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Research article

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Abstract

Background

Copper is an essential trace metal with potential interest for cardiovascular effects. Few studies have explored the association between copper and blood pressure in children and adolescents (hereafter referred to as youths).

Method:

We conducted a cross-sectional analysis of 1242 youths aged 8–17 years who participated in the 2011 to 2016 National Health and Nutrition Examination Survey. Using 2017 American Academy of Pediatrics guidelines, Elevated Blood Pressure (EBP) was defined as a mean systolic and/or diastolic blood pressure (BP) ≥ 90th percentile for sex, age, and height for children aged 1–12 years and systolic BP ≥ 120 mmHg or diastolic BP ≥ 80 mmHg for adolescent age 13–17 years. Mean serum copper was 114.17 µg/dL.

Results

After multiple adjustments, dose-response analyses revealed that EBP was associated with progressively higher serum copper concentrations in a nonlinear trend. In comparison with the lowest quartile of serum copper concentrations, the adjusted odds of EBP for the highest quartile was 5.26 (95% confidence interval [CI], 2.76–10.03).

Conclusion

Our results suggested that high serum copper concentrations were significantly associated with EBP in US youths.

Background

Elevated Blood Pressure (EBP) among youths has become an important public health challenge in the United States [1]. Recent epidemiological studies have suggested that accumulation of essential trace metals may play a critical role in the development of hypertension [2–4].

Copper is an essential trace metal with antioxidant properties mediated through many redox enzymes [5]. Overload of this metal leads to Fenton-type redox reactions, resulting in oxidative injury [6]. Results of pathological accumulation of copper are observed in Wilson's disease, a recessively inherited disorder of copper metabolism [7]. Additionally, evidence for the relationship between high copper concentrations and heart failure, Parkinson's disease, and ischemic stroke were found in epidemiological studies [8–10]. However, evidence regarding the association between serum copper and blood pressure (BP) have been
inconsistent or conflicting and most research has been conducted in adult populations [11–14]. A recent study found serum copper to be associated with several cardiovascular disease risk factors in US youths, including increased total cholesterol, glycohemoglobin and fasting insulin. Nevertheless, no association was found with BP levels [15]. However, this study used BP as a continuous variable and ignored the impact of age, sex and height on the definition of blood pressure level in youths.

In the current study, using the data from the National Health and Nutrition Examination Survey (NHANES) 2011–2016, and the new 2017 American Academy of Pediatrics guidelines (AAP) [1], we aimed to evaluate the association and dose-response relationship between serum copper and EBP in US youths.

**Methods**

**Study population**

NHANES is a nationally representative, multistage survey of the noninstitutionalized US civilian population. All participants provided written informed consent (parental consent was obtained for those < 18 years) and NHANES was approved by the National Center for Health Statistics’ Ethics Review Board. Detailed information on NHANES data collection and survey procedures used in this analysis are publicly available and can be found elsewhere [16].

Participants who had completed blood pressure reading and serum copper measure were included in our analyses. Individuals missing important covariates were excluded. The process of data inclusion is presented in Figure 1.

**Study exposure**

Serum specimens were processed, stored at appropriate temperatures (-70 °C), and shipped to the Division of Laboratory Sciences, National Center for Environmental Health, Centers for Disease Control and Prevention, Atlanta, GA for analysis. Inductively coupled plasma dynamic reaction cell mass spectrometry (ICP-DRC-MS) was used for trace level elemental analysis. The isotopes measured included zinc (m/z 64), copper (m/z 65), and selenium (m/z 78) and the internal standard gallium (m/z 71). Serum samples were diluted 1+1+28 with water and diluent containing gallium (Ga) for multi-internal standardization. If you want, you can reference the technical stuff:


**Definition of EBP**

BP measurements were obtained by a certified examiner using an appropriate cuff size and a mercury manometer. Three BP readings were obtained after the participant had rested for 5 minutes in a seated position with feet flat on the floor. Mean systolic and diastolic BP for each participant were calculated from the recorded readings. Using the 2017 American Academy of Pediatrics guidelines, we classified participants as having EBP or normal BP consistent with previous reports [17]. For children aged 1-13
years, EBP was defined as the mean systolic and/or diastolic BP percentile $\geq$ 90th percentile for sex, age, and height. For adolescents aged 13-17 years, EBP was defined as SBP at least 120 mmHg and/or DBP at least 80 mmHg.

