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Concentrations of Polybrominated Diphenyl Ethers (PBDEs) and 2,4,6-Tribromophenol in Human Placental Tissues

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Abstract

Legacy environmental contaminants such as polybrominated diphenyl ethers (PBDEs) are widely detected in human tissues. However, few studies have measured PBDEs in placental tissues, and there are no reported measurements of 2,4,6-tribromophenol (2,4,6-TBP) in placental tissues. Measurements of these contaminants are important for understanding potential fetal exposures, as these compounds have been shown to alter thyroid hormone regulation *in vitro* and *in vivo*. In this study, we measured a suite of PBDEs and 2,4,6-TBP in 102 human placental tissues collected between 2010–2011 in Durham County, North Carolina, USA. The most abundant PBDE congener detected was BDE-47, with a mean concentration of 5.09 ng/g lipid (range: 0.12–141 ng/g lipid; detection frequency 91%); however, 2,4,6-TBP was ubiquitously detected and present at higher concentrations with a mean concentration of 15.4 ng/g lipid (range: 1.31–316 ng/g lipid; detection frequency 100%). BDE-209 was also detected in more than 50% of the samples, and was significantly associated with 2,4,6-TBP in placental tissues, suggesting they may have a similar source, or that 2,4,6-TBP may be a degradation product of BDE-209. Interestingly, BDE-209 and 2,4,6-TBP were negatively associated with age ($r_s = -0.16$; $p = 0.10$ and $r_s = -0.17$; $p = 0.08$, respectively). The results of this work indicate that PBDEs and 2,4,6-TBP bioaccumulate in human placenta tissue and likely contribute to prenatal exposures to these environmental contaminants. Future studies are needed to determine if these joint exposures are associated with any adverse health measures in infants and children.

Keywords

Brominated flame retardants; polybrominated diphenyl ethers; thyroid hormone; placenta

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Introduction

Polybrominated diphenyl ethers (PBDEs) have been used as additive flame retardants for decades in a variety of applications from polyurethane foams to high-impact polystyrene (HIPS). The presence of PBDEs in consumer products has led to their accumulation in indoor environments, and subsequent human exposure via inadvertent ingestion and/or inhalation of dust particles^{1,2}. Particular attention has been given to a PBDE commercial mixture known as pentaBDE, which had a primary application in polyurethane foam used in residential furniture^{3,4}. Studies have documented higher serum concentrations of PBDEs associated with pentaBDE in the US population relative to other regions of the world, likely due to the higher use of this mixture in residential furniture to meet a regional (state of California) flammability standard⁵. While the use of pentaBDE has now been banned or phased-out throughout the world, many older products in the home still contain these flame retardants, which will continue to leach into the indoor environment during the product lifetime. As a result, human exposure to PBDEs will continue for years to come, especially with the use of recycled foams and plastics in consumer products that may contain these phased-out chemicals. As such, PBDEs continue to be measured in human tissues such as serum, breast milk, umbilical cord blood, and placental tissues, suggesting that prenatal exposures to PBDEs occurs during pregnancy, and continues during infancy via breast feeding⁶⁻⁹.

In contrast, 2,4,6-tribromophenol (2,4,6-TBP) is widely used as an industrial chemical with an estimated US production volume of 4500 to 23,000 tonnes in 2006¹⁰. 2,4,6-TBP has multiple applications, including use as an antifungal agent (e.g. as a replacement for pentachlorophenol) in wood applications, as a reactive brominated flame retardant (BFR), and as an intermediate in the production of other BFRs. 2,4,6-TBP can also be formed as a result of the photolytic degradation of tetrabromobisphenol-A (TBBPA), a widely used reactive BFR, and during the synthesis of 1,2-bis (2,4,6-tribromophenoxy) ethane (BTBPE)¹¹. In addition to the anthropogenic sources of 2,4,6-TBP, there are natural sources of 2,4,6-TBP and other bromophenols from marine organisms and algae¹². Few toxicity studies have examined the effects of 2,4,6-TBP in animal models. One study examined oral exposure to 2,4,6-TBP in adult zebrafish and observed reproductive toxicity in addition to perturbed gonadal morphology when exposed to spiked food at concentrations of 3300 ug/g dw¹³. Only a few studies have examined environmental levels and human exposure to 2,4,6-TBP. It has been measured in marine sediments at an average concentration of 3.02 ng/g dry weight and in riverine systems at 0.66 ng/g dry weight¹⁴. 2,4,6-TBP has also been measured in the indoor environment of Japanese homes, with indoor house dust concentrations ranging from 15–30 ng/g and indoor air concentrations between 220–690 pg/m³¹⁵. Very few biomonitoring studies have included 2,4,6-TBP in the analyses of human tissues such as serum, cord blood, and/or breast milk. One Japanese study collected maternal serum and umbilical cord blood from a cohort of 16 mothers in 2006 for analysis of BFRs and PCBs. This study measured 2,4,6-TBP in maternal blood at a concentration of 22 pg/g wet weight and in cord blood at a concentration of 37 pg/g wet weight¹⁶. BFRs were also evaluated in Norwegian individuals working in electronics dismantling facilities, where 2,4,6-TBP was measured in plasma ranging from 0.17 to 81 ng/g lipid¹⁷. In a study measuring BFRs in a

