Association between blood metals mixtures concentrations and cognitive performance, and effect modification by diet in older US adults

Nasser Laouali1,*, Tarik Benmarhnia2, Bruce P. Lanphear1,6, Jennifer Weuve1, Michael Mascari3, Marie-Christine Boutron-Ruault4, Youssef Oulhote5

Background: Chronic exposure to heavy metals has been associated with adverse neurological outcomes in older adults. Inflammatory processes are suspected as an underlying pathway by which metals exert their neurotoxicity. In parallel, a diet rich in antioxidant and anti-inflammatory components may protect against chronic inflammation.

Objectives: We examined the associations of blood concentrations of lead, cadmium, and manganese as a mixture with cognitive performance in older US adults and potential modification of these associations by diet as measured by the Healthy Eating Index 2015 (HEI-2015) and the Adaptable Dietary Inflammatory Index (ADII).

Methods: We used data on 1,777 adults ≥60 years old from the US National Health and Nutrition Examination Survey (NHANES; 2011–2014). We derived the ADII and the HEI-2015 from two nonconsecutive 24-hour diet recalls. Cognitive performance was measured by the Consortium to Establish a Registry for Alzheimer’s Disease (CERAD) Word Learning subtest, the animal fluency test, and the Digit Symbol Substitution Test (DSST). We also conducted a composite z-score reflecting overall cognitive performance. We used quantile g-computation to evaluate the joint associations of a mixture of metals with cognitive performance test scores. We also evaluated effect modification by sex and diet quality indices using Cochran Q tests.

Results: The median (interquartile range) of blood metals were 0.38 μg/L (0.35), 14.70 μg/L (11.70), and 8.74 μg/L (4.06) for cadmium, lead, and manganese, respectively. Increasing blood concentrations of all metals by one quartile was associated with a decrease in overall cognitive performance (−0.04; 95% confidence interval [CI] = −0.09, 0.02), CERAD (−0.04; 95% CI = −0.12, 0.03), animal fluency (−0.02; 95% CI = −0.11, 0.06), and DSST (−0.05; 95% CI = −0.11, 0.02) test scores. These associations were more pronounced in adults with high pro-inflammatory or low-diet quality and null or positive though imprecise associations in participants with a high anti-inflammatory. These associations also varied by sex with inverse associations in men and positive associations in women.

Conclusions: Our findings suggest that adherence to an antioxidant and anti-inflammatory diet may prevent blood metals adverse cognitive effects among older adults. If confirmed, strategies based on diet could provide a potential complementary and efficient approach to counteract effects of environmental pollutants.

Keywords: Adapted Dietary Inflammatory Index; Cadmium; Epidemiology; Healthy Eating Index; Lead; Manganese; National Health and Nutrition Examination Survey

Introduction

Aging is a naturally occurring physiological process that is characterized by both physical and cognitive decline. Cognitive decline is a process characterized by a gradual deterioration from mid-adulthood onwards in cognitive functions including processing speed, reasoning, and memory and executive functions.1 In addition to advanced age, other factors may contribute to declines in cognitive performance. Gurland et al.2 estimated that 25% of inter-individual variability is due to genetic factors, leaving a large unexplained proportion potentially due to modifiable risk factors, including gene-environment interactions.3 With the increases in life expectancy worldwide,

What this study adds

Chronic exposure to heavy metals has been associated with adverse neurological outcomes in older adults and inflammatory processes are suspected as an underlying pathway. In parallel, a diet rich in antioxidant and anti-inflammatory components may protect against chronic inflammation. However, to the best of our knowledge, no study has evaluated the potential modification of the associations between metals and adverse neurological outcomes by dietary scores. In this cross-sectional study, we found that an antioxidant and anti-inflammatory diet may mitigate adverse cognitive effects of metals among older adults. This result provides a potential complementary and efficient approach to counteract effects of environmental pollutants.
there is a strong interest in identifying modifiable factors that accelerate cognitive decline. As such, recent research has focused on understanding environmental factors, including exposure to neurotoxic metals and lifestyle factors, such as dietary quality.

Heavy metals, such as lead (Pb), manganese (Mn), and cadmium (Cd), are naturally present in our environment. Overall, cross-sectional and longitudinal studies suggest that cumulative or long-term exposure to these metals, measured where they are stored, was associated with cognitive decline and impaired cognitive function in older adults. However, the evidence is less consistent when it comes to measures of current exposure, such as measurements in blood. Although the underlying mechanisms are not completely understood, metals may exert their neurotoxicity via inflammatory processes in the brain. Several heavy metals are involved in the oxidant/antioxidant mechanisms through the production of free radicals, thus leading to oxidative stress, impairment of endothelial cell function, and the initiation of inflammatory processes. On the other hand, inflammation and oxidative stress modifies the effect of cadmium exposure on markers of systemic inflammation and oxidative stress.

Although heavy metals and dietary intake are both involved in the regulation of inflammation and cognitive performance, their contribution to cognitive performance in older adults is poorly understood. Recent findings suggest a potential protective effect of folic acid, an antioxidant component of the diet; against effects of phthalate metabolites, pesticides and air pollutants on autistic traits and intellectual abilities in children, or on the risk of preterm birth. However, to our knowledge, no study evaluated similar patterns for other environmental contaminants on cognitive performance among the elderly. In this investigation, we hypothesized that a high-diet quality or adherence to an anti-inflammatory dietary pattern may attenuate inflammation and subsequently reduce the metals-induced neurotoxicity. Therefore, we sought to examine the associations of a mixture of metals with the cognitive performance in US older adults and the potential effect modification by overall diet quality as measured by the Healthy Eating Index 2015 (HEI-2015) and the Adapted Dietary Inflammatory Index (ADII).

