Association of PM2.5 and Its Components With Lengths of Hospital Stay for Hand Foot and Mouth Disease in Children

Wei Li  
children's hospital of Nanjing medical university  https://orcid.org/0000-0003-0769-5805

Jieguo Wang  
Children's Hospital of Nanjing Medical University

Kai Zhou  
Children's Hospital of Nanjing Medical University

Ye Tian  
Children's Hospital of Nanjing Medical University

Feiran Wei  
Southeast University Zhongda Hospital

Mingzhi Zhang (✉ mingzhizhang@njmu.edu.cn)  
Nanjing Medical University  https://orcid.org/0000-0001-6429-0041

Xu Wang  
Children's Hospital of Nanjing Medical University

Research

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Abstract

Background

Hand foot and mouth disease (HFMD) is a public health concern. Studies on air pollution and the lengths of hospital stay (LOS) of HFMD are scarce.

Methods

We characterized the clinic demographic features of 5135 hospitalized HFMD in Nanjing, China from 2012 to 2017, and analyzed the association between short-term exposure to PM$_{2.5}$ as well as its components (OM, BC, SO$_4^{2-}$, NH$_4^+$, NIT, SOIL and SS) and the LOS of HFMD.

Results

Among them 98.62% were aged 0–6 years old, and 3772 (73.46%) were hospitalized for seven days or less. The LOS of HFMD children was different in different ages, illness onset years and illness onset seasons ($P$<0.01). For per IQR increase in PM$_{2.5}$ concentrations, LOS of HFMD increased by 0.52 (0.33, 0.71), 0.50 (95% CI, 0.31–0.69) and 0.46 (95% CI, 0.28–0.65) day in adjusted models at lag 3 days, lag 7 days and lag 14 days, respectively. In addition, per IQR increase of BC, SO$_4^{2-}$, NH$_4^+$, NIT and SOIL were also significantly associated with the LOS of HFMD.

Conclusions

Our findings corroborated the hypothesis that short-term PM$_{2.5}$ exposure was associated with increased the LOS of HFMD, and its components (BC, SO$_4^{2-}$, NH$_4^+$, NIT and SOIL) of PM$_{2.5}$ might play a key role in the prolonged LOS of HFMD.

Introduction

Hand foot and mouth disease (HFMD) is a public health concern [1, 2]. It is caused by a variety of human intestinal viruses, most commonly by enterovirus 71 (EV71) and coxsackievirus A16 (CVA16) [3–5], and is a common infectious disease in infants and young children under 5 years of age with fever and rash or herpes on hands, feet, mouth and other parts[6]. This disease is distributed globally and widespread in Asia, especially in the East and South Asia[7–11]. In May 2008, China made HFMD as a statutory infectious disease, and the National Health and Health Committee of China officially issued the "Hand foot and mouth disease Treatment Guide (2018 Edition)". During 2008–2015, a total of 398,010 HFMD patients with >2 episodes were reported among children in China [12]. The general cases lasting for 7 to 10 days, however, some theses severe cases need prolonged hospitalization, the longer lengths of
hospital stay (LOS) of HFMD will increase the chance of infection in the hospital and increases the cost of hospitalization.

Due to the long survival period of HFMD related intestinal virus in ambient air, air pollutants may play a critical role in HFMD epidemics[13]. Previous studies have indicated that air pollutants are significantly related to HFMD incidence in children[14]. As PM$_{2.5}$ is one of the major air pollutants in the world and severe to health, good air quality might protect against HFMD[15]. Improving air quality, especially decreasing PM$_{2.5}$ could decrease the risk of HFMD outbreaks[16]. PM$_{2.5}$ is a complex mixture of inorganic and organic compounds, the main components of PM$_{2.5}$ include organic matter (OM), black carbon (BC), sulfate (SO$_4^{2-}$), ammonium (NH$_4^+$), nitrate (NIT), SOIL, sea salt (SS), trace element oxides, and others[17–19]. Most of the research indicated that an exceedance of PM$_{2.5}$ level of the standards could lead to varying degrees of health hazards based on PM$_{2.5}$ chemical constituents[20]. But these studies have focused on adult hospital admissions or mortality, however, few studies have focused on the association between PM$_{2.5}$ and its chemical components and the LOS of HFMD.

