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Use of the Adaptive LASSO Method to Identify PM$_{2.5}$ Components Associated with Blood Pressure in Elderly Men: The Veterans Affairs Normative Aging Study

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**Background:** PM$_{2.5}$ (particulate matter ≤ 2.5 μm) has been associated with adverse cardiovascular outcomes, but it is unclear whether specific PM$_{2.5}$ components, particularly metals, may be responsible for cardiovascular effects.

**Objectives:** We aimed to determine which PM$_{2.5}$ components are associated with blood pressure in a longitudinal cohort.

**Methods:** We fit linear mixed-effects models with the adaptive LASSO penalty to longitudinal data from 718 elderly men in the Veterans Affairs Normative Aging Study, 1999–2010. We controlled for PM$_{2.5}$ mass, age, body mass index, use of antihypertensive medication (ACE inhibitors, non-ophthalmic beta blockers, calcium channel blockers, diuretics, and angiotensin receptor antagonists), smoking status, alcohol intake, years of education, temperature, and season as fixed effects in the models, and additionally applied the adaptive LASSO method to select PM$_{2.5}$ components associated with blood pressure. Final models were identified by the Bayesian Information Criterion (BIC).

**Results:** For systolic blood pressure (SBP), nickel (Ni) and sodium (Na) were selected by the adaptive LASSO, whereas only Na was selected for diastolic blood pressure (DBP). An interquartile range increase (2.5 ng/m$^3$) in 7-day moving-average Ni was associated with 2.48 mmHg (95% CI: 1.45, 3.50 mmHg) increase in SBP and 2.22 mmHg (95% CI: 1.69, 2.75 mmHg) increase in DBP, respectively. Associations were comparable when the analysis was restricted to study visits with PM$_{2.5}$ below the 75th percentile of the distribution (12 μg/m$^3$).

**Conclusions:** Our study suggested that exposure to ambient Ni was associated with increased blood pressure independent of PM$_{2.5}$ mass in our study population of elderly men. Further research is needed to confirm our findings, assess generalizability to other populations, and identify potential mechanisms for Ni effects.

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PM$_{2.5}$ components and blood pressure

718 participants with 1,567 observations had examinations between March 1999 and October 2010. Of the 718 participants, 235 (33%) had one visit, 195 (27%) had two visits, and 288 (40%) had three or more visits.

**Blood pressure measurements.** During a clinical visit, a physician uses a standard mercury sphygmomanometer with a 14-cm cuff to measure blood pressure for the subject while he is sitting, including systolic blood pressure (SBP) and fifth-phase diastolic blood pressure (DBP) in each arm to the nearest 2 mmHg. We used the means of the left and right arm measurements as a subject’s SBP and DBP.

**Environmental data.** Daily ambient PM$_{2.5}$ and its components were measured at the stationary ambient monitoring site at the Harvard University Countway Library (Kang et al. 2010), using the tapered element oscillating microbalance (TEOM 1400a; Rupprecht & Patashnick Co.) and the energy dispersive X-ray fluorescence spectrometer (Epsilon 5; PANalytical), respectively. The monitoring site is 1 km from the clinical examination site.

We obtained daily temperature data from Boston Logan airport weather station.

**Statistical analysis.** We used 7-day moving-average concentrations for PM$_{2.5}$ and the 11 components—K, S, Se, Al, Si, Fe, Ni, V, Cu, Zn, and Na—because previous studies have suggested that PM averaging over that time period is strongly associated with blood pressure (Mordukhovich et al. 2009; Wilker et al. 2010; Zanobetti et al. 2004). We focused on these components because their concentration levels are mostly above the method detection limits and they are representative of different PM sources (Hopke et al. 2006).

