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Outdoor PM\textsubscript{2.5}, Ambient Air Temperature, and Asthma Symptoms in the Past 14 Days among Adults with Active Asthma

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**BACKGROUND:** Relationships between air quality and health are well-described, but little information is available about the joint associations between particulate air pollution, ambient temperature, and respiratory morbidity.

**OBJECTIVES:** We evaluated associations between concentrations of particulate matter ≤ 2.5 μm in diameter (PM\textsubscript{2.5}) and exacerbation of existing asthma and modification of the associations by ambient air temperature.

**METHODS:** Data from 50,356 adult respondents to the Asthma Call-back Survey from 2006–2010 were linked by interview date and county of residence to estimates of daily averages of PM\textsubscript{2.5} and maximum air temperature. Associations between 14-day average PM\textsubscript{2.5} and the presence of any asthma symptoms during the 14 days leading up to and including the interview date were evaluated using binomial regression. We explored variation by air temperature using similar models, stratified into quintiles of the 14-day average maximum temperature.

**RESULTS:** Among adults with active asthma, 57.1% reported asthma symptoms within the past 14 days, and 14-day average PM\textsubscript{2.5} ≥ 7.07 μg/m\textsuperscript{3} was associated with an estimated 4–5% higher asthma symptom prevalence. In the range of 4.00–7.06 μg/m\textsuperscript{3} of PM\textsubscript{2.5}, each 1-μg/m\textsuperscript{3} increase was associated with a 3.4% [95% confidence interval (CI): 1.1, 5.7] increase in symptom prevalence; across categories of temperature from 1.1 to 80.5°F, each 1-μg/m\textsuperscript{3} increase was associated with increased symptom prevalence (1.1–44.4°F: 7.9%; 44.5–58.6°F: 6.9%; 58.7–70.1°F: 2.9%; 70.2–80.5°F: 7.3%).

**CONCLUSIONS:** These results suggest that each unit increase in PM\textsubscript{2.5} may be associated with an increase in the prevalence of asthma symptoms, even at levels as low as 4.00–7.06 μg/m\textsuperscript{3}.

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**Introduction**

Ambient particulate matter (PM) pollution accounted for an estimated 3.1 million deaths and 3% of global disability-adjusted life years in 2010 (Liu et al. 2012). Health studies examining the effects of PM exposures have identified links between PM and new-onset asthma (Young et al. 2014), respiratory symptoms (Balmes et al. 2014; Young et al. 2014), hospitalizations and emergency department visits (Bell et al. 2008; Belleudi et al. 2010; Dominici et al. 2006), and death (Dominici et al. 2000; Samet et al. 2000; Zanobetti and Schwartz 2009) among adults. The biologic effects of particulate air pollution on respiratory health are determined largely by the size and composition of the particulate air pollution, deposition of the particles in the respiratory tract, and the immunologic response to the particles (Koren 1995). Mechanisms by which exposure to particulate air pollution may exacerbate respiratory health among individuals with asthma include oxidative stress, airway inflammation, and hyperresponsiveness of the airways (Bernstein et al. 2004; Li et al. 2003; Nel et al. 2001). In recognition of the importance to public health of the effects of particulate air pollution exposures, standards such as the California Air Resources Board Air Quality Standards, European Union Directives, U.S. National Ambient Air Quality Standards, and World Health Organization Air Quality Guidelines have been used throughout the world to establish ambient air quality standards (Vahslng and Smith 2012).

A growing body of epidemiologic literature also provides initial evidence that exacerbations of adult asthma may be associated with ambient meteorological conditions, and in particular with temperature extremes (Beard et al. 2012; Fitzgerald et al. 2014; Michelozzi et al. 2009). However, the mechanisms through which outdoor temperature exposures may plausibly affect the respiratory tract remain unclear. Potential mechanisms include effects of temperature or other ambient conditions on dehydration and hyperosmolarity of the airways, which may directly induce exacerbations of asthma by triggering bronchoconstriction (Anderson and Daviskas 2000). Proposed indirect effects focus on exposures associated with temperature; for example, well-described associations between ozone (O\textsubscript{3}) and airway inflammation, particularly among adults with asthma (Hernandez et al. 2010; Khatri et al. 2009; Koren 1995), raise the possibility that the observed associations between temperature and exacerbations of asthma may be attributed, at least in part, to changes in the production of ground-level O\textsubscript{3} that are correlated with temperature and other meteorological factors (Baur et al. 2004; Camaleri et al. 2007). Higher temperatures are also associated with increased air pollutant emissions; for example, hot spells can lead to escalated use of air conditioning, thereby increasing demand on electricity-generating units, which can in turn lead to increased emissions of oxides of nitrogen (NO\textsubscript{x}) on hot days (He et al. 2013). The proposed pathways between temperature and exacerbations of asthma may also be modified by adaptive behaviors such as air conditioning use or avoidance of outdoor activities (Andrade et al. 2011). In the absence of clearly delineated mechanisms by which temperature is associated with mortality across the entire range of ambient outdoor temperatures, numerous recent studies have been designed to evaluate interactions between particulate air pollution and temperature that affect mortality, including respiratory, cardiovascular, and all-cause mortality.
PM$_{2.5}$ and asthma exacerbations among adults