Methods

Design and participants

We used data from the US National Health and Nutrition Examination Survey (NHANES). The NHANES is an ongoing survey conducted by the Centers for Disease Control and Prevention (CDC) that uses a representative sample of noninstitutionalized civilians in the United States; selected by a complex, multistage, stratified, clustered probability design. Trained staff collected information on participants by interviews and physical examinations. The interview includes background information such as sociodemographic, dietary, and health-related questions. The examination component consists of medical and physiological measurements, and laboratory tests. The National Center for Health Statistics Ethics Review Board approved all NHANES protocols, and all survey participants completed a consent form. The detailed protocol on NHANES methodology and data collection is available on https://www.cdc.gov/nchs/nhanes/index.htm. For this study, we included participants with data on blood metals concentrations (n = 2,068). We excluded participants with missing data on diet (n = 291). The final study population included 1,777 adults ≥60 years old with data on dietary intake who participated in the cognitive function test of the 2011–2012 and 2013–2014 cycles.

Measurements of trace metals in blood

The methodological details on the laboratory analyses are described in detail on the NHANES website. Briefly, whole blood specimens were frozen (~30 °C), stored, and shipped for analysis to the Division of Laboratory Sciences, National Center for Environmental Health of the CDC. Cadmium (µg/L), lead (µg/dL), and manganese (µg/L) concentrations were measured in whole blood using inductively coupled plasma mass spectrometry. Coupling radio frequency power into a flowing argon stream seeded with electrons creates the plasma. Predominant species in the plasma are positive argon ions and electrons. Diluted whole blood samples are converted into an aerosol using a nebulizer inserted within a spray chamber. A portion of the aerosol is transported through the spray chamber and then through the central channel of the plasma, where it experiences temperatures of 6,000–8,000 °K. This thermal energy ionizes and fragments the sample. The ions, along with the argon, enter the mass spectrometer through an interface that separates the inductively coupled plasma, operating at atmospheric pressure (approximately 760 torr), from the mass spectrometer, operating at approximately 10⁻⁴ torr. The lower limits of detection (LOD) in 2011–2012 and 2013–2014 cycles were 0.16 µg/L and 0.10 µg/L for cadmium, 0.25 µg/dL and 0.07 µg/dL for lead, and 1.06 µg/L and 0.99 µg/L for manganese. Measures below LOD were set by NHANES as LOD/√2. In 2011–2012 and 2013–2014 cycles, LOD constituted 31% and 29% for cadmium, 1% and 0% for lead, and 0% in both cycles for manganese.

Evaluation of cognitive performance

Cognitive performance was measured by the Consortium to Establish a Registry for Alzheimer’s Disease (CERAD) Word List Learning and Recall Test, the Animal Fluency test, and the Digit Symbol Substitution Test (DSST). The CERAD Word List Learning and Recall Tests were designed to assess immediate and delayed learning ability for new verbal information. For the learning trials, participants were instructed to read aloud 10 unrelated words, one at a time, as they were presented. Immediately following the presentation of the words, participants recalled as many words as possible. In each of the three learning trials, the order of the 10 words is changed. The delayed word recall occurred after the other two cognitive exercises (Animal Fluency and DSST) were completed (approximately 8–10 minutes from the start of the word learning trials). The maximum score possible on each trial is 10. The Animal Fluency test examines categorical verbal fluency, a component of executive function in which participants were asked to name as many animals as possible in 1 minute. The DSST, a performance module from the Wechsler Adult Intelligence Scale, was used to assess processing speed, sustained attention and working memory. The exercise was conducted using a paper form that has a key at the top containing nine numbers paired with symbols. Participants have 2 minutes to copy the corresponding symbols in the 133 boxes that adjoin the numbers. The score is the total number of correct matches. Higher scores represent better cognitive function for all tests. In addition, we calculated a composite z-score reflecting the overall cognitive performance as suggested by Li et al. This composite cognitive z-score was the average of the standardized scores of the CERAD Word List Learning and Recall Test, the DSST, and the Animal Fluency test. All cognitive test scores were standardized to keep the same scale as the composite score. Higher scores represent better cognitive function for all tests.
Dietary data and diet quality indices
Dietary intake was collected from the two NHANES cycles using 24-hour dietary recalls. The first 24-hour recall interview was conducted at a Mobile Examination Center using a standard set of measuring guides to assist in estimating portion sizes. The second dietary recall, which was collected by telephone, was scheduled 3 to 10 days later. We derived the ADII and the HEI-2015 from the mean daily intakes of foods/beverages, energy, and nutrients of these two nonconsecutive 24-hour dietary recalls.

