Environmental exposure

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Ambient air pollution and risk of respiratory infection among adults: evidence from the multiethnic study of atherosclerosis (MESA)

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ABSTRACT

Background Air pollution may affect the risk of respiratory infection, though research has focused on uncommon infections or infections in children. Whether ambient air pollutants increase the risk of common acute respiratory infections among adults is uncertain, yet this may help understand whether pollutants influence spread of pandemic respiratory infections like COVID-19.

Objective To estimate the association between ambient air pollutant exposures and respiratory infections in adults.

Methods During five study examinations over 12 years, 6536 participants in the multiethnic study of atherosclerosis (MESA) reported upper respiratory tract infections, bronchitis, pneumonia or febrile illness in the preceding 2 weeks. Using a validated spatiotemporal model, we estimated residential concentrations of ambient PM1.5, NOx, and NO2 for the 2–6 weeks (short-term) and year (long-term) prior to each examination.

Results In this population aged 44–84 years at baseline, 10%–32% of participants reported a recent respiratory infection, depending on month of examination and study region. PM1.5, NOx, and NO2 concentrations over the prior 2–6 weeks were associated with increased reporting of recent respiratory infection, with risk ratios (95% CIs) of 1.04 (1.00 to 1.09), 1.15 (1.10 to 1.20) and 1.21 (1.10 to 1.33), respectively, per increase from 25th to 75th percentile in residential pollutant concentration.

Conclusion Higher short-term exposure to PM1.5 and traffic-related pollutants are associated with increased risk of symptomatic acute respiratory infections among adults. These findings may provide an insight into the epidemiology of COVID-19.

INTRODUCTION

Exposure to ambient air pollutants is a well-established risk factor for several adverse respiratory outcomes.1–3 Long-term exposure to particulate and gaseous pollutants has been associated with negative impacts on pulmonary function in adults and children,4 5 increased risk of asthma development,6 increased severity of chronic obstructive pulmonary disease (COPD) symptoms,7 faster progression of emphysema8 and incidence of malignant neoplasms of the lung.9

Key messages

► Is exposure to ambient air pollutants associated with respiratory infection in adults?
► An increase in residential concentration of PM1.5, NOx, and NO2 from the 25th to the 75th percentile in the previous 2–6-weeks period was associated with 4%, 15% and 21% higher risk of reporting a respiratory infection.
► These findings indicate that fine particulate and traffic-related air pollutants may increase the risk of respiratory infection among adults.

Short-term exposure is associated with worsening of pulmonary function in adults,10 11 aggravation of asthma and bronchitis in adults and children12 13 and increased incidence, hospitalisation and mortality due to respiratory tract infections.14–17

Short and long-term exposure to air pollutants may impair host immune defenses, resulting in increased susceptibility to respiratory infections.18 19 Most prior research on air pollutants and respiratory infections has focused on impacts among children20 21 or individuals with cystic fibrosis.22–24 However, the link between pollutant exposure and acute respiratory infections among adults has received comparatively less attention, even as mortality rates due to lower respiratory infections (LRI) among children under 5 years have declined globally without corresponding improvements in the adult population.25

Acute respiratory infections among adults are common and represent a major source of disease burden.26 The number of LRI-linked deaths among people above 70 years old rose from nearly 750000 to 1.1 million annually between 2000 and 2016.25 Acute respiratory infections account for approximately 5 million emergency department visits in the USA yearly, and for the largest share (41%) of antibiotics prescribed in the adult outpatient


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setting. Among individuals older than 5 years in 2016, there were approximately 11.6 million LRI episodes in the USA, and about 61 million hospitalisations globally. The ongoing COVID-19 pandemic and ageing global population highlight the urgent need to better understand modifiable risk factors of respiratory infections in the adult population.

