Serum Polybrominated Biphenyls (PBBs) and Polychlorinated Biphenyls (PCBs) and Thyroid Function among Michigan Adults Several Decades after the 1973–1974 PBB Contamination of Livestock Feed

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BACKGROUND: In 1973–1974, Michigan residents were exposed to polybrominated biphenyls (PBBs) through an accidental contamination of the food supply. Residents were enrolled in a registry assembled after the incident, and they and their children participated in follow-up studies to assess subsequent health outcomes.

OBJECTIVES: We evaluated associations between serum PBBs and polychlorinated biphenyls (PCBs) and markers of thyroid function among Michigan adults.

METHODS: Serum concentrations of four PBB and four PCB congeners were measured at least once in 753 adults, including 79 women who participated in a 2004–2006 study and 683 women and men with follow-up during 2012–2015. Participants completed questionnaires on health conditions (including physician-diagnosed thyroid disease), behaviors, and demographics. Thyroid hormones were measured in a subset without thyroid disease (n = 551). In multivariable linear regression models, PBB and PCB congener concentrations, on both the volume (nanogram/milliliter) and lipid (nanogram/gram lipid) basis, were assessed in relation to thyroid hormones. Logistic regression models were used to estimate associations between serum PBBs and PCBs and thyroid disease.

RESULTS: Thyroid disease was common (18% overall; 25% among women). Among women, all odds ratios (ORs) for PBB-153 and thyroid disease were positive for quintiles above the reference level, but estimates were imprecise and were without a monotonic increase. For an interquartile range (IQR) increase in PBB-153 (0.43 ng/mL), the OR (any thyroid disease) = 1.12; (95% CI: 0.83, 1.52) (n = 105 cases); for hypothyroidism, OR = 1.35 (95% CI: 0.86, 2.13) (n = 49 cases). There were 21 cases of thyroid disease in men [OR = 0.69 (95% CI: 0.33); 1.44 for an IQR increase (0.75 ng/mL) in serum PBB-153]. PCB congeners were statistically significantly associated with greater total and free thyroxine and total triiodothyronine among women and with total and free triiodothyronine among men in lipid-standardized models.

CONCLUSIONS: We found some evidence to support associations of PBBs and PCBs with thyroid disease and thyroid hormone levels. https://doi.org/10.1289/EHP1302

Introduction

Polybrominated biphenyls (PBBs) and polychlorinated biphenyls (PCBs) are classes of persistent organic pollutants that were used as flame retardants in the manufacturing of plastics and electronics (ATSDR 2004) and as coolants in electrical transformers and capacitors (ATSDR 2000), respectively. Although PBBs and PCBs have not been in production in the United States since the 1970s, exposure to these compounds is still widespread owing to their environmental and biological persistence (Xue et al. 2014). In addition, because of their structural similarities and shared toxicological features with other ubiquitous chemical flame retardants, such as polybrominated diphenyl ethers (PBDEs) and others that are currently produced, research on these compounds may shed light on the potential health effects of exposure to their newer derivatives (Birnbaum and Staskal 2004). Experimental evidence suggests that exposure to halogenated compounds such as PBBs and PCBs may interfere with the endocrine system and specifically with thyroid function (Allen-Rowlands et al. 1981; Brouwer et al. 1999; Byrne et al. 1987), but the body of literature among humans has yielded inconsistent results (as reviewed by Hagmar 2003; Salay and Garabrant 2009).

In 1973–1974, millions of people from across the state of Michigan were exposed to PBBs through an accidental contamination of the food supply (Brilliant et al. 1978; Wolff et al. 1982). The Velsicol Chemical Company shipped PBB mixtures instead of livestock feed supplement to the Farm Bureau Services feed mill; these mixtures were then distributed to farms and retail outlets throughout the state and were subsequently incorporated into cattle, chicken, pig, and sheep feed (Carter 1976; Fries 1985). Due to the common practice of eating locally raised livestock and livestock products in this area, people were exposed through direct consumption of the tainted products including meat from the animals, eggs, and dairy products (Kay 1977). In subsequent generations, people were exposed through breastfeeding and in utero transfer (Joseph et al. 2009). Directly following the episode, the Michigan Department of Health enrolled approximately 6,000 individuals in a registry to measure baseline PBB and PCB concentrations and to monitor long-term health outcomes (Landrigan et al. 1979). Persons eligible for inclusion in the registry lived on or received food from farms quarantined by the Michigan Department of Agriculture after the contamination was discovered or were Velsicol workers and members of their households. This cohort of exposed individuals and their children has now been followed for >40 y.

Evidence of possible health effects has been noted in these registry participants and in their children, specifically among...
women of reproductive age: registry members with greater PBB concentrations (compared with those with lower concentrations) were found to have increased odds of miscarriage (Small et al. 2011), shorter menstrual cycles and longer menstrual periods (Davis et al. 2005), earlier menarche in girls (Blanck et al. 2000), and later pubertal development in boys (Small et al. 2009). In conjunction with evidence from studies suggesting that women with thyroid dysfunction are at increased risk of miscarriage (Liu et al. 2014), menstrual function perturbations such as anovulation and menorrhagia (Krassas et al. 1999), and altered timing of pubertal development (Doufas and Mastorakos 2000), the results from previous studies in the Michigan cohort (Blanck et al. 2000; Davis et al. 2005; Small et al. 2009; Small et al. 2011) have motivated the hypothesis that the thyroid may be a target for PBB toxicity in humans. Animal data provide biologic support for this hypothesis, with studies consistently showing decreases in total thyroxine (T4) and triiodothyronine (T3) (Allen-Rowlands et al. 1981; Byrne et al. 1987; Gupta et al. 1983) and increases in thyroid weights (Ringer and Polin 1977; Ringer 1978; Werner and Sleigh 1981) associated with PBB exposure. However, to our knowledge, only two epidemiologic investigations of the association between PBBs and thyroid function have been published, and both involved the Michigan livestock feed contamination (Bahn et al. 1980; Yard et al. 2011). Bahn et al. (1980) reported that men who were occupationally exposed to PBBs were more likely to have prevalent hypothyroidism [high thyroid-stimulating hormone (TSH) and low total and free T4 outside of population references] as well as elevated levels of thyroid peroxidase antibodies compared with age-matched controls without occupational exposure to PBBs (4 cases/35 exposed individuals vs. 0 cases/89 controls). A subsequent study conducted among members of the original Michigan PBB Registry who were exposed through consumption of contaminated food examined associations between enrollment PBB and PCB levels and self-reported thyroid disease, which was obtained through follow-up that occurred in intervals between 1991 and 2006 (Yard et al. 2011). This nested case–control study found that among women (212 cases, 844 controls) and men (47 cases, 1,761 controls), thyroid disease was not associated with tertiles of enrollment PBB or PCB concentrations compared with those with concentrations below the limit of detection (LOD). However, these two epidemiologic studies were limited by small sample size, particularly in the case of Bahn et al. (1980); in the Yard et al. (2011) study, limitations included early exposure assessment methodologies that were used to determine baseline PBB and PCB concentrations at the time of the contamination incident and a lack of other biomarker measurements including thyroid hormones.

