Response to the comments

Reviewer one

Atmospheric particulates (aerosols) has significant influence on air quality, human health and regional climate changes. Large uncertainties in estimating the aerosol direct radiative forcing exist due to the uncertainties in the aerosol optical properties which were related to the aerosol emissions, profiles, compositions and mixing states. Thus, it's very important to figure out the temporal and spatial distributions of the aerosols in the regions with high aerosol loadings to better accessing their radiative forcing and regional/global climate effects. This study, which examines the spatial and temporal variations of the PM10, PM2.5 and PM1 over 24 sites in China by using a 9 years near continuous PM data measured by GRIMM180 instruments, is important to some extent and the paper has potential. I recommend the manuscript being published in the journal after the revision listed below being addressed.

Comments:

1. Re-typesetting the manuscript. Also, the manuscript still remains poorly written throughout the whole manuscript and requires many corrections.

Response: The manuscript has been edited by a native English-speaking expert at LucidPapers (http://lucidpapers.com/). A certification is attached.

2. Page 15320, line 9 and line 11: It should be "the ratio of PM2.5 to PM10" and "the ratio of PM1 to PM2.5".

Response: It has been revised to "The ratios of $PM_{2.5}$ to PM_{10} showed a clear increasing trend from northern to southern China, because of the substantial contribution of coarse mineral aerosol in northern China. The ratios of PM_1 to $PM_{2.5}$ were higher than 80% at most stations".

3. Page 15320, line 16-17: The authors should show the readers what they found while just what they did.

Response: The brief conclusion was added at the end of the abstract: "Bimodal and unimodal diurnal variation patterns were identified at urban stations. The investigation of meteorological factors effects reveals the emission variation possibly dominates the long-term PM concentration trend; meanwhile meteorological factors play a leading role during a short period".

4. Page 15323, line 9: The authors indicate the uncertainty of GRIMM in (Grimm and Eatough) was 9.9%, how about in China? Is it also 9.9%?

Response: The comparison result of hourly average PM2.5 concentrations simultaneously monitored by GRIMM and TEOM at SDZ (around Beijing, China) in February 2010 shows the good linear relationship between GRIMM and TEOM PM2.5 concentrations (Zhao et al., 2011) (Fig. 1). Also an early study in Beijing downtown during the summer 2004 concluded that GRIMM measurements have shown to reproduce very well all the TEOM-FDMS variations (wet and dry periods) suggesting that optical measurements could be used derive PM2.5 and could also account for semi-volatile material in aerosols (Sciare et al., 2007). There is no specific uncertainty value given by above studies, and such uncertainty value is not constant in different regions and periods. The point is to give the reference evidences that the GRIMM data are acceptable for PM measurement compared with traditional used instruments such as TEOM.

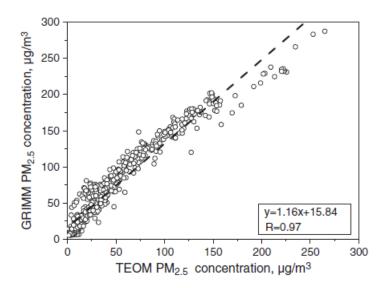


Fig. 1. GRIMM vs. TEOM PM2.5 concentration, 1h average, SDZ, February 2010 (Zhao et al., 2011)

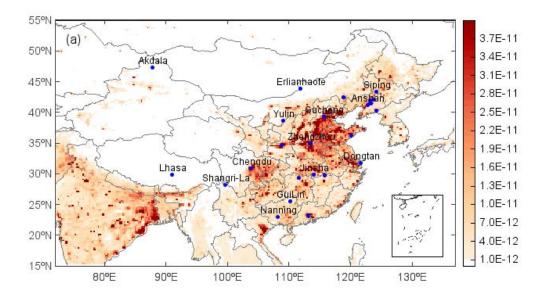
The above two references has been added in revised manuscript: "The GRIMM measurement in Beijing, China have shown the good linear relationship with TEOM and suggesting that optical measurements could be used derive PM2.5 and could account for semi-volatile material in aerosols (Sciare et al., 2007; Zhao et al., 2011)".

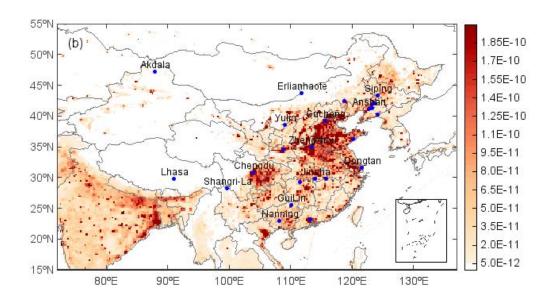
5. Page 15324, line 1: The authors should present some emission results in China form publications (e. g. Q. Zhang et al., 2009) when explaining the reasons.

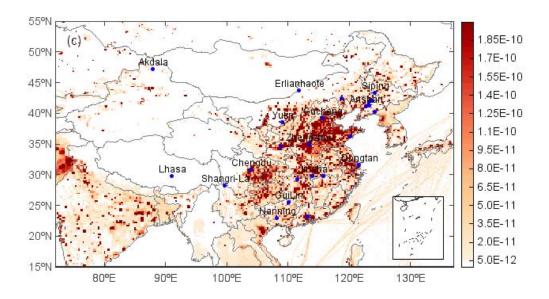
Response: Some emission results in China have been added in revised manuscript. Detailed description could be found in below response.

6. Again, it's obvious that the results being analyzed is too simple. The authors should provide evidences (from the similar studies in publications/references or from self-analysis) to make the readers more clearly.

Response: We have enhanced the analysis part with especially more emission results in China. In revised manuscript, section 3.6 was changed to "Emission and meteorological influences" with emission data analysis based on HTAP harmonized emissions database (http://iek8wikis.iek.fz-juelich.de/HTAPWiki/WP1.1). Anthropogenic emission distributions of BC, PM2.5, SO2 and NO2 in 2010 were presented in figure 2 with similar spatial pattern. PM loadings in China were generally similar to this emission pattern. For example, most PM pollutant stations located in highest emission region of HBP. Therefore, PM loadings were controlled by anthropogenic emission amount in mid-east China.







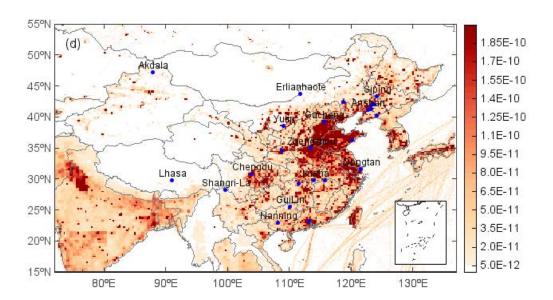
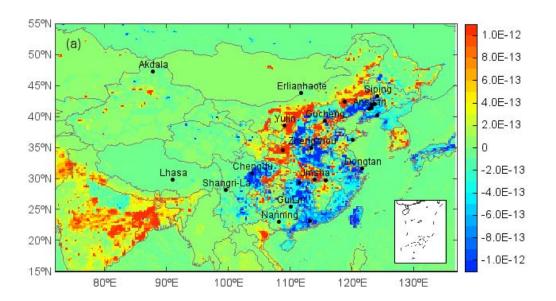
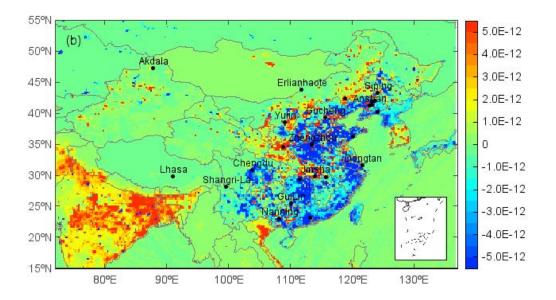


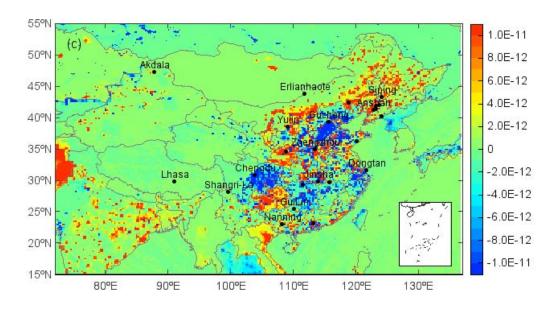
Fig. 2. Anthropogenic emission distributions at 0.1 degree \times 0.1 degree resolution of (a) BC, (b) PM2.5, (c) SO2 and (d) NOx (units: kg m⁻² s⁻²) based on HTAP_v2 dataset

The emission trends in China during 2005-2010 (Wang et al., 2014) were also presented to explain the long-term PM trends. The emissions of SO2 and PM2.5 in East Asia decreased by 15 and 12 %, meanwhile the emissions of NOx and NMVOC increased by 25 and 15 % (Wang et al., 2014). Also spatial distributions of emission difference between 2010 and 2008 were studied using HTAP_v2 emission dataset (Fig. 3), so the PM vs emission variation in different regions could be studied. Although there is no published emission data after 2010, it was believed that the emission was greatly controlled after end of 2013 with the issue of "Action Plan for the Control of Air Pollution" document. It can explain the general decrease trend in

2014. The conclusion of emission influence on long-term PM pattern could be supported from above analysis. More detailed description was added in revised manuscript.







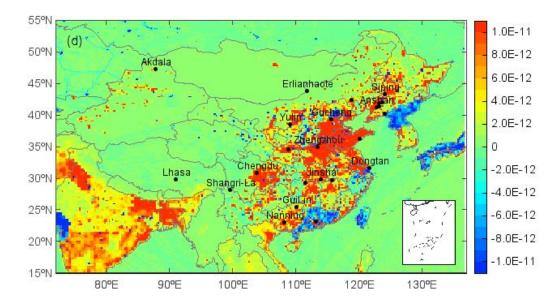


Fig. 3. Emission difference between 2010 and 2008 at 0.1 degree \times 0.1 degree resolution of (a) BC, (b) PM2.5, (c) SO2 and (d) NOx (units: kg m⁻² s⁻²) based on HTAP v2 dataset

7. In addition to diurnal and seasonal cycles, did the PM have any periodic in China.

Response: There is no obvious general long-term periodic of PM in China from this 9-year data study.

Reviewer two

This manuscript presented PM concentrations over China measured by China Atmosphere Watch Network. Although PM2.5 measurements are available over China since 2013, the measurement data presented in this paper are valuable because it covers longer period (2006-2014) and provides PM1/PM2.5/PM10 information. This dataset would be very helpful for understanding the evolution of PM concentration over China. The manuscript could be published in ACP after the following concerns are addressed.

Comments:

8. Sect. 2. The authors claimed that GRIMM measurements are in good agreement with TEOM. I am curious if the authors compared GRIMM with other instruments under heavy pollution condition in China. Referring to comparison in western countries is not very convincing.

Response: This comment has been answered in comment 4.

9. P15326, L6-16, more comparison for the same period (2006-2014) would be meaningful.

Response: More comparison for the same period (2006-2014) has been added in the revised manuscript.

"The average PM_{10} and $PM_{2.5}$ concentrations were 23.9 and 16.3 μg m⁻³ for the period 2008-2009 in Netherlands (Janssen et al., 2013). The 20 European study areas observation results from ESCAPE project between October 2008 and April 2011 showed PM_{10} and $PM_{2.5}$ have similar spatial pattern with low concentrations in Northern Europe and high concentrations in Southern and Eastern Europe (Eeftens et al., 2012)"

10. P15328, Sect. 3.4, this could be the most important section in the manuscript. In this section, inter-annual variations of PM2.5 concentrations of individual sites are presented one by one. It would be very boring the international readers who don't familiar with Chinese cities. The overall PM trend over China and the driven forces behind the trend are missing in this section.

Response: There is no overall PM trend over whole China except the decrease trend in 2014, so the trend analysis was done in different areas of China, such as HBP, northeast China, southern China and so on. Such summary sentence has been added in revised manuscript. The driven forces behind the trend were analyzed by emission and meteorological analysis in section 3.6. (See response of comment 6)

11. P15330, Sect. 3.5, I would like to see diurnal variations in PM1 and PM10 concentrations and if they are similar to PM2.5.

Response: The diurnal variations in PM1 and PM10 concentrations are similar to PM2.5 (Fig. 4). An example figure for stations of Zhengzhou, Xian and Gucheng is given below. So we add a sentence of "The diurnal variations in PM_1 and PM_{10} concentrations are similar to $PM_{2.5}$ at most stations" in revised manuscript.