While current estimates attribute the majority of the excess burden of disease from ambient air pollutants to cardiovascular causes, a potentially large and poorly estimated fraction of that burden is due to respiratory illnesses. The relative paucity of evidence on the association between air pollution and respiratory infections has recently been noted. We examined the association

### Table 1  Baseline characteristics of study participants in the multiethnic study of atherosclerosis (MESA), by report of recent respiratory infection in the first five study examinations*

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>All participants, n=6536 (100.0%)</th>
<th>Participants who reported an infection at any examination, n=3167 (48.5%)</th>
<th>Participants who reported no infection at all examinations, n=3369 (51.5%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years), mean (SD)</td>
<td>62.0 (10.2)</td>
<td>61.2 (10.1)</td>
<td>62.8 (10.3)</td>
</tr>
<tr>
<td>Race</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>White</td>
<td>2516 (38.5)</td>
<td>1157 (36.5)</td>
<td>1359 (40.3)</td>
</tr>
<tr>
<td>Chinese-American</td>
<td>787 (12.0)</td>
<td>332 (10.5)</td>
<td>455 (13.5)</td>
</tr>
<tr>
<td>African-American</td>
<td>1786 (27.3)</td>
<td>898 (28.4)</td>
<td>888 (26.4)</td>
</tr>
<tr>
<td>Hispanic</td>
<td>1447 (22.1)</td>
<td>780 (24.6)</td>
<td>667 (19.8)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>3445 (52.7)</td>
<td>1765 (55.7)</td>
<td>1680 (49.9)</td>
</tr>
<tr>
<td>Male</td>
<td>3091 (47.3)</td>
<td>1402 (44.3)</td>
<td>1689 (50.1)</td>
</tr>
<tr>
<td>Education level</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Less than high school</td>
<td>1169 (17.9)</td>
<td>585 (18.5)</td>
<td>584 (17.3)</td>
</tr>
<tr>
<td>High school</td>
<td>1187 (18.2)</td>
<td>570 (18.0)</td>
<td>617 (18.3)</td>
</tr>
<tr>
<td>Some college or technical certificate</td>
<td>1858 (28.4)</td>
<td>905 (28.6)</td>
<td>953 (28.3)</td>
</tr>
<tr>
<td>College or graduate degree</td>
<td>2322 (35.5)</td>
<td>1107 (35.0)</td>
<td>1215 (36.1)</td>
</tr>
<tr>
<td>Married/living with a partner</td>
<td>3947 (60.4)</td>
<td>1850 (58.4)</td>
<td>2097 (62.2)</td>
</tr>
<tr>
<td>Body mass index, mean (SD)</td>
<td>28.3 (5.5)</td>
<td>28.7 (5.6)</td>
<td>28.0 (5.3)</td>
</tr>
<tr>
<td>Ever smoked</td>
<td>3485 (53.3)</td>
<td>1697 (53.6)</td>
<td>1788 (53.1)</td>
</tr>
<tr>
<td>Current smoker</td>
<td>741 (11.4)</td>
<td>414 (13.1)</td>
<td>327 (9.8)</td>
</tr>
<tr>
<td>Employed</td>
<td>2838 (43.4)</td>
<td>1429 (45.1)</td>
<td>1409 (41.8)</td>
</tr>
<tr>
<td>Annual family income</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;$20000</td>
<td>1597 (24.6)</td>
<td>789 (25.1)</td>
<td>808 (24.2)</td>
</tr>
<tr>
<td>20 000–$39 999</td>
<td>1756 (27.1)</td>
<td>880 (28.0)</td>
<td>876 (26.2)</td>
</tr>
<tr>
<td>$40 000–$74 999</td>
<td>1653 (25.5)</td>
<td>816 (26.0)</td>
<td>837 (25.1)</td>
</tr>
<tr>
<td>≥$75 000</td>
<td>1474 (22.7)</td>
<td>654 (20.8)</td>
<td>820 (24.5)</td>
</tr>
<tr>
<td>Own home or pay mortgage</td>
<td>4413 (67.7)</td>
<td>2045 (64.8)</td>
<td>2368 (70.5)</td>
</tr>
<tr>
<td>Excellent self-reported general health status</td>
<td>1465 (46.9)</td>
<td>639 (44.1)</td>
<td>826 (49.3)</td>
</tr>
<tr>
<td>No health insurance</td>
<td>464 (7.2)</td>
<td>233 (7.4)</td>
<td>231 (6.9)</td>
</tr>
<tr>
<td>On medicaid</td>
<td>616 (9.5)</td>
<td>304 (9.7)</td>
<td>312 (9.3)</td>
</tr>
<tr>
<td>Diabetic per ADA 2003 criteria</td>
<td>1052 (16.1)</td>
<td>537 (17.0)</td>
<td>515 (15.3)</td>
</tr>
<tr>
<td>Has metabolic syndrome</td>
<td>2485 (38.0)</td>
<td>1235 (39.0)</td>
<td>1250 (37.1)</td>
</tr>
<tr>
<td>Moderate and vigorous physical activity, met minutes/week, median (IQR)</td>
<td>3870 (1950–7080)</td>
<td>4065 (2018–7328)</td>
<td>3765 (1905–6840)</td>
</tr>
<tr>
<td>Intentional physical activity, met minutes/week, median (IQR)</td>
<td>810 (105–1950)</td>
<td>788 (88–1875)</td>
<td>840 (158–2085)</td>
</tr>
<tr>
<td>Self-reported emphysema</td>
<td>73 (1.1)</td>
<td>42 (1.3)</td>
<td>31 (0.9)</td>
</tr>
<tr>
<td>Self-reported asthma</td>
<td>253 (3.9)</td>
<td>142 (4.5)</td>
<td>111 (3.3)</td>
</tr>
</tbody>
</table>

