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1	Urban-rural and sex differences in cancer incidence and
2	mortality and the relationship with PM <sub>2.5</sub> exposure: An
3	ecological study in the southeastern side of Hu line
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20	
21	Abstract: This study investigates the urban-rural and sex differences in the increased risks
22	of the ten most common cancers in China (released by the National Central Cancer Registry
23	of China) related to high PM <sub>2.5</sub> concentration in the southeastern side of Hu line. Pearson

24 correlation coefficient is estimated to reveal how the cancers closely associated with PM<sub>2.5</sub> long-term exposure. Then linear regression is conducted to evaluate sex- and area-specific 25 26 increased risks of those cancers from high level PM<sub>2.5</sub> long-term exposure. The results show that, with the increase of every 10  $\mu$ g/m3 of annual mean PM<sub>2.5</sub> concentration, the relative 27 28 risks of lung cancer incidence and mortality increase 15% and 23% for males, and 22% and 24% for females in rural area, respectively. The relative risk of ovarian cancer incidence 29 increases 9% for females in urban area. For prostatic cancer, the relative risk of incidence 30 increases 17% for males in urban area. For leukemia, in rural area, the relative risks of 31 incidence and mortality increase 22% and 19% for females, respectively. While the relative 32 risk of mortality increases 9% for males; and incidence increases 6% for females in urban 33 34 area. The results demonstrated that, with PM<sub>2.5</sub> exposure, the risks of ovarian and prostatic 35 cancer increase significantly in urban area, while lung cancer and leukemia increase significantly in rural area. Moreover, the elevated risks of lung cancer and leukemia with 36 37 higher PM<sub>2.5</sub> exposure seems more significant for female. This study suggests that the carcinogenic effects of PM<sub>2.5</sub> have obvious sex and urban-rural differences. 38

Keywords: Urban-rural difference; Sex difference; Relative risk; PM<sub>2.5</sub>; Cancer incidence;
Cancer mortality.

41

### 42 **1. Introduction**

The threat to the public health caused by the exposure to fine particulate matter and air pollution has attracted more and more attentions from the public, governments and health organizations worldwide [1-3]. Recent studies show that air pollution has become a major global health risk factor [4-7], particulate matter with aerodynamic diameter less than 2.5 μm (PM<sub>2.5</sub>) shortens life expectancy due to its health impact on morbidity and mortality [8-10], especially for lung cancer [11,12] and cardiovascular diseases [13-15]. Cohen et al. [16] indicated that 4.2 million deaths in 2015 were caused by the exposure to PM<sub>2.5</sub>, and more than 1.1 million deaths were in China. WHO issued a  $PM_{2.5}$  guideline value of annual mean 10 µg/m<sup>3</sup>, and interim targets (IT) level 1, 2 and 3 of 35, 25 and 15 µg/m<sup>3</sup>. At IT-1 level, a 15% higher long-term mortality risk is reported relative to the guideline level [17]. However, IT-1 level is not achieved at most of the areas of China, and population-weighted mean of PM<sub>2.5</sub> concentration in Chinese cities were 61 µg/m<sup>3</sup> at the year of 2013 [18, 19]. Therefore, the health impact caused by PM<sub>2.5</sub> exposure has become an urgent issue in China [20-24].

The studies in the developed areas revealed that air pollution caused by PM<sub>2.5</sub> was a 56 serious threat to human health in various aspects, i.e. its enhancement of cardiopulmonary 57 diseases [25-31], premature birth and low birth weight [32-35], and systemic diseases [36-58 39]. In China, many studies showed that short-term exposure of PM<sub>2.5</sub> was also associated 59 60 with the rise of hospital emergency-room visits, cardio-respiratory diseases and mortality in 61 city areas [40-48]; while cohort studies show that ambient particulate matter can increase the 62 risks of total, cardiovascular and respiratory mortality [49-53]. There are two main causes of 63 these health hazards: one is that the fine particles in PM<sub>2.5</sub> are small enough to arrive a large part of human organs (including the respiratory system, the circulatory system, and the 64 reproductive system), and another is that there are numerous kinds of hazardous substances 65 in the PM<sub>2.5</sub>, i.e. carcinogenic polycyclic aromatic hydrocarbons (PAHs), heavy metals (such 66 as lead, mercury, chromium and cadmium), and pathogenic microorganisms (such as bacteria, 67 68 viruses and fungi) [54-56]. However, most of these studies only focused on the possible diseases caused by the former while ignore some other possible diseases caused by the later. 69 Thus, to understand the health effects of PM<sub>2.5</sub> exposure in China, we should not only choose 70 71 some potential diseases to study, but also instead screen all the most common diseases, since the PM<sub>2.5</sub> concentration in China has been at the high level for a long time and the main 72 hazardous substances in PM<sub>2.5</sub> such as PAHs and heavy metals probably increase the risks of 73 74 several kinds of common disease [57-60].

