Systematic Reviews and Meta- and Pooled Analyses

Who is More Affected by Ozone Pollution? A Systematic Review and Meta-Analysis

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Ozone is associated with adverse health; however, less is known about vulnerable/sensitive populations, which we refer to as sensitive populations. We systematically reviewed epidemiologic evidence (1988–2013) regarding sensitivity to mortality or hospital admission from short-term ozone exposure. We performed meta-analysis for overall associations by age and sex; assessed publication bias; and qualitatively assessed sensitivity to socioeconomic indicators, race/ethnicity, and air conditioning. The search identified 2,091 unique papers, with 167 meeting inclusion criteria (73 on mortality and 96 on hospitalizations and emergency department visits, including 2 examining both mortality and hospitalizations). The strongest evidence for ozone sensitivity was for age. Per 10-parts per billion increase in daily 8-hour ozone concentration, mortality risk for younger persons, at 0.60% (95% confidence interval (CI): 0.40, 0.80), was statistically lower than that for older persons, at 1.27% (95% CI: 0.76, 1.78). Findings adjusted for publication bias were similar. Limited/suggestive evidence was found for higher associations among women; mortality risks were 0.39% (95% CI: −0.22, 1.00) higher than those for men. We identified strong evidence for higher associations with unemployment or lower occupational status and weak evidence of sensitivity for racial/ethnic minorities and persons with low education, in poverty, or without central air conditioning. Findings show that some populations, especially the elderly, are particularly sensitive to short-term ozone exposure.

age; air pollution; effect modifiers; hospitalization; mortality; ozone; sex

Abbreviations: CI, confidence interval; CVD, cardiovascular disease; SES, socioeconomic status.

Ozone is a common air pollutant associated with adverse health outcomes, including mortality (1). In the recently published Global Burden of Disease (GBD) Study, Lim et al. (2) estimated almost 2.5 million disability-adjusted life years attributable to ozone in 2010 worldwide. The Global Burden of Disease Study is the most comprehensive research on health burdens worldwide to date, based on a 5-year project involving about 500 researchers. However, the analysis assumed that all persons have identical sensitivity to ozone, by applying a single concentration-response function (3). The authors noted that such assumptions obscure potentially vast differences in health risks across regions and populations. Estimating ozone risks for different populations requires evidence regarding which subpopulations are most sensitive.

Here we refer to “sensitive populations” broadly as individuals or communities with higher health risk due to susceptibility or vulnerability. “Susceptibility” is often used to describe elevated health risk due to biological or other intrinsic factors, such as sex or genetics, whereas “vulnerability” often refers to higher risk from nonbiological or external factors, such as socioeconomic status (SES) or occupation (4). Sensitivity may relate to modified exposure (e.g., different risks by air conditioning status) or different health responses from the same exposure across individuals (e.g., different risks by sex). These population characteristics or factors are also called effect modifiers of the ozone-health relationship. To date, there is no consensus on who is more or less sensitive to health impacts associated with short-term exposure to ozone.

While evidence that ozone adversely affects health is strong and consistent (5–8), results regarding susceptibility and vulnerability of exposed populations are inconclusive. Such evidence would aid efforts to quantify health risks across heterogeneous populations, such as the Global Burden of Disease Study. Understanding which populations are
sensitive could inform knowledge on credible pathological mechanisms. Assessing susceptibility to air pollutants is a priority research area for the US Environmental Protection Agency (9, 10), which sets regulations with a margin of safety for sensitive individuals. Physicians would benefit from information on which patients are most likely to be

Table 1. Scientific Evidence on Populations That May Be Sensitive to Associations of Ozone Exposure With Mortality and Hospital Admissions, 1988–2013

