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The changing health effects of air pollution exposure for respiratory diseases: a multicity study during 2017–2022

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Abstract

Background Multifaceted SARS-CoV-2 interventions have modified exposure to air pollution and dynamics of respiratory diseases. Identifying the most vulnerable individuals requires effort to build a complete picture of the dynamic health effects of air pollution exposure, accounting for disparities across population subgroups.

Methods We use generalized additive model to assess the likely changes in the hospitalisation and mortality rate as a result of exposure to PM2.5 and O_3 over the course of COVID-19 pandemic. We further disaggregate the population into detailed age categories and illustrate a shifting age profile of high-risk population groups. Additionally, we apply multivariable logistic regression to integrate demographic, socioeconomic and climatic characteristics with the pollution-related excess risk.

Results Overall, a total of 1,051,893 hospital admissions and 34,954 mortality for respiratory disease are recorded. The findings demonstrate a transition in the association between air pollutants and hospitalisation rates over time. For every 10 μ g/m³ increase of PM2.5, the rate of hospital admission increased by 0.2% (95% CI: 0.1–0.7%) and 1.4% (1.0–1.7%) in the pre-pandemic and dynamic zero-COVID stage, respectively. Conversely, O₃-related hospitalization rate would be increased by 0.7% (0.5–0.9%) in the pre-pandemic stage but lowered to 1.7% (1.5–1.9%) in the dynamic zero-COVID stage. Further assessment indicates a shift of high-risk people from children and young adolescents to the old, primarily the elevated hospitalization rates among the old people in Lianyungang (RR: 1.53, 95%CI: 1.46, 1.60) and Nantong (RR: 1.65, 95%CI: 1.57, 1.72) relative to those for children and young adolescents. Over the course of our study period, people with underlying diseases would have 26.5% (22.8–30.3%) and 12.7% (10.8–14.6%) higher odds of having longer hospitalisation and over 6 times higher odds of deaths after hospitalisation.

Conclusions Our estimates provide the first comprehensive evidence on the dynamic pollution-health associations throughout the pandemic. The results suggest that age and underlying diseases collectively determines the

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disparities of pollution-related health effect across population subgroups, underscoring the urgency to identifying the most vulnerable individuals to air pollution.

Keywords Air pollution, Respiratory disease, Health risks, Jiangsu, Epidemiology

Background

Dynamics and burdens of respiratory diseases are shaped by interacting climatic and societal processes, including exposure to air pollution and human mobility pattern. Mounting evidence suggests that widespread non-pharmacal interventions during early pandemic quickly transformed social mixing and reduced pollution-attributable mortality [1–5]. To make robust assessment of such health benefits, there is a pressing need to critically analyse multifaceted health outcomes such as hospitalisations and mortality over a longer term.

Identifying the most vulnerable individuals to air pollution is a major public health concern. Given the age disparities in social activities and thus air pollution exposure, it is likely that air pollution may have disproportionately affected different age groups [6]. Recognizing the role of immunity, air pollution threatens to pose a dual burden to individuals with pre-existing diseases [7, 8]. This motivates effort to elucidate variations of pollutionrelated health outcomes across population subgroups. To date, many studies on respiratory diseases have reported findings on the association between ambient air pollution and hospital admissions and mortality [9–11]. Yet, heterogeneities across age groups and the overlapping burden for those with underlying diseases remain unknown.

The aim of this multicity study is to estimate age-stratified associations between air pollution exposure and health outcomes for respiratory diseases, such that recognizing potential changes between these associations during the COVID-19 pandemic. To this end, we first sought to quantify the health effects of air pollution exposure, both for hospital admissions and mortality. Additionally, we identified vulnerable population subgroups by assessing age differences in pollution-health associations and the excess risk of having underlying diseases.

Methods

Study population

Our study is based in 13 cities of Jiangsu province across a broad range of climatic conditions and sociodemographic structures (Supplemental Fig. 1). Individual-level records of respiratory diseases related-hospital admissions and mortality from Jan 1, 2017 to Dec 31, 2019 (defined as "pre-pandemic stage") and from Jan 1, 2020 to Dec 31, 2022 (defined as "dynamic zero-COVID stage") were retrieved from the surveillance system of Jiangsu Provincial Center for Disease Control and Prevention. The associated characteristics, including age, sex, postcode of residence, date of hospitalisation and mortality, diagnosis, and underlying diseases, were available for each individual. The underlying diseases considered include coronary heart disease, stroke, hypertension, chronic obstructive pulmonary disease, diabetes, chronic kidney disease, cancer, and immune defect.