In the analysis, we controlled for continuous variables age, body mass index (BMI; computed as weight (in kilograms) divided by height (in square meters)), years of education, linear and quadratic terms of mean temperature of visit day, and categorical variables use of each class of antihypertensive medication (ACE inhibitors, non-ophthalmic beta blockers, calcium channel blockers, diuretics, and angiotensin receptor antagonists), smoking status (three categories: never, former, current smoker), alcohol intake (whether the participant takes two or more drinks per day; yes or no), and season (four categories: defined as spring; March–May; summer: June–August; fall: September–November; winter: December–February) regardless of statistical significance because these variables have been shown to predict cardiovascular health (Mordukhovich et al. 2009; Schwartz et al. 2012). In addition, we adjusted for potential confounding of associations with PM$_{2.5}$ components by PM$_{1.5}$ mass (Mostofsky et al. 2012). All variables were measured at each visit. We forced these covariates to be included in the models and estimated their fixed effects with no penalization.

Selecting important predictors from a large list of correlated predictors is difficult, and most methods are empirical. Approaches such as stepwise methods ignore stochastic errors inherited in the stages of variable selection (Fan and Li 2001) and can yield falsely narrow confidence intervals (Harrell 2001). To improve on this, we applied the adaptive LASSO (least absolute shrinkage and selection operator) method to select important component(s) that may be associated with blood pressure from those 11 PM$_{2.5}$ components. Briefly, the LASSO is a regression shrinkage and selection approach that applies an $l_1$ penalty to the component regression coefficients. This penalty essentially minimizes the sum of squared errors subject to the sum of the absolute values of the coefficients being less than a given value (Tibshirani 1996). The adaptive LASSO is a later version of the LASSO, which uses weights for penalizing different coefficients in the $l_1$ penalty and enjoys the oracle properties, which means, given that the true model depends only on a subset of the predictors, this selection procedure is able to identify the right subset model and satisfies asymptotic normality (Fan and Li 2001; Zou 2006). Because subjects had repeated measurements, we fit linear mixed-effects models with random subject-specific intercepts to capture the correlation among different measurements within the same subject, as follows:

$$Y_i = X_i \alpha + Z_i \beta + \mu_i + \epsilon_i$$

where, $Y_i$ is the blood pressure level (SBP or DBP) of subject $i$, $X_i = (X_{i1}, \ldots, X_{i27})^T$ is a vector of PM$_{2.5}$ mass and other covariates, $Z_i = (Z_{i1}, \ldots, Z_{i11})^T$ is a vector of PM$_{2.5}$ components, $\mu_i$ is the random intercept. Hence, $\alpha$ indicates the fixed effects of PM$_{2.5}$ mass and other covariates $X_i$ and $\beta$ is the penalized effects of PM$_{2.5}$ components $Z_i$ that are given by the adaptive LASSO.

First, we used the ordinary linear mixed-effects (LME) model to obtain non-zero coefficients ($\beta_{\text{non}}$) for each component, and computed the adaptive weight as its inverse ($w = 1/|\beta_{\text{non}}|$). Heuristically, this allows us to give less weight in the penalty to variables whose standardized regression coefficients are large, because they are more likely to be predictors. When using the adaptive LASSO, we assign a non-negative penalty parameter, $\lambda$, to determine how strongly we penalize, or restrict, the magnitude of the PM$_{2.5}$ components regression coefficients. When $\lambda$ is equal to 0, there is no shrinkage, and the model is just the ordinary mixed-effects regression of the fixed covariates and all components; when it is large enough, there is maximum shrinkage, yielding a model that includes fixed covariates only (all component coefficients equal to 0); when $\lambda$ takes some value in between, some coefficients are 0, and the model is a penalized model. Components with non-zero coefficients are “selected” by the adaptive LASSO. In this way, the method chooses PM$_{2.5}$ components that may be associated with the outcomes. We ran the models across that range of $\lambda$s—from no shrinkage to maximum shrinkage—and chose the $\lambda$ having the smallest Bayesian Information Criterion (BIC) (Schwarz 1978). Last, we used the mixed-effects model with fixed covariates and selected components only, to obtain the estimated effects and corresponding 95% confidence intervals (CIs).

