

## Polybrominated Diphenyl Ethers in Dust from the Vicinity of Power Generation Plants in Nigeria: Distribution, Risks Assessment and Source Apportionment

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### ABSTRACT

Polybrominated diphenyl ethers (PBDEs) constitute a class of widespread environmental contaminants that pose significant health risks. Therefore, this study investigated the distribution, risks and sources of PBDEs in dusts from power generation plants in Nigeria. A total of 24 dust samples were collected from four power plants. Samples were collected for six months covering both wet and dry seasons. Dusts were extracted for PBDEs following the United State Environmental Protection Agency Method 3550C using a 1:1 mixture of dichloromethane and n-hexane. Thereafter, the extracts were cleaned-up and analyzed for PBDEs using a gas chromatograph-mass spectrometer. The results indicated that the mean concentrations of  $\Sigma 39$  PBDEs in the dusts ranged from 17.5 to 54.6 ng g<sup>-1</sup> for wet season and 55.4 to 157 ng g<sup>-1</sup> for dry season. There were significant spatio-temporal variations in the concentrations of PBDEs in the dust. The PBDEs have higher concentrations during the dry season than the wet season. The hazard index and total cancer risk values indicated that there is neither adverse non-carcinogenic risk nor carcinogenic risk from exposure to PBDEs in these dusts. Source apportionment indicated that the PBDEs in the dusts were from the use of deca- and penta-BDE commercial formulations used for dielectric insulating fluids for transformers, capacitors, cable insulators etc. It is hereby recommended that the Federal Ministry of Environment should establish environmental policy and enforceable regulatory frameworks for the control of PBDEs in order to minimize the levels environmental PBDEs and the risk to human and ecosystems.

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### INTRODUCTION

Polybrominated diphenyl ethers (PBDEs) constitute a class of widespread environmental contaminants that pose significant health risks. These compounds are industrially produced chemicals that have been utilized since the 1970s as flame retardants in plastics, textiles, furniture, and electric and electronic devices (Nakajima *et al.*, 2022, Bramwell *et al.*, 2016). Three major commercial formulations of PBDEs have been manufactured: Penta-BDE comprising of PBDEs 47, 99, 100 and 153, Octa-BDE comprising primarily PBDEs 153, 183, 197, 207, and 196 and Deca-BDE is made up of PBDE 209. The Penta-BDE mixtures has been widely utilized in polyurethane foam for mattresses, furniture, and upholstery as well as electronic equipment, including fax machine, computers, and televisions, while octa-BDE have added to thermoplastic resins, particularly acrylonitrile butadiene styrene (ABS) plastics and deca-BDE is incorporated into high-impact polystyrene (HIPS) used in production of casings for electrical and electronic equipment, as well as in clothing fabric, building materials, and polypropylene drapery (Jin *et al.*, 2022). PBDEs possess molecular structures and physicochemical properties that closely resemble those of other persistent organic pollutants (POPs), including polychlorinated biphenyls (PCBs), polychlorinated dibenzo-p-dioxins, and polychlorinated dibenzofurans (PCDD/Fs). They can readily leach from polymeric material in which they are incorporated into the environment during manufacturing, use, and disposal because they are not chemically bound to the polymer matrix through covalent bond (Iwegbue *et al.*,

2025, Gevao *et al.*, 2011). As POPs, PBDEs are highly resistant to degradation and can persist in various environmental matrices, including air, dust, water, soil, and sediments (Yang *et al.*, 2025, Ohoro *et al.*, 2021, Han *et al.*, 2016). They also have a strong tendency to bioaccumulate in living organisms, including humans. Numerous studies have demonstrated that exposure to PBDEs is associated with adverse effects on the endocrine system, neurodevelopment, reproductive health, and immune function (Gomes *et al.*, 2024, Lin *et al.*, 2024). Owing to these environmental and health concerns, PBDEs have been designated as priority-controlled substances under the Stockholm Convention on POPs, leading to stringent regulations and restrictions on their production and use in several major jurisdictions, including the European Union, China, and United State (Yang *et al.*, 2025). Despite regulatory prohibition, human exposure to these compounds remains ongoing, attributable to their persistent nature and their continued occurrence in older consumer goods still in circulation (Iwegbue *et al.*, 2025, Wu *et al.*, 2021).

