As is shown in the figure, the average O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> values during the modeling period within the inner-ring road region is generally lower than 8 and 0.6 respectively, which indicates that in the urban area, the ozone is more sensitive to VOCs. However, in the southeast of Shanghai, the area surrounding Dianshan Lake, and East Chongming District, which represents suburban areas, the average O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> values are higher than 10 and 0.6 respectively, showing that ozone is more sensitive to NO<sub>x</sub> emissions. Thus, to control ozone pollution in Shanghai, different emission control strategies should be made with consideration of the characteristics of different regions.

2.3 Ozone sensitivity at rural and urban sites

The formation mechanism of ozone indicates that the ozone sensitivity in a specific region depends on both the

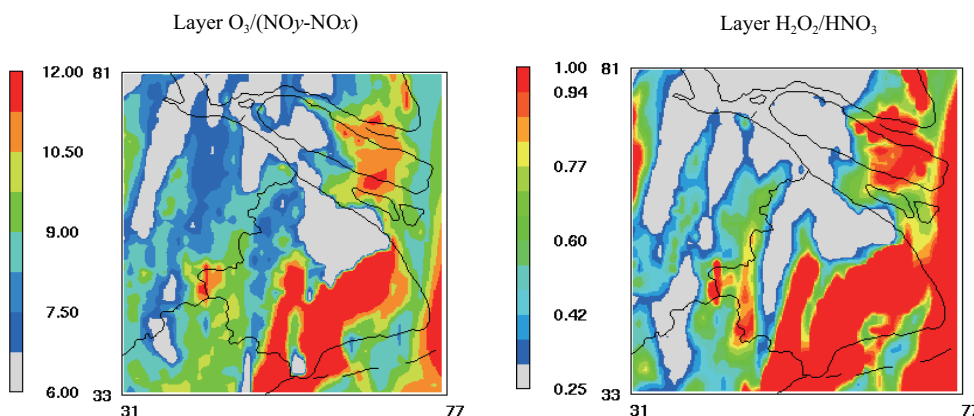


Fig. 7 Modeled distribution of two ozone formation sensitivity indicators in July, 2007.

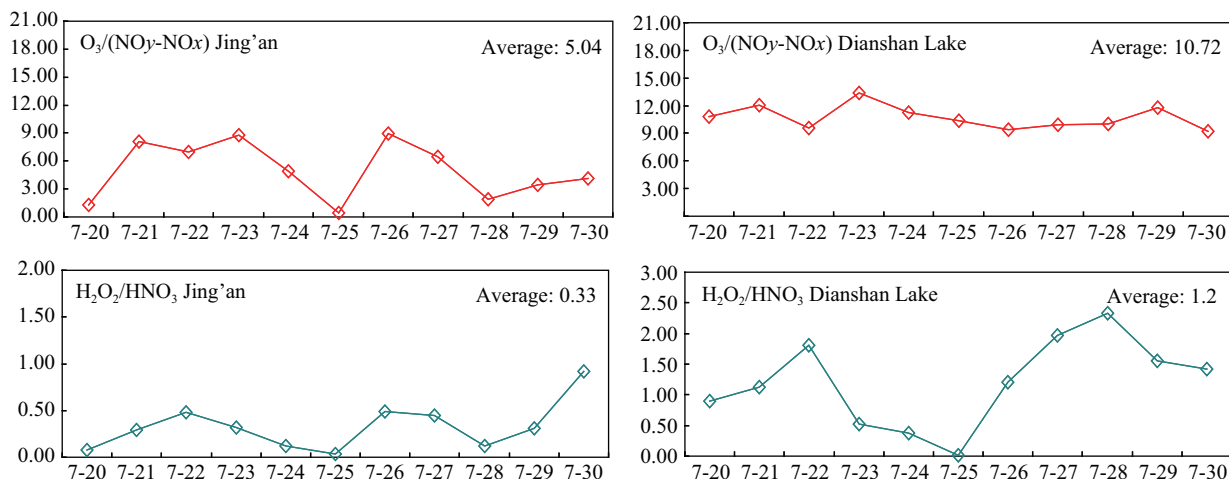


Fig. 8 Modeled indicators at Dianshan Lake and Jing'an sites at 14:00 during July 20–30, 2007.