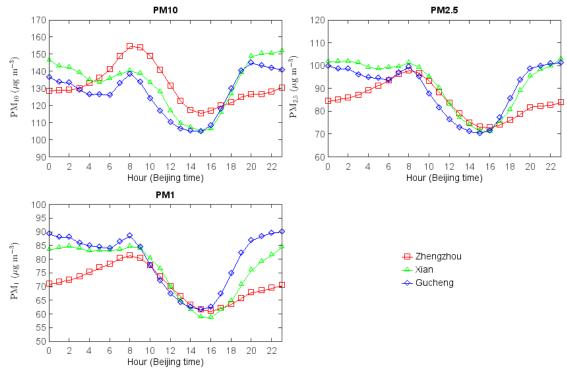


Fig. 4. Diural variations in PM10, PM2.5 and PM1 at Zhengzhou, Xian and Gucheng

12. P15332, L16-17, the authors concluded that "emission variation must to be considered for long-term trend analysis especially in rapid developing countries." Emission data should be used in the discussion to support their arguments.

Response: The emission variation data have been provided for this analysis. Please see the response to comment 6.

13. Figures. Fig. 1 and Fig. 2 could be combined into one figure.

Response: Fig. 1 and Fig. 2 were combined as below one figure.

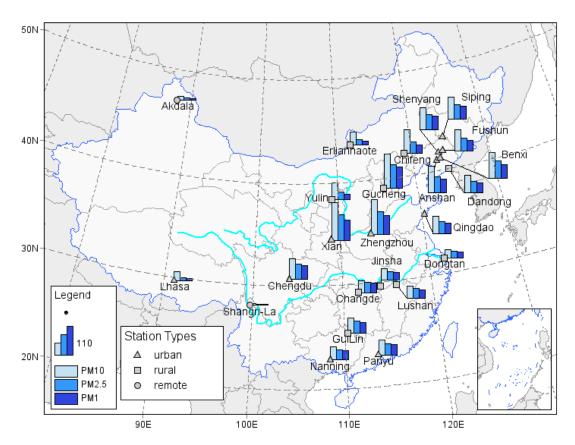


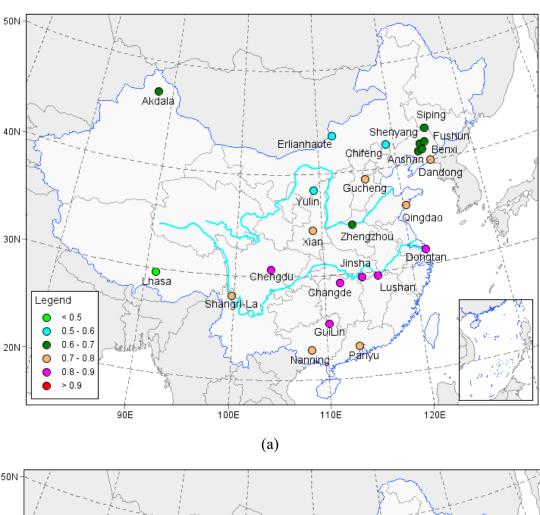
Fig. 5. Map showing PM observation stations and their bar charts of average PM_{10} , $PM_{2.5}$ and PM_1 concentrations ($\mu g \ m^{-3}$)

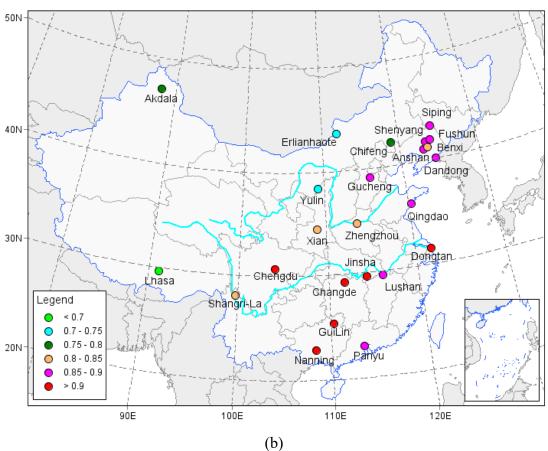
14. Figure 3 could be removed because it doesn't provide additional information than Table 2.

Response: We agree and have removed figure 3 in revised manuscript.

15. PM1/PM2.5 ratios should be also presented in Figure 5.

Response: PM1/PM2.5 ratios have been presented in Figure 5 as below.





Reference

- Eeftens, M., Tsai, M.-Y., Ampe, C., Anwander, B., Beelen, R., Bellander, T., Cesaroni, G., Cirach, M., Cyrys, J., Hoogh, K. d., Nazelle, A. D., Vocht, F. d., Declercq, C., Dedele, A., Eriksen, K., Galassi, C., Grazuleviciene, R., Grivas, G., Heinrich, J., Hoffmann, B., Iakovides, M., Ineichen, A., Katsouyanni, K., Korek, M., Krämer, U., Kuhlbusch, T., Lanki, T., Madsen, C., Meliefste, K., Mölter, A., Moslerm, G., Nieuwenhuijsen, M., Oldenwening, M., Pennanen, A., Probst-Hensch, N., Quass, U., Raaschou-Nielsen, O., Ranzi, A., Stephanou, E., Sugiri, D., Udvardy, O., Vaskövi, É., Weinmayr, G., Brunekreef, B., and Hoek, G.: Spatial variation of PM_{2.5}, PM₁₀, PM_{2.5} absorbance and PMcoarse concentrations between and within 20 European study areas and the relationship with NO₂ Results of the ESCAPE project. *Atmos. Environ.* **62**, 303-317, 2012.
- Janssen, N. A. H., Fischer, P., Marra, M., Ameling, C., and Cassee, F. R.: Short-term effects of PM_{2.5}, PM₁₀ and PM_{2.5-10} on daily mortality in the Netherlands. *Sci. Total Environ.* **463-464**, 20-26, 2013.
- Sciare, J., Cachier, H., Sarda-Este've, R., Yu, T., and Wang, X.: Semi-volatile aerosols in Beijing (R.P. China): Characterization and influence on various PM2.5 measurements. *J. Geophys. Res.* **112**, D18202, doi:10.1029/2006JD007448, 2007.
- Wang, S. X., Zhao, B., Cai, S. Y., Klimont, Z., Nielsen, C. P., Morikawa, T., Woo, J. H., Kim, Y., Fu, X., Xu, J. Y., Hao, J. M., and He, K. B.: Emission trends and mitigation options for air pollutants in East Asia. *Atmos. Chem. Phys.* 14, 6571-6603, 2014.
- Zhao, X., Zhang, X., Pu, W., Meng, W., and Xu, X.: Scattering properties of the atmospheric aerosol in Beijing, China. *Atmos. Res.* **101**, 799-808, 2011.

Certificate



Reference number: 2015-110201 Date: 09 November 2015

Contact author: Yaqiang Wang

Manuscript: Spatial and temporal variations in the concentrations of PM10, PM2.5 and PM1 in China

This document certifies that the above-detailed manuscript was edited by a native English-speaking expert at LucidPapers on the date stated.

Following the editing process, the editor's overall assessment is that:

The manuscript will be ready for consideration by the target journal once the edits have been checked and approved/rejected as necessary.	
The manuscript may require modifications to the text in response to the editor's comments/queries, but further professional editing is unlikely to be needed.	\square
The manuscript may require modifications to the text in response to the editor's comments/queries, and further professional editing may be needed.	
The manuscript may require modifications to the text in response to the editor's comments/queries, and further professional editing is more than likely going to be needed.	
The manuscript requires modifications to the text in response to the editor's comments/queries, and further professional editing is strongly recommended.	

Signed:

Colin Smith Chief Editor LucidPapers

Email: colin.smith@lucidpapers.com
Website: http://www.lucidpapers.com

Address:

85 Cotmore Close Thame Oxfordshire OX9 3ND United Kingdom

Spatial and temporal variations of the concentrations

- $_2$ of PM₁₀, PM_{2.5} and PM₁ in China
- 3

- 4 Y. Q. Wang^{1*}, X. Y. Zhang^{1*}, J. Y. Sun¹, X. C. Zhang², H. Z. Che¹, Y. Li¹
- 5 6
 - [1] {Laboratory of Atmospheric Chemistry, Chinese Academy of Meteorological
- 7 Sciences, Beijing, China}
- 8
- 9 [2]{Meteorological Observation Center, Beijing, China}
- 10 Correspondence to: Y. Q. Wang (<u>wangyq@cams.cma.gov.cn</u>), X. Y. Zhang
- 11 (xiaoye@cams.cma.gov.cn)
- 12

Abstract:

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

31

32

33

34

Concentrations of PM₁₀, PM_{2.5} and PM₁ were monitored at 24 stations of CAWNET (China Atmosphere Watch Network) stations from 2006 to 2014 using GRIMM 180 dust monitors. The highest particulate matter (PM) concentrations were observed at the stations of Xian, Zhengzhou and Gucheng, on the Guanzhong Plain and the Hua Bei Plain (HBP). The second highest PM concentrations were observed in northeast China, followed by southern China. According to the latest air quality standards of China, 14 stations reached the PM₁₀ standard, and only 7 stations, mainly rural and remote stations, reached the PM_{2.5} standard. The <u>ratios of PM_{2.5} and to PM₁₀ ratios</u> showed a clear increasing trend from northern to southern China, because of the substantial contribution of coarse mineral aerosol in northern China. The ratios of PM₁ and to PM_{2.5} ratios were higher than 80% at most stations. PM concentrations tended to be highest in winter and lowest in summer at most stations, and mineral dust impacts influenced the results in spring. A decreasing interannual trend was observed in on the HBP and in southern China from for the period 2006 to 2014, but an increasing trend occurred at some stations in northeast China. Also diurnal variationsof PM concentrations and meteorological factors effects were investigated. Bimodal and unimodal diurnal variation patterns were identified at urban stations. The investigation of emission and meteorological factors effects reveals bBoth emissions and meteorological variations dominate the long-term PM concentration trend; meanwhile while meteorological factors play a leading role during a short period in the short-term.

- **Keywords:** particulate matter, observation, spatial and temporal spatiotemporal
- 36 variation

39

40

41

42

43

44

45

46

47

48

49

50

51

52

53

54

55

56

57

58

59

1. Introduction

Tropospheric aerosols are important because of their strong influence on the climate system through both direct and indirect effects. These include the direct effect of scattering and absorbing radiant energy, and the indirect effect of modifying the microphysical properties of clouds, and hence their radiative properties and lifetime (Haywood and Boucher, 2000). They also attract attention because of their adverse effects on visibility impairment (Watson, 2002) and human health (Delfino et al., 2005; Pope III and Dockery, 2006). Therefore, the spatial and temporal variation of aerosols is essential to understand, but remains a complex subject because of their relatively short lifetimeephemeral nature and the complexity of aerosol their physical and chemical properties (Ramanathan et al., 2001). Particle size is considered a key parameter to define the impact of particulate matter (PM) on human health; specifically, fine PM (PM_{2.5} and PM₁) poses a greater health risk than coarse PM (PM₁₀) (Oberdörster et al., 2005). There have been numerous network-based observation studies of the PM_{2.5} concentration and chemical composition in North America and Europe. For example, based on a dataset across 19 Canadian sites, most of the PM_{2.5} concentrations were found to be below 26 μg m⁻³, and PM_{2.5} accounted for 49% of the measured PM₁₀ (Brook et al., 1997). Meanwhile, Eldred et al. (1997) reported that PM_{2.5} and PM₁₀ particulate concentrations measured at 42 sites in of the Interagency Monitoring of Protected Visual Environments (IMPROVE) network over the 1993 seasonal year (March 1993 to February 1994)

showed the PM_{2.5} concentration had a large gradient from west to east in the US, 60 averaging 3 μg m⁻³ in most of the west compared with 13 μg m⁻³ in the Appalachian region (Eldred et al., 1997). Another study, based on 143 IMPROVE sites of 62 63 **IMPROVE** in the year 2001, showed that sulfates, carbon and crustal material were 64 responsible for most of the measured PM_{2.5} at the majority of sites in the US (Malm et al., 2004). The temporal variation and spatial distribution of PM_{2.5} concentrations 65 66 have also been reported in Switzerland (Gehrig and Buchmann, 2003), Austria (Gomiscek et al., 2004), and six central and eastern European countries (Houthuijs et 67 al., 2001). 68 As a country with a rapidly developing economy, China has suffered from a 70 serious air pollution problem in recent years due to substantial increases in energy consumption and other related production of large amounts of aerosols and precursor gas emissions (Zhang et al., 2009). At the coarse end of the spectrum (PM₁₀), tThe 72 73 spatial distribution and interannual variation of PM₁₀-concentrations was has been 74 comprehensively studied using a dataset accumulated from 86 Chinese cites (Qu et al., 75 2010). Furthermore, Tthe chemical compositions of PM₁₀ samples were investigated 76 at 16 sites over China, and the result indicated a dominant scattering feature of aerosols in China (Zhang et al., 2012). Existing Network-based studies of network 77 PM_{2.5} observations have, however, been limited to certain seasons in a single year 78 (Cao et al., 2012), and most other research has focused on one or more of the largest cities (He et al., 2001; Wang et al., 2002; Wang et al., 2006; Wei et al., 1999; Yao et 80 al., 2002; Zhao et al., 2009; Zheng et al., 2005). The focus on PM_{2.5} needs to improve,

61

69

71

79

not least because the growing problem of heavy haze problem encourageshas

compelled the Chinese government to pay more-greater attention to PM_{2.5} monitoring and air quality standards. Indeed, Tthe Ministry of Environmental Protection of China issued new ambient air quality standards in 2012, among which the PM_{2.5}

concentration was the first to be included. Subsequently, the construction of a network of national environmental PM_{2.5} monitoring stations network began in 2013.