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Statistically Significant Results</th>
<th>Lack of Statistically Significant Evidence</th>
<th>Evidence of Association</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>No. of Studies</td>
<td>Reference No(s.)</td>
</tr>
<tr>
<td>Sex</td>
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<tr>
<td>Mortality</td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher risk in women</td>
<td>2</td>
<td>22, 23</td>
</tr>
<tr>
<td>Hospitalization</td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher risk in women</td>
<td>5</td>
<td>24–28</td>
</tr>
<tr>
<td></td>
<td>Higher risk in men</td>
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<td>28</td>
</tr>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mortality</td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher risk with higher age</td>
<td>11</td>
<td>22, 23, 45, 71, 72, 89–94</td>
</tr>
<tr>
<td></td>
<td>Lower risk with higher age</td>
<td>4</td>
<td>30–32, 93</td>
</tr>
<tr>
<td></td>
<td>Community data</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>Hospitalization</td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher risk with lower age</td>
<td>9</td>
<td>28, 33–36, 38–40, 84</td>
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<tr>
<td>Race/Ethnicity</td>
<td>Mortality</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Community data</td>
<td>1</td>
<td>42</td>
</tr>
<tr>
<td></td>
<td>Higher risk for higher minority population</td>
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<td>41</td>
</tr>
<tr>
<td>Hospitalization (individual data)</td>
<td>Mortality</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher for minority populations</td>
<td>2</td>
<td>27, 135</td>
</tr>
<tr>
<td></td>
<td>Lower for minority populations</td>
<td>1</td>
<td>27</td>
</tr>
<tr>
<td>Education</td>
<td>Mortality</td>
<td></td>
<td></td>
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<tr>
<td></td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Lower risk with higher education</td>
<td>1</td>
<td>45</td>
</tr>
<tr>
<td></td>
<td>Higher risk for unknown education than for known education</td>
<td>1</td>
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<td>Higher risk for known education than for unknown education</td>
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<tr>
<td>Hospitalization</td>
<td>Community data</td>
<td>0</td>
<td>2</td>
</tr>
</tbody>
</table>

Table continues
affected. Lack of scientific evidence on how different populations may respond differently to ozone also hinders understanding of health impacts from climate change (11).

We systematically reviewed population-based studies regarding which persons are most sensitive to risk of mortality and risk of hospital admission from short-term exposure to ozone. Meta-analysis was performed to generate overall risk estimates by subpopulation. Systematic reviews with meta-analyses can help decision-makers, physicians, and researchers integrate findings and identify consistencies in the scientific literature (12).

**METHODS**

We performed a systematic search in the National Library of Medicine’s MEDLINE/PubMed database (13). Search terms and exclusion criteria were designed to identify population-based studies on health impacts of short-term exposure (i.e., a day or a few days) to ambient ozone with regard to mortality, hospital admissions, and emergency room visits in adults (see Web Appendix, available at http://aje.oxfordjournals.org/). Study designs other than population-based research (e.g., chamber studies) were excluded. Both single-city and multicity studies were included.

Article titles and abstracts were reviewed in relation to exclusion criteria, and the full texts of remaining articles were then reviewed. Web Figure 1 provides a flow diagram of the search. The systematic review and meta-analysis were performed with consideration of PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) and MOOSE (Meta-Analysis of Observational Studies in Epidemiology) guidelines (14, 15).

We extracted information on each study’s time frame, location, population description, sample size, ozone metric (e.g., 24-hour average), study design (e.g., time series), lag period between exposure and health responses, health outcome, and statistical approach to assessment of ozone sensitivity (e.g., stratification). We then performed data extraction. Although we focused on populations’ sensitivities to ozone, we noted

<table>
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</thead>
<tbody>
<tr>
<td></td>
<td>No. of Studies</td>
<td>Reference No(s)</td>
<td>No. of Studies</td>
</tr>
<tr>
<td><strong>Income</strong></td>
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<td></td>
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</tr>
<tr>
<td>Mortality</td>
<td>Community data</td>
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<td>Community data</td>
</tr>
<tr>
<td>Employment/Occupation</td>
<td>Individual data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher risk for lower-level employment</td>
<td>3</td>
<td>31, 45, 46</td>
</tr>
<tr>
<td></td>
<td>Higher risk for higher-level employment</td>
<td>1</td>
<td>46</td>
</tr>
<tr>
<td></td>
<td>Community data</td>
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<td></td>
</tr>
<tr>
<td>Higher risk with higher unemployment</td>
<td>2</td>
<td>29, 41</td>
<td></td>
</tr>
<tr>
<td><strong>Poverty</strong></td>
<td>Community data</td>
<td>0</td>
<td>Community data</td>
</tr>
<tr>
<td></td>
<td>Higher risk for higher-poverty areas</td>
<td>1</td>
<td>36</td>
</tr>
<tr>
<td><strong>Air Conditioning</strong></td>
<td>Community data</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Higher risk with lower AC prevalence</td>
<td>3</td>
<td>6, 41, 42</td>
</tr>
<tr>
<td></td>
<td>Higher risk with higher AC prevalence</td>
<td>2</td>
<td>22, 42</td>
</tr>
</tbody>
</table>