Health outcomes

Outcomes of interest differ between analyses on population and individual levels. At the population level, outcomes of interest included the daily number of hospital admissions and deaths for respiratory diseases. While a patient's vulnerability was assessed by using the length of hospitalisations and death after hospitalisation. For simplicity, we categorized the patient's length of hospital stay as less or longer than the median length among all individuals; and distinguished whether individuals died or recovered after hospitalisation.

Air pollution exposure

We obtained daily air temperature at 2 m (m) and relative humidity from the fifth-generation reanalysis (ERA5) provided by the European Center for Medium-Range Weather Forecasts (ECMWF) [12]. Additionally, we collected daily 1 km (km) PM2.5 and 10 km O_3 data from the Tracking Air Pollution in China (TAP) platform (http://tapdata.org.cn) which provides a near real-time distribution of air pollutants [13, 14]. By identifying the nearest grid point to the geographic location of individual's residence, we further extracted the time-series data of local climate and pollution conditions.

Statistical analyses

We conducted descriptive statistics for the city-stratified rate of hospitalisation and mortality for respiratory diseases. These rates were defined by the number of hospital admissions and deaths per 10,000 people. We quantified the changes of the rates by using the ratio of the rate in pre-pandemic and dynamic zero-COVID stages. All subsequent analyses that investigated potential changes in pollution-related health outcomes were done separately for the pre-pandemic and dynamic zero-COVID stages.

We started with evaluating the overall association between air pollution exposure and different health outcomes in an entire population. To do so, we used generalized additive model (GAM) to collate effects of pollution exposure and weather conditions on the rate of hospitalisations and mortality. Note that we focused on short-term effects of PM2.5 and O₃ and assumed linear associations between pollutant predictors and health outcomes accordingly. We additionally included daily mean temperature and humidity as confounding factors to account for their non-linear effects on both hospitalisations and mortality. Therefore, two-pollutant models were established with the equation as follows:

$$Y_{i,j} = \alpha_{i,j} + \beta_1 P_{i-l1,j}^1 + \beta_2 P_{i-l1,j}^2 + b \left(T_{i-l2,j} \right) + c \left(RH_{i-l2,j} \right) + \gamma_j C_j$$
(1)

where $Y_{i,j}$ is the number of hospital admissions or mortality on day i in city j. The parameter $\alpha_{i,j}$ is the overall intercept. $P_{i-l1,j}^1$ and $P_{i-l1,j}^2$ is PM2.5 and O₃ on single day, l1 (categorized as 0, 1, 2, or 3 days) prior to hospital admission or deaths, respectively; β_1 and β_2 are coefficients of two pollutant predictors respectively. $b(T_{i-l2,i})$ and $c(RH_{i-l2,j})$ is a smooth function accounting for daily mean temperature and humidity on cumulative days, 12, (categorized as 0-7, 0-14, 0-21, or 0-28 days) prior to the outcomes of interest, respectively. All models were adjusted for spatial heterogeneity by using a categorical variable to distinguish across cities, C_i . We fitted models by using records in the dynamic zero-COVID stage and used Generalized Cross-Validation (GCV) to determine the optimal lag effects. Assuming that these optimal lag effects were the same for all subsequent analyses, we further applied models to estimate pollution-health associations at the pre-pandemic stage.

Next, we extended the overall estimates to evaluate the age difference in the association between pollution factors and rate of hospitalisations and mortality. The age groups of the study population was children and young adolescents (aged 0–14 years old), working-age people (aged 15–64 years old), and the old population (over 65 years old). We considered records of hospital admissions among all three age groups; we restricted the analyses of mortality to working-age people and the old as deaths among children and young adolescents were very rare. GAMs were stratified by age groups and applied to each city using the following equation:

$$Y_{i,a} = \alpha_{i,a} + \beta_1 P_{i-lopt1,j}^1 + \beta_2 P_{i-lopt1,j}^2 + b \left(T_{i-lopt2,j} \right) + c \left(R H_{i-lopt2,j} \right) + \gamma A$$
(2)

where $Y_{i,a}$ is the number of hospital admissions or mortality on day i in age group a. The parameter $\alpha_{i,a}$ is the overall intercept. $P_{i-lopt1,j}^1$ and $P_{i-lopt1,j}^2$ are PM2.5 and O₃ on the optimal time lag (opt1); while $b(T_{i-lopt2,j})$ and $c(RH_{i-lopt2,j})$ are smooth functions for the multi-day lagged effect (lopt2) of temperature and humidity. Age groups are included as a categorical variable, A.