Canadian Inuit population from Nunavik, Quebec, plasma samples contained a geometric mean 2,4,6-TBP concentration of 9.4 µg/kg lipid, however, these concentrations were not correlated with PBDE concentrations¹⁸. Finally, Qiu et al. measured mean 2,4,6-TBP concentrations of 5.6 ng/g lipid in fetal plasma and 0.8 ng/g lipid in maternal plasma¹⁹.

PBDEs and 2,4,6-TBP share a chemical structure that is similar to endogenous thyroid hormones (THs), and have been demonstrated to disrupt TH homeostasis either *in vitro* or in animal exposure studies^{20,21}. Concentrations of PBDEs in human serum have also been found to be significantly correlated with circulating levels of THs in adults, and are associated with adverse neurodevelopmental outcomes in children^{22,23}. Early childhood represents a developmental period that is vulnerable to endocrine disruption. Development is a hormonally-regulated growth process that is sensitive to perturbations by environmental contaminants, like PBDEs and 2,4,6-TBP. The *in utero* stage of development also represents a highly vulnerable period of fetal growth that may be even more sensitive to endocrine disruption due to the underdeveloped nature of the fetus' detoxification pathways, in addition to the myriad different growth and developmental processes that are occurring throughout gestation.

The placenta acts to facilitate the materno-fetal transfer of nutrients, gas, waste, and hormones throughout gestation and can act as a protective barrier against toxins and environmental contaminants²⁴. In the case of PBDEs, passive diffusion and/or active uptake of these chemicals into the placenta occurs, and the placenta can act as a repository for these lipophilic chemicals. For example, one study looked at mother-child pairs in China and compared the placental transfer characteristics of various environmental endocrine disruptors, including PBDEs. Their results indicated that PBDEs can be transferred across the placenta from maternal circulation, and eventually reach the fetus²⁵. Additionally, Frederiksen et al. used an experimental *ex vivo* human placenta perfusion system to show the differences in transplacental transfer of PBDEs based on degree of bromination²⁶. Thus there is a need to better understand the accumulation of these contaminants in placental tissues, in order to understand fetal exposures. In this study, we present our findings from the analysis of 102 human placental tissues that were collected in North Carolina, USA. Tissue samples were analyzed for a suite of PBDEs and 2,4,6-tribromophenol in order to increase our understanding of exposures during pregnancy and their accumulation within the placenta.

Materials and Methods

Participant recruitment

Participants were recruited from within an observational prospective cohort study assessing the joint effect of social, environmental, and host factors on pregnancy outcomes (the Healthy Pregnancy, Healthy Baby (HPHB) Study conducted by the Children's Environmental Health Initiative)^{27,28}. The HPHB study enrolled pregnant women from the Duke Obstetrics Clinic and the Durham County Health Department Prenatal Clinic at the Lincoln Community Health Center in Durham, NC. Our analyses included a subset of women from the HPHB study that delivered at the Duke University Medical Center between March 2010 and December 2011. The intentional study design was to oversample women

attending the Lincoln Community Health Clinic, in order to explore disparities in pregnancy outcomes by comparing African-American women with good outcomes to those with poor outcomes. As a result, the study population is predominantly African-American women with a lower socioeconomic standing and low educational attainment relative to the general US population. All aspects of this study were carried out in accordance with a human subjects research protocol approved by the Duke University Institutional Review Board.

Sample Collection

Consenting women had placenta tissue subsamples taken at the time of delivery at the Duke University Medical Center. Tissues (approximately 5–20 g) were stored in screwtop cryovials at -80°C until analysis.

Chemicals

All solvents used for the analysis were HPLC-grade or better. A fluorinated BDE standard, 2,3',4,4',6-tetrabromodiphenyl ether (FBDE-69)(Chiron Inc., Trondheim, Norway), ^{13}C labeled 2,2',3,4,5,5'-hexachlorinated diphenyl ether (CDE-141) (Cambridge Isotope Laboratories, Andover, MA), and labeled ^{13}C -2,2',3,3',4,4',5,5',6,6'-decabromodiphenyl ether (BDE-209) were used as internal and recovery standards for the BFR extractions. PBDE calibration standards were purchased from Accustandard and 2,4,6-tribromophenol was purchased from Cambridge Isotope Laboratories, Andover, MA.