*Units are n (%) unless otherwise specified. ADA, American Diabetic Association.
between long- and short-term exposure to particulate matter less than 2.5 microns in diameter (PM$_{2.5}$) and oxides of nitrogen (NO$_x$ and NO$_2$) and recent occurrence of upper respiratory tract infections (URTI), bronchitis, pneumonia and febrile illness in a diverse, well-characterised cohort of adults undergoing follow-up in the multiethnic study of atherosclerosis (MESA).

**METHODS**

MESA is a prospective cohort study of 6814 white, African American, Hispanic and Chinese American individuals recruited from six regions in the USA (Baltimore, MD; New York City, NY; Los Angeles County, CA; Chicago, IL; Winston-Salem, NC and St. Paul, MN) primarily to investigate the progression of subclinical and clinical cardiovascular disease. MESA air is an ancillary study to examine the role of air pollutants in these outcomes. Recruitment and baseline clinical examinations occurred between July 2000 and August 2002. By January 2012, four additional study examinations had been completed. At the first MESA examination in 2000–2002, participants were 44–84 years old and did not have clinical cardiovascular disease. We used data from the first five MESA examinations to quantify the association between long-term and short-term exposure to ambient PM$_{2.5}$, NO$_x$ and NO$_2$ and the occurrence of respiratory infections in the 2 weeks prior to each study examination. We excluded any observations from study visits at which a participant was taking oral steroids.

**Patient and public involvement**

The study design, analysis, interpretation of results and writing or editing of this research paper were made without patient involvement.

**Exposure estimation**

The approach to air pollutant exposure estimation in MESA has been described in detail. A hierarchical high-resolution spatiotemporal model is used to predict outdoor concentrations at each participant’s residence for each 2 weeks period from January 1999 through the end of participant follow-up. Participants provided residential address information for the entire study duration. The model integrates data from US Environmental Protection Agency Air Quality System regulatory monitors, three types of MESA air-specific monitoring campaigns conducted in the study communities and at homes of study participants and hundreds of geocovariates in a universal kriging and spatial smoothing framework. The median (range) cross-validated $R^2$ values of the prediction models across study regions were 0.79 (0.54–0.85) for PM$_{2.5}$, 0.72 (0.50–0.92) for NO$_x$, and 0.78 (0.45–0.90) for NO$_2$. For primary analyses, timescales for assessing associations were selected a priori as the average exposure 1 year before an MESA examination (long-term exposure) and the 4 weeks starting from 14 days before an examination date (short-term exposure). We also conducted sensitivity analyses using average exposures in the 2-weeks period covering 14–28 days prior to an examination.

**Outcome assessment**

At each of five study examinations, participants were asked whether they had experienced or been diagnosed with a cold or influenza, febrile illness, sinus infection, bronchitis or pneumonia in the preceding 2 weeks. We constructed a composite binary variable indicating the occurrence of any of the symptoms or diagnoses.