To date, however, little information is available about the joint associations between particulate air pollution, ambient temperature, and respiratory morbidity. Data from the Centers for Disease Control and Prevention’s (CDC’s) Environmental Public Health Tracking Network and the Behavioral Risk Factor Surveillance System (BRFSS) adult Asthma Call-back Survey provide a unique opportunity to further explore such a possibility by evaluating associations between ambient PM and asthma symptoms among adults with asthma in a large and geographically diverse sample of adults. In the present study, we combined modeled county-level estimates of ambient concentrations of PM $\leq$ 2.5 μm in aerodynamic diameter (PM$_{2.5}$), ambient concentrations of O$_3$, precipitation, and air temperature with individual-level characteristics of adults with asthma to evaluate associations between PM$_{2.5}$ and asthma exacerbations in the United States during the period 2006–2010 and to describe the extent to which the observed associations may vary by air temperature. To accomplish these objectives, our study included two major sets of analyses. First, we conducted a main analysis of associations between PM$_{2.5}$ and asthma exacerbations in the United States that you had asthma?” The Asthma Call-back question: “Have you ever been told by a physician, nurse, or other health professional that you had asthma?” The Asthma Call-back Survey respondents. The sample was generated by pooling data collected from Asthma Call-back Surveys conducted in 2006, 2007, 2008, 2009, and 2010. The pooled sample included 74,209 respondents from 42 geographic areas of the United States (40 states, the District of Columbia, and Puerto Rico). We limited our analysis to respondents with active asthma ($n = 56,509; 76\%$). We then excluded respondents for whom county- and date-linked air quality data were unavailable ($n = 3,200, 41\%$ to 71\% in 2006, 36\% to 72\% in 2007, 35\% to 68\% in 2008, 36\% to 66\% in 2009, and 31\% to 67\% in 2010 (Mirabelli et al. 2014; National Asthma Control Program 2011a, 2011b, 2012). The Asthma Call-back Survey is exempt from Institutional Review Board (IRB) review at the CDC; state-specific IRB requirements apply to each of the participating states, the District of Columbia, and Puerto Rico. The protocol for the present analysis was reviewed and determined to be exempt from IRB review at the CDC.

**Study Sample**

For this analysis, we present results based on a sample of 50,356 Asthma Call-back Survey respondents. The sample was generated by pooling data collected from Asthma Call-back Surveys conducted in 2006, 2007, 2008, 2009, and 2010. The pooled sample included 74,209 respondents from 42 geographic areas of the United States (40 states, the District of Columbia, and Puerto Rico). We limited our analysis to respondents with active asthma ($n = 56,509; 76\%$). We then excluded respondents for whom county- and date-linked air quality data were unavailable ($n = 3,200, 41\%$ to 71\% in 2006, 36\% to 72\% in 2007, 35\% to 68\% in 2008, 36\% to 66\% in 2009, and 31\% to 67\% in 2010 (Mirabelli et al. 2014; National Asthma Control Program 2011a, 2011b, 2012). The Asthma Call-back Survey is exempt from Institutional Review Board (IRB) review at the CDC; state-specific IRB requirements apply to each of the participating states, the District of Columbia, and Puerto Rico. The protocol for the present analysis was reviewed and determined to be exempt from IRB review at the CDC.

**Methods**

**Asthma Call-back Survey**

We conducted these analyses using data from the BRFSS adult Asthma Call-back Survey for the years 2006–2010. The BRFSS is a state-level survey of the adult civilian, noninstitutionalized population $\geq$ 18 years of age that is conducted annually in the United States (CDC 2009). BFRSS survey interviews are conducted throughout the year (i.e., January through December). The Asthma Call-back Survey is a follow-up telephone survey conducted approximately 2 weeks after the BRFSS among respondents who indicated that they have ever had asthma. Respondents reported ever having had asthma by responding “yes” to the following question: “Have you ever been told by a doctor, nurse, or other health professional that you had asthma?” The Asthma Call-back Survey was administered to 10,801 respondents in 2006; 15,245 respondents in 2007; 15,007 in 2008; 15,403 in 2009; and 17,753 in 2010. In the participating areas included in our analysis, the Council of American Survey and Research Organization response rates for the Asthma Call-back Survey ranged from 41\% to 71\% in 2006, 36\% to 72\% in 2007, 35\% to 68\% in 2008, 36\% to 66\% in 2009, and 31\% to 67\% in 2010 (Mirabelli et al. 2014; National Asthma Control Program 2011a, 2011b, 2012). The Asthma Call-back Survey is exempt from Institutional Review Board (IRB) review at the CDC; state-specific IRB requirements apply to each of the participating states, the District of Columbia, and Puerto Rico. The protocol for the present analysis was reviewed and determined to be exempt from IRB review at the CDC.