BFR Analysis and Lipid Determination

Extractions were performed using between 2 and 17 grams of placenta tissue, depending on the sample and the amount collected during delivery. Tissues underwent 24 hours of lyophilization in order to completely dry the samples. The freeze-dried tissue samples were then homogenized into a fine powder with a pre-cleaned mortar and pestle before adding 15 mL of 1:1 hexane/dichloromethane (DCM) and letting the samples sit overnight, in order to allow for full solvent penetration. Samples were spiked with 1 ng of FBDE-69 and ^{13}C -BDE-209 as internal standards. All glassware used for BFR analysis were cleaned by muffle furnace, in addition to triple-rinsing with hexane, DCM, and methanol solvents in order to minimize background contamination. Samples then underwent 20 minutes of water bath sonication followed by centrifugation, after which the solvent was decanted to a separate tube. The extraction step was then repeated twice (three times total), and the solvent extracts were combined in a clean 50 mL glass centrifuge tube. Following extraction, the samples were blown down under a gentle stream of N_2 to a volume of 1 mL. A small aliquot of the extract was used for gravimetric lipid analysis and the remaining extract was passed through acidified silica columns for sample clean-up. Deactivated silica (4.0 g) was acidified using 40% by mass H_2SO_4 , shaken, and loaded into a glass chromatography column. The columns were pre-cleaned by rinsing with hexane and acetone and then conditioned with 15 mL of the elution solvent mix. The extract was then loaded on to the column and eluted using 30 mL of 80:20 hexane/DCM. Sample extracts were then blown down under a gentle stream of N_2 gas to a final volume of 100 μL . Samples were transferred to 200 μL glass vial inserts and spiked with 1 ng of ^{13}C -CDE-141 as a recovery standard. Finally, PBDEs and 2,4,6-TBP were identified and quantified using authenticated standards and gas chromatography with electron capture negative ion mass spectrometry (GC/ECNI-MS).

Quality Control/Quality Assurance

Laboratory blanks (e.g. sodium sulfate) were included with each batch of tissue sample extractions beginning with lyophilization (one batch includes 10 tissues samples plus two lab blanks). All sample values were blank subtracted and MDLs were calculated as three times the standard deviation of the lab blank values for each analyte. Individual values were normalized to the measured lipid content of each tissue sample used for the extraction procedure to yield a final value in ng/g lipid.

Labeled internal standards were used as surrogates and internal standards in all samples and included F-BDE-69 and ^{13}C -BDE-209 as internal standards (spiked prior to extraction) and ^{13}C -CDE-141 as a recovery standard (spiked prior to GC/MS analysis). The recovery of the internal standards was calculated for all tissue samples and laboratory blanks in order to assess the recovery efficiency of the extraction methods. The mean recovery in the lab blanks for FBDE-69 was $82.5 \pm 14\%$, while mean sample recovery was $60 \pm 12\%$.

Additionally, the BFR extraction method was validated using Standard Reference Material (SRM) 1947 (NIST, Gaithersburg, MD). SRM 1947 is a Lake Michigan fish homogenate with certified concentrations of PBDEs. The BFR extraction procedure previously described was used on a triplicate set of SRM 1947 samples. The concentrations of PBDE congeners of interest (BDE-47, -66, -99, -100, -153, and 154) were measured at 99%–116% of the certified values. Recovery of 2,4,6-TBP was evaluated by spiking 10 ng into a laboratory blank (in triplicate) and carrying it through the method. Recoveries averaged 87% ($\pm 29\%$).

Statistical Analysis

Statistical analyses were performed using JMP Pro 11. ΣBDE was calculated by summing all PBDE congeners including BDE-47, -99, -100, -153, -154, and -209, while ΣBFR includes all PBDE congeners plus 2,4,6-TBP. Only analytes with $\geq 50\%$ detection frequency were included in statistical analyses. Values below MDL were assigned a value equal to one-half the detection limit for statistical analyses. Preliminary analyses (Shapiro-Wilkes Test) indicated that the PBDE data were not normally distributed. As such, Spearman rank sum correlation analyses were used to assess the relationships between PBDE congeners in placenta and to assess their relationship with maternal age. It is important to note that the BFR concentrations were not significantly and positively associated with lipid content; however, we conducted all statistical analyses with both wet weight and lipid normalized concentrations. Results were similar using both methods. We present statistics using lipid normalized concentrations to facilitate comparison with other studies. Alpha < 0.05 was considered statistically significant.