**Covariates**

Demographic information included age, gender, race (Mexican American, other Hispanic non-Hispanic White, non-Hispanic Black, and other race) and family monthly poverty level index category. Other covariates included body mass index (BMI), serum cotinine, physical activity, energy intake and other trace metals (serum selenium, serum zinc, urinary manganese, urinary lead, urinary strontium, urinary arsenic and urinary mercury). BMI was calculated as measured weight in kilograms divided by measured height in meters squared, and BMI percentiles were calculated based on the CDC’s BMI-for-age sex-specific growth charts. BMI categories were defined as follows: “normal,” BMI < 85th percentile; “overweight” and “obese,” BMI $\geq$ 85th percentile. Physical activity was assessed by the amount of television, video game and computer usage daily and was classified as high ($\leq$ 2) and low ($> 2$) in concordance with previous report [18]. Energy intake data were obtained from two 24-hour dietary recall interviews and calculated as an average of 2-day energy intake. Serum cotinine, the primary proximal metabolite of nicotine, is generally regarded as the marker of exposure to environmental tobacco smoke.

**Statistical analysis**

Differences between groups were tested by the Chi-square test for categorical data and the independent Student t-test, Mann-Whitney-U test, analysis of variance, or the Kruskal-Wallis test for continuous data, as appropriate. Binary logistic regression models were performed to estimate the association between serum copper and EBP. In multivariate logistic regressions, model 1 adjusted for age and sex, model 2 further adjusted for race, family monthly poverty level category, physical activity, BMI, total energy intake, serum cotinine, and other trace metals. We assessed for collinearity between adjustment variables by calculating variance inflation factors. The dose-response relationship was conducted by restricted cubic spline with three knots. We performed tests for linear trend by entering the median value of each category of copper as a continuous variable in the models. All analyses were performed using Stata 15.1 and R 3.3.0 software. All reported probabilities ($p$-values) were two-sided with $p < 0.05$ considered as significant.

**Results**

Overall, the study cohort included a total of 1242 US youths aged 8–17 years at baseline. Table 1 presents the characteristics of the study participants. The mean serum copper concentrations was 114.17 µg/dL. The overall prevalence of EBP was 14.0%. EBP was more likely to occur in males (132 versus 42, $P < 0.01$) and in Non-Hispanic Black participants ($P < 0.01$). Participants with EBP tend to be older (12.9 versus 12.0 years, $P = 0.01$), overweight and obese ($P < 0.01$), consumed more calories (2152 versus 1878 kcal, $P < 0.01$) and had lower amount of physical activity ($P = 0.01$) compared to those without EBP. There was no difference in serum copper concentrations between genders (male versus female, 114.3 versus 114.1 µg/dL, $P = 0.89$). Non-Hispanic blacks had higher mean serum copper
concentrations compared to non-Hispanic whites, Mexican-Americans and other Hispanic (122.4, 112.1, 110.9 and 109.5 µg/dL, respectively, $P<0.01$). Mean serum copper concentrations were higher in participants with EBP compared to those without EBP (125.5 versus 112.3 µg/dL, $P<0.01$).
<table>
<thead>
<tr>
<th></th>
<th>Total (n = 1242)</th>
<th>1st (n = 312)</th>
<th>2nd (n = 309)</th>
<th>3rd (n = 312)</th>
<th>4th (n = 309)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>624 (50.2)</td>
<td>168 (53.8)</td>
<td>135 (43.7)</td>
<td>138 (44.2)</td>
<td>183 (59.2)</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Race/Ethnicity</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td>&lt; 0.001</td>
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<tr>
<td>Mexican American</td>
<td>261 (21.0)</td>
<td>60 (19.2)</td>
<td>81 (26.2)</td>
<td>66 (21.2)</td>
<td>54 (17.5)</td>
<td></td>
</tr>
<tr>
<td>Other Hispanic</td>
<td>120 (9.7)</td>
<td>30 (9.6)</td>
<td>33 (10.7)</td>
<td>39 (12.5)</td>
<td>18 (5.8)</td>
<td></td>
</tr>
<tr>
<td>Non-Hispanic White</td>
<td>300 (24.2)</td>
<td>78 (25.0)</td>
<td>87 (28.2)</td>
<td>72 (23.1)</td>
<td>63 (20.4)</td>
<td></td>
</tr>
<tr>
<td>Non-Hispanic Black</td>
<td>357 (28.7)</td>
<td>75 (24.0)</td>
<td>63 (20.4)</td>
<td>81 (26.0)</td>
<td>138 (44.7)</td>
<td></td>
</tr>
<tr>
<td>Other Race</td>
<td>204 (16.4)</td>
<td>69 (22.1)</td>
<td>45 (14.6)</td>
<td>54 (17.3)</td>
<td>36 (11.7)</td>
<td></td>
</tr>
<tr>
<td>Family monthly poverty level category</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.001</td>
</tr>
<tr>
<td>&lt; 1.30</td>
<td>579 (46.6)</td>
<td>144 (46.2)</td>
<td>123 (39.8)</td>
<td>138 (44.2)</td>
<td>174 (56.3)</td>
<td></td>
</tr>
<tr>
<td>1.31 ~ 1.85</td>
<td>174 (14.0)</td>
<td>57 (18.3)</td>
<td>51 (16.5)</td>
<td>39 (12.5)</td>
<td>27 (8.7)</td>
<td></td>
</tr>
<tr>
<td>&gt; 1.85</td>
<td>438 (35.3)</td>
<td>105 (33.7)</td>
<td>126 (40.8)</td>
<td>117 (37.5)</td>
<td>90 (29.1)</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>12.2 (2.9)</td>
<td>13.5 (2.5)</td>
<td>12.2 (2.9)</td>
<td>11.7 (3.0)</td>
<td>11.3 (2.9)</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>BMI</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Normal weight</td>
<td>753 (60.6)</td>
<td>243 (77.9)</td>
<td>198 (64.1)</td>
<td>177 (56.7)</td>
<td>135 (60.6)</td>
<td></td>
</tr>
<tr>
<td>Overweight and obese</td>
<td>489 (39.4)</td>
<td>69 (22.1)</td>
<td>111 (35.9)</td>
<td>135 (43.3)</td>
<td>174 (39.4)</td>
<td></td>
</tr>
<tr>
<td>Energy intake (kcal/day)</td>
<td>1920 (635)</td>
<td>2000 (681)</td>
<td>1900 (661)</td>
<td>1870 (559)</td>
<td>1890 (630)</td>
<td>0.063</td>
</tr>
</tbody>
</table>

