

# Toxicity of Formaldehyde, Polybrominated Diphenyl Ethers (PBDEs) and Phthalates in Engineered Wood Products (EWPs) from the Perspective of the Green Approach to Materials: A Review

Ozge Cemiloglu Ulker<sup>a</sup> and Onur Ulker<sup>b,\*</sup>

Adhesives, flame-retardant chemicals, and paints are used in engineered wood products (EWPs) to increase some of the properties of wood. Most of the engineered wood composites, including plywood, particleboard, and fiberboard, used as furniture components contain formaldehyde resins as an adhesive. The International Agency for Research on Cancer (IARC) added formaldehyde to the list of human carcinogens (Group 1) in 2004. Flame-retardant chemicals are semi-volatile organic compounds that can migrate from the products to the air. There are developmental neurotoxic effects from flame-retardant additives, among which polybrominated diphenyl ethers (PBDEs) are commonly used in EWPs. The flexibility and durability of plastics are increased using phthalates, which are a class of synthetic chemicals, by adding them to the polyvinyl chloride (PVC) that is used in the wood-plastic composites (WPC). Formaldehyde, PBDEs, and phthalates are toxicants that are commonly present in value-added furniture products. This review summarized the toxic effects of these chemicals from the aspect of human health and from the perspective of green products.

*Keywords: Format; Engineered wood products; Formaldehyde; PBDEs; Phthalates*

*Contact information: a: Ankara University, School of Pharmacy, Department of Toxicology, P.O. Box 06100, Ankara, Turkey; b: Kirikkale University, Department of Interior Architecture and Environmental Design, P.O. Box 71450, Kirikkale, Turkey;*

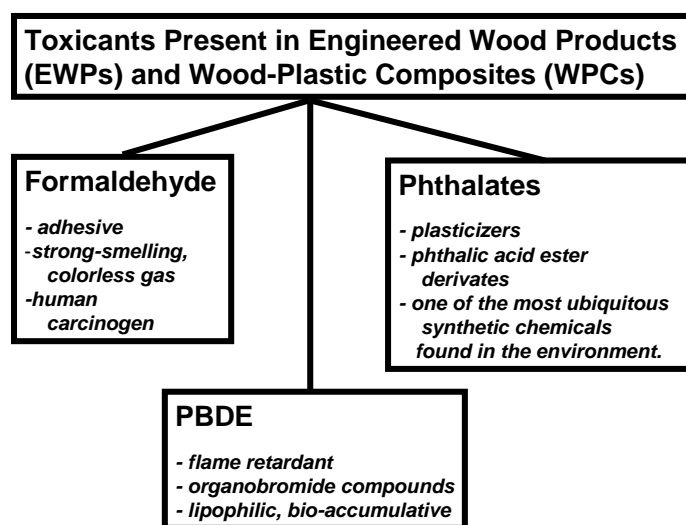
*\* Corresponding author: ulker79o@hotmail.com*

## INTRODUCTION

Engineered wood products (EWPs) can be used for different purposes in various applications. Wood composite materials are produced as an alternative to solid wood materials and provide some advantages due to both their economy and their various technical advantages. They are mainly preferred in the manufacturing of furniture products used in kitchens, bathrooms, lockers, and other places in houses and offices. The EWP industry has been significantly growing to meet the high production capacity that is a result of new technology and rising living standards. Wood-based materials are economical and easy to process at their beginning. Consequently, wood-based panels are preferred by kitchen and design companies. In addition, they can be easily, quickly, and homogeneously produced in different sizes, and the surface of the products can be layered with different colors and patterns. This property provides diversity for the desired features and aesthetics. Medium-density fiberboard (MDF), hardwood plywood, and particleboard are three types of EWPs (Milner and Woodart 2016). Wood-plastic composites (WPCs), mixtures of wood fibers with plasticizers, are one of the developing areas in the wood industry. Flame-

retardant chemicals, formaldehyde-based adhesives, and phthalates are used for enhancing overall product quality and durability. However, these chemicals are also emitted during the production process. Therefore, there is a great health concern associated with furniture manufactured from wood-based materials due to the different chemicals used in their production process.

There has been a decrease in indoor air quality (IAQ) due to the emissions in the indoor air from these compounds in furniture; the IAQ is important for human health. According to the World Health Organization (WHO), human beings spend more than 80% of their lifetime indoors, including living and working spaces (WHO 2000; Weschler 2009). The recent increase in illness frequency, such as allergies and asthma, is being attributed to indoor air pollution. Therefore, the IAQ of residential units and workplaces is a serious concern (Cincinelli and Martellini 2017). Formaldehyde, polybrominated diphenyl ethers (PBDEs), and phthalates are the widely present toxicants in EWPs and WPCs.



**Fig. 1.** The widely present toxicants in EWPs and WPCs

At room temperature, formaldehyde is a strong-smelling, colorless gas. Binders for wood composites composed of formaldehyde-based resins were first used between 1900 and 1930. The first particleboard was produced in Bremen, Germany during World War II. It has been used for furniture manufacturing since 1950 as a significant alternative to solid wood. Reports on the adverse health effects of formaldehyde started to be published in the mid-1960s (Salthammer *et al.* 2010). Formaldehyde was studied by the former International Agency of Research on Carcinogens (IARC) Working Groups in 1982, 1987, 1995, and 2004. Formaldehyde was classified as ‘probably carcinogenic’ to humans prior to 2004. In 2004, the IARC changed its decision on formaldehyde and classified it as a human carcinogen (IARC 2004).

The formaldehyde emission from EWPs and wood buildings is one of the main reasons for poor IAQ. To be able to carry out risk assessment, it is necessary to measure and control the emitted formaldehyde to the indoor air. Various standard test methods include the American ASTM D 5582 (the desiccator method), ASTM D 6007–2, and ASTM E1333 (the environmental chamber method); European EN120 (the perforator extraction method), EN 717–1 (the environmental chamber method), EN 717–2 (the gas analysis

method), EN717–3 (the flask method), ENV 13419–2 (the field and laboratory emission cell method), and the Japanese JIS A1460 (the desiccator method) are proposed for measuring formaldehyde emission by the governments (Song *et al.* 2015).

Polybrominated diphenyl ethers (PBDEs) are organobromide compounds. Since the 1970s, PBDEs have been added to EWPs as flame retardants (Vonderheide *et al.* 2008). Approximately 98% of the global market for one of PBDE congeners named penta-BDE was in North America in 1999. Additionally, penta-BDE was added in the US to furniture before 2006. Information on alternatives to penta-BDE was given in a regional survey in the US. Although alternatives are available and used to replace penta-BDE in many countries, their possible adverse effects on the environment and human health should be considered. PBDEs are persistent organic compounds; they are resistance to degradation and have lipophilic, bio accumulative properties. The studies observed the emission of PBDEs from materials that had been treated with PBDE congeners. PBDEs can bind to dust, so they are commonly found in indoor air and dust. Recently estimating PBDEs concentration in indoor dust takes much attention as a marker for making risk assessment. Chromatography-mass spectrometry (GC-MS) has been employed for detecting PBDE congeners and their concentration in dust by many researchers (Fromme *et al.* 2014; Bennett *et al.* 2015; Afafe and Martincigh 2015; Civan and Kara 2016).