Results

Population characteristics

Participant demographics are summarized in Table 1. Sixty-eight percent of the women in the study were non-Hispanic black. Most (58%) women were relatively young, between the ages of 18–24 years old (range 18–40). This was the first pregnancy for 45.5% of the women. Of all participants, 43.6% reported completing high school, and less than 10% of

the women had private health insurance. Recruitment for this study used English literacy as an exclusion criteria, so the demographics of this study population are not entirely reflective of the population of women visiting the Prenatal Clinic at the Lincoln Community Health Center in Durham, NC. The population of women who most commonly use this prenatal clinic are Hispanic, while the women included in this study are predominantly non-Hispanic black women with a lower socioeconomic standing.

BFRs

Detection frequencies for BDE-47, -100, -99, -154, -153, -209, and 2,4,6-TBP were all greater than 50% and are presented in Table 2 along with the range and distribution of concentrations measured. It is interesting to note that 2,4,6-TBP was detected in 100% of the samples and constituted 47.8% of the Σ BFR concentration measured in tissues. The most prominent PBDE measured was BDE-47, representing 34% of Σ BDE burden. The geometric mean concentration of 2,4,6-TBP was 15.4 ng/g lipid (range: 1.31 – 316 ng/g lipid), while the geometric mean concentration of BDE-47 was 5.09 ng/g lipid (range: 0.12 – 141 ng/g lipid). The PBDE congener ranking profile from highest geometric mean concentration to lowest geometric mean concentration is: BDE-47, -209, -153, -99, -100, -154 (Figure 1). Given the relatively homogenous distribution of our population, we were underpowered to examine associations between BFR exposures and race/ethnicity. However, we did examine associations with age. Interestingly, BDE-209 and 2,4,6-TBP were negatively associated with maternal age, $r_s = -0.16$ ($p = 0.10$) and $r_s = -0.17$ ($p = 0.08$), respectively, although again the associations did not reach statistical significance at $p < 0.05$. The remaining PBDE congeners showed no suggestion of associations with maternal age ($p > 0.20$).

Associations between PBDEs and 2,4,6-TBP

Correlation analyses are summarized in Table 3. All BFRs were significantly ($p < 0.001$) and positively correlated with each other and with Σ BFR concentrations. BDE-100 showed the strongest correlation ($r_s = 0.89$) with Σ BDE content followed by BDE-47 ($r_s = 0.84$). Interestingly, 2,4,6-TBP was significantly associated with all PBDE congeners. For example, 2,4,6-TBP showed a moderately strong correlation with BDE-209 ($r_s = 0.58$; $p < 0.001$; Figure 2).

Discussion

This is the first study to measure both PBDEs and 2,4,6-TBP in human placenta tissues, and observe a suggestive negative association with maternal age. It has often been assumed that the relative tissue concentrations of environmental contaminants within the placenta can be representative of fetal exposures for some contaminants²⁹. In fact, numerous studies have examined the relationships of environmental contaminants, such as PBDEs, within maternal serum, umbilical cord blood, and placenta tissues. The results of these studies indicate that transplacental transfer (TPT) of PBDEs does occur, and leads to fetal exposure during gestation^{30–33}. For example, a study by Frederiksen et al. found that PBDE exposure in the indoor environment, specifically from house dust ingestion, is linked to PBDE concentrations in maternal and umbilical cord plasma, which are additionally correlated with

the PBDE concentrations measured in the paired placental tissues. This study also showed a decreased rate of transport of PBDE congeners across the placenta with increasing degree of bromination²⁹. These results illustrate the placental transfer of PBDEs following maternal exposure to house dust and/or dietary sources of PBDEs, and also show that PBDEs are transferred to the fetal compartment during gestation. However, to our knowledge, no studies have examined TPT of 2,4,6-TBP, which should be addressed in future studies.

Additional research has also been conducted using a human *ex vivo* placenta perfusion system to study the kinetics and placenta transfer characteristics of BDE-47, -99, and -209. Significant accumulation was observed for all PBDE congeners tested, with placental transfer of BDE-47 being faster and more extensive than BDE-99 and BDE-209²⁶. These results indicate that *in utero* exposure to PBDEs occurs during gestation as a result of placental transfer, with higher rates of transfer and exposure for the lower brominated congeners. In contrast to these results, Chen et al. measured higher ratios between fetal cord blood and maternal placenta (F/M ratio) for PBDEs with a higher degree of bromination, suggesting that TPT increases with increasing degree of bromination³⁴. In addition, the ability of a chemical to bind to plasma transporter proteins will likely affect TPT characteristics. In the case of PBDEs, which have chemical structures similar to that of THs, their ability to bind TH transport proteins such as transthyretin (TTR) and/or TH membrane transporters such as OATPSs, MCTs, and LATs, may affect their TPT properties³⁵. Different compounds exhibit different partitioning and transport behaviors, and the exact mechanisms of TPT are not fully understood, however, the presence of contaminants found in both maternal and fetal circulation is clearly indicative of fetal exposure.