In the present study, we investigated the association between PBB and PCB exposure and thyroid function among individuals in the Michigan PBB Registry with the addition of key biologic measures assessed using more sensitive methods than those used in previous investigations, including current serum PBB, PCB, and lipid concentrations; circulating thyroid hormone concentrations; and in a subset, urinary iodine concentrations. The purpose of this study was to examine the association between serum PBB and PCB concentrations and measures of thyroid function in adults from the Michigan cohort, including investigation of overt thyroid disease, thyroid hormone concentrations, and in a subset of women, whether these associations were stronger among the iodine deficient.

**Methods**

**Data Collection**

The original PBB registry consisted of a heterogeneous group: residents of farms that were classified as quarantined by the Michigan Department of Health based on PBB levels in any sample of meat, eggs, or milk that were above a specified limit; those who had received meat, eggs, or dairy products directly from quarantined premises in 1973 or 1974; workers who had been occupationally exposed to PBBs in a chemical manufacturing plant in Michigan and members of their households; residents on farms with low-level PBB contamination; “self-referred” individuals who had either resided on farms identified as being contaminated by PBBs at levels below quarantine limits or who had eaten food produced on such farms; and “self-referred” volunteers who had no documented connection with contaminated farm premises but likely consumed contaminated farm products that were distributed throughout the state (Landrigan et al. 1979). Offspring of women in the original registry were subsequently enrolled (oral communication, H. Humphrey, environmental epidemiologist, Michigan Department of Public Health, November 1995).

Participants were recruited for this study from the original PBB Registry in two phases (see Figure S1). In the first phase (2004–2006), the Michigan Department of Health recruited women from the PBB registry (from all subgroups) for a blood draw, targeting women who were particularly highly exposed to PBB based on their enrollment levels. These women also completed a computer-assisted telephone interview close to the time of the blood draw that included questions about medical history (particularly with regard to endocrine-sensitive end points, such as thyroid disease), current medication use, demographics, and other lifestyle information. The second phase started in 2012: Members from all subgroups of the original PBB Registry were mailed invitations to attend community meetings held throughout the state, where study recruitment was conducted between 2012 and 2015. Community meetings were also advertised in the local press; thus, individuals from the original registry as well as others attended and subsequently participated in the study. Eligibility criteria to participate in this phase of recruitment of the study were at least one of the following: individuals who lived in the state of Michigan during the time of the contamination (1973–1974) or who were offspring of those who lived in the state during that time. Participation in the study entailed a blood draw that was used to determine PBB and PCB concentrations; completion of online health questionnaires, which included details on medical history (including reporting of physician-diagnosed thyroid disease and age at diagnosis), current medication use, behaviors, and demographics; and anthropometric measurements (height and weight) conducted by study staff, from which body mass index (BMI) was derived. BMI was missing for 34% (n = 255) because a substantial proportion of individuals did not participate in the physical measurements.

Directly following their overall study participation (blood draw and questionnaire), subsets of women from Phase 1 and Phase 2 were separately recruited to participate in a study measuring menstrual cycle function. Those who were premenopausal, who were not pregnant or lactating, who were not taking hormonal medications, and who were never diagnosed with a reproductive-area cancer or received cancer treatment were eligible. Women collected daily first-morning-urine samples over three menstrual cycles, and the samples were analyzed to obtain iodine concentrations.

A small proportion of individuals (6%) participated in the study more than once (n = 36, 2 times; n = 6, 3 times) over the course of the two recruitment phases by attending multiple recruitment events; thus, their blood was drawn for exposure assessment each time. Repeat participants consisted of the following: those who participated between 2004 and 2006 and again between 2012 and 2015 (n = 9; four of these participated in 2004–2006 and twice between 2012 and 2015), and those who
Thyroid Disease

Thyroid disease status and age at diagnosis were derived from self-report of physician-diagnosed disease on the questionnaire (n = 128), and we additionally classified individuals who had TSH and total or free T4 outside of clinical reference ranges (TSH = 0.3–5.0 μU/mL; total T4 = 5–12 μg/dL; free T4 = 0.7–2.0 ng/dL) as having thyroid disease (n = 7) (Braverman and Cooper 2012). Two women who participated in both Phase 1 and Phase 2 developed thyroid disease between these time points and were included as noncases and as cases in the analysis, respectively. Thyroid disease included any thyroid disorder including hypothyroidism; hyperthyroidism; other diagnoses including Hashimoto’s thyroiditis, Graves’ disease, and thyroid nodules; and instances where the participant did not specify the type. Among women, analyses were repeated restricting the outcome to hypothyroidism.

Exposure Analyses

Blood samples were collected via venipuncture, processed to isolate the serum, and stored at -20°C until analysis. In batches between March 2013 and June 2015, serum samples were analyzed for target PBB (77, 101, 153, and 180) and PCB (118, 138, 153, 180) congeners using gas chromatography–tandem mass spectrometry (GC-MS/MS; Agilent Technologies; Agilent 7890A gas chromatograph coupled to an Agilent 7000B tandem mass spectrometer) at the Laboratory for Exposure Assessment and Method Development in Environmental Research (LEADER) at Emory University’s Rollins School of Public Health. Detailed laboratory methods have been described previously (Marder et al. 2016). Briefly, samples were subjected to liquid-liquid extraction followed by solid-phase extraction before instrumental analysis using GC-MS/MS in multiple reaction monitoring mode. Quantification was performed using isotope-dilution calibration covering a concentration range of 0.005–12.5 ng/mL. LODs, which were extrapolated from standard injections, were defined as the lowest standard concentration at which the signal:noise ratio was >3. Estimated detection limits for all target compounds ranged from 0.001–0.006 ng/mL. PCB-118 concentrations could not be reported for 99 samples (12.4%), including all samples collected in Phase 1, due to unstable retention times on the laboratory instrument, which was caused by clipping of the peak such that it was outside the acquisition window.

For analyses, PCB congeners were considered individually as well as summed by the degree of ortho-substitution (Safe 1993) based on the hypothesis that chemical structure is a main determinant of PCB-induced thyroid disruption. The di-ortho chlorinated congeners (PCB-138, 153, and 180) were summed (∑PCB), and PCB-118, as the only mono-ortho chlorinated congener measured, was considered separately. PBB-153 was the only PBB congener detected in >10% of the samples; therefore, no groupings of PBB congeners were considered.

Lipid Analyses

Total serum triglyceride content was measured using an Abnova Triglyceride Quantification Assay Kit (Abnova Corporation), and total cholesterol content was measured using a Cayman Cholesterol Assay Kit (Cayman Chemical Company) according to the manufacturers’ instructions. Total lipids were calculated using conventional methods based on these individual lipid components (Phillips et al. 1989).