The Adapted Dietary Inflammatory Index (ADII) was constructed to measure the inflammatory potential of the diet. Briefly, the ADII index proposed by Van Woudenbergh et al. was used in combination with the updated dietary components inflammatory weights designed by Shivappa et al. Instead of the weights proposed by Cavicchia et al., these updated inflammatory weights, based on 3 additional years of published data (2008–2010, inclusive), resulted in a doubling of the total number of articles scored. The ADII was based on a nutritional rationale: first, the inflammatory weights of dietary components are multiplied by the standardized energy-adjusted intake, which acts to reduce between-person variation; second, the intakes of all components are standardized; third, the ADII calculation did not include alcoholic beverages such as beer, wine, and liquor, total fat, and energy, to avoid overestimation of the inflammatory effects of ethanol, fat, and energy. A total of 24 of the 35 possible dietary components were used for the ADII calculation (carbohydrates, proteins, alcohol, fibers, cholesterol, saturated fatty acids, monounsaturated fatty acids, omega 3, omega 6, niacin, thiamin, riboflavin, vitamin B6, vitamin B12, iron, magnesium, zinc, vitamin A, vitamin C, vitamin D, vitamin E, folate, beta carotene, and caffeine). A positive ADII score indicates an anti-inflammatory diet and negative values correspond to a pro-inflammatory diet.

The Healthy Eating Index 2015 (HEI-2015) was developed to measure overall diet quality. Briefly, HEI-2015 is a composite measure of conformance to the 2015–2020 Dietary Guidelines for Americans and has been well validated in the US population. The adequacy components (maximum score) were total fruits (5), whole fruits (5), total vegetables (5), greens and beans (5), whole grains (10), dairy (10), total protein foods (5), seafood and plant proteins (5), and fatty acids (ratio of the sum of polyunsaturated and monounsaturated fatty acids to saturated fatty acids, 10). The moderation components were refined grains (10), sodium (10), added sugars (10), and saturated fats (10). HEI-2015 is a 100-point scale, with a higher score indicating a better quality of overall diet.

Covariates
We included participants’ age (continuous; years), race/ethnicity (non-Hispanic White, non-Hispanic Black, Mexican American, other Hispanic, or other/multiracial), sex (male/female), language of the sample person interview instrument (English, Spanish), educational level (less than high school, high school, higher than high school, and in number of years), smoking status (current, former and never), and family income to poverty ratio (in quartiles) in the analyses. There were no missing values for all variables except for poverty to income ratio (8%); thus, an “unknown” category was created to keep the same number of participants in all models. Models were adjusted for age, sex, marital status, race/ethnicity, family income to poverty ratio, language of the sample person interview instrument, educational level, and diet quality evaluated by the healthy eating index. All these covariates were included in the models as potential confounders and were selected a priori based on previous literature.

Statistical analyses
Means and SDs, and medians and interquartile ranges (IQRs) were calculated for cognitive performance tests scores and blood metals concentrations and also presented by population characteristics. All blood metals concentrations were log10-transformed to reduce the influence of outliers.

We used quantile g-computation to evaluate the joint mixture association of the three blood metals concentrations with the cognitive performance tests scores. This method is a parametric, generalized linear model-based implementation of g-computation. Quantile g-computation is adapted from weighted quantile sum regression, but unlike weighted quantile sum, this method is not constrained to a single direction of effect, does not require splitting of the dataset, and is more robust against unmeasured confounders. In addition, quantile g-computation yields a simple and computationally efficient approach to estimating associations between a mixture of exposures and a health outcome of interest, when we may not be able to rule out confounding or we may be uncertain about the effect direction of some exposures in the mixture. The quantile g-computation allows us to estimate the expected difference in the cognitive performance test scores when simultaneously increasing all blood metals concentrations by one quantile, conditional on covariates. We used quartiles and derived 95% confidence intervals (CIs) of the estimates using the bootstrap method (N = 200). We assigned each metal a weight that was then used in a multivariable linear regression model for each cognitive test score, conditional on covariates listed above. The weights can either be positive or negative and reflect the individual contribution of each metal to the overall mixture effect. We also present metals independent effects on cognitive test scores using traditional multivariable regressions in Supplementary Table 1.

We also investigated sex-specific estimates in stratified analyses because prior studies reported sex-specific associations between metals concentrations and cognitive function.40,41 We also examined dietary modification of the association between metals mixtures and cognitive test scores. To examine differences between estimates from different sex- and dietary-specific strata, we used the method developed by Payton et al. This method allows examination of overlap between confidence intervals or standard error intervals to test hypotheses about the difference between estimates of dietary scores strata. We estimated 4-year survey weights for each participant by dividing the 2-year survey weights by 2 (the number of 2-year cycles NHANES 2011–2012 and 2013–2014) and used them in all analyses to account for the unequal probabilities of inclusion and response rates.

Sensitivity analyses
We ran three sensitivity analyses. First, we performed unweighted analyses that did not account for the survey design because the weighted method is inefficient analysis due to the large variability in assigned weights. The unweighted analysis yields correct estimates when models are adjusted for the auxiliary variables (i.e., age, sex, and ethnicity). Second, we tested effect modification using another alternative definition of the dietary inflammatory index proposed by Shivappa et al.. This score is based on the same inflammatory weights and foods components as in the ADII; however, the foods components were not energy-adjusted to reduce the between-person variation. Finally, the main analysis was restricted to never smokers.

Results
Descriptive statistics
The mean age of participants at screening was 69.5 years (SD = 6.7 years), with slightly fewer men (49%) than women (51%). The mean scores of CERAD, Animal Fluency, DSST, and the overall cognitive performance were 46.4 (SD = 17.1), 16.7 (SD = 5.4), 24.7 (SD = 6.4), and 87.8 (SD = 24.3), respectively. In general, women, non-Hispanic White, highly educated, younger (<69 years), and participants who were in the 4th quartile groups of...
HEI-2015 scores and poverty to income ratio, had higher scores of overall and specific cognitive performance tests. Additionally, participants in the 4th quartile of ADII score—which corresponds to a higher anti-inflammatory diet—had higher test scores compared to participants in the 1st quartile (Table 1).