To date, only two other studies have measured PBDEs in placenta tissue samples from US populations^{9,36}. Additionally, placenta tissues from China, Japan, and European countries including Spain, Denmark, and Finland have been evaluated for PBDEs^{31,37-40}. The results from these studies are summarized in Table 4. The median value for Σ PBDEs₄₋₇ (tetra-through hepta-substituted congeners) in this present study was 13.8 ng/g lipid, and is similar to levels reported for placentae from individuals living near a Chinese e-waste site (19.5 ng/g lipid and 19.4 ng/g lipid), as well as in another US cohort that had a smaller sample size (n=42; 23.7 ng/g lipid). However, these values are much higher than the average Σ PBDE concentrations found in European samples (1.09 ng/g lipid), as well as placentae from Japan (0.25 ng/g lipid)⁴¹.

These current findings align with previous studies that have reported higher concentrations of PBDE congeners associated with the pentaBDE mixture in human samples from North America. North American concentrations are generally one to two orders of magnitude higher than those measured in European and Asian populations as a result of differences in fire regulatory standards, chemical regulatory and policy frameworks, and overall use and exposure to PBDEs⁴². In Europe, Japan, and China, BDE-209 is often found to be the most abundant congener, accounting for more than 50% of the total concentrations in human placentae^{31,39,43}. In the US, however, BDE-47 is the most prevalent congener measured in human and other biological tissues, and this is consistent with the higher concentrations of BDE-47 measured in indoor dust in the US compared to other countries. It is interesting to note that in this study, BDE-209 concentrations measured in placental tissues were

approximately equal to measurements made in Chinese samples, twice as high as the Danish samples, and eight times higher than Japanese samples. Furthermore, the measured values for Σ PBDEs in American placentae from this study are relatively similar to those measurements found in samples from individuals living and working in Chinese e-waste recycling towns. E-waste dismantling and processing is an occupation that typically involves significant human exposure to flame retardants due to their higher contact with electronic components containing flame retardant chemicals. The concentration of PBDEs measured in placenta tissues from both populations suggests that the same level of PBDE exposure and accumulation occurs between the general US population and Chinese e-waste recycling town inhabitants, despite the stark discrepancy in their exposure scenarios, and likely exposure pathways. However, lower brominated PBDEs that are more commonly found in pentaBDE applications such as polyurethane foams sold in North America, were measured in high concentrations in the Chinese e-waste worker samples, despite the fact the pentaBDE has limited use in electronics. These may be the result of metabolic and/or abiotic debromination of BDE-209 and other higher brominated PBDEs that are more widely used in electronics and plastics.

In the current study, we also observed a suggestive negative association with maternal age for BDE-209 and 2,4,6-TBP, which to our knowledge, has not been observed previously. The explanation for this negative relationship is unclear, but may relate to differences in exposure based on difference in behavior with age (e.g. time spent in various micro-environments). The fact that both BDE-209 and 2,4,6-TBP were negatively associated with maternal age, and that both were correlated with each other, suggests that they may share a similar source (e.g. electronics). Usually, PBDE congeners within a single commercial mixture are more strongly correlated with one another than between commercial mixtures. However, our results are partially in agreement with a recent assessment of placental PBDE concentrations that measured significant correlations between BDE-209 and BDE-28, -47, -99, and -183, but not between BDE-209 and BDE-100, -153, and -154⁴⁴. There are currently no other studies that have measured 2,4,6-TBP in human placenta tissue. The specific applications of 2,4,6-TBP as a reactive flame retardant remain unclear; however it appears that exposure to 2,4,6-TBP is common within our study population since it was detected in 100% of samples. 2,4,6-TBP was found to have a positive correlation with all PBDEs quantified, but was strongest for BDE-99 and BDE-209. 2,4,6-TBP and BDE-209 are not commonly analyzed together in biological tissues, likely due to the lower awareness of 2,4,6-TBP as an environmental contaminant of interest. As stated earlier, this unique relationship may be a result of these chemicals having related sources of exposure, or it may be indicative of a metabolic pathway that transforms BDE-209 into 2,4,6-TBP.

Previous work has explored the *in vitro* endocrine-disrupting potency of 2,4,6-TBP, as well as other BFRs, using a wide variety of assays. 2,4,6-TBP was found to be a potent inhibitor of estradiol sulfotransferase (ESULT) activity along with TBBPA and 6-OH-BDE-47, while the PBDEs showed much higher IC₅₀ value (half maximal inhibitory concentration, or the concentration at which the enzyme activity is diminished by 50%) and/or no ESULT inhibition, indicating that ESULT inhibition potency is determined largely in part by the presence of a hydroxylated aromatic group. 2,4,6-TBP was also shown to be a very potent

thyroxine competitor in the transthyretin (TTR)-binding assay, with a TTR-binding affinity 10.2 greater than the natural ligand, thyroxine⁴⁵. Additionally, the ability of 2,4,6-TBP to inhibit thyroid hormone SULT activity in pooled human liver cytosol was evaluated, and 2,4,6-TBP was shown to have an IC₅₀ value of 8.3 nM, which was more potent than any of the hydroxylated PBDEs profiled⁴⁶. However, more research is necessary to understand the sources of 2,4,6-TBP in the indoor environment, as well as to examine potential adverse effects from exposure to 2,4,6-TBP among the general population.