75 PAHs in PM<sub>2.5</sub> are suspected to be a predisposing factor of breast cancer because of its disruption of BRCA-1 gene expression in estrogen receptor [61]. Parikh et al. [62] also 76 77 conclude that PAHs in PM<sub>2.5</sub> have a significant impact on the increased incidence of female breast cancer in urban areas. Further, BRCA-1 has been confirmed to be associated with 78 79 ovarian cancer [63]. Therefore, it could be deduced that cancer could be a manifestation of the health effects of PM<sub>2.5</sub> in the regions with large population density, since PAHs can almost 80 always be detected in PM<sub>2.5</sub> in these regions [64]. Moreover, we need to consider the 81 geographical and sex factors, because there are spatial differences in the PM<sub>2.5</sub> concentration 82 and compositions (mainly between urban and rural area) as well as sex differences in the 83 sensitivity to toxic substances. Thus, to make a detailed assessment of the health effects 84 85 caused by PM<sub>2.5</sub> in China, firstly, we need to find out which cancers closely associated with 86 high PM<sub>2.5</sub> concentration; and then, we need to evaluate sex- and area-specific increased risks of those cancers from high level  $PM_{2.5}$  long-term exposure. In this study, we first investigated 87 88 the association between the ten most common cancers incidence and mortality with PM2.5 concentration to find out which cancers are closely relative to PM<sub>2.5</sub> by using the time series 89 90 data of yearly incidence and mortality of the ten most common cancers and the annual mean 91 PM<sub>2.5</sub> concentration in China from 2000 to 2011. Secondly, we estimated the sex- and areaspecific increased cancer incidence/mortality risks from long-term exposure to high PM<sub>2.5</sub> 92 93 concentration by using spatiotemporal series data of the southeastern side of Hu line from 2006 to 2009. Finally, we studied the urban-rural and sex differences in the increased risks 94 of incidence and mortality for the ten most common cancers from long-term exposure to high 95 PM<sub>2.5</sub> concentration. The southeastern side of Hu line [65] was selected as our research region, 96 97 because this part of China has larger population density and more developed social economic 98 level, which mean the air pollution caused by  $PM_{2.5}$  is very serious and the data collected are 99 also more consistent with the actual situation.

### 101 **2. Materials and methods**

102 The data collection method and statistical method were presented in section 2.1 and 103 section 2.2, respectively.

104 *2.1. Data* 

Cancer incidence and mortality data as well as gridded PM<sub>2.5</sub> concentrations and
 population data could be collected according to the following methods.

107 2.1.1. Cancer incidence and mortality data

108 The data used in this paper are:

109 1) Cancer statistics reported in the work of Chen *et al.* [49]. This is a time series dataset 110 spanning from 2000 to 2011. It consists of cancer incident and mortality of 22 registries 111 covering 44.4 million population. This dataset was employed to conduct Pearson correlation 112 coefficient analysis to investigate the relationship between cancer incidence and mortality 113 and PM<sub>2.5</sub> exposure.

114 2) Chinese cancer registry annual report (2009-2012) issued by the National Central Cancer Registry of China (NCCR). Data collected from the report spanning from 2006 to 115 116 2009. This is a spatiotemporal series dataset containing cancer incident and mortality data 117 reported from 34 local population-based cancer registries located in the area southeast of Hu line known as an area with greater population density and more developed economy (as 118 119 presented in Figure 1). The total 34 cancer registries including 16 and 18 sites for urban and 120 rural areas respectively. This dataset was used to perform linear regression to estimate the area-specific and sex-specific increased risks of cancer incidence and mortality from long-121 122 term exposure to high PM<sub>2.5</sub> concentration.