**Covariate data**

Detailed demographic, anthropometric, lifestyle and medical history data were collected at each MESA clinical examination using self-administered and technician-administered standardised questionnaires. We used data on age (continuous), study region, educational attainment (up to high school or more than high school), sex (male or female), race/ethnicity (white, African American, Hispanic or Chinese American), body mass index in kg/m$^2$ (<25, 25–30 or ≥30), current smoking status (yes or no), marital status (married or not married), annual family income in US dollars (<20 000; 20 000–39 999; 40 000–74 999; ≥75 000), number of people living in the household, number of children in the household, employment status (employed, retired or unemployed/homemaker), home ownership (outright owner, paying a mortgage and renting or other arrangement), having health insurance (yes or no), diagnosis of diabetes mellitus (based on the 2003 American Diabetic Association fasting criteria algorithm), having metabolic syndrome based on National Cholesterol Education Program guidelines and self-reported doctor-diagnosed asthma or emphysema. Potential confounding by neighbourhood socioeconomic status (NSES) was addressed using an NSES index derived from variables abstracted from the 2000 US Census. As earlier described, this index was obtained by summing $z$-scores calculated for six NSES variables (median value of owner-occupied housing units, proportion of residents with at least a high school diploma, percentage with at least a bachelor’s degree, median household income, percentage with household income <50 000 dollars and percent with professional/managerial occupations).

**Statistical analysis**

We applied modified Poisson regression with a sandwich linearised estimator of variance to estimate the association between occurrence of respiratory infections in the 2 weeks preceding each of five MESA clinical examinations and average long-term and short-term residential concentrations of ambient PM$_{2.5}$, NO$_x$ and NO$_2$, adjusting for potential confounders. In contrast to logistic regression, this method for longitudinal binary outcomes
provides a direct measure of the risk ratio and is appropriate when outcomes are common.

We modelled long-term and short-term associations independently, accounting for weather, seasonality and long-term trends. In long-term models, we included smooth functions of calendar time (natural cubic splines with two degrees of freedom per year (df/year)) to account for secular temporal trends. In short-term models, we also used smooth functions of time (splines with six df/year) to account for seasonal and other time-varying influences on infection rates, and additionally included smooth functions of temperature and humidity (four df splines) to deal with potential confounding by weather.

We assessed whether the association between pollutant exposure and recent respiratory infection varied by strata defined by sex, race, current smoking status, age at baseline, diabetes, education level and NSES. Because modification of effects across regions by NSES has been observed in prior MESA studies,38 we examined models with study region–NSES interactions. Since occurrence of a respiratory infection may be influenced by community-wide levels of infection, we did a secondary analysis to examine if the association between reported infection and ambient pollutant concentration varied by whether the week of the study examination was classified by the Centers for Disease Control (CDC) as meeting the state-level threshold for influenza activity, defined as region-specific periods of two or more consecutive weeks in which each week accounted for more than 2% of the season’s total number of specimens that tested positive for influenza in public health laboratories.40 We also conducted a posthoc assessment of whether level of pollutant exposure modified the risk of infection during high vs low influenza activity periods.

Sensitivity analyses included estimation of associations separately for each of the five study examinations, and restriction to participants who were present at all examinations. Because febrile symptoms may reflect both respiratory and non-respiratory illness, we conducted an analysis excluding that symptom subcategory. We also evaluated extended models containing self-reported general health status and levels of moderate and vigorous physical activity (metabolic equivalents of minutes/week) for the baseline visit because these covariates were not available at all visits.

RESULTS

After excluding 278 participants who were missing one or more covariates at all MESA examinations, our analytical data set contained 6536 participants at baseline (table 1). We excluded 347 observations from 79 participants, on study visits at which a participant was taking oral steroids. The mean (SD) age at baseline was 62.0 (10.2) years. At baseline, 3445 (52.7%) participants were female, 2516 (38.5%) were white, 741 (11.4%) were current smokers and 2322 (35.5%) had at least a college education. Among Hispanic and African American participants, females and current smokers, a higher fraction reported an infection on at least one study examination than those reporting no infection at all visits (table 1).

By the fifth examination, 4547 participants who met inclusion criteria were still undergoing follow-up, of whom 4197 contributed to the analysis at all five examinations. Baseline characteristics of participants included in the analysis at all visits compared with the 2339 not
The proportion of participants reporting a recent respiratory infection at each examination was 20.8%, 18.7%, 18.2%, 17.5% and 13.5%, respectively. The prevalence of infection varied between 10% and 32% depending on study region and month of visit, with lower rates during the warmer months (figure 1). The average concentrations of PM$_{2.5}$, NO$_x$ and NO$_2$ varied by region, but generally declined over the study period (figure 2). There was substantial within-region and between-region variability in pollutant concentrations (figure 3).

