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Publication Date
2011-03-25
Effects on Occupants of Enhanced Particle Filtration in a Non-Problem Office Environment: A Double-Blind Crossover Intervention Study

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June 15, 1998

Key Words: indoor air quality, sick building syndrome, intervention, filtration, ventilation

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Acknowledgments -- We thank building occupants and staff for their generous cooperation, and Mary Prince, Charles Mueller, and Caroline Portman for assistance with data collection. This work was supported by: the National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention; the Assistant Secretary of Energy Efficiency and Renewable Energy, Office of Building Technology, State and Community Systems of the U.S. Department of Energy under contract No. DE-AC03-76SF00098; and the Office of Research and Development, U.S. Environmental Protection Agency.
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Background

Workers in indoor environments often complain of symptoms, such as eye and nose irritation, headache, and fatigue, which improve away from work. Exposures causing such complaints, sometimes referred to as sick building syndrome, generally have not been identified. Evidence suggests these worker symptoms are related to chemical, microbiological, physical, and psychosocial exposures not well characterized by current methods (Mendell, 1993). Most research in this area has involved cross-sectional studies, which are limited in their abilities to show causal connections. Experimental studies have also been conducted which, by changing one factor at a time to isolate its effects, can demonstrate benefits of an environmental intervention even before exposures or mechanisms are understood (Mendell, 1993).

This study was prompted by evidence that particulate contaminants may be related to acute occupant symptoms and discomfort (Mendell, 1993; Leinster, 1990). The objective was to assess, with a double-blind, double crossover intervention design, whether improved removal of small airborne particles by enhanced central filtration would reduce symptoms and discomfort.

Methods

Methods are described in detail elsewhere (Fisk et al., 1998). To summarize: The experimental study spaces were two separate floors, with separate (variable air volume) ventilation systems equipped with conventional air filters, within a large office building in St. Louis. Occupants had sporadic complaints about comfort, but not about symptoms. Filtration was enhanced on alternate floors weekly, for four weeks in August 1996, by replacing conventional filters (estimated 3% efficiency at 0.3 microns) with highly efficient particle filters (estimated ≥95% efficiency at 0.3 microns) in ventilation systems. Building occupants and staff
were blinded to the schedule and nature of the interventions. Study staff blinded to the
intervention schedule performed all questionnaire handling and analyses.

An initial background questionnaire collected demographic and other personal
information. During the four-week crossover study, questionnaires distributed Thursday for
completion either Thursday or Friday afternoon assessed outcomes including: environmental
comfort, performance indicators, and symptom severity that day. The questionnaire also assessed
efficacy of the blind. Environmental measurements (described in Fisk et al. [1998]) included
temperature, humidity, carbon dioxide concentration, and ventilation rate (potentially requiring
adjustment in analyses) and concentrations of airborne particles, endotoxin, ergosterol, and beta-
1,3-glucans (potentially affected by the intervention and not used to adjust the analyses).

Intervention effectiveness was assessed by repeated measures ANOVA analysis models.
Because outcome variables were skewed, alternative statistical models were also considered:
Poisson regression, over-dispersed Poisson regression, and repeated-measures logistic regression.

Results

Eighty percent (392) of eligible participants returned the initial questionnaire (72% on
floor 2; 84% on floor 4). Weekly response rates averaged 63% over the four crossover weeks;
usable questionnaires averaged 58%. Of respondents, 60% were female; 29% were under 40
years old; 61% were White and 26% were African-American; 25% were managers or
supervisors; and 55% had never smoked. Workers on the two floors were demographically
similar.

Initial symptom prevalences were average for U.S. buildings [unpublished data, US EPA BASE
Study]. Carbon dioxide concentrations and ventilation rates were typical for US office buildings,
with ventilation rates nearly constant. Microbiological parameters measured were very low. Temperature and relative humidity were mostly within ASHRAE comfort limits (22.8°C-26.1°C). Enhanced filtration reduced concentration of the smallest particles measured, 0.3-0.5 microns, by about 95%, reduced 1 to 2 micron particles by about 50%, and reduced particles larger than 2 microns by about 10%. Enhanced filtration did not change ventilation rate, carbon dioxide concentrations, or supply air flow, and had no effect on endotoxin air concentrations. Air levels of ergosterol and beta-1,3-glucans were too low to characterize.