The median (interquartile range) of blood metals were 0.38 μg/L (0.35), 14.70 μg/L (11.70), and 8.74 μg/L (4.06) for cadmium, lead, and manganese, respectively. Cd concentrations were higher in women and older participants (≥ 80 years) and in participants who were in the 1st quartile groups of the HEI-2015 and the ADII scores (Table 2). Pb concentrations were higher in men and older participants (≥ 80 years) and in participants who were in the 1st quartile groups of the HEI-2015 and the ADII scores (Table 2). In contrast to other metals, blood Mn concentrations were higher in participants in the 4th quartile groups of the HEI-2015 and the ADII scores.

**Associations between blood metals mixture and cognitive performance scores**

In mixture analyses using quantile G-computation, increasing blood concentrations of all metals by one quartile was associated with a decrease in overall cognitive performance (β = –0.04; 95% CI = –0.09, 0.02), CERAD (β = –0.05; 95% CI = –0.12, 0.03), and DSST (β = –0.05; 95% CI = –0.11, 0.02) test scores (Figure 1). No pattern of associations was observed for Animal Fluency test scores (β = –0.02; 95% CI = –0.10, 0.06). The individual weights for each chemical mixture component, as determined by the quantile G-computation approach, were presented in Figure S1 (http://links.lww.com/EE/A173). Lead is the only metal that had a positive weight in each of the cognitive tests and of the two metals that have negative weights, cadmium had the highest percentage except for the DSST test.

In sex-stratified analyses, we observed stronger negative associations in men and slightly positive associations in women. For instance, increasing blood concentrations of all metals by one quartile was associated with lower overall cognitive performance (β = –0.10 SD; 95% CI = –0.17, –0.02) and the DSST test scores (β = –0.17 SD; 95% CI = –0.27, –0.07) in men, whereas the associations in women were positive ([β = 0.02 SD; 95% CI = –0.05, 0.09] and [0.07 SD; 95% CI = –0.01, 0.15], P of heterogeneity: 0.03 and < 0.001, respectively, for overall and DSST test scores). Using traditional multivariable regressions to analyze each metal separately, Cd and Mn showed negative associations with the overall cognitive performance scores, CERAD, and DSST. Again, the associations were stronger in men for Pb and Mn with the DSST score, and Pb with the overall cognitive performance scores; P of heterogeneity < 0.10 for all (see Supplementary Table 1; http://links.lww.com/EE/A173).

**Table 1.**

Means of the overall cognitive function and the specific cognitive performance tests according to characteristics of study participants, NHANES 2011–2014

| Characteristics of participants | Overall cognitive function | DSST test | CERAD test | Animal fluency test |
|---------------------------------|-----------------------------|-----------|------------|---------------------|
|                                 | Means (SD)                  | Means (SD) | Means (SD) | Means (SD)         |
| **Gender**                      |                             |           |            |                     |
| Male                            | 869 (49)                    | 83.7 (23.1)| 23.3 (6.1) | 43.6 (16.2)        | 16.8 (5.5) |
| Female                          | 908 (51)                    | 91.7 (24.7)| 26.0 (6.4) | 49.3 (17.5)        | 16.6 (6.3) |
| **Age at screening (years)**    |                             |           |            |                     |
| 60–69                           | 957 (54)                    | 93.5 (24.4)| 26.0 (6.0) | 49.8 (17.4)        | 17.6 (6.6) |
| 70–79                           | 543 (31)                    | 84.6 (22.8)| 24.0 (6.4) | 44.3 (16.2)        | 16.3 (6.2) |
| ≥80                             | 277 (15)                    | 74.4 (20.0)| 21.1 (6.3) | 38.7 (14.1)        | 14.6 (4.4) |
| **Education level**             |                             |           |            |                     |
| Less than 9th grade             | 199 (11)                    | 62.0 (16.0)| 20.6 (5.9) | 27.1 (11.0)        | 14.3 (4.3) |
| 9–11th grade                   | 232 (13)                    | 75.0 (20.0)| 23.1 (6.1) | 37.3 (13.9)        | 14.7 (4.9) |
| High-school graduate            | 431 (24)                    | 84.8 (21.5)| 24.4 (6.1) | 44.7 (15.0)        | 15.7 (4.9) |
| Some college or AA degree       | 496 (28)                    | 95.4 (21.3)| 26.0 (6.1) | 51.8 (14.9)        | 17.6 (6.3) |
| College graduate or above       | 417 (24)                    | 101.3 (22.1)| 26.3 (6.4) | 56.0 (15.1)        | 19.1 (5.7) |
| **Poverty to income ratio**     |                             |           |            |                     |
| Quartile 1                      | 408 (23)                    | 76.2 (22.6)| 23.2 (6.4) | 37.7 (16.0)        | 15.3 (5.2) |
| Quartile 2                      | 410 (23)                    | 82.8 (21.7)| 24.0 (6.2) | 42.7 (15.2)        | 16.1 (4.9) |
| Quartile 3                      | 406 (23)                    | 92.6 (21.5)| 25.5 (6.3) | 50.1 (14.6)        | 17.1 (6.3) |
| Quartile 4                      | 413 (23)                    | 101.2 (23.1)| 26.2 (6.5) | 56.3 (15.8)        | 18.8 (6.6) |
| Missing                         | 141 (8)                     | 82.7 (24.8)| 24.1 (6.3) | 42.9 (18.1)        | 15.6 (4.9) |
| **Race/Hispanic origin**        |                             |           |            |                     |
| Mexican American                | 137 (8)                     | 79.7 (22.7)| 23.5 (5.8) | 40.1 (16.9)        | 16.1 (4.6) |
| Other Hispanic                  | 181 (10)                    | 75.3 (22.8)| 22.9 (6.1) | 36.8 (16.7)        | 15.7 (4.9) |
| Non-Hispanic White              | 897 (50)                    | 94.0 (24.0)| 25.0 (6.6) | 51.0 (16.1)        | 18.0 (6.6) |
| Non-Hispanic Black              | 417 (24)                    | 80.6 (22.8)| 24.7 (6.3) | 40.7 (15.9)        | 15.2 (6.1) |
| Other race—including multiracial| 145 (8)                     | 93.6 (19.6)| 26.1 (5.9) | 52.3 (14.8)        | 15.2 (4.6) |
| **ADII**                        |                             |           |            |                     |
| Quartile 1                      | 445 (25)                    | 87.0 (23.0)| 24.9 (6.3) | 45.5 (15.9)        | 16.7 (6.5) |
| Quartile 2                      | 443 (25)                    | 83.6 (24.9)| 24.0 (6.8) | 43.2 (17.3)        | 16.3 (6.3) |
| Quartile 3                      | 444 (25)                    | 85.7 (24.1)| 24.3 (5.3) | 45.4 (17.2)        | 16.1 (6.1) |
| Quartile 4                      | 445 (25)                    | 94.9 (23.6)| 25.5 (6.2) | 51.6 (16.7)        | 17.8 (6.6) |
| **HEI-2015**                    |                             |           |            |                     |
| Quartile 1                      | 445 (25)                    | 85.3 (24.5)| 24.5 (6.7) | 44.3 (16.7)        | 16.4 (6.5) |
| Quartile 2                      | 444 (25)                    | 84.8 (22.7)| 24.1 (6.0) | 44.5 (16.2)        | 16.3 (6.1) |
| Quartile 3                      | 444 (25)                    | 88.5 (24.9)| 24.6 (6.4) | 47.0 (17.6)        | 16.9 (6.5) |
| Quartile 4                      | 444 (25)                    | 92.6 (24.2)| 25.5 (6.4) | 49.8 (17.2)        | 17.3 (5.4) |