In a sensitivity analysis, we omitted study visits with PM$_{2.5}$ below the 75th percentile of the distribution (12 $\mu g/m^3$).

Data cleaning was performed with SAS 9.3 (SAS Institute Inc.), and data analysis was performed with R 3.1.2 (R Core Team 2015).

**Results**

Table 1 summarizes the characteristics of study population. Subjects in this study were elderly men, with a mean (± SD) age of 73 ± 7 years at the first visit. Average SBP and DBP

<table>
<thead>
<tr>
<th>Variable</th>
<th>First visit (n = 718)</th>
<th>All visits (n = 1,567)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SBP (mmHg)</td>
<td>131.6 ± 16.7</td>
<td>128.1 ± 17.6</td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>75.9 ± 9.9</td>
<td>71.9 ± 10.3</td>
</tr>
<tr>
<td>Age (years)</td>
<td>72.8 ± 6.8</td>
<td>74.7 ± 6.8</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>28.2 ± 4.0</td>
<td>28.0 ± 4.1</td>
</tr>
<tr>
<td>Education (years)</td>
<td>14.6 ± 2.8</td>
<td>14.6 ± 2.8</td>
</tr>
<tr>
<td>n(%)</td>
<td>197 (27)</td>
<td>540 (34)</td>
</tr>
<tr>
<td>Use of ACE inhibitors</td>
<td>213 (30)</td>
<td>555 (35)</td>
</tr>
<tr>
<td>Use of non-ophthalmic beta blockers</td>
<td>104 (14)</td>
<td>265 (17)</td>
</tr>
<tr>
<td>Use of calcium channel blockers</td>
<td>150 (21)</td>
<td>381 (24)</td>
</tr>
<tr>
<td>Use of diuretics</td>
<td>36 (5)</td>
<td>124 (8)</td>
</tr>
<tr>
<td>Use of angiotensin receptor antagonists</td>
<td>28 (4)</td>
<td>47 (3)</td>
</tr>
<tr>
<td>Former smokers</td>
<td>488 (68)</td>
<td>1,049 (67)</td>
</tr>
<tr>
<td>Two or more drinks per day</td>
<td>143 (20)</td>
<td>299 (19)</td>
</tr>
</tbody>
</table>
DBP at the first visit were 132 ± 17 mmHg and 76 ± 10 mmHg, respectively.

PM$_{2.5}$ and component concentrations are shown in Table 2. 7-day moving-average PM$_{2.5}$ across all study visits had a mean of 10 ± 3.7 μg/m$^3$, with an interquartile range (IQR) of 4.3 μg/m$^3$. S accounted for the largest proportion of the total PM$_{2.5}$ concentration (10.4%), followed by Na (1.9%). The average concentration of Ni was 3.1 ± 2.5 ng/m$^3$, and it only accounted for 0.03% of the mass concentration.

Table 2. Mean PM$_{2.5}$ mass and component concentrations across all study visits.

<table>
<thead>
<tr>
<th>Pollutant</th>
<th>Mean ± SD</th>
<th>IQR</th>
<th>Proportion of PM$_{2.5}$ (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PM$_{2.5}$ (μg/m$^3$)</td>
<td>10.0 ± 3.7</td>
<td>4.3</td>
<td></td>
</tr>
<tr>
<td>Component (ng/m$^3$)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fe</td>
<td>68.1 ± 24.2</td>
<td>21.5</td>
<td>0.7</td>
</tr>
<tr>
<td>K</td>
<td>39.2 ± 24.6</td>
<td>16.9</td>
<td>0.4</td>
</tr>
<tr>
<td>S</td>
<td>1039.1 ± 513.2</td>
<td>554.1</td>
<td>10.4</td>
</tr>
<tr>
<td>Al</td>
<td>51.8 ± 27.8</td>
<td>21.1</td>
<td>0.5</td>
</tr>
<tr>
<td>Si</td>
<td>76.7 ± 51.1</td>
<td>38.4</td>
<td>0.8</td>
</tr>
<tr>
<td>Ni</td>
<td>3.1 ± 2.5</td>
<td>2.5</td>
<td>0.03</td>
</tr>
<tr>
<td>V</td>
<td>3.5 ± 2.3</td>
<td>2.6</td>
<td>0.04</td>
</tr>
<tr>
<td>Cu</td>
<td>3.5 ± 1.2</td>
<td>1.5</td>
<td>0.04</td>
</tr>
<tr>
<td>Zn</td>
<td>11.4 ± 6.0</td>
<td>5.8</td>
<td>0.1</td>
</tr>
<tr>
<td>Si</td>
<td>0.2 ± 0.3</td>
<td>0.3</td>
<td>0.002</td>
</tr>
<tr>
<td>Na</td>
<td>190.7 ± 72.4</td>
<td>92.8</td>
<td>1.9</td>
</tr>
</tbody>
</table>