For all outcomes, positive scores indicate adverse effects, and negative scores indicate beneficial effects (Table). Temperature-adjusted ANOVA models for the effects of enhanced filtration on 13 outcomes showed that 10 outcomes improved slightly, although only two changes were statistically significant. With enhanced filtration, five of seven symptoms improved slightly and two worsened slightly, none statistically significantly (p-values=0.27 - 0.81). All performance indicators (confusion scale, fatigue scale, self-assessed productivity) showed small improvements (p-values=0.009, 0.13, and 0.11). Two environmental dissatisfaction variables, excess warmth and stiffness, improved slightly (p-values= 0.10, 0.006) but dryness worsened slightly (p-value =0.07). Increasing temperature, even within the comfort range, was strongly related to increases in most adverse outcomes; e.g., headache severity increased 1.7 units with a 1°C increase. Relative humidity was omitted from models because it was highly correlated with temperature. Results from alternative analysis models were similar to ANOVA results. Analyses showed that respondents were not aware of the specific times or nature of the intervention.
Table -- Results of Filtration Crossover Intervention, adjusted for mean temperature

<table>
<thead>
<tr>
<th>Outcomes (Range)</th>
<th>Outcome</th>
<th>Mean</th>
<th>Change (p-value) with Enhanced Filtration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Symptom severity</td>
<td>eyes- dry, itching, or irritated</td>
<td>6.2</td>
<td>0.07 (0.81)</td>
</tr>
<tr>
<td>$(0=none, to 25=very severe)$</td>
<td>nose- stuffy or congested</td>
<td>6.9</td>
<td>-0.29 (0.37)</td>
</tr>
<tr>
<td></td>
<td>throat- dry or irritated</td>
<td>5.7</td>
<td>-0.08 (0.80)</td>
</tr>
<tr>
<td></td>
<td>chest tightness</td>
<td>3.8</td>
<td>-0.14 (0.53)</td>
</tr>
<tr>
<td></td>
<td>headache</td>
<td>6.2</td>
<td>-0.26 (0.44)</td>
</tr>
<tr>
<td></td>
<td>fatigue or tiredness</td>
<td>8.7</td>
<td>-0.21 (0.51)</td>
</tr>
<tr>
<td></td>
<td>skin- dry, itchy, or irritated</td>
<td>5.3</td>
<td>0.25 (0.27)</td>
</tr>
<tr>
<td>Performance</td>
<td>mental confusion (5-item scale)</td>
<td>1.9</td>
<td>-0.08 (0.009)</td>
</tr>
<tr>
<td>indicators</td>
<td>scale</td>
<td>2.8</td>
<td>-0.06 (0.13)</td>
</tr>
<tr>
<td>$(1=not at all, to 5=extremely)$</td>
<td>fatigue (5-item scale)</td>
<td>3.6</td>
<td>-0.07 (0.07)</td>
</tr>
<tr>
<td></td>
<td>less productivity</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Environmental</td>
<td>too warm</td>
<td>2.4</td>
<td>-0.09 (0.07)</td>
</tr>
<tr>
<td>comfort</td>
<td>too stuffy</td>
<td>2.4</td>
<td>-0.16 (0.07)</td>
</tr>
<tr>
<td>$(1=not at all, to 5=extremely)$</td>
<td>too dry</td>
<td>2.1</td>
<td>0.09 (0.07)</td>
</tr>
</tbody>
</table>
Discussion

Although chance effects could be excluded only for the confusion scale and environmental stuffiness, enhanced particle filtration in this building was associated with small improvements in 10 of 13 worker outcome measures. Findings of only slight beneficial effects from removing small airborne particles in this building may not apply to buildings with higher symptom levels, lower ventilation rates, or significant microbiologic contamination.

Temperatures, not experimentally manipulated, were generally within the comfort range; however, the positive relation seen between temperatures and most adverse outcomes suggests substantial occupant benefits from thermal control even within the accepted comfort range.

Study strengths included the double-blind, crossover intervention design and analyses of changes within subjects. Although uncontrolled factors varying over time may have biased findings, among measured factors only temperature was related to outcomes, and adjustment for temperature was made in analyses. Participants were not randomly allocated to experimental groups, but the crossover design reduced potential resulting bias. If effects of enhanced filtration began after or lasted more than a week, findings could have been distorted. Health outcomes potentially related to particles other than acute symptoms were not assessed. Studying a building without excess symptoms or contaminants reduced the chance of finding filtration benefits; however, finding small effects in such a building, if replicated, will be more generalizable than results in a heavily contaminated building. Similar blinded, controlled, crossover studies can assess whether enhanced filtration or other reversible interventions produce larger benefits in buildings with suspected particulate contamination or higher symptom levels.
References


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