In unadjusted models, the risk ratios (95% CIs) for the associations between short-term PM$_{2.5}$, NO$_x$ and NO$_2$ exposure and report of recent respiratory infection were 1.07 (1.03 to 1.13), 1.23 (1.20 to 1.26) and 1.28 (1.17 to 1.39), respectively. In fully adjusted models, higher average short-term PM$_{2.5}$, NO$_x$ and NO$_2$ exposure was associated with statistically significantly greater risk of reporting a recent respiratory infection at that visit (PM$_{2.5}$: RR 1.04 (95% CI: 1.00 to 1.09), NO$_x$: 1.15 (1.10 to 1.20) and NO$_2$: 1.21 (1.10 to 1.33) per interquartile range differences of 5.7 µg/m$^3$, 33.5 ppb and 14.4 ppb, respectively) (figure 4). Results for examination-specific associations were similar, although slightly attenuated and with gradual loss of precision over subsequent examinations (figure 4 and online supplemental table S2). We did not observe statistically significant associations between report of respiratory infection and average long-term exposure (1 year before examination) to any of the pollutants. Results from a short-term exposure sensitivity analysis using average concentrations over a 2-week period before study examination were very similar to those from the 4-week average analysis (online supplemental figure S2) and online supplemental table S2). Additionally, our data did not show evidence of statistically significant modification of the association between short-term exposure to PM$_{2.5}$, NO$_x$ and NO$_2$ and risk of infection by any of the characteristics we examined (online supplemental table S3).
Quantitative influenza data could only be obtained from one metropolitan area (Los Angeles), and only for the period starting October 2006. Based on these limited data, we did not find clear evidence that the risk for recent respiratory infection is modified by whether the week of the study visit was classified by the CDC as an influenza week, a designation that denotes a heightened level of circulating influenza-causing pathogens as

Figure 3 Range of estimated $NO_x$, $NO_2$, and $PM_{2.5}$ exposures at study participant residential locations over the year (long-term) and the 2–6 weeks (short-term) interval preceding a clinical examination in MESA, by study site. BAL, Baltimore, MD; CHI, Chicago, IL; LA, Los Angeles, CA; NYC, New York City, NY; STP, St Paul, MN; W-S, Winston Salem, NC. MESA, multiethnic study of atherosclerosis.

Figure 4 Risk ratios and 95% CIs for associations between respiratory infection and pollutant exposure in the 2–6 weeks (A) and year (B) before interview, for an IQR increase in average pollutant exposure. Results are from model adjusted for study site, race/ethnicity, sex, age, body mass index, marital status, education level, smoking status (ever/never smoked), employment status, annual income, neighbourhood level socioeconomic index, home ownership, number of household members, having health insurance, whether on medicaid, diabetes diagnosis based on ADA criteria and splines of calendar time (and temperature and relative humidity for short-term exposure analysis). ADA, American Diabeteic Association.
tested in public health laboratories across the country (online supplemental table S4). Although our data suggested an increased risk of infection for participants with high average short-term exposure to PM$_{2.5}$ and NO$_x$ during high-relative to low-influenza activity periods in Los Angeles, the CIs are wide due to the reduced sample size in this subset of examinations (online supplemental table S5). Additionally, there was no effect modification by NSES, based on models that included study region by NSES interaction terms.

Restricting the sample to the 4197 participants who completed all five study visits and were eligible for the study at each visit, we obtained very similar results (online supplemental table S6) and online supplemental figure S3). When we excluded febrile illness, the results remained the same (online supplemental table S7). Models that additionally included physical activity and self-reported health status did not materially alter the results.

**DISCUSSION**

In a large well-characterised multiethnic cohort observed on up to five occasions over a 12-year period, average exposure to PM$_{2.5}$, NO$_x$ and NO$_2$ in the 2–6 weeks period preceding a clinic visit was associated with report of a recent respiratory infection. The analysis benefited from a validated exposure assessment method developed for this cohort, taking into account changes in participant address and using study-specific monitoring. The model incorporated both regulatory agency and study-specific monitoring to provide refined spatial and temporal contrasts within and between study regions. These findings suggest that short-term variations in fine particulate and traffic-related air pollution may influence susceptibility to acute respiratory infections among adults.