NO<sub>x</sub>, VOCs concentrations, and the reactivity characterization of each VOC compound. Quantifying the ozone sensitivity with the O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> ratios takes both factors into consideration. High ratio of both the indicators represents that the VOCs catalyst is over abundant, and the control strategy favors NO<sub>x</sub> control. However, low ratio of the indicators shows that the NO<sub>x</sub> is higher than the optimum ozone production level, and the control strategy favors VOCs control.

Both O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> ratios for the two representative locations in Shanghai were calculated during high ozone concentration episodes. Figure 8 gives the O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> indicators at Dianshan Lake and Jing'an sites at 14:00 from July 21–30, 2007. From this figure, it can be seen that the O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) value at Jing'an site varies from 0 to 9, with an average value of 5.04, where O<sub>3</sub> is obviously sensitive to VOCs according to Sillman's measure. On the other hand, most of the H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> values at the Jing'an site are between 0–0.5, with the average value of 0.33, which also shows that the Jing'an site belongs to VOCs control areas. However, the Dianshan Lake site is quite different from Jing'an. The O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) values at Dianshan Lake are between 9–13, with the average of 10.72, where O<sub>3</sub> is significantly sensitive to NO<sub>x</sub>. And most of the H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> values at the Dianshan Lake site are between 0.5–2.5, with the average value of 1.2, which also indicates that the Dianshan Lake site belongs to NO<sub>x</sub>-sensitive areas. From the modeling results, it can be concluded that the urban area in Shanghai belongs to VOCs control areas, while the rural area tends to be NO<sub>x</sub>-sensitive.

The ozone sensitivity analysis with application of O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> ratio in this study compared well with other studies in the literature. This result agrees with the work of Geng et al. (2008), who applied a chemical mechanism model (NCAR-MM) to assess the ozone sensitivity in Shanghai, and found that the ozone formation is clearly under VOC-sensitive regime in the urban city of Shanghai, where the O<sub>3</sub> production is strongly depressed by high NO<sub>x</sub> concentrations, resulting in lower O<sub>3</sub> concentration in center of the city than in rural areas.

### 3 Conclusions

This article applies the MM5-CMAQ modeling system to a regional air pollution study in the Yangtze River Delta with Shanghai in the center. The model performance study shows that the system can well reflect the air pollution situation in this region. Modeling results indicate that the ozone concentration shows an obvious regional difference. The ozone concentration in rural areas (taking Dianshan Lake site as the example) is higher than that in the urban area (taking Jing'an site as the example).

The design of control strategies for surface ozone has been impeded by limited observations of O<sub>3</sub>-NO<sub>x</sub>-VOC sensitivity (Sillman, 1999). This article uses a model-based indicator method to characterized O<sub>3</sub>-NO<sub>x</sub>-VOC sensitivity in Shanghai. Two indicators including O<sub>3</sub>/(NO<sub>y</sub>-NO<sub>x</sub>) and H<sub>2</sub>O<sub>2</sub>/HNO<sub>3</sub> show that ozone is more sensitive to VOCs in urban areas like Jiang'an, and tends to be NO<sub>x</sub> sensitive in rural areas like Dianshan Lake.

Shanghai and the Yangtze River Delta are presently undergoing tremendous economic growth, and the threats of high regional pollutant emissions and high ozone pollution are very real. However, the ozone pollution issue may have different characteristics in different regions. Thus, measures to reduce the ozone concentrations should be carefully made.