Globally, the release of contaminants into the environment has been driven primarily by anthropogenic activities, including traffic and transportation systems, petroleum refining, industrial manufacturing and services, power generation plants, and gas flaring, among others. Of these processes, power generation plants have been identified as the foremost contributor to global environment pollution (Okedere *et al.*, 2021). In Nigeria, power generation from thermal plants has relied almost exclusively on the

combustion of the country's abundant natural gas reserves (Adesanmi *et al.*, 2019., Igbokwe *et al.*, 2015). The consumption of this resource, along with the resultant emissions into the environment causes contaminants including PBDEs to adhere to dust particles, which accumulate on surfaces of items and diffuse into various environmental compartments (Adesanmi *et al.*, 2019).

Dust serves as a complex mixture of small particles that collects and holds both synthetic and naturally occurring contaminants (Iwegbue *et al.*, 2025). Power generation plants release dust containing PBDEs which people (workers, nearby communities) are exposed to through multiple pathways such as ingestion, inhalation and dermal contacts. This exposure raises significant health concerns, particularly because these chemical compounds are toxic and individuals (workers) spend considerable time in such vicinity. Research across the United States and other nations, such as Nigeria, has consistently identified elevated concentrations of PBDEs in household dust, particularly in residential environments (Iwegbue *et al.*, 2025, Emebu *et al.*, 2024, Araujo *et al.*, 2024, Choi *et al.*, 2023, Qui, *et al.*, 2023, Zandona *et al.*, 2022, Ohoro *et al.*, 2021, Adeyi *et al.*, 2020, Oloruntoba *et al.*, 2019, Iwegbue *et al.*, 2019). This finding is significant because it demonstrated that dust inhalation represents the primary pathway through which people are exposed to PBDEs. According to Ohoro *et al.* (2021), dust contact may account for a substantial portion of approximately 82% of people's total daily exposure to these harmful chemicals. Most studies on PBDEs in dust have focused on environments such as

electronic waste (e-waste) recycling sites, electronic repair workshops, schools/offices, and household settings, both globally and in Nigeria. However, there is a paucity of information on the occurrence of PBDEs in dust from the vicinity of power generation plants. Existing studies conducted around power generation facilities have primarily investigated metals (e.g. Luqman *et al.*, 2024, Kumari *et al.*, 2023, Zhang *et al.*, 2020, Idowu *et al.*, 2020, Adeyi and Torto, 2014), PCBs (e.g. Li *et al.*, 2023, Aganbi *et al.*, 2019; Folarin *et al.*, 2018; Gedik and Imamoglu, 2011) and polycyclic aromatic hydrocarbons (PAHs) (Zou *et al.*, 2021; Yan, 2020; Gune *et al.*, 2019; Sushkova *et al.*, 2019) and with little or no attention given to PBDEs contamination. This knowledge gap underscores the need for the present study, which aimed to investigate the spatiotemporal distribution, assess potential risks and source apportionment of PBDEs in dust collected from the vicinity of power generation plants in Nigeria.

## MATERIALS AND METHODS

### Description of Study Area

The study area comprised Benin, Calabar, Sapele and Gbarain power plants located in Edo, Cross River, Delta and Bayelsa States respectively in south-south, Nigeria (Figure 1). The information regarding the study area is shown in Table 1. Other information on the study area such as climatic conditions etc have been previously documented (Tesi *et al.*, 2025; Tesi *et al.*, 2022; Tesi *et al.*, 2020a, b; Iwegbue *et al.*, 2021).