**Comparison with previous studies**

Much of the research assessing the association between air pollution and respiratory disease has focused on LRI outcomes among children.$^{41,42}$ Reports have also highlighted the link between air pollution and mortality, hospitalisation and emergency department visits, especially due to pneumonia, asthma and exacerbations of COPD, among adults.$^{43-47}$ However, the evidence base for the role of air pollutants in the acute manifestation of more common respiratory symptoms and infections among ambulatory adults in the general population is more limited. In a case–crossover study of 57,000 patients in Beijing, PM$_{2.5}$, PM$_{10}$, NO$_2$, SO$_2$ and CO were associated with outpatient visits for URTI, acute bronchitis and community-acquired pneumonia.$^{48}$ A retrospective time series analysis in Hong Kong found significant associations between elevated exposures to PM$_{10}$ ozone and especially NO$_x$ and general outpatient clinic visits for URTI.$^{49}$ Another Chinese time series study of 1.4 million patient visits for respiratory diseases that included pneumonia and acute bronchitis, reported associations with higher outdoor PM$_{2.5}$, PM$_{10}$, NO$_2$ and ozone.$^{50}$ In a less-polluted part of China, PM$_{2.5}$, PM$_{10}$ and NO$_2$ exposures, over the past 7 days were associated with outpatient visits for respiratory disease in a 4-year period.$^{31}$

Our study reflects the context of multiple US cities with comparatively lower pollution levels, and leverages highly resolved exposure information and covariate data collected over a long period in which exposure levels varied substantially.

We observed strong temporal trends in both pollutant concentrations and in the rates of illness reporting among participants. Therefore, we deployed strong statistical controls (adjustment for calendar time and seasonality) which greatly reduced potential for these trends to influence the results. It could be argued that this is over-adjustment, if in fact these observed long-term secular trends of reduced reporting of respiratory infections were caused by declines in air pollution. In that case, our estimates of effect may be attenuated relative to the true magnitude of effect.

Acute respiratory infections are major drivers of primary healthcare and antibiotic drug utilisation, suggesting that pollution control may have wide-ranging individual and community health benefits. In a recent analysis, daily rates of telephone calls and e-mails, as well as actual provider visits and emergency department visits related to respiratory symptoms were linked to increased PM$_{2.5}$ levels on that day, with 2.61% more calls and e-mails and 2.63% more provider visits per 100,000 health insurance programme members per day.$^{52}$ The same syndromic surveillance approach also demonstrated increased respiratory-related healthcare utilisation for increases in ozone concentrations.$^{53}$ An advantage of this design, shared with our approach, is the ability to capture subacute symptoms for which a person may not have otherwise sought medical care.

Evidence pointing to a role for air pollutants in morbidity and mortality due to the novel pandemic coronavirus caused by SARS-CoV-2 continues to emerge, although most reports still drawn from ecological study designs because detailed individual-level data are not yet available.$^{54-56}$ Among older adults in the USA, a nationwide study based on county-level COVID-19 death rates suggested that a 1 µg/m$^3$ higher PM$_{2.5}$ exposure was associated with an 11% increase in COVID-19 mortality.$^{57}$ In Europe, multiple studies have also provided preliminary indications of positive area-level associations between air pollution (PM$_{2.5}$ and NO$_x$) and COVID-19 severity, including a report associating an additional 1 µg/m$^3$ of PM$_{2.5}$ at the municipality-level with nine extra COVID-19 cases, three extra hospital admissions and one extra death.$^{58-60}$ Initial investigations using limited individual-level data also point to an increase of approximately 7% in mortality rate for each additional 1 µg/m$^3$ in long-term PM$_{2.5}$ exposure.$^{61}$ While high-quality representative individual-level data remain scarce, this rapidly evolving body of literature suggests that air pollutants may exacerbate the severity of COVID-19, especially in the presence of more common respiratory symptoms and infections among ambulatory adults.
of other key risk factors such as advanced age, obesity, and comorbid health conditions.

