

# **A modelling study of air quality impact of odd-even day traffic restriction scheme before, during and after the 2008 Beijing Olympic Games**

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**Abstract.** Systematic air pollution control measures were designed and implemented to improve air quality for the 2008 Beijing Olympics. This study focuses on the evaluation of the air quality impacts of a short-term odd-even day traffic control scheme (TRS) implemented before, during and after the Games, based on modelling simulation by a well validated urban-scale air quality model. Concentration levels of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> were predicted for the pre- (10<sup>th</sup>-19<sup>th</sup>, July), during- (20<sup>th</sup> July-20<sup>th</sup> September) and post-TRS (21<sup>st</sup>-30<sup>th</sup>, September) periods, based on the on-line monitored traffic flows on a total of 334 road segments constituting the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> Ring Roads (RR) and the major Linkage Roads (LRs) that were subject to the TRS policy and distributed around the main urban area of Beijing, and on the hourly sequential meteorological data from a representative Observatory. Subsequently, we used the predictions and observations at a roadside air quality monitoring site to evaluate the model, based on a widely used statistical framework

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for model evaluation, as well as on the dependence of model performance on time-of-the-day and on wind direction, and the model predictions turned out satisfactory. Results showed that daily average concentrations on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and LRs during the TRS period decreased significantly, by about 35.8%, 38.5%, 34.9% and 35.6% for CO, about 38.7%, 31.8%, 44.0% and 34.7% for PM<sub>10</sub>, about 30.3%, 31.9%, 32.3% and 33.9% for NO<sub>2</sub>, and about 36.7%, 33.0%, 33.4% and 34.7% for O<sub>3</sub>, respectively, compared with the pre-TRS period. Besides, hourly average concentrations were also reduced significantly, particularly for the morning and evening peaks for CO and PM<sub>10</sub>, for the evening peak for NO<sub>2</sub>, and for the afternoon peak for O<sub>3</sub>. Consequently, both the daily and hourly concentration level of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> conformed to the CNAAQS (China National Ambient Air Quality Standards) Grade II during the Games. Besides, a notable ozone weekend effect was revealed for the pre- and post-TRS periods, and was virtually removed for the during-TRS period. In addition, notable reduction of concentration levels were achieved in different regions of Beijing in response to the TRS policy, with the air quality in the downwind northern and western regions improved most significantly. The TRS policy was therefore effective in improving short-term air quality in Beijing during the Games.

## **1. Introduction**

Beijing, the capital city of China and the host city of the 2008 Olympic Games, has a population of over 18 million and meanwhile suffers serious air pollution constituted

by high concentration levels of PM<sub>10</sub>, CO, SO<sub>2</sub> and NO<sub>2</sub>. Particularly, PM<sub>10</sub>, with its high daily and annual concentrations on average, remained the primary air pollutant in Beijing since 2000 (Beijing EPB, 2009). The PM<sub>10</sub> annual average concentration in Beijing has been staying at a high level, fluctuating around 160 µg/m<sup>3</sup> during the period of 2001-2006, and began to decrease in the following two years as a result of various control measures, with the annual average concentration in 2008 still exceeding 120 µg/m<sup>3</sup> (Beijing EPB, 2009), which was about 20% higher than the China National Ambient Air Quality Standards (CNAAQS) Grade II (100 µg/m<sup>3</sup>) and six times the latest World Health Organization (WHO) Air Quality Guidelines (WHO, 2005). Of the major anthropogenic sources of atmospheric particulate matters in the mega cities (e.g. Beijing, Shanghai, Guangzhou) in China, on-road vehicular emissions is an important and perhaps the fastest growing one (Chan and Yao, 2008). Although the source contributions of motor vehicles to PM<sub>10</sub> and PM<sub>2.5</sub> in Beijing revealed by the PMF (positive matrix factorization) methodology were only 5% (Xie et al., 2008) and 6% (Song et al., 2006b), respectively, motor vehicles were a major contributor to ambient concentrations of nitrogen dioxide (NO<sub>2</sub>)/nitrogen oxides (NO<sub>x</sub>) and carbon monoxide (CO), according to a conclusion that on-road vehicle source had contributed 76.5% and 68.4% of the CO and NO<sub>x</sub> concentrations, respectively, in urban atmosphere of Beijing in 1995 (Hao et al., 2001). Moreover, traffic congestion and traffic-related air pollution has been a serious issue in urban Beijing with the vehicle population increasing dramatically at a daily rate of over one

thousand, reaching over 3.5 million by the end of 2008. The booming economic prosperity and substantial increase of vehicle population has resulted in the exponential growth of vehicular emissions of CO, PM<sub>10</sub>, NO<sub>x</sub>, VOC (volatile organic compounds) (Cai and Xie, 2007), as well as speciated VOC emissions containing highly reactive and toxic pollutants in the atmosphere of Beijing (Cai and Xie, 2009). In order to improve effectively the air quality and traffic condition in Beijing, a Sino-Italian environmental protection program based on Intelligent Traffic System and Traffic Air Pollution (ITS-TAP) monitoring was launched between the Italian Ministry for the Environment and Territory and Beijing Municipal Government in 2005. Moreover, the Beijing Municipal Government had committed to the international society that air quality in Beijing would be improved and be better than before, satisfying the CNAAQS and WHO Air Quality Guidelines during the 2008 Olympic Games. To fulfill the air quality commitment during the Games, the government implemented a list of control measures, including greening of bare land by afforestation both in Beijing and some surrounding provinces, enhancing the utilization of natural gas to replace coal for heat supply and residential cooking, closing or relocating heavy industrial polluters (e.g., the Capital Steel Company), reducing local power generation by importing electricity from the surrounding areas, promoting abatement and removal technologies for sulfur dioxide and particulate matter from industrial point sources, suspending construction activities as well as imposing strict control of VOC evaporation at gas stations (Wang et al., 2009a). In

addition, air pollution sources were strictly controlled in the surrounding provinces of Tianjin, Hebei, Shanxi, Neimenggu and Shandong to prevent regional contribution through long-range transportation, based on the evidences obtained in previous studies (Streets et al., 2007 and Wang et al., 2008a). Also, five control schemes for controlling vehicular emissions were implemented: issuance of new automobile emissions standards; decommissioning of high emissions vehicles, buses and taxis; recovery of fuel vapors at pumping stations and from tankers; banning of non-local heavy duty diesel trucks within the Beijing Administrative area; and control of emissions from small stationary diesel generators (COC, 2008). Meanwhile, Beijing took the lead in China to adopt the Euro-IV emission standard to reduce vehicle emissions of air pollutants five months before the Olympic Games. Particularly, an odd-even day traffic restriction scheme (TRS), a control measure that was demonstrated to be effective in Atlanta (Fang et al., 2009), was enforced for two months from 20<sup>th</sup> July to 20<sup>th</sup> September, 2008, to help ease congestion and improve air quality during the Olympics and Paralympics. Consequently, the UAB had a distinct characteristic of source emissions compared to usual periods, as emissions from sectors other than on-road vehicles were expected to decrease significantly, with on-road vehicles becoming the major source for air pollution in that particular period, due to their huge population even after the TRS policy was in effect. Previously, Cheng et al. (2008) demonstrated that the four-day traffic restrictions in Beijing during the Sino-African Summit in early November 2006 resulted in significant

temporary reductions in concentrations of NO<sub>x</sub> and particulates in the city. Besides, Westerdahl et al. (2009) conducted in-situ measurements and concluded that a four-day traffic control experiment from 17<sup>th</sup>-20<sup>th</sup> August, 2007 conducted by the Beijing Government as a pilot to test the effectiveness of the proposed odd-even day TRS was effective in reducing extreme concentrations that occurred at both on-road and ambient environments. Wang et al. (2009a) evaluated the air quality impacts of the 2008 Beijing Olympic Games, focusing on the measurement of on-road black carbon emission factors and the evaluation of reduction of black carbon concentrations. Wang et al. (2009b) conducted a modelling analysis, aiming to assess the effectiveness of various emission restrictions implemented during the 2008 Beijing Olympics on the ozone air quality at a rural site of Beijing during the period. As the control measures for sectors other than on-road vehicles were constantly in effect and remained the same for the pre-, during- and post-TRS periods, this study intends to seize this unique opportunity to study the air quality impact and effectiveness of the TRS policy, by adopting an integrated urban-scale modelling system with online-monitored data of on-road traffic flows at a high temporal resolution of two seconds from the ITS-TAP system, and based on the analysis of the short-term air quality variation near the simulated roads in response to the TRS policy.

Model-based simulation has been one of the major tools for air quality assessment, air pollution diagnosis and evaluation of pollution control policies at the urban, regional and global levels. A few dispersion models have been widely applied to

simulate urban air pollution from traffic-related emissions, e.g. OSPM (Berkowicz, 2000; Kukkonen et al., 2001; Assael et al., 2008; Berkowicz et al., 2008), CALINE (Levitin et al., 2005; Yura et al., 2007), CALPUFF (Wang et al., 2006), ISCST3 (Elbir, 2002; Ying et al., 2007; Sharma and Chandra, 2008). ADMS-Urban, a well tested and intensively validated quasi-Gaussian dispersion air quality model, which is widely used for regulatory purposes in the UK and used in the investigation and assessment of air pollution mitigation and control strategies in many cities of China (Carruthers et al., 1994; Bennett and Hunter, 1997; McHugh et al., 1997; Carruthers et al., 1999; Carruthers et al., 2000a; Carruthers et al., 2001; Riddle et al., 2004; McHugh et al., 2005; Hirtl and Baumann-Stanzer, 2007), has recently been validated by the Ministry of Environmental Protection of China as one of the three recommended dispersion models (the others are the US EPA's CALPUFF and AERMOD) for air quality impact assessment (MEPC, 2009), and by a commercial system of street-level air quality forecasting which was launched in Beijing in July, 2008 (BeijingAir, 2008). Moreover, ADMS-Urban has some features especially tuned to obtain best performance on urban areas (CERC, 2003), which is mostly due to its advantages over other recommended models, like the up to date parameterization of atmospheric boundary layer structure based on the Monin-Obukhov length and the boundary layer height, and the Gaussian concentration distributions in stable and neutral conditions, but non-Gaussian vertical distributions in convective conditions to take into account the skewed structure of the

vertical component of the turbulence. Furthermore, it is particularly convenient to set up on-road emissions, and to define the road source geometry and the output presentation in an accurate way, by means of the nested GIS tool (Arcview or Mapinfo). Thus, ADMS-Urban was adopted in this study to simulate and assess the air quality impact in UAB in response to the TRS policy before, during and after the 2008 Beijing Olympic Games. Firstly, the modelled results were evaluated with the measurement data from a typical roadside air quality monitoring site. Subsequently, the ambient concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> at a height of 1.5 metres were calculated for the periods of pre-TRS (10<sup>th</sup>–19<sup>th</sup> July, 2008), during-TRS (20<sup>th</sup> July–20<sup>th</sup> September, 2008) and post-TRS (21<sup>st</sup>–30<sup>th</sup> September, 2008), respectively, based on traffic flows on the dense network of 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> Ring Roads (RR) and Linkage Roads (LRs, including major intercity expressways and intercity roads between RRs), followed by an assessment of the temporal and spatial variation of air quality impacts in response to the TRS policy.

## **2. Methods and data**

### **2.1 Description of study domain, urban and background monitoring sites, and receptors**

The study domain, the main UAB, covers the 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> RR, which were 32.7, 48.0 and 65.3 kilometers long, and had six, six to eight and eight lanes, respectively, as well as the major LRs mainly having eight lanes. These simulated roads were the main roads of UAB with the majority of traffic flows in it. About 62% of Beijing is



mountainous area located in the west, the north and the northeast. Thus, the local wind field has a clear diurnal variation, with northeasterly and southeasterly winds dominating in the daytime and southeasterly wind dominating in the night. The Chegongzhuang (CGZ) air quality monitoring site, which was located five kilometers west of the West 2<sup>nd</sup> RR and was on a corner (geographical coordinates: 39°55'53"N, 116°19'38"E) of a crossroad where a five-lane North-South oriented street and a six-lane West-East oriented street intersected with high traffic flows, had a sampling height of about 4.5 metres from ground and had been well maintained with routine calibration of the measurement equipment by the Beijing Municipal Environmental Monitoring Centre during the Games as a traffic monitoring site. This site provided the model evaluation data of hourly concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> for 10<sup>th</sup>-20<sup>th</sup> July and 10<sup>th</sup>-20<sup>th</sup> August, 2008. To assure the accuracy of the model predictions, it is important to account for significant underlying, or 'background' levels of pollutants in the atmosphere, and to account for any sources of pollution that are not otherwise included in the model run. Therefore, we selected the background diurnal profiles of CO, PM<sub>10</sub> and NO<sub>2</sub> for the whole evaluation period from Dingling (DL) air quality monitoring station, which was approximately 42 kilometers north from the CGZ air quality monitoring site and was in the suburban area with very few motor vehicles. Background measurements from DL could reflect well the real situation of the air quality in the UAB during the Games, which was expected to be influenced mainly by local on-road vehicles, with a minor contribution from other

190 well controlled sources like power plants and polluting industrial plants, construction  
191 sites and gas stations both in Beijing and the surrounding provinces during the Games  
192 (Wang et al., 2009a). Therefore, it is reasonable to make comparisons between  
193 modelled predictions and measurement data at CGZ for the model evaluation. For  
194 O<sub>3</sub>, a major secondary air pollutant, we used the measurement data from Shangdianzi  
195 (SDZ) regional atmospheric background monitoring site, one of the four regional  
196 atmospheric background monitoring sites in China and located about 150 kilometres  
197 northeast of Beijing (40°39'N, 117°07'E). Measurement data from the SDZ  
198 background measurement site are free of influence by motor vehicles and represent  
199 the background characteristic of atmospheric constituents in northern regions of China  
200 including Beijing (Liu et al., 2007). Therefore, this site is suitable for providing the  
201 background O<sub>3</sub> measurement data for this modelling study. Due to the lack of the  
202 measurement data from this site for the summer in 2008, we have adopted the  
203 measurement data in SDZ for the summer periods in 2004 (Liu et al., 2006) and in  
204 2006 (Liu et al., 2008) as a substitute, considering the generally accepted  
205 understanding of the relatively constant feature of background concentrations within a  
206 short time period. Finally, we used the averaged daily profiles for July, August and  
207 September based on the 2004 and 2006 data from SDZ as the background O<sub>3</sub>. To  
208 reflect the variation of air quality in the areas near the dense network of simulated  
209 roads, 31 representative receptors located along the 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> RR and the LRs  
210 were chosen, where the hourly concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> were

simulated for the pre-TRS, during-TRS and post-TRS periods. Besides, receptors located in different regions of the UAB were used to capture the air quality variation in different parts of the UAB in response to the TRS policy. The road network of Beijing and the road sources of RR and LRs distributed in the study domain, including the locations of the CGZ, DL and SDZ air quality monitoring sites and the representative receptors, are shown in Figure 1.

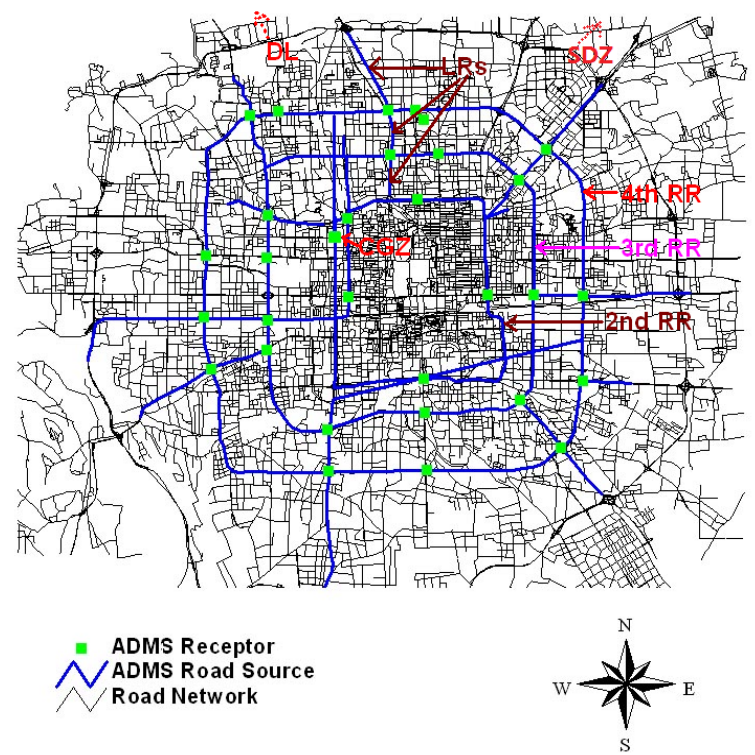


Fig. 1. Map of the road network of UAB, road sources of RR and LRs, CGZ, DL and SDZ air quality monitoring sites, and representative receptors distributed in the study domain.