Figure 1 shows the relationship between BIC, a criterion for model selection and $\lambda$, the adaptive LASSO penalty parameter. For SBP models, the model with the smallest BIC had $\lambda = 4$ and Ni and Na as the only two among the 11 PM$_{2.5}$ components (i.e., K, S, Se, Al, Si, Fe, Ni, V, Cu, Zn, Na) with non-zero coefficients, whereas all component coefficients were zero when $\lambda = 9$. For DBP models, the model with the smallest BIC had $\lambda = 13$ and Ni as the only component with a non-zero coefficient, whereas all component coefficients were zero when $\lambda = 30$.

In models fitted using only the selected components, we found that an IQR increase (2.5 ng/m$^3$) in 7-day moving-average Ni was associated with a 2.48-mmHg (95% CI: 1.45, 3.50 mmHg) increase in SBP and a 2.22-mmHg (95% CI: 1.69, 2.75 mmHg) increase in DBP, respectively. To compare with other studies, we also estimated the effects of PM$_{2.5}$ mass: Every 10-μg/m$^3$ increase in 7-day moving-average PM$_{2.5}$ was associated with a 1.36-mmHg (95% CI: –1.67, 4.39 mmHg) increase in SBP and a 0.61-mmHg (95% CI: –0.85, 2.07 mmHg) increase in DBP, respectively.

LASSO coefficient paths for SBP and DBP are shown in Figure 2. Each component coefficient is expressed as the change in mean SBP or DBP per 1-μg/m$^3$ increase in the 7-day moving-average concentration of the PM$_{2.5}$ component. Each curve indicates the rate at which the component coefficient shrinks toward zero as $\lambda$ increases. When $\lambda = 0$, all components have non-zero coefficients.

Table 3 shows the comparison of results from the main analysis and the sensitivity analysis that was restricted to data from study visits with PM$_{2.5}$ concentrations below the 75th percentile of the distribution (12 μg/m$^3$). We found that the estimated coefficients of Ni for both SBP and DBP in the sensitivity analysis were comparable with
those in the main analysis, and their statistical significance remained. That is, Ni was associated with SBP and DBP even when overall PM$_{2.5}$ concentrations were restricted to < 12 μg/m$^3$.

Ni in ambient air is considered a marker of oil combustion; other sources of Ni include coal combustion, nickel metal refining, sewage sludge incineration, and manufacturing facilities [U.S. Environmental Protection Agency (EPA) 2000]. A number of toxicological studies examined the effects of ambient Ni on cardiovascular health. In a mouse model of atherosclerosis, mice had acute changes in heart rate and heart rate variability when exposed to concentrated fine PM (average concentration of Ni was 43 ng/m$^3$, and there were Ni peaks at ~175 ng/m$^3$) (Lippmann et al. 2006). Another animal study showed that Ni inhalation caused a decrease of 75 bpm in maximal heart rate at the concentration of 1.3 mg/m$^3$ and a decrease of 100 bpm at 2.1 mg/m$^3$ in rats (Campion et al. 2001). Moreover, Ni was reported to induce increases in pulmonary protein leakage and perivascular and peribronchial inflammation in both normotensive and spontaneously hypertensive rats that were intratracheally instilled with 1.5 μmol/kg of NiSO$_4$·6H$_2$O in saline (Kodavanti et al. 2001). A similar study found alterations in heart rate variability (HRV) related to PM exposure were Ni-dependent in spontaneously hypertensive rats after adjustment for HRV responses in control rats (Chuang et al. 2013).

