**Background:** Bisphenol A (BPA) is a common chemical used in the manufacture of polycarbonate plastics and epoxy resins, and > 93% of U.S. adults have detectable levels of urinary BPA. Recent animal studies have suggested that BPA exposure may have a role in several mechanisms involved in the development of cardiovascular disease (CVD), including weight gain, insulin resistance, thyroid dysfunction, endothelial dysfunction, and oxidative stress. However, few human studies have examined the association between markers of BPA exposure and CVD. Peripheral arterial disease (PAD) is a subclinical measure of atherosclerotic vascular disease and a strong independent risk factor for CVD and mortality.

**Objective:** We examined the association between urinary BPA levels and PAD in a nationally representative sample of U.S. adults.

**Methods:** We analyzed data from 745 participants in the National Health and Nutritional Examination Survey 2003–2004. We estimated associations between urinary BPA levels (in tertiles) and PAD (ankle–brachial index < 0.9, n = 65) using logistic regression models adjusted for potential confounders (age, sex, race/ethnicity, education, smoking, body mass index, diabetes mellitus, hypertension, urinary creatinine, estimated glomerular filtration rate, and serum cholesterol levels).

**Results:** We observed a significant, positive association between increasing levels of urinary BPA and PAD before and after adjusting for confounders. The multivariable-adjusted odds ratio for PAD associated with the highest versus lowest tertile of urinary BPA was 2.69 (95% confidence interval: 1.02; 7.09; p-trend = 0.01).

**Conclusions:** Urinary BPA levels were significantly associated with PAD, independent of traditional CVD risk factors.

**Keywords:** ankle–brachial index, bisphenol A, CVD, NHANES, peripheral arterial disease.

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those who had smoked > 100 cigarettes in their lifetimes were considered former smokers if they answered negatively to the question “Do you smoke now?” and current smokers if they answered affirmatively.

Rigorous procedures with quality control checks were used in blood collection, and details about these procedures are provided in the NHANES laboratory/medical technologists procedures manual (NCHS 2010d). We used the modified hexokinase method to measure serum glucose levels at the University of Missouri Diabetes Diagnostic Laboratory (Columbia, MO, USA). Diabetes mellitus was defined, based on the recent guidelines of the American Diabetes Association (2011), as a serum glucose level of > 126 mg/dL after fasting ≥ 8 hr or of > 200 mg/dL for those who fasted < 8 hr before their NHANES visit, a glycosylated hemoglobin value of > 6.5%, or a self-reported current use of oral hypoglycemic medication or insulin. Seated SBP and DBP were measured using a mercury sphygmomanometer according to the American Heart Association and Seventh Joint National Committee recommendations (Chobanian 2003), and up to three measurements were averaged. Patients were considered hypertensive if they reported current use of blood pressure–reducing medication or had SBPs of > 140 mmHg or DBPs of > 90 mmHg (NCHS 2010b).

In NHANES 2003–2004, serum creatinine measurements were conducted at the Collaborative Laboratory Services (Ottumwa, IA, USA) using the Beckman Coulter Synchro LX20 chemistry analyzer (Beckman Coulter, Fullerton, CA, USA) using the Jaffe rate method (kinetic alkaline picrate). Coefficients of variation ranged from 1.5% to 4.3% (NCHS 2010c). In a calibration sub-study, serum creatinine assays were performed on 190 stored specimens from NHANES 2003–2004 at the Cleveland Clinic laboratory (Cleveland, OH, USA) using the Roche coupled enzymatic assay that was traceable to gold-standard reference methods, including an isotope dilution mass spectrometric method for serum creatinine using standard reference methods [NIST standard reference material (SRM) 967; National Institute of Standards and Technology, Gaithersburg, MD, USA] and confirmed by analysis of CAP LN-24 (College of American Pathologists Creatinine Accuracy Calibration Verification/Linearity Survey) linearity set based on NIST-assigned values (Selvin et al. 2007). There were no significant differences in results between these two measurements, and therefore it was concluded that there was no correction necessary for serum creatinine values in NHANES 2003–2004 (Selvin et al. 2007). The glomerular filtration rate was estimated (eGFR) from serum creatinine using the four-variable Modification of Diet in Renal Disease study equation (Levey et al. 2006) as follows:

\[
eGFR = 175 \times \left( \frac{\text{serum creatinine in mg/dL}}{1.154} \right) \times (\text{age in years})^{-0.203} \times (0.742 \text{ if female}) \times (1.21 \text{ if black}).
\]

Chronic kidney disease was defined as an eGFR of < 60 mL/min/1.73 m², consistent with definitions of the National Kidney Foundation Kidney Disease Outcome Quality Initiative (KDOQI; National Kidney Foundation 2002) working group and the Kidney Disease Improving Global Outcomes (KDIGO) (Levey et al. 2006).