In our study, we characterized the clinic demographic features of hospitalized HFMD in Nanjing, China, focusing on age, gender, illness onset year, illness onset season, as well as geographical patterns. We evaluated HFMD cases exposure to PM$_{2.5}$ and its main components, aimed to evaluate the association between short-term exposure to PM$_{2.5}$ and its components and the LOS of HFMD, to test the hypothesis that specific chemical component of PM$_{2.5}$ is responsible for PM$_{2.5}$ associated the LOS of HFMD.

**Methods**

**Study Design and Participants**

**Population**

The clinical data collected from Children's Hospital of Nanjing Medical University between January 2012 and December 2017 were analyzed retrospectively. 5135 hospitalized Nanjing children with HFMD were diagnosed by a professional pediatrician. Their medical histories were abstracted from medical records. The study was approved by the Ethics Committee of Children's Hospital of Nanjing Medical University. Data used in this work were anonymous, and no individually identifiable information was available here.

**Clinic demographic features of participants**

After admission, a face-to-face interview was conducted by the professional pediatrician to collect data on children's characteristics, including age, gender, illness onset season and illness onset year. All HFMD cases were diagnosed, excluding other infectious diseases. The LOS was recorded for every participant.

**Modeling PM$_{2.5}$ and its components concentrations**
The geocoded residential addresses of all 5135 HFMD cases were linked to the average modeled concentrations of PM$_{2.5}$ and its constituents. We evaluated the residential children exposure to PM$_{2.5}$ and its seven main constituents (OM, BC, SO$_4^{2-}$, NH$_4^+$, NIT, SOIL and SS) 3, 7 and 14 days prior to hospitalization using the V4.CH.02 product of the Dalhousie University Atmospheric Composition Analysis Group (ACAG) [21, 22].

**Statistical analysis**

First, we estimated the characteristics of age, gender, illness onset season (spring (from March to May), summer (from June to August), autumn (from September to November) and winter (from December to February)), illness onset year and LOS on 5135 hospitalized children living in Nanjing with HFMD. Cases’ residential addresses were collected from the hospital medical records. The geographic location of children at home was converted to latitude and longitude and described in Figure 1. Then we explore age, gender, illness onset season and illness onset year parameters by specific LOS. Correlation between each component of PM$_{2.5}$ was calculated by Pearson correlation analysis. Multiple linear regression models were performed to examine the associations between (1) per interquartile range (IQR) increase in PM$_{2.5}$ and its components and the LOS of HFMD in lag 3 days; (2) per IQR increase in PM$_{2.5}$ and its components and the LOS of HFMD in lag 7 days; (3) per IQR increase in PM$_{2.5}$ and its components and the LOS of HFMD in lag 14 days. Model 1 was unadjusted, Model 2 was adjusted for age, gender, illness onset year and illness onset season. FDR corrections were conducted to correct the p-values due to the multiple comparisons. All analyses were performed using R software (version 3.6.1, R Core Team 2019).

**Results**

**Study population characteristics**

We described the characteristics of 5135 hospitalized HFMD children. The majority were boys (64.32%). The mean age was 1.99 years (standard deviation [SD]: 1.48) (Table 1), and 5064 (98.62%) were aged 0–6 years old. Among all patients, 3772 (73.46%) were hospitalized for seven days or less. There are two epidemic cycles in summer and autumn. There were no deaths, and all of them were discharged from the hospital.
Table 1
Characteristics of 5135 hospitalized children living in Nanjing with hand foot mouth disease.

| Characteristic           | N (%) or Mean±SD |
|-------------------------|------------------|
| Age (year)              | 1.99±1.48        |
| ≤1                      | 2664 (51.88)     |
| 1-6                     | 2400 (46.74)     |
| >6                      | 71 (1.38)        |
| Gender                  |                  |
| boy                     | 3303 (64.32)     |
| girl                    | 1832 (35.68)     |
| Illness onset year      |                  |
| 2012                    | 762 (14.80)      |
| 2013                    | 1048 (20.41)     |
| 2014                    | 919 (17.90)      |
| 2015                    | 836 (16.28)      |
| 2016                    | 964 (18.77)      |
| 2017                    | 606 (11.80)      |
| Illness onset season    |                  |
| spring                  | 882 (17.18)      |
| summer                  | 1606 (31.28)     |
| autumn                  | 1646 (32.05)     |
| winter                  | 1001 (19.49)     |
| Hospital length of stay (day) |  |  |
| ≤7                      | 3772 (73.46)     |
| 7-14                    | 1241 (24.17)     |
| >14                     | 122 (2.38)       |