Abbreviation: AC, air conditioning.
results for other potential modifiers of the association between ozone and health. We classified each potential modifier as individual-level (e.g., individual’s SES), community-level (e.g., county’s unemployment rate), or temporal (e.g., day’s temperature or season). Results were divided by health outcome (mortality, hospital admissions), grouping emergency department visits with hospitalizations. For multicity studies providing city-specific results, we examined each city’s results separately.

No studies were excluded for quality considerations; however, information on study design and presentation of results was summarized for the following characteristics: description of model structure (e.g., model structure provided, general description), consideration of co-pollutants, single or multiple lags, rationale for selected lag, exposure approach (e.g., ambient monitors), presentation of ozone sensitivity results (e.g., tables, figures), and specification of International Classification of Diseases codes and source of health data. For studies included in the meta-analysis, we extracted information on ozone-related health risk (e.g., relative risk), measure of uncertainty (e.g., confidence interval), increment of pollution for estimates of the association (e.g., 10 µg/m³), and ozone metric (e.g., 24-hour average).

Meta-analysis by random-effects modeling (16), which addresses heterogeneity in the actual effects across studies, was performed separately for sex and age groups. Population characteristics that were defined differently by study (e.g., employment defined as “percent unemployed” vs. “occupation”) could not meaningfully be combined quantitatively. Meta-analyses were considered when at least 5 studies using individual-level data were available. If studies presented estimates for multiple lags, we used the key lag presented by the study authors or the single-day lag closest to the health outcome (i.e., lag 0 if available). If multiple models had been utilized in a given study (e.g., different sets of confounding variables), we used results from the main model as presented by the original study authors. For multicity studies, city-specific estimates were included separately when available. Results based on categorical exposures (e.g., the highest quartile of exposure vs. the lowest quartile) were not incorporated into the meta-analysis. Results reported in various forms were converted to regression coefficients and their standard errors for pooling. Estimates based on specific 8-hour periods (e.g., 10 AM–6 PM) were combined with those for daily 8-hour maximum. Values from other ozone metrics were converted to daily 8-hour maximum based on standard ratios, although true ratios vary (17). Estimates presented in µg/m³ were converted to parts per billion (ppb), assuming standard temperature and pressure. Overall estimates were calculated for a 10-ppb increase in daily 8-hour maximum. We calculated the uncertainty parameter (I²) representing the percentage of total variance in study-specific results explained by heterogeneity (18). We assessed publication bias with Egger’s test for asymmetry (19) and adjusted for publication bias using the “trim and fill” method (20). Analyses were performed in R, version 2.15.2 (R Foundation for Statistical Computing, Vienna, Austria). We generated forest plots and funnel plots for each meta-analysis.

A priori, we identified population characteristics potentially relevant to ozone sensitivity: sex, age, race/ethnicity, SES indicators (education, income, employment/occupation, and poverty), and air conditioning. For these characteristics, we synthesized the overall evidence using categories loosely based on those established by Institute of Medicine committees (21) and applied by government agencies and researchers. The original categories used by the Institute of Medicine and other groups were: sufficient evidence of a causal relationship; sufficient evidence of an association; limited/suggestive evidence of an association; inadequate/insufficient evidence to determine whether an association does or does not exist; and limited/suggestive evidence of no association (21). Because our study focused on modification of associations for epidemiologic studies, we altered these categories to the following, in increasing order of certainty: no evidence of ozone sensitivity, weak evidence of ozone sensitivity, limited/suggestive evidence of ozone sensitivity, and strong evidence of ozone sensitivity. The overall state of scientific evidence for each population characteristic was assigned a category based on the quality and quantity of studies providing consistent and significant evidence in comparison with conflicting findings. For sex and age, we also considered meta-analysis results. These categories allowed qualitative synthesis of evidence for population characteristics for which meta-analysis was not viable.