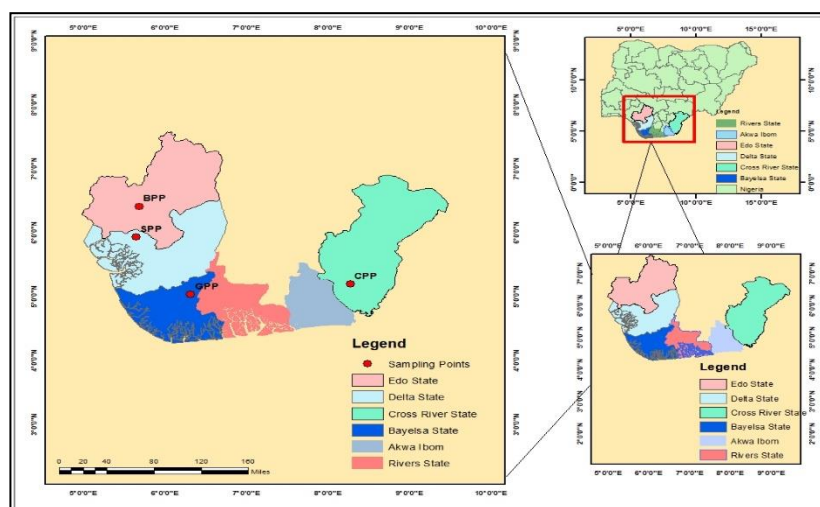


Figure 1: Map of study area

Table 1: Information Regarding the Sampled Power Generation Plants

| POWER PLANTS                      | COORDINATES             | LOCATION                       | TYPE                     | CAPACITY | STATUS      | YEAR COMPLETION | OF |
|-----------------------------------|-------------------------|--------------------------------|--------------------------|----------|-------------|-----------------|----|
| Calabar Power Plant (CPP)         | 5°11'21'' N 8°16'25'' E | Calabar, Cross River State     | Simple cycle gas turbine | 561MW    | Operational | 2014            |    |
| Benin (Ihovbor) Power Plant (BPP) | 6°24'20'' N 5°41'00'' E | Benin City, Edo State          | Simple cycle gas turbine | 450MW    | Operational | 2013            |    |
| Sapele Power Plant (BPP)          | 5°55'40'' N 5°38'41'' E | Sapele, Delta State            | Simple cycle gas turbine | 450MW    | Operational | 2012            |    |
| Gbarain (GPP) Power Plant         | 5°1'52'' N 6°18'5'' E   | Koroama-Yenagoa, Bayelsa State | Simple cycle gas turbine | 254MW    | Operational | 2017            |    |

### Reagents

High purity dichloromethane, n-hexane, a PBDE standard mixture of 39 congeners (BDE-1, 2, 3, 7, 8, 10, 11, 12, 13, 15, 17, 25, 28, 30, 32, 33, 35, 37, 47, 49, 66, 71, 75, 77, 85, 99, 100, 116, 118, 119, 126, 137, 153, 154, 155, 166, 181, 183,

and 209), alumina and silica gel were purchased from Isotope Laboratories Inc., (Cambridge, MA, USA).

### Sample Collection

Dust samples were collected with a brush and a stainless steel dust pan. At each power plant, composite sampling was

adopted. That is, dust samples were collected from different points and pooled together to form a composite sample. Dust samples were collected for a period of six (6) months i.e. August to October, 2024 for wet season and January to March, 2025 for dry season. Thus, a total of 24 dust samples were collected. Dust samples were kept in amber glass bottles. Samples were labeled properly, kept in ice chest and transported to the laboratory. Dust samples were sieved through < 63µm sieve before storing in the refrigerator until analysis.

### Extraction, Cleanup and Quantification of PBDEs

The PBDEs in the dusts were extracted by ultrasonication (EPA Method 3550C) using n-hexane/dichloromethane (DCM) (1:1 v/v) at 30 °C for 30 minutes and filtered. The extraction was repeated twice. Thereafter the extracts were combined and reduced to 1 mL by using a rotary evaporator. Then sodium sulphate was added to the extract to remove traces of water and then cleaned in a column of alumina (Al<sub>2</sub>O<sub>3</sub>) and silica gel (SiO<sub>2</sub>). The PBDEs was washed down from the column with 40 mL of DCM. The solution was concentrated to approximately 2 mL, and kept in a vial until analysis. The PBDEs in the extracts were quantified by means of a gas chromatograph equipped with mass spectrometry (HP 6890 Palo Alto, C A, USA).

### Quality Assurance and Quality Control

All the reagents used were of analytical grade. The effectiveness of the analytical procedure for extraction of the PBDEs was evaluated by employing method blanks and recovery studies. None of the 39 PBDEs congeners investigated were found in the method blanks (n=3). The percentage recoveries of PBDEs congeners from the spike matrix were in the range of 87.3 to 92.8 %. The external calibration approach was used to quantify the PBDE levels in each sample. The concentrations of the blank samples that result in a signal over noise ratio of three and ten, respectively, were used to calculate the limits of detection (LODs) and quantification (LOQs). The LODs ranged from 0.01 to 1.0 ng g<sup>-1</sup> while the LOQs ranged from 0.03 to 3.0 ng g<sup>-1</sup>. The r<sup>2</sup> of the calibration curves obtained from the plot of peak areas against known PBDEs concentrations varied from 0.9996 to 0.9999.

### Statistical Analysis

The statistical analysis of data was done using the IBM-SPSS software (version 27). The Kruskal Wallis and Mann-Whitney U tests were used to determine if there were significant spatio-temporal variations in the concentrations of PBDEs in the dusts. The relationship between PBDEs

homologues was established by means of Pearson's correlation coefficient. Principal component analysis (PCA) was utilized for source identification of PBDEs in the dusts.