## 2.2 ADMS-Urban model set up

The ADMS-Urban model, developed by Cambridge Environmental Research Consultants (CERC, 2009), is a quasi-Gaussian dispersion model for the dispersion

simulation of pollutants released from industrial, domestic and road traffic sources in urban areas. Particularly, ADMS-Urban has an advantage over other urban-scale dispersion models as it characterizes the boundary layer structure based on the Monin-Obukhov length and boundary layer height rather than the more imprecise characterization achieved with the Pasquill-Gifford stability parameter (CERC, 1999).

Various parameters which need to be set for running ADMS-Urban include surface roughness, the latitude of the modelling area, minimum Monin-Obukhov length, the chemical reaction scheme, the meteorological data and the height of recorded wind, the background data, the source data including the road width, elevation, and the canyon heights in case of modeling street canyons, the time varying factors for weekdays, Saturdays and Sundays, the road geometry indicated by the geographical coordinates of the constituting nodes, and the output parameters like the pollutants included, the number of grids for the modelling area and the heights of concentrations to be calculated for the modelling domain and the receptors, respectively.

#### 2.2.1 Descriptive parameters, chemical reaction scheme and deposition

Descriptive parameters were used to describe the modelled area, for example, the surface roughness characterised the surrounding area in terms of the effects it would have on wind speed and turbulence, which is one of the key components of the modelling. The values chosen for some representative parameters for the modelling domain is given in Table 1.

Table 1.

Parameters	Value
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Surface roughness (m)	1.0
Latitude of the modelling area ( °)	40
Minimum Monin-Obukhov length (m)	30
Pollutants included	NO <sub>x</sub> , NO <sub>2</sub> , CO, PM <sub>10</sub> and O <sub>3</sub>
Number of grids	10000
Horizontal resolution	0.35km×0.35km
Height of concentrations to be calculated (m)	1.5

The value of surface roughness was chosen based on some local studies (Hu, 1994; Zhang and Chen, 1997; Lu et al., 2002) and on the recommended value for modelling big cities, which is 1.0 metre. The value of 30 metres was chosen as the minimum Monin-Obukhov length as recommended by ADMS-Urban for cities.

ADMS-Urban has two chemistry options to calculate NO<sub>2</sub> and O<sub>3</sub> concentrations. The first option uses the empirical function (Derwent and Middleton, 1996). The second one is based on a chemistry scheme known as the Generic Reaction Set (GRS) (Venkatram et al., 1994), which is a semi-empirical model that simplifies photochemical reactions using a set of parameters derived from observations. The GRS scheme can work when background NO<sub>x</sub>, NO<sub>2</sub> and O<sub>3</sub> data, as well as hourly sequential meteorological data including cloud cover are available. In ADMS-Urban, the primary NO<sub>2</sub>/NO<sub>x</sub> emissions fraction is assumed constant (5%) with the GRS scheme, and the NO<sub>2</sub> photo-dissociation coefficient is calculated using cloud cover data, in place of the radiation data. In this study, we adopted the GRS scheme to simulate the important reactions involving NO<sub>x</sub>, VOC and O<sub>3</sub>, since it has been demonstrated that the GRS scheme produced better results than the other option (D-M) (Vardoulakis et al., 2007).

The dry deposition process was considered by the dry deposition module of

ADMS-Urban, which is actually a resistance model dependent of the pollutant species, the nature of the surface and the wind speed (CERC, 2009), while the wet deposition process was ignored during the modelling since the dry air and lack of rain during the evaluation period were not favorable for wet deposition of pollutants (Wang et al., 2008c).

#### 2.2.2 Emissions, road geometry, traffic flows, fleet compositions and emission factors

By implementation of high intensity of administrative regulations for emission control, the natural, industrial, agricultural, construction, biomass burning and residential emissions were expected to be reduced significantly and become sparse, and the compilation of a complete emission inventory including the scarce and scattered emissions from sectors other than on-road vehicles was very difficult, and the inaccuracy of modelling could be considerable if all sources were considered. Furthermore, emissions from other minor roads were not considered either, partly because traffic flows on these roads were relatively less than those on the monitored main roads, and partly because emissions from the minor roads were difficult to quantify accurately due to the lack of an online continuous monitoring system for traffic flows. Most importantly, variations in air quality near the simulated roads, which were reflected by the receptors located near the monitored main roads distributed around the UAB, were focused on and analyzed, to fulfill the objective of evaluating the air quality impact and effectiveness of the TRS policy, rather than analyze the air quality variations throughout the UAB, which benefited from all

control measures including the TRS policy. Therefore, emissions considered in this study only involved with the vehicular emissions from the dense network of monitored main roads, with additional background information representing contributions from other sources of pollution that were not included in the model run. We believe this approach is a good substitute for a complete emission inventory that was difficult to compile accurately and could probably raise large uncertainty, and is reasonable to assure the credibility of modelling results and conclusions, particularly when the primary objective of this study is concerned, although we admit that the major drawback of not using a complete emission inventory in the simulations might cause some inaccuracy in the absolute values of the predictions, particularly in the areas near other ignored minor roads or away from the monitored major roads.

The 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> RR, as well as the major LR were separated into a total of 334 road segments, the lengths of which were automatically identified by Arcview, a nested GIS (Geographical Information System) software of ADMS-Urban, by recognizing the geographical coordinates of the road nodes constituting each segment. The width of each road segment was manually input based on field surveys. Unlike another study that focused on the traffic-related air pollution within twelve typical street canyons of Beijing using the OSPM model (Wang and Xie, 2009), this study focuses on the modelling of traffic-related air pollution around the UAB, and thus the street canyon effect was not modelled, with further consideration on the fact that the majority of roads in the UAB were not typical street canyons, of which the building

heights on both sides were expected to be at least three times of the road width.

Traffic flows on each of the road segments of the 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> RR were intensively monitored automatically every two seconds from 10<sup>th</sup> July, 2008 to 30<sup>th</sup> September, 2008 by the ITS-TAP system. The high temporal resolution traffic flow data were further processed to summarize the hourly traffic flow and the hourly average running speed on each of the monitored road segments for the whole assessment period.

For the quality assurance and control of the traffic data, we first screened out the abnormal maximum values of monitored driving speed, which were recorded as 240 km/h. We regarded this to be impossible and abnormal, with the confirmation by technical experts of the ITS-TAP system. Further, these abnormal maximum speeds were replaced by the average of the normal speeds on the adjacent road segments. Moreover, the major problem of the monitored traffic flow data was the missing data on certain road segments for some periods of time. Accordingly, we treated those hourly sequential data sets with a fraction of missing data over 10% by replacing them with the average of the traffic flows on the nearest upstream and downstream road segments, which had less than 10% of missing data. In addition, we eliminated the few days (6<sup>th</sup>, 13<sup>th</sup>, 14<sup>th</sup>, 24<sup>th</sup> and 28<sup>th</sup>, September) when large quantities of missing data were found (data missing by about 43%, 36%, 47%, 84% and 73%, respectively, for those days), due to the malfunction of monitoring cameras of the ITS-TAP system on many road segments during the days. In this way, we assured that the monitored



driving speed was proper, and traffic flows were relatively complete, with a possible underestimation of less than 10%.

The on-line monitored traffic flow data consisted of two vehicle categories: long vehicles which were constituted of buses and heavy duty vehicles, and short vehicles, which were constituted of passenger cars and light duty vehicles. The further split-up of the fleet on each of the road segments into these four categories were based on the surveyed fleet composition of the manual traffic counts conducted by technical assistants at the roadsides of 290 road segments in urban Beijing in 2004.

Three different sets of time-varying factors for diurnal traffic flows were identified separately for weekdays, Saturdays and Sundays, based on average traffic counts of all the road segments for each hour of the day throughout the assessment period. These profiles were used to derive time-varying emission rates (g/km/s) on all road segments for weekdays, Saturdays and Sundays, respectively, which are thought to reflect variations in diurnal emissions due to road traffic congestion or improvement in traffic conditions.

Continuous emissions of CO, PM<sub>10</sub>, NO<sub>x</sub> and VOC from vehicles on the monitored roads were provided every hour for the model run. According to the definition of emission rates, as expressed by Equation (1), the emissions, lengths of road segments and travelling time of vehicles are needed, to calculate the emission rates of CO, PM<sub>10</sub>, NO<sub>x</sub> and VOC on each road segment. More specifically, emission rates can be calculated based on four parameters: emission factors (*EF*), vehicle population (*VP*),

lengths of road segments ( $L$ ) and driving speed of vehicles ( $v$ ), as shown by Equation (4), based on Equations (2-3). Traffic flows derived from the ITS-TAP system included the vehicle population of various vehicle types and the corresponding driving speeds on each segment. Emission factors of each pollutant from various vehicle categories were calculated by COPERT, a European emission factor model (EEA, 2000), based on the average running speed, ambient temperature and fuel property for a given vehicle category. The travelling time ( $t$ ) on each road segment was calculated based on the speeds and road lengths that were automatically identified by Arcview, a nested GIS (Geographical Information System) software of ADMS-Urban.

$$ER_s = \frac{E_s}{L_s \times t_s} \quad (1)$$

$$E_s = \sum_i EF_i \times VP_i \times L_s \quad (2)$$

$$t_s = \frac{L_s}{v_s} \times 3600 \quad (3)$$

$$ER_s = \frac{\sum_i EF_i \times VP_i \times v_s}{3600 \times L_s} \quad (4)$$

Where:  $ER_s$  is the emission rate on road segment  $s$ , expressed in  $g/km/s$ ;  $E_s$  is the emission on road segment  $s$ ;  $L_s$  is the length of road segment  $s$ ;  $t_s$  is the time used to travel on the road segment  $s$ ;  $EF_i$  is the emission factor of vehicle type  $i$ ;  $VP_i$  is the vehicle population of vehicle type  $i$ ; and  $v_s$  is the running speed of vehicles on road segment  $s$ .

Using the established methodology (Cai and Xie, 2007), emission factors of

passenger cars, light duty vehicles, heavy duty vehicles and buses at a typical urban running speed of 20 km/h in Beijing were calculated, as shown by Table 2. Particularly, emission factors of each vehicle category on each road segment were updated in accordance with the temporal variation of running speed to improve the accuracy of source emission estimation.

Table 2. Emission factors (g/km) for passenger cars, light duty vehicles, heavy duty vehicles and buses at a typical urban running speed of 20 km/h in Beijing, calculated by COPERT and from literature reports.

	PM <sub>10</sub>	CO	NO <sub>x</sub>	VOC
Passenger cars	0.015a	5.99	0.44	0.35
Light duty vehicles	0.015a	11.28	0.82	0.47
Heavy duty vehicles	2.94a	2.22	2.72	0.91
Buses	2.94a	4.76	10.54	0.87

a: from Wang et al. (2001), as COPERT assumes gasoline vehicles do not emit particles.

As for ozone, a secondary air pollutant, its model-predicted concentrations over the simulated domain were based on the background measurement from SDZ background monitoring site and the results of the GRS chemical mechanism operated by ADMS-Urban, given the information of precursor pollutant emissions of NO<sub>x</sub> and VOC.

### 2.2.3 Meteorological data

Transport and dispersion of air pollutants in the atmosphere are influenced by regional weather patterns. Operating on one hour intervals, meteorological data for

ADMS-Urban input files include wind speed, wind direction (including anemometer height), cloud cover, precipitation and temperature, which were used by the meteorological processor of ADMS-Urban to calculate the parameters for use in the model such as boundary layer depth and Monin-Obukhov length. The complete hourly meteorological data during the assessment period were obtained from the Beijing Capital International Airport Meteorology Observatory (ZBAA), which is about 20 km northeast from the Northeast 4<sup>th</sup> Ring Road and is known to be representative of the wind fields for UAB (Wang et al., 2009a). The daytime (8 am and 2 pm) and nighttime (8 pm and 2 am) wind roses during the period of 10<sup>th</sup> July – 30<sup>th</sup> September, 2008 are shown by Figure 2, which reveals that northeasterly and southeasterly winds dominated in the daytime while southeasterly wind dominated in the nighttime. Consequently, the UAB was dominated by the southeasterly and northeasterly winds in the whole day during the assessment period. Besides, the majority of wind speeds during the assessment period was less than 5.0 m/s, which indicate that the dissipation of air pollutants tended to be prolonged. It was also noted that the percentage of calm wind conditions was very low (0.5%) during the assessment period, which is in favor of ADMS-Urban's performance, since modelling of traffic sources in low wind speed conditions can lead to over-predictions and is sensitive to the minimum value of wind speed used (Carruthers et al., 2000b).

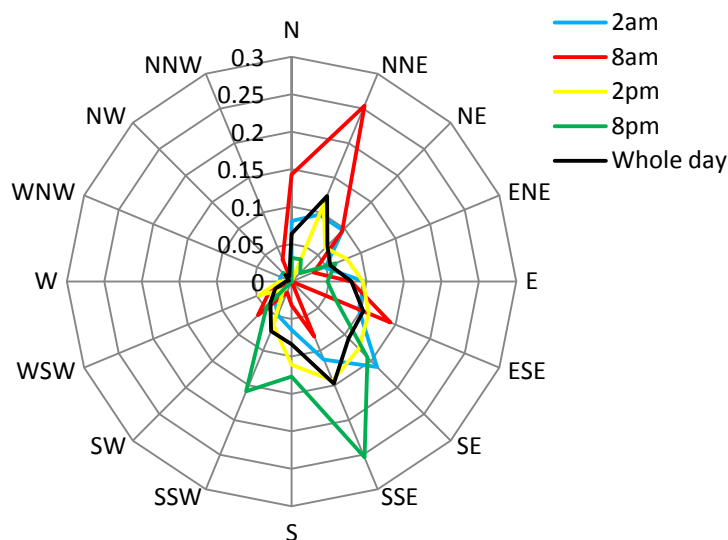


Fig. 2. Wind roses of wind directions at 2 am, 8 am, 2 pm, 8 pm and the whole day from July to September, 2008. The radius indicates the frequency of wind observed in each direction.

### 2.3 Model evaluation

The evaluation of the model performance was carried out by comparing the observed hourly concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> at CGZ monitoring site with model predictions for the periods of July 10<sup>th</sup>-20<sup>th</sup> and August 10<sup>th</sup>-20<sup>th</sup>, using the statistics recommended by Hanna et al (1991, 1993), which have been adopted as a common model evaluation framework for the European Initiative on “Harmonization within Atmospheric Dispersion Modelling for Regulatory Purposes” (Olesen, 2001). The statistical performance measures include (a) the fractional bias (FB) showing the tendency of the model to overpredict or underpredict; (b) the normalized mean square error (NMSE) showing the overall accuracy of the model; (c) the geometric mean bias (MG) showing the mean relative bias and indicating systematic errors; (d) the

geometric variance (VG) showing the mean relative scatter and reflecting both systematic and random errors; (e) the Pearson correlation coefficient (R) describing the degree of association between observed concentrations and model results; and (f) the fraction of predictions within a factor of two of observations (FAC2). These statistical measures are defined as follows:

$$FB = \frac{\overline{C_o} - \overline{C_p}}{0.5(\overline{C_o} + \overline{C_p})} \quad (5)$$

$$NMSE = \frac{(\overline{C_o} - \overline{C_p})^2}{\overline{C_o} \overline{C_p}} \quad (6)$$

$$MG = \exp(\overline{\ln C_o} - \overline{\ln C_p}) \quad (7)$$

$$VG = \exp[\overline{(\ln C_o - \ln C_p)^2}] \quad (8)$$

$$R = \frac{(\overline{C_o} - \overline{C_o})(\overline{C_p} - \overline{C_p})}{\sigma_{C_p} \sigma_{C_o}} \quad (9)$$

$$FAC2 = \text{fraction of data that satisfy } 0.5 \leq \frac{C_p}{C_o} \leq 2.0 \quad (10)$$

Where:  $C_p$  are model predictions;  $C_o$  are observations;  $\overline{C}$  is the average over the dataset; and  $\sigma_C$  is the standard deviation over the dataset.