Phthalates are a group of chemicals that are used for making plastics more flexible. They were developed in the 1920s, and in 1931 the phthalate industry started to expand with the development of di-2-ethylhexyl phthalate (DEHP). Studies about phthalates on health effects in the human population were first published in the 1970s. After phthalate exposure, the chemical is rapidly metabolized and excreted in urine. Scientists can estimate the quantity of phthalates that have entered bodies by detecting and measuring phthalate metabolites in urine samples. Exposure to phthalates is widespread in the US population according to the findings of the Centers for Disease Control and Prevention (CDC). Based on these findings, the US Consumer Product Safety Commission (US CPSC) proposed new phthalate requirements for children's products in 2008 (US CPSC 2008). Phthalate plasticizers are emitted from the product into the indoor air slowly based on their low volatility properties, because they are not chemically bound to the product materials (Koch *et al.* 2006). In addition, they can diffuse into water, soil, house dust, food, living organisms, and other media, especially in conditions involving heat. Phthalates are major indoor pollutants because they mainly bind to dust particles in the home. Various chamber systems such as the FLEC (Field and Laboratory Emission Cell) (Wolkoff 1996) and CLIMPAQ (Chamber for Laboratory Investigations of Materials, Pollution and Air Quality) (Gunnarsen *et al.* 1993) have been used for detecting the concentrations of phthalates congeners. It is possible to analyze different congeners in the dust by GC-MS (Clausen *et al.* 2002).

This review aims to summarize the chronic (systemic) toxic effects of these chemicals by ingestion or inhalation of the contaminated indoor air from the aspect of human health and the perspective of green products.

## CURRENT STUDIES ON FORMALDEHYDE; PBDEs AND PHTHALATES

### Chronic Toxic Effects of Formaldehyde Resins

Durable and water-resistant resins are used for gluing together many layers of wood and adhesives during the production of EWPs. Due to their low cost and high performance, urea formaldehyde, phenol formaldehyde, and melamine formaldehyde are used frequently in the wood industry. However, highly toxic formaldehyde is emitted into indoor environments during the production and post-production processes. Generally, the formaldehyde concentration in a normal home environment is 0.02 mg/m<sup>3</sup> to 0.06 mg/m<sup>3</sup> (IARC 2006). Temperature, humidity, ventilation rate, building age, use of the product, presence of flammable sources, and smoking habits affect the formaldehyde level in an indoor environment. The air formaldehyde concentration was found to be higher during the summer compared to winter due to the difference in ultraviolet (UV) light intensity and humidity. Minimal risk levels (MRLs) of inhalation exposure to formaldehyde are 0.05 ppm for acute exposure (14 days or less), 0.01 ppm for sub-chronic exposure (15 days to 364 days), and 0.003 ppm for chronic exposure (1 year or more) (ATSDR 1999). The WHO investigated the health effects of formaldehyde at different concentrations by bringing together all the research data available. Formaldehyde concentrations of 0.1 mg/m<sup>3</sup> and 0.38 mg/m<sup>3</sup> in indoor air are recommended for preventing sensory irritation and eye irritation, respectively, in the general population. The WHO reported that the highest acceptable concentration is 0.21 mg/m<sup>3</sup> to protect human health from long-term effects, including cancer (WHO 2010).

### *Toxicokinetic properties of formaldehyde*

Formaldehyde is produced in all cells as a metabolic intermediate product and is formed during the normal metabolism of glycine, serine, methionine, and choline, as well as the N-, S-, and O-methyl compound demethylation process. The endogenous formaldehyde level is multidimensional (ATSDR 1999). Formaldehyde is easily absorbed orally and *via* inhalation, but the dermal absorption is poor. Endogenous and exogenous formaldehyde, metabolized by formaldehyde dehydrogenase (FDH), is initially converted into formate and CO<sub>2</sub>, and then it is excreted in the form of metabolites (Casanova-Schmitz *et al.* 1984). Formaldehyde, which cannot be metabolized by FDH, may form cross-links between single strands of protein and DNA, or it may be incorporated into a single carbon-mediated metabolic pool through binding to tetrahydrofolate. Formaldehyde may react with proteins and nucleic acids in tissues and bind to a single strand chain to form DNA-adduct products (Bolt 1987).

### *The effect of formaldehyde on the respiratory system*

Human exposure to formaldehyde is mainly by inhalation. Formaldehyde has been suggested to show irritant effects resulting from its cross-linking to the cell DNA and proteins and from the precipitation of proteins after exposure by inhalation (ATSDR 1999).

There are conclusions about the impacts of chronic formaldehyde exposure on pulmonary parameters. In two different studies involving 0.34 ppm to 0.40 ppm and 0.1 ppm to 1 ppm formaldehyde exposure in furniture and wood enterprises, respectively, formaldehyde changed some parameters related to respiration in workers (Alexandersson and Hedenstierna 1989; Khamgaonkar and Fulare 1991). These studies showed the reduction in lung functions, such as forced expiratory volume, forced vital capacity, and flow rate. The workers inhaled 0.3 ppm (0.36 mg/m<sup>3</sup>) of formaldehyde based on these two

studies. Studies on chronic exposure to formaldehyde vapor in workplaces reported similar symptoms as coughing, shortness of breath, and chest tightness (Khamgaonkar and Fulare 1991). It was reported by the Agency for Toxic Substances and Disease Registry (ATSDR) that formaldehyde concentrations below 3 ppm do not greatly affect pulmonary function parameters and do not increase pulmonary hyperreactivity in animals or humans (ATSDR 1999). Researchers explained that because formaldehyde is mainly metabolized in the upper respiratory tract, such as the nasal mucosa, the normal level of formaldehyde exposure in the lower airways is not usually detected. It is reported that formaldehyde, which cannot be metabolized in small amounts, progresses to the trachea, bronchus, and lungs, but its effect is weak (ATSDR 1999). According to animal studies, the lesions in the respiratory system are suggested to be dose-dependent, and formaldehyde toxicity may change with the nasal defense mechanism (Wilmer *et al.* 1987, 1989).

#### *The effect of formaldehyde on the skeletal muscle system*

There are studies suggesting that chronic inhalation of formaldehyde in humans may cause mild side effects such as muscle and joint stiffness in the musculoskeletal system (Holness and Nethercott 1989). However, the occurrence of these complaints has not been associated with formaldehyde, and other unknown factors may contribute (Morgan *et al.* 1986).

#### *The effect of formaldehyde on the endocrine system*

There is little evidence that exposure to formaldehyde orally, dermally, or *via* inhalation affects the human endocrine system. It was reported in some animal studies that formaldehyde did not have any adverse effects on the endocrine system (Rusch *et al.* 1983; Vargová *et al.* 1993). However, it was found in one study that thyroid tissue weight decreased, triiodothyronine (T3) and thyroxine (T4) levels were reduced, and thyroid stimulating hormone (TSH) levels increased in rats treated with 10 mg/kg and 15 mg/kg of formaldehyde for 30 days. Additionally, it has been reported that thyroid follicle activity increased and T4 synthesis capacity is impaired and may cause atrophy with repeated exposure (Patel *et al.* 2003).

#### *The effect of formaldehyde on the eyes*

There are reports of frequent occurrence of eye irritation resulting from the chronic exposure of 0.1 ppm formaldehyde contained in indoor air (Holness and Nethercott 1987; Ritchie and Lehnen 1987).