In this paper, we firstly present a long-term PM₁₀, PM_{2.5} and PM₁ monitoring dataset from 2006 to 2014, based on 24 stations of CAWNET (China Atmosphere Watch Network), operated by the China Meteorological Administration (CMA). The spatial pattern of average PM concentration levels and the relationships among them

are reported. In addition, their seasonal and interannual variations are presented.

2. The near real-time PM dataset

The PM₁₀, PM_{2.5} and PM₁ concentrations were monitored at 24 <u>CAWNET</u> stations of <u>CAWNET</u> from 2006 to 2014 using GRIMM dust monitor EDM 180 instruments with 31 different size channels at a flow rate of 1.2 L/min. The instrument was is designed to measure <u>the particle</u> size distribution and particulate mass, based on a light scattering measurement of individual particles in the sampled air. GRIMM-developed protocols were used to convert the measured size number distribution to a mass concentration consistent with U.S. Environmental Protection Agency protocols for measuring <u>particulate matterPM</u> <u>based onusing the</u> aerodynamic diameter. A

nonvolatile and semi-volatile components are included in the measurement result 104 105 (Grimm and Eatough, 2009). The GRIMM-instruments were calibrated annually using 106 a calibration tower that permitted powder injection (on demand) of aerosol particles in 107 a wide size range of 0.2–30 µm. The operation was fully computer-controlled and 108 permitted access to one to three spectrometers in comparison to one reference "mother 109 unit". The 5-min averaged PM₁₀, PM_{2.5} and PM₁ concentrations were recorded at each station and transported to the CMA information center hourly in near real-time. 110 111 The PM concentration data-results from the GRIMM instruments were compared to the results those from tapered element oscillating microbalances (TEOM) 112 113 instruments in reported in a number of other some studies (Grimm and Eatough, 2009; Hansen et al., 2010). The instruments were in good agreement, e.g. linear regression 114 of with the Grimm vs. the FDMS TEOM data in from Rubidoux (California, USA) 115 yielded a slope of $1.10 \pm \pm 0.05$, with an intercept of $-3.9 \pm 4.2 \mu \text{g m}^{-3}$, and the 116 an uncertainty was of 9.9% (Grimm and Eatough, 2009). The Furthermore, GRIMM 117 and TEOM measurements in Beijing, China have shown thea good close linear 118 119 relationship with TEOM and, suggesting that optical measurements couldcan be used 120 to derive PM_{2.5} and could account for semi-volatile material in aerosols (Sciare et al., 2007; Zhao et al., 2011). 121 The 24 PM observation stations are described detailed in Table 1, and a map of 122 their distribution is given in Figure 1. Most of the stations were located in East China, 123

an area of high population density and fast rapid economic development;

therefore, meaning the PM emitted from human activities was mainly recorded. The

124

125

批注 [LP1]: Please confirm that this information inserted by the editor is correct.

批注 [LP2]: Please check that the changes made here by the editor have not affected the intended meaning.

带格式的:下标

stations were classified as urban/suburban, rural and remote stations, according to their location. Unlike rural stations, remote stations were located in areas far away from regions of strong anthropogenic emissions, and thus natural emissions and long-range transport of anthropogenic air pollution were the main sources of PM at these stations.

3. Results and discussion

3.1. Average PM₁₀, PM_{2.5} and PM₁ levels in China

The averaged PM concentration values are presented in Table 2, and their A distributions bar chart map of the averaged PM concentrations is presented in Figure 12. The highest PM₁₀, PM_{2.5} and PM₁ concentrations were observed at the stations of Xian (135.4, 93.6 and 77.0 µg m⁻³, respectively), Zhengzhou (131.7, 84.8 and 71.0 µg m⁻³, respectively) and Gucheng (127.8, 89.7 and 79.4 µg m⁻³, respectively), which are located in the most polluted areas of the Hua Bei Plain (HBP) and the Guanzhong Plain. Although Gucheng is a rural site, it is located in the rapidly urbanizing urbanization area around Beijing, and is therefore subjected to associated large quantities of air pollution pollutants emissions. These areas were also identified by Zhang et al. (Zhang et al., 2012) as having experienced Region—II—with similar visibility changes and large visibility loss in the past 40 years. The stations all recorded very high coarse and fine PM concentrations, implying high emissions of both primary emitted mineral particles and secondary formatted anthropogenic

concentrations compared with inland cities in on the HBP.

The PM concentrations were also high in northeast China, which is an established industrial base area. The ensemble average values of the five urban stations of Ansan, Shenyang, Benxi, Fushun and Shiping were 88.8, 58.4 and 49.8 µg m⁻³, for PM₁₀, PM_{2.5} and PM₁₂ respectively. Dandong is a rural station with relatively low PM concentrations.

The similarity among the PM values for Chifeng, Erlianhaote and Yulin is due to their location, far from regions of intensive economic development, and is but also strongly affected by sand and dust storms, given their stations are located adjacent proximity to dust source areas. Thus, the average PM₁₀ concentrations were much higher than the PM_{2.5} and PM₁ concentrations at these sites. For example, the average PM₁₀ concentrations were at Chifeng, which is surrounded by sandy land, was 88.0 µg m⁻³, compared with 42.4 and 32.6 µg m⁻³ for PM_{2.5} and PM₁, respectively at Chifeng, which is surrounded by Horqin Sandy Land and Onqin Daga-Sandy Land.

Chengdu, the capital of Sichuan Province, is located in the Sichuan Basin, which is also another highly polluted area, with hHigh aerosol optical depth and low visibility, due to the poor dispersion conditions and heavy local industrial emissions, have been reported for this site (Li et al., 2003; Luo et al., 2001; Zhang et al., 2012), due to the poor dispersion conditions and heavy local industrial emissions. In the present study, Tthe average PM₁₀, PM_{2.5} and PM₁ concentrations were 78.0, 59.5 and 52.7 µg m⁻³, respectively.

There are three stations in the South China area: Panyu, located in Guangzhou City, the capital of Guangdong Province, which is the center of the Pearl River Delta region; Nanning, the capital of Guangxi Province; and Guilin, a famous tourist city, also located in Guangxi Province. The ensemble average PM concentrations of these three sites were 55.8, 43.1 and 38.8 μ g m⁻³ for PM₁₀, PM_{2.5} and PM₁, respectively. Significant visibility loss and relatively high PM₁₀ concentrations have been observed over the middle and lower reaches of the Yangtze River after the 1980s due to the rapid economic development in thethat has taken place in this region (Qu et al., 2010; Zhang et al., 2012). Although there was no urban site available in for this study to characterize help quantify the high PM concentrations in this region, the background conditions and temporal variance could be determined from the rural site data. Dongtan, near Shanghai City, is located on Chongming Island, with where there were low PM concentrations (31.9, 27.4 and 24.8 µg m⁻³ for PM₁₀, PM_{2.5} and PM₁, respectively) due to the substantial influence of clean sea air mass. The ensemble average PM concentrations of for Lushan, Changde and Jinsha were 44.3, 37.2 and 33.6 μ g m⁻³ for PM₁₀, PM_{2.5} and PM₁, respectively. Lhasa, the capital of Tibet Autonomous Region, is located in the center of the Tibetan Plateau at a very high altitude of 3663 m. The PM_{2.5} and PM₁ concentrations in Lhasa were low, with average values of 14.0 and 9.6 µg m⁻³, respectively, because of its relatively small population and few industrial emissions. However, the average PM_{10} concentration was 37.7 µg m⁻³, mainly due to the high amounts of fugitive dust

from dry and bare land and the impacts of regional dust storm events (Chen et al.,

170

171

172

173

174

175

176

177

178

179

180

181

182

183

184

185

186

187

188

189

190

2013). As a result, minerals represent the major component are the main constituent of 192 193 aerosol samples in this area (Zhang et al., 2012). 194 The lowest PM concentration values were observed in the two remote sites of 195 Akadala and Shangri-La. The lower altitude and stronger contribution of soil dust at Akadala (Qu et al., 2009), located in a dry region, leads to higher PM concentrations 196 than at the Shangri-La site. 197 According to the latest air quality standards of China (annual averaged PM₁₀ and 198 PM_{2.5} concentrations of 70 and 35 µg m⁻³), 14 stations reached the PM₁₀ standard, and 199 while only 7 stations, mainly rural and remote stations, reached the PM_{2.5} standard. 200 201 The ratio of substandard (daily averaged PM₁₀ or PM_{2.5} concentrations that exceed the 202 standard values) days to total observation days at each station was calculated using the standard daily averaged PM₁₀ and PM_{2.5} concentrations of 150 and 75 $\mu g m^{-3}$ 203 (Table 2). Substandard days of PM₁₀ and PM_{2.5} represented more than 30% and 50% 204 205 of the total period at the three most polluted sites (Xian, Zhengzhou and Gucheng). The PM_{2.5} substandard days ratios at five other stations (Chengdu, Anshan, Shenyang, 206 207 Benxi and Siping) were also larger than 20%. 208 Average PM₁₀, PM_{2.5} and PM₁ concentrations at urban/suburban stations in this study were 83.6, 56.3 and 48.3 µg m⁻³, respectively. Meanwhile, 7the values were 209 54.8, 36.3 and 30.8 μ g m⁻³ at rural stations, and 11.9, 7.5 and 6.1 μ g m⁻³ at remote 210 stations. They are All values were much higher than the results from other countries. 211 For example, Tthe observed PM concentration data observed in Canada between 1984 212 to 1993 showed the average PM_{2.5} concentrations were was 14.1 and 10.7 $\mu g m^{-3}$ at 213

urban and rural stations, respectively (Brook et al., 1997). THe average PM_{2.5} values were 3 to 13 μg m⁻³-from west to east US observed from across the IMPROVE network in 1993 (most stations were located in rural areas) were 3 to 13 μg m⁻³in-1993 (Eldred et al., 1997). The Observations in Swiss-Switzerland from 1998 to 2001 showed the average PM₁₀ and PM_{2.5} concentrations at urban/suburban stations were of 27.7 and 20.1 μg m⁻³, respectively (Gehrig and Buchmann, 2003). In Austria, in 1998, the Aannual means of mass concentrations of PM₁₀, PM_{2.5} and PM₁ were inthe order of around 28, 20 and 16 µg m⁻³, respectively, at the urban sites, _and a little bitslightly lower at the rural sites in Austria in 1998 (Gomiscek et al., 2004). The average PM₁₀ and PM_{2.5} concentrations were 23.9 and 16.3 µg m⁻³, respectively, for the period 2008—2009 in the Netherlands (Janssen et al., 2013). Between October 2008 and April 2011, Tthe 20 European study areas observation results from of the European ESCAPE project between October 2008 and April 2011 showed PM₁₀ and PM_{2.5} havewith similar spatial patterns; withspecifically, low concentrations in Northern Europe and high concentrations in Southern and Eastern Europe (Eeftens et al., 2012). With the rapid urbanization and corresponding increase in the traffic and energy consumption in India, the ambient concentrations of fine particulate PM in India are also high in India. For example, the measurements at in New Delhi during the August to December 2007 showed the concentrations of PM₁₀, PM_{2.5} and PM₁ were ranged from 20 to 180 μg m⁻³ during the monsoon season, and from 100 to 500 μ g m⁻³ during the winter (Tiwari et al., 2012).