According to Table 2, the results showed that the LOS of HFMD children was different in different ages, illness onset years and illness onset seasons (P< 0.01). The LOS of HFMD between gender was not significantly different (P=0.91).
Table 2
Parameters by specific length of hospital stay among 5135 children.

| Characteristic       | Length of hospital stay(days) | \( p \)  |
|----------------------|-------------------------------|---------|
|                      | \( \leq 7 \)(n=3772) | 7-14 (n=1241) | >14 (n=122) |
| Age (year)           |                               |         |
| \( \leq 1 \)         | 2019 (53.53)                 | 793 (63.90) | 31 (25.41)   |
| 1-6                  | 1693 (44.88)                 | 448 (36.10) | 90 (73.77)   |
| >6                   | 60 (1.59)                    | 0 (0)     | 1 (0.01)     |
| Gender               |                               |         |
| boy                  | 2430 (64.42)                 | 793 (63.90) | 80 (65.57)   |
| girl                 | 1342 (35.58)                 | 448 (36.10) | 42 (34.43)   |
| Illness onset year   |                               |         |
| 2012                 | 487 (12.91)                  | 233 (18.78) | 42 (34.43)   |
| 2013                 | 779 (20.65)                  | 243 (19.58) | 26 (21.31)   |
| 2014                 | 644 (17.07)                  | 247 (19.90) | 28 (22.95)   |
| 2015                 | 690 (18.29)                  | 142 (11.44) | 4 (3.28)     |
| 2016                 | 749 (19.86)                  | 201 (16.20) | 14 (11.48)   |
| 2017                 | 423 (11.21)                  | 175 (14.10) | 8 (6.56)     |
| Illness onset season |                               |         |
| spring               | 614 (16.28)                  | 245 (19.74) | 23 (18.85)   |
| summer               | 1042 (27.62)                 | 507 (40.85) | 57 (46.72)   |
| autumn               | 1343 (35.60)                 | 278 (22.40) | 25 (20.49)   |
| winter               | 773 (20.49)                  | 211 (17.00) | 17 (13.93)   |

Distributions Pm2.5 And Its Components Exposure

Table 3 shows the average distribution of children exposure to PM\(_{2.5}\) and its components (OM, BC, SO\(_4^{2-}\), NH\(_4^+\), NIT, SOIL, and SS) 3, 7 and 14 days prior to hospitalization. During 3 days before hospitalization, the median concentrations was 50.00µg/m\(^3\) of PM\(_{2.5}\), 50.00µg/m\(^3\) and 50.02µg/m\(^3\) for 7 days and 14
days before hospitalization, respectively. The detailed distributions of PM$_{2.5}$ and its components for 3, 7 and 14 days before hospitalization are presented in Table 3. The correlations between PM$_{2.5}$ and its components were shown in Figure S1, Figure S2 and Figure S3.
Table 3
The distribution of exposure to PM$_{2.5}$ and its components within different periods.