Hormone Analyses

Participants without a history of thyroid disease at the time of blood draw and who reported no current thyroid medication use were tested for thyroid hormones. Thyroid hormones (TSH, total and free T4, and total and free T3) were analyzed at the Emory Clinical Translational Research Laboratory with a Beckman Coulter Access II chemiluminescent immunoassay analyzer (Beckman Coulter). Analyses were conducted as directed by the manufacturer. Daily quality control (QC) samples were run before and after all study samples, and blinded controls were run throughout. If QC samples were more than two standard deviations from the expected values, the assay was repeated.

Urinary Iodine Analyses

A subset of women (n = 28 from Phase 1 and n = 52 from Phase 2) also participated in a study examining menstrual cycle function, which involved daily collection of urine samples for three months directly following their blood draw for this study. Women were eligible for this study if they were premenopausal, were not pregnant or lactating, were not currently taking hormonal medication, and had never had a reproductive cancer or treatment for any cancer. Because it has been shown that 10 to 14 urinary iodine measurements are needed to estimate individual iodine status with 20% precision owing to the high degree of within-person variability in spot urine samples (Andersen et al. 2008; König et al. 2011), we pooled equal volumes from 14 consecutive urine samples for each woman and measured iodine and creatinine concentrations in the pooled sample. The samples selected for pooling were those that were collected closest in time to the blood draw to approximate iodine status at the time thyroid hormones were measured. One woman participated twice such that she had two iodine concentrations (Phase 1 in 2004 and Phase 2 in 2013).

Iodine was determined in diluted samples by inductively coupled plasma–mass spectrometry (ICP-MS) and incorporated an octopole collision cell to reduce interference from polyatomic species. Quantitation was based on the use of internal standards, and a calibration curve was generated using a matched synthetic matrix. Urinary creatinine was measured using a colorimetric method based on the modified Jaffé reaction (Taussky 1954). Creatinine levels were used to adjust urinary iodine concentrations for urine flow rate. The lower limits of quantification were 100 μg creatinine/L and 20 μg iodine/L (NMS Labs).

Statistical Analyses

The distribution of exposure was explored, and geometric means (GMs) were calculated for PBB-153, for each PCB congener (PCB-118, 138, 153, and 180), and for ∑PCB (di-ortho) by thyroid disease status and for each covariate stratum. To estimate the association between serum PBBs and PCBs and thyroid disease prevalence, we fit logistic regression models with PBBs and PCBs considered as natural log–transformed continuous variables (to reduce the influence of extreme outliers), and we report associations for interquartile range (IQR) increases in ln-transformed concentrations as well as for quintiles. IQRs and quintiles were calculated separately for women and men. Models were adjusted for the following covariates that were identified a priori based on the literature and on our conceptualized directed acyclic graph (see Figure S2): lipids, age (18–29, 30–39, 40–49, 50–59, 60–69), and several other covariates as specified in the Methods section.
≥60 y), study phase (2004–06; 2012–15), and smoking status (current, former, never). Those who reported thyroid disease before the contamination episode were excluded from analyses with PBB-153 because these cases were likely due to other factors; however, they were included in all PCB analyses. In addition, all analyses were stratified by sex owing to the different PCB and PBB exposure distributions and to known differences in thyroid physiology between women and men as well as to greater baseline risks of thyroid dysfunction among women compared with men (Adlersberg and Burrow 2002). We did not control for BMI because BMI may be a consequence of overt thyroid dysfunction (Dale et al. 2001; Tzotzas et al. 2000): Hypothyroid and hyperthyroid patients often present with weight gain or loss, respectively (Fox et al. 2008). In models for hypothyroidism (among women only), those missing thyroid disease type were excluded from the model, and those with other specified disease types (e.g., hyperthyroidism) were considered to be noncases.

Among adults with no history of physician-diagnosed thyroid disease or current thyroid medication use, we fit multivariable linear regression models to estimate the associations between serum PBB and PCB concentrations and thyroid hormones. We fit separate models for each thyroid hormone and for each PBB and PCB congener with a detection frequency >90% (PBB-153 and PCB-118, 138, 153, and 180) as well as for ∑PCB (di-ortho), both as natural log-transformed continuous variables and as deciles. Deciles were chosen a priori because for PBB-153 concentrations, the 10th percentile represents background levels in the general U.S. population per NHANES (GM = 0.01 ng/mL) (CDC 2004). TSH was natural log-transformed for normalization because there is evidence to suggest that TSH concentrations change on the logarithmic scale as opposed to the additive scale and are lognormally distributed (Demers and Spencer 2003). All regression models with hormone outcomes were controlled for lipids, age (18–29, 30–39, 40–49, 50–59, ≥60 y), current smoking status (yes, no), and time of blood collection (morning, midday, afternoon/evening). Study phase was not controlled for because it was not associated with the outcome. In the case of missing covariate data, all analyses were conducted using complete case analysis. Associations were evaluated by statistical significance at p < 0.05.

In analyses of PBB-153, measurements below the LOD were imputed using a distribution-based maximum likelihood technique (Chen et al. 2011; Lubin et al. 2004) whose parameters (μ, σ²) were estimated assuming a lognormal probability distribution and based on the proportion of measures below the LOD. To incorporate uncertainty in the maximum likelihood estimates, we generated 10 sets of distribution parameters and imputed values below the LOD based on each set of parameters, creating 10 complete data sets (Chen et al. 2011; Lyles et al. 2001). Finally, we used the SAS procedure PROC MIANALYZE to generate summary regression coefficients, standard errors, and 95% confidence intervals (CIs) (Little and Rubin 2002).

Owing to the small proportion of repeat participants, in all analyses (for thyroid hormones as well as for thyroid disease), we used generalized estimating equations with robust standard errors to account for the potential correlated nature of their outcomes. Lastly, in all our analyses, we expressed serum PBBS and PCBs on the wet weight (volume) basis (nanograms/milliliter serum) and controlled for lipids as a covariate based on the potential for serum lipids to act as a confounder because of its association with time of day (likely due to recently eaten meals), which is also associated with thyroid hormones, specifically TSH (see Figure S3); as well as increased statistical flexibility by expressing it as a separate term instead of within the denominator of the PCB or PBB concentration itself (Schisterman et al. 2005). However, we repeated all analyses with PBBs and PCBs expressed on the lipid basis (nanograms/gram lipid).

Finally, among the subset for whom iodine was measured (n = 79 individuals, 80 observations), we explored the potential for effect measure modification of the relationship between PBBs and PCBs and thyroid hormones by iodine status by fitting models stratified by iodine status. p-Values were obtained by fitting linear regression models with a cross-product term and assessing the p-value of the interaction term at α = 0.05. Because analysis using the cut-off for deficient iodine (<100 μg/g creatinine (WHO 1994)) was limited by sample size (n = 14 iodine deficient), we used the alternative of <150 μg/g creatinine, which is considered low iodine (Caldwell et al. 2011).

All statistical analyses were performed using SAS (version 9.4; SAS Institute Inc.).