#### *The effect of formaldehyde on the nervous system*

Men exposed to formaldehyde vapor at concentrations of 1 ppm and higher for 5.5 h may display symptoms such as headache and fatigue (Bach *et al.* 1990). Neuropsychological improvement may decrease in those showing deliberate symptoms, such as sleep, seizures, unconsciousness, and coma, following oral exposure to formaldehyde in humans (Eells *et al.* 1981; Burkhart *et al.* 1990; Köppel *et al.* 1990; Marceaux *et al.* 2008). In addition, formaldehyde at non-toxic concentrations has reduced the expression of tyrosine hydroxylase, which limits dopamine synthesis and therefore increases dopamine-like effects (Lee *et al.* 2008). There are considerable toxic effects on the nervous system from exposure to high doses of formaldehyde; however, there may be moderate effects at low doses in long-term exposure. Formaldehyde is predicted to be a possible neurotoxic agent for chronic exposure (Pitten *et al.* 2000).

### *The effect of formaldehyde on the reproductive system*

Low birth weight infants and spontaneous abortions are seen in pregnant women who are exposed to formaldehyde *via* their occupation (IARC *et al.* 2006). Birth defects have been reported in babies of two mothers who were exposed to formaldehyde from household goods (Thrasher and Kilburn 2001). Most of the current epidemiological studies report that formaldehyde in women woodworkers, laboratory staff, and beauticians has delayed conception and increased the risk of spontaneous abortion.

### *The effect of formaldehyde on the immune system*

There is conflicting data on the effects of formaldehyde exposure on the immune system in both humans and animals. Formaldehyde is suggested to cause immune system activation, and Immunoglobulin E (IgE) antibodies specific for formaldehyde were detected after inhalation in a few studies (Dykewicz *et al.* 1991; Wantke *et al.* 2000). In a school's interior air with a formaldehyde level that does not exceed 0.08 ppm, 38% of the children in the 8 years of age group had higher IgE levels (Wantke *et al.* 2000). The number of T and B lymphocytes significantly decreased in those living in furnished houses with formaldehyde release (Thrasher *et al.* 1987).

### *The effect of formaldehyde on cancer formation*

Exposure to formaldehyde increases the risk of cancer in the respiratory tract, colon, skin, bone, prostate, bile duct, liver, kidney, bladder, and lymphatic system (*e.g.*, leukemia and Hodgkin's lymphoma) (Hansen and Olsen 1995; ATSDR 1999; Luce *et al.* 2002; Collins 2004; Collins and Lineker 2004; IARC 2006; Hauptmann *et al.* 2009). Exposure to formaldehyde orally, dermally, or *via* inhalation causes carcinogenic effects in experimental animals (Dalbey 1982; Kerns *et al.* 1983; Rusch *et al.* 1983; Takahashi *et al.* 1986; Iversen 1988; Wilmer *et al.* 1989; Soffritti *et al.* 2002). The IARC classified formaldehyde as a Group 1 carcinogen (human carcinogen) because of the conclusions of the studies (IARC 2004; Cogliano *et al.* 2005). Most importantly, there is enough evidence from animal studies and strong mechanistic evidence from humans exposed to formaldehyde to classify it as a carcinogen (IARC 2004).

## **Alternatives to Formaldehyde-based Adhesives**

Recently limitations have been introduced due to the toxic effects of formaldehyde that have been mentioned above. In many countries, wood composite industries try to reduce and control formaldehyde emissions from the EWPs. All green furniture products, including salvaged, refurbished, or remanufactured furniture products, must not contain formaldehyde in concentrations greater than 50 ppb (RPN 2013).

The aim of these studies was to increase the utilization of environmentally friendly raw materials in the wood-based composite industry to eliminate or reduce the use of urea formaldehyde resin (Sulaiman *et al.* 2013). Kannerth *et al.* (2009) investigated possibly producing particleboard by using animal protein glue. Several researchers have thought to use proteins as binders for hot-pressed boards (Pizzi 2006; Müller *et al.* 2007; Migneault *et al.* 2011; Nikvash *et al.* 2012; Nasir *et al.* 2013; Pizzi 2014; Zhang *et al.* 2015).

Low-emitting resins are not formaldehyde-free, but they can be regarded as a safer alternative. Phenol-formaldehyde is the most common, low-emitting resin that is used frequently for exterior panels and other non-decorative applications. The red-black color of this resin is a disadvantage when visibly used. Additionally, formaldehyde is still emitted

in low concentrations and needs proper ventilation to maintain the near-undetectable formaldehyde concentration levels (Emery 1986).

Polymer methylene diphenyl diisocyanate (pMDI) resins are a popular alternative to formaldehyde. These resins contain polyurethanes. The Leadership in Energy and Environmental Design (LEED) recognizes MDI-based materials as "low-emitting;" however pMDI itself has resulted in clear indications of nasal toxicity (US EPA 1998). The EWPs made with MDI cannot be considered natural. Another problem with MDI resins is their considerably high prices.

Soybeans and various types of starches included among the green and environmentally friendly materials could potentially be used to produce composite panels. Several studies examined the use of starch as a binder for hot-pressed boards (Müller *et al.* 2007; Lamaming *et al.* 2013). Corn, potato, rice, wheat, and sago plants can provide starch. The chemical structure of starch, a carbohydrate, can be differentiated by the amylase and amylopectin it contains. Amylase is the linear  $\alpha$ -(1→4) linked glucan, and amylopectin is an  $\alpha$ -(1→4) linked glucan with 4.2% to 5.9%  $\alpha$ -(1→6) branch linkages. The various modifications, such as esterification, etherification, oxidation, and crosslinking of starch, have been evaluated and well-documented (El Mansouri *et al.* 2007). These modifications yield different types of starch, *e.g.*, starch xanthate, dialdehyde starch, carboxymethyl starch, and hydroxyethyl starch.

Additionally, researchers have studied the utilization of starch in the biodegradable thermoplastic field in addition to its food industry application. A main concern when using starch in panels is the dimensional stability of the test particleboard samples. Starch is a hygroscopic material even when modified by using glutaraldehyde. The water absorption and thickness swelling of the panels in the study by Amini *et al.* (2013) were unsatisfactory and did not meet the minimum requirements. A main concern when using starch in panels is that the dimensional stability of the sample were unsatisfactory and did not meet the minimum requirements of the JIS A 5908 (2003) standard (Amini *et al.* 2013). Water-repellent materials should be added to improve the dimensional stability of the panels for moisture, or the particleboards produced should be for dry condition usage only. Water absorption and thickness swelling of the samples can be enhanced through heat and chemical treatments. Better dimensional properties of the products can be obtained by using approximately 1% wax by commercial panel manufacturers (Amini *et al.* 2013). The physical and mechanical properties of formaldehyde-free plywood prepared from corn starch-tannin adhesives with phenol-formaldehyde plywood, which has already been commercialized, were compared in a study by Moubarik *et al.* (2010). They used hexamine, a non-toxic aldehyde hardener, for hardening the tannin. This yielded good and comparable mechanical properties for passing the international standards. A combination of lignin and tannin additives has additionally been suggested and has shown promise as wood panel adhesives (Pizzi 1983; Navarrete *et al.* 2010).

One of the newer natural adhesives, oil palm starch (OPS), can potentially be used in the particleboard industry. The use of OPS modified with epichlorohydrin for particleboard production was studied by Sulaiman *et al.* (2013). It can possibly be produced as a green adhesive due to the high starch content in the oil palm trunk (Noor *et al.* 2000). Epichlorohydrin is a colorless liquid that can react with many types of compounds due to the one epoxide ring and one chlorine atom in its molecular structure. A highly resistant starch is formed *via* the modification of starch with epichlorohydrin that could be used as a component for water-resistant adhesives in the paper industry or the food industry (National Toxicology Program 2011). Through the reaction of starch with epichlorohydrin,

glycerol is formed by ether linkages between the hydroxyl groups and the cross-links in the starch (Jyothi *et al.* 2007). Particleboards made with OPS with epichlorohydrin met the JIS for mechanical strength (internal bond, modulus of rupture, and modulus of elasticity) compared to those manufactured with native oil palm starch. However, the minimum requirements of the JIS were not met for the water absorption and thickness swelling. Applying wax on the surface of panels can possibly solve this problem. Abuarra *et al.* (2014) suggest that for dry applications, such as indoor furniture, gum arabic could be used as a particleboard binder. Another particleboard adhesive from oil palm trunks could be the bacterially produced natural polyesters, such as polyhydroxyalkanoates (Baskaran *et al.* 2012).