|                       | Mean±SD | 5th  | 25th | 50th | 75th | 95th | Range          |
|-----------------------|---------|------|------|------|------|------|----------------|
| Three days before hospitalization (µg/m$^3$) |         |      |      |      |      |      |                |
| PM$_{2.5}$            | 53.99±18.79 | 28.90 | 38.00 | 50.00 | 66.85 | 90.00 | (21.00, 118.90) |
| OM                   | 8.40±7.13   | 2.30  | 3.60  | 5.30  | 11.10 | 25.50 | (1.70, 34.10)   |
| BC                   | 3.78±1.55   | 1.70  | 2.60  | 3.40  | 4.90  | 6.60  | (1.40, 8.70)    |
| SO$_4^{2-}$           | 12.62±5.26  | 6.20  | 8.80  | 12.00 | 14.90 | 21.43 | (4.90, 38.30)   |
| NH$_4^+$              | 8.65±2.69   | 4.80  | 6.80  | 8.40  | 10.20 | 13.70 | (3.50, 19.60)   |
| NIT                  | 13.63±7.26  | 4.80  | 7.90  | 12.00 | 18.01 | 27.70 | (3.40, 41.70)   |
| SOIL                 | 5.38±5.82   | 0.10  | 0.50  | 4.00  | 8.60  | 14.30 | (0.10, 38.80)   |
| SS                   | 0.26±0.18   | 0.10  | 0.10  | 0.20  | 0.30  | 0.70  | (0.10, 1.50)    |
| One week before hospitalization (µg/m$^3$) |         |      |      |      |      |      |                |
| PM$_{2.5}$            | 53.94±18.59 | 29.00 | 38.00 | 50.00 | 66.60 | 90.00 | (21.00, 118.90) |
| OM                   | 8.36±7.03   | 2.30  | 3.60  | 5.31  | 10.90 | 25.00 | (1.70, 34.10)   |
| BC                   | 3.78±1.54   | 1.72  | 2.60  | 3.41  | 4.90  | 6.59  | (1.40, 8.70)    |
| SO$_4^{2-}$           | 12.66±5.28  | 6.20  | 8.89  | 12.00 | 15.00 | 21.70 | (4.90, 38.30)   |
| NH$_4^+$              | 8.66±2.68   | 4.90  | 6.80  | 8.40  | 10.20 | 13.70 | (3.50, 19.60)   |
| NIT                  | 13.61±7.21  | 4.90  | 7.90  | 12.00 | 18.00 | 27.60 | (3.40, 41.70)   |
| SOIL                 | 5.35±5.73   | 0.18  | 0.50  | 4.00  | 8.54  | 14.20 | (0.10, 38.80)   |
| SS                   | 0.27±0.18   | 0.10  | 0.10  | 0.20  | 0.30  | 0.70  | (0.10, 1.50)    |
| Two weeks before hospitalization (µg/m$^3$) |         |      |      |      |      |      |                |
| PM$_{2.5}$            | 53.83±18.20 | 29.00 | 38.60 | 50.20 | 66.13 | 89.13 | (24.00, 118.90) |
| OM                   | 8.26±6.83   | 2.40  | 3.60  | 5.33  | 10.57 | 24.49 | (1.80, 34.10)   |
| BC                   | 3.76±1.50   | 1.75  | 2.60  | 3.42  | 4.86  | 6.50  | (1.40, 8.70)    |
| SO$_4^{2-}$           | 12.75±5.30  | 6.30  | 8.95  | 12.00 | 15.10 | 21.89 | (4.90, 38.30)   |
| NH$_4^+$              | 8.67±2.68   | 4.95  | 6.80  | 8.50  | 10.20 | 13.69 | (3.60, 19.60)   |
|                     | Mean±SD   | 5th    | 25th   | 50th   | 75th   | 95th   | Range       |
|---------------------|-----------|--------|--------|--------|--------|--------|-------------|
| NIT                 | 13.56±7.10| 4.99   | 7.93   | 12.00  | 17.80  | 27.50  | (3.40, 41.70)|
| SOIL                | 5.31±5.55 | 0.20   | 0.60   | 4.00   | 8.46   | 14.00  | (0.10, 38.80)|
| SS                  | 0.27±0.18 | 0.10   | 0.11   | 0.20   | 0.30   | 0.70   | (0.06, 1.18) |
| Hospital length of stay (day) |          |        |        |        |        |        |             |
| LOS                 | 7.00±3.20 | 3.00   | 5.00   | 6.00   | 8.00   | 14.00  | (1.00, 44.00)|

The association between PM$_{2.5}$ and its components exposure and the LOS of HFMD

As show in Figure 2 and Table S1, Table S2 and Table S3, the crude and adjusted estimate values and 95% confidence interval (95% CI) per IQR of PM$_{2.5}$ increase on the hospital LOS was 0.15 (0.02, 0.28) and 0.52 (0.33, 0.71) at lag 3 days. For lag 7 days, per IQR increase of PM$_{2.5}$ corresponded to a 0.15 (95% CI, 0.02–0.29) and 0.50 (95% CI, 0.31–0.69) increase in the LOS in crude and adjusted models, respectively. For lag 14 days, an IQR increase in PM$_{2.5}$ concentrations corresponded to a 0.17 (95% CI, 0.04–0.30) and 0.46 (95% CI, 0.28–0.65) increase in the hospital LOS, respectively.