Results

Population Characteristics

A total of 778 adults had their blood drawn and concurrently completed a health questionnaire during either 2004–2006 or 2012–2015 (or both in some cases). Among them, 3% (n = 26) were missing information on thyroid disease diagnoses and were subsequently excluded, leaving 753 individuals for this analysis (801 samples). This study sample consisted of 79 individuals (79 samples) from Phase 1 (all women), 683 individuals (722 samples) from Phase 2 (385 women and 298 men); 9 individuals (all women) participated in both phases including 5 women who participated twice (once in 2004–2006 and once again in 2012–2015) and 4 women who participated three times (once in 2004–06 and twice between 2012 and 2015). Among those who participated between 2012 and 2015 only, 21 and 1 women participated 2 and 3 times, respectively, and 10 and 1 men participated 2 and 3 times, respectively.

The study population was >99% white and had a high prevalence of overweight and obesity (69%) and ever smoking (40%) (Table 1). Most of the samples were collected between 2012 and 2015 (91%) and were from women (all samples from 2004–2006 and 60% overall). The majority (72%) were initially exposed to PBBs directly by consuming contaminated food (42% before the age of 16 y), with the remaining 28% likely exposed in utero, through breastfeeding, or both, because they were born after the acute contamination episode (after May 1974).
Table 1. Population characteristics and serum polybrominated biphenyl (PBB) and polychlorinated biphenyl (PCB) concentrations by thyroid disease status among Michigan adults 2004–2015.

| Characteristic | Total study sample | Thyroid disease | No thyroid disease | p-Value<sup>b</sup> |
|----------------|-------------------|-----------------|-------------------|------------------|
| Exposures [GM; ng/mL (ng/g lipid)] | | | | |
| PBB-153<sup>c</sup> | 0.17 (26.3) | 0.22 (33.2) | 0.16 (25.0) | 0.07 (0.09) |
| PCB-118 | 0.05 (7.9) | 0.08 (11.5) | 0.05 (7.3) | <0.01 (<0.01) |
| PCB-138 | 0.17 (25.8) | 0.22 (33.3) | 0.16 (24.4) | <0.01 (<0.01) |
| PCB-153 | 0.19 (29.0) | 0.24 (36.8) | 0.18 (27.5) | <0.01 (<0.01) |
| PCB-180 | 0.14 (20.8) | 0.18 (26.5) | 0.13 (19.7) | <0.01 (<0.01) |
| ΣPCB (di-ortho)<sup>d</sup> | 0.51 (76.9) | 0.65 (97.7) | 0.48 (73.0) | <0.01 (<0.01) |
| Sex [n (%)] | | | | |
| Female | 455 (60.4)<sup>e</sup> | 113 (24.8) | 344 (75.6) | <0.01 |
| Male | 298 (39.6)<sup>e</sup> | 22 (7.4) | 276 (92.6) | |
| Age (y) at blood draw [n (%)]<sup>f</sup> | | | | |
| 18–29 | 100 (13.3) | 3 (3.0) | 97 (97.0) | <0.01 |
| 30–39 | 176 (23.4) | 19 (10.8) | 157 (89.2) | |
| 40–49 | 153 (20.3) | 32 (20.9) | 121 (79.1) | |
| 50–59 | 150 (19.9) | 33 (22.0) | 117 (78.0) | |
| ≥60 | 186 (24.7) | 49 (26.3) | 137 (73.7) | |
| Age (y) at PBB contamination [n (%)] | | | | |
| ≥16 at contamination | 226 (30.0) | 58 (25.7) | 168 (74.3) | <0.01 |
| 0–15 at contamination | 315 (41.8) | 61 (19.4) | 256 (81.3) | |
| Born after contamination | 212 (28.2) | 16 (7.5) | 196 (92.5) | |
| Study years [n (%)] | | | | |
| 2004–2006 only<sup>g</sup> | 70 (9.3) | 4 (5.7) | 66 (94.3) | <0.01 |
| 2012–2015 only<sup>g</sup> | 674 (89.5) | 128 (19.0) | 546 (81.0) | |
| 2004–2006 and 2012–2015<sup>g</sup> | 9 (1.2) | 3 (33.3) | 8 (88.9) | |
| Race/ethnicity [%] | | | | |
| Non-Hispanic white | 712 (96.1) | 128 (18.0) | 586 (82.3) | 0.92 |
| Hispanic white | 22 (3.0) | 3 (13.6) | 19 (86.4) | |
| Other | 7 (0.9) | 1 (14.3) | 6 (85.7) | |
| Missing | 12 | 3 | 9 | |
| Body mass index<sup>h</sup> [n (%)] | | | | |
| ≤Normal | 155 (31.1) | 16 (10.3) | 140 (90.3) | 0.02 |
| Overweight | 156 (31.3) | 20 (12.8) | 137 (87.2) | |
| Obese | 190 (38.2) | 42 (22.1) | 148 (77.9) | |
| Missing | 255 | 57 | 198 | |
| Smoking status [%] | | | | |
| Current smokers | 120 (16.1) | 14 (11.7) | 106 (88.3) | 0.11 |
| Former smokers | 182 (24.4) | 35 (19.2) | 147 (80.8) | |
| Never smokers | 444 (59.5) | 86 (19.4) | 360 (81.1) | |
| Missing | 7 | 0 | 7 | |

Note: GM, Geometric mean. Total study sample: n = 753 individuals, n = 801 samples.
<sup>a</sup>Includes those who specified physician-diagnosed thyroid disease [n = 128 (107 women; 21 men)] and those who had thyroid hormone concentrations outside clinically normal ranges [n = 113 (6 women; 1 man)]; includes 8 women and 1 man who reported thyroid disease before PBB contamination; includes 13 (9.1%) hyperthyroid (13 individuals), 66 (46.2%) hypothyroid (60 individuals), 17 (11.9%) other types (15 individuals), and 47 (32.9%) not specified (47 individuals).
<sup>b</sup>Includes those born before 1973, 77% were still above the LOD (compared with 99% and 100% of those ages 0–15 y and ≥16 years at the contamination, respectively). Men were more highly exposed to PBBs and to PCB-138, 153, and 180 than women, and this was particularly true for PBB-153 (GM = 0.24 ng/mL vs. 0.14 ng/mL). There was no consistent pattern between PBB-153 and PCB congener concentrations and BMI categories.

Based on laboratory parameters. Those with thyroid disease had higher GM serum PBB-153 and PCB-118, 138, 153, and 180 concentrations (wet weight and nanograms/gram lipid) than those without thyroid disease.

Geometric mean serum PBB and PCB concentrations increased monotonically with age at the time of blood draw in the cohort as a whole (Table 3; see also Table S1) and in samples from those without thyroid disease who were tested for thyroid hormones (see Table S2). This correlation between age and exposure was also clear when comparing GM serum PBB-153 and PCB congener levels across strata of age at PBB contamination. However, although those born after the contamination incident had lower PBB-153 concentrations than those born before 1973, 77% were still above the LOD (compared with 99% and 100% of those ages 0–15 y and ≥16 years at the contamination, respectively). Men were more highly exposed to PBBs and to PCB-138, 153, and 180 than women, and this was particularly true for PBB-153 (GM = 0.24 ng/mL vs. 0.14 ng/mL). There was no consistent pattern between PBB-153 and PCB congener concentrations and BMI categories.

