

1 **Impacts of emission reduction and meteorological conditions on air**  
2 **quality improvement during the 2014 Youth Olympic Games in**  
3 **Nanjing, China**

4

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18

19 **Abstract**

20 As the holding city of the 2nd Youth Olympic Games (YOG), Nanjing is highly  
21 industrialized and urbanized facing with several air pollution issues. In order to ensure  
22 better air quality during the event, the local government took great efforts to control  
23 the emissions from pollutant sources. However, air quality can still be affected by  
24 synoptic weather, not only emission. In this paper, the influences of meteorological  
25 factors and emission reductions were investigated using observational data and  
26 numerical simulations with WRF/CMAQ. During the YOG holding month (Aug.,  
27 2014), the observed hourly mean concentrations of SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and  
28 O<sub>3</sub> were 11.6 μg/m<sup>3</sup>, 34.0 μg/m<sup>3</sup>, 57.8 μg/m<sup>3</sup>, 39.4 μg/m<sup>3</sup>, 0.9 mg/m<sup>3</sup>, and 38.8 μg/m<sup>3</sup>,  
29 respectively, which were below China National Ambient Air Quality Standard (Level

30 2). However, model simulation showed that the weather conditions such as weaker  
31 winds during the holding time were adverse for better air quality, and could increase  
32 SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub> and CO by 17.5%, 16.9%, 18.5%, 18.8%, 7.8% and 0.8%,  
33 respectively. Taking account of local emission abatement only, the simulated SO<sub>2</sub>,  
34 NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub> and CO was decreased by 24.6%, 12.1%, 15.1%, 8.1% and 7.2%,  
35 respectively. Consequently, stringent emission control measures can reduce the  
36 concentrations of air pollutants in short term, and emission reduction is the very  
37 important factor for the air quality improvement during the YOG, which has set up a  
38 good example in air protection for important social events.

39 **KEY WORDS:** Youth Olympic Games; Emission reduction; Meteorological  
40 conditions; WRF/CMAQ; Nanjing

41

## 42 **1 Introduction**

43 As located in the economically developed Yangtze River Delta (YRD) region of  
44 China, Nanjing successfully hosted the second Youth Olympic Games (YOG) during  
45 16 - 28 Aug., 2014. Nanjing is a highly urbanized city and its GDP ranked the 12<sup>th</sup> of  
46 all the cities in China by 2013 (National Bureau of Statistics of China, 2014). Due to  
47 fast urbanization and industrialization, heavy motor vehicles and construction dust,  
48 Nanjing has long been suffered from air pollution (Dong et al., 2013; Chen et al.,  
49 2015).

50 In order to realize the promise of “Green YOG”, the local government had taken  
51 a series of measures to reduce emissions of air pollutants. The preparatory work  
52 started from 1 Jul., 2014. Besides, the local government performed the work plan  
53 for stringent environmental quality assurance from 1 Aug. (National Bureau of  
54 Statistics of China, 2014). The controlled emissions include 5 major categories:  
55 industry, power plants, traffic, VOC product-related sources and others. Some local  
56 petrochemical, chemical and steel industries were forced to limit or halt the  
57 production. Coal-combustion enterprises were required to use high-quality coals with  
58 low sulfur content and ash content. And vehicles with heavy pollution called “yellow  
59 label buses” were prohibited in Nanjing during 10-28 Aug.. Oil loading and unloading

60 operations were strictly controlled. All construction processes in the city were forced  
61 to stop. The ground surface with bare soil was covered.

62 It is well known that air quality can be affected by both meteorological factors  
63 and pollutant emissions. Many cases verified that both emission abatement and  
64 weather conditions can influence the improvement of local air quality. Emission  
65 control has been taken in many social events, like Beijing Olympic Games in 2008  
66 and Shanghai Expo in 2010. Xing et al. (2011) suggested that emission controls  
67 benefit for pollutants reduction, but meteorological effects can be either ways at  
68 different locations. Cermak and Knutti (2009), Wang et al. (2009b, 2010) and Xing et  
69 al. (2011) reported that typical meteorological conditions accounted more for air  
70 improvement during 2008 Beijing Olympics than emission reductions. Zhou et al.  
71 (2010) concluded that control measures for transportation resulted in a reduction of  
72 44.5% and 49.0% in daily CO and NO<sub>x</sub> emission from motor vehicles during the 2008  
73 Olympics. Cai et al. (2011) and Wang et al. (2009a) also studied the transportation  
74 controls on improving air quality during Beijing Olympic Games. Okuda et al. (2011)  
75 argued that sources control during Beijing Olympics significantly reduced PM<sub>10</sub>, NO<sub>2</sub>  
76 and SO<sub>2</sub>, but did not effectively reduce PM<sub>2.5</sub>. Streets et al. (2007) proposed that local  
77 sources controlling is inadequate for heavily populated, urbanized, and industrialized  
78 city, regional air quality management is in urgent need. Lin et al. (2013) applied  
79 monitoring data to analyze the weather impacts on air quality of the World Expo in  
80 YRD and concluded that high frequency of marine winds during the Expo had a  
81 positive effect on the air quality of coastal cities, but a negative effect on some inland  
82 cities in YRD. Satellite data reflected that the tropospheric NO<sub>2</sub> column, aerosol  
83 optical thickness (AOT), and CO concentration dropped by 8%, 14% and 12%,  
84 respectively over Shanghai during the Expo period, compared to the past three years  
85 (Hao et al., 2011). Liu et al. (2013) compared the contributions of long-term and  
86 short-term emission control via CMAQ simulation and investigated their effects on air  
87 quality in Guangzhou during the Asian Games. Xu et al. (2013) concluded that PM<sub>2.5</sub>  
88 was mainly emitted from anthropogenic sources other than biogenic sources and  
89 indicated that cutting down anthropogenic emissions could decrease PM<sub>2.5</sub> effectively.

90 Dong et al. (2013) found that independent NO<sub>x</sub> emission reduction would strengthen  
91 O<sub>3</sub> as a side effect in YRD. Chen et al. (2015, 2017) studied the source apportionment  
92 of size-fractionated particles in Nanjing, and found that fugitive and construction dust  
93 decreased significantly in YOG.

94 There have been some studies on air quality during the 2nd YOG (Ding et al.,  
95 2015; Chen et al., 2017; Zhou et al., 2017), but few work focused on the relative  
96 contributions of meteorology and control efforts. This study takes the air quality  
97 monitoring data and applies WRF/CMAQ model to estimate the effect of  
98 meteorological factors and emission reduction on air quality of Nanjing during the  
99 2nd YOG. Data and model descriptions as well as simulation scenarios are described  
100 in Section 2. Section 3 examines the characteristics of six major air pollutants (SO<sub>2</sub>,  
101 NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub>) and compares their concentrations during the YOG  
102 with those a year ago (Aug., 2013) and the months without emission reduction (Jul.  
103 and Sept., 2014). Besides, this section discusses the separate effect of weather  
104 conditions and emission abatement qualitatively and quantitatively based on the  
105 simulation results. Section 4 summaries the main conclusions, emphasizes the  
106 important factor of the air quality promotion during the YOG, and provides some  
107 advice for ensuring pleasant future air quality.

108

## 109 **2 Methodology**

### 110 2.1 Data description

111 Hourly observed air quality data during Jul.-Sept. 2014 and Aug. 2013 of two  
112 representative stations was collected from Nanjing Environmental Monitoring Center  
113 (<http://222.190.111.117:8023/>). Both of the two stations are state controlling air  
114 monitoring sites. The data quality assurance and quality control procedures for  
115 monitoring strictly follow the national standards (State Environmental Protection  
116 Administration of China, 2006). Caochangmen (CCM) Station (118.75° E, 32.06° N)  
117 locates in Gulou District, the city center of Nanjing. Gulou District is the center of  
118 economy, politics, culture and education in Nanjing, where gathers many East China's  
119 high-end industrial and corporate headquarters. Besides, over 90% provincial

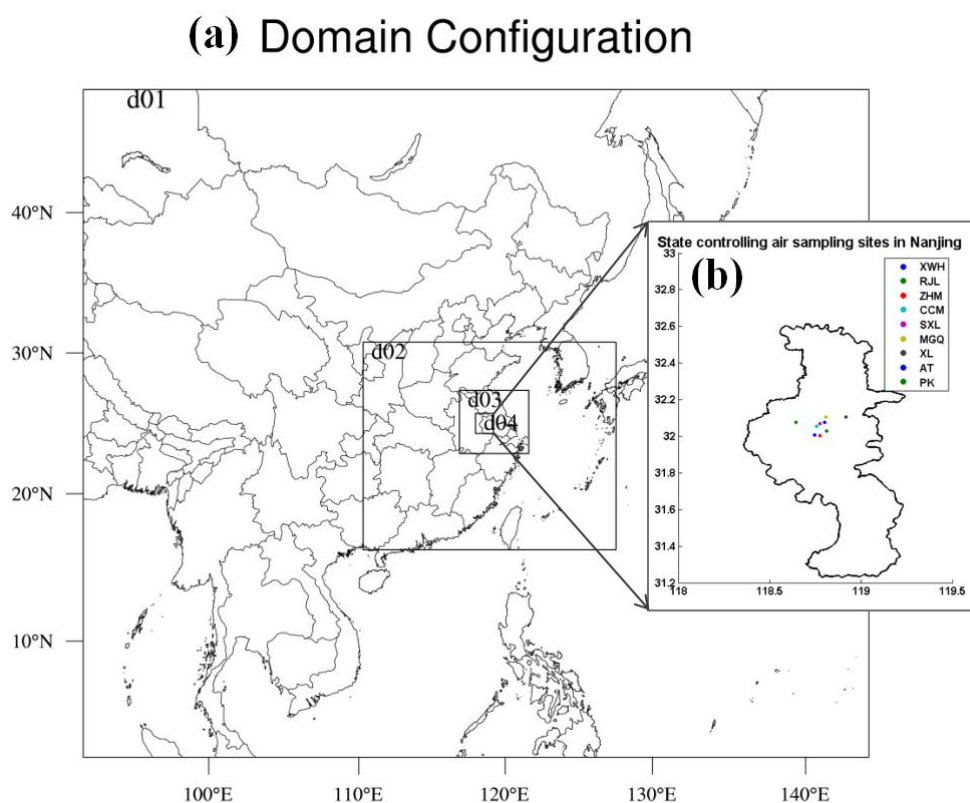
120 authorities, more than 20 colleges and universities, and more than 120 institutes  
121 situate in Gulou District. It's the most populated area in Nanjing, with lively  
122 commercial hub and heavy traffic. Thus, CCM station was chosen to represent the  
123 urban status of Nanjing. The other site calls Xianlin (XL) Station ( $118.92^{\circ}$  E,  $32.11^{\circ}$   
124 N ), which locates in Qixia District, the suburb of Nanjing. Compared to Gulou  
125 District, Qixia District is much more sparsely populated with few traffic congestion  
126 problems. Thus, XL station was chosen to represent the suburban status of Nanjing.