Both of the two datasets used in this study were validated by NCCR based on the Guidelines for Chinese Cancer Registration and International Agency for Research on Cancer/International Association of Cancer Registries (IARC/IACR) data-quality criteria.



Figure 1. Maps of the contributing cancer registries and geographic regions in China from the
Chinese cancer registry annual report (2009-2012). The blue line indicates the Hu line, which
marks a drastic difference in the distribution of China's population. The southeast of Hu line is
known as an area with greater population density and more developed economy.

131

## 132 2.1.2. Gridded PM<sub>2.5</sub> concentrations, population and per capita GDP data

Because the ground monitoring of  $PM_{2.5}$  were not reported until 2012 in China,  $PM_{2.5}$ concentration estimated from AOD was used as an alternative. The  $0.1^{\circ} \times 0.1^{\circ}$  gridded annual mean  $PM_{2.5}$  concentrations for the period of 2000-2011 were obtained from Atmospheric Composition Analysis Group (http://fizz.phys.dal.ca/~atmos/martin/?page\_id=140). The area covered by each grid is approximately equal to the area covered by a cancer registration site. Therefore a corresponding gridded  $PM_{2.5}$  concentration can be associated with each cancer registry.

The permanent resident population data for each cancer registry were obtained from the 5<sup>th</sup> and 6<sup>th</sup> national population census conducted in 2000 and 2010, respectively. The population were then linearly interpolated and extrapolated to the whole period of 2000-2011. 143 The annual per capita GDP data are obtained from National Bureau of Statistics of China
144 (http://data.stats.gov.cn/easyquery.htm?cn=C01).

145 2.2. Statistical Methods

146 2.2.1. The exposure level of  $PM_{2.5}$ 

147 For each cancer registry, the population-weighted annual mean  $PM_{2.5}$  exposure 148 concentration could be calculated by Eq. (1):

149 
$$PWEL = \sum (Pi \times Ci) / \sum Pi$$
(1)

150 Where PWEL (population-weighted exposure level) is the population-weighted mean  $PM_{2.5}$ 

151 exposure concentration ( $\mu$ g/m<sup>3</sup>), and Pi and Ci are the population (10 thousands) and PM<sub>2.5</sub>

152 concentration ( $\mu$ g/m<sup>3</sup>) at each grid of the cancer registration site, respectively.

# 153 2.2.2. The association analysis of the time series dataset

In order to find out which cancers are closely associated with PM2.5 long-term exposure, 154 the correlation between cancers incidence (and mortality) and the population-weighted PM<sub>2.5</sub> 155 exposure concentration were evaluated. As a comparison, the correlation between cancers 156 incidence (and mortality) and per capita GDP were also calculated. Therefore, the Pearson 157 158 correlation coefficients (R) between the time series of population-weighted annual mean PM<sub>2.5</sub> concentration (or per capita GDP) and incidence (or mortality) for the ten most 159 common cancers were calculated and used to determine whether or not the two are related 160 161 by comparing with the critical value of R when p < 0.05. R can be determined by Eq. (2):

162 
$$\mathbf{R} = \frac{\sum_{i=1}^{n} (X_i - \bar{X})(Y_i - \bar{Y})}{\sqrt{\sum_{i=1}^{n} (X_i - \bar{X})^2} \sqrt{\sum_{i=1}^{n} (Y_i - \bar{Y})^2}}$$
(2)

163 Where R is the Pearson correlation coefficients, n = 12, is the number of the years.  $X_i$  and  $Y_i$ 164 are the cancer incidence (or mortality) and the population-weighted PM<sub>2.5</sub> exposure 165 concentration (or per capita GDP) of the *i*<sup>th</sup> year, respectively.  $\overline{X} = \frac{\sum_{i=1}^{n} X_i}{n}$  and  $\overline{Y} = \frac{\sum_{i=1}^{n} Y_i}{n}$ , are the arithmetic average of X and Y, respectively.