**Associations between PBBs/PCBs and Thyroid Disease**

Among those with thyroid disease (n = 135), the following types were specified [or were apparent based on thyroid assays...
were all >1, although all estimates were imprecise (Table 4). The
in PBB-153 (Table 4), and in lipid-standardized models,
signiﬁcantly related to thyroid disease based on thyroid hormone levels, the mean total and free
T₃ decreased with age, and GM TSH concentrations were higher
among those who were iodine deﬁcient (<100 µg/g creatinine).
Men were much less likely to report being diagnosed with
thyroid disease (n = 22, 7.4%) including 1 case diagnosed before the PBB contamination incident and 1 case classiﬁed
based on thyroid hormone levels versus a self-reported diagno-
sis. Estimated aORs for thyroid disease in association with IQR increases in wet-weight serum PBB-153 and PCB concentrations were imprecise, with inverse associations estimated for
PBB-153 [for an IQR increase (0.75 ng/mL), aOR = 0.69 (95% CI:
0.3, 1.4)] and ∑PCB (di-ortho) [for an IQR increase (0.81 ng/mL),
aOR=0.93 (95% CI: 0.4, 2.1)], and a positive association with PCB-118 [for an IQR increase (0.06 ng/mL), aOR=1.50 (95% CI: 0.9, 2.5)] (Table 5).

Thyroid Hormones
Among 551 participants who did not report thyroid disease at the
time of their blood draw and were not classiﬁed as having thyroid
disease based on thyroid hormone levels, the mean total and free
T₃ decreased with age, and GM TSH concentrations were higher
in samples collected in the morning versus afternoon or evening
(see Table S6). In addition, among those for whom urinary iodine was measured, TSH and total T₃ concentrations were greater among those who were iodine deﬁcient (<100 µg/g creatinine), although differences were minimal across strata of low iodine
(<150 µg/g creatinine). Lastly, among those who reported hormonal medication use and the few who were

(n = 7): 13 (9.6%) hyperthyroid (10 women, 3 men), 60 (44.4%)
hypothyroid (52 women, 8 men), 15 (11.1%) other types (14
women, 1 man), and 47 (34.8%) not speciﬁed (37 women, 10 men).
The average age at thyroid disease diagnosis was 35 (median = 33, range = 10–74) for women and 42 (median = 40, range = 14–77) for men. Among women, estimated adjusted odds ratios (aORs) for thyroid disease in association with IQR increases in ln-transformed serum PBB-153, PCB-118, and ∑PCB (di-ortho) concentrations were above the null [e.g., OR = 1.1 (95% CI: 0.77, 1.68) (Table 4)]. When exposures were
compared with the first (8 cases); OR = 2.19 (95% CI: 0.67, 7.23)
for the ﬁfth quintile (11 cases)] (Table 4). ORs for hypothyroid-
ism in association with lipid-adjusted PBB-153 (nanograms/gram
lipid) were consistent with those for wet-weight concentrations
(seetable S3).

Men were much less likely to report being diagnosed with
thyroid disease (n = 22, 7.4%) including 1 case diagnosed before the PBB contamination incident and 1 case classiﬁed
based on thyroid hormone levels versus a self-reported diagno-
sis. Estimated aORs for thyroid disease in association with IQR increases in wet-weight serum PBB-153 and PCB concentrations were imprecise, with inverse associations estimated for
PBB-153 [for an IQR increase (0.75 ng/mL), aOR = 0.69 (95% CI:
0.3, 1.4)] and ∑PCB (di-ortho) [for an IQR increase (0.81 ng/mL),
aOR=0.93 (95% CI: 0.4, 2.1)], and a positive association with PCB-118 [for an IQR increase (0.06 ng/mL), aOR=1.50 (95% CI: 0.9, 2.5)] (Table 5).

Thyroid Hormones
Among 551 participants who did not report thyroid disease at the
time of their blood draw and were not classiﬁed as having thyroid
disease based on thyroid hormone levels, the mean total and free
T₃ decreased with age, and GM TSH concentrations were higher
in samples collected in the morning versus afternoon or evening
(see Table S6). In addition, among those for whom urinary iodine was measured, TSH and total T₃ concentrations were greater among those who were iodine deﬁcient (<100 µg/g creatinine), although differences were minimal across strata of low iodine
(<150 µg/g creatinine). Lastly, among those who reported hormonal medication use and the few who were

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Table 3. Serum polybrominated biphenyl (PBB) and polychlorinated biphenyl (PCB) (nanograms/milliliter serum and nanograms/gram lipids) concentrations by population characteristics in Michigan adults, 2004–2015.

| Characteristics                            | PBB-153 (GM) *ng/mL (ng/g lipid) | PCB-118 * (GM) ng/mL (ng/g lipid) | Σ PCB (di-ortho) * (GM) ng/mL (ng/g lipid) |
|-------------------------------------------|----------------------------------|----------------------------------|------------------------------------------|
| Sex                                       | Male: 0.24 (38.0)                | Female: 0.14* (21.0)*            | Male: 0.05 (7.8)                         | Female: 0.47* (68.6)*                      |
| Age (y) at blood draw                     | 18–29: 0.01* (1.9)*              | 30–39: 0.08 (13.3)               | 40–49: 0.25 (35.6)                        | 50–59: 0.44 (62.1)                         | ≥60: 0.53 (81.9) |
|                                          |                                  |                                  |                                          |                                        | 0.09 (14.2)                                                                 |
|                                          |                                  |                                  |                                          |                                        | 1.03 (157.2)                                                                 |
| Smoking status                            | Current smokers: 0.09* (13.6)*   | Former smokers: 0.28 (42.8)      | Never smokers: 0.17 (26.4)               | 0.04* (5.6)*                             | 0.43* (66.1)* |
|                                          |                                  |                                  |                                          |                                        | 0.64 (97.4)                                                                 |
|                                          |                                  |                                  |                                          |                                        | 0.48 (73.1)                                                                 |

Note: Data not applicable; GM, Geometric mean. Total: n = 753 individuals, n = 801 samples. [See Table S1 for information on individual PCB congeners (PCB-138, 153, and 180).]

*p < 0.01; **p < 0.05; ***p < 0.10; p-values computed using Student’s t-tests or analysis of variance (ANOVA) using natural log-transformed PBB and PCB levels for comparison between categories.

*GMs for PBB-153 computed after multiple imputation for observations <LOD.

PCB-118 not reported for 32 (2.4%) of samples owing to unstable retention time on laboratory instrument; includes all samples from 2004–2006 (all women; n = 79) and 20 samples from 2012–2015 (10 women, 10 men).