In some recent studies, the utilization of bamboo green was suggested for developing new wood composite panels. Song and coworkers investigated the effect of different bamboo green/wood fiber mixture ratio, different hot-pressing temperature, and hot pressing duration time on the physical and mechanical properties of the composite panels. The optimum ranges were found for these variables as 35% to 49%, 173 °C to 198 °C, and 111 s to 134 s, respectively. It was suggested that over the optimum ranges, raising these values negatively impacted the physical and mechanical properties of the panels (Song *et al.* 2018a,b). Hubbe *et al.* (2018) published a recent review about reconstituted lignocellulose-based materials without the addition of formaldehyde. The hypothesis was that the development of strength is based on links in the chain (molecular contact, mechanical contact, chemical bonding, and structural integrity).

### Chronic Toxic Effects of PBDEs

Additives to hinder fire and flames in EWPs, such as PBDEs, have been used. The National Health and Nutrition Examination Survey (NHANES) reported that 100% of the US population has at least one PBDE-type chemical in their blood at detectable levels (Sjödin *et al.* 2014). There are three types of commercial PBDE mixtures: deca-BDE, penta-BDE, and octa-BDE. However, the most commonly used is deca-BDE (Hites 2004; Vonderheide *et al.* 2008). Lower brominated compounds have more bioaccumulative properties than higher brominated compounds. Polybrominated diphenyl ethers release bromine radicals at high temperatures (Hooper and McDonald 2000). The ATSDR has established a minimal risk level (MRL) of 0.006 mg/m<sup>3</sup> deca-BDE for inhalation exposure at an intermediate duration. Problems in the nervous, reproductive, and endocrine systems are the main health problems following PBDE exposure. Due to the developmental toxic effects of PBDEs, children have the highest serum levels of PBDEs (Lunder *et al.* 2010). In addition, they are the most susceptible group because of frequent hand-to-mouth behavior, and they spend time on the floor (Jones-Otazo *et al.* 2005). In December 2009, the main US importer of deca-BDE guaranteed to end production, sale, and import by the end of 2013 (US EPA 2013, 2017). Octa-BDE and penta-BDE have been banned for over 15 years in the EU (EEC Directive 2003/11/EC 2003). On February 7, 2017, the EU Commission published a new regulation to add flame-retardant deca-BDE to the Registration, Evaluation, and Authorization of Chemicals (REACH) Annex XVII restricted substances' list (EU Commission Regulation 227/2012 2017). After March 2, 2019, the substances and mixtures are not allowed to be manufactured or sold in the EU market (EU Commission 2017).



### *Toxicokinetic properties of PBDEs*

Inhalation or ingestion of PBDE-contaminated household dust in an indoor air environment is the main human exposure to PBDE. The PBDEs in humans and laboratory animals are metabolized *via* oxidative hydroxylation and converted to hydroxylated PBDEs. These metabolites are shown to be in human biological fluids and breast milk (Athanasiadou *et al.* 2008; Lacorte and Ikonomou 2009; Qiu *et al.* 2009; Rydén *et al.* 2012; Wang *et al.* 2012; Yu *et al.* 2012; Butryn *et al.* 2015; Caspersen *et al.* 2016; Parry *et al.* 2018). The uptake of PBDEs from breast milk is shown in animal and human studies. Additionally, it was confirmed that PBDEs are transferred across the placenta (Frederiksen *et al.* 2010). The half-lives of PBDEs in humans are much longer than in animals. These differences make it difficult to extrapolate data from animals to humans. However, human studies support a comprehensive understanding that higher-brominated PBDEs are more easily eliminated than the lower-brominated PBDE conjugates. The concentration in the body fluids and the breakthrough rates can vary for different congeners (ATSDR 2017).

### *The effect of PBDEs on the development of the nervous system*

Fetuses and newborns are the most vulnerable to PBDE exposure. Brain development is one of the most sensitive endpoints for PBDE toxicity. Many epidemiological studies show the effect of PBDEs on the developing nervous system in infants (Chen *et al.* 2014; Herbstman and Mall 2014; Ding *et al.* 2015; Donauer *et al.* 2015; Chevrier *et al.* 2016; Martin *et al.* 2017; Vuong *et al.* 2017a,b,c; Gibson *et al.* 2018). A decreased IQ and increased occurrence of attention deficit hyperactivity disorder (ADHD) have been correlated with the PBDE concentrations in infant serum, maternal cord blood, and/or breast milk (ATSDR 2017). Additionally, animal studies and *in vitro* three-dimensional brain models report the development of neurotoxicity, including neurobehavioral changes and altered protein and gene expression levels (Hogberg *et al.* 2016; Dorman *et al.* 2018; Zhang *et al.* 2018). These animal and epidemiologic studies indicate that the development of the nervous system can be a target of concern, especially for lower-brominated PBDE exposure. Although the toxicity of PBDE exposure can potentially affect the development of the nervous system, evidence is too limited to determine the specific neurotoxic effects in adults by both human and animal studies (ATSDR 2017).

It is essential for proper neurodevelopment and healthy brain development in the fetus and early childhood to have thyroid hormones. Thyroid hormones regulate cell migration, differentiation, and proliferation during development; therefore, one of the mechanisms of the developmental neurotoxic effects of PBDEs in children may be related to developing the endocrine system (Jacobson *et al.* 2016; ATSDR 2017). Polybrominated diphenyl ethers have structural similarities to the two major thyroid hormones, thyroxine (T<sub>4</sub>), and triiodothyronine (T<sub>3</sub>) (Ibhazehiebo *et al.* 2011). When PBDEs enter organisms, they can replace the thyroid hormones, which can result in hypothyroidism. Severe hypothyroidism (*e.g.*, decreased T<sub>3</sub> and/or T<sub>4</sub> levels) in young children can lead to many neurological problems, such as poor auditory, fine motor, and executive processing skills, language and memory deficits, and intellectual disabilities (Porterfield and Hendrich 1993; Zoeller and Rovet 2004; Williams 2008). The potential for PBDEs and their metabolites to interfere with thyroid function has been the topic of extensive research in recent years. There are few epidemiological studies on children (Gascon *et al.* 2011; Xu *et al.* 2014). In a study that was conducted in the USA between 2011 and 2012 by Jacobson *et al.* (2016), thyroid hormones were evaluated in serum samples of 80 children (ages 1 to 5) from the

southeastern United States. Serum levels of the seven PBDE congeners were also measured in this group. Their results suggest that exposure to PBDEs during childhood sub-clinically changes thyroid hormone function, and the disruption can cause hypothyroidism. There are many animal studies compatible with this study that show a decrease in serum T4 and/or T3 in puppies or other offspring following gestational or lactational exposure to PBDEs (Thuvander and Darnerud 1999; Hallgren and Darnerud 2002; Zhou *et al.* 2002; Stoker *et al.* 2004). According to this data, the developing thyroid can be a target of concern, especially for lower-brominated PBDE exposure.