127

## 128 2.2 Model description

129 The integrated modeling system WRF/CMAQ was employed in this study.  
130 Community Multiscale Air Quality (CMAQ) is a third-generation regional air quality  
131 model developed by the Environmental Protection Agency of USA (USEPA). It  
132 incorporates a set of up-to-date compatible modules and control equations for the  
133 atmosphere, and can fully consider complicated physical and chemical processes  
134 (Byun and Schere, 2006; Foley et al., 2010). Many applications have proven that  
135 CMAQ is a reliable tool in simulating air quality from city scale to mesoscale (Xing  
136 et al., 2011; Dong et al., 2013; Liu et al., 2013; Xu et al. 2013; Shu et al., 2016).  
137 Community Multiscale Air Quality (CMAQ v4.7.1, Binkowski and Roselle, 2003)  
138 model includes the 2005 Carbon Bond gas-phase mechanism (CB05) (Yarwood et al.,  
139 2005) and the fourth-generation CMAQ aerosol module (AERO4) (Byun and Schere,  
140 2006). And it was applied to simulate the pollutant distribution over Nanjing in this  
141 paper. Weather Research and Forecasting (WRF) is a new generation of mesoscale  
142 weather forecast model and assimilation system, developed by the National Center for  
143 Atmospheric Research (NCAR). It has been widely applied in China and shows a  
144 good performance in all kinds of weather forecasts (Jiang et al., 2008, 2012; Xu et  
145 al.,2013; Liao et al., 2014, 2015; Xie et al., 2014, 2016; Li et al., 2016; Shu et al.,  
146 2016). WRF v3.2.1 (Skamarock et al., 2008) model was run to provide meteorological  
147 fields for CMAQ. Four nested domains were set for both models, with horizontal  
148 resolutions of 81km, 27km, 9km, 3km, and the innermost domain covering Nanjing  
149 (Fig. 1). For all domains, 23 vertical sigma layers from the surface to the top pressure

150 of 100 hpa was set, with about 10 layers in the planetary boundary layer. The detail  
 151 dynamic parameterization in WRF as well as the physical and chemical schemes of  
 152 CMAQ applied in this research were the same as those in Shu et al. (2016)'s work and  
 153 were proven to have good performance. As for the innermost domain, Nanjing  
 154 Environmental Protection Bureau chooses the local 9 state controlling air monitoring  
 155 sites (See Fig. 1, Table 1) to represent the whole Nanjing (NJ) city. In conformity with  
 156 this, the 9 sites in domain 4 were chosen to represent the whole Nanjing when  
 157 analyzing the impacts of weather conditions and emission reduction.  
 158



159  
 160 **Fig. 1.** Modeling domains and state controlling air monitoring sites in Nanjing. ((a) The four nested  
 161 modeling domains at 81km (d01: East Asia), 27km (d02: East China), 9km (d03: Yangtze River Delta),  
 162 and 3km (d04: Nanjing), (b) Locations of 9 sites in Nanjing).

163

164 **Table 1**

165 The air monitoring sites in Nanjing

Site	Abbreviation	Location
Xuanwuhu Station	XWH	32.08° N, 118.80° E
Ruijinlu Station	RJL	32.03° N, 118.82° E
Zhonghuamen Station	ZHM	32.00° N, 118.76° E

Caochangmen Station	CCM	32.06° N, 118.75° E
Shanxilu Station	SXL	32.07° N, 118.77° E
Maigaoqiao Station	MGQ	32.11° N, 118.81° E
Xianlin Station	XL	32.11° N, 118.92° E
Aoti Station	AT	32.01° N, 118.74° E
Pukou Station	PK	32.07° N, 118.64° E

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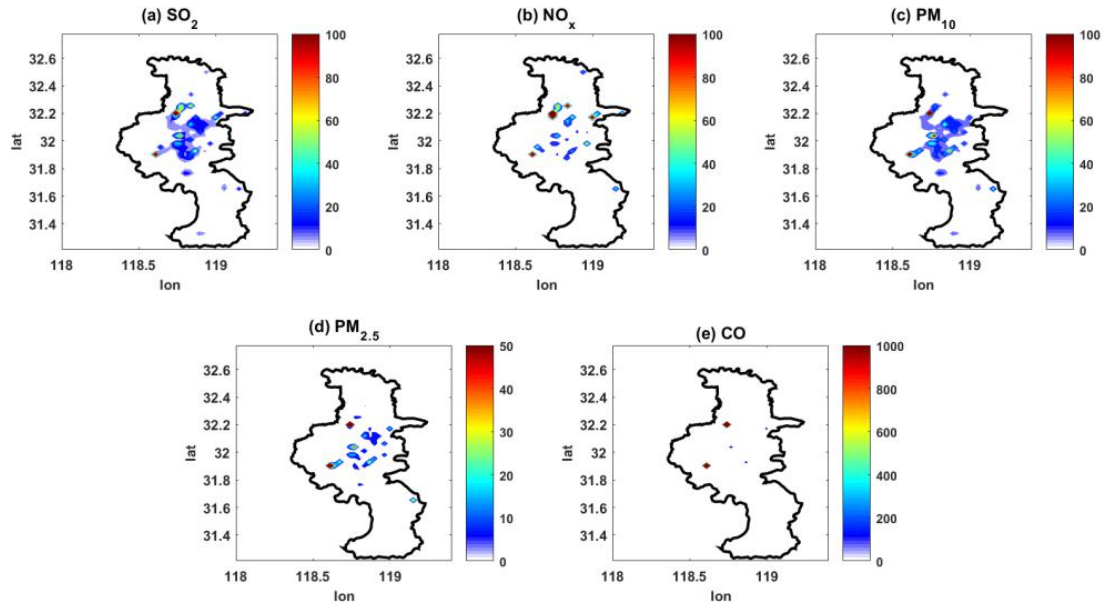
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### 167 2.3 Emissions and simulation scenarios

168 In this study, Multi-resolution Emission Inventory for China (MEIC v1.2,  
169 <http://www.meicmodel.org/>) with a resolution of  $0.25^\circ \times 0.25^\circ$  was employed to  
170 provide the anthropogenic emissions for species including SO<sub>2</sub>, NO<sub>x</sub>, CO, NMVOC,  
171 NH<sub>3</sub>, CO<sub>2</sub>, PM<sub>2.5</sub>, PM<sub>10</sub>, BC, and OC, which form 4 sectors as industry, power plants,  
172 transportation, and residential.

173 For the innermost domain, the local emission inventory before and after emission  
174 control was used with a horizontal resolution of 3km  $\times$  3km. The base year of the  
175 local emission is 2012. According to the local emission control program, 5 major  
176 categories: industry, power plants, traffic, VOC product-related sources and others  
177 were in the control list. In Aug. 2014, all coal-combustion enterprises must use  
178 high-quality coals with low sulfur content less than 0.5% and ash content less than  
179 13%. Besides, the local government ordered over 100 local petrochemical, chemical  
180 and steel enterprises to cut or halt their production. Moreover, heavy pollution  
181 vehicles were prohibited in Nanjing during 10-28 Aug. 2014 to reduce traffic  
182 emission. To reduce emissions of volatile organic compounds, loading and unloading  
183 oil operations were prohibited at the docks in Nanjing section of Yangtze River.  
184 What's more, local construction work was halted during Aug. 2014. With these efforts,  
185 the emission would be cut by 25.0% for SO<sub>2</sub>, 15.0% for NO<sub>x</sub>, 42.8% for PM<sub>10</sub>, 36.2%  
186 for PM<sub>2.5</sub>, and 20.0% for CO. The spatial distributions of emission reduction for  
187 different species were showed in Fig. 2. For SO<sub>2</sub>, NO<sub>x</sub>, PM<sub>10</sub> and PM<sub>2.5</sub>, they centered  
188 in the middle of Nanjing city and for CO, it centered in several points.

189



190  
191 **Fig. 2.** Emission reduction in domain 4 ((a) SO<sub>2</sub>, (b) NO<sub>x</sub>, (c) PM<sub>10</sub>, (d) PM<sub>2.5</sub>, (e) CO (unit: t/month)).

192  
193 The simulated period was from 27 Jul. to 1 Sept. (China standard time, CST), but  
194 only the holding month (1 Aug. to 31 Aug.) was focused on for discussion. In order to  
195 better understand the influence of meteorological factors and emission abatement,  
196 three experiments were carried out. Exp. 1 used the weather conditions during Aug.  
197 2014 (CST) and the emission inventory after reduction. Exp. 2 used the same weather  
198 conditions as Exp. 1 with the emission inventory before reduction. Exp. 3 used the  
199 same inventory as Exp. 2 and the weather conditions during Aug. 2013 (CST).  
200 Besides, Exp. 2 acted as the control experiment. Therefore, Exp. 1 and Exp. 2 were  
201 performed to investigate the influence of emission reduction on pollutant levels.  
202 Similarly, Exp. 2 and Exp. 3 were conducted to understand the impact of meteorology  
203 on air quality.