PCB-138 + PCB-153 + PCB-180.

*n = 36 individuals participated twice, and n = 6 individuals participated three times over time; thus, cell counts sum to >753 and percents sum to >100%.

*n = 255 missing BMI, n = 7 missing smoking status.

*Assessed at time of blood draw.

Table 4. Adjusted odds ratios (aORs) and 95% confidence intervals (95% CI) for the association between polybrominated biphenyl (PBB)-153, polychlorinated biphenyl (PCB)-118 and ΣPCB (nanograms/milliliter) and thyroid disease among Michigan females, 2004–2015.

| Exposure | Total | n (%) | aOR (95% CI) | n (%) | aOR (95% CI) |
|----------|-------|-------|--------------|-------|--------------|
| PBB-153  |       |       |              |       |              |
| PCB-118  |       |       |              |       |              |
| ΣPCB (di-ortho)  |       |       |              |       |              |

Note: IQR, interquartile range; LOD, limit of detection; Q1–Q5, quintiles 1 through 5. Models for PBB-153 and ΣPCB (di-ortho) control for lipids, age, study phase, and smoking status; models for PCB-118 do not control for study phase because all samples from Phase 1 were not reportable owing to instrumentation error [see Table S4 for information on individual PCB congeners (PCB-138, 153, and 180).]

Includes those who specified physician-diagnosed thyroid disease (n = 107) and those who had thyroid hormone concentrations outside clinically normal ranges (n = 6); includes 10 hyperthyroid, 52 hypothyroid, 14 other types, and 37 not specified.

PCB-153 analyses exclude 8 women who were diagnosed with thyroid disease before the contamination event; PCB-118 analyses exclude 88 women (89 samples) who had missing values owing to unstable retention time on laboratory instrument; ΣPCB (di-ortho) analyses include all observations.

*Includes 30 women participated multiple times, so cell counts may not sum to total.

*Hypothyroidism analyses exclude 37 observations owing to lack of information on thyroid disease type (32 in PBB-153 model); these are excluded from denominator of percent calculations.

All IQRs and quintiles calculated among women only.

PCB-153 concentrations below the LOD were imputed using a distribution-based multiple imputation approach.

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Among women, the positive associations between PCBs and each thyroid hormone (estimated for total and free T4 and T3 concentrations) were generally stronger among women. The association with free T4 was of particular interest; it was imprecise. Serum levels of PCBs but not of PBB-153 were associated with greater total and free T3, although estimated differences were larger and were only statistically significant on the lipid basis (Table 6; see also Table S7) [i.e., a natural log–unit increase in ΣPCB (di-ortho) was associated with 0.16 μg/dL greater total T3 (95% CI: −0.03, 0.35), 0.03 ng/dL greater free T3 (95% CI: 0.01, 0.04), and 3.97 ng/dL greater total T3 (95% CI: 0.53, 7.40)]. The estimates for PCB-118 were greater than those for the other measured PCB congeners in relation to differences in total T3 and T4 concentrations [a natural log–unit increase in PCB-118 was associated with 0.23 μg/dL (95% CI: 0.04, 0.43) greater total T4 and 4.94 ng/dL (95% CI: 2.05, 7.83) greater total T3]. Among men, PCB congeners were associated with greater total and free T3, although estimated differences were larger and were only statistically significant on the lipid basis (Table 6; see also Table S7). Overall, all estimates corresponded to very small relative differences (1–5%) when considered in conjunction with the distribution of circulating levels of each thyroid hormone (estimated for total and free T4 and T3 from Table S8 by dividing each beta coefficient by its corresponding median thyroid hormone value).

When PBB-153 and PCBs were expressed as deciles, the results were largely consistent with the results of the continuous natural log–transformed analyses (Figure 1; see also Tables S9–S12). Among women, the positive associations between PCBs and total and free T3 largely remained; again, estimates were largest for PCB-118 (see Tables S9 and S11). For PCB-118 and total T4 among women, estimates increased from the first to the second decile and subsequently leveled off without further increases in the upper deciles, whereas estimates for total T3 continued to increase throughout the upper deciles. However, among men, there was some evidence for a positive association between PBB-153 and total T4 concentrations, which was not observed in analyses with PBB-153 expressed as a natural log–transformed continuous variable, although estimates for the highest deciles were close to the null (see Table S10).

### Associations between PBBs/PCBs and Thyroid Hormones

In linear regression models with the exposures expressed as natural log–transformed continuous variables, PCB congeners (and other compounds included in the analyses) were associated with modest differences in various hormones (Table 6). Among women, PCBs (modeled on either the volume or the lipid basis) were associated with higher total and free T4 and total T3 concentrations (Table 6; see also Table S8) [i.e., a natural log–unit increase in ΣPCB (di-ortho) was associated with 0.16 μg/dL greater total T4 (95% CI: −0.03, 0.35), 0.03 ng/dL greater free T4 (95% CI: 0.01, 0.04), and 3.97 ng/dL greater total T4 (95% CI: 0.53, 7.40)]. The estimates for PCB-118 were greater than those for the other measured PCB congeners in relation to differences in total T3 and T4 concentrations [a natural log–unit increase in PCB-118 was associated with 0.23 μg/dL (95% CI: 0.04, 0.43) greater total T4 and 4.94 ng/dL (95% CI: 2.05, 7.83) greater total T3]. Among men, PCB congeners were associated with greater total and free T3, although estimated differences were larger and were only statistically significant on the lipid basis (Table 6; see also Table S7). Overall, all estimates corresponded to very small relative differences (1–5%) when considered in conjunction with the distribution of circulating levels of each thyroid hormone (estimated for total and free T4 and T3 from Table S8 by dividing each beta coefficient by its corresponding median thyroid hormone value).

When PBB-153 and PCBs were expressed as deciles, the results were largely consistent with the results of the continuous natural log–transformed analyses (Figure 1; see also Tables S9–S12). Among women, the positive associations between PCBs and total and free T3 largely remained; again, estimates were largest for PCB-118 (see Tables S9 and S11). For PCB-118 and total T4 among women, estimates increased from the first to the second decile and subsequently leveled off without further increases in the upper deciles, whereas estimates for total T3 continued to increase throughout the upper deciles. However, among men, there was some evidence for a positive association between PBB-153 and total T4 concentrations, which was not observed in analyses with PBB-153 expressed as a natural log–transformed continuous variable, although estimates for the highest deciles were close to the null (see Table S10).

### Associations between PBBs/PCBs and Thyroid Hormones Stratified by Iodine Status

Finally, in the women with urinary iodine measurements (n = 80), 18% (n = 14) were iodine deficient (<100 μg/g creatinine), and 54% (n = 43) had low iodine (<150 μg/g creatinine) (see Table S4). Heterogeneity in the estimates was detected based on urinary iodine concentrations <150 μg/g creatinine (Table 7). ΣPCB (di-ortho) was positively associated with TSH concentrations among 43 women classified as having low urinary iodine (<150 μg/g creatinine): a 24.6% increase in TSH associated with a natural log–unit increase in ΣPCB (95% CI: 3.0, 49.2%); in contrast, among the 37 women classified as having sufficient iodine (≥150 μg/g creatinine), ΣPCB was inversely associated with TSH [−22.9% (95% CI: −39.3%, −1.0%) for a natural log–unit increase in ΣPCB; interaction p-value <0.01)]. The pattern of associations was similar for PCB-118, but stratum-specific estimates were closer to the null and were not significantly different from each other (interaction p = 0.32).