Several epidemiological studies have provided evidence of cardiovascular effects of Ni. A national study conducted in 106 U.S. counties reported that associations between PM$_{2.5}$ concentrations and cardiovascular and respiratory hospitalizations were stronger when Ni was high (Bell et al. 2009). Zanobetti et al. (2009) examined associations of PM$_{2.5}$ with emergency hospital admissions in 26 U.S. communities and found that Ni significantly modified the association between PM$_{2.5}$ mass and hospital admissions for cardiac diseases and myocardial infarctions. A recent study found a significant association between ischemic heart disease mortality and Ni based on data from the American Cancer Society (Lippmann et al. 2013). On the other hand, Zhou et al. (2011) failed to find cumulative effects from lag 0 to lag 2 of Ni in Detroit, Michigan, or Seattle, Washington. A more recent nationwide study that included 75 U.S. cities did not observe any effect modification of Ni in the PM$_{2.5}$-mortality association (Dai et al. 2014).

Discussion

In this study, we used the adaptive LASSO shrinkage method to choose PM$_{2.5}$ components that might be related to blood pressure in a cohort of elderly men. We found that 7-day moving-average concentrations of Ni and Na were associated with SBP, and 7-day moving-average Ni concentration was also associated with DBP. This association persisted when restricted to data from study visits with PM$_{2.5}$ concentrations < 12 μg/m$^3$.

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There are several possible reasons for the differences in these epidemiological studies. First, Ni concentrations are usually lower than the method detection limits, which makes it difficult to determine whether associations are present (Burnett et al. 2000). New York counties had particularly high levels of Ni (a mean of 19.0 ng/m$^3$ Ni in New York fine PM vs. a mean of 1.9 ng/m$^3$ Ni in national fine PM) due to combustion of residual oil-fired power plants and ocean-going ships (Lippmann et al. 2006). In a reanalysis of the National Morbidity, Mortality, and Air Pollution Study (NMMAPS) data, Dominici et al. (2007) found evidence of effect modification by Ni, which was consistent with the results of Lippmann et al. (2006); however, the effect modification of Ni on the PM–mortality association was much weaker and no longer statistically significant when New York counties were excluded from the analysis. In the two studies that did not find significant associations, Ni had a relatively low level. For example, the national mean concentration of Ni was 2.5 ng/m$^3$ in the study by Dai et al. (2014). Given the substantial differences in Ni concentrations, it is conceivable that studies conducted in other places or nationally may not be able to observe the same health effects of Ni as the New York studies did. In our study, Ni had an average concentration of 3.1 ng/m$^3$, which was higher than the national mean but still much lower than New York levels (Dominici et al. 2007; Lippmann et al. 2006). Furthermore, it is possible that Ni interacts with other PM components to pose an increased risk to health. Campen et al. (2001) reported evidence of a synergistic interaction between Ni and V, both of which are markers of PM from oil combustion. Hence, the heterogeneous composition of PM in different locations might lead to different estimated effects of Ni.

Na also was selected, in addition to Ni, when the adaptive LASSO method was applied to identify PM$_{2.5}$ components associated with SBP. There is limited literature on the effects of ambient Na on cardiovascular health. Zanobetti et al. (2009) documented that Na$^+$ modified the relationship between PM$_{2.5}$ and emergency hospital admissions for cardiac diseases.