It is well known that the first trimester of pregnancy is a critical period for fetal neurodevelopment, and that these neurodevelopmental processes are largely driven by the action of THs⁴⁷. Animal exposure studies with PBDEs have shown permanent effects on spontaneous motor behavior (eg. hyperactivity) and decreased performance in learning and memory tests, implicating PBDEs as developmental neurotoxicants and endocrine disruptors^{48,49}. Additionally, PBDEs have been shown to disrupt TH homeostasis; therefore, the presence of PBDEs in the placenta may impact the materno-fetal transfer of THs during gestation, leading to the disruption of TH-mediated processes in the fetal compartment. Furthermore, growing epidemiological evidence show associations between prenatal exposure to PBDEs and subsequent neurodevelopmental deficits measured in children^{50,23,51}. Overall, flame retardant levels should continue to be closely monitored in the placenta, as well as their potential effects on fetal TH status and neurodevelopment.

One potential shortcoming of this study is the subsampling technique used in the collection of the placental tissue samples. Placental samples were collected at delivery and then sub-sampled to share among various studies. Therefore it was impossible to collect a whole placenta and homogenize the sample prior to sub-sampling. The placenta is a large, highly vascularized, heterogeneous organ. As a result, inconsistent or non-standardized subsampling techniques of the placenta organ may result in differences in our measurements of BFRs. For example, subsamples taken from the highly vascularized central region of the organ may contain different concentrations of BFRs than a peripheral subsample that has different vasculature and adipose composition. Normalization to lipid content may help control for some of these differences. However, we observed no significant correlations between BFR concentrations on a wet weight basis (ng/g ww) with percent lipid. But despite the inconsistencies in subsampling, the median values of PBDE concentrations from this study agree with PBDE measurements from previous studies^{9,34,44}.

Results from this study indicate that PBDEs and 2,4,6-TBP bioaccumulate in human placenta tissues, and provide insight into fetal BFR exposure during pregnancy. This study also characterizes BFR exposures in a population of women from low socioeconomic backgrounds and represents a unique subpopulation of understudied women in the US that are not typically represented in other exposure studies. These data may provide useful comparisons to other study populations from different regions with different ethnic and socioeconomic backgrounds, and further our understanding of the exposure patterns across the US. Future studies should also consider investigating associations between adverse health outcomes and exposures to these mixtures of BFRs given their reported effects on endocrine function.

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Abbreviations

2,4,6-TBP	2,4,6-tribromophenol
BFR	brominated flame retardant
DI	deiodinase
GC/ECNI-MS	electron capture negative ion mass spectrometry
HIPS	high-impact polystyrene
LC-MS/MS	liquid chromatography tandem mass spectrometry
PBDE	polybrominated diphenyl ether
SPE	solid phase extraction
SULT	sulfotransferase
THs	thyroid hormones
TPT	transplacental transfer
TTR	transthyretin

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Highlights

- A suite of PBDEs and 2,4,6-TBP were measured in 102 placenta tissue samples.
- BDE-209 was detected in more than 50% of the samples.
- 2,4,6-TBP was found in the highest concentrations in placenta tissue.
- 2,4,6-TBP was significantly correlated with PBDEs.
- BDE-209 and 2,4,6-TBP were suggested to be negatively associated with maternal age.