**County-level Estimates of Environmental Variables**

For this analysis, we linked data from the Asthma Call-back Survey with county-level estimates of environmental variables. In the United States, counties (or equivalent entities such as boroughs, parishes, and independent cities) are the legally defined political and administrative units within each state (U.S. Census Bureau 1994). Among the geographic areas included in our analysis, the number of counties (or equivalent entities, hereafter referred to as “counties”) ranged from 1, in the District of Columbia, to 254, in Texas; in total, we linked environmental data from 2,253 U.S. counties for this analysis. Daily estimates of PM$_{2.5}$ and O$_3$ generated using a Bayesian space-time Downscaler fusion model (Berrocal et al. 2012). The Downscaler modeling approach combines output from the Community Multi-scale Air Quality (CMAQ) model (Byun and Schere 2006; Foley et al. 2010) with measurements from the U.S. Environmental Protection Agency’s (EPA’s) Air Quality System (CDC 2013; U.S. EPA 2014) to yield air quality predictions at specific geographic locations. The CMAQ model is a multi-pollutant, multi-scale chemical transport model used to generate air quality predictions at user-defined spatio-temporal scales, taking into account land use, chemical transport, chemistry, emission processes, land use, and weather (Byun and Schere 2006; Foley et al. 2010). Descriptions of the theory, development, and initial evaluation of the downscaler model have been published previously (Berrocal et al. 2010a, 2010b; Foley et al. 2010). Daily predictions of 24-hr average PM$_{2.5}$ concentrations in micrograms/cubic meter and daily maximum 8-hr average O$_3$ concentrations in parts per billion were generated at 2010 U.S. census-tract centroid locations (U.S. Census Bureau 2012) using Downscaler software (Heaton et al. 2012). In addition, daily county-level estimates of PM$_{2.5}$ and O$_3$ were generated using a population-weighted approach in which census-tract population counts were used to weight daily census tract–level PM$_{2.5}$ and O$_3$ predictions (Ivy et al. 2008; Vaidyanathan et al. 2013). County-level estimates of precipitation in millimeters and ambient air temperature in degrees Fahrenheit (°F) were generated using meteorological predictions from the North American Land Data Assimilation System Phase 2 model (Mitchell et al. 2004) at 0.125° spatial resolution (i.e., 14 × 14 km).

At the time these analyses were conducted, estimates of environmental variables were not available for interview dates in 2011 and, as noted above, all survey respondents for whom environmental data were missing were excluded from analysis. For each of the remaining respondents in our analysis, we assigned 14-day estimates of PM$_{2.5}$ and O$_3$ concentrations estimated for the 14-day period leading up to and including the date on which the Asthma Call-back Survey interview was conducted. For PM$_{2.5}$, we assigned a county-level average of the daily 24-hr average PM$_{2.5}$ concentrations estimated for the 14-day period leading up to and including the day of the interview. Without an a priori hypothesis about the best metric of PM$_{2.5}$ to use, we categorized the distribution of 14-day county-level average PM$_{2.5}$ using three metrics: a) quartiles (quartile 1, 4.00–7.06 μg/m$^3$; quartile 2, 7.07–8.97 μg/m$^3$;
Mirabelli et al.

Estimates of O₃, precipitation, and temperature were each assigned using county-level averages of values estimated for the 14-day period leading up to and including the day of the interview. For O₃, we assigned a county-level average of the daily 8-hr maximum O₃ concentrations estimated for the 14-day period leading up to and including the day of the interview. Our final models included O₃ parameterized using deciles of the distribution, with the 6th decile (range: 38.6–41.4 ppb) as the referent category; estimated precipitation following categories: 0.0 mm, 0.1–1.1 mm, 1.2–3.0 mm, and ≥ 3.1 mm, with 0.0 mm as the referent category; and temperature were each assigned using county-level average of values estimated for the National Climatic Data Center (Karl and Knight 1985) a single continuous measure of PM₂.₅.