214

215

216

217

218

219

220

221

222

223

224

225

226

227

228

229

230

231

232

233

3.2. Relationships between PM₁₀, PM_{2.5} and PM₁ concentrations

235

236

237

238

239

240

241

242

243

244

245

246

247

248

249

250

251

252

253

254

255

256

257

The squared correlation coefficient (R^2) values of the linear fit between PM₁₀ and PM_{2.5} and between PM₁ and PM_{2.5} are given in Table 2. Higher values indicate that the two PM size bins were closer matched in terms of their sourcesmay have more identical source characteristics. At most stations, the R² values between PM₁ and PM_{2.5} were higher than the values between PM_{2.5} and PM_{10.5} This is because PM₁ and PM_{2.5} both belong to fine particle size bins, which are normally emitted from the same sources. For example, the R^2 values were 0.7857 between PM_{2.5} and PM₁₀, and 0.9689 between PM₁ and PM_{2.5}, at Gucheng (Figure 3). Correlation coefficient analysis is sensitive to outliers, and thus sand storm events may have impacted upon the results considerably, due to abnormally high concentration values. There were four strong dust storm event days at Akdala in 2012, on April 21 and 22, and May 9 and 20, which resulted in the four outliers points shown in Figure 24a, and the low R^2 value of 0.5346 between PM₁ and PM_{2.5}. The value increased to 0.9406 when the four outliers points were removed (Figure 24b). Similar results were also observed at Yulin and Erlianhaote around dust storm source regions (Table 2). The average values of the daily PM_{2.5}/PM₁₀ and PM₁/PM_{2.5} ratios are listed in Table 2. The spatial distribution $\frac{\text{map}}{\text{of}}$ of the average PM_{2.5}/PM₁₀ ratios (Figure 3a5) shows lower ratio-values in northern China, influenced by Asian sand and dust storms (Wang et al., 2008; Zhang et al., 2003). The values were also influenced by fugitive dust due to the low precipitation amounts in northern China, especially at Lhasa, Erlanhaote, Yulin and Chifeng, with ratios values of less than 0.6. The ratios values at the stations in northeast China were between 0.6 and 0.7, except at Dandong where

the value was 0.71. The values were also low at Zhengzhou and Akdala, at 0.68 and 0.67, respectively. The highest ratio value was 0.9 at Dongtan, and the other stations with ratios values higher than 0.8 were Chengdu, Changde, Guilin, Jinsha and Lushan. The values were between 0.7 and 0.8 at other stations. The PM₁/PM_{2.5} ratios (Figure 3b) showed a similar spatial distribution, but the values were higher than PM_{2.5}/PM₁₀. The lowest ratio value of 0.6 was also observed at Lhasa, and the values at most stations in southern China were greater than or equal to 0.9.

3.3. Seasonal variations

The seasonal variations of PM₁₀ concentrations (Figure 46a) show that winter and spring were the most polluted seasons at all sites except Lushan, where the highest value was observed in autumn. This result is consistent with a previous study of PM₁₀ variation across China from 2000 to 2006 (Qu et al., 2010). The higher winter concentrations were caused by higher emissions during the cold season from heating, and more stagnant weather conditions with a lower planetary boundary layer. The opposite conditions and more precipitation due to the summer monsoon resulted in the lowest PM₁₀ concentration values in summer. Spring is the dust storm season in East Asia (Qian et al., 2004; Wang et al., 2008; Zhou and Zhang, 2003), which leads to high PM₁₀ concentrations in dust source regions and downwind areas in northern China. For example, the PM₁₀ concentrations in spring were much higher than other seasons at the dust source sites of Yulin and Erlianhaote.

For PM_{2.5}, winter was still the most polluted season of at most sites, while the contribution of spring decreased substantially in northern China (Figure <u>4</u>6b). This

trend can be further observed from the PM_1 distribution (Figure 6c); hence, the average PM_1 concentration of in spring was lowest at Yulin, Xian, Zhengzhou, Gucheng and Benxi. The seasonal variation patterns were very similar for PM_{10} , $PM_{2.5}$ and PM_1 at the sites in southern China.

A spatial distribution map of the seasonal average $PM_{2.5}/PM_{10}$ ratios is given in Figure 6d. For the reasons given above, lower $PM_{2.5}/PM_{10}$ ratios were observed in spring at the northern China sites, while the seasonal variation was not significant at the southern China sites.

3.4 Interannual variations

The interannual variations of PM_{2.5} at various stations are-is presented in Figure 57. Significant decreasing trends were observed at the HBP stations of Zhengzhou and Gucheng (Figure 57a). The annual averaged PM_{2.5} concentrations decreased from 123.4 to 65.2 μg m⁻³ at Zhengzhou, and from 101.0 to 69.1 μg m⁻³ at Gucheng, during 2006–2014. At Zhengzhou, the lowest value of 63.7 μg m⁻³ occurred in 2012, and this level was maintained in subsequent years; however, at Gucheng, the value increased suddenly in 2012 to 95.1 μg m⁻³ and then declined rapidly during 2013 and 2014. At Xian, the annual averaged PM_{2.5} concentrations decreased from 2006 to 2009, increased until 2011, and then decreased again until 2014 (Figure 57a).

For the stations in northeast China, a significant increasing trend of the PM_{2.5} concentrations was observed at Shenyang and Benxi from 2006 to 2013, followed by a decrease in 2014 (Figure 57b). The peak value at Shenyang was especially high in 2013 at 123.1 μg m⁻³, while the values were less than 60 μg m⁻³ in the other years.

The highest values were observed in 2009 at Anshan and Dandong, but the lowest values was were in 2014 at Anshan and 2010 at Dandong. A general decreasing trend was observed at Siping, with a few fluctuations. At Fushun, the value decreased from 2006 to 2011 and then increased to 2013, followed by a slight decrease in 2014.

For the stations along the middle and lower reaches of the Yangtze River, a common trend was a clearly lower PM_{2.5} value in 2014 than in 2013, but the general variation trend was not significant (Figure <u>5</u>7c). A peak value of 33.7 μg m⁻³ was observed in 2012 at Dongtan, followed by a decrease to 24.12 μg m⁻³ over the subsequent two years. At Jinsha and Changde, the highest value was in 2013, while it was in 2009 at Lushan.

For the stations in southern China, a general decreasing trend was observed, with obvious fluctuations (Figure $\underline{57}$ d). Panyu is a typical station in the centre of the Pearl River Delta economic area of China. The PM_{2.5} value decreased from 64.6 μ g m⁻³ in 2006 to 41.6 μ g m⁻³ in 2014, and the lowest value was 36.4 μ g m⁻³ in 2010. A similar trend was observed in Gulin, with a stronger fluctuation from 2010 to 2012. At Nanning, a peak value occurred in 2010 and the lowest value of 28.5 μ g m⁻³ was observed in 2012.

The general observations of Generally, the PM_{10} and PM_1 interannual variation trends were similar to the that of $PM_{2.5}$ at most stations. For example, a similar trend and fluctuations were observed at the stations presented in Figure 8 and Figure 7a. A difference in the trend was observed at Zhengzhou from 2013 to 2014, with a significant increasing trend of PM_{10} and decreasing trend of PM_1 .

3.5 Diurnal variations

324

325

326

327

328

329

330

331

332

333

334

335

336

337

338

339

340

341

342

343

344

345

346

The average diurnal variations of PM_{2.5} at various stations are is presented in Figure 79. Pronounced diurnal variations of PM_{2.5} were was observed in at most urban sites, with an obvious morning peak at around 7:00 to 8:00 a.m. and an afternoon valley between 2:00 and 4:00 p.m. At some stations, an evening peak can could be recognized at around 7:00 to 9:00 p.m. (Siping, Benxi, Fushun, Anshan, Guilin and Panyu) or midnight (Gucheng, Xian). This bimodal pattern was also observed in Beijing city (Zhao et al., 2009). - The A unimodal pattern, without an evening peak, can-could be identified at some other stations (Zhengzhou, Shengyang and Nanning). In urban areas, the morning and evening peaks are contributed to by enhanced anthropogenic activity during rush hour, and the afternoon valley is mainly due to a higher atmospheric mixing layer, which is benefit beneficial for air pollution diffusion. Panyu station is on top of a 140 m hill at the edge of Guangzhou city, so aged and mixing aerosols were observed with a weakly urban diurnal variation pattern. Similar with to Panyu station, the rural stations along the middle and lower reaches of the Yangtze River have showed no typical urban diurnal variation pattern (Figure 79c). The diurnal variations in PM₁ and PM₁₀ concentrations arewas similar to that of PM_{2.5} at most stations.

带格式的: 下标 **带格式的:** 下标 **带格式的:** 下标

$\textbf{3.6} \; \underline{\textbf{Emission and m}} \\ \textbf{Meteorological influences}$

PM loadings wereare controlled by both emissions and meteorological conditions. Even Mmineral dust emissions from deserts and VOCvolatile organic compound (VOC) emissions from vegetation are even controlled by meteorological factors, e.-g., wind speed and temperature. The major source of air pollution in China

347	was contributed fromis anthropogenic emissions, especially with the rapid economic
348	development that has taken place in recent years with rapidly economic development.
349	In this caseAs such, the averaged PM concentration pattern wasis determined
350	decided largely by emissions, but meteorological factors also play an important role by
351	impactionaffecting pollutants diffusion and deposition.
352	The distributions of the Aanthropogenic emissions distributions of black carbon
353	(BC), PM _{2.5} , SO ₂ and NO ₂ in 2010, based on the HTAP v2 harmonized emissions
354	database (http://iek8wikis.iek.fz-juelich.de/HTAPWiki/WP1.1), wereare presented in
355	#Figure 8. The emissions data offor the East Asia domain were contributed
356	fromsupplied by the MICS-Asia project. The spatial distributions of the species
357	haveshow a consistent pattern with the high emissions regions of the HBP,
358	Guanzhong Plain, Sichuan Basin, middle and lower reaches of the Yangtze River,
359	Pearl River Delta region, and the industr yial region of northeast China, which is
360	generally similar withto the PM loadings pattern infor China (Figure 1). For example,
361	most PM pollutant stations subjected to PM pollution are located in the highest
362	emissions region of the HBP. ItThis indicateds that averaged PM loadings wereare
363	controlled by the quantity of anthropogenic emissions amount in midcentral-eastern
364	China.
365	The emission-trends in emissions for China during 2005—2010 (Wang et al.,
366	2014b) showed thethat emissions of SO ₂ and PM _{2.5} in East Asia decreased by 15%
367	and 12 %, respectively, meanwhile the emissions of NOx and non-methane NMVOCs
368	increased by 25% and 15-%, respectively. Driven by emission-changes in emissions,

	7) () () () () () () () () () ({ 带格式的: 下标
369	PM _{2.5} concentrations decreased by 2–17 μg m ⁻³ inover most of the North China Plain,	
370	the Yangtze River Delta and the Pearl River Delta (Zhao et al., 2013). Which This	
371	could explain the general decreaseing trend efwith respect to PM during 2006—2010	
372	at most stations (Figure 5). The Sepatial distributions of emissions differences	
373	between 2010 and 2008 offor BC, PM _{2.5} , SO ₂ and NO ₂ wereare plotted in fFigure 9,	
374	based on the HTAP_v2 emission dataset, so the PM vs emission variation in different	
375	regions could be studied. BC emissions decreased from 2008 to 2010 in most regions	
376	of east China, except the provinces of Hebei, Shanxi, Hubei, Jiangxi and Inter	
377	Mongolia-provinces (Figure 9a). More areas of China showed a reduction of PM _{2.5}	带格式的: 下标
378	emissions, except Shanxi and Hubei provinces (Figure 9b). The SO ₂ emissions	带格式的: 下标
379	difference (Figure 9c) has showed a similar pattern withto that of BC but with an	
380	increaseing trend identified apparent in northeast China. NOx emissions increased in	
381	most regions of midcentral-eastern China, except in the provinces of Guangdong,	
382	Zhejiang and Taiwan-provinces (Figure 9d). †This trend was driven by the rapid	
383	increase growth of industry and transportation, and combined with inadequate control	
384	strategies (Wang et al., 2014b).	
385	Although there isare no published emissions data after 2010, it wasis believed	
386	that the emissions werehave to a certain extent been greatly controlled well aftersince	
387	the end of 2013, with the issuearrival of China's "Action Plan for the Control of Air	
388	Pollution" document. HThis eancould explain the general decreaseing trend infor the	
389	year 2014 at most stations (fFigure 5).	
390	Central-eastern China experienced severe haze events in January 2013, with a	

regionally stable planetary boundary layer and low mixing height (Wang et al., 2014a). The daily averaged PM_{2.5} concentrations and the meteorological factors of wind speed and relative humidity data infor this period at Zhengzhou, Shenyang and Nanning were are plotted in Figure 10. Zhengzhou is located in this haze region, with and <u>experienced</u> very high PM_{2.5} concentrations, especially from Jan. 6 to 15. The wind speed variation was negatively related with PM_{2.5} concentrations. The rapid increasing increase of in PM_{2.5} concentrations from Jan. 1 to 6 was corresponding ed with the rapid decreasing decrease of in wind speed at during the same period. Also, the big strong wind speed in on Jan. 24 resulted in the low PM_{2.5} concentration. Shenyang and Nanning are not located in this severe haze region, but still suffered some fine particle pollutantPM days in thisthat month. The A negative correlation 带格式的:下标 between PM_{2.5} and wind speed were was also observed at Shengyang and Nanning. In general, relative humidity (RH) variation was positively related with the PM_{2.5} concentrations if no precipitation occurred. Otherwise, the high RH with precipitation correspondences corresponded to low PM concentrations due to wet deposition. For In terms of interannual variation, the negative correlation between PM_{2.5} concentrations and wind speed, and the positive correlation between PM2.5 concentrations and relative humidity, cannot could not be well identified (Figure 11). 带格式的:下标 Although thea generally similar variation trend offor the PM₁₀ concentrations and relative humidity was observed inat Zhengzhou, but it cannot bethis was not found 带格式的:下标 inat other stations. The PM_{2.5} concentration in 2014 was lower than in 2013, but the relative humidity was much higher and the wind speed was much lower. The