Previous measures of BPA in biological matrices involved techniques such as gas chromatography or high performance liquid chromatography (Ye et al. 2005). To achieve enhanced sensitivity and selectivity over previous methods, in the current NHANES, measures of environmental phenols were derivatized to alkyl or acyl derivatives before gas chromatography–mass spectrometry analysis (NCHS 2010a). The lower limit of detection for BPA concentrations was 0.36 ng/mL.

**Main outcome of interest: PAD.** For study subjects with at least one arm and weighing ≤ 400 lb, supine SBP was measured with blood pressure cuffs on the right arm, compressing the brachial artery and the two posterior tibial arteries (NCHS 2010a). For subjects 40–59 years of age, two measurements were taken at each site and averaged, and for patients ≥ 60 years of age, one measurement was taken at each site. For subjects with conditions precluding the measurement of the right arm, left brachial artery SBP was taken. Left and right ABI values were calculated as the ratio of left and right ankle SBP, respectively, to arm SBP. The smallest of the left and right ABI measurements was used. Patients with ABIs ≥ 1.5 (p = 3) are expected to have severe arterial rigidity and were excluded from the present analyses (Newman et al. 1993). For these analyses, PAD was defined as ABI < 0.9 (American Diabetes Association 2011; Selvin and Erlinger 2004). Because of the selection criteria for ABI measurements in NHANES (subjects ≥ 40 years of age), subjects who had ABI measurements taken were significantly older than the general NHANES cohort but were otherwise similar with respect to demographic factors (data not shown).

**Statistical analysis.** Urinary BPA was categorized into tertiles (< 1.4 ng/mL, 1.4–3.5 ng/mL, > 3.5 ng/mL). We hypothesized that high BPA levels were associated with PAD. Odds ratios (ORs) and 95% confidence intervals (CIs) for PAD in association with BPA were calculated by taking the lowest tertile (tertile 1) as the referent, using multivariable logistic regression models. We used two models: one adjusted only for age and sex, and a multivariable model that also adjusted for race/ethnicity (non-Hispanic whites, non-Hispanic blacks, Mexican Americans, others), education (< high school, high school, > high school), household income (< $25,000, $25,000–54,999, ≥ $55,000), smoking status (never, former, current), pack-years of smoking, alcohol intake (nondrinker, moderate drinker, heavy drinker), BMI (normal, overweight, obese), hypertension (present, absent), diabetes (present, absent), urinary creatinine (milligrams per deciliter), eGFR (milliliters per minute per 1.73 m²), and total cholesterol (milligrams per deciliter). Trends in the OR of PAD across increasing urinary BPA categories were determined by modeling BPA as an ordinal variable. We examined effect modification by performing stratified analysis by categories of sex, race/ethnicity, BMI, diabetes mellitus, and hypertension. We also conducted formal statistical tests for interaction by including multiplicative cross-product interaction terms in the multivariable logistic regression models; p < 0.1 was interpreted as a statistically significant interaction. Sample weights that accounted for the unequal probabilities of selection, oversampling, and non-response were applied for all analyses using SAS (version 9.2; SAS Institute Inc., Cary, NC, USA) and SUDAAN software (version 10; RTI International, Research Triangle Park, NC, USA); SEs were estimated using the Taylor series linearization method.

**Results.** The median (interquartile range) of urinary BPA levels was 2.30 (3.60) ng/mL; the 25th, 50th, and 75th percentile values were 1.00, 2.30, and 4.60 ng/mL, respectively. Participants with higher levels of BPA were more likely to be men, non-Hispanic blacks, and former or current smokers and had higher levels of urinary creatinine. Table 1 shows these characteristics for the study population by tertiles of urinary BPA levels. Overall, we observed a positive association between increasing BPA levels and PAD on the basis of the age- and sex-adjusted model and the multivariable-adjusted model, with significant (p < 0.05) trend tests (Table 2). This positive association persisted when BPA was analyzed as a continuous variable (1-SD increase in log-transformed BPA).