### 204 205 **3 Results and discussions**

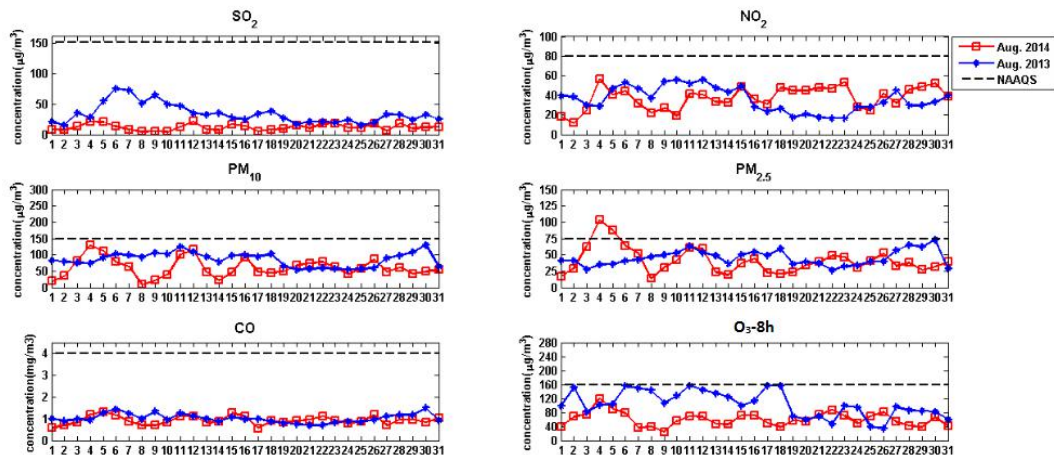
#### 206 **3.1 Observed air quality during the YOG**

207 In Aug. 2014, emission sources including 5 major categories were strictly  
208 reduced, and the air quality had great promotion compared to Aug. 2013. Firstly, air  
209 quality was good during the Games in accordance with China's National Ambient Air  
210 Quality Standards (NAAQS) (Ministry of Environmental Protection of the People's



211 Republic of China, 2012) (Fig. 3, Fig. 4). The hourly mean concentrations of  
 212 pollutants at the two sites during Aug. 2014 are 11.6  $\mu\text{g}/\text{m}^3$  for  $\text{SO}_2$ , 34.0  $\mu\text{g}/\text{m}^3$  for  
 213  $\text{NO}_2$ , 57.8  $\mu\text{g}/\text{m}^3$  for  $\text{PM}_{10}$ , 39.4  $\mu\text{g}/\text{m}^3$  for  $\text{PM}_{2.5}$ , 0.9  $\text{mg}/\text{m}^3$  for CO, and 38.8  $\mu\text{g}/\text{m}^3$   
 214 for  $\text{O}_3$ . Secondly, as showed in Table 2 and Table 3, the mean concentrations of the six  
 215 major species dropped by 64.7% for  $\text{SO}_2$ , 29.8% for  $\text{PM}_{10}$ , 9.8% for  $\text{PM}_{2.5}$ , 8.9% for  
 216 CO and 31.7% for  $\text{O}_3$  at CCM station, while 50.0% for  $\text{SO}_2$ , 18.6% for  $\text{NO}_2$ , 32.8%  
 217 for  $\text{PM}_{10}$ , 4.1% for  $\text{PM}_{2.5}$ , and 31.7% for  $\text{O}_3$  at XL station. Besides, the smaller  
 218 standard deviation (std) of  $\text{SO}_2$ ,  $\text{NO}_2$ , CO and  $\text{O}_3$  revealed that concentrations of these  
 219 air pollutants varied more steadily in Aug. 2014. Actually, the drop of pollutant  
 220 concentrations could be caused mainly by meteorological conditions or emission  
 221 reduction, which will be discussed based on model simulations in Section 3.2 and  
 222 Section 3.3.

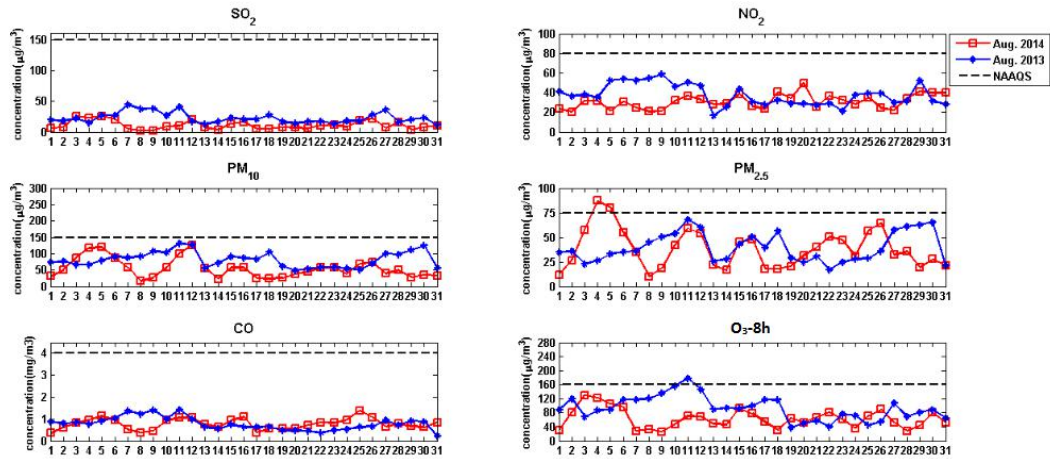
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225 **Fig. 3.** Day-to-day variations in  $\text{SO}_2$ ,  $\text{NO}_2$ ,  $\text{PM}_{10}$ ,  $\text{PM}_{2.5}$ , CO and  $\text{O}_3$ -8h at CCM station in Aug. 2013  
 226 and Aug. 2014 (CST). Observed data in Aug. 2013 and Aug. 2014 are indicated in blue and red,  
 227 respectively. NAAQS are indicated in black dotted line.

228



229  
 230 **Fig. 4.** Day-to-day variations in SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub>-8h at XL station in Aug. 2013 and  
 231 Aug. 2014 (CST). Observed data in Aug. 2013 and Aug. 2014 are indicated in blue and red,  
 232 respectively. NAAQS are indicated in black dotted line.

233

234 **Table 2**

235 Statistical analysis of hourly data in Aug. 2013 and Aug. 2014 at CCM station (The unit is µg/m<sup>3</sup>  
 236 except CO (mg/m<sup>3</sup>))

Species	Time	Max	Min	Mean	Median	Std	Δ
SO <sub>2</sub>	Aug. 2013	169.0	1.0	33.7	27.0	23.7	
	Aug. 2014	72.0	2.0	11.9	10.0	7.8	-64.7%
NO <sub>2</sub>	Aug. 2013	111.0	1.0	35.4	32.0	19.4	
	Aug. 2014	110.0	1.0	37.3	35.0	18.6	5.0%
PM <sub>10</sub>	Aug. 2013	213.0	19.0	86.0	84.0	29.5	
	Aug. 2014	198.0	6.0	60.4	54.0	36.6	-29.8%
PM <sub>2.5</sub>	Aug. 2013	123.0	10.0	45.2	43.5	16.2	
	Aug. 2014	165.0	3.0	40.7	36.0	23.8	-9.8%
CO	Aug. 2013	3.1	0.4	1.0	0.9	0.4	
	Aug. 2014	2.2	0.3	0.9	0.9	0.3	-8.9%
O <sub>3</sub>	Aug. 2013	198.0	1.0	56.9	42.0	46.2	
	Aug. 2014	150.0	9.0	38.9	34.0	22.6	-31.7%

237 Δ : The change percentage of species in Aug. 2014 based on Aug. 2013.

238

239 **Table 3**

240 Statistical analysis of hourly data in Aug. 2013 and Aug. 2014 at XL station (The unit is µg/m<sup>3</sup> except  
 241 CO (mg/m<sup>3</sup>))

Species	Time	Max	Min	Mean	Median	Std	Δ
SO <sub>2</sub>	Aug. 2013	139.0	0.0	22.8	19.0	16.1	
	Aug. 2014	71.0	1.0	11.4	8.0	10.4	-50.0%
NO <sub>2</sub>	Aug. 2013	129.0	0.0	37.7	32.0	21.7	

	Aug. 2014	95.0	7.0	30.7	27.0	15.0	-18.6%
PM <sub>10</sub>	Aug. 2013	215.0	0.0	82.1	79.0	32.4	
	Aug. 2014	196.0	6.0	55.2	47.0	35.9	-32.8%
PM <sub>2.5</sub>	Aug. 2013	122.0	0.0	39.7	37.5	18.9	
	Aug. 2014	157.0	3.0	38.0	34.0	24.1	-4.1%
CO	Aug. 2013	3.2	0.0	0.8	0.7	0.4	
	Aug. 2014	2.0	0.3	0.8	0.7	0.3	<0.1%
O <sub>3</sub>	Aug. 2013	193.0	0.0	56.6	44.0	37.5	
	Aug. 2014	148.0	2.0	38.7	32.0	28.3	-31.7%

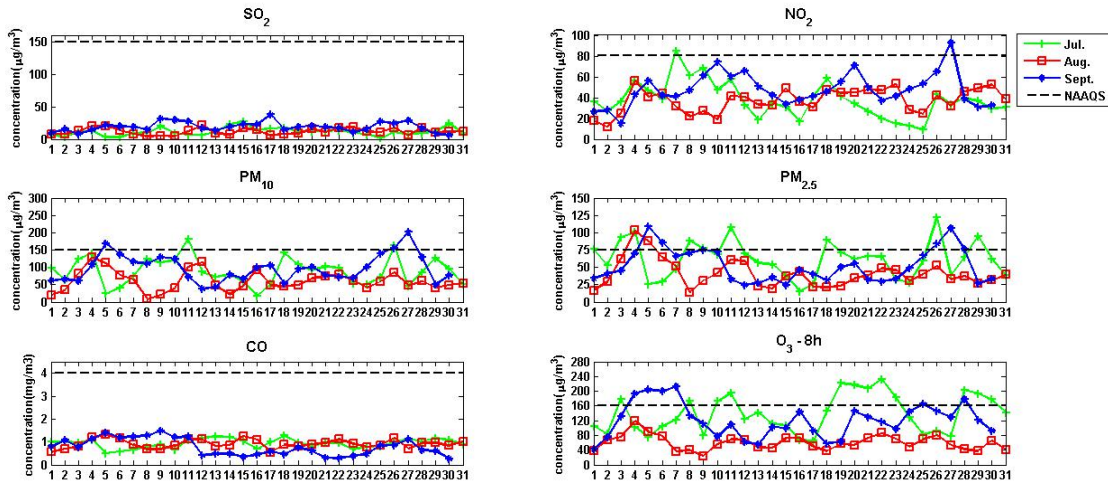
242  $\Delta$  : The change percentage of species in Aug. 2014 based on Aug. 2013.