167 2.2.3. The regression analysis of the spatiotemporal series dataset

The spatiotemporal series data were derived from 34 cancer registries (including 16 168 169 urban sites and 18 rural sites) over a 4-year period, and a sample of 136 (including 64 for urban sites and 72 for rural sites) PM<sub>2.5</sub>-cancer incidence (and PM<sub>2.5</sub>-cancer mortality) data 170 were obtained. All the annual data at each of sites were firstly categorized at a bin of 5  $\mu$ g/m<sup>3</sup> 171 172 population-weighted annual mean PM<sub>2.5</sub> concentration, and then bin-averaged incidence, mortality and population-weighted annual mean PM2.5 concentration were calculated. With 173 174 the bin-averaged data, linear regression was performed to reveal the relation between 175 incidence/mortality of the ten most common cancers and population-weighted annual mean PM<sub>2.5</sub> concentration, then to obtain the increased risks of cancers incidence and mortality 176 with every 10  $\mu$ g/m<sup>3</sup> increment of population-weighted annual mean PM<sub>2.5</sub> concentration. 177 178 Accordingly, the errors can be reduced and the regression precision can be improved by using the average values of PM2.5 exposure level, cancer incidence and mortality in each bin instead 179 180 of original ones.

181

## 182 **3. Results and Discussion**

183 The results of this study were firstly presented in this section, then, the results were184 discussed in details.

3.1. The association between the time series of PM<sub>2.5</sub> and incidence (and mortality) of the ten
most common cancers during 2000-2011

The correlation coefficients between the time series of population-weighted annual mean 187 PM<sub>2.5</sub> concentration and incidence (and mortality) of the ten most common cancers for both 188 189 males and females were presented in Figure 2. The parameter-per capita GDP was added as a contrast, because economy is an important interference factor which can not only contribute 190 191 to environmental deterioration but also the public health condition. In fact, studies have reported that some high cancers incidence were stimulated by the economic development 192 [54], and two main reasons were speculated: firstly, environmental degradation usually 193 follows with economic development [55], the results in this study also validated this (R<sub>PM2.5</sub>-194 per capita GDP is 0.594, means the population-weighted annual mean PM<sub>2.5</sub> concentration is 195 positively related with per capita GDP between 2000 and 2011); secondly, more mature 196 197 cancer screening methods were applied with economic development, and advanced methods 198 can diagnose cancers earlier to reduce the missed diagnosis cases, and finally result in increased incidence [56]. For example, the incidence of female thyroid cancer increased year 199 200 by year may be a false alarm due to the improvement of medical technology [57].

According to Figure 2, it could be concluded that the incidence of six cancers in the ten 201 202 most common cancers are significantly positive associated with population-weighted annual 203 mean PM<sub>2.5</sub> concentration and per capita GDP both for male (including prostatic cancer, leukemia, brain/central nervous system (CNS) cancer, pancreatic cancer, bladder cancer, and 204 205 colorectal cancer) and female (including lung cancer, breast cancer, colorectal cancer, uterine 206 cancer, ovarian cancer and cervical cancer). On the Contrary, there are only two cancers whose mortality are significantly positive associated with population-weighted annual mean 207 208 PM<sub>2.5</sub> concentration and per capita GDP, both for male (including leukemia and prostatic cancer) and female (ovarian cancer and cervical cancer). 209



Figure 2. The correlation coefficient (R) between population-weighted annual mean  $PM_{2.5}$ concentrations (and per capita GDP) and cancer incidence, between population-weighted annual mean  $PM_{2.5}$  concentrations (and per capita GDP) and cancer mortality for males and females. The R = ± 0.576 lines are also shown in the figure as orange dashed lines (when |R| > 0.576, P

217 Consequently, the negative correlation of cancer- $PM_{2.5}$ / GDP presented in Figure 2 can 218 be attributed to the health effects of economic development, while the positive correlation 219 was associated with the environmental deterioration ( $PM_{2.5}$  is one of the most dangerous 220 pathogenic contaminants) or/and more mature cancer screening methods.