In the unadjusted model, we noted that positive associations of per IQR increase of OM, SO$_4^{2−}$, NH$_4^+$ and SS with the LOS of HFMD at lag 3 days, 7 days and 14 days, there were no association between BC, NIT and SOIL with the LOS of HFMD at lag 3 days, 7 days and 14 days. After adjusting for age, gender, illness onset year and illness onset season, we found that per IQR increase of BC, SO$_4^{2−}$, NH$_4^+$, NIT and SOIL were significantly associated with the LOS of HFMD at lag 3 days, 7 days and 14 days, but OM and SS were not associated with the LOS of HFMD at lag 3 days, 7 days and 14 days. The detailed data of the relationship between PM$_{2.5}$ and its components and the LOS of HFMD at lag 3 days, 7 days and 14 days were showed in Table S1, Table S2 and Table S3.

Discussion

Pollutants such as PM$_{2.5}$ are polluting the environment seriously, causing numerous health problems[23]. Many time-series studies have used PM$_{2.5}$ as an exposure indicator[24]. However, HFMD hospitalization caused by PM$_{2.5}$ has been little reported so far. To our knowledge, the topic of the clinic demographic features by the LOS is scarce in Chinese children. Long LOS will not only seriously reduce the turnover rate of scarce bed resources in the hospitals and reduce the throughput of patients, but also increase the financial burden on the families of children[25, 26]. Analyzing the LOS of HFMD is important for hospital
administrators to formulate countermeasures, improve hospital work efficiency, and control excessive medical expenses. Therefore, we described the clinic demographic features and the LOS of HFMD in children's hospital of Nanjing medical university during 2012–2017 in Nanjing, China. Meanwhile, we conducted the relationship between the exposure of PM$_{2.5}$ and its components and the LOS of HFMD.

The incidence of HFMD in China has been reported to be 1-2 per 1,000 people[27]. In our research, the incidence of hospitalized HFMD peaked in children aged 1 year and then decreased with age, over 98% of hospitalized HFMD cases occur in children younger than 6 years of age, which was consistent with the findings in other studies[28–30]. Therefore, measures must be taken to prevent HFMD in these key population groups. There was a difference in gender-specific hospital admission, the same results were reported in previous study[31–33], Although infection rates between males and females are comparable, males are more likely to develop symptoms, more involved in the propagation of outbreaks, and more likely to be brought in for medical care than females[30], the reason for the differences observed in gender is not known exactly. The cases of hospitalized HFMD tended to arise in the warmer season (summer and autumn) of the year. Nanjing has strong sunshine, high temperature and heavy rainfall in summer and autumn, and it has a more serious greenhouse effect. We estimate the sunshine, temperature and humidity might explain the seasonality of HFMD[34, 35].

The cases of hospitalized HFMD were the highest in 2013, and then slightly decreased from 2014 to 2017, especially in 2017, the number of hospitalized HFMD was the lowest. The promulgations of the guidelines for HFMD can reduce the number of inpatients through standardized diagnosis and treatment in outpatient. Meanwhile, the guidelines introduce measures for personal protection, family protection, kindergarten or school prevention and management, thus reducing the incidence of hand foot and mouth disease[36]. EV71 infections are one of the main etiological agents of HFMD, on December 2015, the China Food and Drug Administration (CFDA) approved the first inactivated EV71 whole virus vaccine for preventing severe HFMD. EV71 vaccination could decrease HFMD incidence significantly among children aged two to five years[37]. In children, the EV71 vaccine elicited EV71-specific immune response, less EV71-associated HFMD cases have been observed. One real-world study provided evidence of EV71 vaccine effectiveness for preventing EV71 and "other" viruses associated with HFMD[38].