AA indicates associate of arts.
index and Figure 3 for the healthy eating index. The associations were more pronounced in the lower quartiles of the dietary inflammatory scores, which represent, high pro-inflammatory diet. We found null or positive though imprecise associations in participants with a high anti-inflammatory diet, especially the third quartile of the ADII score. This pattern of associations was more apparent in men. For example, blood metal mixture concentrations were positively associated with the overall cognitive test scores among men with more pro-inflammatory diet, while among those with low-diet quality, the association was observed for the associations between blood metal mixture concentrations and the animal fluency. An inverse association was observed in men with high-diet quality (β = –0.22; 95% CI, –0.47, 0.03) for quartile 4, and a positive association in those with the low-quality diet (β = 0.12; 95% CI, –0.08, 0.32 for quartile 1; P of heterogeneity: 0.03).

When the models were unweighted, we observed the same pattern of associations (Figs. S2–S4; http://links.lww.com/EE/A173). However, using an alternative definition of dietary inflammatory index showed different results. The associations between blood metal mixture concentrations and cognitive function were more pronounced in women from quartile 1 of the DII, while these associations were more pronounced in men from quartile 1 of the ADII (Fig. S5; http://links.lww.com/EE/A173). Analyses restricted to nonsmokers (46% of the total sample) showed the same pattern of associations as the analysis on the entire population.

### Discussion

In this cross-sectional study of US adults ≥60 years old, we found that concentrations of metals and metal mixtures were negatively associated with cognitive performance test scores in men but not in women. In addition, we found that an anti-inflammatory diet
Figure 1. Overall and sex-specific adjusted beta coefficients (95% CI) for the overall cognitive function score and the specific cognitive performance tests, for increase in all blood metals by one quartile, NHANES 2011–2014. Model was adjusted for age (years), gender (except for stratified analysis on this variable), race/ethnicity (Mexican American, other Hispanic, Non-Hispanic White, Non-Hispanic Black, other race), language of the sample person interview instrument (English, Spanish), educational level (less than high school, high school, higher than high school), smoking status, marital status, poverty to income ratio (quartiles), and healthy eating index.

Figure 2. Estimates (95% CI) of the associations of a one quartile increase in the metal mixture and overall cognitive function score and specific cognitive performance tests, by sex and ADII score, NHANES 2011–2014. Model was adjusted for age (years), race/ethnicity (Mexican American, other Hispanic, Non-Hispanic White, Non-Hispanic Black, other race), language of the sample person interview instrument (English, Spanish), educational level (less than high school, high school, higher than high school), smoking status, marital status, poverty to income ratio (quartiles), and healthy eating index. All P values for between ADII quartile tests of heterogeneity were >0.10 except for men (P of heterogeneity: 0.09 and 0.07 for the overall cognitive test and the animal fluency test scores, respectively).
may mitigate the effects of metals. To our knowledge, this is the first study to investigate the potential effect of dietary modification of the association between blood metals concentrations and cognitive performance test scores in older adults.