### Human Risk Assessment

The non-carcinogenic risk of PBDEs in the dust was evaluated using the hazard index (HI) while the carcinogenic risk was evaluated using the total cancer risk (TCR) via the three exposure pathways of ingestion, inhalation, and dermal contact. Equations (1) through (4) were used to determine the HI, while equations (5) through (8) were used to determine the overall cancer risks (USEPA, 2009).

$$\text{Hazard index (HI)} = \text{HQ}(\text{Hazard Quotient})_{\text{Ingestion}} + \text{HQ}_{\text{Inhalation}} + \text{HQ}_{\text{Dermal contact}} \quad (1)$$

$$\text{HQ}_{\text{Ingestion}} = \left[ \frac{C \times \text{IngR} \times \text{EF} \times \text{ED}}{\text{BW} \times \text{AT}_{\text{nc}}} \times 10^{-6} \right] / \text{RfDo} \quad (2)$$

$$\text{HQ}_{\text{Inhalation}} = \frac{C \times \text{Inh} \times \text{EF} \times \text{ED} \times \text{RfDi}}{\text{PEF} \times \text{AT}_{\text{nc}}} \quad (3)$$

$$\text{HQ}_{\text{Dermal contact}} = \frac{C \times \text{SA} \times \text{AF} \times \text{ABS} \times \text{EF} \times \text{ED}}{\text{BW} \times \text{AT}_{\text{nc}}} \times 10^{-6} / (\text{RfDo} \times \text{GIABS}) \quad (4)$$

$$\text{Total cancer risk} = \text{Risk}_{\text{Ing}} + \text{Risk}_{\text{Inh}} + \text{Risk}_{\text{Dermal}} \quad (5)$$

$$\text{Risk}_{\text{Ingestion}} = \frac{C \times \text{IngR} \times \text{EF} \times \text{ED} \times \text{CF} \times \text{SFO}}{\text{BW} \times \text{AT}_{\text{ca}}} \quad (6)$$

$$\text{Risk}_{\text{Inhalation}} = \frac{C \times \text{Inh} \times \text{EF} \times \text{ED} \times \text{IUR}}{\text{PEF} \times \text{AT}_{\text{ca}}} \quad (7)$$

$$\text{Risk}_{\text{Dermal contact}} = \frac{C \times \text{SA} \times \text{AF} \times \text{ABS} \times \text{EF} \times \text{ED} \times \text{CF} \times \text{SFO} \times \text{GIABS}}{\text{BW} \times \text{AT}_{\text{ca}}} \quad (8)$$

The definitions of terms and values of variables used for evaluating the human health risk are given elsewhere (Iwegbue et al., 2025; Emoyan et al., 2022). In general, TCR values larger than 1.0 × 10<sup>-6</sup> imply carcinogenic risk, but HI values greater than 1 suggest adverse non-carcinogenic risk (USEPA, 2022).

## RESULTS AND DISCUSSION

### PBDEs Concentrations in the Dust

The concentrations of the PBDEs in the dusts from the power generation plants are shown in Tables 2 and 3 for wet and dry seasons respectively. The mean concentrations of total PBDEs in the dusts ranged from 17.5 to 54.6 ng g<sup>-1</sup> for wet season and 55.4 to 157 ng g<sup>-1</sup> for dry season. There were significant spatiotemporal variations (p < 0.05) in the PBDEs concentrations in the dust. The spatial variations are related to dust dispersion rate, pavement types and frequency of pavement cleaning (Ossai et al., 2021; Khanal et al., 2018). The mean total PBDEs concentrations were in the order of CPP > SPP > BPP > GPP for wet season and SPP > GPP > BPP > CPP for dry season. The highest concentration of total PBDEs was found in March 2025 except for CPP which has the highest concentration in September 2024 (Figure 1). Dry season PBDE levels in the dust were higher than wet season levels. The higher levels of PBDEs during the dry season may be due to a lack of dilution and reduced transport.