In our research, most of inpatient children with HFMD were hospitalized for less than 7 days, followed by 7 to 14 days, and few were hospitalized for more than 14 days. Children aged 1 year, illness onset in summer and 2012 were most likely to be hospitalized for more than 14 days. In 2012, the number of inpatient children with HFMD exceeded 14 days was the largest, followed by 2013 and 2014, then it decreased significantly since 2015, which was related to the gradual improvement of the diagnosis and treatment of HFMD. The number of children hospitalized for more than 14 days in summer may be related to the increase of nosocomial infections caused by high temperature and high humidity in summer[39]. The number of children hospitalized for more than 14 days is also higher in aged 1-6 than that of other age groups. In terms of hospitalization management, more attention should be paid to these individuals.
Previous studies showed that PM$_{2.5}$ has a higher influence on hospital admission than other air pollutants[40]. PM$_{2.5}$ was positively associated with LOS among children[41]. There was a short-term increase in hospital admission rates associated with PM$_{2.5}$ for all of the health outcomes except injuries[42]. Our study showed that short-term exposure to PM$_{2.5}$ was positively associated with the LOS of HFMD. Our findings were broadly consistent with those studies[16, 43]. It has been demonstrated that exposure to PM$_{2.5}$ could adversely affect vascular endothelial function, the activity of the sympathetic nervous system, and systemic inflammation, leading to vasoconstriction, increased plasma viscosity, and a risk of blood clotting and thrombosis[44]. These adverse effects can exacerbate the child’s symptoms, therefore increase the LOS of HFMD.

In addition, in our research, SO$_4^{2-}$ was the most significantly associated component of PM$_{2.5}$ with the LOS of HFMD followed by NH$_4^+$, SOIL, NIT and BC. SO$_4^{2-}$ and NH$_4^+$ were the secondary pollutants and mainly concentrated in power plants dust, motor vehicle exhaust and construction dust, generally higher values were found during summer and spring months. SO$_4^{2-}$ is mostly regionally transported in the summer, various studies have linked sulfate exposure with adverse respiratory and cardiovascular effects, as well as mortality[17].

The BC component of PM$_{2.5}$ consists of soot, charcoal, char, and other light absorbing refractory matter. Although the health impacts of BC have been extensively studied[17, 45], its association with HFMD is not as well characterized. In Nanjing, we found an expected association between BC exposure and the LOS of HFMD, our research has enriched the epidemiological information on the health impacts of BC on HFMD. NIT is mainly a secondary particle found in the atmosphere, unlike the adverse health effects of BC exposure, NIT exposure have been less explored [46].

The SS originate Cl$^-$, Na$^+$, and Mg$^{2+}$ and is variable during winter months because of unsettled weather conditions in winter season[47], in our study, the cases of hospitalized HFMD tended to arise in the warmer season. The OM is a highly complex mixture of hundreds of compounds such as organic carbon, polycyclic aromatic hydrocarbons, alkanes, and fatty acids, but the health effects of OM remain largely uncharacterized, we did not find the association between OM exposure and the LOS of HFMD, these findings need to be further confirmed.

Our study had advantages. We focused on the epidemiologic features of the LOS of HFMD in the Chinese population and the relationship between the exposure of PM$_{2.5}$ and its components and the LOS of HFMD. This is a topic that has not received much attention and will provide a basis for hospital management. PM$_{2.5}$ exposure is associated with the increased LOS of HFMD, and its components (BC, SO$_4^{2-}$, NH$_4^+$, NIT and SOIL) of PM$_{2.5}$ might play a key role in the prolonged LOS of HFMD. Our findings call for greater awareness of environmental protection and the implementation of effective measures to improve the quality of air, which may reduce the risks of adverse effects on children. Policy changes to reduce outdoor air pollutant exposure may lead to improved HFMD outcomes and substantial savings in healthcare spending.
Our study had limitations. Some covariates are unmeasured, such as the socioeconomic and educational status of the parents. Although our observational study was completed in a representative city in China and lasted for 7 years, the sample size included is still relatively small, and larger sample size is needed to support the research results. In our study, we focus on PM$_{2.5}$, other air pollutants such as nitrogen dioxide (NO$_2$), ozone (O$_3$) and carbon monoxide (CO) might also contribute to the LOS of HFMD.

**Conclusion**

Short-term PM$_{2.5}$ and its components (BC, SO$_4^{2-}$, NH$_4^+$, NIT and SOIL) exposure might increase the LOS of HFMD. Improving the quality of air could help to reduce the burden of HFMD.