391

392

393

394

395

396

397

398

399

400

401

402

403

404

405

406

407

408

409

410

411

interannual variation of PM concentrations eannot could not be explained justeonsideringsolely by meteorological factors, although the a recent model simulation
for years of the period 2004—2012 with fixed anthropogenic emissions at fixed at the
values for the year 2006 indicated that the variations of in meteorological fields were
found to dominant dominated the interannual variations of in aerosols in China (Mu
and Liao, 2014). For IL ong-term-variation, both emissions and meteorological factors
play important roles; on it It reveals the emission variation possibly dominates the
long term PM concentration trend; mean while in the short-term, meteorological
factors play a leading role—at least in the absence of during a short period without
significant emissions changes.

4 Conclusion

Spatial and temporal trends in PM pollution were examined using PM₁₀, PM_{2.5} and PM₁ concentration data at 24 stations from 2006 to 2014. Relatively high PM concentrations were observed at most stations. There were 14 stations that reached the PM₁₀ annual air quality standard, but only 7 stations, mostly rural and remote stations, reached the PM_{2.5} annual air quality standard of China. The highest PM concentrations were observed at the stations in on the HBP and Guanzhong Plain. In addition, the percentage value of substandard days of PM_{2.5} was greater than 50%, indicating very serious air pollution in these regions. PM pollutants are also a serious problem in the industrial regions of northeast China and the Sichuan basin. The PM concentrations were relatively lower in the southern areas of China, but the averaged PM_{2.5} concentration was still higher than the national standard.

As Given they are both fine particles, PM₁ and PM_{2.5} were more closely correlated than PM_{2.5} and PM₁₀. The correlations were sensitive to the effect of outlier data at those stations heavily impacted by dust storm events. More dust aerosol was observed in northern China, and thus the PM_{2.5}/PM₁₀ ratios increased from less than 0.6 to around 0.9 when moving from north to south China.

Pronounced seasonal variations were observed at most stations, with the highest concentrations in winter and lowest concentrations in summer. PM₁₀ concentrations

concentrations in winter and lowest concentrations in summer. PM₁₀ concentrations were also high in spring, due to the contribution of dust storm events, especially at those stations near to dust source regions. For PM_{2.5} and PM₁, spring was a relatively low concentration season, especially at the stations in northern China. Also, low PM_{2.5}/PM₁₀ ratios were observed in spring in northern China.

An interannual decreasing trend was observed in the HBP and southern China from 2006 to 2014, but an increasing trend occurred at some stations in northeast China, and no significant trend could be found over the middle and lower reaches of the Yangtze River. Annual-averaged PM concentrations were lower in 2014 than 2013 at most stations, which may indicate an improvement in air quality following the "Action Plan for the Control of Air Pollution" document issued by the Chinese government in September 2013.

Bimodal and unimodal diurnal variation patterns were identified at urban stations.

The A negative correlation between PM concentrations and wind speed was found in a for the short-term-period, but variations in emissions variation must to be considered for long-term trend analysis analyses, especially in rapidly developing countries.

This network-based observation dataset provides the longest continuous record of fine particle concentrations in China, but it features a limited number of stations and an uneven spatial distribution. Importantly, there is no representative city site in the Yangtze River delta region, which is considered an important haze area in China. The emissions sources and meteorological factors influencing PM spatial and temporal patterns in China still require further study.

Δckn	owle	daei	ments
ACNII	OWIE	uue	HIGHLS

This work was supported by grants from the National Key Project of Basic Research

(2014CB441201), the National Natural Science Foundation of China (41275167) and

the Chinese Academy of Meteorological Sciences (2013Z007). It was also supported

by the Climate Change Collaborative Innovation Center and the CMA Innovation

Team of Haze-fog Observation and Forecasts.

References:

- Brook, J. R., Dann, T. F., and Burnett, R. T.: The relationship among TSP, PM10, PM2.5, and Inorganic constituents of atmospheric particulate matter at multiple Canadian locations. J. Air Waste
- 474 Manage. Assoc. 47, 2-19, 1997.
- Cao, J.-J., Shen, Z.-X., Chow, J. C., Watson, J. G., Lee, S.-C., Tie, X.-X., Ho, K.-F., Wang, G.-H., and Han, Y.-M.: Winter and Summer PM2.5 Chemical Compositions in Fourteen Chinese Cities. J.
- 477 Air Waste Manage. Assoc. 62, 1214-1226, 2012.
- Chen, S., Huang, J., Zhao, C., Qian, Y., Leung, R., and Yang, B.: Modeling the transport and radiative
 forcing of Taklimakan dust over the Tibetan Plateau: A case study in the summer of 2006. J.
- 480 Geophys. Res. 118, 797-812, 2013.
- Delfino, R. J., Sioutas, C., and Malik, S.: Potential role of ultrafine particles in associations between airborne particle mass and cardiovascular health. Environ. Health Persp. 113, 934-946, 2005.
- 483 Eeftens, M., Tsai, M.-Y., Ampe, C., Anwander, B., Beelen, R., Bellander, T., Cesaroni, G., Cirach, M.,
- 484 Cyrys, J., Hoogh, K. d., Nazelle, A. D., Vocht, F. d., Declercq, C., Dedele, A., Eriksen, K.,
- 485 Galassi, C., Grazuleviciene, R., Grivas, G., Heinrich, J., Hoffmann, B., Iakovides, M.,
- 486 Ineichen, A., Katsouyanni, K., Korek, M., Krämer, U., Kuhlbusch, T., Lanki, T., Madsen, C.,
- 487 Meliefste, K., Mölter, A., Moslerm, G., Nieuwenhuijsen, M., Oldenwening, M., Pennanen, A.,
- 488 Probst-Hensch, N., Quass, U., Raaschou-Nielsen, O., Ranzi, A., Stephanou, E., Sugiri, D.,
- Udvardy, O., Vaskövi, É., Weinmayr, G., Brunekreef, B., and Hoek, G.: Spatial variation of
- 490 PM_{2.5}, PM₁₀, PM_{2.5} absorbance and PMcoarse concentrations between and within 20 European
- 491 study areas and the relationship with NO₂ Results of the ESCAPE project. Atmos. Environ.
- 492 62, 303-317, 2012.
- Eldred, R. A., Cahill, T. A., and Flocchini, R. G.: Composition of PM2.5 and PM10 aerosols in the IMPROVE network. J. Air Waste Manage. Assoc. 47, 194-203, 1997.
- Gehrig, R., and Buchmann, B.: Characterising seasonal variations and spatial distribution of ambient
 PM10 and PM2.5 concentrations based on long-term Swiss monitoring data. Atmos. Environ.
- 497 37, 2571-2580, 2003.
- Gomiscek, B., Hauck, H., Stopper, S., and Preining, O.: Spatial and temporal variations of PM₁, PM_{2.5},
 PM₁₀ and particle number concentration during the AUPHEP project. Atmos. Environ. 38,
- 500 3917-3934, 2004.
- Grimm, H., and Eatough, D. J.: Aerosol Measurement: The Use of Optical Light Scattering for the
 Determination of Particulate Size Distribution, and Particulate Mass, Including the
- Semi-Volatile Fraction. J. Air Waste Manage. Assoc. 59, 101-107, 2009.
- Hansen, J. C., Woolwine, W. R., Bates, B. L., Clark, J. M., Kuprov, R. Y., Mukherjee, P., Murray, J. A.,
- Simmons, M. A., Waite, M. F., Eatough, N. L., Eatough, D. J., Long, R., and Grover, B. D.:
- Semicontinuous PM_{2.5} and PM₁₀ mass and composition measurements in Lindon, Utah, during winter 2007. J. Air & Waste Manage. Assoc. 60, 346-355, 2010.
- Haywood, J., and Boucher, O.: Estimates of the direct and indirect radiative forcing due to tropospheric aerosols: A review. Rev. Geophys. 38, 513-543, 2000.
- He, K., Yang, F., Ma, Y., Zhang, Q., Yao, X., Chan, C. K., Cadle, S., Chan, T., and Mulawa, P.: The characteristics of PM2.5 in Beijing, China. Atmos. Environ. 35, 4959-4970, 2001.
- 512 Houthuijs, D., Breugelmans, O., Hoek, G., Vaskövi, É., Miháliková, E., Pastuszka, J. S., Jirik, V.,
- 513 Sachelarescu, S., Lolova, D., Meliefste, K., Uzunova, E., Marinescu, C., Volf, J., Leeuw, F. d.,

- Wiel, H. v. d., Fletcher, T., Lebret, E., and Brunekreef, B.: PM10 and PM2.5 concentrations in
- Central and Eastern Europe: results from the Cesar study. Atmos. Environ. 35, 2757-2771,
- 516 2001.
- Janssen, N. A. H., Fischer, P., Marra, M., Ameling, C., and Cassee, F. R.: Short-term effects of PM_{2.5},
- PM_{10} and $PM_{2.5-10}$ on daily mortality in the Netherlands. Sci. Total Environ. 463-464, 20-26,
- 519 2013.
- 520 Li, C., Mao, J., Lau, K.-H. A., Chen, J.-C., Yuan, Z., Liu, X., Zhu, A., and Liu, G.: Characteristics of
- 521 distribution and seasonal variation of aerosol optical depth in eastern China with MODIS
- 522 products. Chinese Sci. Bull. 48, 2488-2495, 2003.
- $523 \qquad Luo, Y., Lu, D., Zhou, X., Li, W., and He, Q.: Characteristics of the spatial distribution and yearly$
- variation of aerosol optical depth over China in last 30 years. J. Geophys. Res. 106,
- 525 14501-14513, 2001.
- Malm, W. C., Schichtel, B. A., Pitchford, M. L., Ashbaugh, L. L., and Eldred, R. A.: Spatial and
- 527 monthly trends in speciated fine particle concentration in the United States. J. Geophys. Res.
- 528 109, D03306, doi:10.1029/2003JD003739, 2004.
- 529 Mu, Q., and Liao, H.: Simulation of the interannual variations of aerosols in China: role of variations in
- 530 meteorological parameters. Atmos. Chem. Phys. 14, 9597-9612, 2014.
- Oberdörster, G., Oberdörster, E., and Oberdörster, J.: Nanotoxicology: An Emerging Discipline
- Evolving from Studies of Ultrafine Particles. Environ. Health Persp. 113, 823-839, 2005.
- Pope III, C. A., and Dockery, D. W.: Health effects of fine particulate air pollution: Lines that connect.
- J. Air Waste Manage. Assoc. 56, 709-742, 2006.
- Qian, W. H., Tang, X., and Quan, L. S.: Regional characteristics of dust storms in China. Atmos.
- 536 Environ. 38, 4895-4907, 2004.
- Qu, W.-J., Zhang, X.-Y., Arimoto, R., Wang, Y.-Q., Wang, D., Sheng, L.-F., and Fu, G.: Aerosol
- 538 background at two remote CAWNET sites in western China. Sci. Total Environ. 407,
- 539 3518-3529, 2009.
- Qu, W. J., Arimoto, R., Zhang, X. Y., Zhao, C. H., Wang, Y. Q., Sheng, L. F., and Fu, G.: Spatial
- distribution and interannual variation of surface PM_{10} concentrations over eighty-six Chinese
- 542 cities. Atmos. Chem. Phys. 10, 5641-5662, 2010.
- Ramanathan, V., Crutzen, P. J., Kiehl, J. T., and Rosenfeld, D.: Aerosols, climate, and the hydrological
- 544 cycle. Science 294, 2119-2124, 2001.
- 545 Sciare, J., Cachier, H., Sarda-Este've, R., Yu, T., and Wang, X.: Semi-volatile aerosols in Beijing (R.P.
- China): Characterization and influence on various PM2.5 measurements. J. Geophys. Res. 112,
- 547 D18202, doi:10.1029/2006JD007448, 2007.
- 548 Tiwari, S., Chate, D. M., Pragya, P., Ali, K., and Bisht, D. S.: Variations in Mass of the PM10, PM2.5
- and PM1 during the Monsoon and Winter at New Delhi. Aerosol and Air Quality Research 12,
- 550 20-29, 2012.
- Wang, G., Huang, L., Gao, S., Gao, S., and Wang, L.: Measurements of PM₁₀ and PM_{2.5} in urban area of
- Nanjing, China and the assessment of pulmonary deposition of particle mass. Atmos. Environ.
- 553 48, 689-695, 2002.
- Wang, H., Xu, J., Zhang, M., Yang, Y., Shen, X., Wang, Y., Chen, D., and Guo, J.: A study of the
- meteorological causes of a prolonged and severe haze episode in January 2013 over
- central-eastern China. Atmospheric Environment 98, 146-157, 2014a.
- 557 Wang, S. X., Zhao, B., Cai, S. Y., Klimont, Z., Nielsen, C. P., Morikawa, T., Woo, J. H., Kim, Y., Fu, X.,