Other Covariates

Demographic covariates used in this analysis include age, educational attainment, race/ethnicity, and sex. Cigarette smoking status was categorized as current smoker, former smoker, and lifetime nonsmoker. The states represented in our final sample were grouped into eight U.S. climate regions classified by the National Climatic Data Center (Karl and Koss 1984; National Climatic Data Center 2014): central (Illinois, Indiana, Missouri, Ohio, West Virginia), east north central (Iowa, Michigan, Wisconsin), northeast (Connecticut, District of Columbia, Maine, Maryland, Massachusetts, New Hampshire, New Jersey, New York, Pennsylvania, Rhode Island, Vermont), northwest (Oregon, Washington), south (Kansas, Louisiana, Mississippi, Oklahoma, Texas), southeast (Alabama, Florida, Georgia, Virginia), southwest (Arizona, Colorado, New Mexico, Utah), west (California, Nevada), and west north central (Montana, Nebraska, North Dakota).

The urbanicity of each county in our final sample was categorized as rural, suburban, or urban using 2013 Economic Research Service rural-urban continuum codes published by the U.S. Department of Agriculture (USDA 2013), with counties not adjacent to a metro area categorized as rural, counties adjacent to a metro area categorized as suburban, and counties in metro areas categorized as urban.

Statistical Analysis

Demographic characteristics and smoking status are presented for the sample of respondents and the weighted population estimate. Weighted population estimates were generated using adjusted sampling weights applied to account for BRFS and Asthma Call-back Survey nonresponse and unequal sampling probabilities. Annual Asthma Call-back Survey sampling weights were provided with the Asthma Call-back Survey data. Because we pooled data collected from 2006 through 2010 and because the number of geographic areas with Asthma Call-back Survey respondents varied from year to year, we readjusted the sampling weights in each geographic area by dividing the annual Asthma Call-back Survey weights by the number of years for which data were available (Mirabelli et al. 2014). All descriptive analyses were performed using survey procedures (e.g., PROC SURVEYFREQ) for the analysis of complex survey data in SAS v.9.3 (SAS Institute, Inc.).

Using additive binomial models specified with an identity link, we estimated the prevalence of asthma symptoms during the past 14 days for the entire weighted population estimate and across categories of individual- and county-level characteristics, accounting for complex survey sampling. Associations between PM₂.₅ and asthma symptoms during the past 14 days were evaluated using PROC SURVEYREG in SAS v.9.3, with standard errors generated by adapting a two-step approach developed and published by Natarajan and colleagues designed to generate robust variance estimates to fit the binomial error distribution of our data (Natarajan et al. 2008; Slade et al. 2012). Because we conducted our analyses using statistical software designed to analyze complex survey data, all models accounted for the state-based sampling approach and our pooling of 5 years of survey data. Models were adjusted for individual-level covariates (age, educational attainment, race/ethnicity, sex, and smoking status) and county-level covariates (O₃, precipitation, region, temperature, and urbanicity). After performing these analyses, we conducted two additional analyses in which we replaced the measure of ambient air temperature, initially based on the mean of the distribution of daily maximum temperatures during the 14-day period, with the median of the distribution and with the mean apparent temperature during the 14-day period.

To evaluate the extent to which the observed associations between PM₂.₅ and asthma symptoms in the past 14 days varied by temperature, we stratified adjusted models into quintiles of the distribution of daily maximum ambient air temperature [11.1–14.4°F (−17.2 to 6.8°C), 14.5–58.6°F (6.9–16.8°C), 58.7–70.1°F (14.9–21.1°C), 70.2–80.5°F (21.2–26.9°C), and 80.6–112.4°F (27.0–44.7°C)].

Measures of association are presented as adjusted percent differences (PDs) with 95% confidence intervals (CIs). For analyses in which PM₂.₅ was evaluated using quartiles, PDs are interpreted as arithmetic differences in the prevalence of asthma symptoms in PM₂.₅ quartiles 2, 3, and 4 compared with the prevalence of asthma symptoms in referent quartile 1. For analyses of linear spline segments and analyses in which PM₂.₅ was included as a single continuous measure, PDs are interpreted as the change in prevalence of asthma symptoms per 1 μg/m³ increase in PM₂.₅. All analyses were conducted using SAS v9.3 (SAS Institute, Inc., Cary, North Carolina).

Results

Characteristics of the 50,356 adults with active asthma and the weighted population estimate are shown in Table 1. Among adults with active asthma, an estimated 57.1% reported asthma symptoms within the past 14 days. Variations in percentages of adults reporting asthma symptoms within the past 14 days were observed across categories of respondent characteristics as well as by U.S. climate region and county-level urbanicity.