#### *The effect of PBDEs on the reproductive system*

In limited studies, reproductive effects have been reported in men associated with PBDE exposure. Akutsu *et al.* (2008) reported a significant correlation between increased serum HxBDE-153 one of the PBDE congeners concentration and reduced sperm concentration ( $r = -0.841$ ,  $p = 0.002$ , Fig. 2) and reduced testis size ( $r = -0.764$ ,  $p = 0.01$ ). On the other hand, Albert and co-workers found no associations between human adult exposure to phthalates and sperm concentration (Albert *et al.* 2018). In a prospective cohort study in Michigan and Texas, the altered parameters of semen quality were found to be associated with BDEs (Robledo *et al.* 2015). Exposure to PBDEs during childhood may disrupt the reproductive system development, with the male reproductive system being a target of concern, especially for lower-brominated PBDE exposure in the light of the adequate animal data. However, there is not enough data available to determine whether PBDE exposure in children will cause reproductive function alterations. In contrast, the data for the female reproductive system in human and animal studies is inconsistent (ATSDR 2017).

#### *The effect of PBDEs on cancer formation*

The risk of cancer associated with PBDE exposure by inhalation or ingestion has been investigated in many human cohort and animal studies. These studies suggest that there is no evidence in humans and limited evidence in animals for the carcinogenicity of PBDEs (ATSDR 2017). There are limited case-control studies. Hoffman *et al.* (2017) suggested that exposure to flame retardants in the home, particularly BDE-209 and Tris(2-chloroethyl) phosphate (TCEP), may be associated with the occurrence of papillary thyroid cancer. He *et al.* (2018) reported that exposure to PBDE may play a role in the occurrence and development of breast cancer. The animal studies indicate a significant increase in incidences of hepatocellular adenomas and carcinomas in mice and neoplastic liver nodules in rats ( $P \leq 0.01$ ). (Dunnick *et al.* 2018).

### **Alternatives to PBDEs**

Due to the health concerns of PBDEs and the limitations for manufacturers, there is a growing interest in new, flame-retardant chemicals with inherent flame resistance that do not contain halogen or any additives. Some furniture companies are requesting PBDE-free polyurethane foam from their manufacturers. The Environmental Protection Agency (EPA) published a final report in 2014 titled "An Alternatives Assessment for the Flame Retardant Decabromodiphenyl Ether (DecaBDE)." In this report, the EPA presents detailed hazard information for 29 alternative substances (US EPA 2014). Unfortunately, the long-term fate of these alternative chemicals in the environment is not yet known. It is suggested that halogenated polymer compounds can generate halogenated dioxin and furan during combustion.

Recently, interest in proposing environmentally friendly and safer flame retardants has increased. The development of flame retardants from renewable resources has become more important in the past five years (Sonnier *et al.* 2018).

The German government determined that ammonium polyphosphate, aluminum trihydroxide, and red phosphorus are less problematic in the environment. Inorganic flame-retardant compounds, such as phosphate compounds (*e.g.*, ammonium polyphosphate, di-ammonium phosphate, and melamine phosphate), boron compounds (*e.g.*, boric acid, borax, and boric oxide), nitrogen compounds, and hydroxide compounds, of aluminum or magnesium are thought to be environmentally friendly. During combustion, these inorganic flame retardants do not release dioxins or halogen acids as by-products. Therefore, these chemicals are becoming much more practical due to being safer alternatives in the case of a fire (Sharma *et al.* 2013; Özdemir *et al.* 2017).

The presence of N-P bonds in the phosphate compounds (*e.g.*, di-ammonium phosphate, ammonium polyphosphate, and melamine phosphate) makes them thermally stable. Volatile constituents are formed through the pyrolyzation of nitrogen compounds, and when they reach the gas phase, they act as free radical interceptors. A combination of phosphorus and nitrogen, compared to only phosphorus compounds or nitrogen compounds, can provide strong flame retardancy. During combustion, the nitrogen and phosphorus combination covers the outer layer of the substrate with a nonflammable char (Jin *et al.* 2017).

Boron salts are used as the major non-fixing flame retardant for wood products. They are water-soluble, and the water evaporates after treatment, leaving the salts inside of the wood material. Boric acid acts *via* both physical and chemical mechanisms to impart flame resistance. Physically, boric acid is formed by boron derivatives and first prevents oxygen diffusion, then it prevents the exothermic combustion reactions from propagation. Chemically, boric acid increases the amount of char formed on acid-catalyzed dehydration reactions during wood pyrolysis (Yu *et al.* 2017). Compounds containing boron act with an endothermic, stage-wise water release. When these compounds are heated, the mixture dissolves in its own water of hydration. Then, it swells to form a frothy substance before losing water and finally fusing into a clear melt. Adequate protection is given to cellulosic elements with this treatment system in protected non-ground contact situations. Flame retardants containing boron have been developed as alternatives to traditional flame retardants (*e.g.*, antimony oxide) because they are cheaper and less toxic. Another flame retardant used frequently is zinc borate. It produces non-volatile products in the presence of flame, promotes char formation, and releases water (Sharma *et al.* 2013; Jin *et al.* 2017; Terzi *et al.* 2018). The primary toxic health effect associated with inhalation exposure of humans to boron compounds is acute respiratory irritation in short exposure durations (up to 47 min) (ATSDR 2010). On the other hand, boron compounds were classified as toxic to reproduction (Cat 2) by the European Union, which means that they may impair fertility and may cause harm to the fetus. This classification based on the oral animal exposure studies (SCCS 2010).

### Chronic Toxic Effects of Phthalates

There is a mixture of wood, thermoplastics, and some additives in WPCs with a wood content between 50 wt% to 80 wt% (Clemons 2002). Recently, wood fiber reinforced polyvinyl chloride (PVC) has become more popular due to its acceptable mechanical properties, long lifetime, moisture and fungus resistance, recyclability, and wood-like surface performance (Clemons 2002). PVC is suggested as a thermoplastic resin to urea-

formaldehyde due to the adhesive properties (Song *et al.* 2017). PVC is not a thermally stable polymer, but dimethyl phthalate can be used to add thermoplasticity to the wood. Phthalates are a class of synthetic chemicals commonly used in various consumer products to increase the flexibility and durability of plastics by adding it to PVC. In addition, they are popular due to their relatively low cost, low volatility, and ability to create elastic materials. There have been concerns about their effects on the environment and human health since the early 1980s. Currently, phthalates are one of the most ubiquitous synthetic chemicals found in the environment. Over 11 billion pounds of phthalates are annually produced worldwide. Phthalates are derived from phthalic acid esters. The seven phthalates extensively used in consumer products are benzyl butylphthalate (BBP or BzBP), dibutyl phthalate (DBP), di-(2-ethylhexyl) phthalate (DEHP), diisodecyl phthalate (DIDP), diisononyl phthalate (DINP), dioctyl phthalate (DnOP), and di-n-butyl phthalate (DnBP) (The Lowell Center for Sustainable Production 2011). Different phthalate esters have different uses due to their different chemical and physical properties. Most of the DEHP and BBzP produced are used in PVC products, *e.g.*, PVC flooring. The DnBP is widely used as a plasticizer in cellulose plastics, in latex adhesives, and as a solvent for certain dyes (Bornehag *et al.* 2005)

Exposure to phthalates has been associated with several adverse human health effects. They can impact the fetus developing *in utero* by crossing the placental barrier. Values can reach as high as 70 µg/kg per day for daily intake of phthalates (Net *et al.* 2015). Many phthalates are classified as reproductive and developmental toxicants, and they may also have effects as endocrine disrupters. The US EPA classifies BBP and DEHP as possible and probable human carcinogens, respectively. However, their toxicity varies depending on the structure of phthalate (US EPA 2012).