The trace metals examined here had been previously evaluated for their effect on cognitive function. For example, in a study in an older adults population with the same NHANES data reported an inverse association between blood Cd concentrations and cognitive function scores. Two other studies in older adults in Brazil and China reported inverse associations between blood Cd concentrations and cognitive function. Manganese and lead have also been associated with diminished cognitive function in older adults. A recent meta-analysis showed that higher blood Pb concentrations are associated with poorer verbal abilities, visuospatial abilities, memory, attention, and psychomotor function. For the association between adult’s blood Mn and cognitive function, few studies reported inverse associations. Only a few of these studies have explored sex-specific associations and none evaluated the potential effect modification by diet in the association between metals and cognitive scores. One of the main differences of these studies compared to ours is that they focused on specific metals and did not take into account the simultaneous exposure to several metals and the potential joint associations of these metals with cognitive function.

Inflammation and oxidative stress are two key mechanisms for metal neurotoxicity. Acute high-level exposure or long-term persistent low-level exposure to Cd interferes with the antioxidant defense system and leads to oxidative stress in neuronal cells and brain endothelial cells. The mechanism by which Pb exposure affects cognition in older adults has yet to be revealed, but the existing reports suggest that the presence of Pb in the brain causes a potential pro-inflammatory effect on the central nervous system and neuronal death may be related to the production of various cytokines and chemokines. For Mn, the underlying mechanisms include oxidative stress and mitochondrial dysfunction, autophagy dysregulation, accumulation of intracellular toxic metabolites, and apoptosis. Excess manganese can impair manganese superoxide dismutase activity, thus increasing reactive oxygen species production and eventually leading to oxidative stress and mitochondrial dysfunction.

A number of studies have provided evidence that diet plays a role in the pathogenesis of many neurological diseases such as Alzheimer’s disease, Parkinson’s disease, and multiple sclerosis. In our study, we observed that adherence to an anti-inflammatory diet can attenuate the potential adverse effect of higher metals concentrations on cognitive performance test scores. In older adults, no study has evaluated this potential effect modification of diet in the association between blood metals concentrations and cognitive performance test scores.

Trace elements exhibit human toxicity and disease through inflammation and oxidative stress. Thus, it is conceivable that nutrients that can contribute to cellular oxidative stress also could exacerbate or amplify environmental toxicity. On the other hand, nutrients that have antioxidant or anti-inflammatory activity could reduce or prevent compromised health or disease induction from environmental pollutants. For example, antioxidants such as vitamin C supplementation in lead-exposed animals significantly reduce blood, liver, and renal lead levels and associated biochemical changes. The main hypothesis is that vitamin C decreases the intestinal absorption of lead, possibly by reducing ferric iron to ferrous iron in the duodenum and increase the availability of iron, which competes with lead for intestinal absorption. Folic acid supplementation can lower blood arsenic concentrations and plant-derived bioactive nutrients can lower cardiovascular and cancer risks linked to pollutant exposures. Data also indicate that bioactive food components such as polyphenols and omega-3 polyunsaturated fatty acids may scavenge the reactive oxygen species and free radicals before they can activate pathways in pathogenesis of diseases and prevent or decrease toxicant-induced inflammation.

Our study has several notable strengths, including the analysis of a nationally representative sample, the use of objective...
Our findings suggest that blood concentrations of metals mixture was negatively associated with cognitive performance test scores among older adults in the United States, especially in men. This is in line with previous findings of adverse associations between specific metals and some domains or overall cognitive test scores in predominantly male studies. This result contributes to the existing literature on the health effects of short-term exposure to metals, rather than concurrent measurements in blood that only reflect recent exposure. However, we do not have data on all these metals measured in the organs where they are stored as a proxy for long-term exposure in this wave of the NHANES study. Overall, future studies should include longitudinal data, particularly on diet, and consider metals biomarkers of chronic exposures as a mixture and thus better characterize the associated risk.

Conclusions

Acknowledgments

We thank the Philippe Foundation for their contribution to and ongoing support of scientific and medical activities and research by facilitating Franco-American exchanges.