Summary statistics describing the distributions of PM₂.₅, O₃, temperature, and precipitation as well as Pearson’s correlation coefficients indicating correlations between the four measures are shown in Table 2. Despite the small magnitudes of the correlations, all pairwise correlation coefficients were statistically significant at < 0.05. The distributions of PM₂.₅, O₃, precipitation, and temperature for the weighted population estimate of nearly 18.0 million adults with active asthma and...
the estimated percentages of adults with active asthma reporting asthma symptoms in the past 14 days across categories of each environmental measure are shown in Figure 1. Across quartiles of PM$_{2.5}$, the percentages of adults reporting asthma symptoms in the past 14 days were 54.2% ± 1.2 in quartile 1, 58.4% ± 1.0 in quartile 2, 58.3% ± 1.0 in quartile 3, and 56.7% ± 1.1 in quartile 4. Table 3 shows estimates of the difference in the percentage of adults with active asthma who experienced symptoms in the past 14 days; the values were generated using unadjusted models and models adjusted for individual- and county-level covariates. Broadly speaking, estimates generated using models adjusted for individual-level covariates were similar to those generated using unadjusted models, and estimates generated using models adjusted for county-level covariates were similar to those generated using fully adjusted models. In fully adjusted models, when PM$_{2.5}$ exposure was parameterized using indicators of PM$_{2.5}$ quartile, the estimated prevalences of asthma symptoms in the past 14 days were typically 4–5% higher in quartiles 2, 3, and 4 than the prevalence in quartile 1. In models adjusted for individual- and county-level covariates in which PM$_{2.5}$ was parameterized as four linear spline segments, the pattern observed in our main analysis was also observed across quintiles of temperature. Spline segment 1 was associated with positive point estimates of the PD per microgram/cubic meter of PM$_{2.5}$ across categories of temperature ranging from 1.1°F to 80.5°F (–17.2 to 26.9°C) [1.1–44.4°F (–17.2 to 6.8°C): 7.9%; 44.5–58.6°F (6.9–14.7°C): 6.9%; 58.7–70.1°F (14.8–21.1°C): 2.9%; 70.2–80.5°F (21.2–26.9°C): 7.3%]. Per-microgram/cubic meter changes in symptom prevalence in spline segments 2 through 4 were consistent with the null value of 0.0%. Minimal variation was observed across categories of temperature when PM$_{2.5}$ was evaluated as a single continuous measure.