### Discussion

In this study of 753 adults from a cohort historically exposed to PBBs, thyroid disease was prevalent (18.0%), particularly among women (24.8%). Among women, there was some evidence of a positive association between serum PBB-153 concentrations and thyroid disease, specifically hypothyroidism, although the association did not increase with increasing exposure, and estimates were imprecise. Serum levels of PCBs but not of PBB-153 were associated with subtle differences in thyroid hormone concentrations, which were generally stronger among women.

Compared with NHANES data from 2003–2004, this cohort had substantially higher exposure to PBB-153, with 59% having levels above the national 95th percentile (0.19 ng/mL; 27.2 ng/g lipid) (CDC 2017). In contrast, the distribution of PCB exposure was more similar to national levels (median PCB-153 and PCB-180 = 20.8 and 18.0 ng/g lipid, respectively) (CDC 2017). Theoretically, this observation could be consistent with the observed burden of thyroid disease among women in this population (24.8%), which was somewhat greater than expected (18.2%).
Table 6, β-coefficients with 95% confidence intervals (95% CI) from regression models for associations of ln-transformed polybrominated biphenyl (PBB) and polychlorinated biphenyl (PCB) congeners (nanograms/milliliter) with thyroid hormones among Michigan adults without thyroid disease (N=551 individuals, 571 samples), 2004–2015. 

| Exposure          | Women | Men |
|-------------------|-------|-----|
| Total T4 (μg/dL)  | β (95% CI) | β (95% CI) |
| PBB-118           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PBB-138           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PCB-153           | 0.02 (−0.02, 0.06)  | −0.02 (−0.06, 0.02) |
| PCB-180           | 0.03 (−0.03, 0.09)  | −0.01 (−0.08, 0.06) |
| Note: T4 models fit among 342 women (362 samples) and among 208 men (208 samples). All models controlled for lipids, age at blood draw (indicator variables for each decade), time of blood collection (indicator variables for morning, midday, afternoon, and midnight), and region (indicators for 6 regions). | | |
| Free T4 (μg/dL)   | β (95% CI) | β (95% CI) |
| PBB-118           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PBB-138           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PCB-153           | 0.02 (−0.02, 0.06)  | −0.02 (−0.06, 0.02) |
| PCB-180           | 0.03 (−0.03, 0.09)  | −0.01 (−0.08, 0.06) |
| Note: Free T4 models fit among 342 women (362 samples) and among 208 men (208 samples). All models controlled for lipids, age at blood draw (indicator variables for each decade), time of blood collection (indicator variables for morning, midday, afternoon, and midnight), and region (indicators for 6 regions). | | |
| Total T3 (ng/dL)  | β (95% CI) | β (95% CI) |
| PBB-118           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PBB-138           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PCB-153           | 0.02 (−0.02, 0.06)  | −0.02 (−0.06, 0.02) |
| PCB-180           | 0.03 (−0.03, 0.09)  | −0.01 (−0.08, 0.06) |
| Note: T3 models fit among 342 women (362 samples) and among 208 men (208 samples). All models controlled for lipids, age at blood draw (indicator variables for each decade), time of blood collection (indicator variables for morning, midday, afternoon, and midnight), and region (indicators for 6 regions). | | |
| Free T3 (pg/mL)   | β (95% CI) | β (95% CI) |
| PBB-118           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PBB-138           | −0.03 (−0.08, 0.02) | 0.02 (−0.01, 0.05) |
| PCB-153           | 0.02 (−0.02, 0.06)  | −0.02 (−0.06, 0.02) |
| PCB-180           | 0.03 (−0.03, 0.09)  | −0.01 (−0.08, 0.06) |
| Note: Free T3 models fit among 342 women (362 samples) and among 208 men (208 samples). All models controlled for lipids, age at blood draw (indicator variables for each decade), time of blood collection (indicator variables for morning, midday, afternoon, and midnight), and region (indicators for 6 regions). | | |

Note: T4, thyroxine; T3, triiodothyronine; TSH, thyroid-stimulating hormone; ln, natural log. β-coefficients should be interpreted as follows: a natural log increase in a given congener is associated with a multiplicative change in TSH of e\(^{\beta}\). In terms of changes in PBB and PCB levels not logged an additive change in each hormone of \(100\times e^{\beta}\) should be interpreted. All PBB congeners measures below the LOD were multiply imputed (n=10) based on a lognormal probability distribution whose parameters were determined by maximum likelihood estimation.
and 27 y old on average at PBB exposure, respectively. In the present study, among those born before the contamination incident, the average age at exposure was 15 (13 for women and 19 for men), and we included those born after the incident as well. Therefore, it may be possible that our study population was more vulnerable to thyroid disruption given the earlier age at initial PBB exposure. Another important difference is the exposure assessment methodology. The Yard et al. (2011) study used enrollment PBB and PCB levels, which were measured in the early 1970s soon after the contamination episode, whereas we were able to quantify individual PBB and PCB congeners with increased sensitivity particularly at low levels owing to a detection limit three orders of magnitude lower than in older methods (Needham et al. 1981). Finally, we also measured thyroid hormones, which allowed us to include those with undiagnosed thyroid diseases (n = 7) in addition to those identified by self-report, whereas Yard et al. (2011) included self-reported cases only.

One alternative explanation for the observed association between PBBS and thyroid disease among women is that exposure and disease may both be related to participation, thereby potentially inducing an association (Hernán et al. 2004). Because this cohort was historically exposed to PBBS and many had received information about their PBB levels in the past, more highly exposed individuals may have been more motivated to participate in this study. Additionally, individuals with diagnosed health conditions may have been more likely to participate. If both of these factors affected participation, an association between PBBS and disease might be expected.

A second possible explanation for our findings is that thyroid function may influence serum PBB and PCB levels. Although these chemicals are long-lived in human tissue, the most biologically relevant exposure parameter for making causal inference would be at some point before the disease process started. In our study, we measured PBBS and PCBs after disease diagnosis, and because the thyroid regulates metabolism, including actions that metabolize halogenated compounds (Park et al. 2012; Safe 1994; Takahashi et al. 2010), it may be plausible that disease status could affect serum levels.