In the study, the maximum level of 7-day moving-average PM$_{2.5}$ concentration was 34.3 μg/m$^3$, whereas daily PM$_{2.5}$ peaked at 44.8 μg/m$^3$ with a 99th percentile of 34 μg/m$^3$. Hence, we identified associations in a study population that was usually exposed to PM$_{2.5}$ concentrations below the current U.S. EPA daily ambient standard of 35 μg/m$^3$ (U.S. EPA 2012). Associations with Ni were similar when we excluded observations with 7-day moving-average PM concentration > 12 μg/m$^3$. Our findings may suggest stricter air quality standards.

To date, many studies have investigated the biological mechanisms of the adverse effects of inhalation exposures to PM on cardiovascular diseases. Brook et al. (2010) summarized three potential pathways: a) inducing pulmonary oxidative stress and inflammation via the release of proinflammatory mediators or vasoactive molecules; b) interacting with lung receptors or nerves to perturb systemic autonomic nervous system balance or heart rhythm; or c) PM or PM components being transmitted into the systemic circulation. Metals are typical PM components. It has been documented that metals can enhance lung inflammation and injury (Ghio and Devlin 2001; Schaumann et al. 2004), which may be attributed to the metal-catalyzed oxygen stress via non-nitric oxide pathways (Dye et al. 1997). Nevertheless, mechanisms of cardiovascular effects of Ni have not been fully established. Previous studies have shown that metals in particles (e.g., Ni, V) could induce the activation of transcription factor NF-kB (nuclear factor kB: a family of proteins that regulates DNA transcription in cellular responses such as immune, inflammatory response, and apoptosis), cell apoptosis, and cell cycle regulation (Chen and Shi 2002; Goebeler et al. 1995; Quay et al. 1998). Although the clinical relevance is unclear, our finding that an IQR increase in Ni was associated with a 2.48/2.22-mmHg increase in blood pressure may imply elevated risks of cardiovascular outcomes induced by Ni.

The major strengths in the study are as follows: First, we used a novel approach, the adaptive LASSO, to investigate the relationship between PM$_{2.5}$ components and health outcomes. This method has advantages over

| Table 3. Comparison of estimated coefficients of Ni in the main analysis and in the sensitivity analysis where study visits with 7-day moving-average PM$_{2.5}$ > 12 μg/m$^3$ were excluded. |
|---|---|---|---|
| | SBP | | DBP |
| | Coefficient | p-Value | Coefficient | p-Value |
| Main analysis (n = 1,567) | 0.988 | < 0.001 | 0.888 | < 0.001 |
| Sensitivity analysis (n = 1,201) | 1.149 | < 0.001 | 1.104 | < 0.001 |
conventional approaches. Typically, researchers examined effects of components by including all components in models or by using conventional selection procedures, such as stepwise selection. Linear regression with all components included may fail to detect any association because the collinearity among components reduces power, and conventional selection methods make no guarantee to select the right variables asymptotically. Second, to our knowledge, this is the first longitudinal cohort study to examine the effects of PM-related metals on blood pressure. The study population was geographically stable, well described, and followed up since enrollment in 1963. Third, we had daily concentrations of PM metals for >10 years. In previous studies, especially large national/multi-city studies, researchers usually followed up since enrollment in 1963. Third, because the collinearity among components included may fail to detect any association with mortality in the United States: a multicity reanalysis of the NMMAPS data. Environ Health Perspect 115:1701–1702; doi:10.1289/ehp.10737.


Dai et al. 2014; Krall et al. 2013; Zanobetti et al. 2009) and hence had to face the challenge in lack of data.

On the other hand, there are several limitations in the study. Due to the use of stationary measures of PM2.5 components, we were unable to capture the personal exposures of our subjects. Another limitation of our study is the potential measurement errors in blood pressure, because blood pressure was measured only once at each study visit. Last, because the study population was limited to elderly men, most of whom were Caucasian, our findings cannot be directly generalized to women, younger men, or more diverse populations of elderly men. Subjects voluntarily continue to participate in the ongoing NAS study, so there may be volunteer bias if healthier people are more likely to participate. Also, there would be survivor bias if people who stay in the study are healthier than other people.

REFERENCES