Table 1. Characteristics of the study sample and the population estimate, with percentages reporting asthma symptoms in the past 14 days.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Survey sample</th>
<th>Weighted population estimate</th>
<th>Asthma symptoms$^a$</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>$n$</td>
<td>$n^*$</td>
<td>Percent (95% CI)</td>
</tr>
<tr>
<td>Total</td>
<td>50,356</td>
<td>17,963</td>
<td>57.1 ± 0.5</td>
</tr>
<tr>
<td>Age, years</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>18–34</td>
<td>5,430</td>
<td>5,597</td>
<td>54.2 ± 0.5</td>
</tr>
<tr>
<td>35–44</td>
<td>6,653</td>
<td>3,318</td>
<td>56.7 ± 1.1</td>
</tr>
<tr>
<td>45–54</td>
<td>11,237</td>
<td>3,515</td>
<td>59.1 ± 0.9</td>
</tr>
<tr>
<td>55–64</td>
<td>13,075</td>
<td>2,833</td>
<td>59.5 ± 1.0</td>
</tr>
<tr>
<td>65–99</td>
<td>13,961</td>
<td>2,701</td>
<td>60.4 ± 0.8</td>
</tr>
<tr>
<td>Educational attainment</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Less than high school</td>
<td>4,795</td>
<td>1,817</td>
<td>63.2 ± 1.6</td>
</tr>
<tr>
<td>Graduated high school</td>
<td>13,372</td>
<td>4,628</td>
<td>60.2 ± 1.2</td>
</tr>
<tr>
<td>College 1–3 years/technical school</td>
<td>15,062</td>
<td>5,120</td>
<td>59.5 ± 1.0</td>
</tr>
<tr>
<td>College ≥ 4 years</td>
<td>17,127</td>
<td>6,288</td>
<td>51.4 ± 0.8</td>
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<td>Race/ethnicity</td>
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<tr>
<td>White, non-Hispanic</td>
<td>42,143</td>
<td>13,415</td>
<td>58.3 ± 0.6</td>
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<tr>
<td>Black, non-Hispanic</td>
<td>3,105</td>
<td>1,656</td>
<td>55.4 ± 1.9</td>
</tr>
<tr>
<td>Other, non-Hispanic</td>
<td>2,812</td>
<td>1,161</td>
<td>61.8 ± 2.5</td>
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<tr>
<td>Hispanic</td>
<td>2,296</td>
<td>1,731</td>
<td>48.8 ± 2.4</td>
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<tr>
<td>Sex</td>
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<tr>
<td>Female</td>
<td>36,995</td>
<td>11,244</td>
<td>58.0 ± 0.6</td>
</tr>
<tr>
<td>Male</td>
<td>13,361</td>
<td>6,719</td>
<td>55.8 ± 1.0</td>
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<tr>
<td>Smoking status</td>
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<tr>
<td>Current smoker</td>
<td>9,261</td>
<td>3,473</td>
<td>69.8 ± 1.1</td>
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<td>Former smoker</td>
<td>17,085</td>
<td>4,877</td>
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<tr>
<td>Lifetime nonsmoker</td>
<td>24,010</td>
<td>9,513</td>
<td>51.3 ± 0.8</td>
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<tr>
<td>U.S. climate region</td>
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<tr>
<td>Central</td>
<td>5,901</td>
<td>2,687</td>
<td>60.8 ± 1.2</td>
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<tr>
<td>East north central</td>
<td>4,575</td>
<td>1,401</td>
<td>61.2 ± 1.2</td>
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<td>Northeast</td>
<td>13,481</td>
<td>4,582</td>
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<td>Northwest</td>
<td>5,943</td>
<td>741</td>
<td>56.9 ± 1.2</td>
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<tr>
<td>South</td>
<td>6,478</td>
<td>2,226</td>
<td>62.1 ± 1.6</td>
</tr>
<tr>
<td>Southeast</td>
<td>3,442</td>
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<td>50.9 ± 1.6</td>
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<tr>
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<td>3,778</td>
<td>1,029</td>
<td>60.7 ± 1.8</td>
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<tr>
<td>West</td>
<td>2,587</td>
<td>2,676</td>
<td>46.0 ± 1.7</td>
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<tr>
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<td>4,171</td>
<td>198</td>
<td>57.9 ± 1.5</td>
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<tr>
<td>Urbanicity</td>
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<tr>
<td>Rural</td>
<td>7,329</td>
<td>1,007</td>
<td>63.1 ± 1.5</td>
</tr>
<tr>
<td>Suburban</td>
<td>9,238</td>
<td>2,122</td>
<td>65.1 ± 1.2</td>
</tr>
<tr>
<td>Urban</td>
<td>33,789</td>
<td>14,834</td>
<td>55.6 ± 0.6</td>
</tr>
</tbody>
</table>

Notes: CI, confidence interval; SE, standard error of the mean.

*in the past 14 days.

Table 2. Summary statistics and correlation coefficients describing the distributions of PM$_{2.5}$, O$_3$, precipitation, and temperature.

<table>
<thead>
<tr>
<th>Exposure</th>
<th>Average daily mean ± SD$^a$</th>
<th>Percentile$^a$ 25th</th>
<th>Percentile$^a$ 50th</th>
<th>Percentile$^a$ 75th</th>
<th>PM$_{2.5}$</th>
<th>O$_3$</th>
<th>Precipitation</th>
<th>Temperature[°F/°C]</th>
</tr>
</thead>
<tbody>
<tr>
<td>PM$_{2.5}$ (μg/m$^3$)</td>
<td>50.356</td>
<td>9.4 ± 3.1</td>
<td>71</td>
<td>9.0</td>
<td>11.4</td>
<td>0.03</td>
<td>–0.02</td>
<td>–0.02</td>
</tr>
<tr>
<td>O$_3$ (ppb)</td>
<td>50.356</td>
<td>30.7 ± 9.7</td>
<td>31.2</td>
<td>38.6</td>
<td>45.7</td>
<td>1</td>
<td>–0.11</td>
<td>0.57</td>
</tr>
<tr>
<td>Precipitation (mm)</td>
<td>50.356</td>
<td>2.8 ± 0.8</td>
<td>0.8</td>
<td>2.0</td>
<td>3.7</td>
<td>1</td>
<td>–0.03</td>
<td>1.0</td>
</tr>
<tr>
<td>Temperature[°F/°C]</td>
<td>50.356</td>
<td>62.9 ± 17.2</td>
<td>48.5</td>
<td>64.6</td>
<td>77.9</td>
<td>1</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Notes: O$_3$, ozone; PM$_{2.5}$, particulate matter ≤ 2.5 μm in diameter; SD, standard deviation.