- Xu, J. Y., Hao, J. M., and He, K. B.: Emission trends and mitigation options for air pollutants
 in East Asia. Atmos. Chem. Phys. 14, 6571-6603, 2014b.
- Wang, X., Bi, X., Sheng, G., and Fu, J.: Chemical composition and sources of PM10 and PM2.5 aerosols in Guangzhou, China. Environ. Monit. Assess. 119, 425-439, 2006.
- Wang, Y. Q., Zhang, X. Y., Gong, S. L., Zhou, C. H., Liu, H. L., Niu, T., and Yang, Y. Q.: Surface
 observation of sand and dust storm in East Asia and its application in CUACE/Dust. Atmos.
 Chem. Phys. 8, 545-553, 2008.
- 565 Watson, J.: Visibility: Science and Regulation. J. Air Waste Manage. Assoc. 52, 628-713, 2002.
- Wei, F., Teng, E., Wu, G., Hu, W., Wilson, W. E., Chapman, R. S., Pau, J. C., and Zhang, J.: Ambient
 concentrations and elemental composition of PM₁₀ and PM_{2.5} in four Chinese cities. Environ.
 Sci. Technol. 33, 4188-4193, 1999.
- Yao, X., Chan, C. K., Fang, M., Cadle, S., Chan, T., Mulawa, P., He, K., and Ye, B.: The water-soluble
 ionic composition of PM2.5 in Shanghai and Beijing, China. Atmos. Environ. 36, 4223-4234,
 2002.
- Zhang, Q., Streets, D. G., Carmichael, G. R., He, K. B., Huo, H., Kannari, A., Klimont, Z., Park, I. S.,
 Reddy, S., Fu, J. S., Chen, D., Duan, L., Lei, Y., Wang, L. T., and Yao, Z. L.: Asian emissions
 in 2006 for the NASA INTEX-B mission. Atmos. Chem. Phys. 9, 5131-5153, 2009.
- Zhang, X. Y., Gong, S. L., Zhao, T. L., Arimoto, R., Wang, Y. Q., and Zhou, Z. J.: Sources of Asian dust
 and role of climate change versus desertification in Asian dust emission. Geophys. Res. Lett.
 30, 2272 10.1029/2003GL018206, 2003.
- Zhang, X. Y., Wang, Y. Q., Niu, T., Zhang, X. C., Gong, S. L., Zhang, Y. M., and Sun, J. Y.:
 Atmoshperic aerosol compositions in China: spatial/temporal variability, chemical signature,
 regional haze distribution and comparisons with global aerosols. Atmos. Chem. Phys. 12,
 779-799, 2012.
- Zhao, B., Wang, S., Dong, X., Wang, J., Duan, L., Fu, X., Hao, J., and Fu, J.: Environmental effects of
 the recent emission changes in China: implications for particulate matter pollution and soil
 acidification. Environ. Res. Lett. 8, 024031, doi:10.1088/1748-9326/8/2/024031, 2013.
- Zhao, X., Zhang, X., Pu, W., Meng, W., and Xu, X.: Scattering properties of the atmospheric aerosol in Beijing, China. Atmos. Res. 101, 799-808, 2011.
- Zhao, X., Zhang, X., Xu, X., Xu, J., Meng, W., and Pu, W.: Seasonal and diurnal variations of ambient
 PM2.5 concentration in urban and rural environments in Beijing. Atmos. Environ. 43,
 2893-2900, 2009.
- Zheng, M., Salmon, L. G., Schauer, J. J., Zeng, L., Kiang, C. S., Zhang, Y., and Cass, G. R.: Seasonal
 trends in PM2.5 source contributions in Beijing, China. Atmos. Environ. 39, 3967-3976, 2005.
- Zhou, Z. J., and Zhang, G. C.: Typical severe dust storms in northern China during 1954-2002. Chinese
 Sci. Bull. 48, 2366-2370, 2003.

595

Stations	Latitude	Longitude	Altitude	Start	Description		
	(°N)	N) (° E) (m) Time					
Zhengzhou	34.78	113.68	99.0	1/2006	Urban, in the center of Zhengzhou city, 56 m building.		
Chengdu	30.65	104.04	496.0	3/2006	Urban, in the center of Chengdu		
Chenguu					city, 91 m building.		
Xian	34.43	108.97	363.0	1/2006	Urban, in northern margin of Xian city, 20 km north of center of Xian city, 4 m sampling container.		
Nanning	22.82	108.35	84.0	1/2006	Urban, in Nanning city, 140 m hill.		
Anshan	41.05	123.00	78.3	10/2007	Urban, in Anshan city, 10 m building.		
Shenyang	41.76	123.41	110.0	10/2007	Urban, in Shenyang city, 15m building.		
Benxi	41.19	123.47	185.4	10/2007	Urban, in Benxi city, 12 m building.		
Fushun	41.88	123.95	163.0	10/2007	Urban, in Fushun city, 10 m building.		
Qingdao	36.07	120.33	77.2	3/2007	Urban, in Qingdao city, top of Fulongshan hill.		
Lhasa	29.67	91.13	3663.0	1/2006	Urban, in Lhasa city, 7 m building.		
Siping	43.18	124.33	165.4	3/2007	Urban, in Siping city, 4 m sampling container.		
Panyu	23.00	113.35	5.0	1/2006	Suburban, in Panyu district of Guangzhou city, 140 m hill.		
Gucheng	39.13	115.80	15.2	1/2006	Suburban, 38 km southwest of Baoding city, within area of rapid urbanization, 8 m building.		
Chifeng	42.27	118.97	568.0	3/2007	Rural, suburbs of Chifeng city, 4 m sampling container.		
Dandong	40.05	124.33	13.9	3/2007	Rural, suburbs of Dandong city, 4 m sampling container.		
Erlianhaote	43.65	111.97	965.9	3/2007	Rural, suburbs of Erlianhaote city, 4 m sampling container.		
Yulin	38.43	109.20	1135.0	1/2006	Rural, 10 km north of Yulin city, at the southeastern edge of Mu Us desert.		
Jinsha	29.63	114.20	416.0	4/2006	Rural, 105 km north of Wuhan city, 8 m building.		
Guilin	25.32	110.30	164.4	1/2006	Rural, north margin of Guilin		

					city, meteorological observation
					field.
Lushan	29.57	115.99	1165.0	1/2006	Rural, Kuniubei peak of Mount
	29.37	113.99	1105.0		Lu.
Changde	29.17	111.71	563.0	1/2006	Rural, 18 km northwest from
					Changde city, 8 m building.
Danatan	31.50	121.80	10.0	5/2009	Rural, east of Chongming island
Dongtan					near Shanghai.
Akdala	47.12	87.97	562.0	9/2006	Remote, 55 km west of Fuhai
		87.97		9/2006	county, 10 m building.
Shangri-La	28.02	00.72	2590.0	10/2006	Remote, 12 km northeast of
		99.73	3580.0	10/2006	Shangri-La county.

each station.									
	Av	eraged P	M						
Stations	COI	concentrations			CD	PM _{2.5}	DN# /	D ² (D) 4	D ² (D) 4
	$(\mu g m^{-3})^a$			SB ratio (PM ₁₀) ^b	SB ratio (PM _{2.5}) ^b	PM _{2.5} /PM ₁₀	PM ₁ /	R^2 (PM _{2.5}	`
	PM_{10}	PM _{2.5}	PM_1	(PM ₁₀)	(PM _{2.5})	/PIVI ₁₀	PM _{2.5}	to PM ₁₀)	to PM _{2.5)}
Zhengzhou	131.7 (84.4)	84.8 (47.4)	71.0 (40.5)	0.31	0.51	0.68	0.84	0.68	0.91
Chengdu	78.0 (72.5)	59.5 (42.2)	52.7 (35.4)	0.11	0.27	0.83	0.91	0.76	0.94
Xian	135.4 (97.3)	93.6 (67.3)	77.0 (55.6)	0.34	0.52	0.73	0.83	0.77	0.93
Nanning	51.2 (56.3)	38.4 (24.7)	34.9 (22.2)	0.01	0.08	0.77	0.91	0.52	0.97
Anshan	97.8 (62.9)	60.9 (42.9)	52.3 (39.0)	0.17	0.25	0.65	0.85	0.72	0.98
Shenyang	85.0	59.1	50.8	0.11	0.25	0.69	0.85	0.88	0.97
Benxi	(58.2) 97.6	(42.7) 66.7	(36.7) 54.8	0.13	0.30	0.69	0.82	0.81	0.94
Fushun	(57.4) 80.3	(45.0) 50.1	(36.4) 42.8	0.07	0.17	0.66	0.85	0.64	0.97
Qingdao	(54.2) 64.8	(31.7) 47.3	(28.3) 41.1	0.05	0.17	0.76	0.86	0.76	0.95
Lhasa	(52.1) 37.7	(34.0) 14.0	(30.5) 9.6	0.01			0.66		
	(30.8) 58.7	(10.7) 44.5	(8.6) 39.7		0.00	0.40		0.72	0.94
Panyu	(33.1) 127.8	(24.4) 89.7	(22.1) 79.4	0.02	0.12	0.77	0.89	0.95	0.98
Gucheng	(75.1)	(53.0)	(48.8)	0.31	0.54	0.71	0.87	0.79	0.97
Siping	83.3 (54.3)	55.4 (35.2)	48.5 (32.5)	0.10	0.22	0.68	0.86	0.71	0.96
Chifeng	88.0 (68.9)	42.4 (33.1)	32.6 (27.8)	0.17	0.14	0.51	0.75	0.72	0.92
Dandong	66.8 (44.0)	45.6 (24.8)	39.3 (21.3)	0.03	0.11	0.71	0.86	0.64	0.90
Erlianhaote	49.1 (80.2)	22.0 (22.6)	15.9 (14.7)	0.03	0.03	0.51	0.72	0.71	0.61
Yulin	66.6 (67.1)	31.2 (21.0)	22.4 (15.9)	0.06	0.03	0.54	0.72	0.54	0.61
Jinsha	42.0 (38.6)	33.6 (24.1)	30.5 (21.9)	0.01	0.06	0.85	0.90	0.63	0.89
GuiLin	57.6	46.5	41.7	0.04	0.15	0.85	0.90	0.70	0.96

	(50.5)	(30.8)	(27.1)						
Lushan	45.4	37.8	33.2	0.01	0.09	0.85	0.86	0.91	0.95
	(32.7)	(27.9)	(26.7)	0.01					
Changde	45.7	40.3	37.0	0.01	0.12	0.89	0.91	0.93	0.96
	(33.8)	(29.1)	(27.5)	0.01					
Dongtan	31.9	27.4	24.8	0.01	0.06	0.90	0.90	0.92	0.96
	(34.0)	(25.9)	(23.8)						
Akdala	17.1	9.8	7.7	0.00	0.00	0.67	0.79	0.80	0.53
	(57.6)	(13.7)	(6.9)	0.00					
Shangri-La	6.8	5.2	4.5	0.00	0.00	0.76	0.81	0.94	0.99
	(6.3)	(5.3)	(5.0)	0.00				0.24	

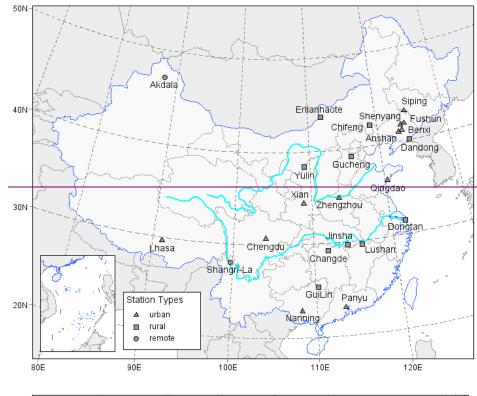
^aArithmetic mean value with standard deviation in parentheses.