Further, we sought to assess the vulnerability of people with underlying diseases. We used multivariable logistic regression to examine the role of having underlying diseases on the odds of longer hospital stay and death after hospitalisation. Models were adjusted for confounding factors, including demographic characteristics (i.e. age and sex), geographic location of residence (rural or urban) and local weather conditions. We additionally accounted for the potential difference in the excess risk among individuals with different number of underlying diseases. We addressed this by evaluating the odds of hospitalisation and mortality for individuals with single disease as compared to those with double or more diseases.

Sensitivity analyses

To validate our findings and insights, we first investigated variations in the pollution-health associations across age groups and cities. Furthermore, we examined potential uncertainty that may arise from the alternative classification of patients with underlying diseases. These individuals were segmented into three categories with single, double and multiple underlying diseases. Recognizing the control measures during the early pandemic and the potential modification of the pollution-related health effects, we make additional assessment of the association between pollutants and health outcomes in the dynamic zero-COVID stage by excluding the records in the first six months of 2020. All analyses were conducted with R version 4.3.0 and ArcGIS 10.8.1.

Results

Study population

We retrieved 1,051,893 records of hospital admissions and 34,954 of mortality for respiratory disease during 2017–2022 in Jiangsu Province, China. The rate of hospitalisations and mortality differs significantly between the pre-pandemic and dynamic zero-COVID stages (Supplemental Fig. 2 and Supplemental Fig. 3); yet, spatial heterogeneity of such changes is observed (Fig. 1).

Overall pollution-health associations

We first identified the overall association between air pollution exposure in the three days before hospital admission and hospitalisation rate for respiratory diseases (Table 1). For every 10 μ g/m³ increase of PM2.5 and O₃, the rate of hospital admission in the pre-pandemic stage increased by 0.2% (95% CI: 0.1–0.7%) and 0.7% (0.5–0.9%), respectively. Mortality of respiratory diseases correlates with pollutants exposure on the previous day (Table 1). It is estimated that the rate of mortality would increase by 1.8% (1.3–2.2%) with every 10 μ g/m³ increase of O₃.

Statistically significant associations between pollutants and health outcomes are also seen in the dynamic zero-COVID stage. We found similar positive association between PM2.5 and hospitalisation rate, with a slightly higher effect (1.4% (1.0-1.7%)) increase in hospitalisation



Fig. 1 Hospital admissions and mortality for respiratory diseases across 13 cities in Jiangsu province. Cities are coloured by the relative rate of (A) hospitalisations and (B) mortality, i.e. the number of hospitalisations and death per 10,000, in pre-pandemic and dynamic zero-COVID stages

 Table 1
 Assessment of the associations between pollutant

 exposure and rate of hospitalisation and mortality in prepandemic and dynamic zero-COVID stages

	Pre-pandemic stage	Dynamic zero- COVID stage
Hospitalisation rate		
PM2.5 (lag 3d)	1.002* (1.001,1.007)	1.014* (1.010, 1.017)
O ₃ (lag 3d)	1.007* (1.005,1.009)	0.983* (0.981, 0.985)
Mortality rate		
PM2.5 (lag 1d)	0.995 (0.989, 1.001)	0.996 (0.991, 1.003)
O ₃ (lag 1d)	1.018* (1.013, 1.022)	1.012* (1.006, 1.018)

Median and 95% credible intervals of the rate ratio is presented. Statistically significant results (p<0.05) are marked with *

rate) compared to pre-pandemic estimates. It is noted that the pattern of association between O_3 and hospitalisations changes, yielding a 1.7% (1.5–1.9%) lower rate of hospitalisation for every 10 µg/m³ increase of O_3 . Additionally, we observed significant association between O_3 and mortality. It shows that O_3 -related increase in the mortality rate would be lowered to 1.2% (0.6–1.8%) in the dynamic zero-COVID stage. Restricting analyses to subpopulations and geographic areas yield greater variations of the associations among age groups and cities (Supplemental Tables 1–2).