#### *Toxicokinetic properties of phthalates*

The inhalation of a contaminated indoor air environment and the ingestion of contaminated food during packaging are the main human exposures to phthalates. In addition, young children are exposed to phthalates through childcare products, teething toys, and other products containing phthalates. Studies suggest that there is 100% oral absorption because phthalates, such as DEHP and DINP, can dissolve in saliva and become absorbed during mouthing teething (Müller *et al.* 2003). The main metabolites of phthalates are DEHP, monobutyl phthalate (MBP), mono-benzylphthalate (MBzP), mono-(3-carboxypropyl) phthalate (MCP), mono-ethylphthalate (MEP), mono-(2-ethyl-5-carboxypentyl) phthalate (MECPP), mono-(2-ethylhexyl) phthalate (MEHP), mono-(2-ethyl-5-hydroxyhexyl) phthalate (MEHHP), mono-(2-ethyl-5-oxohexyl) phthalate (MEOHP), and mono-isobutyl phthalate (MiBP) (Frederiksen *et al.* 2007). Higher lipophilic compounds have long alkyl chains and can easily enter body systems (Koch *et al.* 2004, 2005, 2012, and 2013; Koch and Angerer 2007; Koch and Calafat 2009; Leng *et al.* 2014). Glucuronidation helps phthalate metabolites to be excreted *via* urine (Silva *et al.* 2003; Samandar *et al.* 2009).

#### *The effect of phthalates on the liver system*

According to a study by Ganning *et al.* (1984), phthalates change the function and structure of the liver. Phthalates can induce mitochondria, enzymes, and peroxisomes that participate in  $\beta$ -oxidation and fatty acid transport (Ganning *et al.* 1984). The prolonged administration of phthalate esters causes an accumulative effect on the liver (Ganning *et al.* 1987; Beliles *et al.* 1989).

### *The effect of phthalates on the reproductive system*

Certain phthalates are reproductive toxins in the male groups of animal studies, but the human studies are more limited. In a human study, an inverse association between urine MBP levels and sperm motility and concentration was reported among 168 men (Duty *et al.* 2003). This result was confirmed in a follow-up study that included an additional 295 men (Hauser *et al.* 2006). In another study, workers were exposed to high concentrations of DEHP and DBP in a PVC flooring plant. The levels of these phthalate metabolites in urine samples were inversely associated with free testosterone levels, which means a high urine metabolite level results in decreasing free testosterone levels (Fong *et al.* 2015). In a case-control study, a female group that was diagnosed with endometriosis (n = 92) between 1996 and 2006 was compared with a population-based control group (n = 195) in the Pacific Northwest of the US. Endometriosis is a hormonally mediated disease among reproductive-aged females. In this study, urine phthalate metabolite concentrations measured in the cases and in population-based controls showed that exposure to phthalates might increase the risk of endometriosis (Upson *et al.* 2013).

### *The effect of phthalates on the endocrine system*

It was demonstrated that phthalates can affect the thyroid signaling system in experimental animals, but the human studies with adults remain limited. In a human study conducted at the Massachusetts General Hospital between 2000 and mid-2004, the urine MEHP levels among 408 men were inversely associated with free T4 and total T3, but not TSH (Meeker *et al.* 2007). In girls and women, phthalate exposure is additionally associated with altered thyroid function (Meeker and Ferguson 2011).

### *The effect of phthalates on the development of the endocrine system*

The relationship between the urine concentration of seven different phthalates and thyroid hormones was investigated in a cohort study on children. No association was found between the urinary phthalate metabolites and TT<sub>4</sub>, FT<sub>4</sub>, and TSH in boys. However, a significant negative association was found between T<sub>3</sub> and phthalate metabolites in girls (P < 0.05). In 845 children (both girls and boys), a significant inverse association was found between the urinary phthalate metabolites and TT<sub>3</sub> and FT<sub>3</sub> (P < 0.05) (Boas *et al.* 2010). Additionally, DEHP is significantly related with insulin resistance in Mexican American adolescents (P < 0.02) (Trasande *et al.* 2013). In a cohort study of participating mothers and children (n = 345) at the Center for the Health Assessment of Mothers and Children of Salinas (CHAMACOS), the 10 urinary phthalate metabolite concentrations were measured twice in the mothers during pregnancy and in their children between 5 and 12 years of age as well. According to their results, exposure to certain phthalates *in utero* is associated with the risk for childhood obesity and increased BMI during childhood (Harley *et al.* 2017).

### *The effect of phthalates on the development of the nervous system*

Higher levels of DEHP, DBP, and BBzP have been found in the urine samples of children. Children throughout the world are exposed to phthalates and therefore are at risk. For example, DBP is strongly associated with ADHD (Garner *et al.* 2013). Additionally, urine phthalate metabolites (*e.g.*, MMP, MEP, MBP, MBzP, MEHP, 5-oxo-MEHP, and 5-OH-MEHP) are related to a decrease in neurocognitive functions and intelligence through postnatal exposure (Sathyanarayana *et al.* 2008). One study suggests that DEHP metabolites can impact attention deficit disorder (ADD), and that girls are effected by both

ADD and learning defects more than boys (Diamond 2015). In addition, DEHP metabolites are shown to be related to autism (Testa *et al.* 2012).

### **Alternatives to Phthalates**

Several compounds, including citrates and phosphates, have been identified as alternatives to phthalates. However, the potential effects of these alternative plasticizers on human health and the environment have not yet been studied thoroughly. These compounds are not chemically bound to polymers and can be emitted from the products into the air. Some of these plasticizers may cause eye, skin, and respiratory irritation. However, most of the data are from animal studies. The studies are new, and only a few epidemiologic studies have been conducted on these materials; thus, it is hard to predict their long-term health effects. The animal studies show that these materials can have toxic effects on the kidneys, liver, spleen, testes, and uterus, meaning that phthalate-free WCPs are not considered safe (Van Vliet *et al.* 2011).

Recently, the technology for using natural-based plasticizers has rapidly grown due to their low toxicity and low accumulation properties. This group of plasticizers contains epoxidized triglyceride vegetable oils from linseed oil, soybean oil, sunflower oil, castor oil, or fatty acid esters (Baltacioglu and Balkose 1999; Pedersen *et al.* 2008). Researchers and industries have increased their work to develop new, bio-based materials from renewable and biodegradable sources. This is related to the search for natural-based plasticizers to reduce the use of conventional plastic products (Vieira *et al.* 2011). Bioplastics are sourced from bioresources (*i.e.*, a renewable resource) and/or are biodegradable. Among the different bioplastics, polyhydroxyalkanoates (PHAs) have received considerable attention as new biopolymers due to their high functionality. In addition, they are truly biodegradable and are non-toxic to the environment (Vandi *et al.* 2018). They are microbiologically produced, and the choice of substrate, bacteria, and fermentation conditions determines their material properties, ranging from rigid thermoplastics to stretchy elastomers (Dietrich *et al.* 2017). For the manufacture of WPCs, PHA biopolymers are considered particularly suitable. Initial developments on wood-PHA composites indicate that properties like commercially available PVC-based WPCs are feasible (Chan *et al.* 2016, 2017, and 2018). The use of naturally based polymer films depends on several parameters, including mechanical properties such as resistance to water, barrier requisites (*e.g.*, water vapor), O<sub>2</sub> and CO<sub>2</sub> permeability, flexibility and strength, optical quality (*e.g.*, gloss and opacity), and other issues like cost, functional attributes, and availability.