Possible mechanisms

The mechanisms through which pollutants potentially increase the risk of respiratory infection are still being studied. Particulate and gaseous pollutants are thought to cause respiratory injury through induction of oxidative stress and inflammation in the airways.\textsuperscript{18-62} Particles are removed through exhalation, mucociliary clearance or phagocytosis by alveolar macrophages; or they may be translocated through respiratory epithelial surfaces.\textsuperscript{9-63} Particle deposition and clearance vary by individual-level factors such as age, sex, smoking status, concomitant conditions like asthma and COPD, activity levels and breathing patterns, which may therefore affect the risk of developing pollutant-related disease.\textsuperscript{9} Particles that remain in the respiratory tract can cause inflammation, oxidative stress, inhibition of mucociliary clearance mechanisms, disruption of epithelial tight junctions and impairment of macrophage antimicrobial phagocytic activity, potentially enhancing susceptibility to respiratory infections.\textsuperscript{18-64} There is mounting in vitro and in vivo evidence for the pulmonary immunotoxicity of fine particulate matter, although a clear understanding of the underlying mechanisms remains elusive.\textsuperscript{65} Particulate matter appears to enhance \textit{Streptococcus pneumoniae} biofilm growth and colonisation of human middle ear epithelium cells in vitro, and to increase pneumococcal colonisation of the nasopharynx in an animal model.\textsuperscript{66}

Oxides of nitrogen are absorbed in both the upper respiratory tract and pulmonary regions, and their reactions with surface epithelial fluids may result in oxidative stress, while absorbed components may cause deep lung irritation and immunomodulation of host response to infection, possibly increasing susceptibility or worsening the severity of existing infection.\textsuperscript{67} In the Swiss Cohort Study on Air Pollution and Lung and Heart Disease in Adults (SAPALDIA), increased exposure to \(\text{NO}_2\) was linked to higher risk of respiratory problems leading to doctor visits among participants with underlying COPD and chronic bronchitis.\textsuperscript{68} This suggests that particulate and traffic-related pollutants may elevate the virulence of secondary pathogenic agents in the event of concurrent exposure in the human respiratory tract, partly through inhibition of immune function. Some investigators have hypothesised that air pollution may increase the mortality risks associated with the novel epidemic Severe Acute Respiratory Syndrome Coronavirus-2.\textsuperscript{69}

Strengths and limitations

This study has some limitations. Sufficiently detailed CDC influenza activity data for our period of interest were not readily available for most study regions, reducing our ability to assess the interesting potential interactive role of community-level influenza activity on the pollutant-illness relationship. The outcomes were self-reported, although trained research staff conducted the interviews consistently over follow-up. In addition, we used outdoor residential pollutant concentrations that do not take into account the actual microenvironments in which participants spend time. Importantly, we utilised high-quality exposure predictions in an ethnically and geographically diverse and well-characterised cohort that permitted ample confounder control and exploitation of fine spatiotemporal variations in exposure.

CONCLUSION AND RECOMMENDATIONS

Our study provides evidence that higher exposure to PM\(_{2.5}\), \(\text{NO}_x\) and \(\text{NO}_2\) contributes to an increased risk of common but mostly self-limited respiratory infections. The potential burden of these illnesses—in lost productivity, healthcare utilisation and antibiotic use—suggests that control of particulate and traffic-related pollution may provide substantial individual and societal benefit. In light of the increased attention to respiratory infections in the time of the COVID-19 pandemic, these results also may provide an insight into the potential role of pollutants in modifying the risk of novel viral infections. Given the known deleterious effects of pollutants on the respiratory system, including increasing the risk of infections, and preliminary linkages to COVID-19 morbidity and mortality, it is suggested that reducing pollutant exposure may prevent severe illness and death due to COVID-19. Because respiratory infections, possibly including COVID-19, will remain major health problems among adults, two recommendations can be inferred from our study. First, it may be beneficial for older adults who spend substantial amounts of time in the home to utilise air purification equipment such as filters. Second, multipronged regulatory actions to control pollution at the societal level remain critical public health priorities including science-based adjustments of air quality standards and concerted efforts and resources in areas with high concentrations of particulate and traffic-related pollutants.

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REFERENCES