RESULTS

The search identified 2,470 published articles, with 2,091 unique papers. After exclusions, 169 papers remained. Two peer-reviewed agency reports that duplicated published articles were omitted. Of the remaining 167 papers, 73 examined mortality and 96 examined hospitalizations and emergency department visits, including 2 studies that considered both mortality and hospitalization. Web Tables 1 and 2 provide information on the studies of mortality and hospital admission, respectively. Although we focused on certain population characteristics for ozone sensitivity, Web Tables 1 and 2 also show results for other potential modifiers considered in these studies. The most represented country was the United States (21% of studies), while 12% of the studies were in Taiwan and 11% were in Canada. Web Figure 2 gives study characteristics regarding analytical decisions and the presentation of study design and results.

Table 1 presents scientific evidence for selected population characteristics and provides our conclusions on ozone sensitivity. The table notes the numbers of studies that found statistically significant evidence of ozone sensitivity (and in which direction the evidence pointed) and those that did not. Below we provide evidence for ozone sensitivity with regard to sex, age, race/ethnicity, SES indicators, and air conditioning, including the results of meta-analyses for sex and age. Meta-analysis was not applied to other population characteristics because of few estimates or substantial heterogeneity in how characteristics were defined.

Modification of the association by sex

Thirty-six studies examined ozone sensitivity by sex, all using individual-level data. Risk estimates were generally higher for women, with 6 studies finding this result (2 for mortality (22, 23), 4 for hospitalization (24–27)) and 1 finding
higher hospitalization associations for women during the warm season and for men during the cool season or all year (28). Twenty-nine studies identified no significant results. We conducted meta-analysis for total mortality separately by sex, based on 9 pairs of risk estimates from 9 studies (Table 2, Figures 1 and 2). For women, a 10-ppb increase in daily 8-hour ozone concentration was associated with a 1.12% (95% confidence interval: 0.62, 1.63) increase in mortality—slightly higher than the estimate for men at 0.73% (95% CI: 0.40, 1.07) (Figure 3). The ozone-mortality relative risk was 0.39% (95% CI: −0.22, 1.00) higher for women than for men. Estimates adjusted for publication bias were similar. We found limited/suggestive evidence of higher associations in women than in men.

Table 2. Results From a Meta-Analysis of Associations Between Ozone Exposure and Mortality, by Sex, 1988–2013

<table>
<thead>
<tr>
<th></th>
<th>I² (Uncertainty Parameter)</th>
<th>95% CI</th>
<th>P Value (Egger’s Regression Test)</th>
<th>Overall Risk Estimate</th>
<th>95% CI</th>
<th>Overall Estimate Adjusted for Publication Bias,</th>
<th>95% CI</th>
<th>No. of Adjusted Estimates</th>
</tr>
</thead>
<tbody>
<tr>
<td>Men</td>
<td>26.4</td>
<td>0.0, 65.6</td>
<td>0.02</td>
<td>0.73</td>
<td>0.40, 1.07</td>
<td>0.64</td>
<td>0.31, 0.98</td>
<td>2</td>
</tr>
<tr>
<td>Women</td>
<td>64.7</td>
<td>27.8, 82.7</td>
<td>0.003</td>
<td>1.12</td>
<td>0.62, 1.63</td>
<td>1.12</td>
<td>0.62, 1.63</td>
<td>0</td>
</tr>
<tr>
<td>Women vs. men</td>
<td>0.39</td>
<td>−0.22, 1.00</td>
<td>0.48</td>
<td>−0.13, 1.09</td>
<td>0</td>
<td>0</td>
<td></td>
<td>0</td>
</tr>
</tbody>
</table>

Abbreviation: CI, confidence interval.

a Percentage of variance in observed study results explained by heterogeneity.
b Percent increase in risk for a 10-ppb increase in 8-hour ozone concentration.
c Percent increase in ozone risk estimates for women compared with men.