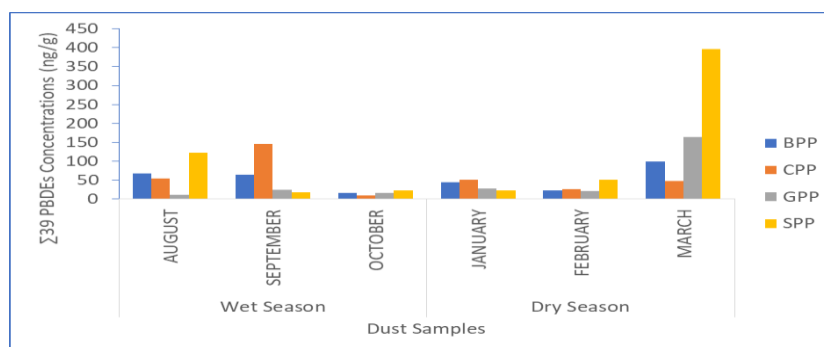


Figure 1: Monthly variations of PBDEs in the dust

The concentrations of PBDEs obtained in this study were similar to those reported for urban outdoor dusts in Nigeria, China and Vietnam irrespective of the number of PBDEs determined and the number of dust samples analyzed. In Nigeria, Iwegbue *et al.* (2025) reported a concentration range of 0.54 -60.4 ng g<sup>-1</sup> in urban dust of Port-Harcourt, Nigeria. In China, Wu *et al.* (2021) and Wu *et al.* (2015) reported a range of 53.5-213 ng g<sup>-1</sup> and 6.71-342 ng g<sup>-1</sup> respectively while Yu *et al.* (2012) reported a range of 8.7-3116 ng g<sup>-1</sup>. Moreover, Anh *et al.* (2018) reported a range of 0.91-56 ng g<sup>-1</sup> in outdoor dust from Vietnam.

#### PBDEs Occurrence Pattern in the Dust

The occurrence pattern of PBDEs in the dusts from the PGPs are displayed in Figure 2. The occurrence pattern of PBDEs homologues in the dust followed the order of penta-BDE > tri-BDE > hexa-BDE > tetra-BDE > hepta-BDE > deca-BDE > mono-BDE for wet season and penta-BDE > hexa-BDE > tri-BDE > di-BDE > tetra-BDE > hepta-BDE > deca-BDE > mono-BDE for dry season.

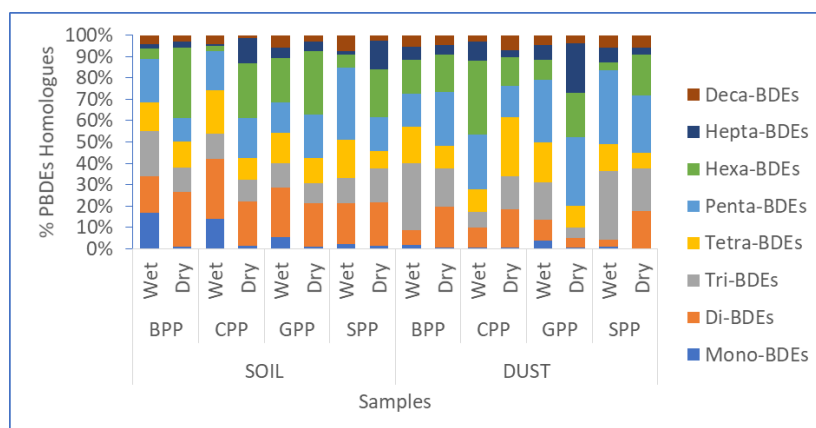


Figure 2: Homologues distribution of PBDEs in the dusts

The concentrations of the mono-BDEs (sum of BDE-1, BDE-2 and BDE-3) ranged from 0.22 to 0.84 ng g<sup>-1</sup> and constituted 0.4 to 3.6% of the total PBDEs. The concentrations of the individual mono-BDE congeners were in the order of BDE-3 > BDE-2 > BDE-1. The concentrations of the di-BDEs (sum of BDE-7, BDE-8, BDE-10, BDE-11, BDE-12, BDE-13, and BDE-15) ranged from 1.77 to 26.9 ng g<sup>-1</sup> and constituted 3.4 to 19.1% of the total PBDEs. The concentrations of the individual di-BDE congeners were in the order of BDE-10 > BDE-11 > BDE-15 > BDE-13 > BDE-8 > BDE-12 > BDE-7. The concentrations of the tri-BDEs (sum of BDE-17, BDE-25, BDE-28, BDE-30, BDE-32, BDE-33, BDE-35 and BDE-37) ranged from 3.0 to 31.3 ng g<sup>-1</sup> and constituted 4.8 to 32.1% of the total PBDEs. The concentrations of the individual tri-BDE congeners were in the order of BDE-17 > BDE-28 > BDE-35 > BDE-25 > BDE-33 > BDE-37 > BDE-30 > BDE-32. The concentrations of the tetra-BDEs (sum of BDE-47, BDE-49, BDE-66, BDE-71, BDE-75, and BDE-77) ranged from 3.29 to 19.2 ng g<sup>-1</sup> and constituted 7.5 to 27.5% of the total PBDEs. The concentrations of the individual tetra-BDE congeners were in the order of BDE-77 > BDE-71 > BDE-47 > BDE-49 > BDE-75 > BDE-66. The concentrations of the penta-BDEs (sum of BDE-85, BDE-99, BDE-100, BDE-116, BDE-118, BDEs-119 and BDE-126) ranged from 5.10 to 41.7 ng g<sup>-1</sup> and constituted 15.0 to 34.5% of the total PBDEs. The concentrations of the individual penta-BDE