 $^{^{}b}$ The ratio of substandard days (daily averaged PM_{10} or $PM_{2.5}$ concentrations that exceed the standard values) to total observation days.

606 Figure Captions 批注 [LP3]: Please note that the 607 captions have been edited alongside 608 Figure 1. Map showing the PM observation stations and their bar charts of for their their respective figures, below. Once 609 average PM₁₀, PM_{2.5} and PM₁ concentrations (μg m⁻³). Distribution of the PM checked and approved, the final observation stations used in this study. 610 versions will need to be copied to this Figure 2. Map showing bar charts of average PM₁₀, PM_{2.5} and PM₁ concentration (µg-611 separate list. m⁻³) distributions at the observation stations. 612 **带格式的:**缩进:左 悬挂缩进:7.2 字符 左侧: 0 厘米, Figure 3. Scatterplots of (a) PM_{2.5} versus PM₁₀ and (b) PM₄ versus PM_{2.5} at Gueheng. 613 Figure 24. Scatterplots of PM₁ versus PM_{2.5} (a) with and (b) without data from the 614 615 strong sand and dust storm at Akdala. **带格式的:**缩进:左侧: 0 厘米, 悬挂缩进: 7.2 字符 Figure 35. Spatial distribution of the average ratios of (a) PM_{2.5}/PM₁₀ and (b) 616 617 PM₁/PM_{2.5}. Spatial distribution of the average ratios of PM_{2.5}/PM₁₀. Figure 46. Spatial distribution of the seasonal average concentrations (µg m⁻³) of (a) 618 619 PM_{10} , (b) $PM_{2.5}$, (c) PM_1 and (d) ratios of $PM_{2.5}/PM_{10}$. Figure 57. Interannual variations of PM_{2.5} concentrations at the stations (a) on the 620 HBP and Guanzhong Plain, (b) in northeast China, (c) along the middle and 621 622 lower reaches of the Yangtze River, and (d) in southern ChinaInterannual variations of PM_{2.5} concentrations at (a) the stations in the HBP and 623 Guanzhong Plain, (b) the stations in northeast China, (c) the stations along the 624 middle and lower reaches of the Yangtze River, and (d) the stations in southern 625 626 Figure 68. Interannual variations of (a) PM₁₀ concentrations and (b) PM₁ 627 628 concentrations at Zhengzhou, Xian and Gucheng. Figure 79. Diurnal variation of PM_{2.5} concentrations at the stations (a) on the HBP and 629 630 Guanzhong Plain, (b) in northeast China, (c) along the middle and lower reaches of the Yangtze River, and (d) in southern ChinaDiurnal variations of 631 PM_{2.5} concentrations at (a) the stations in the HBP and Guanzhong Plain, (b) 632 the stations in northeast China, (c) the stations along the middle and lower-633 reaches of the Yangtze River, and (d) the stations in southern China. 634 Figure 8. Anthropogenic emission distributions at a resolution of $0.1^{\circ} \times 0.1^{\circ}$, based on 635 HTAP v2 dataset: (a) BC; (b) PM_{2.5}; (c) SO₂; (d) NOx (units: kg m⁻² 636 s⁻²). Anthropogenic emission distributions at 0.1 degree × 0.1 degree re 637 of (a) BC, (b) PM_{2.5}, (c) SO₂ and (d) NOx (units: kg m⁻² s⁻²) based on 下标 带格式的: 638 带格式的:下标 HTAP v2 dataset. 639 Figure 9. Emissions differences between 2010 and 2008 at a resolution of $0.1^{\circ} \times 0.1^{\circ}$, 640 based on HTAP v2 dataset: (a) BC; (b) PM_{2.5}; (c) SO₂; (d) NOx (units: kg m⁻¹ 641 s⁻²)Emission difference between 2010 and 2008 at 0.1 degree × 0.1 degree 642 resolution of (a) BC, (b) PM_{2.5}, (c) SO₂ and (d) NOx (units: kg m⁻² s⁻²) based 带格式的: 下标 643 带格式的: 下标 on HTAP v2 dataset. 644 645 Figure 10. Daily averaged PM_{2.5} concentrations vs wind speed and relative humidity data at 646 Zhengzhou, Shenyang and Nanning in January 2013. 647 Figure 11. Interannual variations of PM₁₀, PM_{2.5} and PM₁ vs wind speed and relative

648

649 650 humidity at Zhengzhou and Nanning.



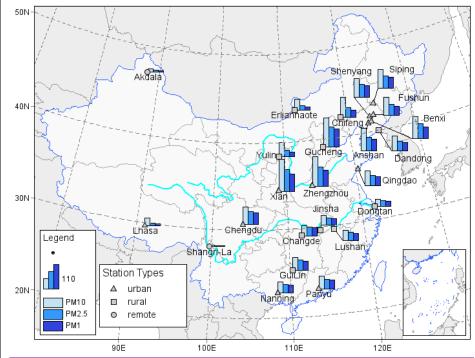


Figure 1. Map showing the PM observation stations and their bar charts offor their average PM_{10} , $PM_{2.5}$ and PM_{1} concentrations ($\mu g \ m^{-3}$). Distribution of the PM observation stations used in this study.



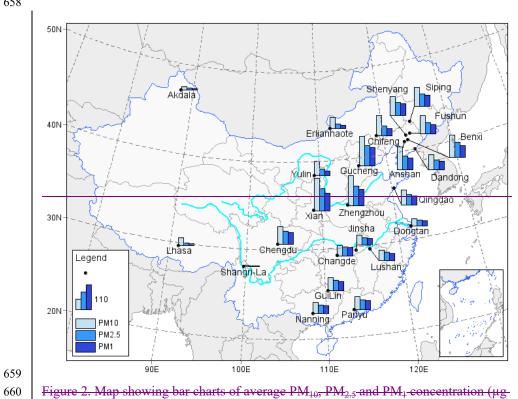
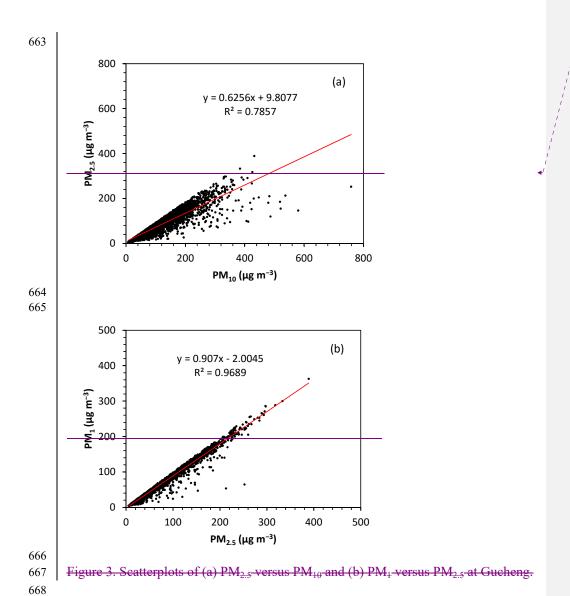
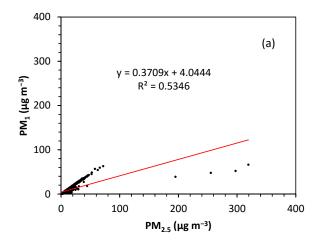


Figure 2. Map showing bar charts of average PM₁₀, PM_{2.5} and PM₁ concentration (µgm⁼³) distributions at the observation stations.



带格式的: 左



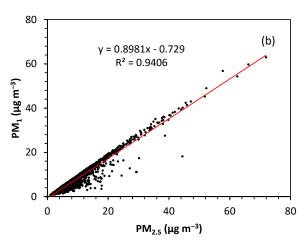
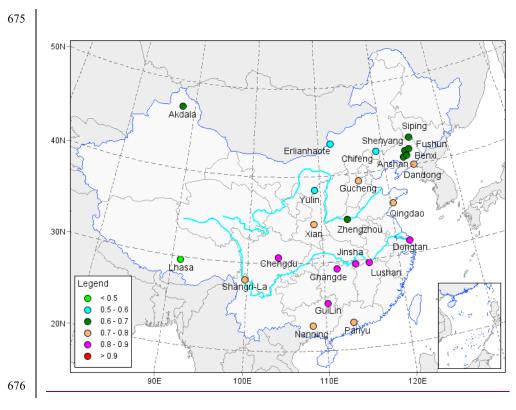


Figure $\underline{2}4$. Scatterplots of PM_1 versus $PM_{2.5}$ (a) with and (b) without data from the strong sand and dust storm at Akdala.



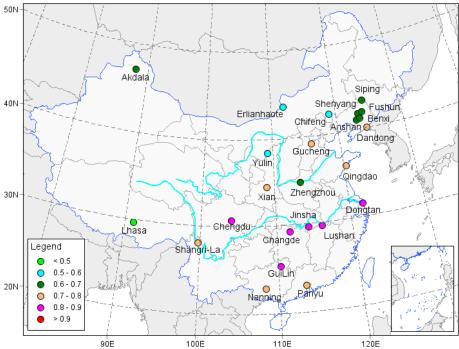
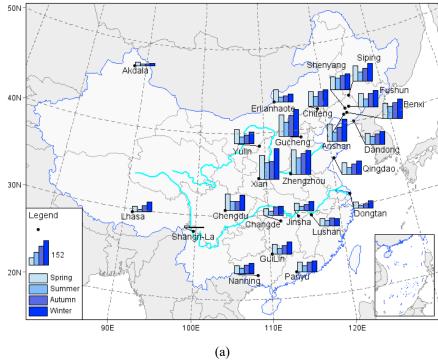
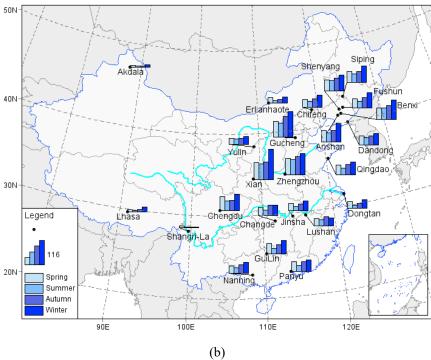
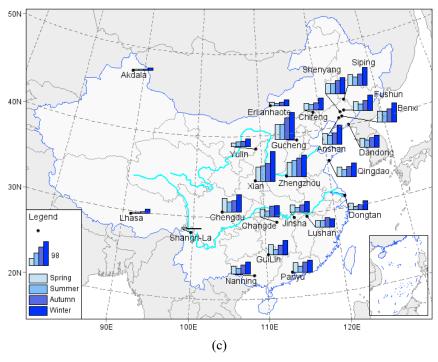


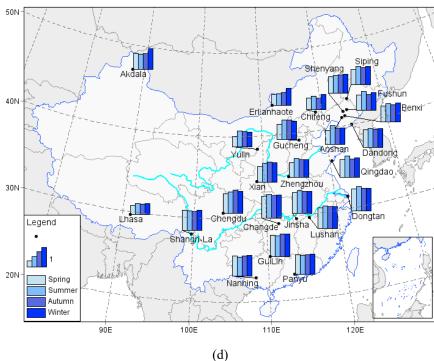
Figure 35. Spatial distribution of the average ratios of (a) PM_{2.5}/PM₁₀ and (b)

 $\underline{PM_1/PM_{2.5}}$. Spatial distribution of the average ratios of $\underline{PM_{2.5}}$.



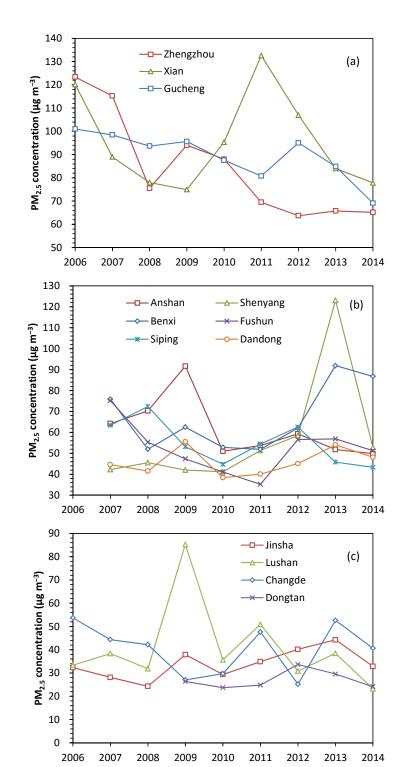






691 | Figure 64. Spatial distribution of the seasonal average concentrations ($\mu g \ m^{-3}$) of (a) 692 | PM₁₀, (b) PM_{2.5}, (c) PM₁ and (d) ratios of PM_{2.5}/PM₁₀.