Modification of the association by age

Ozone sensitivity by age was investigated in 46 mortality studies and 43 hospitalization studies. Categorization of age differed by study. All studies used individual-level data on age, except for 1 mortality study that found no evidence that risk varied by communities’ percentage of persons over age 75 years (29). Of mortality studies using individual-level data for age, 10 found risk increases with age, and 1 additional study found higher or lower risks for older persons depending on season and cause of death (Table 1). Three studies finding lower risk for older persons were based on subpopulations (e.g., survivors of myocardial infarction) (30–32).

Figure 1. Percent increase in risk of total mortality for a 10-ppb increase in 8-hour ozone concentration for published studies included in a meta-analysis, by sex, 1988–2013. The open points represent individual study estimates; the closed points and dashed vertical lines represent overall estimates from the meta-analysis. Horizontal lines represent 95% confidence intervals.

Figure 2. Estimates of the association between ozone exposure and total mortality for published studies included in a meta-analysis, by sex, 1988–2013. Estimates show the percentage increase in risk per 10-ppb increase in 8-hour ozone concentration. The open points represent individual study estimates; the closed points represent “missing studies” for which data were derived from the trim-and-fill method to adjust for publication bias. Solid vertical lines represent overall estimates based on study results; dashed lines represent overall estimates adjusted for publication bias.
Of the 43 hospitalization studies on age, higher associations were observed for older persons in 13. Five studies found mixed results, with the highest risk appearing in an age group other than the youngest or oldest (33, 34), or different results on age by time lag (35), outcome and cause (36), or sex (28). Four studies finding lower risks for older persons were for subpopulations (e.g., asthma emergency department visits for women during the warm season) (37–40).

We performed meta-analysis by age category (younger and older populations) for total, cardiovascular disease (CVD), and respiratory mortality and for CVD, respiratory, and asthma hospitalizations (Table 3, Figures 3–5, Web Figures 3–12). For older populations, we used the oldest age category available in each study. Younger populations were considered those most closely matching adults, excluding elder populations. Some studies compared estimates for older populations with persons of all ages, in which case we combined estimates for “all ages” with younger populations and performed sensitivity analysis excluding these studies.

A 10-ppb increase in daily 8-hour ozone concentration was associated with a 0.60% (95% CI: 0.40, 0.80) increase in total mortality for younger persons and a 1.27% (95% CI: 0.76,
1.78) increase for older persons. The ozone-mortality risk was 0.66% (95% CI: 0.12, 1.12) higher for older populations than for younger populations (Table 3). Sensitivity analysis removing studies comparing “all ages” with older populations was based on 22 pairs of estimates from 20 studies. Estimates were higher for older populations at 0.87% (95% CI: 0.61, 1.13) as compared with 0.50% (95% CI: 0.24, 0.76) for younger populations. Meta-analysis results were higher, but not statistically so, for older persons than for younger persons for CVD mortality, CVD hospital admissions, and respiratory hospital admissions (Web Figures 3–10). Results for respiratory mortality or asthma hospitalization were lower for older groups than for younger groups but were not statistically different (Table 3). Results from Egger’s test did not suggest heterogeneity (P > 0.05), except for ozone–respiratory mortality estimates. We found strong evidence of higher ozone associations for mortality in older populations.

### Modification of the association by race/ethnicity

Eight studies examined ozone sensitivity by race/ethnicity. Two found higher risks for minority populations, 1 with

<table>
<thead>
<tr>
<th>Table 3. Results From a Meta-Analysis of Associations Between Ozone Exposure and Mortality and Hospital Admission, by Age, 1988–2013</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>No. of Estimate Pairs</strong></td>
</tr>
<tr>
<td>Total mortality</td>
</tr>
<tr>
<td>Younger persons</td>
</tr>
<tr>
<td>Older persons</td>
</tr>
<tr>
<td>Older vs. younger</td>
</tr>
<tr>
<td>CVD mortality</td>
</tr>
<tr>
<td>Younger</td>
</tr>
<tr>
<td>Older</td>
</tr>
<tr>
<td>Older vs. younger</td>
</tr>
<tr>
<td>Respiratory mortality</td>
</tr>
<tr>
<td>Younger</td>
</tr>
<tr>
<td>Older</td>
</tr>
<tr>
<td>Older vs. younger</td>
</tr>
<tr>
<td>CVD hospitalization</td>
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<tr>
<td>Younger</td>
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<tr>
<td>Older</td>
</tr>
<tr>
<td>Older vs. younger</td>
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<tr>
<td>Asthma hospitalization</td>
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<tr>
<td>Younger</td>
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<tr>
<td>Older</td>
</tr>
<tr>
<td>Older vs. younger</td>
</tr>
</tbody>
</table>