congeners were in the order of BDE-85 > BDE-100 > BDE-99 > BDE-118 > BDE-1116 > BDE-119 > BDE-126. The concentrations of the hexa-BDEs (sum of BDE-137, BDE-153, BDE-154, BDE-155, and BDE-166) ranged from 1.69 to 30.6 ng g<sup>-1</sup> and constituted 3.7 to 34.6% of the total PBDEs. The concentrations of the individual hexa-BDE congeners were in the order of BDE-166 > BDE-155 > BDE-153 > BDE-137 > BDE-154. The concentrations of the hepta-BDEs (sum of BDE-181, and BDE-183) ranged from 1.21 to 16.5 ng g<sup>-1</sup> and constituted 3.3 to 23.2% of the total PBDEs. The concentrations of the individual hepta-BDE congeners were in the order of BDE-183 > BDE-181. The concentrations of the deca-BDE (BDE-209) ranged from 0.77 to 8.81 ng g<sup>-1</sup> and constituted 2.9 to 6.8% of the total PBDEs. On average, the penta-, hexa- and tri-BDEs were the top three dominant PBDE homologues in the dust. The lower brominated PBDEs (mono- to penta-BDEs) were dominant in both seasons (Figure 3). This might have significant environmental and health implications such as environmental degradation (debromination), increased ecological risk, and higher mobility. The concentrations of the LB-PBDEs (sum of mono- to penta-BDEs) in the dust ranged from 13.8 to 112 ng g<sup>-1</sup> and constituted 52.4 to 83.5% of the total PBDEs while the concentrations of the HB-PBDEs ranged from 3.67 to 44.5 ng g<sup>-1</sup> and constituted 16.4 to 47.6% of the total PBDEs.

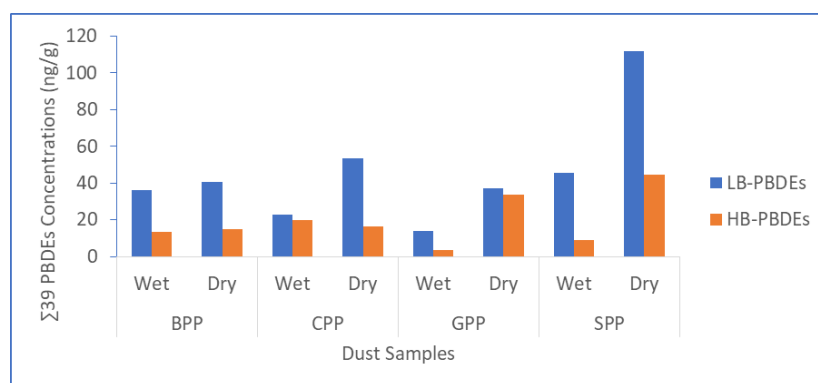


Figure 3: Lower brominated vs higher brominated PBDEs in dust

### Human Health Risks of PBDEs

The computed HI for children and adults' exposure to the PBDEs in the dusts from the PGP are shown in Table 4. The HQ values were as follows:  $HQ_{Ing} > HQ_{Derm} > HQ_{Inh}$  for both children and adults' exposure. The HI values ranged from  $4.08 \times 10^{-2}$  to  $3.67 \times 10^{-1}$  for children and  $5.66 \times 10^{-3}$  to  $5.08 \times 10^{-2}$  for adults in both seasons. The HI values adults were lower than those of children. This implied that children will be more prone to risk compared to adults. The HI values were

< 1. This indicates there were no adverse non-carcinogenic risk for human exposures to PBDEs in the dust. Moreover, the TCR computed for children and adults' exposure to PBDEs in the dusts from the PGP are shown in Table 5. The TCR values ranged from  $2.00 \times 10^{-6}$  to  $1.80 \times 10^{-5}$  for children and  $1.87 \times 10^{-7}$  to  $1.68 \times 10^{-6}$  for adults in both seasons. The TCR values were  $< 1 \times 10^{-6}$ . This indicates absence of carcinogenic risk for human exposures to PBDEs in the dust.