243

244 Analogously, when comparing the observational data in Aug. 2014 with that in  
245 Jul. and Sept. 2014 (the months before and after the most aggressive abatement), the  
246 concentrations of most species also decreased obviously. As presented in Fig. 5 and  
247 Fig. 6, without abatement, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub> and O<sub>3</sub> were likely to exceed NAAQS,  
248 especially for PM<sub>2.5</sub> and O<sub>3</sub>. As shown in Table 4 and Table 5, compared with Jul.  
249 2014, the concentration of NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub> dropped by 0.7%, 31.8%,  
250 33.7%, 1.1%, and 52.8%, respectively at CCM station in Aug. 2014, while the  
251 concentration of SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub> decreased by 15.8%, 39.6%,  
252 34.6%, 7.1%, and 50.7%, respectively at XL station in Aug. 2014. Without emission  
253 control, the concentration of air pollutants rebounded in Sept. 2014. Compared to  
254 Aug., the concentration of SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub> and O<sub>3</sub> increased by 37.4%, 19.8%,  
255 37.6%, 22.3%, and 47.2%, respectively at CCM station in Sept. 2014 (Table 4), while  
256 the concentration of SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub> increased by 24.6%, 21.8%,  
257 28.7%, 17.7%, 4.9%, and 39.9%, respectively at XL station in Sept. 2014 (Table 5). In  
258 addition, for most species, the standard deviation was the lowest in Aug., which meant  
259 that the potential of extreme events was the least in Aug.. Assume that the weather  
260 conditions in Jul., Aug., Sept., 2014 were nearly similar, it can be estimated that  
261 emission sources could be the major impact factor of explaining the concentration  
262 changes during the three months. These results proved that concentrations of most  
263 species decreased and had less potential in extreme events after aggressive emission  
264 abatement. However, the concentration became higher without emission control.

265 Section 3.3 would further discuss the change of pollutant concentration with and  
 266 without emission reduction based on model simulation.

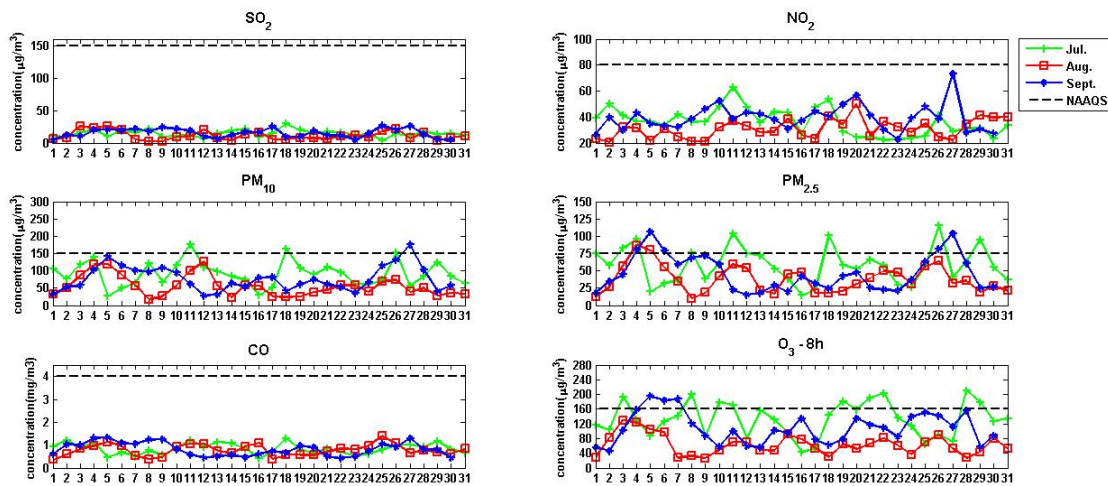
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268

269 **Fig. 5.** Day-to-day variations in SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub>-8h at CCM station in Jul., Aug.  
 270 and Sept. 2014 (CST). Observed data in Jul., Aug. and Sept. 2014 are indicated in green, red and blue,  
 271 respectively. NAAQS are indicated in black dotted line.

272



273

274 **Fig. 6.** Day-to-day variations in SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO and O<sub>3</sub>-8h at XL station in Jul., Aug. and  
 275 Sept. 2014 (CST). Observed data in Jul., Aug. and Sept. 2014 are indicated in green, red and blue,  
 276 respectively. NAAQS are indicated in black dotted line.

277

278 **Table 4**

279 Statistical analysis of hourly data in Jul. - Sept. 2014 at CCM station (The unit is µg/m<sup>3</sup> except CO  
 280 (mg/m<sup>3</sup>))

Species	Month	Max	Min	Mean	Median	Std	Δa	Δb
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	Jul. 2014	83.0	1.0	11.3	9.0	9.8		
SO <sub>2</sub>	Aug. 2014	72.0	2.0	11.9	10.0	7.8	5.1%	-37.4%
	Sept. 2014	70.0	4.0	19.0	18.0	9.9		
	Jul.-Sept. 2014	83.0	1.0	14.0	12.0	9.8		
	Jul. 2014	161.0	1.0	37.5	32.0	28.3		
NO <sub>2</sub>	Aug. 2014	110.0	1.0	37.3	35.0	18.6	-0.7%	-19.8%
	Sept. 2014	151.0	8.0	46.5	42.0	24.5		
	Jul.-Sept. 2014	161.0	1.0	40.2	37.0	24.4		
	Jul. 2014	255.0	6.0	88.5	88.0	50.7		
PM <sub>10</sub>	Aug. 2014	198.0	6.0	60.4	54.0	36.6	-31.8%	-37.6%
	Sept. 2014	243.0	6.0	96.7	90.0	45.8		
	Jul.-Sept. 2014	255.0	6.0	81.7	76.0	47.4		
	Jul. 2014	171.0	1.0	61.5	58.0	33.9		
PM <sub>2.5</sub>	Aug. 2014	165.0	3.0	40.7	36.0	23.8	-33.7%	-22.3%
	Sept. 2014	143.0	3.0	52.4	46.0	27.2		
	Jul.-Sept. 2014	171.0	1.0	51.5	45.0	29.9		
	Jul. 2014	2.7	0.2	0.9	0.9	0.3		
CO	Aug. 2014	2.2	0.3	0.9	0.9	0.3	-1.1%	21.1%
	Sept. 2014	2.1	0.1	0.8	0.7	0.4		
	Jul.-Sept. 2014	2.7	0.1	0.9	0.8	0.4		
	Jul. 2014	281.0	4.0	82.4	69.0	57.6		
O <sub>3</sub>	Aug. 2014	150.0	9.0	38.9	34.0	22.6	-52.8%	-47.2%
	Sept. 2014	240.0	6.0	73.6	61.0	49.2		
	Jul.-Sept. 2014	281.0	4.0	64.7	51.0	49.3		

281 Δa: The change percentage of species in Aug.2014 based on Jul. 2014.

282 Δb: The change percentage of species in Aug. 2014 based on Sept. 2014.

283

284 **Table 5**

285 Statistical analysis of hourly data in Jul. - Sept.2014 at XL station (The unit is μg/m<sup>3</sup> except CO

286 (mg/m<sup>3</sup>))

Species	Month	Max	Min	Mean	Median	Std	Δa	Δb
	Jul. 2014	61.0	1.0	14.5	12.0	10.3		
SO <sub>2</sub>	Aug. 2014	71.0	1.0	11.4	8.0	10.4	-21.2%	-24.6%
	Sept. 2014	75.0	1.0	15.1	14.0	10.3		
	Jul.-Sept. 2014	75.0	1.0	13.7	11.0	10.4		
	Jul. 2014	123.0	9.0	36.4	33.0	18.9		
NO <sub>2</sub>	Aug. 2014	95.0	7.0	30.7	27.0	15.0	-15.8%	-21.8%
	Sept. 2014	127.0	11.0	39.2	36.0	18.7		
	Jul.-Sept. 2014	127.0	7.0	35.4	32.0	18.0		
	Jul. 2014	300.0	4.0	91.3	85.0	48.9		

PM <sub>10</sub>	Aug. 2014	196.0	6.0	55.2	47.0	35.9	-39.6%	-28.7%
	Sept. 2014	226.0	9.0	77.3	70.0	40.3		
	Jul.-Sept. 2014	300.0	4.0	74.5	64.0	44.6		
	Jul. 2014	158.0	2.0	58.2	51.0	34.8		
PM <sub>2.5</sub>	Aug. 2014	157.0	3.0	38.0	34.0	24.1	-34.6%	-17.7%
	Sept. 2014	144.0	3.0	46.2	38.0	29.0		
	Jul.-Sept. 2014	158.0	2.0	47.4	40.5	30.7		
	Jul. 2014	2.0	0.3	0.8	0.8	0.4		
CO	Aug. 2014	2.0	0.3	0.8	0.7	0.3	-7.1%	-4.9%
	Sept. 2014	2.8	0.3	0.8	0.7	0.4		
	Jul.-Sept. 2014	2.8	0.3	0.8	0.7	0.4		
	Jul. 2014	238.0	2.0	78.4	67.0	55.6		
O <sub>3</sub>	Aug. 2014	148.0	2.0	38.7	32.0	28.3	-50.7%	-39.9%
	Sept. 2014	226.0	2.0	64.4	54.0	46.4		
	Jul.-Sept. 2014	238.0	2.0	60.3	48.0	47.7		

287  $\Delta$ a: The change percentage of species in Aug.2014 based on Jul. 2014.

288  $\Delta$ b: The change percentage of species in Aug. 2014 based on Sept. 2014.

289

### 290 3.2 Simulated impact of meteorological conditions

291 In this paper, the model configurations were the same as those set by Shu et al.  
 292 (2016), who has evaluated the model performance of WRF/CMAQ and proved the  
 293 model's reliability in simulating air quality in Nanjing.