Since NMSE accounts for both systematic and random errors, it is helpful to partition NMSE into the component due to systematic errors, NMSEs, and the unsystematic component due to random errors, NMSEu. NMSEs, the minimum NMSE without any unsystematic errors, was defined by Hanna et al (1991) for a given value of FB as:

$$NMSEs = \frac{4FB^2}{4 - FB^2} \quad (11)$$

The above statistical measures, however, do not provide information about the model performance under diurnal variation of weather and traffic conditions. For

that reason, the following qualitative performance measures have also been used: (i) diurnal pollution profiles to identify specific periods of the day when the model had good/poor performance; and (ii) pollution roses to compare predictions and observations at each of the wind directions and thus to show the model performance for different wind directions.

### **3. Results and discussion**

#### **3.1 Model evaluation**

##### **3.1.1 Overall model performance**

The statistical evaluation results for CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> based on the hourly predictions and observations are summarized in Table 3. The model predicted the concentrations of all pollutants reasonably well, with the FAC2 between 0.50 and 0.71. Besides, the FB statistic revealed that ADMS-Urban had a moderate tendency to underestimate NO<sub>2</sub> (FB=0.12) and O<sub>3</sub> (FB=0.31) concentrations, and had a moderate tendency to overestimate CO (FB=-0.22) concentrations and a slight overestimation of PM<sub>10</sub> (FB=-0.0084) concentrations. The model showed satisfactory performance for NO<sub>2</sub> with a NMSE of 0.33, and moderate NMSE for CO, PM<sub>10</sub> and O<sub>3</sub>. Furthermore, the model performances indicated by NMSEs are very satisfactory for all pollutants, which agreed well with MG, the systematic error indicator. Besides, the model had a good performance for CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> concentrations when considering both systematic and random errors as indicated by VG. Although the Pearson correlation coefficient for CO is relatively low (0.34), the R values for PM<sub>10</sub>,

NO<sub>2</sub> and O<sub>3</sub> were much higher. The R, however, is not a very robust measure due to its sensitivity to a few outlier data pairs, and thus Willmott (1982) discourages the use of R, because it does not consistently relate to the accuracy of predictions. Instead, the FAC2, which is not overly influenced by either low or high outliers, is the most robust performance measure.

Table 3. Statistical evaluation of model performance based on hourly predictions and observations at CGZ monitoring site for the period of July 10<sup>th</sup>-20<sup>th</sup> and August 10<sup>th</sup>-20<sup>th</sup>, 2008.

	CO	PM <sub>10</sub>	NO <sub>2</sub>	O <sub>3</sub>
Observed mean (µg/m <sup>3</sup> )	1214.2	84.1	52.3	52.7
Predicted mean (µg/m <sup>3</sup> )	1459.6	84.8	46.5	38.3
FB (ideal value: 0)	-0.22	-0.01	0.12	0.31
FAC2 (ideal value: 100%)	0.71	0.65	0.69	0.50
NMSE (ideal value: 0)	0.76	0.54	0.33	0.71
NMSEs (ideal value: 0)	0.05	0.00	0.01	0.10
MG (ideal value: 1)	0.92	0.81	1.07	3.33
VG (ideal value: 1)	1.57	2.27	1.42	4.51
R (ideal value: 1)	0.34	0.45	0.56	0.68

A scatter plot between the observed and predicted concentrations at CGZ is presented in Figure 3, which illustrates the robust FAC2 statistic for CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub>, respectively. There was a moderate overestimation for CO probably due to the adoption of higher model-calculated emission factors, as shown by Figure 3 (a), although the FAC2 for CO is high (0.71). As shown by Figure 3 (b), predicted and observed PM<sub>10</sub> concentrations agreed very well with each other, with its scatter plot almost symmetrical. Figure 3 (c) reveals that a majority (69%) of predicted concentrations fall within a factor of two of the observations for NO<sub>2</sub>, with a slight



underestimation. The higher  $\text{NO}_2$  observations might derive from the enhanced photochemical reactions of NO with oxidants such as peroxy radicals that were present either in the atmosphere or in vehicle exhaust (Kenty et al., 2007). Ozone was moderately underestimated, as revealed by Figure 3 (d), but was still satisfactorily predicted when taking into account the limitation of the simplified GRS scheme adopted in predicting secondary pollutant like ozone.

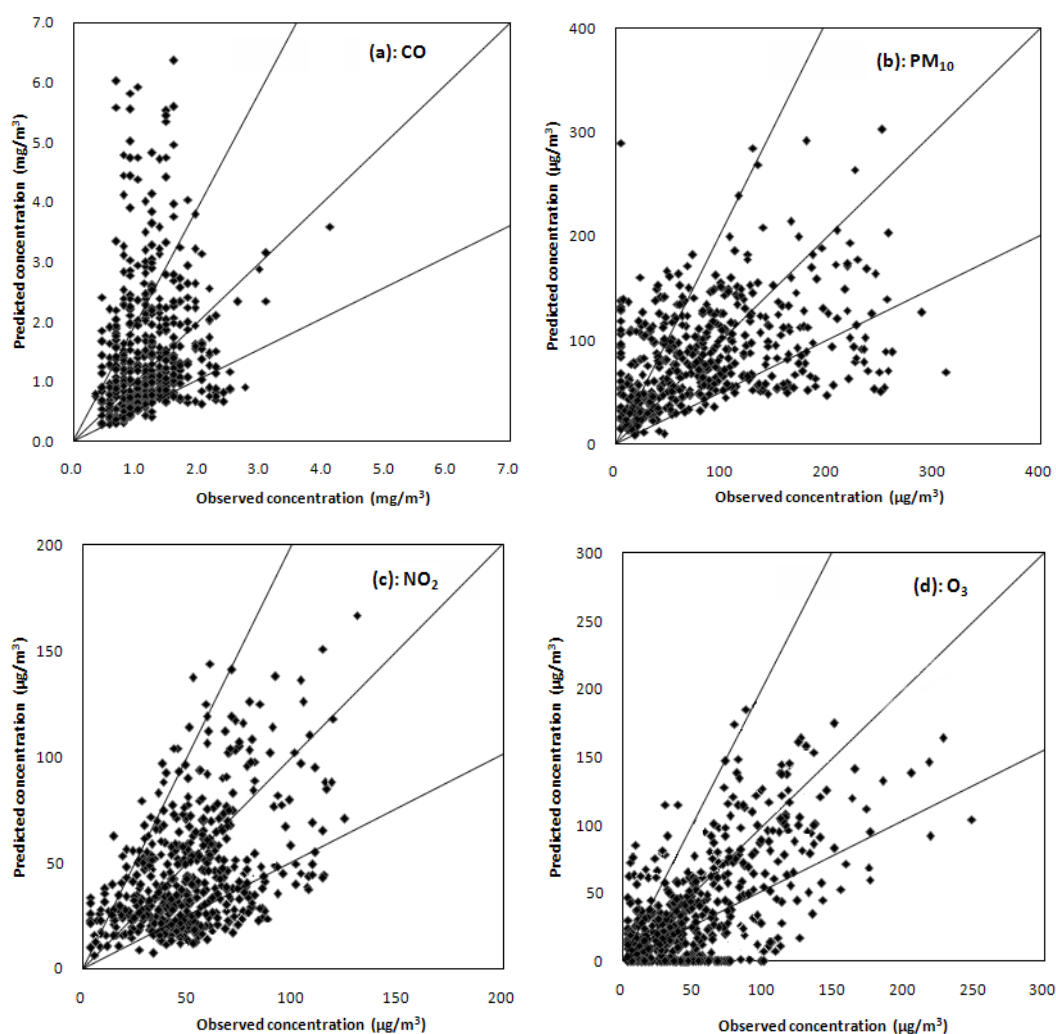


Fig. 3. Scatter plot comparison between observed and predicted hourly concentrations of (a) CO, (b) PM<sub>10</sub>, (c) NO<sub>2</sub> and (d) O<sub>3</sub> for the periods of July 10<sup>th</sup>-20<sup>th</sup> and August 10<sup>th</sup>-20<sup>th</sup>, 2008, at the CGZ air quality monitoring site, for the evaluation of ADMS-Urban model. The lines showing an agreement of predictions

and observations by a factor of two are also presented.

Figure 3 (b) and the statistical evaluation reveal that the model predictions of  $PM_{10}$  concentrations were very good compared to the observations, rather than an underestimation as some previous field measurements (Okuda et al., 2004; Sun et al., 2004), modelling studies (Song et al., 2006a; Wang et al., 2008a) and source apportionment work (Zheng et al., 2005; Song et al., 2006b; Song et al., 2007; Zhang et al., 2007; Wang et al., 2008b; Xie et al., 2008 ) would expect. These good  $PM_{10}$  predictions in this study were credible, primarily due to the distinct characteristic of source emissions during the Games when a series of control measures were taken for sectors other than on-road vehicles, which were expected to decrease significantly the contributions from soil dust, coal or fossil fuel combustion, industry, secondary sources and biomass burning, the identified major sources for  $PM_{10}$  or  $PM_{2.5}$  in Beijing by previous work (Okuda et al., 2004; Sun et al., 2004; Zheng et al., 2005; Song et al., 2006a,b; Zhang et al., 2007; Wang et al., 2008b; Xie et al., 2008). Consequently, on-road vehicles became the remaining major source, with relatively more  $PM_{10}$  emissions and a higher contribution for  $PM_{10}$  during the particular period. Moreover, the  $PM_{10}$  or  $PM_{2.5}$  samples used for the previous source apportionment studies in Beijing were collected with samplers located at the roofs of several sites, of which the heights ranged from one story (about 5 metres) to five stories (about 25 metres) (Zheng et al., 2005; Song et al., 2006b), and even to 40 metres (Zhang et al., 2007). These PM measurements obtained on the roofs of buildings tended to be less influenced by on-road vehicles of which the exhaust tailpipe height was usually below 0.3 meter. Besides, according to a previous study focusing on the vertical profile of PM near major roads, both the  $PM_{10}$  and  $PM_{2.5}$  concentrations decreased substantially on the top of a high-rise residential building compared to their ground-level values

near roadways (Wu et al., 2002). Therefore, the PM measurements used in the previous source apportionment studies tended to underestimate the PM contribution from on-road vehicles, while on-road vehicles considered in this study had a more substantial impact on the PM<sub>10</sub> concentrations, which were predicted at a much lower height of 1.5 metres. Furthermore, the background concentrations of PM<sub>10</sub> from a representative rural monitoring station was added during the modelling, to take into account the contributions from other minor source emissions that were not included specifically in the model run. Thus, the distinct characteristic of source emissions during the particular Olympic period, larger influence of on-road vehicles on predicted PM<sub>10</sub> concentrations in this study than previous measurements, and the use of background data from a rural monitoring station, resulted in reasonably good predictions compared to the observations, rather than an underestimation of predicted PM<sub>10</sub>, and the slight overestimation of PM<sub>10</sub> was therefore not coincidental at the station chosen for model evaluation.

### 3.1.2 Dependence of model performance on time-of-the-day

Diurnal profiles on hourly average have been separately plotted for all pollutants measured and predicted at CGZ, as shown by Figure 4. The model captured well the diurnal variations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub>, and did particularly well for NO<sub>2</sub> and O<sub>3</sub> at predicting the observed peaks. The predicted morning peaks of CO and PM<sub>10</sub> were about three hours earlier than the observed, but were in good agreement with the morning peak of traffic flows. In addition, the temporal variations of predicted CO and PM<sub>10</sub> concentrations changed more dramatically than the observed, with a notably overprediction in peak CO concentrations, which was mainly related to an

overestimation of the CO emission factor adopted by the model, resulting in a relatively larger overestimation of emission rates during the rush hours with higher traffic flows than other periods of the day. Moreover, Figure 4 showed that the model had an excellent performance in predicting the diurnal profile of NO<sub>2</sub> and O<sub>3</sub>, which revealed the satisfactory model performance in reasonably predicting the temporal variation of secondary gases like NO<sub>2</sub> and O<sub>3</sub> with the application of the GRS chemical scheme. Consequently, the model captured the evening peak NO<sub>2</sub> concentrations, and the afternoon peak ozone concentrations accompanied by the trough NO<sub>2</sub> concentrations.

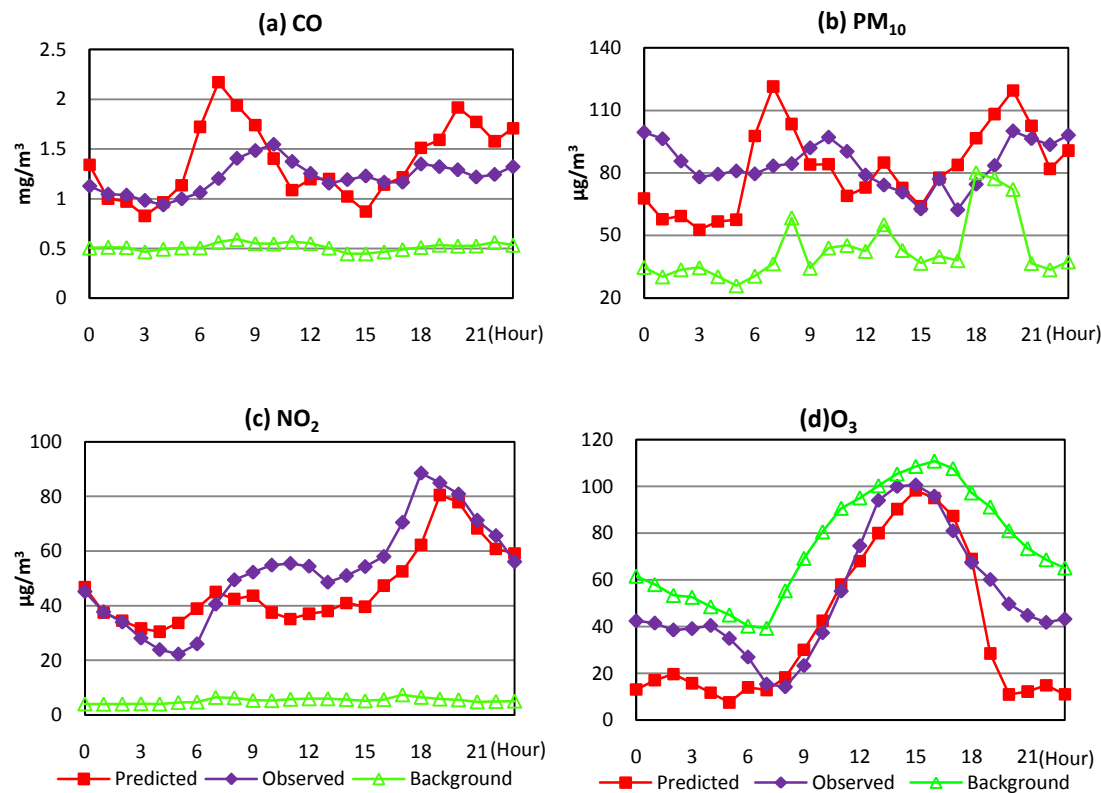


Fig. 4. Model performance on time-of-the-day, in comparison with the diurnal profiles of observed and background concentrations for (a) CO; (b) PM<sub>10</sub>; (c) NO<sub>2</sub> and (d) O<sub>3</sub>, for the periods of July 10<sup>th</sup>-20<sup>th</sup> and August 10<sup>th</sup>-20<sup>th</sup>, 2008.

### 3.1.3 Dependence of model performance on wind direction

Although wind speed has an important influence on the dispersion of pollutant and thus the model predictions of pollutant concentrations, we do not focus on the evaluation of the dependence of model performance on wind speed, because the fundamental factor influencing the diurnal profile of model predictions was the temporal variation of source intensity and subsequent atmospheric chemical reactions, rather than the accompanying wind speed. However, to evaluate the dependence of model performance on wind direction was important to know whether the meteorological station adopted in this study was representative of the wind field of the study domain, and to understand for the right reason the characteristics of spatial distribution of the traffic-related air pollution after the TRS was implemented. Thus, the pollution roses for all pollutants were plotted, as showed by Figure 5, which revealed that the peaks and troughs of the predicted and observed concentrations were virtually in the same directions. This proved that the model well predicted the variance of concentrations in response to the wind direction variation. Besides, Figure 5 also revealed that there was a clear dependence of concentrations on wind directions for all pollutants: high CO concentrations occurred mainly in the downwind southwesterly, northwesterly and southerly directions, with high PM<sub>10</sub> concentrations also occurring in the downwind southwesterly and northwesterly directions. In addition, NO<sub>2</sub> concentration had a weaker dependence on wind directions and distributed more uniformly in each direction, with the peak concentrations mainly

572 occurring in the southwesterly direction.  $O_3$  had a very clear dependence of wind  
 573 direction, with the peak concentrations appearing in the northwesterly and  
 574 southwesterly directions. Therefore, the model captured well the spatial variation of  
 575 the pollutant concentrations in response to the wind direction variation, which  
 576 revealed that the meteorological data from the Airport Meteorological Observatory  
 577 were well representative of the wind field of UAB, in consistence with another study  
 578 using the same meteorological data source for the evaluation of air quality impacts of  
 579 the Games (Wang et al., 2009a).