Our finding that the association between PCBs and TSH varied by iodine status should be interpreted with caution due to the small number of women for whom we had measured urinary iodine. However, similar observations have been reported in several studies with other environmental toxicants (Blount et al. 2006; Jain 2014; Romano et al. 2015). Results from experimental studies have suggested that PBB and iodine may act synergistically on thyroid function, which could be due to inhibition of the

![Figure 1. β-Coefficients and 95% confidence intervals (95% CIs) from regression models for associations of PBB-153, PCB-118, and ΣPCB (di-ortho) deciles (nanograms/milliliter) with thyroid hormones stratified by sex. (A) PBB-153; (B) PCB-118; (C) ΣPCB (di-ortho). Note: D2–10, deciles 2–10; Ref, reference (decile 1); T4, thyroxine; T3, triiodothyronine; TSH: thyroid-stimulating hormone. All models adjusted for lipids, age, time of blood collection, and current smoking status (See S9 and S10 for β-coefficients and 95% CIs.) Sum of PCB 138, 153, and 180. (See the footnotes of Tables S9 and S10 for interpretations of β-coefficients.) Black circles denote women, and white circles denote men.](image-url)
incorporation of iodine into thyroidal amino acids (e.g., monoiodotyrosine), which are precursors of thyroid hormones (Allen-Rowlands et al. 1981; Sleight et al. 1978). In humans, the recently proposed “multiple-hit hypothesis” suggests that individuals with multiple thyroid “stresors” may be particularly susceptible to environmental thyroid disruption (Webster et al. 2014, 2015). Although we detected a difference in estimates across strata based on 150 μg/g creatinine [rather than the WHO (1994) iodine deficiency cut-off of 100 μg/g creatinine], this is still considered to be low iodine status (Caldwell et al. 2011) and may result in thyroidal stress.

In studies involving lipophilic chemicals, there has been considerable debate about how to statistically handle serum lipids (Chevrier 2013; O’Brien et al. 2015; Schisterman et al. 2005). Lipophilic compounds, such as PCBs, expressed on the lipid basis (nanograms/gram lipid) are correlated with the amount stored in body fat (Brown and Lawton 1984; Hirai et al. 2012), and because PCBs accumulate in lipids, this may be the most biologically relevant parameter for adjusting for lipids. However, this method mathematically constrains the form of lipids to be in the denominator of the PCB term (as nanograms/gram lipid) (Schisterman et al. 2005). In contrast, expressing PCBs on the wet weight (volume) basis (nanograms/milliliter serum) and controlling for lipids as a covariate allows for more flexible modeling and still accounts for the variability in serum lipid concentrations (Schisterman et al. 2005). Schisterman et al. (2005) showed through simulations that expressing compounds on the lipid basis induced more bias, consistently in the negative direction, than controlling for lipids as a covariate across various possible causal scenarios. Based on these findings, in our primary analysis, we considered PBBs and PCBs on the volume basis and controlled for lipids as a covariate, although we also performed all analyses on the lipid basis as well. More recently, O’Brien et al. (2015) published a simulation study similar to that of Schisterman et al. (2005) that concluded that modeling PCBs on the lipid basis incurred less bias than other methods and that adjusting for lipids as a covariate tended to bias effect estimates away from the null. In our study, we observed some differences in our results across these two methods. Expressing PBB-153 and PCBs on the lipid basis resulted in associations with thyroid disease that were mostly up and away from the null compared with results on the volume basis. Similarly, among men, we observed larger differences (more positive) in total and free T3 in relation to PCB congeners when expressed on the lipid basis than when expressed on the volume basis. These results are contrary to what may have been expected based on both the Schisterman and O’Brien studies and could be due to chance or to differences between the causal scenarios presented in these papers and the underlying causal relationships in our study. More research is needed to understand whether lipid-standardized PCB levels truly reflect adipose tissue levels or body burden as well as how accounting for lipids may affect epidemiological analyses.

This study benefited from several strengths. First, we were able to measure several important biomarkers, including a full
thyroid panel. We observed greater TSH concentrations among blood samples collected in the morning, which is consistent with its known diurnal pattern (Braverman and Cooper 2012). In addition, we also observed greater total T4 and T3 concentrations among women who reported current hormonal medication use or were pregnant, consistent with estrogen-induced increased sialylation of thyroxine-binding globulin (TBG), which leads to greater concentrations of TBG and thyroid hormones (Surks and Sievert 1995). These observations for multiple hormones support the validity of our thyroid hormone measurements. We also measured levels of several PBB and PCB congeners and lipids in serum as well as in a subset; iodine concentrations in longitudinally collected and pooled urinary samples. Although the subset with urinary iodine measurements was small, the design of pooling samples offers a methodological advantage over other studies that have used spot urine samples due to the reduction in the variability in iodine measurements for each individual (König et al. 2011). In addition, we had long-term follow-up on a cohort that was historically exposed to a flame-retardant mixture, which is significant for two reasons. First, it allowed for increased time for health conditions to manifest, including for those in a subsequent generation. Second, environmental PBB exposure occurred for a fixed and limited time, which allows us to infer that external exposure occurred before the outcome(s) of interest even though we assessed blood levels of PBBs and thyroid hormones at the same time, and because of its persistence, we were unable to infer when the potentially biologically relevant exposure may have occurred.

Our study was also subject to several limitations. First, a large proportion of participants (35%) who reported a history of thyroid disease did not specify what type they had been diagnosed with, which led to a loss of power and statistical precision when...
attempts to analyze specific thyroid disease subtypes such as hypothyroidism. Having access to medical records as opposed to our assessment of self-reported physician-diagnosed thyroid disease would have improved our ability to test associations with these different subtypes and would have allowed us to confirm self-reported diagnoses. Additionally, given that thyroid function comprises a variety of hormonal biomarkers, it is difficult to discern between potentially important hormonal changes and the influence of random error, multiple testing, or uncontrolled bias. Lastly, our study was limited by small sample size, particularly for our assessment of thyroid disease, a heterogeneous outcome, in relation to PBBs and PCBs. Nevertheless, we note that this is still one of the largest studies to date on these exposures and thyroid function.

Although PBBs and PCBs are no longer in production, exposure to PCBs remains widespread in human serum owing to their long biologic half-lives as well as to their persistence in the food chain (Sjödin et al. 2014; Wattigney et al. 2015; Xue et al. 2014). Furthermore, they share structural similarities with their replacements such as PBDEs and with newer flame retardants currently being produced, such as tetrabromobisphenol A and hexabromocyclododecane (Birnbaum and Staskal 2004). This makes the study of these older chemicals still relevant and important, as they may serve as a model for exposure to other persistent lipophilic halogenated compounds.

Conclusions

In this cohort of individuals historically exposed to PBBs, we found continued evidence of elevated levels of exposure to PBB-153 compared with national estimates, as well as a higher than expected prevalence of self-reported thyroid disease, particularly in women. We found some evidence to suggest that PBB-153 concentrations were associated with thyroid disease among women and possibly with hypothyroidism specifically. Lastly, modest thyroid hormone differences were observed in relation to serum PCB concentrations, and there was a suggestion that this may have varied by iodine status. Many decades after an acute exposure to flame retardants, this population remains highly exposed to PCBs, which may have contributed to the prevalence of thyroid disease.

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