### **CLOSING COMMENTS**

The emission of formaldehyde, PBDEs, and phthalates from wood-based building products and EWPs is one of the main reasons of poor IAQ. There are various test methods and mathematical methods used for estimating the emission. These calculations are required for making risk assessment. If we can capture the relevant information about exposure assessment, then we may be able to utilize the knowledge to predict the possibility of toxicity.

The use of these chemicals, which are associated with health and environmental concerns, can be minimized by using green alternatives that require fewer or no chemicals using some surface modification techniques and by using natural-based adhesives or bio-

plasticizers that have a lower toxicity. This approach recently has motivated research on the development of natural-based EWPs in various academic and industrial areas. The use of low-toxicity EWPs has become more attractive. However, both the mechanical and physical properties of natural-based EWPs still need to be improved.

As stated for flame-retardant compounds, alternatively produced chemicals may not always be safe, especially with long-term exposure. Regulations for these chemicals should be reevaluated by considering the toxic effects.

The production and consumption of MDF and particleboard that contains formaldehyde resin is quite common in both developing and undeveloped countries. This situation causes health concerns, especially in children and adults. The use of green alternatives in place of these preferred products will be an important step in the protection of public health.

Children are more susceptible to the toxic effects of chemicals that are used in EWPs. Long-term exposure to these chemicals during fetal and child development may cause permanent alterations in the nervous system, endocrine system, and reproductive system.

The amount of toxic chemicals in recycled products is increasing during the recycling processes for EWPs and WPCs produced by traditional methods. Green products are a good alternative to avoid this situation. One solution is using recycling processes in the furniture industry.

There are many parameters that are affected by the mechanical and physical properties of naturally based EWPs. These parameters are classified into four classes and called “links” by Hubbe *et al.* (2018). By changing these parameters, it is possible to manufacture natural-based EWPs that meet the requirements stated in international standards.

In the future due to environmental and toxicological concerns will become increasingly important, and the production and usage of green alternatives to these chemicals will rapidly increase.

## ACKNOWLEDGEMENTS

The authors wish to thank Kırıkkale University (Kırıkkale, Turkey), the Department of Interior Architecture and Environmental Design, Oklahoma State University (Stillwater, USA), the Department of Natural Resource Ecology and Management (NREM), and Ankara University, School of Pharmacy (Ankara, Turkey), the Department of Toxicology.

## REFERENCES CITED

- Abafe, O. A., and Martincigh, B. S. (2015). “Polybrominated diphenyl ethers and polychlorinated biphenyls in indoor dust in Durban, South Africa,” *Indoor Air* 25(5), 547-556. DOI: 10.1111/ina.12168
- Abuarra, A., Hashim, R., Bauk, S., Kandaiya, S., and Tousi, E. T. (2014). “Fabrication and characterization of gum Arabic bonded *Rhizophora* spp. particleboards,” *Mater. Design* 60, 108-115. DOI: 10.1016/j.matdes.2014.03.032

- Akutsu, K., Takatori, S., Nozawa, S., Yoshiike, M., Nakazawa, H., Hayakawa, K., Makino, T., and Iwamoto, T. (2008). "Polybrominated diphenyl ethers in human serum and sperm quality," *Bull. Environ. Contam. Toxicol.* 80, 345-350. DOI: 10.1007/s00128-008-9370-4
- Albert, O., Huang, J. Y., Aleksa, K., Hales, B. F., Goodyer, C. G., Robaire, B., Chevrier, J., and Chan, P. (2018). "Exposure to polybrominated diphenyl ethers and phthalates in healthy men living in the greater Montreal area: A study of hormonal balance and semen quality," *Environ. Int.* 116, 165-175. DOI: 10.1016/j.envint.2018.04.012
- Alexandersson, R., and Hedenstierna, G. (1989). "Pulmonary function in wood workers exposed to formaldehyde: A prospective study," *Arch. Environ. Health* 44(1), 5-11. DOI: 10.1080/00039896.1989.9935865
- Amini, M. H. M., Hashim, R., Hizirolu, S., Sulaiman, N. S., and Sulaiman, O. (2013). "Properties of particleboard made from rubberwood using modified starch as binder," *Compos. Part B-Eng.* 50, 259-264. DOI: 10.1016/j.compositesb.2013.02.020
- Athanasiadou, M., Cuadra, S. N., Marsh, G., Bergman, A., and Jakobsson, K. (2008). "Polybrominated diphenyl ethers (PBDEs) and bioaccumulative hydroxylated PBDE metabolites in young humans from Managua, Nicaragua," *Environ. Health Persp.* 116(3), 400-408. DOI: 10.1289/ehp.10713
- Bach, B., Pedersen, O. F., and Molhave, L. (1990). "Human performance during experimental formaldehyde exposure," *Environ. Int.* 16(2), 105-113. DOI: 10.1016/0160-4120(90)90150-5
- Baltacıoğlu, H., and Balköse, D. (1999). "Effect of zinc stearate and/or epoxidized soybean oil on gelation and thermal stability of PVC-DOP plastigels," *J. Appl. Polym. Sci.* 74(10), 2488-2498. DOI: 10.1002/(SICI)1097-4628(19991205)74:10<2488::AID-APP18>3.0.CO;2-B
- Baskaran, M., Hashim, R., Said, N., Raffi, S. M., Balakrishnan, K., Sudesh, K., Sulaiman, O., Arai, T., Kosugi, A., Mori, Y., *et al.* (2012). "Properties of binderless particleboard from oil palm trunk with addition of polyhydroxyalkanoates," *Compos. Part B-Eng.* 43(3), 1109-1116. DOI: 10.1016/j.compositesb.2011.10.008
- Beliles, R., Salinas, J. A., and Kluwe, W. M. (1989). "A review of di(2-ethylhexyl) phthalate (DEHP) risk assessments," *Drug Metab. Rev.* 21(1), 3-12. DOI: 10.3109/03602538909029952
- Bennett, D. H., Moran, R. E., Wu, X., Tolve, N. S., Clifton, M. S., Colón, M., Weathers, W., Sjödin, A., and Hertz-Picciotto, I. (2015). "Polybrominated diphenyl ether (PBDE) concentrations and resulting exposure in homes in California: Relationships among passive air, surface wipe and dust concentrations, and temporal variability," *Indoor Air* 25(2), 220-229. DOI: 10.1111/ina.12130
- Boas, M., Frederiksen, H., Feldt-Rasmussen, U., Skakkebaek, N. E., Hegedus, L., Hilsted, L., Juul, A., and Main, K. M. (2010). "Childhood exposure to phthalates: Associations with thyroid function, insulin-like growth factor I, and growth," *Environ. Health Persp.* 118(10), 1458-1464. DOI: 10.1289/ehp.0901331
- Bolt, H. M. (1987). "Experimental toxicology of formaldehyde," *J. Cancer Res. Clin.* 113(4), 305-309. DOI: 10.1007/BF00397713
- Bornhag, C. G., Lundgren, B., Weschler, C. J., Sigsgaard, T., Hagerhed-Engman, L., and Sundell, J. (2005). "Phthalates in indoor dust and their association with building characteristics," *Environ. Health Persp.* 113(10), 1399-1404. DOI: 10.1289/ehp.7809
- Burkhart, K. K., Kulig, K. W., and McMartin, K. E. (1990). "Formate levels following a formalin ingestion," *Vet. Hum. Toxicol.* 32(2), 135-137.