In a supplementary analysis, we examined the association between BPA levels and PAD after excluding n = 10 subjects with very high BPA values (levels > 30 ng/mL). The results were found to be essentially similar. The multivariable OR of PAD associated with 1 SD of log-transformed BPA was 1.60 (1.16–2.21). We also examined the association between increasing tertiles of BPA and self-reported CVD, and the magnitude of association was found to be weaker than that for PAD.
Compared to tertile 1 of BPA (referent), the multivariable-adjusted OR (95% CI) of self-reported CVD was 1.09 (0.18–6.60) for tertile 2, and 1.83 (1.01–3.31) for tertile 3 (p-trend = 0.02).

Discussion

In a large multiethnic, nationally representative sample, we found that increasing serum BPA levels were strongly associated with PAD. The observed association was independent of the confounding factors of smoking, BMI, alcohol intake, diabetes mellitus, hypertension, and serum cholesterol level. Our study adds to the emerging evidence suggesting a role for environmental exposure to BPA in cardiovascular disease in humans (Lang et al. 2008; Melzer et al. 2010). Furthermore, because PAD and low ABI are markers of atherosclerosis, our findings suggest that potential effects of BPA on atherosclerosis may be a mechanism underlying the previously reported association between BPA exposure and self-reported CVD (Lang et al. 2008; Melzer et al. 2010).

BPA is an environmental chemical used as a constituent monomer in polycarbonate plastics, which are used extensively in drink containers and food packaging and in the production of oxidants used in the lining of canned goods (Vandenberg et al. 2009). Exposure to BPA is believed to be mainly through dietary intake with additional exposure through water, dental sealants, inhalation of household dusts, and exposure through skin (Vandenberg et al. 2009). Recent studies from NHANES data have documented that > 93–95% of the general U.S. population has measurable concentrations of BPA metabolites in their urine (Calafat et al. 2005, 2008).

Several lines of recent evidence suggest that an association between urinary BPA levels and PAD may be biologically plausible. Animal studies have suggested that BPA exposure may have a role in CVD development through several mechanisms, including the role of BPA in weight gain and obesity development potentially through its action on adipocytes (Masuno et al. 2005; Stegeman et al. 1995) and elevations in lipids (Marmugi et al. 2012). However, there are few studies in humans for comparison.

Two of the few human studies (Lang et al. 2008; Melzer et al. 2010) reported positive associations between urinary BPA levels and self-reported CVD. Also, BPA levels were found to be associated with abnormal liver function enzymes and higher levels of fasting glucose, insulin, and HOMA-IR (homeostasis model of assessment—insulin resistance) (Lang et al. 2008; Melzer et al. 2010), all points suggesting that BPA may have a role in CVD. It is in this context that our results on objectively measured ABI to define PAD are important. In the present study, we have demonstrated a significant, positive association between higher BPA levels and PAD that persisted after adjustment for multiple confounders.

The main strengths of our study include its nationally representative sample and use of rigorous study methods to collect the data and the availability of extensive data on confounders (NCHS 2010a, 2010b). The main study limitation is that the present study is cross-sectional in nature, therefore making it impossible to confirm that exposure preceded the outcome. Another limitation is the possibility of residual confounding. For example, despite adjusting for education and