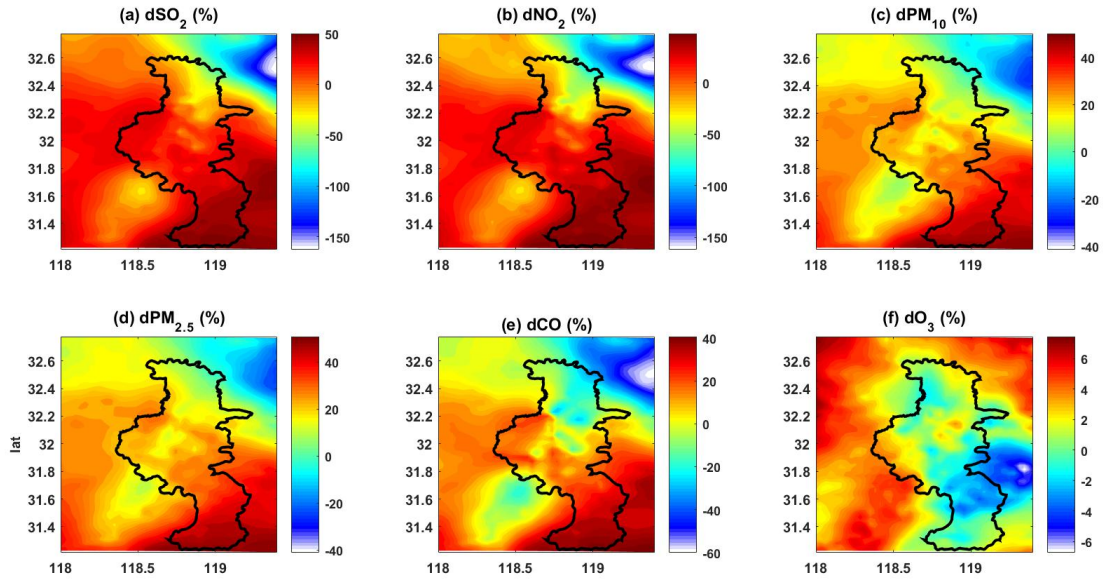
294 As we know, meteorology is an important impact factor on air quality. Good  
 295 diffusion conditions can alleviate air pollution in the short term (Cermak and Knutti,  
 296 2009; Wang et al., 2009b). In this premise, if two experiments (Exp. 2 and Exp. 3) use  
 297 the same emission inventory but different weather conditions, it can be concluded that  
 298 the higher concentrations may result from poor meteorological conditions. According  
 299 to model simulation, Exp. 2 exhibited higher pollutant concentrations for all species in  
 300 most part of Nanjing as shown in Fig. 7. For SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO, and O<sub>3</sub>,  
 301 their concentrations were increased by 17.5%, 16.9%, 18.5%, 18.8%, 7.8% and 0.8%  
 302 during Aug. 2014 compared to Aug. 2013. Additionally, the contributions of  
 303 meteorological conditions to primary and secondary particulate matters differed (See  
 304 Fig. 8). Secondary PM<sub>10</sub> (PM<sub>10s</sub>) was raised by 21.5%, while primary PM<sub>10</sub> (PM<sub>10p</sub>)  
 305 rose by 12.6% during Aug. 2014 compared to Aug. 2013. And secondary PM<sub>2.5</sub>

306 (PM<sub>2.5s</sub>) was increased by 21.5%, while primary PM<sub>2.5</sub> (PM<sub>2.5p</sub>) was enhanced by  
307 9.5%. Thus, the weather conditions had a slightly greater impact on secondary fine  
308 particulate matters. Moreover, for SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO, PM<sub>10p</sub>, PM<sub>10s</sub>, PM<sub>2.5p</sub>,  
309 and PM<sub>2.5s</sub>, there were some small decreasing areas in the northeast Nanjing (Fig. 7  
310 and Fig. 8). In domain 4, the simulated predominant wind was northeast wind in Aug.  
311 2014, while that was southeast wind in Aug. 2013. So, the diffusion condition of  
312 northeast Nanjing might be better in Aug. 2014 and resulted in small decrease in these  
313 areas.

314 The overall increasing pollutant levels in Aug. 2014 suggested that the diffusion  
315 conditions in Aug. 2014 were worse than those in Aug. 2013. Focus on the weather  
316 conditions during the YOG holding period (16-28 Aug., 2014) and the same period in  
317 2013, the simulated hourly mean 10-m wind speed in Nanjing was 1.5 m/s larger in  
318 16-28 Aug., 2013 than that of 16-28 Aug., 2014 (Fig. 9). Also, the simulated 2-m  
319 temperature was 2.0 K higher in 16-28 Aug., 2013 than that of 16-28 Aug., 2014 (Fig.  
320 9). Also, the simulated planetary boundary layer height (PBLH) was higher in Aug.  
321 2013, especially in 16-28 Aug., and it was 27.5 m higher than that of 16-28 Aug.,  
322 2014 (Fig. 9). Larger wind speed and higher PBLH benefited the diffusion of air  
323 pollutants. Warming on the ground surface was conducive to the promotion of  
324 convective instability and was also good for the vertical dilution and diffusion of  
325 pollutant. Thus, the diffusion conditions in 16-28 Aug. 2013 were better than those in  
326 16-28 Aug. 2014. Rather worse meteorological conditions in 16-28 Aug. 2014 implied  
327 that abatement controls might play a more important role in improving air quality in  
328 YOG compared with the same period in 2013. What's more, relative humidity, cloud  
329 cover and shortwave solar radiation all affect ozone chemical reaction (Katrakou et  
330 al., 2011; Pu et al., 2017). The generation and photochemical reaction of surface  
331 ozone depends on the availability of solar radiation. During the heat wave period, less  
332 relative humidity leads to less cloud cover, which could result in more net downward  
333 shortwave solar radiation and more production of O<sub>3</sub> (Pu et al., 2017). For ordinary O<sub>3</sub>,  
334 its production also corresponded well to the cloud cover. As shown in Fig. 9, more  
335 relative humidity resulted in more cloud cover in northern and eastern Nanjing during

336 16-28 Aug., 2013, which resulted in less O<sub>3</sub> in Aug. 2013, but more O<sub>3</sub> in Aug. 2014  
 337 (Fig. 7).

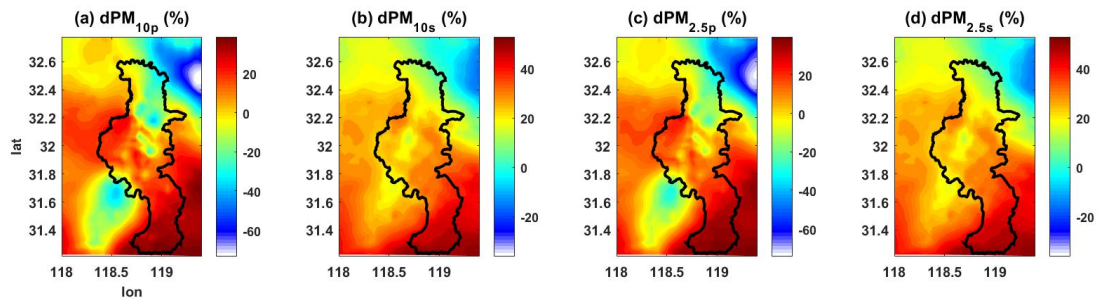
338



339

340 **Fig. 7.** Influence of meteorology on hourly mean concentrations of air pollutants in Aug. 2014  
 341 compared with Aug. 2013. (Black thick lines draw the outline of Nanjing. Picture a - f are hourly  
 342 average values of impact percentage ( $d_{\text{species}}(\%) = (\text{Exp. 2} - \text{Exp. 3}) / \text{Exp. 2} * 100\%$ ) of SO<sub>2</sub>, NO<sub>2</sub>,  
 343 PM<sub>10</sub>, PM<sub>2.5</sub>, CO, and O<sub>3</sub>, respectively.).

344

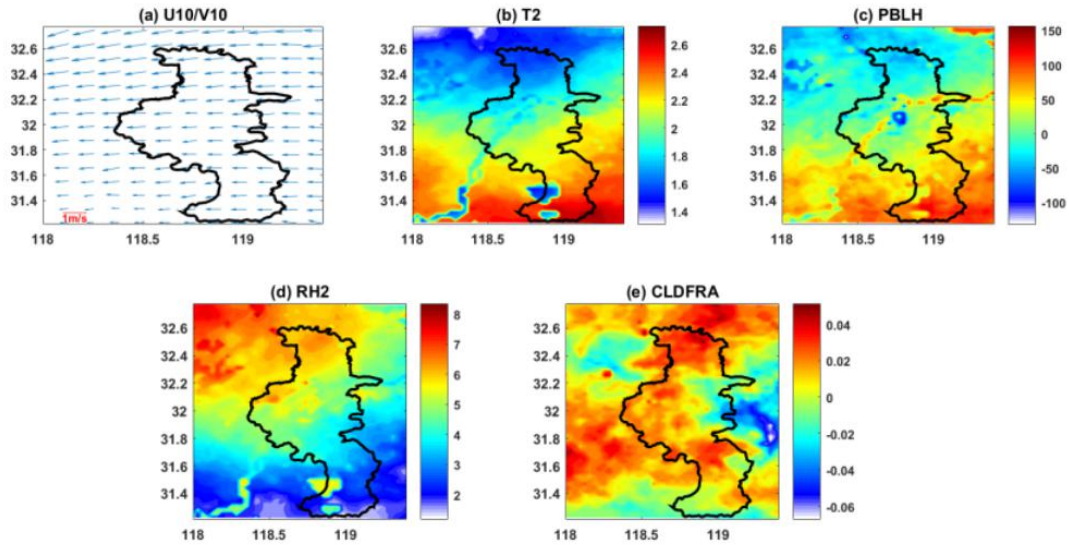


345

346 **Fig. 8.** Influence of meteorology on hourly mean concentrations of primary and secondary particulate  
 347 matters in Aug. 2014 compared with Aug. 2013. (Black thick lines draw the outline of Nanjing. Picture  
 348 a - d are hourly average values of impact percentage ( $d_{\text{species}}(\%) = (\text{Exp. 2} - \text{Exp. 3}) / \text{Exp. 2} * 100\%$ )  
 349 of PM<sub>10p</sub>, PM<sub>10s</sub>, PM<sub>2.5p</sub>, and PM<sub>2.5s</sub>, respectively.)

350





351  
 352 **Fig. 9.** Bias of simulated hourly mean meteorological conditions during the YOG. (Bias =  
 353 Meteorological Factors in 16-28 Aug., 2013 - Meteorological Factors in 16-28 Aug., 2014. (a) Bias of  
 354 Wind at 10m during 16-28 Aug. (unit: m/s), (b) Bias of temperature at 2m during 16-28 Aug. (unit: K),  
 355 (c) Bias of planetary boundary layer height during 16-28 Aug. (unit: m), (d) Bias of relative humidity at  
 356 2m during 16-28 Aug. (unit: %), (e) Bias of cloud fraction during 16-28 Aug.).