221 More concretely, significant positive association between cancer incidence and per capital GDP reveals that there is an increased risk from environmental deterioration (mainly 222 223 from  $PM_{2,5}$ ) and / or more mature cancer screening methods, while significant positive association between cancer mortality and per capital GDP was irrelevant to the screening 224 225 methods. Meanwhile, negative association shows that there is a decreased risk because of some healthy factors along with a higher per capital GDP such as a lower smoking rate. 226 227 Therefore, the negative association between cancer and  $PM_{2.5}$  should be attributed to the 228 health effects of GDP and the positive correlation between PM<sub>2.5</sub> and GDP. And the positive association between cancer incidence and PM<sub>2.5</sub> were caused by two reasons: one is that PM<sub>2.5</sub> 229 pollution increases the risk of cancer incidence, and another is that more mature cancer 230 231 screening methods increase the cancer incidence. Meanwhile, there is only one reason for the 232 positive association between cancer mortality and PM<sub>2.5</sub>, and it is that PM<sub>2.5</sub> pollution can increase the risk of cancer mortality. 233

234 A lower smoking rate was one of the most important protector of health [66]. According to the work of Research on National Health Services in Global Adult Tobacco Survey 235 236 between 1993 and 2010, smoking rates in China are declining (see Table 1), which is the possible cause of the decreasing incidence of lung cancer, esophagus cancer, liver cancer and 237 238 stomach cancer. Meanwhile, Figure 2 also shows that the incidence and mortality of male 239 lung cancer were not correlated with  $PM_{2.5}$  concentration, and the results was inconsistent with literature [14, 20, 21] due to the smoke rate data. The positive correlation between 240 female lung cancer incidence and population-weighted annual mean PM<sub>2.5</sub> concentration 241 demonstrates the effect of PM<sub>2.5</sub> exposure on lung, while the hazards by PM<sub>2.5</sub> exposure to 242 lung for male have been papered over. A possible explanation is that the effect of  $PM_{2.5}$  on 243 244 lung is not noticeable on people with high smoking rate (male) but significant on those with low smoking rates (female), because smoking is much more harmful compared with PM<sub>2.5</sub> 245 [25]. 246

 Table 1. Smoking rates for residents over 15 years old (%)

Voor		Male		Female			
Tear	Total	Urban	Rural	Total	Urban	Rural	
2010	52.9	49.2	56.1	2.4	2.6	2.2	
1993	59.3	56.8	60.3	5.0	6.2	4.5	

To summarize, for the ten most common cancers, the incidence of six cancers and the mortality of two cancers are closely related with PM<sub>2.5</sub> exposure. The involved organs include cardiovascular system, respiratory system, reproductive system, digestive system and hematopoietic system.

3.2. The association between the spatiotemporal series of PM<sub>2.5</sub> and incidence/mortality of
the ten most common cancers during 2006-2009

255 In order to investigate the urban-rural and sex differences of the health effects of PM<sub>2.5</sub> 256 exposure, the association between the spatiotemporal series of PM<sub>2.5</sub> and incidence (and 257 mortality) of the ten most common cancers during 2006-2009 were analyzed. According to results in section 3.1, cancer incidence was increased both by PM<sub>2.5</sub> pollution and mature 258 259 screening methods, while cancer mortality was only increased by PM<sub>2.5</sub> pollution, therefore, 260 in order to exclude the impact of mature screening methods, we select the cancers whose mortality are significantly positive related with PM<sub>2.5</sub> as our study subjects in this section. As 261 a special case, lung cancer was also analyzed in this section because it is fully demonstrated 262 263 to be affected by  $PM_{2.5}$  exposure [11].

Table 2. The number of data in each PM<sub>2.5</sub> concentration range

 Bins of PM <sub>2.5</sub> concentration	Urban area	Rural area
$(\mu g/m^3)$	(numbers of data)	(numbers of data)
 21-25	0	4
26-30	0	0
31-35	1	7
36-40	5	3

41-45	9	8	
46-50	15	6	
51-55	6	10	
56-60	15	9	
61-65	7	16	
66-70	2	4	
71-75	3	2	
76-80	0	3	
81-85	1	0	
Total	64	72	

The annual mean of  $PM_{2.5}$  concentration at most of the sites falls into the range of 36-60 µg/m<sup>3</sup>, and only 9% are below 35 µg/m<sup>3</sup> (IT-1 level), 92% of which are from rural area. Furthermore, there are about 20% of urban areas and 35% of rural areas with  $PM_{2.5}$ concentration more than 60 µg/m<sup>3</sup> respectively, which means more rural population were exposed to high levels  $PM_{2.5}$  concentration. It is worth pointing out that the annual mean  $PM_{2.5}$  concentration was not weighted by population because the area of  $0.1^{\circ} \times 0.1^{\circ}$  grid is almost equal to the area covering by a cancer registry.