Abbreviations: BMI, body mass index.
Table 2 presents the odds ratios of EBP based on quartiles of serum copper concentrations. In crude logistic regression analyses, high levels of serum copper were significantly associated with increased risk of EBP. After adjustment for age and sex (model 1), serum copper concentrations had a significantly positive association with EBP. After further adjustment for race, family monthly poverty level category, physical activity, BMI, total energy intake, serum cotinine, and other trace metals (model 2), the results remained stable and statistically significant. Dose-response relationship between serum copper and EBP were shown in Fig. 2. In restricted cubic spline model, excessive serum copper increased the risk of EBP in a linear manner ($P = 0.029$).
Table 2
Weighted Odds Ratios and 95% Confidence Intervals for Elevated Blood Pressure According to Quartiles of Serum Copper.

<table>
<thead>
<tr>
<th>Serum copper (µg/dL)</th>
<th>Events/Cases</th>
<th>OR(95%CI)</th>
<th>Crude</th>
<th>Model1&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Model 2&lt;sup&gt;b&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Quartile 1</td>
<td>27/312</td>
<td>Ref.</td>
<td>Ref.</td>
<td>Ref.</td>
<td>Ref.</td>
</tr>
<tr>
<td>Quartile 2</td>
<td>30/279</td>
<td>1.14 (0.67–1.96)</td>
<td>1.87 (1.05–3.32)</td>
<td>1.80 (0.94–3.43)</td>
<td></td>
</tr>
<tr>
<td>Quartile 3</td>
<td>45/267</td>
<td>1.78 (1.07–2.95)</td>
<td>3.37 (1.84–5.85)</td>
<td>3.70 (1.97–6.95)</td>
<td></td>
</tr>
<tr>
<td>Quartile 4</td>
<td>72/237</td>
<td>3.21 (2.00-5.16)</td>
<td>5.44 (3.21–9.22)</td>
<td>5.26 (2.76–10.03)</td>
<td></td>
</tr>
<tr>
<td>P for trend</td>
<td>NA</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
<td>&lt; 0.001</td>
<td></td>
</tr>
<tr>
<td>Per 1 SD serum copper</td>
<td></td>
<td>1.52 (1.32–1.75)</td>
<td>1.76 (1.50–2.06)</td>
<td>1.69 (1.38–2.07)</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: CI, confidence interval; OR, odds ratio; NA, not applicable; SD, Standard Deviation.

<sup>a</sup>, Model 1 were adjusted by age and sex.

<sup>b</sup>, Model 2 were adjusted by age, sex, race, family monthly poverty level category, physical activity, body mass index (BMI), total energy intake, serum cotinine, and other trace metals.

*P < 0.05.

**P < 0.01.

**Discussion**

In this study, using a nationally representative large-scale database and updated classification system for pediatric BP, we found that serum copper was associated with increased risk of EBP in the US population aged between 8 to 17 years old. The association was statistically significant despite adjustments for the impact of co-exposure to multiple metals [19, 20].