带格式的: 两端对齐



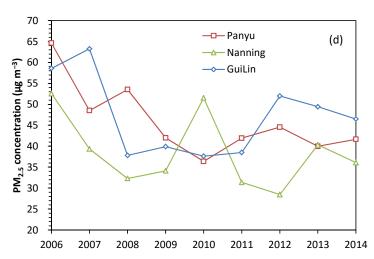


Figure 75. Interannual variations of PM_{2.5} concentrations at the stations (a) the stations in on the HBP and Guanzhong Plain, (b) the stations in northeast China, (c) the stations—along the middle and lower reaches of the Yangtze River, and (d) the stations—in southern China.



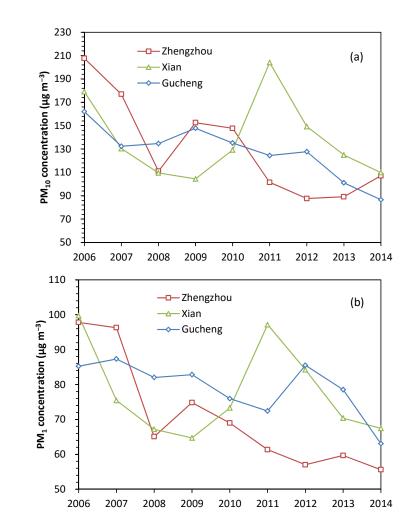
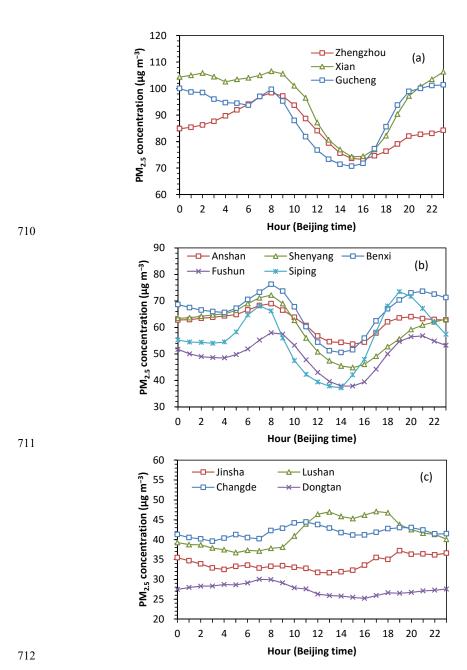


Figure §6. Interannual variations of (a) PM_{10} concentrations and (b) PM_1 concentrations at Zhengzhou, Xian and Gucheng.



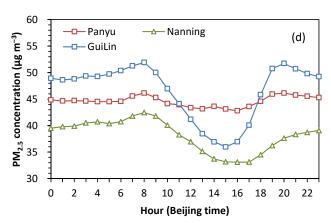
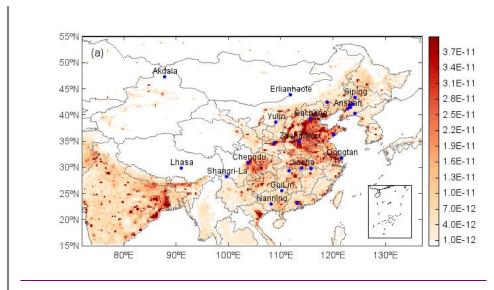
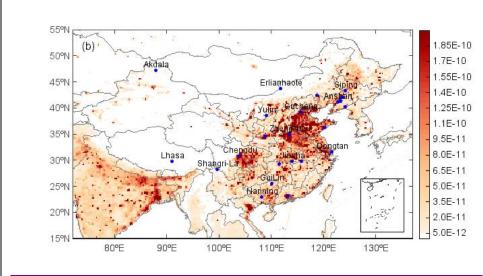
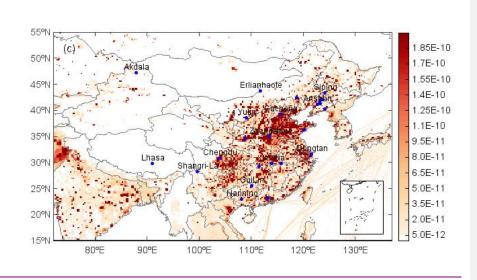


Figure 97. Diurnal variations of $PM_{2.5}$ concentrations at <u>the stations</u> (a) <u>the stations</u> in northeast China, (c) <u>the stations</u> along the middle and lower reaches of the Yangtze River, and (d) <u>the stations</u> in southern China.







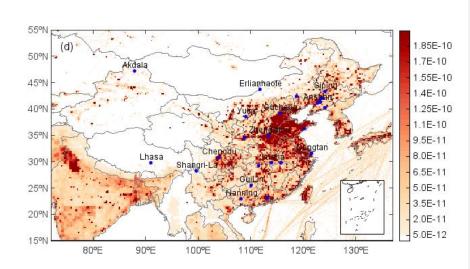
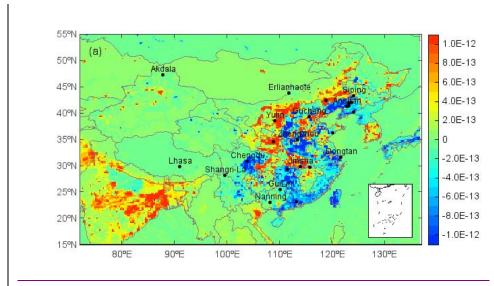


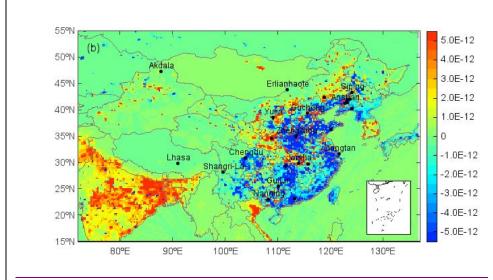
Figure 8. Anthropogenic emission distributions at a resolution of $0.1^{\circ} \times \text{degree} \times 0.1^{\circ}$, based on HTAP_v2 dataset degree resolution of: (a) BC₇; (b) PM_{2.5}; (c) SO₂; and (d) NOx (units: kg m⁻² s⁻²) based on HTAP_v2 dataset.

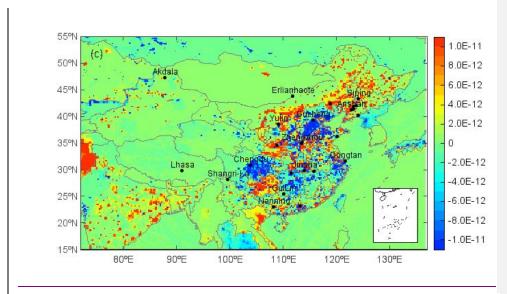
 带格式的:
 左

 带格式的:
 下标

 带格式的:
 下标







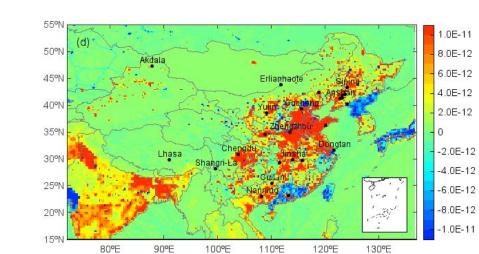
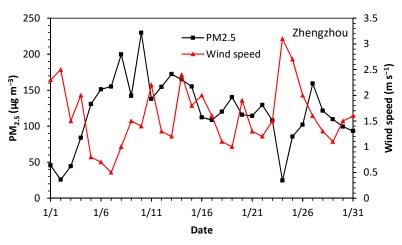


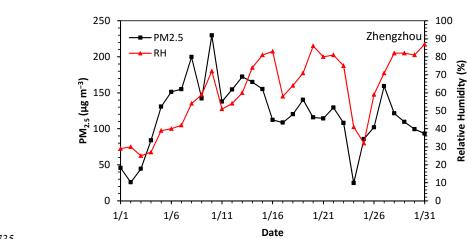
Figure 9. Emissions differences between 2010 and 2008 at a resolution of $0.1^{\circ} \times - \frac{\text{degree} \times 0.1^{\circ}, \text{ based on HTAP}_{v2} \text{ dataset: } \frac{\text{degree resolution of (a) BC;; (b) PM}_{2.5;; (c)}}{\text{SO}_{2}; \text{ and (d) NOx (units: kg m}^{-2} \text{ s}^{-2}) - \frac{\text{based on HTAP}_{v2} \text{ dataset.}}{\text{based on HTAP}_{v2}}$

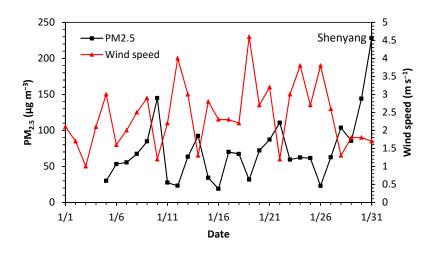
 带格式的: 左

 带格式的: 下标

 带格式的: 下标







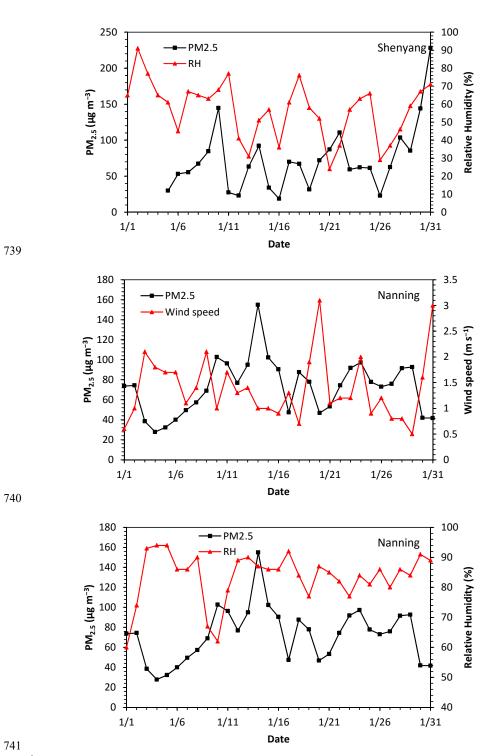
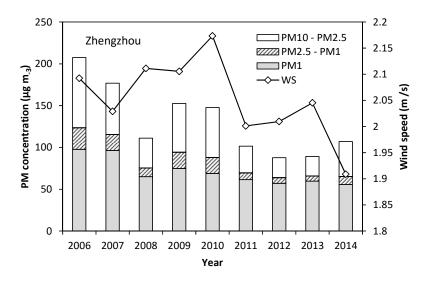
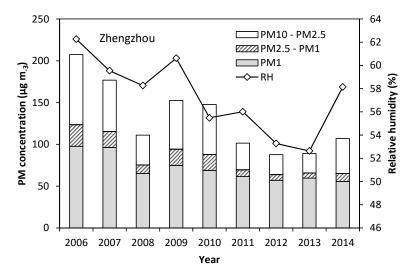
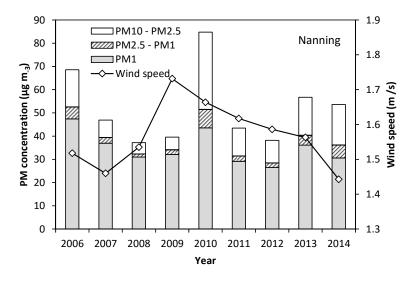


Figure 10. Daily averaged PM_{2.5} concentrations vs wind speed and relative humidity data at Zhengzhou, Shenyang and Nanning in January 2013.









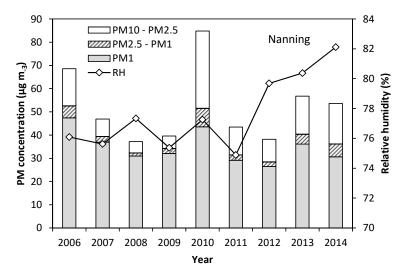


Figure 11. Interannual variations of PM_{10} , $PM_{2.5}$ and PM_1 vs wind speed and relative humidity at Zhengzhou and Nanning.