For each kind of cancer, the regression of cancer with urban male, rural male, urban female and rural female were carried out. The regression formula was presented in Eq. (3):

275

276

 $y = Bx + b_0 \tag{3}$ 

277

278 Where y is cancer incidence or mortality, x is annual mean  $PM_{2.5}$  concentration. All the 279 data need to be preprocessed with the method stated in section 2.2.3.

280

**Table 3.** The R<sup>2</sup> and significance of regression model for cancer incidence and mortality.

	Url	ban	Rural		
Site	Male	Female	Male	Female	
	Incidence Mortality	Incidence Mortality	Incidence Mortality	Incidence Mortality	

Lung	0.616*	0.576*	0.307	0.387	0.766*	0.861*	0.805*	0.671*
Ovary	/	/	0.747*	0.015	/	/	0.291	0.009
Prostate	0.668*	0.009	/	/	0.048	0.279	/	/
Cervix	/	/	0.561*	0.924*	/	/	0.162	0.087
Leukaemia	0.021	0.269	0.454*	0.236	0.198	0.415*	0.429*	0.742*

281 '\*' indicates P < 0.05 and the regression is significant.

282

According to Table 3, there is not always a linear relationship between the incidence and mortality of the selected five cancers and  $PM_{2.5}$  concentration. For the same cancer, there are significant sex and urban-rural differences in the health effects of  $PM_{2.5}$ . And for different cancers,  $PM_{2.5}$  has different hazards in urban and rural areas. The regression relationships were employed to identify the trends and patterns, as presented Figure 3.





Figure 3. The linear regression trendlines for the five cancers. The formulas in the chart are theregression formulas of the corresponding regression trendlines.

Accordingly, it could be estimated that the increased relative risks (RRs) of the five cancers incidence and mortality for every 10  $\mu$ g/m<sup>3</sup> increment of annual mean PM<sub>2.5</sub> concentration, compared with which is at 35  $\mu$ g/m<sup>3</sup> can be determined according to Eqs.4-5.

294 
$$RR_{I}=10B/I_{0}$$
 (4)

(5)

295 
$$RR_{M} = 10B/M_{0}$$

Where  $RR_1$  and  $RR_M$  are the increased relative risk for incidence and mortality respectively, and I<sub>0</sub> and M<sub>0</sub> are incidence and mortality when annual mean PM<sub>2.5</sub> concentration is at 35  $\mu g/m^3$  (IT-1 proposed by WHO) respectively, B is the coefficient of the corresponding formula.

300 **Table 4.** The RRs of cancer incidence and mortality for every 10  $\mu$ g/m<sup>3</sup> increment of annual mean 301 PM<sub>2.5</sub> concentration, compared with which is at 35  $\mu$ g/m<sup>3</sup>.

	Urban				Rural			
Site	Male		Fen	Female M		ale	Female	
	Incidence	Mortality	Incidence	Mortality	Incidence	Mortality	Incidence	Mortality
Lung	-8%	-9%	Non	Non	15%	23%	22%	24%

Ovary	/	/	9%	Non	/	/	Non	Non
Prostate	17%	Non	/	/	Non	Non	/	/
Cervix	/	/	-13%	-14%	/	/	Non	Non
Leukaemia	Non	Non	6%	Non	Non	9%	22%	19%

302 'Non' represents there is no significant risk.

303

Table 4 shows that, there are obvious urban-rural and male-female distinctions for the 304 health impact of  $PM_{2.5}$  exposure. More specifically, the risks of  $PM_{2.5}$  exposure are higher in 305 urban area for ovarian cancer and prostatic cancer; while for lung cancer and leukaemia, the 306 307 risks are higher in rural area. Moreover, the risks of male lung cancer and cervical cancer in urban decrease along with every 10 µg/m<sup>3</sup> increment of annual mean PM<sub>2.5</sub> concentration, 308 and this should be an erroneous conclusion, since that recent years in developed areas when 309 310 annual mean PM<sub>2.5</sub> concentration kept growing, and the declined smoking rate was a dominant factor for diminished lung cancer; and more advanced medical techniques inhibited 311 cervical cancer. 312