In animal experiments, plasma copper concentrations were significantly higher in hypertensive rats [21]. Other animal studies have demonstrated implanting a copper cuff results in neointimal thickening in response to vascular injury [22], while copper chelators inhibit the development of vascular inflammation and new intima formation in response to vascular injury [23, 24]. In humans, evidence linking serum copper and blood pressure has been inconsistent. Most research conducted thus far has been in adults. Taneja, S.K et al [25], conducted a study with 500 adult participants which noted higher levels of copper in urine in hypertensive subjects (2.70 ± 0.10 versus 5.14 ± 0.15 mg/dL). Conversely, Li Y et al [26], found
no difference. Additionally, Kim MH et al [27]. found that copper intake was inversely associated with SBP and DBP in 258 adult subjects. There is a dearth of studies investigating the association between copper and blood pressure in youths. In a cross-sectional study conducted among 1427 US youth, no association was found between serum copper concentrations and blood pressure[15]. Another study among European children suggested a positive association between serum copper and diastolic blood pressure [28]. However, a primary deficit of these studies was that they did not use blood pressure percentiles to define hypertension, which is the current standard for delineating normotension from hypertension in youths. Using the 2017 hypertension diagnostic criteria, our results indicate that high serum copper concentrations increase the risk of EBP in youths.

The mechanisms behind the association between serum copper and BP remain unclear. It has been reported that Reactive oxygen species (ROS) disrupts the balance of oxidation and antioxidant systems. Excessive copper can affect the activity of ROS [29]. The resultant increase in superoxide may lead to the impairment and dysfunction of endothelial structure, and consequently result in the development of EBP [30]. Furthermore, high copper concentrations can reduce myosin-ATPase activity, resulting in calcium overload [31]. Increased Ca$^{2+}$ in smooth muscular layer of vessels may cause an increase in the arterial wall tension finally resulting in EBP [32].

As suggested by previous literature, serum copper appears to reflect the status of copper nutrition in both depleted and replete populations. Pediatric reference intervals for serum copper concentrations are not well established [33]. Only a few studies have measured serum copper concentrations in children and adolescents [33]. The reference values of serum copper concentrations in US were higher than the counterparts in Mexico and Egypt and was the same as in China. In our study, the average serum copper concentrations in participants with EBP was 125.5 ± 32.8 µg/dL and in those with normal BP was 112.3 ± 24.2 µg/dL, which were within the US reference range. We provide the first evidence of a correlation between serum copper and EBP in US youths.

Our study has several strengths. This is the first study to explore the relationship between serum copper and EBP since the standards for pediatric blood pressure were updated in 2017. In addition, we investigated the dose-response relationship between serum copper and EBP. Furthermore, we considered the simultaneous impacts of various metals in our analyses to capture the complex nature of environmental exposure as a whole.

However, this study also has some limitations. First, this is a cross-sectional study, and it is difficult for us to determine causality. Second, although a number of potential confounders were controlled, we cannot rule out the possibility of unmeasured confounding factors. Third, the complicated interactions of many trace metals with copper warrants further investigation.

Conclusions
In conclusion, our study has found a positive and non-linear association between serum copper and elevated blood pressure in US children and adolescents. We hope that this study can help policy makers develop safer reference ranges in US youths. Further prospective studies are needed to confirm our results and elucidate the mechanisms involved.

**Abbreviations**

EBP Elevated blood pressure

BP Blood pressure

NHANES National Health and Nutrition Examination Survey

AAP American Academy of Pediatrics guidelines

ICP-DRC-MS Inductively coupled plasma dynamic reaction cell mass spectrometry

Ga gallium

BMI body mass index

ROS Reactive oxygen species

**Declarations**

**Ethics approval and consent to participate**

All participants provided written informed consent (parental consent was obtained for those < 18 years) and NHANES was approved by the National Center for Health Statistics’ Ethics Review Board.

**Consent for publication**

Not applicable.

**Availability of data and materials**

Anonymized data are available from the National Center for Health Statistics (https://www.cdc.gov/nchs/nhanes/about_nhanes.htm) or from the corresponding author on reasonable request.

**Competing interests**

The authors declare that they have no competing interests.

**Author’s Contribution**
Changsong Liu collected the data and drafted the manuscript; Yanfen Liao and Zongyuan Zhu revised the manuscript critically for important intellectual content; Lili Yang and Qin Zhang analyzed and interpreted the data; Li Li were the designer of the manuscript and approved to submit the manuscript finally.

Acknowledgements

We would like to thanks for the participants who contributed their data.

References