$^a$Based on unweighted survey data.
Discussion
Our findings suggest that among adults with asthma, the prevalence of self-reported asthma symptoms during the past 14 days was 4–5% higher among respondents with a 14-day average concentration of PM$_{2.5}$ > 7.07 μg/m$^3$ than the prevalence among respondents with 14-day averages in the range of 4.00–7.06 μg/m$^3$. In this lowest category of PM$_{2.5}$, each 1 μg/m$^3$ increase in PM$_{2.5}$ was associated with a 3% increase in the prevalence of asthma symptoms. Taken together, our results raise the possibility that among individuals with asthma, exacerbations may begin to increase at relatively low levels of ambient PM$_{2.5}$. Stratification of these results across categories of temperature suggests that these findings may be driven largely by effects observed at temperatures < 80.6°F (27.0°C).

These results expand on previous research that evaluated relationships between PM$_{2.5}$ and asthma symptoms across quartiles of the distribution of PM$_{2.5}$ (Mirabelli et al. 2015) by exploring additional metrics of PM$_{2.5}$.
and refining the statistical models used to estimate the exposure–outcome relationships. Previously, we reported on quartiles of the PM$_{2.5}$ distribution and found higher percentages of adults with asthma symptoms in the past 14 days in quartiles 2, 3, and 4 compared with quartile 1 (Mirabelli et al. 2015). In considering relationships between PM$_{2.5}$ and asthma symptoms using four linear spline segments, the present analysis supports and expands our earlier findings by suggesting that each unit increase in PM$_{2.5}$ may be associated with a measurable increase in the prevalence of asthma symptoms in the past 14 days, even at levels as low as 4.00–7.06 μg/m$^3$. Indeed, using results from our fully adjusted linear spline segment model, PD estimates generated at the median of each spline segment support

**Figure 2.** Associations between particulate matter ≤ 2.5 μm in diameter (PM$_{2.5}$) and the prevalence of asthma symptoms in the past 14 days among adults with active asthma across quintiles of air temperature. Results are shown for temperature category-stratified models in which PM$_{2.5}$ is parameterized as quartiles (A), four linear spline segments (B), and a single continuous measure (C). All models are adjusted for individual- and county-level covariates. Formula for conversion from degrees Fahrenheit ($T_F$) to degrees Celsius ($T_C$): $T_C = (T_F - 32) \times (5/9)$. 
the results of our analysis using quartiles. For example, where models using quartiles of PM$_{2.5}$ generated PDs of 4.4%, 4.7%, and 4.9% in quartiles 2, 3, and 4, respectively, models using linear spline segments generated PDs of 4.5%, 5.2%, and 5.7% (data not shown). Both models suggest that among individuals with asthma, exacerbations may begin to increase at levels of PM$_{2.5} < 7.07$ $\mu$g/m$^3$. The strength of this association at levels between 4.00 and 7.06 $\mu$g/m$^3$ would have been missed had we only evaluated the relationship between PM$_{2.5}$ and asthma symptoms using a single continuous measure of PM$_{2.5}$, which generated a 0.5% increase in the prevalence of asthma symptoms with each 1 $\mu$g/m$^3$ increase in PM$_{2.5}$.

At the present time, there are few population-based studies of PM$_{2.5}$ and asthma morbidity outcomes with which to contrast our results. Following an evaluation of associations between daily minimum temperature and respiratory hospitalizations and emergency department visits, investigators reported that ambient concentrations of PM$_{10} \leq 10$ $\mu$m in aerodynamic diameter (PM$_{10}$) modified the association between temperature and respiratory hospitalizations but not emergency department visits, and that the observed associations with increasing temperatures were more pronounced when concentrations of PM$_{10}$ also increased (Ren et al. 2006). Combined with convincing evidence of the effects of ambient PM exposures on the respiratory health of adults [Health Effects Institute (HEI) Panel on the Health Effects of Traffic-Related Air Pollution 2010], our findings that the magnitudes of the associations between PM$_{2.5}$ and asthma symptoms are not constant across the range of 4.00–19.98 $\mu$g/m$^3$ of PM$_{2.5}$ and that the associations vary across categories of ambient air temperature support and extend previous evidence that ambient particulate matter pollution and temperature may act jointly to affect respiratory health in manners that may be evident before death (Ren et al. 2006).