References

1. Milte CM, Ball K, Crawford D, McNaughton SA. Diet quality and cognitive function in mid-aged and older men and women. BMC Geriatr. 2019;19:361.
2. Gurland BJ, Page WF, Plassman BL. A twin study of the genetic contribution to age-related functional impairment. J Gerontol A Biol Sci Med Sci. 2004;59:839–863.
3. Livingston G, Sommerlad A, Orgeta V, et al. Dementia prevention, intervention, and care. Lancet. 2017;390:2673–2734.
4. Li H, Wang Z, Fu Z, et al. Associations between blood cadmium levels and cognitive function in a cross-sectional study of US adults aged 60 years or older. BMJ Open. 2018;8:e020533.
5. Vlaski T, Jordakieva G, Gnamts B, et al. Blood lead levels and cognitive functioning: a meta-analysis. Sci Total Environ. 2019;668:678–684.
6. Mohajerzadeh Delgado-Saborit JM, Sly PD, et al. Environmental chronic exposure to metals and effects on attention and executive function in the general population. Sci Total Environ. 2020;705:135911.
7. Angeloni C, Businaro R, Vauzour D. The role of diet in preventing and reducing cognitive decline. Curr Opin Psychiatri. 2020;34:432–438.
8. Chen P, Miah MR, Aschner M. Metals and neurodegeneration. F1000Res. 2016;5:F1000 Faculty Rev–F1000 Faculty 366.
9. Iqbal G, Zada W, Mannan A, Ahmed T. Elevated heavy metals levels in cognitively impaired patients from Pakistan. Environ Toxicol Pharmocol. 2018;60:100–109.
10. Weuve J, Korrick SA, Weisskopf MG, et al. Cumulative exposure to lead in relation to cognitive function in older women. Environ Health Perspect. 2009;117:574–580.
11. Van Winggaard E, Winters PC, Cory-Slechta DA. Blood lead levels in relation to cognitive function in older U.S. adults. NeuroToxicology. 2011;32:110–115.
12. Shih RA, Glass TA, Bandeen-Roche K, et al. Environmental lead exposure and cognitive function in community-dwelling older adults. Neurology. 2006;67:1556–1562.
13. Yu YL, Thijs L, Saenen N, et al. Two-year neurocognitive responses to first occupational lead exposure. Scand J Work Environ Health. 2021;47:233–243.
14. Horton CJ, Weng HY, Wells EM. Association between blood lead level and subsequent Alzheimer’s disease mortality. Environ Epidemiol. 2019;3:e0045.
15. Chibowska K, Baranowska-Bosiacka I, Falkowska A, Gutowska I, Goschorska M, Chlubek D. Effect of lead (Pb) on inflammatory processes in the brain. Int J Mol Sci. 2016;17:E2140.
16. Gouaref I, Kocer EA. Trace elements modulates oxidative stress in type 2 diabetes. Diabetes Food Plan. 2018.
17. Garza-Lombó G, Posadas Y, Quintanar L, Gonsébatt ME, Franco R. Neurotoxicity linked to dysfunctional metal ion homeostasis and xenobiotic metal exposure: redox signaling and oxidative stress. Antioxid Redox Signal. 2018;28:1669–1703.
18. Markesbery WR. Oxidative stress hypothesis in Alzheimer’s disease. Free Radic Biol Med. 1997;23:134–147.
19. Calder PC, Aklwala N, Brouns F, et al. Dietary factors and low-grade inflammation in relation to overweight and obesity. Br J Nutr. 2011;106(suppl 3):S5–S78.
20. Blomquist C, Alvehus M, Berún J, et al. Attenuated low-grade inflammation following long-term dietary intervention in postmenopausal women with obesity. Obesity. 2017;25:892–900.
21. Colacino JA, Arthur AE, Ferguson KK, Rozek LS. Dietary antioxidant and anti-inflammatory intake modifies the effect of cadmium exposure on markers of systemic inflammation and oxidative stress. Environ Res. 2014;131:6–12.
22. Ouelhote Y, Lanphear B, Braun JM, et al. Gestational exposures to phthalates and folic acid, and autistc traits in Canadian children. Environ Health Perspect. 2020;128:27004.
23. Schmidt RJ, Kogan V, Shelton JF, et al. Combined prenatal pesticide exposure and folic acid intake in relation to autism spectrum disorder. Environ Health Perspect. 2017;125:097007.
24. Goodrich AJ, Volk HE, Tarenzi DJ, et al. Joint effects of prenatal air pollutant exposure and maternal folic acid supplementation on risk of autism spectrum disorder. Autism Res. 2018;11:69–80.
25. Loftus CT, Hazlehurst MF, Szapiro AA, et al. Prenatal air pollution and childhood IQ: preliminary evidence of effect modification by folate. *Environ Res*. 2019;176:108505.

26. Li Q, Wang Y, Guo Y, et al. Folic acid supplementation and the association between maternal airborne particulate matter exposure and preterm delivery: a National Birth Cohort Study in China. *Environ Health Perspect*. 2020;128:127010.

27. Morris JC, Heyman A, Moths RC, et al. The Consortium to Establish a Registry for Alzheimer’s Disease (CERAD). Part I. Clinical and neuropsychological assessment of Alzheimer’s disease. *Neurology*. 1989;39:1159–1165.

28. Strauss E, Sherman EMS, Spreen O. *A Compendium of Neuropsychological Tests: Administration, Norms, and Commentary*. American Chemical Society; 2006.

29. Saklofske DH, Schoenberg MR. Wechsler Adult Intelligence Scale (All Versions). In: Kreutzer JS, DeLuca J, Caplan B, eds. *Encyclopedia of Clinical Neuropsychology*. Springer, New York, NY; 2011. https://doi.org/10.1007/978-0-387-39948-3-1073

30. Lauaoli N, Mancini FR, Haji-Louna M, et al. Dietary inflammatory index and type 2 diabetes risk in a prospective cohort of 70,991 women followed for 20 years: the mediating role of BMI. *Diabetologia*. 2019;62:2222–2232.

31. Van Wouwdenbergh GJ, Theofylaktopoulos D, Kuijsten A, et al. Adapted dietary inflammatory index and its association with a summary score for low-grade inflammation and markers of glucose metabolism: the Cohort study on Diabetes and Atherosclerous Maastricht (CODAM) and the Hoorn study. *Am J Clin Nutr*. 2013;98:1533–1542.

32. Shivappa N, Steck SE, Hurley TG, Hussey JR, Hébert JR. Designing and developing a literature-derived, population-based dietary inflammatory index. *Public Health Nutr*. 2014;17:1689–1696.

33. Cavicchia PP, Steck SE, Hurley TG, et al. A new dietary inflammatory index predicts interval changes in serum high-sensitivity C-reactive protein. *J Nutr*. 2009;139:2365–2372.

34. Willett WC, Howe GR, Kushi LH. Adjustment for total energy intake in epidemiologic studies. *Am J Clin Nutr*. 1997;66(suppl 4):1220S–1228S.