Age difference in pollution-health associations

Disaggregation of age groups indicates heterogeneity in pollution-health associations within the population (Fig. 2). Prior to the COVID-19 pandemic, children and young adolescents were at the highest rate of hospitalisation for respiratory diseases across cities, except for Suqian (Fig. 2A). However, such age-stratified associations between pollution and hospital admission shifted in the dynamic zero-COVID stage (Fig. 2B), with substantially elevated rates observed for the old people in Lianyungang (RR: 1.53, 95%CI: 1.46, 1.60) and Nantong (RR: 1.65, 95%CI: 1.57, 1.72) relative to those for children and young adolescents. This increase generates a shift of high-risk people from children and young adolescents to the old. Conversely, we identify that children and the young adolescents turn into the high-risk group with the highest rate of hospitalisation in Suqian (Fig. 2B). Limiting associations to individuals over 14 years shows that mortality rates are generally greater among the old population, with RR values ranging from 1.12 to 2.96, as compared to those for the working-age people across cities and pandemic stages (Fig. 2C–D).

Excess risk of having underlying diseases

After adjusting for age, sex, and residence, patients having underlying diseases correlate with significantly increased odds of longer hospital stay and death after hospitalisation (Table 2). It is estimated that the medium length of hospitalisation over the study period is seven days. Compared with this, our estimates show that people with underlying diseases would have 26.5% (22.8-30.3%) more odds of having longer hospitalisation than those without underlying diseases. Such increase of the odds attributable to underlying diseases are slightly attenuated to 12.7% (10.8-14.6%) in the dynamic zero-COVID stage. Of note, patients with underlying diseases have a strikingly higher odds of deaths after hospitalisation. Over the course of our study period, these patients have odds of deaths that are over 6 times (6.4, (5.6, 7.3)) in the pre-pandemic stage and 6.1 (5.7, 6.6) in the dynamic zero-COVID stage) than those in the reference group.

Further assessment indicates disparities in the odds of deaths among patients with single and multiple underlying diseases (Supplemental Table 3). Compared to patients without underlying disease, our estimates show that having single underlying disease is associated with around 20% and 540% higher odds of longer hospital stay and death after hospitalisation, respectively. These



Fig. 2 Age disparities in pollution-related rate of hospitalisation and mortality. Estimates of (A–B) hospitalisation and (C–D) mortality rate across age groups in pre-pandemic and dynamic zero-COVID stage are presented

Table 2	Odds ratio	of the over	lapping burde	n among individ	luals with underl	ying diseases
						/ /