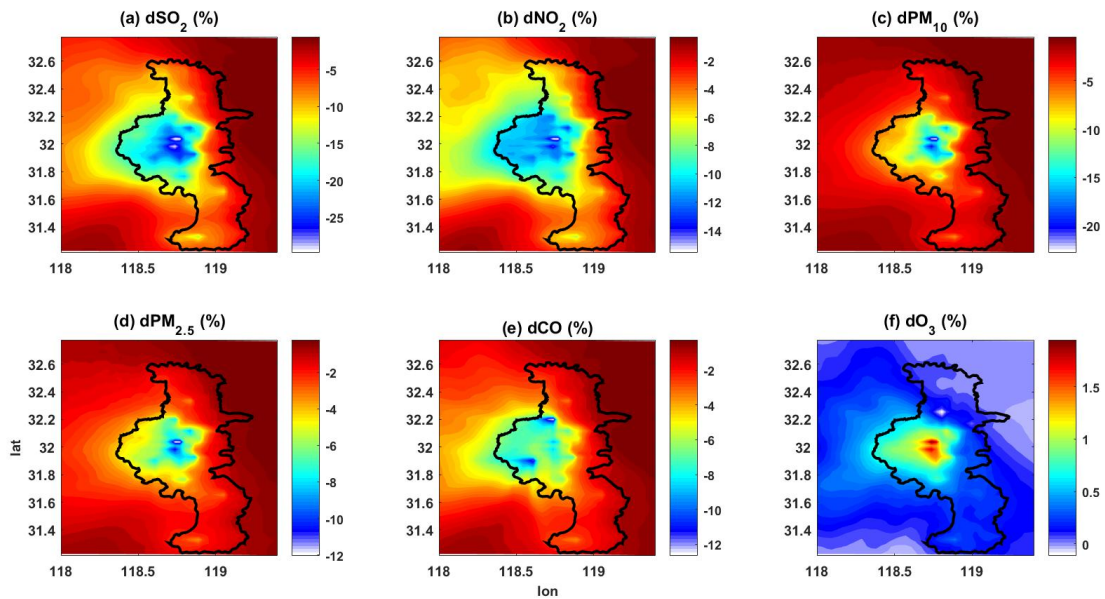
357

### 358 3.3 Simulated impact of emission reduction

359 As for SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, and CO, the distributions of such short-lived  
 360 chemical compositions are largely affected by their sources and sinks. As seen in Fig.  
 361 10, the simulated spatial distributions of concentration changes were uneven, large  
 362 variations were found in the west of Nanjing corresponding to the downwind regions  
 363 of heavy reduction districts (See Fig. 2). Impact percentages ( $ds_{\text{species}} (\%) = (\text{Exp. 1} -$   
 364  $\text{Exp. 2}) / \text{Exp. 2} * 100\%$ ) of species were negative except O<sub>3</sub>, implying that emission  
 365 regulatory efforts were effective on the other species, but counterproductive to O<sub>3</sub>.  
 366 Statistically, the concentrations of SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, and CO in Nanjing were  
 367 reduced by 24.6%, 12.1%, 15.1%, 8.1% and 7.2% during Aug. 2014. As for O<sub>3</sub>, the  
 368 variation was positive (1.3%), especially in the downwind area of NO<sub>x</sub> with heavy  
 369 reduction, which might due to the less titration of O<sub>3</sub> by NO<sub>x</sub> (Liu et al., 2013; Dong  
 370 et al., 2013). For primary and secondary particulate matters, the influence of emission  
 371 reduction varied dramatically. As shown in Fig. 11, PM<sub>10p</sub> was dropped by 39.6%,  
 372 while PM<sub>10s</sub> only declined by 2.9%. And PM<sub>2.5p</sub> was decreased by 26.2%, while  
 373 PM<sub>2.5s</sub> merely cut down by 2.9%. It seems that emission controls had much more

374 impacts on primary pollutants, especially on coarse particulate matters.

375



376

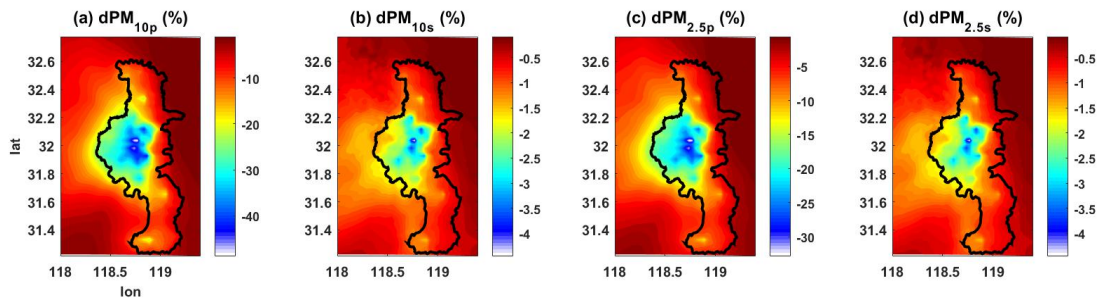
377 **Fig. 10.** Influence of emission reduction on hourly mean concentrations of air pollutants in Aug. 2014.

378 (Black thick lines draw the outline of Nanjing. Picture a - f are hourly average values of impact

379 percentage ( $\text{dspecies (\%)} = (\text{Exp. 1} - \text{Exp. 2}) / \text{Exp. 2} * 100\%$ ) of  $\text{SO}_2$ ,  $\text{NO}_2$ ,  $\text{PM}_{10}$ ,  $\text{PM}_{2.5}$ ,  $\text{CO}$  and  $\text{O}_3$ ,

380 respectively.).

381



382

383 **Fig. 11.** Influence of emission reduction on hourly mean concentrations of primary and secondary

384 particulate matters in Aug. 2014. (Black thick lines draw the outline of Nanjing. Picture a - d are hourly

385 average values of impact percentage ( $\text{dspecies (\%)} = (\text{Exp. 1} - \text{Exp. 2}) / \text{Exp. 2} * 100\%$ ) of  $\text{PM}_{10p}$ ,  $\text{PM}_{10s}$ ,

386  $\text{PM}_{2.5p}$  and  $\text{PM}_{2.5s}$ , respectively.).

387

### 388 3.4 Comparison of simulated meteorological factors and emission reduction

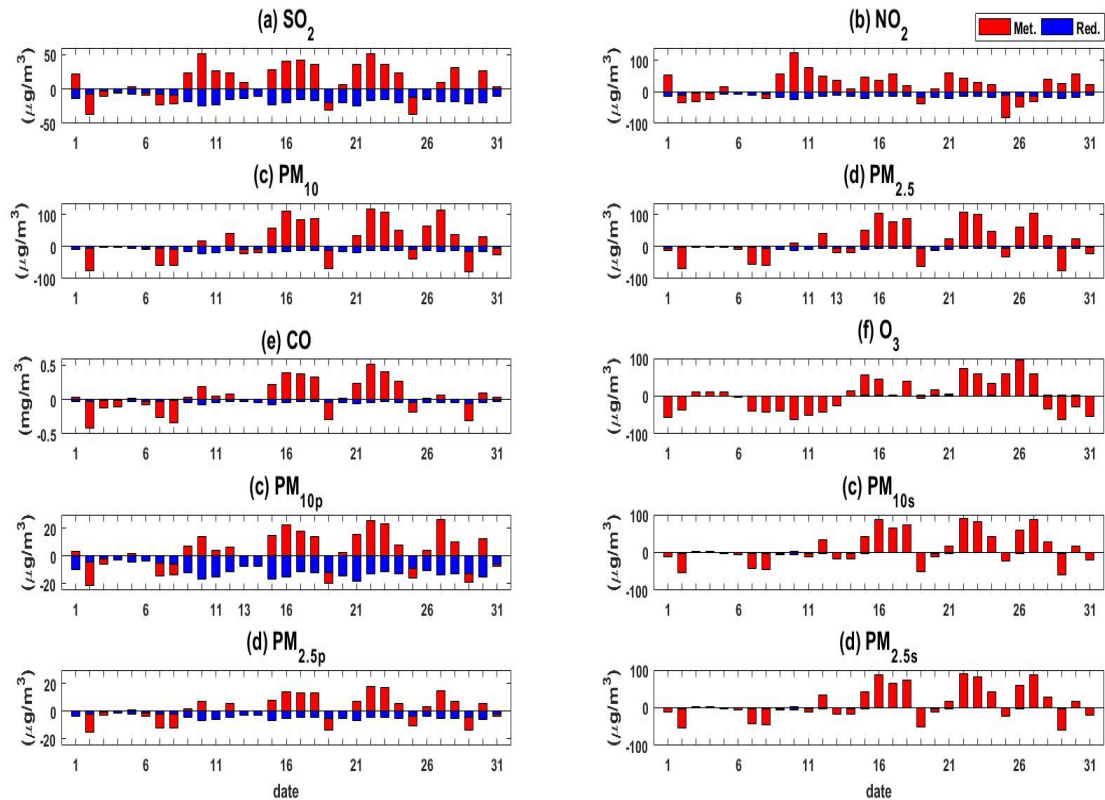
389 Fig. 12 displays the simulated effect of meteorological factors and emission

390 reduction in Nanjing on air quality improvement during Aug. 2014. In general,

391 meteorological conditions played a negative role in air quality promotion in most days,

392 only played a positive role in a few days (See discussion in Section 3.2). On the other

393 hand, emission reduction contributed to the decline of SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, CO,  
 394 PM<sub>10p</sub>, PM<sub>10s</sub>, PM<sub>2.5p</sub>, and PM<sub>2.5s</sub> all the time, especially for primary coarse particulate  
 395 matters. However, reduction of NO<sub>x</sub> could cause a slight rise of O<sub>3</sub>.  
 396



397  
 398 **Fig. 12.** The simulated effect of meteorology and emission reduction on pollutant concentrations in  
 399 Nanjing during 1-31 Aug. , 2014, Met. (Exp. 2-Exp. 3) represents the effect of meteorology, while Red.  
 400 (Exp. 1-Exp. 2) represents the simulated effect of emission reduction.