Table 4: Hazard Index of PBDEs in the dusts

| Children Exposures |     |                   |                   |                    |          | Adults' Exposure  |                   |                    |          |
|--------------------|-----|-------------------|-------------------|--------------------|----------|-------------------|-------------------|--------------------|----------|
|                    |     | HQ <sub>Ing</sub> | HQ <sub>Inh</sub> | HQ <sub>Derm</sub> | HI       | HQ <sub>Ing</sub> | HQ <sub>Inh</sub> | HQ <sub>Derm</sub> | HI       |
| BPP                | Wet | 9.10E-02          | 1.00E-04          | 2.55E-02           | 1.17E-01 | 1.14E-02          | 2.51E-04          | 4.54E-03           | 1.62E-02 |
|                    | Dry | 1.01E-01          | 1.12E-04          | 2.83E-02           | 1.30E-01 | 1.27E-02          | 2.79E-04          | 5.05E-03           | 1.80E-02 |
| CPP                | Wet | 7.76E-02          | 8.55E-05          | 2.17E-02           | 9.94E-02 | 9.69E-03          | 2.14E-04          | 3.87E-03           | 1.38E-02 |
|                    | Dry | 1.28E-01          | 1.41E-04          | 3.58E-02           | 1.64E-01 | 1.60E-02          | 3.53E-04          | 6.38E-03           | 2.27E-02 |
| GPP                | Wet | 3.19E-02          | 3.52E-05          | 8.93E-03           | 4.08E-02 | 3.99E-03          | 8.79E-05          | 1.59E-03           | 5.66E-03 |
|                    | Dry | 1.30E-01          | 1.43E-04          | 3.64E-02           | 1.67E-01 | 1.63E-02          | 3.58E-04          | 6.48E-03           | 2.31E-02 |
| SPP                | Wet | 9.97E-02          | 1.10E-04          | 2.79E-02           | 1.28E-01 | 1.25E-02          | 2.75E-04          | 4.97E-03           | 1.77E-02 |
|                    | Dry | 2.86E-01          | 3.16E-04          | 8.01E-02           | 3.67E-01 | 3.58E-02          | 7.89E-04          | 1.43E-02           | 5.08E-02 |

Table 5: Total Cancer Risk of PBDEs in the Dust

| Table 3: Total Cancer Risk of PDEEs in the Dust |     |                     |                     |                      |          |                     |                     |                      |          |
|-------------------------------------------------|-----|---------------------|---------------------|----------------------|----------|---------------------|---------------------|----------------------|----------|
| Children Exposures                              |     |                     |                     |                      |          | Adults' Exposure    |                     |                      |          |
|                                                 |     | RISK <sub>Ing</sub> | RISK <sub>Inh</sub> | RISK <sub>Derm</sub> | TCR      | RISK <sub>Ing</sub> | RISK <sub>Inh</sub> | RISK <sub>Derm</sub> | TCR      |
| BPP                                             | Wet | 4.46E-06            | 4.92E-09            | 1.25E-06             | 5.71E-06 | 3.04E-07            | 6.71E-09            | 2.22E-07             | 5.33E-07 |
|                                                 | Dry | 4.96E-06            | 5.47E-09            | 1.39E-06             | 6.35E-06 | 3.38E-07            | 7.46E-09            | 2.47E-07             | 5.93E-07 |
| CPP                                             | Wet | 3.80E-06            | 4.19E-09            | 1.06E-06             | 4.87E-06 | 2.59E-07            | 5.72E-09            | 1.90E-07             | 4.54E-07 |
|                                                 | Dry | 6.27E-06            | 6.92E-09            | 1.76E-06             | 8.03E-06 | 4.27E-07            | 9.43E-09            | 3.13E-07             | 7.50E-07 |
| GPP                                             | Wet | 1.56E-06            | 1.72E-09            | 4.37E-07             | 2.00E-06 | 1.07E-07            | 2.35E-09            | 7.79E-08             | 1.87E-07 |
|                                                 | Dry | 6.37E-06            | 7.03E-09            | 1.78E-06             | 8.16E-06 | 4.34E-07            | 9.58E-09            | 3.18E-07             | 7.62E-07 |
| SPP                                             | Wet | 4.88E-06            | 5.39E-09            | 1.37E-06             | 6.26E-06 | 3.33E-07            | 7.34E-09            | 2.44E-07             | 5.84E-07 |
|                                                 | Dry | 1.40E-05            | 1.55E-08            | 3.93E-06             | 1.80E-05 | 9.56E-07            | 2.11E-08            | 6.99E-07             | 1.68E-06 |