Pre-pandemic stage		Dynamic zero-COVID sta	ge
Hospitalisation OR (95%Cl)	Mortality OR (95%Cl)	Hospitalisation OR (95%CI)	Mortality OR (95%CI)
1.015* (1.014, 1.015)	1.085* (1.139, 1.337)	1.019* (1.019, 1.020)	1.084* (1.082, 1.087)
1 (ref)	1 (ref)	1 (ref)	1 (ref)
1.086* (1.062, 1.110)	1.233* (1.139, 1.337)	1.066* (1.051, 1.080)	1.161* (1.108, 1.218)
1 (ref)	1 (ref)	1 (ref)	1 (ref)
1.265* (1.228, 1.303)	6.376* (5.578, 7.326)	1.127* (1.108, 1.146)	6.136* (5.698, 6.618)
1 (ref)	1 (ref)	1 (ref)	1 (ref)
1.149* (1.123, 1.175)	0.362* (0.336, 0.391)	1.033* (1.019, 1.047)	0.521* (0.499, 0.544)
0.994* (0.989, 0.999)	0.989 (0.974, 1.004)	1.007* (1.005, 1.010)	0.971* (0.961, 0.980)
1.015* (1.012, 1.018)	0.952* (0.943, 0.962)	0.994* (0.992, 0.996)	1.012* (1.006, 1.018)
0.994* (0.989, 0.993)	1.002 (0.996, 1.008)	0.998* (0.997, 0.998)	0.979* (0.977, 0.982)
1.005* (1.004, 1.007)	0.998 (0.993, 1.002)	1.006* (1.005, 1.007)	1.004* (1.001, 1.007)
	Pre-pandemic stage Hospitalisation OR (95%CI) 1.015* (1.014, 1.015) 1 (ref) 1.066* (1.062, 1.110) 1 (ref) 1.265* (1.228, 1.303) 1 (ref) 1.149* (1.123, 1.175) 0.994* (0.989, 0.999) 1.015* (1.012, 1.018) 0.994* (0.989, 0.993) 1.005* (1.004, 1.007)	Pre-pandemic stage Hospitalisation OR (95%CI) Mortality OR (95%CI) 1.015* (1.014, 1.015) 1.085* (1.139, 1.337) 1 (ref) 1 (ref) 1.086* (1.062, 1.110) 1.233* (1.139, 1.337) 1 (ref) 1 (ref) 1.265* (1.228, 1.303) 6.376* (5.578, 7.326) 1 (ref) 1 (ref) 1.149* (1.123, 1.175) 0.362* (0.336, 0.391) 0.994* (0.989, 0.999) 0.989 (0.974, 1.004) 1.015* (1.012, 1.018) 0.952* (0.943, 0.962) 0.994* (0.989, 0.993) 1.002 (0.996, 1.008) 1.005* (1.004, 1.007) 0.998 (0.993, 1.002)	Pre-pandemic stage Dynamic zero-COVID state Hospitalisation OR (95%CI) Mortality OR (95%CI) Hospitalisation OR (95%CI) 1.015* (1.014, 1.015) 1.085* (1.139, 1.337) 1.019* (1.019, 1.020) 1 (ref) 1 (ref) 1 (ref) 1.066* (1.052, 1.110) 1.233* (1.139, 1.337) 1.066* (1.051, 1.080) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1.127* (1.108, 1.146) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 1 (ref) 0.362* (0.336, 0.391) 1.033* (1.019, 1.047) 0.994* (0.989, 0.999) 0.989 (0.974, 1.004) 1.007* (1.005, 1.010) 1.015* (1.012, 1.018) 0.952* (0.943, 0.962) 0.994* (0.992, 0.996) 0.994* (0.989, 0.993) 1.002 (0.996, 1.008) 0.998* (0.977, 0.998) 1.005* (1.004, 1.007) 0.998 (0.993, 1.002) 1.006* (1.005, 1.007)

Statistically significant results (p < 0.05) are marked with *

estimates surge, primarily the odds of deaths in the dynamic zero-COVID stage, among those with double or more underlying diseases. Sensitivity analyses confirm our finding of the increased odds of hospitalisation and death attributable to underlying diseases (Supplemental Table 4).

Discussion

Our multicity study fits well in the current context of the exacerbated health effects of air pollution exposure. In light of emerging evidence that pollution-attributable mortality may have reduced during the pandemic [1-5], we provided the first essential evidence on the dynamic pollution-health associations throughout the pandemic. Unlike previous studies focusing solely on mortality, our multicity study in Jiangsu province extends the analysis to include hospitalizations, thereby offering a more multifaceted understanding how pandemic-related interventions may have modulated the influence of pollutants on health outcomes. We found that air pollution exposure generally associates with multifaceted health outcomes, including hospitalisations and mortality, for respiratory diseases. This finding is broadly consistent with other studies done across a variety of settings [15-21]. It is important to note that, over time, there is an opposite shift in the pattern of association between air pollutants and hospitalisation rates. The effect of O₃ on the hospitalisation rate was attenuated which, in turn, may be partially attributed to the lowered level of O_3 during the pandemic. In contrast, risk of PM2.5-related hospitalisation increased in the dynamic zero-COVID stage. Although associations between pollutants and hospital admission evolves, we detected no evidence of deviations from the pre-pandemic association between pollutants exposure and mortality.

When considering the disparities of pollution-related health outcomes among population subgroups, our findings emphasize the importance of two main axes of vulnerability to air pollution exposure: age and underlying diseases. Over the course of the pandemic, the shift in the age profile of high-risk groups may be largely dependent on the relative fraction of hospitalisation across age groups (Supplemental Fig. 3). Specifically, the fraction of hospitalisation for respiratory diseases increased among children and young adolescents but reduced among the old in Suqian. Such trade-off generated the shift of the high-risk group from the old to children and young adolescents. Similarly, people over 65 accounted for the greatest fraction of hospitalisation in Lianyungang and Nantong in the dynamic zero-COVID stage, resulting in high-risk population shifts from children and young adolescents to the elderly. Moving beyond the age burden, estimates of an individual's excess risk would additionally heighten the overlapping burden from air pollution and underlying diseases.