35. Krebs-Smith SM, Pannucci TE, Subar AF, et al. Update of the Healthy Eating Index: HEI-2015. *J Acad Nutr Diet*. 2016;118:1622–1633.

36. Panizza CE, Shvetsov YB, Harmon BE, et al. Testing the predictive validity of the healthy eating index-2015 in the multithoracic cohort: is the score associated with a reduced risk of all-cause and cause-specific mortality? *Nutrients*. 2018;10:452.

37. Reedy J, Lerman JL, Krebs-Smith SM, et al. Evaluation of the Healthy Eating Index-2015. *J Acad Nutr Diet*. 2018;118:1622–1633.

38. Robins JM. Causal models for estimating the effects of weight gain on mortality. *Int J Obes (Lond)*. 2008;32(suppl 3):S15–S41.

39. Keil AP, Buckley JP, O’Brien KM, Ferguson KK, Zhao S, White AJ. A quantile-based g-computation approach to addressing the effects of exposure mixtures. *Environ Health Perspect*. 2020;128:47004.

40. Zhou T, Guo J, Zhang J, et al. Mediterranean diet and the association between air pollution and cardiovascular disease mortality risk. *Circulation*. 2019;139:2798–2804.

41. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.

42. Gao S, Jin Y, Unverzagat FW, et al. Trace element levels and cognitive function in rural elderly Chinese. *J Gerontol A Biol Sci Med Sci*. 2008;63:635–641.

43. Ghazali AR, Kamarulzaman F, Normah CD, et al. Levels of metallic elements and their potential relationships to cognitive function among elderly from Federal Land Development Authority (FELDA) settlement in Selangor Malaysia. *Biol Trace Elem Res*. 2013;153:16–21.

44. Roels HA, Bowler RM, Kim Y, et al. Manganese exposure and cognitive deficits: a growing concern for manganese neurotoxicity. *Neurotoxicology*. 2012;33:872–880.

45. Figueiredo-Pereira ME, Yakushin S, Cohen G. Disruption of the intracellular sulfhydryl homeostasis by cadmium-induced oxidative stress leads to protein thiolation and ubiquitination in neuronal cells. *J Biol Chem*. 1998;273:12703–12709.

46. Tsubota S, Wang HJ, Carey J, Banks W, Ercal N. Effects of lead and cadmium on brain endothelial cell survival, monolayer permeability, and crucial oxidative stress markers in an in vitro model of the blood-brain barrier. *Toxicol*. 2014;2:258–275.

47. Struzynska L, Dabrowska-Bouta B, Koza K, Sulkowski G. Inflammation-like g response in lead-exposed immature rat brain. *Toxict Sci*. 2007;95:156–162.

48. Guilate TR. Manganese neurotoxicity: new perspectives from behavioral, neuroimaging, and neuropathological studies in humans and non-human primates. *Front Aging Neurosci*. 2013;5:23.

49. Peres TV, Schettinger MR, Chen F, et al. “Manganese-induced neurotoxicity: a review of its behavioral consequences and neuroprotective strategies.” *BMC Pharmacol Toxicol*. 2016;17:57.

50. Gunter TE, Gavin CE, Aschner M, Gunter KK. Speciation of manganese in cells and mitochondria: a search for the proximal cause of manganese neurotoxicity. *Neurotoxicology*. 2006;27:765–776.

51. Bakulska KM, Seo YA, Hickman RG, et al. Heavy metals exposure and Alzheimer’s disease and related dementias. *J Alzheimer’s Dis*. 2020;76:1215–1242.

52. Thrh E, Shivappa N, Mann JR, Hébert JR, Wirth MD, Loprinzi PD. Dietary inflammatory index and memory function: population-based national sample of elderly Americans. *Br J Nutr*. 2018;119:552–538.

53. Yang F, Wolk A, Häkansson N, Pedersen NL, Wirdefeldt K. Dietary antioxidants and risk of Parkinson’s disease in two population-based cohorts. *Mov Disord*. 2017;32:1631–1636.

54. Zhao X, Zhang M, Li C, Jiang X, Su Y, Zhang Y. Benefits of vita mins in the treatment of Parkinson’s disease. *Oxid Med Cell Longev*. 2019;2019:942687.

55. Vj AG, Satija NK, Flora SJ. Lead induced disorders in hematopoietic and drug metabolizing enzyme system and their protection by ascorbic acid supplementation. *Biomed Environ Sci*. 1998;11:7–14.

56. Shahlan MG, Mostafa MS, Hassouna MM, El-Nabi SE, El-Refaie A. Amelioration of lead toxicity on rat liver with vitamin C and silymarin supplements. *Toxicology*. 2005;206:1–15.

57. Vahter ME. Interactions between arsenic-induced toxicity and nutrition in early life. *J Nutr*. 2007;137:2798–2804.

58. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between air pollution and cardiovascular disease mortality risk. *Circulation*. 2019;139:1766–1775.

59. Kordas K, Lönnerdal B, Stoltzfus RJ. Interactions between nutrition and manganese exposures in school-age children: a prospective cohort study. *Biol Trace Elem Res*. 2019;139:1766–1775.

60. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.

61. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.

62. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.

63. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.

64. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.

65. Lim CC, Hayes RB, Ahn J, et al. Mediterranean diet and the association between heavy metal exposure and poor working memory and possible mediation effect of antioxidant defenses during aging. *Sci Total Environ*. 2017;575:759–757.