401  
 402 The opposite effects of meteorology and emission abatement on SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>,  
 403 PM<sub>2.5</sub>, CO, PM<sub>10p</sub>, PM<sub>10s</sub>, PM<sub>2.5p</sub> and PM<sub>2.5s</sub> during the whole month were significant  
 404 as statistically listed in Table 6. CCM station represents the urban status, XL station  
 405 represents the suburban status and NJ represents the whole city. Adverse  
 406 meteorological condition was found to raise the concentrations of the six pollutants as  
 407 17.4% for SO<sub>2</sub>, 15.1% for NO<sub>2</sub>, 15.6% for PM<sub>10</sub>, 14.9% for PM<sub>2.5</sub>, 6.4% for CO and  
 408 0.9% for O<sub>3</sub> at CCM station, and 14.1% for SO<sub>2</sub>, 12.4% for NO<sub>2</sub>, 22.4% for PM<sub>10</sub>,  
 409 24.5% for PM<sub>2.5</sub>, 2.3% for CO, and 1.6% for O<sub>3</sub> at XL station. On the contrary,  
 410 emission abatement reduced their levels in most cases, especially in the urban site. It  
 411 seems that the levels of air pollutants reduced more at CCM station compared to XL

412 station. Emission abatement independently led to a 24.3% decrease in SO<sub>2</sub> at CCM  
 413 station, which was 5.1% higher compared to XL station. Moreover, the cutbacks of  
 414 NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub> and CO were 11.7%, 13.9%, 7.5% and 7.0%, respectively at CCM  
 415 station, being larger by 1.0% to 2.0% compared with XL station. Though O<sub>3</sub> under  
 416 emission reduction scenarios resulted in a slightly rise (0.9% to 1.3%) at both sites,  
 417 the effectiveness of emission abatement was remarkable generally. As for primary and  
 418 secondary particulate matters, meteorological factors also played a negative role  
 419 during Aug. 2014, and had slightly more effect on secondary fine particles. Emission  
 420 controls seemed to cause tremendous impact on primary particulate matters,  
 421 especially for primary coarse particles. Emission abatement led to a 38.3% decrease in  
 422 PM<sub>10p</sub> at CCM station, a 33.2% decrease in PM<sub>10p</sub> at XL station, and a 39.6% decrease  
 423 in PM<sub>10p</sub> for the whole city. For secondary particulate matters, including sulfate,  
 424 nitrate, and ammonium salt, emission reduction played rather minor role in cutting the  
 425 pollutants, with merely a 2.4% decrease in PM<sub>10s</sub> and PM<sub>2.5s</sub> at CCM station, a 2.9%  
 426 decrease in PM<sub>10s</sub> and PM<sub>2.5s</sub> at XL station.

427

428 **Table 6**

429 Comparison between the simulated effect of meteorology and emission reduction at CCM and XL  
 430 station

Species	Met. (CCM)	Red. (CCM)	Met. (XL)	Red. (XL)	Met. (NJ)	Red. (NJ)
SO <sub>2</sub>	17.4%	-24.3%	14.1%	-19.2%	17.5%	-24.6%
NO <sub>2</sub>	15.1%	-11.7%	12.4%	-10.2%	16.9%	-12.1%
PM <sub>10</sub>	15.6%	-13.9%	22.4%	-11.9%	18.5%	-15.1%
PM <sub>2.5</sub>	14.9%	-7.5%	24.5%	-6.3%	18.8%	-8.1%
CO	6.4%	-7.0%	2.3%	-5.5%	7.8%	-7.2%
O <sub>3</sub>	0.9%	1.3%	1.6%	0.9%	0.7%	1.5%
PM <sub>10p</sub>	13.2%	-38.3%	5.9%	-33.2%	12.6%	-39.6%
PM <sub>10s</sub>	16.7%	-2.4%	29.4%	-2.9%	21.5%	-2.9%
PM <sub>2.5p</sub>	8.4%	-25.8%	4.9%	-20.1%	9.5%	-26.2%
PM <sub>2.5s</sub>	16.7%	-2.4%	29.4%	-2.9%	21.5%	-2.9%

431 Met.: The change percentage of species in Exp. 2 based on Exp3, represents the effect of meteorology.

432 Red.: The change percentage of species in Exp. 1 based on Exp 2, represents the effect of Nanjing local  
 433 emission reduction.

434

435 The decrease of SO<sub>2</sub> might result from the limit and halt of power plants and  
436 improvement of coal-combustion. The prohibition of heavy pollution vehicles could  
437 contribute to the drop of NO<sub>2</sub> and CO. Also, limiting the production of industries  
438 helped to reduce NO<sub>2</sub> and CO. The cut of particulate matters might be attributable to  
439 the stop of construction process and use of low ash content coal for industry. For  
440 secondary particulate matters, controlling the emission of SO<sub>2</sub> and NO<sub>x</sub> could help to  
441 reduce the formation of sulfate and nitrate, but no control on the emission of NH<sub>3</sub>  
442 could still result in quite a part of ammonium salt. The response of O<sub>3</sub> to emission  
443 control could be due to its non-linearity chemistry (Fu et al., 2012), reducing NO<sub>2</sub>  
444 may have side-effect by increasing O<sub>3</sub> because of the titration effect. On the whole,  
445 the meteorological factors and emission reduction during the YOG had opposite  
446 effects on SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, and CO, and emission reduction indeed played a  
447 very important role in the air quality improvement.

448

#### 449 **4 Summary and conclusions**

450 The air quality during the 2nd YOG was superior according to the current  
451 NAAQS. Both observation and modeling confirmed that stringent emission reductions  
452 were effective to ambient air quality promotion during the Youth Olympic Games,  
453 especially to SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub> and CO. Compared to Aug. 2013, the observed  
454 concentrations in Aug. 2014 were dropped by 64.7% for SO<sub>2</sub>, 29.8% for PM<sub>10</sub>, 9.8%  
455 for PM<sub>2.5</sub>, 8.9% for CO and 31.7% for O<sub>3</sub> at CCM station, while 50.0% for SO<sub>2</sub>,  
456 18.6% for NO<sub>2</sub>, 32.8% for PM<sub>10</sub>, 4.1% for PM<sub>2.5</sub>, and 31.7% for O<sub>3</sub> at XL station. The  
457 simulated impact percentage of emission reductions were -24.6%, -12.1%, -15.1%,  
458 -8.1% and -7.2% for SO<sub>2</sub>, NO<sub>2</sub>, PM<sub>10</sub>, PM<sub>2.5</sub>, and CO, respectively.

459 The meteorological conditions in the holding time of the YOG were inferior to  
460 those of the same period in 2013, with lower temperature and weaker winds. Model  
461 simulations show that unfavorable weather conditions caused higher concentrations  
462 for all species, including O<sub>3</sub> which was increased due to less cloud cover. Besides,  
463 meteorological conditions had slightly more effect on secondary fine particular  
464 matters compared to primary fine particular matters. Emission reduction could cut

465 down the levels of SO<sub>2</sub>, NO<sub>2</sub>, CO and particular matters, especially for primary coarse  
466 particles. Thus, emission reduction control is the very important factor for the air  
467 quality improvement during the YOG.

468 In general, better air quality during the YOG benefit a lot from emission  
469 reduction, which has set up a good example in air protection for important social  
470 events. However, the enhanced concentrations of air pollutants after the YOG (in Sept.  
471 2014) suggest that short-term emission control can only ease air pollution effectively  
472 and temporarily. Long-term control policies are necessary to ensure pleasant future air  
473 quality.

474

## 475 **5 Data availability**

476 The air quality monitoring records are available at <http://222.190.111.117:8023/>.  
477 The Multi-resolution Emission Inventory for China (MEIC) is available at  
478 <http://www.meicmodel.org/>. The WRF model is available at  
479 <http://www2.mmm.ucar.edu/wrf/users/downloads.html>. The CMAQ model is  
480 available at <https://www.cmascenter.org/cmaq/>.

481

482 *Competing interests.* The authors declare that they have no conflict of interest.

483

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493

494 **References**

- 495 Binkowski, F. S. and Roselle, S. J.: Models-3 Community Multiscale Air Quality  
496 (CMAQ) model aerosol component 1. Model description, *J. Geophys. Res.*, 108,  
497 4183, doi:10.1029/2001JD001409, 2003.
- 498 Byun, D. and Schere, K. L.: Review of the governing equations, computational  
499 algorithms, and other components of the models-3 Community Multiscale Air  
500 Quality (CMAQ) modeling system, *Appl. Mech. Rev.*, 59, 51-77,  
501 doi:10.1115/1.2128636, 2006.
- 502 Cai, H. and Xie, S.: Traffic-related air pollution modeling during the 2008 Beijing  
503 Olympic Games: the effects of an odd-even day traffic restriction scheme,  
504 *Science of the total environment*, 409, 1935-1948, doi:  
505 10.1016/j.scitotenv.2011.01.025, 2011.
- 506 Center for Earth System Science, Tsinghua University: Multi-resolution Emission  
507 Inventory for China, <http://www.meicmodel.org/>, 2015.
- 508 Cermak, J. and Knutti, R.: Beijing Olympics as an aerosol field experiment,  
509 *Geophysical Research Letters*, 36, 1-5, doi:10.1029/2009GL038572, 2009.
- 510 Chen P., Wang T., and Hu X.: Chemical Mass Balance Source Apportionment of  
511 Size-Fractionated Particulate Matter in Nanjing, China, *Aerosol & Air Quality  
512 Research*, 15, 1855-1867. doi:10.4209/aaqr.2015.03.0172, 2015.
- 513 Chen, P., Wang, T., Lu, X., Yu, Y., Kasoar, M., Xie, M., and Zhuang, B.: Source  
514 apportionment of size-fractionated particles during the 2013 Asian Youth Games  
515 and the 2014 Youth Olympic Games in Nanjing, China, *Science of the Total  
516 Environment*. 579, 860-870. <http://dx.doi.org/10.1016/j.scitotenv.2016.11.014>,  
517 2017.
- 518 Ding, J., Van, d. A. R. J., Mijling, B., Levelt, P. F., and Hao, N.: NO<sub>x</sub> emission  
519 estimates during the 2014 Youth Olympic Games in Nanjing, *Atmos. Chem.  
520 Phys.*, 15, 9399-9412, doi: 10.5194/acp-15-9399-2015, 2015.
- 521 Dong, X. Y., Gao, Y., Fu, J. S., Li, J., Huang, K., Zhuang, G. S., and Zhou, Y.: Probe  
522 into gaseous pollution and assessment of air quality benefit under sector  
523 dependent emission control strategies over megacities in Yangtze River Delta,

524 China, *Atmos. Environ.*, 79, 841-852, doi:10.1016/j.atmosenv.2013.07.041,  
525 2013.

526 Foley, K. M., Roselle, S. J., Appel, K. W., Bhave, P. V., Pleim, J. E., Otte, T. L.,  
527 Mathur, R., Sarwar, G., Young, J. O., Gilliam, R. C., Nolte, C. G., Kelly, J. T.,  
528 Gilliland, A. B., and Bash, J. O.: Incremental testing of the Community  
529 Multiscale Air Quality (CMAQ) modeling system version 4.7, *Geosci. Model*  
530 *Dev.*, 3, 205-226, doi:10.5194/gmd-3-205-2010, 2010.