Despite the clear signature of age and underlying diseases on individual's vulnerability, we provided essential evidence that sex and location of residence are strongly associated with the increased odds of longer hospitalisation and higher risk of death (Table 2). We identified that male is more likely to have a higher risk of longer hospital admission (8.6% (95% CI 6.2-11%) in pre-pandemic and 6.6% (95% CI 5.1-8.0%) in dynamic zero-COVID stage) relative to females. Similarly, males tend to have 23.3% (13.9-33.7%) and 16.1% (10.8-21.8%) higher risk of death after hospitalisation in two pandemic stages, respectively. This finding is similar to other environmental health studies, highlighting gender difference in the risk of air pollution exposure [22–26]. Additionally, living in urban areas would associate with a 14.9% (12.3-17.5%) increase in the risk of longer hospitalisation but a 63.8% (60.9-66.4%) lowered mortality risk prior to the pandemic. Although this urban-rural difference is narrowed in the dynamic zero-COVID stage, it is mostly attributed to the health inequalities among areas [27]. Inadequate health resources yield the limited access to health facilities for people living in rural areas. In such context, people may not seek care in health facilities, leading to a great reduction of hospital admission and the lowered risk of hospitalisation accordingly. With respect to mortality rate, the limited health resources may delay the treatment of respiratory diseases and therefore contribute to higher odds of death after hospitalisation for the rural population.

The key messages of this study are of great public health significance. We provided fundamental estimates of the dynamic pollution-health associations over time, which makes critical contributions to understanding how pandemic-related interventions may have modulated the influence of pollutants on health outcomes. Importantly, by identifying vulnerable population subgroups, particularly the elderly and those with preexisting conditions, we underscore the disproportionate effects of air pollution. These findings support WHO global air quality guidelines where the elderly and those with underlying diseases bear the greatest burden of air pollution [28]. Additionally, this study based on a multicity approach across 13 cities in Jiangsu province, offers a comprehensive perspective that enhances the generalizability of our findings. In light of these evidence, health authorities will build a complete picture of the heterogeneous health effects of air pollution exposure for respiratory diseases, formulating relevant policies to identify and protect the most vulnerable individuals.

Future studies may build on our findings in several ways. First, we have explicitly estimated air pollution effects on hospitalisation and mortality. Examining across a variety of pollutants and outcomes is essential for differentiating the roles of co-occurring air pollutants on respiratory outcomes. Additionally, assessment of the pollution-related health effects may vary by applying alternative statistical approaches. Evaluating the potential bias of assessment attributable to the statistical approach would be a promising direction for future investigations. Furthermore, our inability to obtain hospitalisation records in 2017–2018 is likely to bias our observed associations which should be validated. Finally, we have reported the excess risk of having underlying diseases. Moving from this finding, prioritizing the most vulnerable subpopulation will require distinguishing contributions across specific underlying diseases.

Abbreviations

ERA5	Fifth-generation reanalysis
ECMWF	European Center for Medium-Range Weather Forecasts
TAP	Tracking Air Pollution in China
GAM	Generalized additive model

Supplementary Information

The online version contains supplementary material available at https://doi.org/10.1186/s12940-024-01083-1.

Supplementary Material 1

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Not applicable.

Author contributions

This study was conceptualised by RL and ZD. The research was supervised by ZD. The data analyses were carried out by SJ, LT and ZL. All authors were involved in the data interpretation. SJ, LT and ZL prepared the manuscript. All authors reviewed and approved the final version of the manuscript. RL and ZD had full access to the data and codes used in this study. All authors have final responsibility for the decision to submit the manuscript for publication.

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Data availability

All data were archived within the surveillance system of the Jiangsu Provincial Center for Disease Control and Prevention. The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Declarations

Ethics approval and consent to participate

Ethics approval for this study was granted through a data sharing agreement from Jiangsu Provincial Center for Disease Prevention and Control.

Competing interests

The authors declare no competing interests.

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