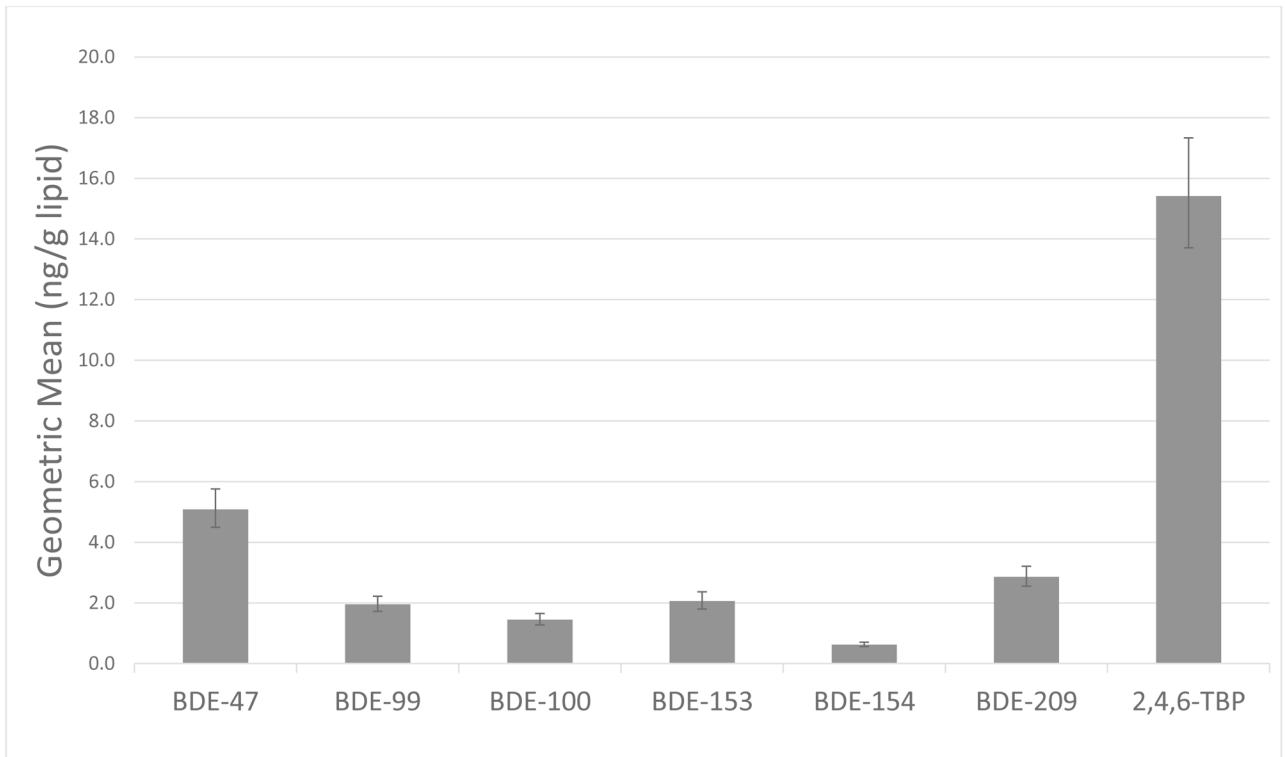


Figure 1. Geometric mean concentrations of BFRs measured in human placenta tissues (n=102; Error bars represent \pm SEM)

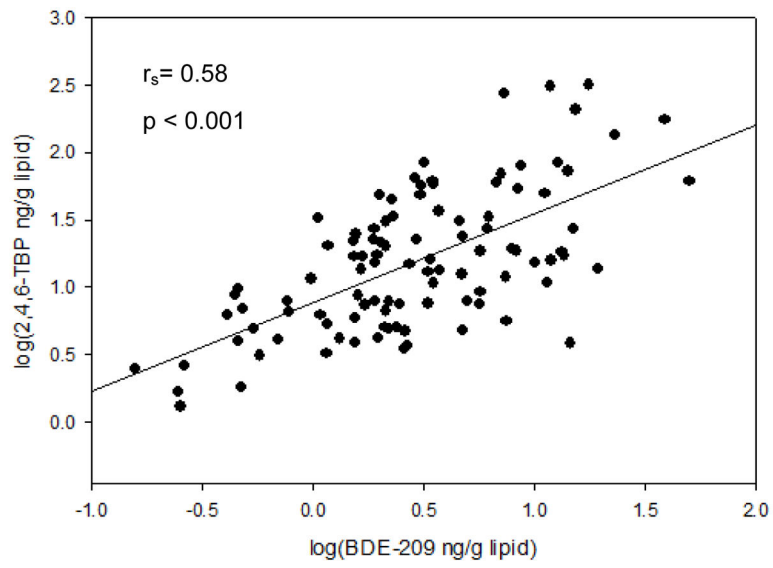


Figure 2.
Scatterplot showing correlation between 2,4,6-TBP and BDE-209

Table 1Cohort characteristics (n=101^{*})

Characteristic	N (%)
Maternal race	
Non-Hispanic white	16 (15.8)
Non-Hispanic black	69 (68.3)
Hispanic	12 (11.9)
Other	4 (4.0)
Maternal age	
18–19	21 (20.8)
20–24	38 (37.6)
25–40	42 (41.6)
Parity	
First birth	46 (45.5)
Male infant	52 (51.5)
Maternal education	
Less than high school	25 (24.8)
High school diploma	32 (31.7)
More than high school	44 (43.6)
Not married	82 (81.2)
Smoked during pregnancy	22 (21.8)
Private health insurance [#]	9 (9.3)

* Demographic data was missing for one individual

[#] No data is available on health insurance for four of the individuals

Table 2

BFR concentrations (ng/g lipid) measured in placental tissues.