531 Fu, J. S., Dong, X. Y., Gao, Y., Wong, D. C., and Lam, Y. F.: Sensitivity and linearity  
532 analysis of ozone in East Asia: the effects of domestic emission and  
533 intercontinental transport, *Journal of the Air and Waste Management Association*,  
534 62, 1102-1114, doi:10.1080/10962247.2012.699014, 2012.

535 Hao, N., Valks, P., Loyola, D., Cheng, Y. F., and Zimmer, W.: Space-based  
536 measurements of air quality during the World Expo 2010 in Shanghai,  
537 *Environmental Research Letters*, 6, doi:10.1088/1748-9326/6/4/044004, 2011.

538 Jiang, F., Wang, T. J., Wang, T. T., Xie, M., and Zhao, H.: Numerical modeling of a  
539 continuous photochemical pollution episode in Hong Kong using WRF-chem,  
540 *Atmos. Environ.*, 42, 8717-8727, doi:10.1016/j.atmosenv.2008.08.034, 2008.

541 Jiang, F., Zhou, P., Liu, Q., Wang, T. J., Zhuang, B. L., and Wang, X. Y.: Modeling  
542 tropospheric ozone formation over East China in springtime, *J. Atmos. Chem.*,  
543 69, 303-319, doi:10.1007/s10874-012-9244-3, 2012.

544 Katragkou E., Zanis P., Kioutsioukis I., Tegoulas I., Melas D., Krüger, B. C., and  
545 Coppola E.: Future climate change impacts on summer surface ozone from  
546 regional climate-air quality simulations over Europe, *Journal of Geophysical*  
547 *Research Atmospheres*, 116, D22307, doi:10.1029/2011JD015899, 2011.

548 Li, M. M., Song, Y., Mao, Z. C., Liu, M. X., and Huang, X.: Impacts of thermal  
549 circulations induced by urbanization on ozone formation in the Pearl River Delta  
550 region, China, *Atmos. Environ.*, 127, 382-392,  
551 doi:10.1016/j.atmosenv.2015.10.075, 2016.

552 Liao, J. B., Wang, T. J., Wang, X. M., Xie, M., Jiang, Z. Q., Huang, X. X., and Zhu, J.  
553 L.: Impacts of different urban canopy schemes in WRF/Chem on regional



554 climate and air quality in Yangtze River Delta, China, *Atmos. Res.*, 145, 226-243,  
555 doi:10.1016/j.atmosres.2014.04.005, 2014.

556 Liao, J. B., Wang, T. J., Jiang, Z. Q., Zhuang, B. L., Xie, M., Yin, C. Q., Wang, X. M.,  
557 Zhu, J. L., Fu, Y., and Zhang, Y.: WRF/Chem modeling of the impacts of urban  
558 expansion on regional climate and air pollutants in Yangtze River Delta, China,  
559 *Atmos. Environ.*, 106, 204-214, doi:10.1016/j.atmosenv.2015.01.059, 2015.

560 Lin Y., Huang, K., Zhuang, G., Fu, J. S., Xu, C., Shen, J., and Chen, S.: Air Quality  
561 over the Yangtze River Delta during the 2010 Shanghai Expo, *Aerosol & Air  
562 Quality Research*, 13, 1655-1666, doi: 10.4209/aaqr.2012.11.0312, 2013.

563 Liu, H., Wang, X. M., Zhang, J. P., He, K. B., Wu, Y., and Xu, J. Y.: Emission controls  
564 and changes in air quality in Guangzhou during the Asian Games, *Atmos.  
565 Environ.*, 76, 81-93, doi:10.1016/j.atmosenv.2012.08.004, 2013.

566 Ministry of Environmental Protection of the People's Republic of China: The second  
567 Summer Youth Olympic Games environmental quality assurance work plan,  
568 Beijing, [http://www.js.xinhuanet.com/2014-07/19/c\\_1111695515\\_2.htm/](http://www.js.xinhuanet.com/2014-07/19/c_1111695515_2.htm/), 2014.

569 National Bureau of Statistics of China: <http://www.stats.gov.cn/>, 2014.

570 Okuda, T., Matsuura, S., Yamaguchi, D., Umemura, T., Hanada, E., Orihara, H.,  
571 Tanaka, S., He, K.B., Ma, Y. L., Cheng, Y., and Liang, L. L.: The impact of the  
572 pollution control measures for the 2008 Beijing Olympic Games on the chemical  
573 composition of aerosols, *Atmos. Environ.*, 45, 2789-2794,  
574 doi:10.1016/j.atmosenv.2011.01.053, 2011.

575 Pu X., Wang T. J., Huang X., Melas D., Zanis P., Papanastasiou D. K., and Poupkou  
576 A.: Enhanced surface ozone during the heat wave of 2013 in Yangtze River Delta  
577 region, China. *Science of the Total Environment*, 603-604, 807-816,  
578 doi:10.1016/j.scitotenv.2017.03.056, 2017.

579 Shu L., Xie M., Wang T. J., Gao D., Chen P. L., Han Y., Li S., Zhuang B. L., Li M. M.:  
580 Integrated studies of a regional ozone pollution synthetically affected by subtrop  
581 ical high and typhoon system in the Yangtze River Delta region, China. *Atmos.  
582 Chem. Phys.*, 16, 15801-15819. doi:10.5194/acp-16-15801-2016, 2016.

583 Skamarock, W. C., Klemp, J. B., Dudhia, J., Gill, D. O., Barker, D. M., Duda, M. G.,

584 Huang, X. Y., Wang, W., and Powers, J. G.: A Description of the Advanced  
585 Research WRF Version 3. NCAR Tech Notes-475+STR, 2008.

586 State Environmental Protection Administration of China: China National  
587 Environmental Protection Standard: Automated Methods for Ambient Air  
588 Quality Monitoring, China Environmental Science Press, Beijing, 2006.

589 Streets, D. G., Fu, J. S., Jang, C. J., Hao, J. M., He, K. B., Tang, X. Y., Zhang, Y. H.,  
590 Wang, Z. F., Li, Z. P., Zhang, Q., Wang, L. T., Wang, B. Y., and Yu, C.: Air  
591 quality during the 2008 Beijing Olympic Games, *Atmos. Environ.*, 41, 480-492,  
592 doi:10.1016/j.atmosenv.2006.08.046, 2007.

593 Wang, X., Westerdahl, D., Chen, L. C., Wu, Y., Hao, J. M., Pan, X. C., Guo, X. B.,  
594 and Zhang, K. M.: Evaluating the air quality impacts of the 2008 Beijing  
595 Olympic Games: On-road emission factors and black carbon profiles, *Atmos.*  
596 *Environ.* 43, 4535-4543, doi:10.1016/j.atmosenv.2009.06.054, 2009a.

597 Wang, W. T., Primbs, T., Tao, S., Zhu, T., and Simonich, S. L.: Atmospheric  
598 particulate matter pollution during the 2008 Beijing Olympics, *Environmental*  
599 *Science & Technology*, 43, 5314-5320, 2009b.

600 Wang, T., Nie, W., Gao, J., Xue, L. K., Gao, X. M., Wang, X. F., Qiu, J., Poon, C. N.,  
601 Meinardi, S., Blake, D., Wang, S. L., Ding, A. J., Chai, F. H., Zhang, Q. Z., and  
602 Wang, W. X.: Air quality during the 2008 Beijing Olympics: secondary  
603 pollutants and regional impact, *Atmos. Chem. Phys.*, 10, 7603-7615,  
604 doi:10.5194/acp-10-7603-2010, 2010.

605 Xie, M., Zhu, K. G., Wang, T. J., Yang, H. M., Zhuang, B. L., Li, S., Li, M. G., Zhu, X.  
606 S., and Ouyang, Y.: Application of photochemical indicators to evaluate ozone  
607 nonlinear chemistry and pollution control countermeasure in China, *Atmos.*  
608 *Environ.*, 99, 466-473, doi:10.1016/j.atmosenv.2014.10.013, 2014.

609 Xie, M., Liao, J., Wang, T., Zhu, K., Zhuang, B., Han, Y., Li, M., and Li, S.: Modeling  
610 of the anthropogenic heat flux and its effect on regional meteorology and air  
611 quality over the Yangtze River Delta region, China, *Atmos. Chem. Phys.*, 16,  
612 6071-6089, doi:10.5194/acp-16-6071-2016, 2016.

613 Xing, J., Zhang, Y., Wang, S. X., Liu, X. H., Cheng, S. H., Zhang, Q., Chen, Y. S.,

614 Streets, D. G., Jang, G., Hao, J. M., and Wang, W. X.: Modeling study on the air  
615 quality impacts from emission reductions and atypical meteorological conditions  
616 during the 2008 Beijing Olympics, *Atmos. Environ.*, 45, 1786-1798,  
617 doi:10.1016/j.atmosenv.2011.01.025, 2011.

618 Xu, H. M., Tao, J., Ho, S. S. H., Ho, K. F., Cao, J. J., Li, N., Chow, J. C., Wang, G. H.,  
619 Han, Y. M., Zhang, R. J., Watson, J. G., Zhang, J. Q.: Characteristics of fine  
620 particulate non-polar organic compounds in Guangzhou during the 16th Asian  
621 Games: Effectiveness of air pollution controls, *Atmos. Environ.*, 76, 94-101,  
622 doi:10.1016/j.atmosenv.2012.12.037, 2013.

623 Yarwood, G., Rao, S., Yocke, M., and Whitten G.: Updates to the Carbon Bond  
624 chemical mechanism: CB05, Final Report to the US EPA, RT-0400675, 2005.

625 Zhou, D., Li, B., Huang, X., Virkkula, A., Wu, H., Zhao, Q., Zhang, J., Liu, Q., Li, L.,  
626 Li, C., Chen, F., Yuan, S., Qiao, Y., Shen, G., and Ding, A.: The Impacts of  
627 Emission Control and Regional Transport on PM<sub>2.5</sub> Ions and Carbon Components  
628 in Nanjing during the 2014 Nanjing Youth Olympic Games, *Aerosol & Air  
629 Quality Research*, 17, 730-740, doi: 10.4209/aaqr.2016.03.0131, 2017.

630 Zhou, Y., Wu, W., Yang, L., Fu, L. X., He, K. B., Wang, S. X., Hao, J. M., Chen, J. C.,  
631 Li, C. Y.: The impact of transportation control measures on emission reductions  
632 during the 2008 Olympic Games in Beijing, China, *Atmos. Environ.*, 44,  
633 285-293, doi:10.1016/j.atmosenv.2009.10.040, 2010.