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### References

- Byun D W, Ching J K S, 1999. Science algorithms of the EPA Models-3 Community Multiscale Air Quality (CMAQ) modeling system. US Environmental Protection Agency Report EPA/600/R-99/030. Research Triangle Park, NC, USA.
- Carmichael G R, Tang Y, Kurata G, Uno I, Streets D G, Thongboon-

- choo N, 2003. Evaluating regional emission estimates using the TRACE-P observations. *Journal of Geophysical Research*, 108(D21), 8810. DOI: 10.1029/2002JD003116.
- Chen J C M, 2002. Aerosol and ozone sensitivity analysis with the community multi-scale air quality (CMAQ) model for the Pacific Northwest. Master Thesis. Washington State University, USA.
- Geng F H, Tie X X, Xu J M, Zhou G Q, Peng L, Gao W et al., 2008. Characterizations of ozone, NO<sub>x</sub> and VOCs measured in Shanghai, China. *Atmospheric Environment*, 42(29): 6873–6883.
- Geng F H, Zhao C S, Tang X, Lu G L, Tie X X, 2007. Analysis of ozone and VOCs measured in Shanghai: A case study. *Atmospheric Environment*, 41(5): 989–1001.
- Huang C, Chen C H, Li L, Cheng Z, Wang H L, Huang H Y et al., 2011. The study of emission inventory on anthropogenic air pollutants and VOC species in the Yangtze River Delta region, China. *Atmospheric Chemistry and Physics Discussions*, 11: 951–983. DOI: 10.5194/acpd-11-951-2011.
- Jia J H, Huang C, Chen C H, Chen M H, Wang H L, Shao M et al., 2009. Emission characterization and ambient chemical reactivity of volatile organic compounds (VOCs) from coking processes. *Acta Scientiae Circumstantiae*, 29(4): 1–8.
- Jiangsu Statistical Yearbook, 2008. China Statistics Press, Beijing, China.
- Lamb R G, 1982. A regional-scale (1000 km) model of photochemical air pollution: Part I. Theoretical formulation, U.S. Environmental Protection Agency Report EPA/600/3-85-035, Research Triangle Park, NC.
- Liu Y, Shao M, Fu L L, Lu S H, Zeng L M, Tang D G, 2008. Source profiles of volatile organic compounds (VOCs) measured in China: Part I. *Atmospheric Environment*, 42(25): 6247–6260.
- Lu J, Wang H L, Chen C H, Huang C, Li L, Cheng Z et al., 2010. The composition and chemical reactivity of volatile organic compounds (VOCs) from vehicle exhaust in Shanghai, China. *Environmental Pollution & Control*, 32(6): 19–26.
- MEP (Ministry of Environmental Protection) of the People's Republic of China, 1996. User's Guide on the Production and Emission Factors of Industrial Pollutants. China Environmental Science Press, Beijing, China.
- Ran L, Zhao C S, Xu W Y, Lu X Q, Han M, Lin W L et al., 2011. VOC reactivity and its effect on ozone production during the HaChi summer campaign. *Atmospheric Chemistry and Physics Discussions*, 11: 8595–8623. DOI: 10.5194/acpd-11-8595-2011.
- Seinfeld J H, Pandis S N, 1998. *Atmospheric Chemistry and Physics from Air Pollution to Climate Change*. John Wiley & Sons, New York.
- Shan W P, Yin Y Q, Zhang J D, Ding Y P, 2008. Observational study of surface ozone at an urban site in East China. *Atmospheric Research*, 89(3): 252–261.
- Shanghai Statistical Yearbook, 2001. China Statistics Press. Beijing, China.
- Shanghai Statistical Yearbook, 2010. China Statistics Press. Beijing, China.
- Sillman S, 1999. The relation between ozone, NO<sub>x</sub>, and hydrocarbons in urban and polluted rural environments. *Atmospheric Environment*, 33(12): 1821–1845.
- Sillman S, He D, 2002. Some theoretical results concerning O<sub>3</sub>-NO<sub>x</sub>-VOC chemistry and NO<sub>x</sub>-VOC indicators. *Journal of Geophysical Research*, 107(D22): 4659. DOI: 10.1029/2001JD001123.
- Sillman S, Logan J A, Wofsy S C, 1990. The sensitivity of ozone to nitrogen oxides and hydrocarbons in regional ozone episodes. *Journal of Geophysical Research*, 95(D2): 1837–1852.