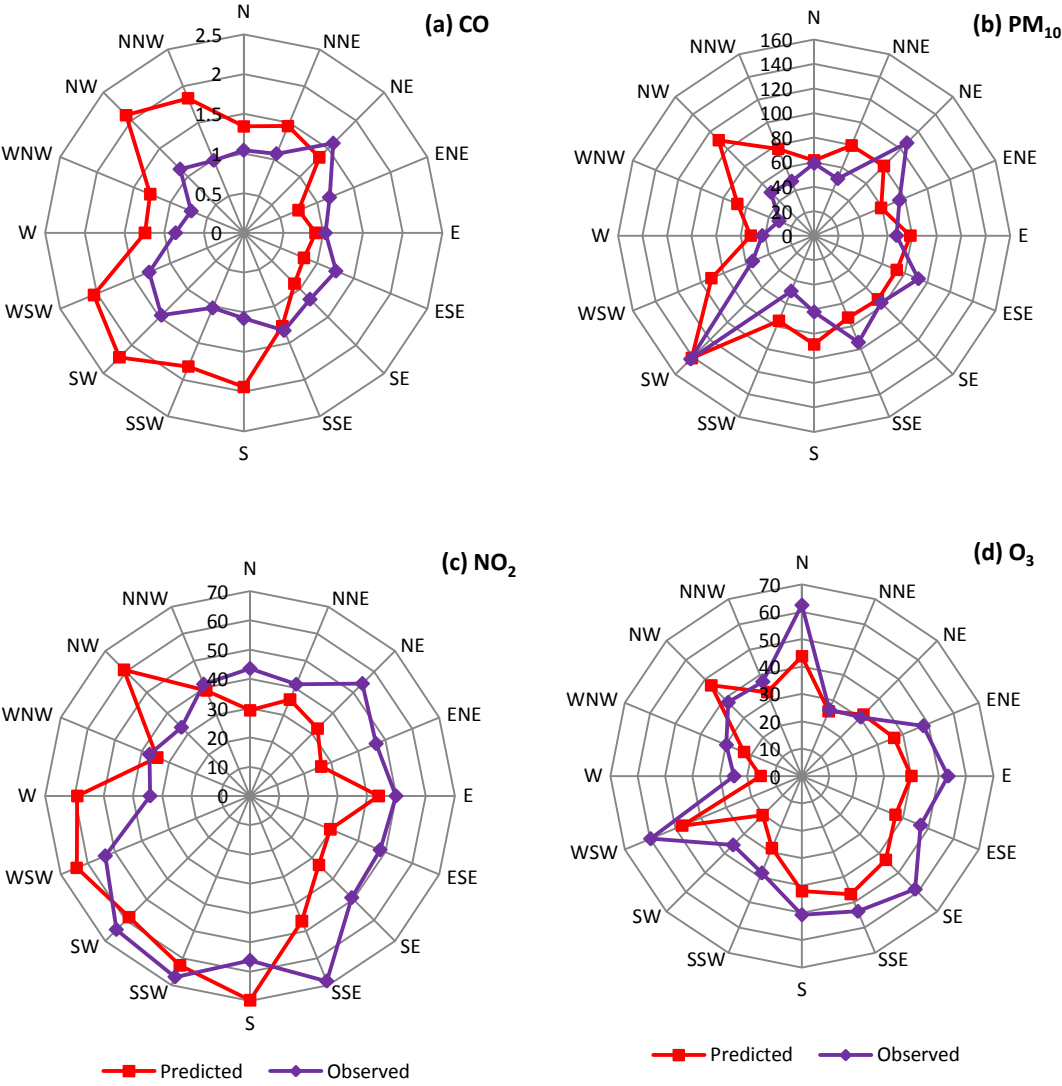


Fig. 5. Comparison of mean predicted and observed pollutant concentration roses for (a) CO, (b) PM<sub>10</sub>, (c) NO<sub>2</sub> and (d) O<sub>3</sub> in different wind directions for the periods of July 10<sup>th</sup>-20<sup>th</sup> and August 10<sup>th</sup>-20<sup>th</sup>, 2008, at the CGZ air quality monitoring site.

Only one traffic-representative monitoring station was included in the model evaluation, because the data from the other one, the Qianmen station set up at a high-traffic environment and maintained by the Beijing Municipal Environmental Monitoring Centre (BMEMC), were incomplete during the evaluation period, with about 70% of the data missing due to malfunction of the monitoring equipment. Besides, as industrial production were restricted, construction sites were shut down, and agricultural activities, particularly biomass burning in rural areas were strictly banned both in Beijing and other eight surrounding provinces (MEP, 2008), the major emissions in the UAB during the Games came from vehicles, and this emission characteristic during the Games was distinct from the usual situation with much more emissions from other sources, under which circumstances the previous modelling studies (Song et al., 2006b and Wang et al., 2008a) were conducted, using additional rural and industrial monitoring stations for the modelling evaluation. Therefore, the overall good model evaluation results in this study based on the CGZ monitoring station were not coincidental due to the station location, but was a true reflection of the particular characteristic of source emissions dominated by vehicles during the Games. Under such a circumstance, a city-level evaluation of the model performance was not conducted in this study, as the particular characteristic of sources emissions during the Games convinced us that other air quality monitoring stations, particularly those usually adopted to reflect air quality impacts from industrial sources or rural emissions, were unnecessary or not suitable to be included for the model evaluation in this study. Furthermore, since the following analysis of

the modelling results was focused on the predictions at the simulated roads and at the receptors, where the traffic-dominant emission characteristic was identified and the air quality responses have been evaluated, using the measurements from the representative traffic monitoring station (CGZ), it is reasonable to believe that the conclusions drawn based on the analysis were credible. Nevertheless, the possible drawback of using one traffic-representative station for model evaluation was that the model predictions in areas away from the simulated roads might have some unknown uncertainty.

### 3.2 Air quality impacts in response to TRS policy before, during and after the Games

Applying the evaluated ADMS model, we predicted the CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> concentrations from the RRs and LRs around UAB before, during and after the Olympic Games, with the aim of evaluating the air quality impacts of the TRS policy. First, the emission reduction in accordance with the decrease of traffic flows was reported; Second, the daily and diurnal variation of pollutant concentrations along the RRs and LRs were compared for the pre-, during- and post-TRS periods, followed by a further assessment of the weekly variation of air quality in response to the TRS policy; Finally, the spatial variation of the traffic-related air pollution for the pre-, during- and post-TRS periods was assessed, to understand the regional differences in air quality impacts of the TRS policy.

#### 3.2.1 Reduction of diurnal average traffic flow and pollutant emissions

The traffic flows on the RRs and LRs had a notable reduction after the TRS policy was implemented on 20<sup>th</sup> July, 2008. The daily average traffic flows of the short and



long motor vehicles for the during-TRS period had a reduction of 26.1% and 11.0%,  
 respectively, compared to the pre-TRS period. Figure 6, which shows the variation  
 of traffic flows on RRs and LRs on daily average and the variation of the diurnal  
 profiles of the total traffic flow on hourly average for the pre-TRS, during-TRS and  
 post-TRS periods, revealed that the TRS policy had a substantial effect on the  
 reduction of both daily average and hourly average traffic flows, with average  
 reduction rates of 22.4%, 26.1%, 27.7% and 27.6% on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs,  
 respectively, and a notably overall reduction of about 26.1% for the total traffic flow.  
 Particularly, the TRS policy had a largest traffic flow reduction of 30.8% during the  
 morning rush hours (7-8 am). On the other hand, both the daily average and hourly  
 average traffic flows on all types of roads during the post-TRS period virtually  
 bounced to the same level as the pre-TRS period, with the double-peak pattern of the  
 diurnal profile of traffic flows in Beijing unchanged by the TRS policy.

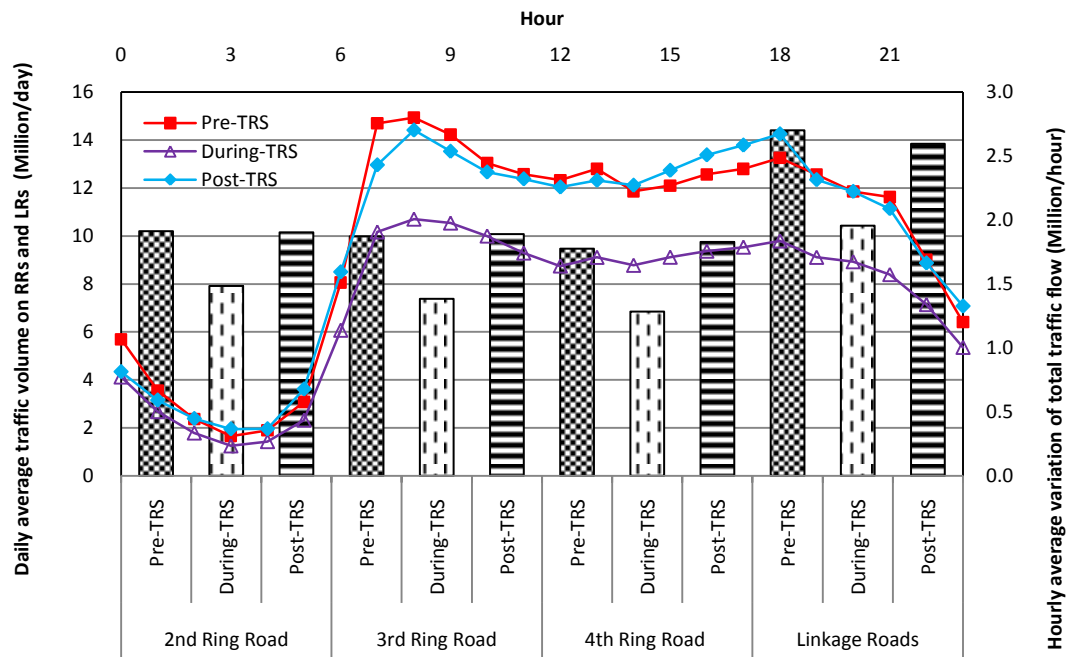


Fig.6. Comparison of traffic flows on 2<sup>nd</sup> RR, 3<sup>rd</sup> RR, 4<sup>th</sup> RR and the LRs on daily average (histograms), and diurnal variation of total traffic flow on hourly average (scatter plots) for the pre-TRS, during-TRS and post-TRS periods, respectively.

With the reduction of traffic flows, the traffic conditions were improved and the traffic running speeds on hourly average increased by about 15% for the during-TRS period, as revealed by the ITS-TAP monitoring network. The raised speed decreased the emission factors of CO, PM<sub>10</sub> and NO<sub>x</sub> for all vehicle types, as calculated by COPERT model. In addition to the reductions of traffic flows and emission factors, the emissions on daily average decreased by about 26.1%, 25.4% and 25.2% for CO, PM<sub>10</sub> and NO<sub>x</sub>, respectively. Consequently, emission rates on the RRs and LRs decreased as well, especially for the 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs.

### 3.2.2 Temporal variation of air quality impacts of the TRS policy

Pollutant concentration levels are directly related to air quality, and thus we focused on the pollutant concentration impacts of the TRS policy to assess its effectiveness on air quality improvement in Beijing during the Games. Concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> in the following results and discussion were predicted at a height of 1.5 meters above the ground.

#### 3.2.2.1 Changes in daily average concentrations for the pre-, during- and post-TRS periods

The daily average concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> had a notable reduction on the RRs and LRs around UAB after the TRS policy was implemented, as shown by Figure 7, which illustrates that the daily average predictions of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub>

665 on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs for the pre-, during- and post-TRS periods. Daily  
 666 average CO concentrations were about 2.1, 2.1, 2.2 and 2.3 mg/m<sup>3</sup> on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup>  
 667 RR and LRs, respectively, for the pre-TRS period, which decreased by about 35.8%,  
 668 38.5%, 34.9% and 35.6% on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and LRs, respectively, for the  
 669 during-TRS period. Consequently, the CO daily concentrations on all road types  
 670 satisfied the CNAAQs Grade II of 4 mg/m<sup>3</sup> for the during-TRS period. The  
 671 pre-TRS daily average concentrations of PM<sub>10</sub> were 120.9, 115.7, 136.1 and 163.2  
 672 µg/m<sup>3</sup> on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs, respectively, which was successfully  
 673 reduced below 150 µg/m<sup>3</sup> by the TRS policy, as shown by Figure 7. For the  
 674 pre-TRS period, 40% of days on the LRs exceeded the daily CNAAQs Grade II of  
 675 150 µg/m<sup>3</sup>, while the PM<sub>10</sub> daily concentrations on the LRs were successfully reduced  
 676 by 34.7%, to satisfy the Grade II limit, with the PM<sub>10</sub> concentrations on the 2<sup>nd</sup>, 3<sup>rd</sup>  
 677 and 4<sup>th</sup> RR reduced by 38.7%, 31.8% and 44.0%, respectively, resulting in satisfaction  
 678 of the Grade II limit for every single day for the during-TRS period. The reduction  
 679 of vehicular emissions of PM<sub>10</sub> due to the TRS policy, which was the major cause for  
 680 PM<sub>10</sub> emission reduction during the Games, as control measures for sources other  
 681 than on-road vehicles remained unchanged and had little extra effect on emission  
 682 reduction, resulted in this reasonably notable effect on the reduction of PM<sub>10</sub>  
 683 concentration, which the previous source apportionment studies would not expect.  
 684 With the bounce of traffic flows during the post-TRS period, concentrations of both  
 685 CO and PM<sub>10</sub> increased notably, with 20% of days on the LRs exceeds the Grade II

limit for  $\text{PM}_{10}$ . The  $\text{NO}_2$  concentrations were reduced by about 30.3%, 31.9%, 32.3% and 33.9% on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs, respectively, to about 52.3, 46.2, 45.9 and 47.0  $\mu\text{g}/\text{m}^3$  during the TRS period, with the daily concentration on every single day during this period conforming to the CNAAQs Grade II of 80  $\mu\text{g}/\text{m}^3$ . Only 10% of days on the 2<sup>nd</sup> RR exceeded the  $\text{NO}_2$  Grade II limit for the post-TRS period, despite a concentration increase of about 9.0%, 13.7%, 17.3% and 18.8% on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs, respectively, during the period.  $\text{O}_3$ , of which the concentration level is usually lower in the source-intensive urban areas than in the rural areas, had a concentration of about 38-52  $\mu\text{g}/\text{m}^3$  in the urban areas during the pre-TRS period, and the  $\text{O}_3$  concentration decreased to about 24-29  $\mu\text{g}/\text{m}^3$  during the TRS period, which was consistent with the findings of Wang et al. (2009b). Unlike other pollutants, the  $\text{O}_3$  concentration continued decreasing for the post-TRS period, to about 23-26  $\mu\text{g}/\text{m}^3$  despite the increase of traffic flow and vehicular emissions. This is mainly because  $\text{O}_3$  concentrations in UAB are produced by local photochemical reactions under VOC-limited conditions (Wang and Li, 2002; Wang et al., 2009c), and higher emission level of NO during the post-TRS period consumed  $\text{O}_3$  and suppressed the accumulation of  $\text{O}_3$  in urban environment of Beijing (named the “titration effect”) (Chou et al., 2006; Sadanaga et al., 2008). Besides, the elevated nitrogen dioxide ( $\text{NO}_2$ ) through the titration effect and reactions of NO with other radicals further reacted with the OH radical and yielded nitric acid ( $\text{HNO}_3$ ), which ended the photochemical reaction chain involved with the OH radical that otherwise

accumulates the ambient  $O_3$  concentration (NARSTO, 2000). Therefore, it was clear that the TRS policy was effective in reducing daily average concentrations of pollutants, ensuring that the air quality in Beijing during the Olympic Games conformed to the 24 hours CNAQS Grade II limits.