### Pearson's Correlation Analysis

The Pearson's correlation coefficients of PBDE homologues in the dust from the PGP are shown in Table 6. The PBDE homologues correlated with each other except mono- and

hepta-BDEs that do not correlate with any other homologue. These implied that, both mono- and hepta-BDEs in the dust have different sources and/or fate in the environment.

Table 6: Pearson's Correlation Coefficient of PBDE Homologues in the Dust

|            | Mono-BDEs | Di-BDEs | Tri-BDEs | Tetra-BDEs | Penta-BDEs | Hexa-BDEs | Hepta-BDEs | Deca-BDEs |
|------------|-----------|---------|----------|------------|------------|-----------|------------|-----------|
| Mono-BDEs  | 1.00      | -0.08   | 0.30     | -0.09      | -0.04      | -0.10     | -0.08      | 0.06      |
| Di-BDEs    |           | 1.00    | 0.75**   | 0.55*      | 0.75**     | 0.82**    | -0.09      | 0.91**    |
| Tri-BDEs   |           |         | 1.00     | 0.41*      | 0.73**     | 0.55*     | -0.18      | 0.87**    |
| Tetra-BDEs |           |         |          | 1.00       | 0.22       | 0.28      | -0.05      | 0.65*     |
| Penta-BDEs |           |         |          |            | 1.00       | 0.82**    | 0.35       | 0.84**    |
| Hexa-BDEs  |           |         |          |            |            | 1.00      | 0.32       | 0.76**    |
| Hepta-BDEs |           |         |          |            |            |           | 1.00       | 0.07      |
| Deca-BDEs  |           |         |          |            |            |           |            | 1.00      |

\*\*Significant at 0.01 level of significance; \*Significant at 0.05 level of significance



### Principal Component Analysis (PCA)

The PCA of PBDEs homologues in the dusts from the power generation plants are shown in Table 7. The PCA for BDE homologues in the dust has three components accounting for 86.732 % of the total variances. Component 1 has high positive loadings for all the PBDE homologues except mono- and hepta-BDEs and explained 54.616 % of the variance. Component 1 is associated with discharges from common

plastics used in electrical and electronic equipment such as computers, televisions, air conditioners, cell phones, remote controls, computer monitors, plumbing pipes (Iwegbue *et al.*, 2025). Component 2 has high positive loadings for hepta-BDE and explained 17.560 % of the variance while component 3 has high positive loadings for mono-BDE with 14.555 % of the total variance.

**Table 7: PCA of PBDEs Homologues in the Dusts**

|                    | Component |        |        |
|--------------------|-----------|--------|--------|
|                    | 1         | 2      | 3      |
| Mono-BDE           | .019      | -.086  | .947   |
| Di-BDE             | .950      | -.062  | -.102  |
| Tri-BDE            | .866      | -.146  | .367   |
| Tetra-BDE          | .616      | -.380  | -.316  |
| Penta-BDE          | .835      | .484   | .052   |
| Hexa-BDE           | .810      | .419   | -.094  |
| Hepta-BDE          | .022      | .904   | -.104  |
| Deca-BDE           | .991      | .014   | .030   |
| Variance (%)       | 54.616    | 17.560 | 14.555 |
| Cumm. Variance (%) | 54.616    | 72.176 | 86.732 |

### CONCLUSION

The study provides information on the occurrence of PBDEs in dust samples from power generation plants in Nigeria. The occurrence pattern indicated that penta-BDEs were the dominant PBDEs in the dust. There were significant spatial and temporal variations in the PBDEs concentrations with higher concentrations observed during the dry seasons. Human risk assessment indicated the absence of health risk from exposure to these dusts. Even though the PBDEs concentrations in the dusts presently pose no health risk, we recommend that the Federal Ministry of Environment should establish environmental policy and enforceable regulatory frameworks for the control of PBDEs in order to minimize the levels environmental PBDEs and the risk to human and ecosystems.

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