| Characteristic                              | Tertile 1 (≤ 1.4 ng/mL) | Tertile 2 (1.4–3.6 ng/mL) | Tertile 3 (> 3.6 ng/mL) |
|--------------------------------------------|-------------------------|---------------------------|-------------------------|
| Unweighted sample size                     | 253                     | 240                       | 252                     |
| Age (years)                                | 56.7 ± 0.6              | 55.6 ± 0.7                | 55.5 ± 0.9              |
| Female (%)                                 | 56.5                    | 54.8                      | 42.7                    |
| Race/ethnicity (%)                         |                         |                           |                         |
| Non-Hispanic whites                        | 76.3                    | 81.4                      | 75.0                    |
| Non-Hispanic blacks                        | 3.7                     | 9.1                       | 14.4                    |
| Mexican Americans and others               | 20.0                    | 9.5                       | 10.6                    |
| Education (%)                              |                         |                           |                         |
| < High school                              | 16.5                    | 19.7                      | 16.8                    |
| High school                                | 24.0                    | 29.5                      | 27.6                    |
| > High school                              | 59.5                    | 50.8                      | 55.6                    |
| Income (%)                                 |                         |                           |                         |
| < $25,000                                  | 20.1                    | 23.3                      | 21.2                    |
| $25,000–$49,999                            | 29.9                    | 40.1                      | 40.0                    |
| ≥ $50,000                                  | 50.0                    | 36.6                      | 38.8                    |
| Smoking status (%)                         |                         |                           |                         |
| Never                                      | 51.7                    | 52.2                      | 39.2                    |
| Former                                     | 30.8                    | 30.1                      | 37.5                    |
| Current                                    | 17.5                    | 17.7                      | 23.3                    |
| Pack-years of smoking                      | 12.8 ± 1.6              | 13.5 ± 1.0                | 17.4 ± 2.2              |
| Alcohol intake (%)                         |                         |                           |                         |
| Nondrinker                                 | 29.9                    | 41.1                      | 39.5                    |
| Moderate drinker                           | 55.1                    | 42.5                      | 40.0                    |
| Heavy drinker                              | 15.0                    | 16.4                      | 20.5                    |
| BMI (%)                                    |                         |                           |                         |
| Normal weight (< 25 kg/m²)                 | 31.5                    | 32.5                      | 23.3                    |
| Overweight (25–30 kg/m²)                   | 38.7                    | 35.0                      | 38.6                    |
| Obese (BMI ≥ 30 kg/m²)                     | 29.8                    | 32.5                      | 38.1                    |
| Diabetes (%)                               | 11.0                    | 14.4                      | 16.1                    |
| Hypertension (%)                           | 43.4                    | 45.9                      | 51.5                    |
| Urinary creatinine (mg/dL)                 | 70.2 ± 5.2              | 120.5 ± 4.3               | 161.8 ± 5.2             |
| eGFR (mL/min/1.73 m²)                      | 81.5 ± 1.3              | 80.5 ± 1.3                | 80.1 ± 1.9              |
| Total cholesterol (mg/dL)                  | 210.9 ± 2.3             | 209.6 ± 3.5               | 207.0 ± 3.3             |
| Urinary BPA (ng/mL)                        | 0.7 ± 0.02              | 2.4 ± 0.05                | 10.0 ± 1.1              |
| PAD (%)                                    | 2.8                     | 4.1                       | 9.1                     |

Data presented are mean ± SE except where indicated.

Table 2. Association between urinary BPA and PAD.

| BPA (ng/mL)       | Unweighted sample size (weighted BPA prevalence) | Age, sex-adjusted OR (95% CI) | Multivariable-adjusted OR (95% CI) |
|-------------------|--------------------------------------------------|--------------------------------|------------------------------------|
| Tertile 1 (≤ 1.4) | 253 (2.8%)                                        | 1 (referent)                   | 1 (referent)                       |
| Tertile 2 (1.4–3.6) | 240 (4.1%)                                      | 1.53 (0.39, 6.04)             | 1.10 (0.22, 5.39)                 |
| Tertile 3 (> 3.6) | 252 (9.1%)                                        | 3.73 (2.03, 6.86)             | 2.69 (1.02, 7.09)                 |
| p-Trend           | < 0.0001                                         | 0.01                           | 0.01                               |
| 1-SD increase in log-transformed BPA       | 1.57 (1.30, 1.90)                                | 1.38 (1.11, 1.72)             |                                    |

*Adjusted for age, sex, race/ethnicity, education, income, smoking status, pack-years of smoking, alcohol intake, BMI, hypertension, diabetes, urinary creatinine, eGFR, and total cholesterol. ** SD of log-transformed BPA = 1.15 ng/mL.
income, the observed association between BPA levels and PAD may be at least partly explained by residual confounding by low socioeconomic status. Future prospective studies are required to confirm or disprove our findings. A third limitation is that because ABI measures were available only in NHANES 2003–2004, we are not able to examine data from the recent NHANES cycles such as NHANES 2005–2006, where BPA levels may be lower. Fourth, the use of urinary BPA levels measured from a single spot sample is likely to result in some misclassification because of the large variability of BPA. Therefore, to provide the best approach for BPA exposure assessment, expert panels have recommended collection of multiple (rather than single) spot urine samples and study designs that consider important exposure contributors such as the time of the day of sampling (e.g., in relation to food consumption) and the time of last urination (World Health Organization/Food and Agriculture Organization of the United Nations Expert Panel 2011). Finally, urinary BPA levels may not be reflective of the free, unconjugated, circulating BPA in blood, which is considered to be the biologically active form. Future studies examining the health effects of BPA should preferably measure this free, unconjugated part of BPA in serum.

Conclusions
In a nationally representative sample of U.S. adults, higher BPA levels were positively associated with PAD after adjusting for confounding factors such as age, sex, smoking, BMI, alcohol intake, diabetes, hypertension, and cholesterol levels. Although our findings must be confirmed in other studies, they provide preliminary evidence that environmental exposure to BPA may contribute to PAD.

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