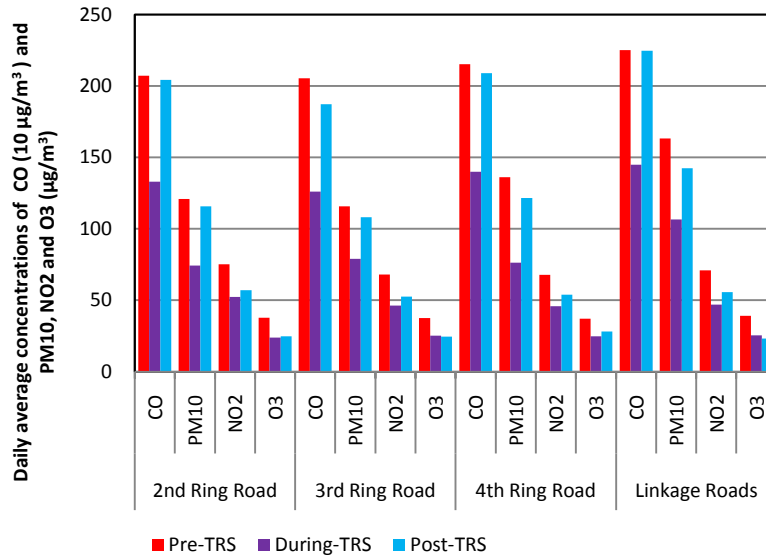


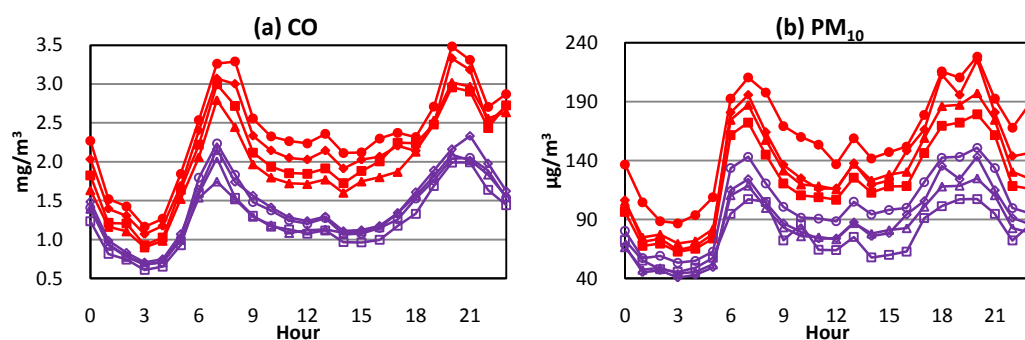
Fig. 7. Comparison of daily average predicted concentrations of CO,  $PM_{10}$ ,  $NO_2$  and  $O_3$  on different road types for the pre-TRS, during-TRS and post-TRS periods.

The predicted highest daily average concentrations of the CO,  $PM_{10}$ ,  $NO_2$  and  $O_3$ , which occurred at a very limited area under certain circumstances, also showed a notable decrease during the TRS period, which was in consistence with the previous study proving the effectiveness in reducing extreme concentrations of a four-day traffic control experiment by the Beijing Municipal Government (Westerdahl et al., 2009). The predicted highest concentrations had decreased from 6.3 to 4.6  $mg/m^3$  for CO, from 635.7 to 498.3  $\mu g/m^3$  for  $PM_{10}$ , from 194.7 to 154.2  $\mu g/m^3$  for  $NO_2$  and from 88.2 to 53.3  $\mu g/m^3$  for  $O_3$ , a reduction of about 27.6%, 21.6%, 20.8% and 39.6%

for CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub>, respectively, in comparison with the pre-TRS period.

### 3.2.2.2 Diurnal variation of hourly average concentration

To further evaluate the effectiveness of the TRS policy on air quality improvement, we focused on the variation of diurnal profiles of hourly average concentration on the RRs and LRs in response to the TRS policy. Figure 8 shows that hourly average concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> on the RRs and the LRs decreased notably, in comparison with the pre-TRS period. The morning and evening peak concentrations of CO on the LRs had decreased from about 3.29 mg/m<sup>3</sup> to about 1.83 mg/m<sup>3</sup>, the largest reduction of 34.9% and 38.0% among all road types, as shown by Figure 8 (a), while the largest reduction of the morning and evening peak concentrations of PM<sub>10</sub> was observed on the 2<sup>nd</sup> RR, accounting for about 37.8% and 40.2%, respectively. Consequently, the hourly CO concentrations were well below the 1-hour CNAQS Grade II of 10 mg/m<sup>3</sup> and the hourly PM<sub>10</sub> concentrations decreased below 150 µg/m<sup>3</sup> on all types of roads. Particularly, the average of evening peak CO, PM<sub>10</sub> and NO<sub>2</sub> concentrations were reduced significantly, by about 36.6%, 36.1% and 32.8%, respectively, which was a solid proof for the air quality improvement during the TRS policy period.



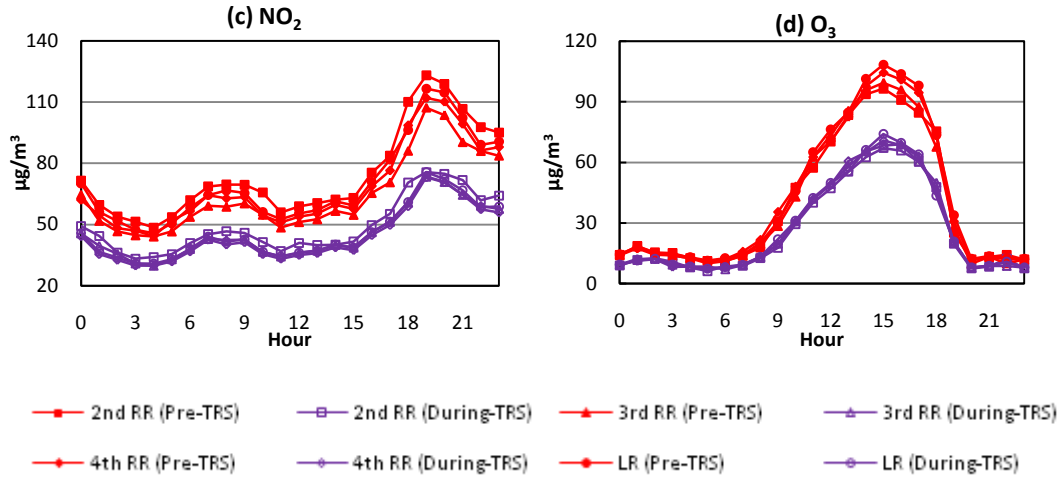


Fig. 8. Comparison of diurnal predicted variations of hourly concentrations of (a) CO; (b)  $\text{PM}_{10}$ ; (c)  $\text{NO}_2$  and (d)  $\text{O}_3$  on 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs for the pre- and during- TRS periods.

For both pre- and during-TRS periods, the diurnal profile of CO and  $\text{PM}_{10}$  took on an obvious double-peak mode, as shown by Figure 8 (a) and (b), with the morning and evening peak concentrations occurring at around 7-8 am and around 8-9 pm, respectively, and the lowest concentrations occurring at around 3 am. While the occurring time of the morning peak concentration was consistent with the morning rush hour of traffic flows, the evening peaks for CO and  $\text{PM}_{10}$  were about two to three hours later than the evening rush hour. Moreover, the evening peak concentrations of CO and  $\text{PM}_{10}$  during the pre-TRS period even exceeded the morning peak, which was in contradiction with the diurnal profile of traffic flows during the period. These delayed and higher evening peak concentrations should be ascribed to the particular evening meteorological conditions in UAB: urban heat island (UHI) frequently occurs in Beijing in the summer night, which tends to produce strong

temperature inversion and higher inversion layer at night. When the UHI occurred, the UHI convergence and the strong temperature inversion at night hindered air dispersion and contributed to the local accumulation of pollutants, resulting in high pollution concentrations at night (Miao et al., 2008; Liu et al., 2006).

The NO<sub>2</sub> concentrations on all roads for the pre-TRS periods, as shown by Figure 8 (c), began to increase in the early morning, reached a peak in about 7-8 am, and came to a trough during the mid-day time, which was related to the traffic and emission rate cycles and similar diurnal variation of NO<sub>2</sub> concentrations were found in the urban area of Cairo (Khoder, 2009). The maximum concentration of NO<sub>2</sub> occurred at about 7-8 pm, which was higher in magnitude than the morning peak. This was partly due to the evening rush hour of traffic with high emission rate of NO<sub>x</sub>, and partly due to the same UHI effect weakening the dispersion conditions and resulting in evening peak concentrations of CO and PM<sub>10</sub>. Moreover, the lower temperature and solar radiation intensity in the evening decreased the photodissociation of NO<sub>2</sub>, a major chemical mechanism for NO<sub>2</sub> loss during the daytime. The TRS policy, as shown by Figure 8(c), did not alter the diurnal variation characteristic of NO<sub>2</sub> concentrations, but significantly reduced the hourly sequential concentrations of NO<sub>2</sub> during the Games. Particularly, the highest hourly average concentrations of 123.1, 107.2, 111.9 and 116.6 µg/m<sup>3</sup> on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs, respectively, which occurred at about 7 pm during the pre-TRS period, were significantly reduced by as much as 38.6%, 31.5%, 34.7% and 32.5%,



778 respectively, to 77.5, 73.4, 73.1 and 75.7  $\mu\text{g}/\text{m}^3$ , respectively, during the TRS period,  
779 which was in good agreement with the tropospheric  $\text{NO}_2$  observations by NASA's  
780 OMI revealing a 30-50% reduction of  $\text{NO}_2$  during the Games (NASA, 2009).  
781 Therefore, the TRS policy was effective in reducing the hourly  $\text{NO}_2$  concentrations,  
782 and consequently the hourly concentrations on all types of roads were far below the  
783 1-hour CNAAQs Grade II of 120  $\mu\text{g}/\text{m}^3$  during the TRS period. As shown by  
784 Figure 8 (d), the  $\text{O}_3$  concentrations on all roads had a similar diurnal profile for both  
785 pre- and during-TRS periods, with the concentrations increasing in the daytime,  
786 reaching the peak at about 3 pm and beginning to decrease rapidly. It was clear that  
787 the TRS policy reduced remarkably the hourly  $\text{O}_3$  concentrations on all types of roads,  
788 and had a notable concentration reduction of the afternoon peak, with the  
789 concentrations on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs decreasing by about 35.4%, 30.7%,  
790 32.2% and 31.8%, respectively. As a result, the hourly  $\text{O}_3$  concentration during the  
791 TRS period was far below the 1-hour CNAAQs Grade II of 160  $\mu\text{g}/\text{m}^3$ . Meanwhile,  
792 it was interesting to find that the diurnal variation of the concentration ratio of  $\text{NO}_2$  to  
793  $\text{O}_3$  ( $[\text{NO}_2]/[\text{O}_3]$ ) for all types of roads was opposite to the diurnal profile of  $\text{O}_3$   
794 concentrations, which revealed that the traffic-related  $\text{NO}_x$  (mostly  $\text{NO}$ ) consumed  $\text{O}_3$ ,  
795 and resulted in an increase of the secondary  $\text{NO}_2$  concentration and a decrease of the  
796  $\text{O}_3$  concentration, a typical VOC-limited characteristic for ozone formation. With  
797 the significant reduction of both primary and secondary pollutant concentration levels,  
798 air quality in UAB had therefore been improved remarkably during the Games in

response to the TRS policy.

### 3.2.2.3 Weekly variation of diurnal profile of pollutant concentrations for the pre-, during- and post-TRS periods

To investigate the day-of-the-week variation of air quality impact in response to the TRS policy, we plotted the diurnal profiles of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> for weekdays, Saturdays and Sundays for the separate periods before, during and after the TRS policy. As shown by Figure 9 (a) and (b), the diurnal profiles for CO and PM<sub>10</sub> remained similar, with a typical two-peak pattern, and had no obvious weekly variation in response to the TRS policy, as the occurring time of the peaks and trough were consistent for weekdays, Saturdays and Sundays for the three separate periods, due to the smooth weekly cycle of diurnal variation of traffic flows. Notably lower levels of CO and PM<sub>10</sub> concentrations on Saturdays and Sundays than weekdays, however, were observed, particularly during the rush hours when peak concentrations occurred, for the pre-, during- and post-TRS periods. Therefore, it was clear that the traffic-related air pollution still presented a typical two-peak diurnal variation pattern, but the weekly cycle had no obvious variation in response to the TRS policy, which was, nevertheless, effective in reducing the hourly concentrations, particularly the peak concentrations of CO and PM<sub>10</sub>.

The day-of-the-week variation of NO<sub>2</sub> diurnal profile, as shown by Figure 9 (c), revealed a similarity in the diurnal variation on weekdays, Saturdays and Sundays, particularly, in the ascending trend of NO<sub>2</sub> concentrations in the daytime and reaching

820 a peak in the evening, despite a notable decrease in the afternoon, for the pre-, during-  
821 and post-TRS periods. Besides, lower concentration levels on Saturdays and  
822 Sundays were observed compared to that on weekdays, which revealed that the TRS  
823 policy did not change the weekly cycle of traffic flows, with heavier traffic load on  
824 weekdays than on the weekends. Meanwhile, Figure 9 (d) revealed a notable  
825 difference between the diurnal profile of  $O_3$  and other pollutants, with relatively  
826 higher concentration levels on Saturdays and Sundays than that on weekdays for the  
827 pre- and post-TRS periods, which was in agreement with the characteristic of the  
828 ozone weekend effect that has been frequently observed in urban areas  
829 (Atkinson-Palombo et al., 2006; Gao and Niemeier, 2007; Murphy et al., 2007; Tang  
830 et al., 2008; Khoder, 2009). The major reason for the observed ozone weekend  
831 effect was the “VOC-limited” feature for ozone formation in UAB: the ozone  
832 concentration tended to increase with the decrease of  $NO_x$  emissions, typically due to  
833 the reduction of traffic flows, as a result of the weakened titration effect of lower  $NO_x$   
834 emissions on the weekends that tended to accumulate ozone concentrations. This  
835 mechanism also explained why a more remarkable ozone weekend effect was  
836 observed for the post-TRS period, as shown by Figure 9 (d): the peak  $O_3$   
837 concentration at around 3 pm on weekdays for the post-TRS period continued to  
838 decrease, compared to the during-TRS period, due to the enhanced titration effect of  
839 increased  $NO_x$  emissions that suppressed the ozone formation on weekdays, while the  
840 peak concentrations on Saturdays and Sundays increased, due to a larger reduction of

841 NO<sub>x</sub> emissions and thus a more weakened titration effect on the weekends during the  
842 post-TRS period, compared to the during-TRS period. Moreover, Figure 9 (d) also  
843 revealed that the TRS policy effectively reduced both the hourly peak and daily  
844 average concentrations of ozone, and meanwhile reduced and virtually removed the  
845 weekend effect of ozone for the during-TRS period, probably due to the smoother  
846 weekly variation of traffic flows resulting from the restriction on traffic.