Abbreviations: CI, confidence interval; CVD, cardiovascular disease.

a Percentage of variance in observed study results explained by heterogeneity.

b Percent increase in risk for a 10-ppb increase in 8-hour ozone concentration.

c Percent increase in ozone risk estimates for older persons compared with younger persons.
individual-level data and 1 with community-level data, and an additional study with individual-level data found higher risk for non-Caucasians for asthma and pneumonia emergency room admissions and higher associations for Caucasians for exacerbation of chronic obstructive pulmonary disease. We conclude that there is weak evidence of higher associations in minority populations. Investigation of ozone sensitivity by race/ethnicity was limited. All 4 hospitalization studies considered a single community (1 in Italy, 3 in the United States). All 4 mortality studies were US multicity studies. Race/ethnicity categorizations were simplistic, with most studies using 2 categories (e.g., black and other (22, 41, 42)). Only 2 studies used 3 or more race/ethnicity categories (43, 44).

**Modification of the association by SES indicators**

We considered ozone sensitivity with the following SES indicators: education, income, employment/occupation, and poverty. Overall, research suggested higher associations with lower SES. The most commonly studied SES indicator was education, with 10 studies evaluating this.
Some studies divided communities’ populations into binary categories (36, 49) or quartiles (44) by poverty status. One study found higher risks of ozone-related emergency department visits for CVD and dysrhythmia in high-poverty areas (36). The remaining studies found no evidence of ozone sensitivity by poverty level. We found weak evidence of higher ozone associations in high-poverty communities.

**Modification of the association by air conditioning**

Four studies examined sensitivity to ozone mortality by the presence of air conditioning (6, 22, 41, 42). Findings were more consistent for central air conditioning than for window air conditioning; there were higher associations with lower levels of central air conditioning in 3 studies (6, 41, 42). The 1 hospitalization study found higher risks with lower air conditioning (48). Analyses were limited by the use of community-level data, measurement of air conditioning prevalence rather than use, and a US focus, with the exception of 1 meta-analysis that included both US and Canadian cities (6). We found weak evidence of higher ozone associations with lower air conditioning prevalence.

**DISCUSSION**

Ozone sensitivities may relate to physiological differences. The structure of the respiratory system changes as we age, with decreased chest wall compliance, respiratory muscle strength, and vital capacity (50). Hormones and structural/morphological differences in the respiratory system may affect differences in risk between men and women (51).

Populations likely differ in terms of how estimated exposure relates to actual exposure. In the studies we identified, most researchers estimated exposure using outdoor levels, whereas actual exposure is affected by occupational exposure and indoor/outdoor activity patterns. A US survey found that 86.9% of participants’ time was spent indoors (52). Activity patterns can differ by sex, age, race/ethnicity, employment, and education (53). The indoor/outdoor ratio of ozone levels varies with air conditioning status (54) and housing structure, which can relate to SES. Further, studies use different approaches to estimate exposure to ozone, such as the nearest monitor value, the average of monitor values over a given area, interpolated values from several monitors, and modeled estimates, and they can take different approaches on whether to estimate ozone levels for days missing monitor values and the nature of such estimation.

Baseline health status, smoking, obesity, occupation, and other health-related factors that may affect vulnerability to ozone are more prevalent in some populations. For example, obesity can vary by race/ethnicity and sex (55, 56). Further, population characteristics are not independent (e.g., SES and race/ethnicity), complicating the ability to disentangle which factors in this complex system are most relevant for ozone sensitivity.