Variable	MDL	Detection Frequency (%)	Geometric Mean	Min	Max	Percentile				
						25th	50th	75th	95th	
PBDEs (n=102)										
BDE-47	0.07	91.2	5.09	0.12	141	2.12	5.05	12.4	37.5	
BDE-99	0.07	68.6	1.95	0.09	223	0.62	1.95	4.43	17.1	
BDE-100	0.02	88.2	1.45	0.03	50.1	0.62	1.65	3.25	11.1	
BDE-153	0.01	93.1	2.06	0.02	513	1.21	2.36	4.15	16.9	
BDE-154	0.01	83.3	0.63	0.01	20.2	0.33	0.74	1.41	3.41	
BDE-209	0.17	52.9	2.86	0.16	50.4	1.55	2.64	6.83	17.3	
ΣPBDEs			17.6	0.54	528	8.71	19.10	34.7	98.7	
Phenolic compound (n=102)										
2,4,6-TBP	0.05	100	15.4	1.31	316	6.25	15.0	32.7	171	
ΣBFRs			37.3	2.18	568	18.3	38.1	75.6	317	

ΣPBDE was calculated by summing all PBDE congeners (i.e. BDE-47, -99, -100, -153, -154, and -209), while ΣBFR includes the sum of all PBDE congeners plus 2,4,6-TBP.

Table 3

Spearman correlation matrix for BFRs (ng/g lipid)

BFR	BDE-47	BDE-99	BDE-100	BDE-153	BDE-154	BDE-209	2,4,6-TBP	ΣPBDEs	ΣBFRs
BDE-47	1.00	0.48 [#]	0.88 [#]	0.58 [#]	0.61 [#]	0.49 [#]	0.50 [#]	0.84 [#]	0.73 [#]
BDE-99		1.00	0.52 [#]	0.43 [#]	0.52 [#]	0.60 [#]	0.66 [#]	0.68 [#]	0.72 [#]
BDE-100			1.00	0.71 [#]	0.71 [#]	0.50 [#]	0.48 [#]	0.89 [#]	0.77 [#]
BDE-153				1.00	0.71 [#]	0.50 [#]	0.38 [#]	0.77 [#]	0.66 [#]
BDE-154					1.00	0.54 [#]	0.50 [#]	0.77 [#]	0.72 [#]
BDE-209						1.00	0.58 [#]	0.73 [#]	0.72 [#]
2,4,6-TBP							1.00	0.58 [#]	0.85 [#]
ΣPBDEs								1.00	0.89 [#]
ΣBFRs									1.00

[#] p<0.001

Table 4

PBDE measurements in human placental tissues from various regions.

Population Location	Year	# of placentae	# of BDE congeners	Median Σ_{3-7} BDEs (ng/g lw [*])	Median BDE-209 (ng/g lw [*])	Lipid (%)	Congener concentration ranking	References
USA	2010–2012	42	10	23.7	NA	NA	47>153=99>209	Nanes et al., 2014 ⁹
USA	2008–2010	102	7	13.8	2.64	1.09	47>209>153=99>100	<i>This study</i>
USA	2007–2008	5	42	NA	NA	NA	47=99>153	Dassanayake et al., 2009 ³⁶
China	2012	69	8	19.4	3.30	NA	28>209>153>183>47	Xu et al., 2015 (e-waste site) ³⁷
China	2012	86	8	1.9	2.08	NA	209>153>28>47>183	Xu et al., 2015 (reference site) ³⁷
China	2012	30	17	9.96	ND	1.42	47>99>153	Chen et al., 2014 ⁴⁰
China	2009–2011	31	9	3	2.64	1.38	209>197>153>47	Zhao et al., 2013 ⁴³
China	2005–2007	130	39	0.54	NA	NA	47>153=99	Ma et al., 2012 ⁵²
China	2005	5	36	19.5	NA	NA	47>153>99	Leung et al., 2010 (e-waste site) ⁵⁵
China	2005	5	36	1.02	NA	NA	47>153>99	Leung et al., 2010 (reference site)
China	NA	6	7	2.73	NA	NA	47	Zhang et al., 2008 ⁵⁴
Japan	NA	10	25	0.25	0.32	3.6	209>47>153	Takastuga et al., 2007
Spain	2003–2004	30	15	0.65	1.0	0.7	209>47>153	Gómara et al., 2007 ⁵¹
Denmark	2007	50	12	1.22	1.14	1.21	209>47=153	Frederiksen et al., 2009 ⁵⁹
Denmark	1997–2001	129	14	1.31	NA	1.09	153=47	Main et al., 2007 ⁵⁸
Finland	1997–2001	56	14	1.18	NA	1.21	47>153	Main et al., 2007 ⁵⁸

* lw = lipid weight based