847

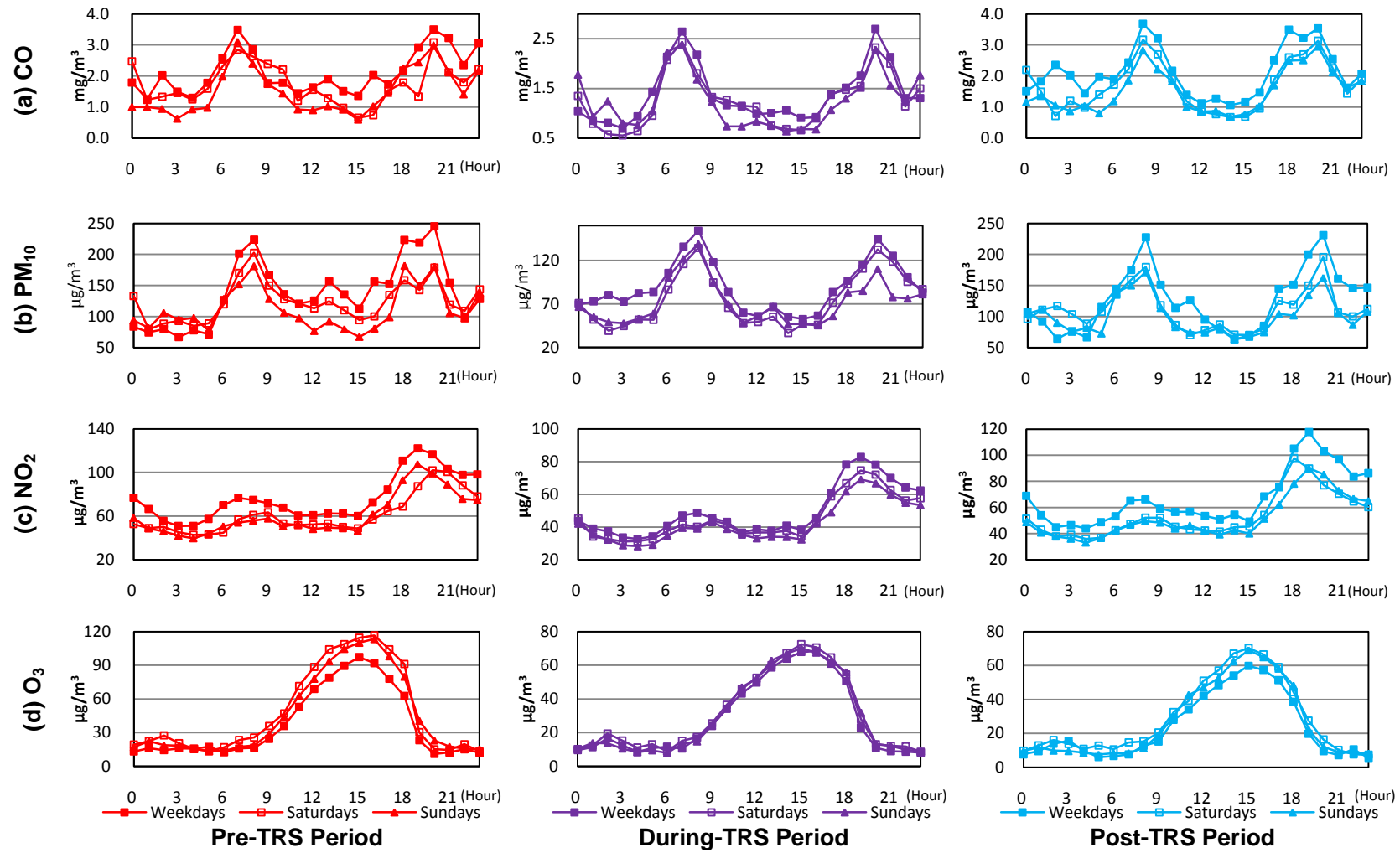


Fig. 9. Day-of-the-week variations of (a) CO; (b) PM<sub>10</sub>; (c) NO<sub>2</sub> and (d) O<sub>3</sub> diurnal profiles in response to the implementation of the TRS policy.

### 3.2.3 Spatial variation of traffic-related air pollution and implication on regional air quality improvement for the pre-, during- and post-TRS periods

To investigate the regional air quality impacts of the TRS policy, we looked into the spatial distribution of the traffic-related air pollution around UAB, as well as the spatial variation of regional air quality improvement in response to the TRS policy.

Figure 10, which illustrates the contour plot of the daily average concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> from on-road motor vehicles on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs, for the pre-, during- and post-TRS periods, respectively, revealed remarkable differences in the regional distribution of traffic-related air pollutant concentrations and notable variation of regional air quality in response to the TRS policy. Before the TRS policy was placed, the highest and lowest CO concentration levels occurred in the upwind eastern regions and the southern areas, respectively, as shown by Figure 10 (a), with the daily average concentrations of the representative receptors located in the downwind northwest and upwind southeast accounting for about 2.56 and 2.02 mg/m<sup>3</sup>, respectively. Particularly, the West 2<sup>nd</sup> RR and the Northwestern LRs between 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> RR had the heaviest traffic and accordingly suffered the highest CO concentration level of above 2.4 mg/m<sup>3</sup>. With the TRS policy, a significant reduction of the CO daily average concentration was achieved, and the previously highest concentration in the north was reduced by 31.2%, to 1.79 mg/m<sup>3</sup>, with the concentration levels in the south, west and east regions decreasing by about 17.6%, 18.2% and 17.6%, respectively. Besides, CO concentrations in a broad area

868 of the northeast, east and southeast outside the 4<sup>th</sup> RR decreased below a level of 0.5  
869 mg/m<sup>3</sup>, mainly due to the prevailing northeast, east and southeast wind that  
870 transported the pollutant to the downwind directions and resulted in a relatively  
871 lower concentrations in these upwind areas during the period. Moreover, the  
872 previously seriously polluted areas in the West 2<sup>nd</sup> RR and the Northwestern LRs  
873 were relieved, with a much smaller area where the concentration level remained  
874 above 2 mg/m<sup>3</sup>. Therefore, the TRS policy was effective in bringing down the  
875 concentration levels in both the most polluted areas and different parts of the urban  
876 area, resulting in a much better air quality during the Games. With the bouncing of  
877 traffic flows almost to the pre-TRS level, the CO concentration increased  
878 dramatically in every part of UAB, and consequently, almost caught up with the  
879 previous level before the TRS policy was placed, with the concentrations in the east,  
880 south, west and north increasing by about 24.2%, 16.0%, 15.2% and 22.2%,  
881 respectively.

882 The spatial distribution of PM<sub>10</sub> concentrations was characterized as linear  
883 distribution of relatively higher pollution along the RR and the LRs, with  
884 concentration levels in the areas between the simulated roads and outside the 4<sup>th</sup> RR  
885 much lower. For the during-TRS period, the PM<sub>10</sub> concentration levels decreased  
886 remarkably for the whole UAB, with the PM<sub>10</sub> concentration in most of the area  
887 reduced below 100 µg/m<sup>3</sup>. Particularly, the previously highly polluted areas in the  
888 West 2<sup>nd</sup> RR, Southwestern 4<sup>th</sup> RR and the Northern LRs with PM<sub>10</sub> concentrations

above  $150 \mu\text{g}/\text{m}^3$  before the TRS shrank significantly during the TRS, as shown by Figure 10 (b). Consequently, the concentrations in the east, south, west and north regions of UAB decreased from about 118.2, 111.9, 160.4 and  $217.7 \mu\text{g}/\text{m}^3$ , respectively, for the pre-TRS period, to about 66.1, 67.6, 105.6 and  $155.2 \mu\text{g}/\text{m}^3$ , respectively, for the during-TRS period. The dramatically decrease of the primary pollutant concentration indicated that the air quality during the Games was improved significant in response to the TRS policy. However, the  $\text{PM}_{10}$  concentration levels increased by about 23.4%, 33.6%, 20.7% and 29.5%, respectively, in the east, south, west and north regions for the post-TRS period. As a result, the areas with  $\text{PM}_{10}$  concentration above  $150 \mu\text{g}/\text{m}^3$  expanded, which proved reversely the effectiveness of the TRS policy on air quality improvement.

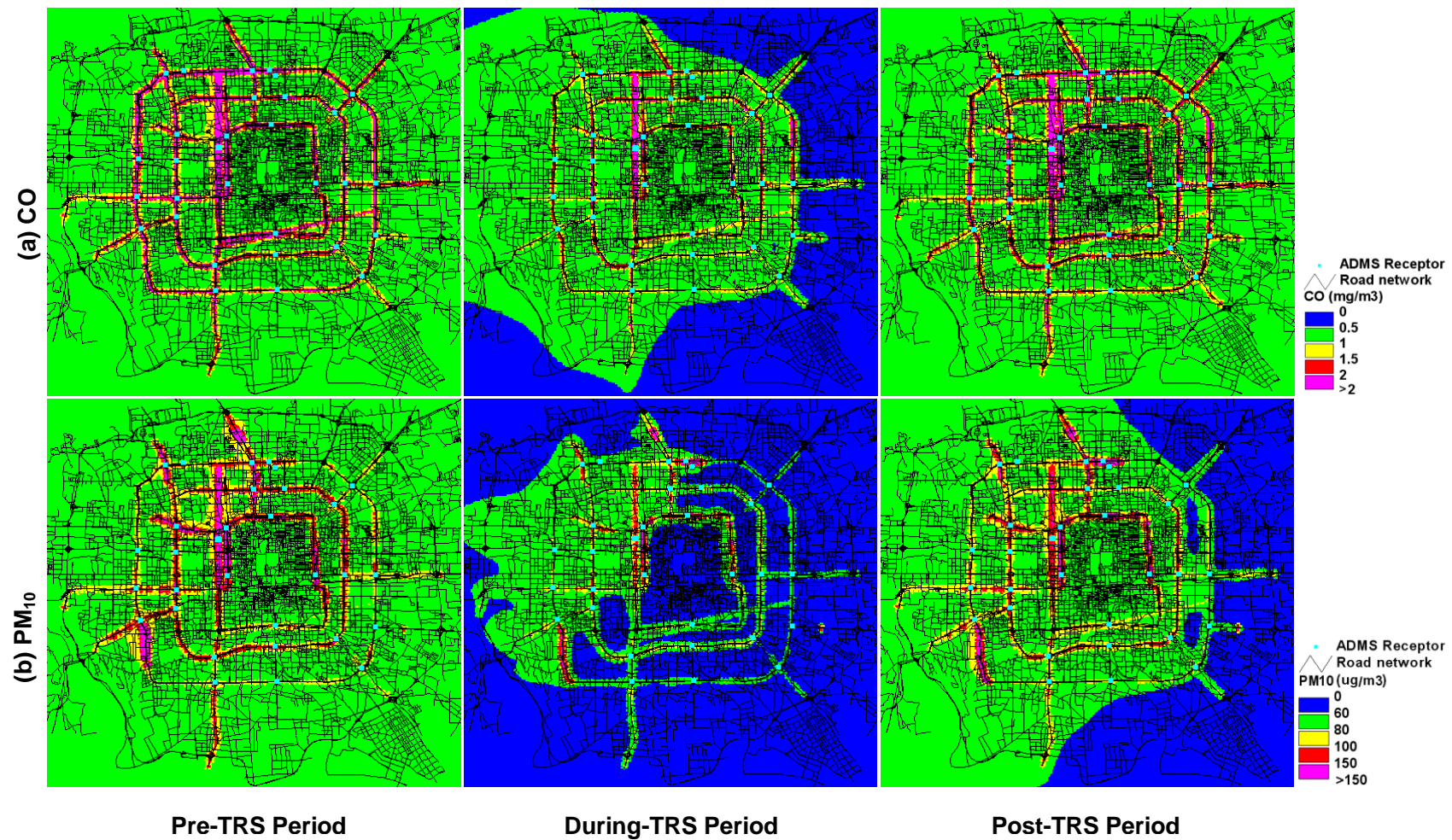
The spatial distribution of  $\text{NO}_2$  concentration was more influenced by the prevailing northeasterly, easterly and southeasterly winds during the pre-TRS period, with the relatively higher daily average concentration of about 76.8 and  $68.0 \mu\text{g}/\text{m}^3$  predicted in the downwind western areas and the northern traffic-heavy areas, respectively. Besides, the daily average  $\text{NO}_2$  concentration along the West 2<sup>nd</sup> RR, the Northern LRs and part of the West 4<sup>th</sup> RR exceeded the CNAAQs Grade II of  $80 \mu\text{g}/\text{m}^3$ . On the contrast, the  $\text{NO}_2$  concentration decreased significantly in response to the TRS period, with the concentration level in most of UAB below  $80 \mu\text{g}/\text{m}^3$ , which was a clear evidence of the effective control of traffic-related air pollution and corresponding remarkable air quality improvement by the TRS policy. Particularly,



better effect on pollution reduction was observed in the previously most severely polluted areas, as the concentration level in these areas were brought down sufficiently to comply with the CNAAQs limited values. With the expiration of the TRS policy, NO<sub>2</sub> pollution with concentration above 80 and even 120 µg/m<sup>3</sup> appeared in a much larger area for the post-TRS period, and NO<sub>2</sub> concentration increased significantly in the downwind western areas, in comparison with the during-TRS period, as shown by Figure 10 (c). The spatial distribution variation of NO<sub>2</sub> concentration in UAB revealed a notable air quality improvement due to reduced on-road vehicular emissions.

The spatial distribution of O<sub>3</sub> concentration was distinct from those of CO, PM<sub>10</sub> and NO<sub>2</sub>, as shown by Figure 10 (d). For the pre-, during- and post-TRS periods, notably lower O<sub>3</sub> concentrations were observed along the RR and the LRs as these areas were the emission sources and O<sub>3</sub>, as a typical secondary air pollutant, had its higher concentration levels in the relatively remote areas off the road sources. Consequently, the O<sub>3</sub> daily average concentration in the near-road areas was well below 60 µg/m<sup>3</sup> for the pre-TRS period and relatively higher concentration levels of 60-80 µg/m<sup>3</sup> and above 80 µg/m<sup>3</sup> covered a large area around and outside UAB where no direct vehicular emissions were generated. Particularly, the severely polluted West 2<sup>nd</sup> RR, Northern LRs and part of the South 4<sup>th</sup> RR areas with high concentration levels of CO, PM<sub>10</sub> and NO<sub>2</sub>, however, had the lowest O<sub>3</sub> concentrations. In response to the TRS policy, the O<sub>3</sub> concentrations in the East,

931 South, West and North regions of UAB decreased significantly from 43.3, 46.6, 36.4  
932 and 43.9  $\mu\text{g}/\text{m}^3$ , respectively, to about 30.7, 31.2, 23.7 and 24.4  $\mu\text{g}/\text{m}^3$ , respectively,  
933 mostly due to the decrease of the precursor emissions of  $\text{NO}_x$  and VOC from on-road  
934 vehicles. Moreover, the  $\text{O}_3$  concentration during the post-TRS period continued to  
935 decrease, despite the increase of vehicular emissions of  $\text{NO}_x$  and VOC. As has  
936 been discussed previously, this phenomenon was mostly due to the “VOC-limited”  
937 characteristic for ozone formation in UAB. Therefore, the spatial variation of  $\text{O}_3$   
938 concentrations in response to the TRS policy also provided solid evidence of the  
939 effectiveness of such a restriction policy on air quality improvement during the  
940 Games.



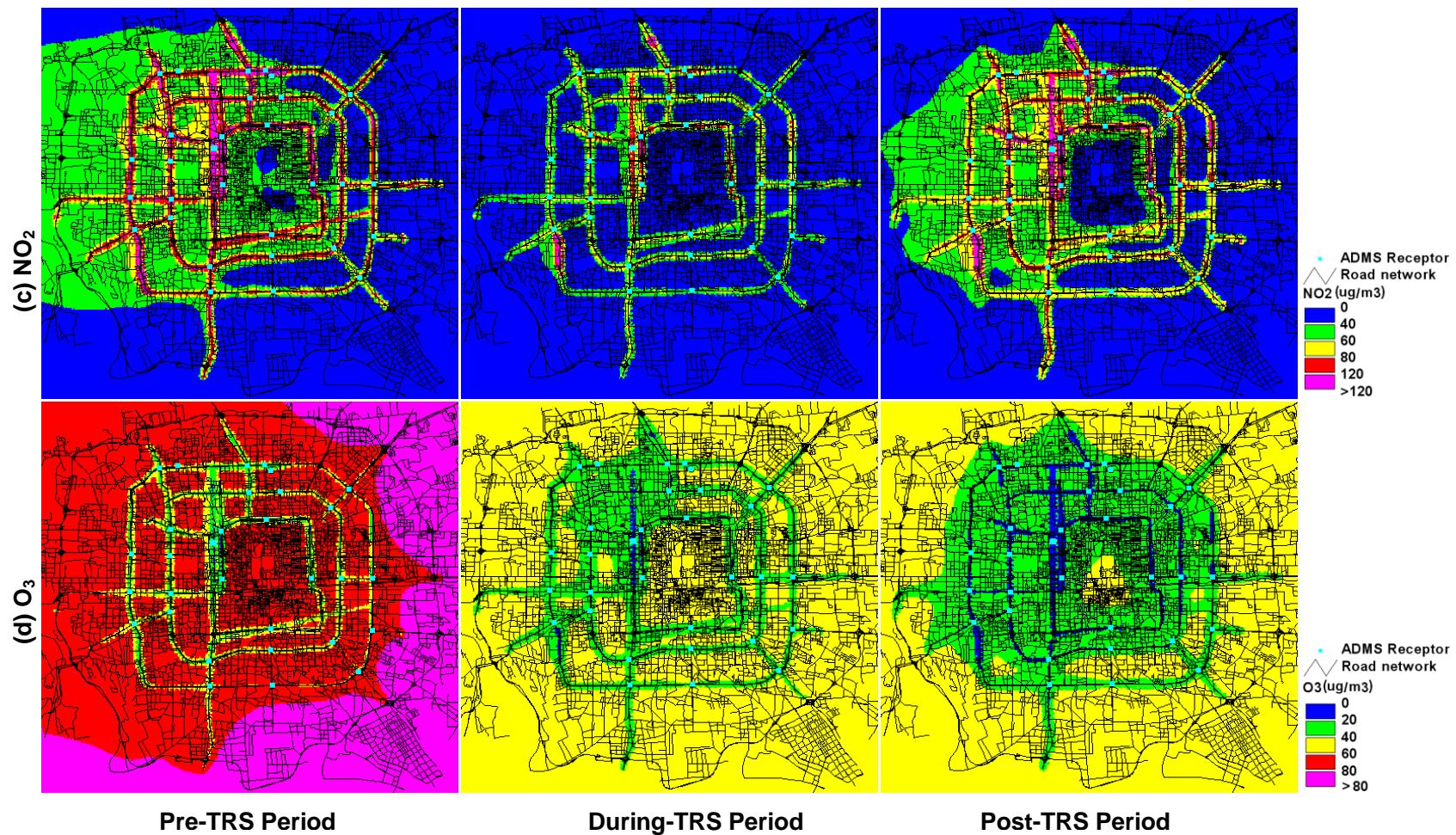


Fig. 10. Spatial distribution of (a) CO; (b) PM<sub>10</sub>; (c) NO<sub>2</sub> and (d) O<sub>3</sub> concentrations on daily average for pre-, during- and post-TRS periods, and the spatial variation of regional air quality impacts in response to the TRS policy.