- Sillman S, Samson P J, 1995. Impact of temperature on oxidant photochemistry in urban, polluted rural and remote environments. *Journal of Geophysical Research*, 100(D6): 11497–11508.
- Streets D G, Bond T C, Carmichael G R, Fernandes S D, Fu Q Y, He D et al., 2003a. An inventory of gaseous and primary aerosol emissions in Asia in the year 2000. *Journal of Geophysical Research*, 108(D21): 8809. DOI: 10.1029/2002JD003093.
- Streets D G, Fu J S, Jang C J, Hao J M, He K B, Tang X Y et al., 2007. Air quality during the 2008 Beijing Olympic Games. *Atmospheric Environment*, 41(3): 480–492.
- Streets D G, Jiang K, Hu X, Sinton J E, Zhang X Q, Xu D et al., 2001. Recent reductions in China's greenhouse-gas emissions. *Science*, 294(5548): 1835–1836.
- Streets D G, Yarber K F, Woo J H, Carmichael G R, 2003b. Biomass burning in Asia: annual and seasonal estimates and atmospheric emissions. *Global Biogeochemical Cycles*, 17: 1099. DOI: 10.1029/2003GB002040.
- Tang W Y, Zhao C S, Geng F H, Peng L, Zhou G Q, Gao W et al., 2008. Study of ozone "weekend effect" in Shanghai. *Science in China Series D: Earth Sciences*, 51(9): 1354–1360.
- Tang X Y, 2004. The characteristics of urban air pollution in China. In: *Urbanization, Energy, and Air Pollution in China*. The National Academies Press, Washington, DC. 47–54.
- Tie X X, Brasseur G P, Zhao C S, Granier C, Massie S, Qin Y et al., 2006. Chemical characterization of air pollution in Eastern China and the Eastern United States. *Atmospheric Environment*, 40(14): 2607–2625.
- U.S. Environmental Protection Agency, 2006. AP-42 Emission Factors, Washington, DC.
- Wang B G, Zhang Y H, Shao M, Zhou Y, Feng Z C, 2008. Sources apportionment of anthropogenic C<sub>2</sub>-C<sub>9</sub> non-methane hydrocarbons in the atmosphere of Guangzhou, China. *Acta Scientiae Circumstantiae*, 28(7): 1430–1440.
- Wang H X, Kiang C S, Tang X Y, Zhou X J, Chameides W L, 2005. Surface ozone: A likely threat to crops in Yangtze delta of China. *Atmospheric Environment*, 39(21): 3843–3850.
- Wang H X, Tang X Y, Wang M L, Yan P, Wang T, Shao K S et al., 2003. The temporal and spatial allocation characteristics of trace gases in the Yangtze River Delta. *Science in China (Series D)*, 33(2): 114–118.
- Wang T, Cheung V T F, Anson M, Li Y S, 2001. Ozone and related gaseous pollutants in the boundary layer of eastern China: overview of the recent measurements at a rural site *Geophysical Research Letters*, 28(12): 2373–2376.
- Wang X S, Li J L, 2002. The contribution of anthropogenic hydrocarbons to ozone formation in Beijing areas. *China Environmental Science*, 22(6): 501–505.
- Xu J, Zhang Y H, 2006. Process analysis of O<sub>3</sub> formation in summer at Beijing. *Acta Scientiae Circumstantiae*, 26(6): 973–980.
- Xu J, Zhang Y H, Wang W, 2006. Numerical study on the impacts of heterogeneous reactions on ozone formation in the Beijing urban area. *Advance in Atmospheric Sciences*, 23(4): 605–614.
- Yuan B, Shao M, Lu S H, Wang B, 2010. Source profiles of volatile organic compounds associated with solvent use in Beijing, China. *Atmospheric Environment*, 44(15): 1919–1926.
- Zhang Q, Streets D G, Carmichael G R, He K B, Huo H, Kannari A et al., 2009. Asian emissions in 2006 for the NASA INTEX-B emission. *Atmospheric Chemistry and Physics*, 9: 5131–5153. DOI: 10.5194/acp-9-5131-2009.
- Zhao C S, Peng L, Sun A D, Qin Y, Liu H L, Li W L et al., 2004. Numerical modeling of tropospheric ozone over Yangtze Delta region. *Acta Scientiae Circumstantiae*, 24(3): 525–533.
- Zhejiang Statistical Yearbook, 2008. China Statistics Press, Beijing, China.