Previous meta-analyses of ozone associations did not focus on sensitive populations, although some examined risks by age. One earlier meta-analysis found a 0.57% (95% CI: -0.26, 1.41) higher total mortality—ozone estimate for older populations than for younger populations (8), a finding confirmed by our significant result of 0.66% (95% CI: 0.12, 1.12). Ozone—respiratory hospitalization estimates were

0.52% higher for older populations than for younger populations in our study, whereas a previous meta-analysis found a 0.15% difference (57); in both meta-analyses, risks were not statistically different by age group. An earlier meta-analysis found a 2.45% lower estimate for the association between ozone exposure and chronic obstructive pulmonary disease hospitalization among older persons than among persons of all ages; findings were not statistically significant and were based on 4 estimates for all ages (57). Other previous meta-analyses of ozone investigated mortality without exploration of sensitive populations (5, 58, 59).

In this analysis, 69% of studies provided results for multiple lag structures. The rationale for the selected lag structure varied and was not presented in all articles (Web Figure 2). Presentation of findings from a single lag structure may result in publication bias (8). Earlier work found that estimates of ozone associations from meta-analysis were consistently higher when based on a single lag than when results for multiple lags were reported (8). In future work, investigators could examine whether risks differ by lag using studies that presented results on multiple lag structures.

Although our findings ascertained sensitive populations, we identified gaps in the scientific literature. The lack of sufficient numbers of studies to perform meta-analysis necessitated qualitative assessment for sensitivity by race/ethnicity, SES indicators, and air conditioning. Categorization of cause of death or hospitalization by means of International Classification of Diseases codes is not perfectly consistent across studies (57). Many studies were designed to investigate issues other than population sensitivities. Studies developed specifically to explore ozone sensitivity may have different study designs. In particular, analysis of sensitivity by race/ethnicity used crude categories. Only 2 studies used 3 or more race/ethnicity categories (43, 44). No studies used individual-level data to investigate sensitivity by education or employment for hospitalizations, or to investigate income or poverty for either health outcome. All air conditioning studies employed prevalence of air conditioning rather than use. Some analysis was hindered further by a US focus. Because of these limitations and publication bias, the absence of evidence for ozone sensitivity should not be interpreted as evidence for the absence of sensitivity.

Many of the differences in estimates of ozone associations among groups were small in terms of the relative percent difference (Tables 2 and 3). However, these values present the relative increase in risk for a given group as compared with another group, and the actual health burden will relate to the baseline level of risk for ozone and the given health outcome for each group. Given the high levels of ozone pollution in many parts of the world and the large populations exposed, small relative differences in the risks for 2 populations could translate to large differences in the public health burden. Further, evidence on differences in sensitivity to ozone can inform policy-makers in their efforts to protect the public from air pollution, the medical community in their efforts to protect patients, and researchers who assess health impacts across different populations, as in the Global Burden of Disease Study.

Ozone levels have generally declined in many industrialized countries, yet over 123 million people in the United States live in areas with ozone concentrations exceeding the levels specified in health-based ozone regulations (60). Concentrations are increasing in much of the developing world with the expanding transportation networks, energy consumption, and industry that accompany urbanization (61). Worldwide, an estimated 470,000 respiratory deaths per year result from ozone exposure (62), and it is estimated that ozone-related deaths in the United Kingdom will increase 71% by 2050 (61). However, the true health burden of ozone is unknown without evidence regarding which populations are most sensitive. Estimates on health consequences from ozone for specific populations, such as in the Global Burden of Disease Study, for the present day (2, 62–64), or under a changing climate (61, 65, 66), often rely on a single or small number of concentration-response functions applied to all persons, although such functions may be based on populations with characteristics quite dissimilar from the population of interest and such characteristics change over time. Our findings indicate that health responses to ozone differ by age and employment/occupation, and possibly by sex, race/ethnicity, and other SES indicators. Evidence on which populations are most sensitive to ozone is needed to inform physicians as to which patients face higher risk, to aid decision-makers who formulate air-quality regulations, and to help sensitive individuals themselves, who may wish to modify their ozone exposure and understand their risk factors. Future studies should investigate ozone sensitivity for the population characteristics identified here, ideally with studies designed for this task.

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REFERENCES


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