### 3.3 Contribution of traffic flow reduction, raise of running speed and meteorological conditions to air quality improvements during the TRS period

It is meaningful to understand deeply the reasons that resulted in the air quality improvement after the TRS policy was implemented, to make sure that the model performed well for the right reasons, and to learn and inherit the useful experiences for the future air quality management in Beijing and other cities of China. Therefore, we focused on a discussion about the air quality impacts of traffic flow reduction, raise of running speed, variation of emission factors of pollutants, as well as the meteorological condition variation during the Games.

The TRS policy had a direct impact on the traffic flow and the running speeds of traffic fleet. As revealed by the ITS-TAP system, the daily average traffic flows of short and long vehicles during the TRS period decreased by 26.1% and 11.0%, respectively, an average reduction of about 24.2% for the total traffic fleet, in comparison with the pre-TRS period. Besides, running speed on the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs increased by 18.5%, 12.8%, 15.3% and 13.8%, respectively, an average 15% increase in driving speeds of total traffic fleet, estimated based on the fractions of traffic flows on these roads after the TRS policy took effect, which had a positive effect on reducing emission factors of pollutants. Figure 11 (a) illustrates the decrease rates of model-calculated emission factors of CO, PM<sub>10</sub> and NO<sub>x</sub> from various vehicle types resulting from the raise of running speed. The PC and LDV, which were mostly gasoline vehicles, had the largest reduction of CO emission factor,

962 accounting for about 10%. The largest reductions of both NO<sub>x</sub> and PM<sub>10</sub> emission  
963 factors were ascribed to diesel HDV, of which the traffic flow accounted for only 1-2%  
964 of the total fleet. Therefore, the reduction of NO<sub>x</sub> emission factor mainly relied on  
965 the other vehicle types, and an overall 3% reduction in NO<sub>x</sub> emission factor was  
966 achieved. The PM<sub>10</sub> emission factors were assumed constant for gasoline PC and  
967 LDV in response to the running speed increase, as there were no domestic studies  
968 available reporting the variation of PM emission factors of gasoline vehicles induced  
969 by speed raise, like from 20 km/h to 23 km/h, and a comprehensive sensitivity  
970 evaluation of the USEPA's MOBILE 6.2 model for PM emission factor found that a  
971 speed increase from 20 km/h to 23 km/h had very little impact on PM emission factor  
972 and concluded that speed has a negligible effect on PM emission factors for gasoline  
973 vehicles, which was unlike the emission factors for CO NO<sub>x</sub> and VOC that were  
974 highly sensitive to speed (Granell et al., 2004), while the HDV and buses had a  
975 reduction of about 10% of PM<sub>10</sub> emission factor. Consequently, the emissions of CO,  
976 PM<sub>10</sub> and NO<sub>x</sub> were reduced by about 26.1%, 25.4% and 25.2%, respectively, and the  
977 daily average concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> decreased by about 25.5%,  
978 35.4%, 32.1% and 34.4%, respectively, in response to the TRS policy. Therefore,  
979 the TRS policy was effective in bringing down traffic-related emissions, by a  
980 dominant contribution of traffic flow reduction, and by an extra bonus of decreasing  
981 emission factors due to the improvement of driving conditions. Taking into account  
982 the variation of wind speed, a major meteorological factor that affected the

traffic-related air pollution, the daily average wind speeds were about 2.52 and 2.53 m/s for the pre- and during-TRS periods, respectively, and had no statistically significant difference ( $p < 0.05$ ). Besides, the diurnal wind speed profile on the hourly average had little variation, particularly in the morning and for the evening rush hours (7-8 pm), as shown by Figure 11 (b). Under the virtually constant wind speed, the TRS policy turned out an effective measure in simultaneously bringing down traffic flow, increasing traffic fluency, and achieving air quality improvement with significantly decreased concentration levels. Therefore, the TRS policy was effective in reducing short-term traffic-related air pollution and improving air quality promptly, which could be considered as a feasible alternative in pollution control and air quality assurance in megacities of China like Beijing under particular or emergency occasions.

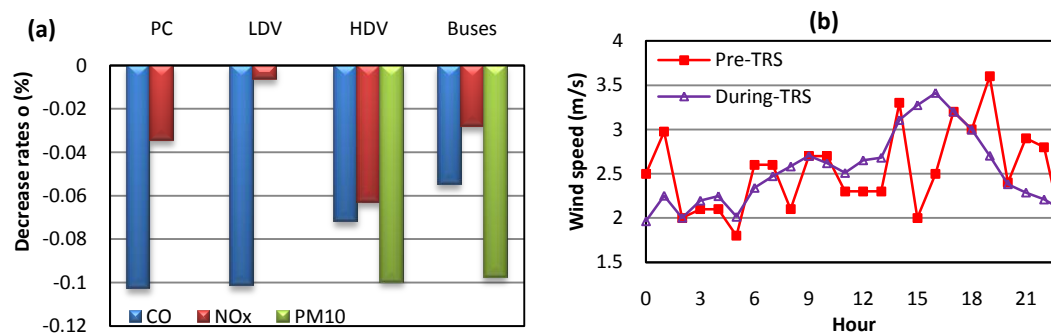


Fig. 11. Air quality impacts during the Games of (a): Reduction rates of emission factors of CO, PM<sub>10</sub> and NO<sub>x</sub> from various vehicle types resulting from the raise of running speed from 20 km/h to 23 km/h, calculated by COPERT model; (b) Variation of wind speed (m/s) on hourly average for the pre-TRS and during-TRS periods.

#### 4. Conclusions

We conducted a modelling evaluation of the air quality impacts of the odd-even day traffic restriction scheme implemented by the Beijing Municipal Government during the 2008 Olympic Games, and proved that this policy was effective in reducing short-term traffic-related air pollution and substantially improved the air quality in the urban area of Beijing during the Games.

The ADMS-Urban was well evaluated with hourly observations from a traffic-representative air quality monitoring station, and produced satisfactory predictions, based on the high temporal resolution on-road traffic flow data retrieved from the ITS-TAP monitoring network covering the 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup> RR and the LRs distributing over the major UAB, and on the hourly meteorological data from a representative Observatory. This study demonstrated that modelling-based air quality evaluation was a reliable approach and was especially useful for simultaneous and intensive assessment of air quality responses at multi-receptor and in different regions of the study domain. However, the possible drawback of using one traffic-representative station for model evaluation in this study was that the model predictions in areas away from the simulated roads might have some unknown uncertainty.

Both daily average and maximum concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> during the pre-TRS period decreased significantly to much lower levels during the TRS period, with the daily average concentrations from on-road vehicles conforming to the CNAAQs Grade II. However, pollutant concentrations of CO, PM<sub>10</sub> and NO<sub>2</sub>



increased for the post-TRS period, with the exception for O<sub>3</sub> concentration, which continued decreasing, mainly due to the “VOC-limited” characteristic for ozone formation in UAB. The bouncing of pollutant concentration levels of CO, PM<sub>10</sub> and NO<sub>2</sub> after the expiration of the TRS policy just reversely reflected the effectiveness of the TRS in improving short-term air quality.

The hourly average, especially the peak concentrations of CO, PM<sub>10</sub>, NO<sub>2</sub> and O<sub>3</sub> were reduced significantly in response to the TRS policy. However, the TRS policy did not change the typical two-peak diurnal variation pattern of the primary pollutants (CO and PM<sub>10</sub>). Besides, no remarkable weekly variation of pollutant concentrations were revealed as a result of the TRS policy, with the concentration levels of CO, PM<sub>10</sub>, NO<sub>2</sub> on weekday generally higher than those on weekends. Particularly, a notable ozone weekend effect with higher concentrations on weekends was revealed, mostly due to the decreased NO<sub>x</sub> emissions on weekends and the “VOC-limited” characteristic for ozone formation in UAB. Meanwhile, a remarkable reduction of peak hour and daily average concentrations of O<sub>3</sub> was achieved in response to the TRS policy.

Notable air quality improvement was revealed around UAB during the Games, which indicated the overall effectiveness of the TRS policy. Particularly, better effect on pollution reduction was observed in the previously most severely polluted areas, where the concentration level decreased sufficiently to comply with the CNAAQs guidelines. Besides, significant air quality improvement was achieved in

the upwind eastern and southern areas in response to the TRS policy. In conclusion, the TRS policy was effective in significantly improving short-term and regional air quality in UAB during the Games, and provided valuable experiences for future temporary and regional control of traffic-related air pollution.

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## References

Assael, M. J., Delaki, M., and Kakosimos, K. E.: Applying the OSPM model to the calculation of  $PM_{10}$  concentration levels in the historical centre of the city of Thessaloniki, *Atmos. Environ.*, 42, 65-77, 2008.

Atkinson-Palombo, C. M., Miller, J. A., and Balling, R. C.: Quantifying the ozone "weekend effect" at various locations in Phoenix, Arizona, *Atmos. Environ.*, 40, 7644-7658, 2006.

BeijingAir: <http://www.beijingairquality.cn>, 2008.

Beijing Municipal Environmental Protection Bureau: Beijing Environmental Bulletin,

1063 <http://www.bjepb.gov.cn>, 2009.

1064 Bennett, M., and Hunter, G. C.: Some comparisons of lidar estimates of peak  
 1065 ground-level concentrations with the predictions of UK-ADMS, *Atmos. Environ.*,  
 1066 31, 429-439, 1997.

1067 Berkowicz, R.: OSPM - a parameterised street pollution model, *Environ. Monit.*  
 1068 *Assess.*, 65, 323-331, 2000.

1069 Berkowicz, R., Ketzel, M., Jensen, S. S., Hvidberg, M., and Raaschou-Nielsen, O.:  
 1070 Evaluation and application of OSPM for traffic pollution assessment for a large  
 1071 number of street locations, *Environ. Modell. Softw.*, 23, 296-303, 2008.

1072 Cai, H., and Xie, S. D.: Estimation of vehicular emission inventories in China from  
 1073 1980 to 2005, *Atmos. Environ.*, 41, 8963-8979, 2007.

1074 Cai, H., and Xie, S. D.: Tempo-spatial variation of emission inventories of speciated  
 1075 volatile organic compounds from on-road vehicles in China, *Atmos. Chem. Phys.*,  
 1076 9, 6983-7002, 2009.

1077 Cambridge Environmental Research Consultants: Validation and sensitivity study of  
 1078 ADMS-Urban for London,  
 1079 [http://www.cerc.co.uk/environmental-research/assets/data/CERC\\_2003\\_ADMS-](http://www.cerc.co.uk/environmental-research/assets/data/CERC_2003_ADMS-Urban_validation_and_sensitivity_study_for_London_10_TR-0191-h.pdf)  
 1080 [Urban\\_validation\\_and\\_sensitivity\\_study\\_for\\_London\\_10\\_TR-0191-h.pdf](http://www.cerc.co.uk/environmental-research/assets/data/CERC_2003_ADMS-Urban_validation_and_sensitivity_study_for_London_10_TR-0191-h.pdf), 2003.

1081 Cambridge Environmental Research Consultants: Modelling dry deposition,  
1082 [http://www.cerc.co.uk/environmental-software/assets/data/doc\\_techspec/CERC\\_](http://www.cerc.co.uk/environmental-software/assets/data/doc_techspec/CERC_)  
1083 [ADMS4\\_P17\\_13.pdf](http://www.cerc.co.uk/environmental-software/assets/data/doc_techspec/CERC_ADMS4_P17_13.pdf), 2009.

1084 Cambridge Environmental Research Consultants, <http://www.cerc.co.uk>, 2009.

1085 Carruthers, D. J., Holroy, D. R. J., Hunt, J. C. R., Weng, W. S., Robins, A. G., Apsley,  
1086 D. D., Thomson, D. J., and Smith, F. B.: UK-ADMS - a new approach to  
1087 modeling dispersion in the earth's atmospheric boundary-layer, *J. Wind. Eng. Ind.*  
1088 *Aerod.*, 52, 139-153, 1994.

1089 Carruthers, D. J., Mckeown, A. M., Hall, D. J., and Porter, S.: Validation of ADMS  
1090 against wind tunnel data of dispersion from chemical warehouse fires, *Atmos.*  
1091 *Environ.*, 33, 1937-1953, 1999.

1092 Carruthers, D. J., Dyster, S., and McHugh, C. A.: Contrasting methods for validating  
1093 ADMS using the Indianapolis dataset, *Int. J. Environ. Pollut.*, 14, 115-121,  
1094 2000a.

1095 Carruthers, D. J., Edmunds, H. A., Lester, A. E., McHugh, C. A., and Singles, R. J.:  
1096 Use and validation of ADMS-Urban in contrasting urban and industrial locations,  
1097 *Int. J. Environ. Pollut.*, 14, 364-374, 2000b.

1098 Carruthers, D. J., Dickson, P., McHugh, C. A., Nixon, S. G., and Oates, W.:  
1099 Determination of compliance with UK and EU air quality objectives from  
1100 high-resolution pollutant concentration maps calculated using ADMS-Urban, *Int.*

1101 J. Environ. Pollut., 16, 460-471, 2001.

1102 CERC. ADMS-Urban User Guide. CERC Limited, Cambridge, February 1999.

1103 Chan, C. K., and Yao, X.: Air pollution in mega cities in China, Atmos. Environ., 42,  
1104 1-42, 2008.

1105 Cheng, Y. F., Heintzenberg, J., Wehner, B., Wu, Z. J., Su, H., Hu, M., and Mao, J. T.:  
1106 Traffic restrictions in Beijing during the Sino-African Summit 2006: aerosol size  
1107 distribution and visibility compared to long-term in situ observations, Atmos.  
1108 Chem. Phys., 8, 7583–7594, 2008.

1109 Chinese Olympic Committee: The temporary traffic control implementations for  
1110 non-Beijing local vehicles during 2008 Beijing Olympic and Paralympic Games,  
1111 <http://www.beijing2008.cn/news/olympiccities/beijing/n214412173.shtml>, 20  
1112 June, 2008.

1113 Chou, C. C. K., Liu, S. C., Lin, C. Y., Shiu, C. J., and Chang, K. H.: The trend of  
1114 surface ozone in Taipei, Taiwan, and its causes: Implications for ozone control  
1115 strategies, Atmos. Environ., 40, 3898-3908, 2006.

1116 McHugh, C., Carruthers, D., Sheng, X.Y.: Using ADMS models for air quality  
1117 assessment and management in China, Cambridge Environmental Research  
1118 Consultants (CERC), Chinese. J. Popul. Resour. Environ., 3, 2005.

1119 Derwent, R.G., and Middleton, D.R.: An empirical function for the ratio  $\text{NO}_2/\text{NO}_x$ .  
1120 Clean Air (NSCA) 26, 57–60, 1996.

1121 Elbir, T.: Application of an ISCST3 model for predicting urban air pollution in the  
 1122 Izmir metropolitan area, *Int. J. Environ. Pollut.*, 18, 498-507, 2002.

1123 European Environment Agency: COPERT III-Computer programme to calculate  
 1124 emissions from road transport, methodology and emission factors (version 2.1),  
 1125 technical report No. 49, 2000.