**Declarations**

**Competing interests:**

The authors declare that they have no competing interests.

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N/A

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Figures
Figure 1

The distribution of geographical location of participants in Nanjing
| Per IQR increase (µg/m³) | β (95% CI) | P   | P-adjusted |
|--------------------------|------------|-----|------------|
| PM2.5                    |            |     |            |
| 3 days before hospitalization | 0.52 (0.33, 0.71) | 5.21E-8 | 1.39E-7    |
| 1 week before hospitalization | 0.50 (0.31, 0.69) | 2.49E-7 | 6.64E-7    |
| 2 weeks before hospitalization | 0.46 (0.29, 0.65) | 1.395E-6 | 3.379E-6   |
| BC                       |            |     |            |
| 3 days before hospitalization | 0.27 (0.07, 0.46) | 6.86E-3 | 0.870E-3   |
| 1 week before hospitalization | 0.24 (0.05, 0.43) | 0.015 | 0.020      |
| 2 weeks before hospitalization | 0.20 (0.00, 0.38) | 0.046 | 0.061      |
| NH₄⁺                     |            |     |            |
| 3 days before hospitalization | 0.46 (0.32, 0.61) | 3.36E-10 | 1.320E-9   |
| 1 week before hospitalization | 0.44 (0.29, 0.58) | 3.742E-9 | 1.499E-8   |
| 2 weeks before hospitalization | 0.40 (0.25, 0.55) | 1.267E-7 | 5.163E-7   |
| NO₂⁻                     |            |     |            |
| 3 days before hospitalization | 0.34 (0.14, 0.55) | 1.158E-3 | 1.849E-3   |
| 1 week before hospitalization | 0.31 (0.11, 0.52) | 3.227E-3 | 5.163E-3   |
| 2 weeks before hospitalization | 0.27 (0.06, 0.49) | 0.013 | 0.020      |
| OM                       |            |     |            |
| 3 days before hospitalization | 0.08 (-0.08, 0.25) | 0.354 | 0.465      |
| 1 week before hospitalization | 0.05 (-0.11, 0.22) | 0.543 | 0.629      |
| 2 weeks before hospitalization | 0.01 (-0.14, 0.17) | 0.856 | 0.856      |
| SO₄²⁻                    |            |     |            |
| 3 days before hospitalization | 0.48 (0.33, 0.62) | 1.206E-10 | 9.330E-10 |
| 1 week before hospitalization | 0.46 (0.32, 0.61) | 7.055E-10 | 5.609E-9  |
| 2 weeks before hospitalization | 0.44 (0.29, 0.56) | 1.467E-8 | 1.109E-7  |
| DUST                     |            |     |            |
| 3 days before hospitalization | 0.31 (0.15, 0.47) | 1.429E-4 | 2.857E-4  |
| 1 week before hospitalization | 0.33 (0.17, 0.49) | 7.346E-5 | 1.468E-4  |
| 2 weeks before hospitalization | 0.38 (0.22, 0.55) | 4.866E-6 | 9.809E-6  |
| SS                       |            |     |            |
| 3 days before hospitalization | -0.01 (-0.13, 0.12) | 0.918 | 0.918      |
| 1 week before hospitalization | -0.02 (-0.15, 0.10) | 0.702 | 0.702      |
| 2 weeks before hospitalization | -0.05 (-0.19, 0.07) | 0.369 | 0.421      |

**Figure 2**

As show in Figure 2 and Table S1, Table S2 and Table S3, the crude and adjusted estimate values and 95% confidence interval (95% CI) per IQR of PM2.5 increase on the hospital LOS was 0.15 (0.02, 0.28) and 0.52(0.33, 0.71) at lag 3 days. For lag 7 days, per IQR increase of PM2.5 corresponded to a 0.15 (95% CI, 0.02–0.29) and 0.50 (95% CI, 0.31–0.69) increase in the LOS in crude and adjusted models, respectively. For lag 14 days, an IQR increase in PM2.5 concentrations corresponded to a 0.17 (95% CI, 0.04–0.30) and 0.46 (95% CI, 0.28–0.65) increase in the hospital LOS, respectively.

**Supplementary Files**

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