Several aspects of our study merit careful consideration when interpreting our findings. First, a comparison of the results shown in Table 1 suggests that the relationship between PM$_{2.5}$ and the prevalence of asthma symptoms in the past 14 days may be confounded by county-level factors, including O$_3$, precipitation, and temperature. Despite including these and other county-level covariates in our analysis, our results may be affected by residual confounding, including within categories of ambient air temperature or by geographic differences within climate regions. We were unable to stratify our results into smaller categories of air temperature. Earlier research designed to estimate the effects of ambient air pollutant exposures on health initially suggested that statistical models that include one or two weather variables, such as temperature and humidity, may not be adequate to fully control for the effects of weather (Pope and Kalkstein 1996). From subsequent research designed to evaluate the extent to which statistical models of the relationship between particulate pollution and mortality that simultaneously account for the effects of weather or temperature may yield biased results if incorrect metrics of weather or temperature are used, investigators reported finding little evidence that the choice of metrics influenced findings (Samet et al. 1998). Results generated from our main models, which included temperature and O$_3$ using indicators for deciles of the distributions and precipitation using indicators for tertiles of the distribution, may not have adequately accounted for the complex relationships between air quality, weather, and health, but they were not notably changed when we considered alternative parameterizations of O$_3$, temperature, and precipitation (not shown). Improvements in our understanding of the complex relationships between air quality, weather, and health would improve our ability to incorporate these relationships, including nonlinear relationships and relationships at temperature extremes (Pope and Kalkstein 1996), into our analyses of the relationship between air quality and asthma exacerbations.

Second, modeled exposures provide estimates of PM$_{2.5}$, O$_3$, precipitation, and temperature for geographic areas in which respondents live. Assigning county-level exposure measures may have resulted in exposure misclassification if respondents’ true exposures were markedly different from those assigned to the counties in which they lived. The modeling approach applied in our study advantageously provided estimates for geographic areas without adequate measurements of air pollutants or meteorological parameters. However, uncertainties associated with estimates of exposures derived using modeled data could result in exposure misclassification, which is not incorporated into our statistical models, and we were unable to take into account a wide range of other air pollutants or PM of other sizes (e.g., coarse or ultrafine). When concentrations of multiple air pollutants are highly correlated, results generated from models that do not include each of the multiple pollutants may be confounded by the effects of other pollutants not considered, and we are unable to evaluate the extent to which our findings may represent relationships between other air pollutants and exacerbations of asthma. Nonetheless, the integration of data from a large, population-based survey of adults with asthma with environmental data estimated at the county level enabled us to evaluate the relationship between county-level PM$_{2.5}$ and asthma symptoms in a large population of U.S. adults.

Our analyses did not account for indoor exposures, filtration of indoor air, air exchange rates, occupational exposures, pollen concentrations, or other factors affecting personal exposure. We did not have information about the extent to which respondents may have been included in two or more annual sample populations. Our analyses did not take into account the timing, including day of the week, month, or season, of the survey interview. In these data, correlations between interview month, U.S. climate region, and temperature prevented our statistical models from converging when all three factors were included. Similarly, imprecise estimates within each state prevented us from stratifying our analysis by state. When using these data, we were unable to evaluate temporal relationships between exposures and asthma symptoms within the 14-day period; therefore, we cannot draw conclusions about the extent to which symptoms may have preceded or followed peak exposures. Furthermore, our analyses did not incorporate information about the changes in exposure to outdoor air or changes in behaviors that may occur at temperature extremes. If any behavioral changes, such as closing windows, using air conditioning, and staying indoors during hot weather, reduced exposures to ambient air pollutants, then our observation of attenuated associations at temperatures $> 80.6^\circ$F (27.0°C) is unsurprising. In addition to the direct effects of exposure to ambient particulate matter pollution, susceptibility to the effects of exposure may be linked to other factors associated with asthma, such as diet and obesity, indoor air quality, medication use, socioeconomic status, and stress (Guarnieri and Balmes 2014). Some populations, such as those living in neighborhoods located near highways or affected by industrial air emissions, may experience exposures higher than county averages and be more vulnerable to the effects of poor air quality. Additional information about other factors (e.g., pollen and other allergens) associated with exacerbation of asthma and about the frequency and severity of the asthma exacerbations would improve our ability to draw conclusions about the role of exposures to PM$_{2.5}$ in ambient air.

**Conclusion**

Despite the limitations described above, our findings extend the present understanding of the health effects of PM$_{2.5}$ by considering self-reported asthma exacerbations rather than more severe outcomes such as hospital encounters or mortality. Health effects associated with PM$_{2.5}$ exposures are wide-ranging and may have measurable...
impacts on outcomes not considered in this analysis, including indicators of asthma severity, symptom frequency, medication use, functional consequences of asthma, and other cardiovascular and respiratory conditions. Our results suggest that the relationship between PM$_{2.5}$ and asthma may not be constant across the entire range of PM$_{2.5}$ concentrations, and as a consequence, when the exposure of interest is PM$_{2.5}$ in ambient air, changes in population-level metrics of asthma exacerbations may occur more noticeably in the range of 4.00–7.06 $\mu$/m$^3$. These findings provide novel information about the importance of county-level air quality for the nearly 18 million adults with asthma represented by the sampled survey population of Asthma Call-back Survey respondents.

**REFERENCES**