1126 Fang, M., Chan, C. K., and Yao, X. H.: Managing air quality in a rapidly developing  
 1127 nation: China, *Atmos. Environ.*, 43, 79-86, 2009.

1128 Gao, H. O., and Niemeier, D. A.: The impact of rush hour traffic and mix on the ozone  
 1129 weekend effect in southern California, *Transport. Res. D-Tr. E.*, 12, 83-98, 2007.

1130 Granell, J., Ho, C., Tang, T., and Roberts, M.: Analysis of MOBILE6.2's PM emission  
 1131 factor estimating function, USEPA's 13th International Emission Inventory  
 1132 Conference "Working for Clean Air in Clearwater",  
 1133 <http://www.epa.gov/ttn/chief/conference/ei13/mobile/granell.pdf>, 2004.

1134 Hanna, S.R., Strimaitis, D.G., and Chang, J.C.: Hazard response modeling uncertainty  
 1135 (A quantitative method), 1991.

1136 Hanna, S.R., Chang, J.C., and Strimaitis, D.G.: Hazardous gas model evaluation with  
 1137 field observations, *Atmos. Environ.*, 27, 2265–2285, 1993.

1138 Hao, J. M., Wu, Y., Fu, L. X., He, D. Q., and He, K. B.: Source contributions to  
 1139 ambient concentrations of CO and NO<sub>x</sub> in the urban area of Beijing, *J. Environ.*  
 1140 *Sci. Heal. A - Toxic/Hazard. Subst. Environ. Eng.*, 36, 215-228, 2001.

1141 Hirtl, M., and Baumann-Stanzer, K.: Evaluation of two dispersion models  
 1142 (ADMS-Roads and LASAT) applied to street canyons in Stockholm, London and  
 1143 Berlin, *Atmos. Environ.*, 41, 5959-5971, 2007.

1144 Hu, F.: Turbulence interval and atmospheric boundary layer [D], Ph.D dissertation (in  
 1145 Chinese), Institute of Atmospheric Physics, Chinese Academy of Sciences, 1994.

1146 Kenty, K. L., Poor, N. D., Kronmiller, K. G., McClenny, W., King, C., Atkeson, T.,  
 1147 and Campbell, S. W.: Application of CALINE4 to roadside NO/NO<sub>2</sub>  
 1148 transformations, *Atmos. Environ.*, 41, 4270-4280, 2007.

1149 Khoder, M. I.: Diurnal, seasonal and weekdays-weekends variations of ground level  
 1150 ozone concentrations in an urban area in greater Cairo, *Environ. Monit. Assess.*,  
 1151 149, 349-362, 2009.

1152 Kukkonen, J., Valkonen, E., Walden, J., Koskentalo, T., Aarnio, P., Karppinen, A.,  
 1153 Berkowicz, R., and Kartastenpaa, R.: A measurement campaign in a street  
 1154 canyon in Helsinki and comparison of results with predictions of the OSPM  
 1155 model, *Atmos. Environ.*, 35, 231-243, 2001.

1156 Levitin, J., Harkonen, J., Kukkonen, J., and Nikmo, J.: Evaluation of the CALINE4  
 1157 and car-fmi models against measurements near a major road, *Atmos. Environ.*,  
 1158 39, 4439-4452, 2005.

1159 Liu, J., Zhang, X.L., Xie, P., Dong, F., Ouyang, J., and Wang, Z.F.: The variation  
 1160 characteristics of trace reactive gases at Shangdianzi regional background

1161 monitoring station, Environ. Chem. (in Chinese), 26 (5): 693-698, 2007.

1162 Liu, J., Zhang, X.L., Xu, X.F., and Xu, H.H.: Comparison analysis of variation  
 1163 characteristics of SO<sub>2</sub>, NO<sub>x</sub>, O<sub>3</sub> and PM<sub>2.5</sub> between rural and urban areas, Beijing,  
 1164 Environ. Science. (in Chinese), 29 (4): 1059-1065, 2008.

1165 Liu, J., Zhang, X.L., Zhang, X.C., and Tang, J.: Surface ozone characteristics and the  
 1166 correlated factors at Shangdianzi atmospheric background monitoring station,  
 1167 Res. Environ. Sciences. (in Chinese), 19 (4): 19-25, 2006.

1168 Liu, X.M., Hu, and F., Li, L.: Analyses on the characteristics of Beijing summer urban  
 1169 heat island and its meteorological fields (in Chinese), J. Grad. School. Chinese.  
 1170 Acad. Sci., 23, 70-76, 2006.

1171 Lu, L.H., Bian, L.G., Cheng, Y.J., Gao, Z.Q., and Wang, Y.: Meteorological  
 1172 Characteristics of the ground layer in Beijing in winter, J. Appl. Meteorol.  
 1173 Science. (in Chinese), 13 (Supl.): 34-42, 2002.

1174 McHugh, C. A., Carruthers, D. J., and Edmunds, H. A.: ADMS-Urban: An air quality  
 1175 management system for traffic, domestic and industrial pollution, Int. J. Environ.  
 1176 Pollut., 8, 666-674, 1997.

1177 Miao, S.G., Chen, F., Margaret, A.L., Mukul, T., Li, Q.C., and Wang, Y.C.: An  
 1178 observational and modeling study of characteristics of urban heat island and  
 1179 boundary layer structures in Beijing, J. Appl. Meteorol. Clim., 48, 484-501,  
 1180 2008.



1181 Ministry of Environmental Protection of the People's Republic of China: Disposition  
 1182 of banning of biomass burning for the air quality assurance during the Games,  
 1183 [http://test.mep.gov.cn/ztbdsjhjr/0865/lsay/200805/t20080529\\_123206.htm](http://test.mep.gov.cn/ztbdsjhjr/0865/lsay/200805/t20080529_123206.htm) (in  
 1184 Chinese), 2008.

1185 Ministry of Environmental Protection of the People's Republic of China, Guidelines  
 1186 for environmental impact assessment - atmospheric environment, pp 24, 2009.

1187 Murphy, J. G., Day, D. A., Cleary, P. A., Wooldridge, P. J., Millet, D. B., Goldstein, A.  
 1188 H., and Cohen, R. C.: The weekend effect within and downwind of Sacramento -  
 1189 Part 1: Observations of ozone, nitrogen oxides, and VOC reactivity, *Atmos.*  
 1190 *Chem. Phys.*, 7, 5327-5339, 2007.

1191 NARSTO: An assessment of tropospheric ozone pollution: A North American  
 1192 perspective, The NARSTO Synthesis Team, Palo Alto, CA, 2000.

1193 NASA, <http://aura.gsfc.nasa.gov/science/china.html>, 2009.

1194 Okuda, T., Kato, J., Mori, J., Tenmoku, M., Suda, Y., Tanaka, S., He, K., Ma, Y.,  
 1195 Yang, F., Yu, X., and Duan, F.: Daily concentrations of trace metals in aerosols  
 1196 in Beijing, China, determined by using inductively coupled plasma mass  
 1197 spectrometry equipped with laser ablation analysis, and source identification of  
 1198 aerosols, *Sci. Total Environ.*, 330 (1-3), 145-158, 2004.

1199 Olesen, H.R.: Ten years of harmonization activities: past, present, and future. 7th Int.  
 1200 Conf. on Harmonisation within Atmospheric Dispersion Modelling for

1201 Regulatory Purposes, Belgirate, Italy, National Environmental Research Institute,  
1202 Roskilde, Denmark, [www.harmo.org](http://www.harmo.org), 2001.

1203 Riddle, A., Carruthers, D., Sharpe, A., McHugh, C., and Stocker, J.: Comparisons  
1204 between fluent and ADMS for atmospheric dispersion modelling, *Atmos.*  
1205 *Environ.*, 38, 1029-1038, 2004.

1206 Sadanaga, Y., Shibata, S., Hamana, M., Takenaka, N., and Bandow, H.:  
1207 Weekday/weekend difference of ozone and its precursors in urban areas of Japan,  
1208 focusing on nitrogen oxides and hydrocarbons, *Atmos. Environ.*, 42, 4708-4723,  
1209 2008.

1210 Sharma, S., and Chandra, A.: Simulation of air quality using an ISCST3 dispersion  
1211 model, *Clean-Soil. Air. Water.*, 36, 118-124, 2008.

1212 Song, Y., Tang, X.Y., Xie, S.D., Zhang, Y.H., Wei, Y.J., Zhang, M.S., Zeng, L.M.,  
1213 and Lu, S.H.: Source apportionment of PM<sub>2.5</sub> in Beijing in 2004, *J. Hazard.*  
1214 *Mater.*, 146, 124-130, 2007.

1215 Song, Y., Zhang, M.S., Cai, X.H.: PM<sub>10</sub> modeling of Beijing in winter, *Atmos.*  
1216 *Environ.*, 40 (22), 4126-4136, 2006a.

1217 Song, Y., Zhang, Y.H., Xie, S.D., Zeng, L.M., Zheng, M., Salmon, L.G., Shao, M.,  
1218 and Slanina, S.: Source apportionment of PM<sub>2.5</sub> in Beijing by positive matrix  
1219 factorization, *Atmos. Environ.*, 40 (8), 1526-1537, 2006b.

1220 Sun, Y.L., Zhuang, G.S., Wang, Y., Han, L.H., Guo, J.H., Dan, M., Zhang, W.J.,

1221 Wang, Z.F. and Hao, Z.P.: The air-borne particulate pollution in  
 1222 Beijing-concentration, composition, distribution and sources, *Atmos. Environ.*,  
 1223 38 (35), 5991-6004, 2004.

1224 Streets, D.G., Fu, J.S., Jang, C.J., Hao, J.M., He, K.B., Tang, X.Y., Zhang, Y.H.,  
 1225 Wang, Z.F., Li, Z.P., Zhang, Q., Wang, L.T., Wang, B.Y., and Yu, C.: Air  
 1226 quality during the 2008 Beijing Olympic Games, *Atmos. Environ.*, 41 (3),  
 1227 480-492, 2007.

1228 Tang, W. Y., Zhao, C. S., Geng, F. H., Peng, L., Zhou, G. Q., Gao, W., Xu, J. M., and  
 1229 Tie, X. X.: Study of ozone "weekend effect" in Shanghai, *Sci China Ser D*, 51,  
 1230 1354-1360, 2008.

1231 Vardoulakis, S., Valiantis, M., Milner, J., and ApSimon, H.: Operational air pollution  
 1232 modelling in the UK-Street canyon applications and challenges, *Atmos. Environ.*,  
 1233 41, 4622-4637, 2007.

1234 Venkatram, A., Karamchandani, P.O., Pai, P., Goldstein, R.: The development and  
 1235 application of a simplified ozone modelling system, *Atmos. Environ.*, 28,  
 1236 3665-3678, 1994.

1237 Wang, B.G., Zhang, Y.H., Wu, Z.Q., Chan, L.Y.: Tunnel test for motor vehicle  
 1238 emission factors in Guangzhou, *Res. Environ. Sci.*, 14, 13-16, 2001.

1239 Wang, H.L., Zhuang, Y.H., Wang, Y., Sun, Y.L., Yuan, H., Zhuang, G.S., and Hao,  
 1240 Z.P.: Long-term monitoring and source apportionment of PM<sub>2.5</sub>/PM<sub>10</sub> in Beijing,

1241 China, *J. Environ. Sci. - China*, 20, 1323-1327, 2008b.

1242 Wang, L. J., Parker, D. B., Parnell, C. B., Lacey, R. E., and Shaw, B. W.: Comparison  
 1243 of Calpuff and ISCST3 models for predicting downwind odor and source  
 1244 emission rates, *Atmos. Environ.*, 40, 4663-4669, 2006.

1245 Wang, L.T., Hao, J.M., He, K.B., Wang, S.X., Li, J.H., Zhang, Q., Streets, D.G., Fu,  
 1246 J.S., Jang, C.J., Takekawa, H., and Chatani, S.: A modeling study of coarse  
 1247 particulate matter pollution in Beijing: Regional source contributions and control  
 1248 implications for the 2008 Summer Olympics, *J. Air. Waste. Manag. Assoc.*, 58  
 1249 (8), 1057-1069, 2008a.

1250 Wang, T. and Xie, S.D.: Assessment of traffic-related air pollution in the urban streets  
 1251 before and during the 2008 Beijing Olympic Games traffic control period, *Atmos.*  
 1252 *Environ.*, 43 (35), 5682-5690, 2009.

1253 Wang, W.X., Chai, F.H., Zhang, Kai., Wang, S.L., Chen, Y.Z., Wang, X.Z., and Yang,  
 1254 Y.Q.: Study on ambient air quality in Beijing for the summer 2008 Olympic  
 1255 Games, *Air. Qual. Atmos. Health.*, 1 (1), 31-36, 2008c.

1256 Wang, X., Westerdahl, D., Chen, L. C., Wu, Y., Hao, J. M., Pan, X. C., Guo, X. B.,  
 1257 and Zhang, K. M.: Evaluating the air quality impacts of the 2008 Beijing  
 1258 Olympic Games: On-road emission factors and black carbon profiles, *Atmos.*  
 1259 *Environ.*, 43 (30), 4535-4543, 2009a.

1260 Wang, X. S., and Li, J. L.: The contribution of anthropogenic hydrocarbons to ozone

1261 formation in Beijing areas (in Chinese), *China Environ Sci*, 22, 501-505, 2002.

1262 Wang, X. S., Li, J. L., Zhang, Y. H., Xie, S. D., and Tang, X. Y.: Ozone source  
 1263 attribution during a severe photochemical smog episode in Beijing, China, *Sci.*  
 1264 *China. Ser. B-Chem.*, 52, 1270-1280, 2009c.

1265 Wang, Y., Hao, J., McElroy, M. B., Munger, J. W., Ma, H., Chen, D., and Nielsen, C.  
 1266 P.: Ozone air quality during the 2008 Beijing Olympics: Effectiveness of  
 1267 emission restrictions, *Atmos. Chem. Phys.*, 9, 5237-5251, 2009b.

1268 WHO: World Health Organization Air Quality Guidelines Global Update,  
 1269 [http://www.euro.who.int/air/activities/20050222\\_2](http://www.euro.who.int/air/activities/20050222_2), 2005.

1270 Willmott, C.J.: Some comments on the evaluation of model performance, *Bull. Amer.*  
 1271 *Meteor. Soc.* 63, 1309–1313, 1982.

1272 Wu, Y., Hao, J.M., Fu, L.X., Wang, Z.S., Tang, U.: Vertical and horizontal profiles of  
 1273 airborne particulate matter near major roads in Macao, China. *Atmos. Environ.*,  
 1274 36 (31), 4907-4918, 2002.

1275 Xie, S. D., Liu, Z., Chen, T., and Hua, L.: Spatiotemporal variations of ambient PM<sub>10</sub>  
 1276 source contributions in Beijing in 2004 using positive matrix factorization,  
 1277 *Atmos. Chem. Phys.*, 8, 2701-2716, 2008.

1278 Ying, G. X., Ma, J., and Xing, Y.: Comparison of air quality management strategies of  
 1279 PM<sub>10</sub>, SO<sub>2</sub>, and NO<sub>x</sub>, by an industrial source complex model in Beijing, *Environ.*  
 1280 *Prog.*, 26, 33-42, 2007.

- 1281 Yura, E. A., Kear, T., and Niemeier, D.: Using CALINE dispersion to assess vehicular  
1282 PM<sub>2.5</sub> emissions, *Atmos. Environ.*, 41, 8747-8757, 2007.
- 1283 Zhang, H.S. and Chen, J.Y.: Estimation of aerodynamic parameters on nonsingle  
1284 horizontal homogeneous underlying surface, *Journal of Applied Meteorological*  
1285 *Science (in Chinese)*, 8 (3): 310-315, 1997.
- 1286 Zhang, W., Guo, J.H., Sun, Y.L., Yuan, H., Zhuang, G.S., Zhuang, Y.H., and Hao,  
1287 Z.P.: Source apportionment for urban PM<sub>10</sub> and PM<sub>2.5</sub> in the Beijing area,  
1288 *Chinese Science Bulletin*, 52 (5), 608-615, 2007.
- 1289 Zheng, M., Salmon, L.G., Schauer, J.J., Zeng, L.M., Kiang, C.S., Zhang, Y.H., and  
1290 Cass, G.R.: Seasonal trends in PM<sub>2.5</sub> source contributions in Beijing, China,  
1291 *Atmos. Environ.*, 39 (22), 3967-3976, 2005.