More concretely, compared to the situation of 35  $\mu$ g/m<sup>3</sup> annual mean PM<sub>2.5</sub> concentration 313 (IT-1), RRs of lung cancer mortality for males and females in rural area increases 23% and 314 24% respectively with every 10 µg/m<sup>3</sup> increase of annual mean PM<sub>2.5</sub> concentration, which 315 is consistent with Turner et al. [67], and there is almost no sex difference in lung cancer 316 mortality in rural area. Different from rural area, the risks of incidence and mortality for male 317 318 (female) lung cancer decline (change insignificantly) with PM<sub>2.5</sub> respectively in urban area, which again suggests that smoking rate decline could be the dominant factor for lung cancer. 319 320 The urban and rural differences of RRs of incidence and mortality for other cancers are as following: with every 10 µg/m<sup>3</sup> increment of annual mean PM<sub>2.5</sub> concentration, the RR of 321 ovarian cancer incidence increases 9% in urban area, which changes insignificantly in rural 322

area; the RR of prostatic cancer incidence increases 17%, which also changes insignificantly in rural area; for male leukemia, only the RR of mortality in rural increase 9%, but for female leukemia, the RRs of incidence and mortality increase 22% and 19% in rural area respectively, and in urban area, only the RR of incidence increases 6%, which suggests that the RR of leukemia from PM<sub>2.5</sub> exposure is more significant in rural area than urban area, and it is more significant for females than males.

329

### **4. Conclusions**

Particulate matter pollution has become an urgent issue in most areas in China because 331 that the annual mean PM<sub>2.5</sub> concentration has remained at a high level greater than 35  $\mu$ g/m<sup>3</sup> 332 for a long period. In this paper we used the data of cancer incidence/mortality and PM<sub>2.5</sub> 333 concentration to carry out a preliminary exploration of the sex and urban-rural differences in 334 the health effects of PM<sub>2.5</sub> pollution in densely populated areas located in the southeastern 335 side of Hu line of China. Pearson correlation coefficient and linear regression were performed 336 to analyze the association between PM2.5 and the incidence (and mortality) of the ten most 337 common cancers. For the ten most common cancers, the incidence of six cancers and the 338 mortality of two cancers are closely related with PM<sub>2.5</sub> exposure. The mainly involved organs 339 include cardiovascular system, respiratory system, reproductive system, digestive system and 340 hematopoietic system. For the same cancer, there is a big gap in RRs of PM<sub>2.5</sub> long term 341 exposure between urban and rural area, and between male and female. For different cancers, 342 343 the hazards of PM<sub>2.5</sub> vary in urban and rural areas. Our results are demonstrated reliable because of the consistency with Turner et al. [67]. 344

All in all, PM<sub>2.5</sub> long term and high concentration exposure sharply raises the risks of some cancers in China, meanwhile, the responses of cancers to PM<sub>2.5</sub> are inconsistent in urban and rural areas, and in different sex. Specifically, the responses of lung cancer and leukaemia to  $PM_{2.5}$  are more significant in rural area, while the responses of ovarian cancer and prostatic cancer are more significant in urban area. In addition, the hazards of  $PM_{2.5}$  on female are more significant. Therefore, urban and rural differences and sex differences should be taken into account in the management of the air pollution and the associated health problem.

352 Our research is the first step of differential study of the health risks of PM<sub>2.5</sub> long term and high concentration exposure in urban-rural areas and different sex, so the interference of 353 354 some other factors is not completely excluded and leads to some abnormal results. A case in 355 point is that the RRs of male lung cancer and cervical cancer in urban area decrease along with PM<sub>2.5</sub> concentration increment, which suggests that there may be other influencing 356 factors leading to the result, such as the declined smoking rate and more advanced medical 357 358 techniques. In order to get more accurate exposure-response relationship between cancers 359 and long-term PM<sub>2.5</sub> exposure, it is necessary to choose typical areas to carry on large-scale prospective cohort study. 360

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364 Data Sharing Statement: The data used in this study all come from published articles,
365 yearbooks and publicly-accessible websites, therefore, all the data are open to everyone. No
366 additional data available.

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conducted data analysis. J. R. led this manuscript; L.C., K.C., and Y.L. gave some useful
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370 **Conflicts of Interest:** The authors declare no conflict of interest.

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