- Butryn, D. M., Gross, M. S., Chi, L. H., Schecter, A., Olson, J. R., and Aga, D. S. (2015). “‘One-shot’ analysis of polybrominated diphenyl ethers and their hydroxylated and methoxylated analogs in human breast milk and serum using gas chromatography-tandem mass spectrometry,” *Anal. Chim. Acta* 892, 140-147. DOI: 10.1016/j.aca.2015.08.026
- Casanova-Schmitz, M., David, R. M., and Heck, H. D. (1984). “Oxidation of formaldehyde and acetaldehyde by NAD<sup>+</sup>-dependent dehydrogenases in rat nasal mucosal homogenates,” *Biochem. Pharmacol.* 33(7), 1137-1142. DOI: 10.1016/0006-2952(84)90526-4
- Caspersen, I. H., Kvale, H. E., Haugen, M., Brantsæter, A. L., Meltzer, H. M., Alexander, J., Thomsen, C., Frøshaug, M., Bremnes, N. M., Broadwell, S. L., *et al.* (2016). “Determinants of plasma PCB, brominated flame retardants, and organochlorine pesticides in pregnant women and 3 year old children in The Norwegian Mother and Child Cohort Study,” *Environ. Res.* 146, 136-144. DOI: 10.1016/j.envres.2015.12.020
- Chan, C. M., Johansson, P., Magnusson, P., Vandi, L. J., Arcos-Hernandez, M., Halley, P., Laycock, B., Pratt, S., and Werker, A. (2017). “Mixed culture polyhydroxyalkanoate-rich biomass assessment and quality control using thermogravimetric measurement methods,” *Polym. Degrad. Stabil.* 144, 110-120. DOI: 10.1016/j.polymdegradstab.2017.07.029
- Chan, C. M., Vandi, L. J., Pratt, S., Halley, P., Richardson, D., Werker, A., and Laycock, B. (2016). “Processing and characterisation of polyhydroxyalkanoate (PHA)-based wood plastic composites: Effect of non-reactive additives,” *Appita J.* 69(4), 352-360. DOI: 10.1016/j.polymdegradstab.2017.07.029
- Chan, C. M., Vandi, L. J., Pratt, S., Halley, P., Richardson, D., Werker, A., and Laycock, B. (2018). “Mechanical performance and long-term indoor stability of polyhydroxyalkanoate (PHA)-based wood plastic composites (WPCs) modified by non-reactive additives,” *Eur. Polym. J.* 98, 337-346. DOI: 10.1016/j.eurpolymj.2017.11.041
- Chen, A., Yolton, K., Rauch, S. A., Webster, G. M., Hornung, R., Sjödin, A., Dietrich, K. N., and Lanphear, B. P. (2014). “Prenatal polybrominated diphenyl ether exposures and neurodevelopment in U.S. children through 5 years of age: The HOME study,” *Environ. Health Persp.* 122(8), 856-862. DOI: 10.1289/ehp.1307562
- Chevrier, C., Warembourg, C., Le Maner-Idrissi, G., Lacroix, A., Dardier, V., Le Sourn-Bissaoui, S., Rouget, F., Monfort, C., Gaudreau, E., Mercier, F., *et al.* (2016). “Childhood exposure to polybrominated diphenyl ethers and neurodevelopment at six years of age,” *NeuroToxicology* 54, 81-88. DOI: 10.1016/j.neuro.2016.03.002
- Cincinelli, A., and Martellini, T. (2017). “Indoor air quality and health,” *Int. J. Env. Res. Pub. He.* 14(11), 1286. DOI: 10.3390/ijerph14111286
- Civan, M. Y., and Kara, U. M. (2016). “Risk assessment of PBDEs and PAHs in house dust in Kocaeli, Turkey: Levels and sources,” *Environmental Science and Pollution Research*, 23(23), 23369-23384. DOI: 10.1007/s11356-016-7512-5
- Clausen, P. A., Hansen, V., Gunnarsen, L., Afshari, A., and Wolkoff, P. (2002). “Emission of phthalates from PVC flooring in two very different test chambers,” *Proceedings: Indoor Air 2*, 932-937.
- Clemons, C. (2002). “Wood-plastic composites in the United States: The interfacing of two industries,” *Forest Prod. J.* 52(6), 10-18.
- Cogliano, V. J., Grosse, Y., Baan, R. A., Straif, K., Secretan, M. B., and El Ghissassi, F. (2005). “Meeting report: Summary of IARC Monographs on formaldehyde, 2-

- butoxyethanol, and 1-*tert*-butoxy-2-propanol,” *Environ. Health Persp.* 113, 1205-1208. DOI: 10.1289/ehp.7542
- Collins, J. J. (2004). “Formaldehyde exposure and leukemia,” *Occup. Environ. Med.* 61(11), 875-876. DOI: 10.1136/oem.2004.014324
- Collins, J. J., and Lineker, G. A. (2004). “A review and meta-analysis of formaldehyde exposure and leukemia,” *Regul. Toxicol. Pharm.* 40(2), 81-91. DOI: 10.1016/j.yrtph.2004.04.006
- Dalbey, W. E. (1982). “Formaldehyde and tumors in hamster respiratory tract,” *Toxicology* 24(1), 9-14. DOI: 10.1016/0300-483X(82)90058-0
- Diamond, A. (2015). “Attention-deficit disorder (attention-deficit/hyperactivity disorder without hyperactivity): A neurobiologically and behaviorally distinct disorder from attention-deficit/hyperactivity disorder (with hyperactivity),” *Dev. Psychopathol.* 17(3), 807-825. DOI: 10.1017/S0954579405050388
- Dietrich, K., Dumont, M. J., Del Rio, L. F., and Orsat, V. (2017). “Producing PHAs in the bioeconomy—Towards a sustainable bioplastic,” *Sustainable Production and Consumption* 9, 58-70. DOI: 10.1016/j.spc.2016.09.001
- Ding, G., Yu, J., Cui, C., Chen, L., Gao, Y., Wang, C., Zhou, Y., and Tian, Y. (2015). “Association between prenatal exposure to polybrominated diphenyl ethers and young children's neurodevelopment in China,” *Environ. Res.* 142, 104-111. DOI: 10.1016/j.envres.2015.06.008
- Donauer, S., Chen, A., Xu, Y., Calafat, A. M., Sjodin, A., and Yolton, K. (2015). “Prenatal exposure to polybrominated diphenyl ethers and polyfluoroalkyl chemicals and infant neurobehavior,” *J. Pediatr.* 166(3), 736-742. DOI: 10.1016/j.jpeds.2014.11.021
- Dorman, D. C., Chiu, W., Hales, B. F., Hauser, R., Johnson, K. J., Mantus, E., Martel, S., Robinson, K. A., Rooney, A. A., Rudel, R., *et al.* (2018). “Polybrominated diphenyl ether (PBDE) neurotoxicity: A systematic review and meta-analysis of animal evidence,” *J. Toxicol. Env. Heal. B* 21(4), 269-289. DOI: 10.1080/10937404.2018.1514829
- Dunnick, J. K., Pandiri, A. R., Merrick, B. A., Kissling, G. E., Cunny, H., Mutlu, E., Waidyanatha, S., Sills, R., Hong, H. L., Ton, T. V., *et al.* (2018). “Carcinogenic activity of pentabrominated diphenyl ether mixture (DE-71) in rats and mice,” *Toxicology Reports* 5, 615-624. DOI: 10.1016/j.toxrep.2018.05.010
- Duty, S. M., Silva, M. J., Barr, D. B., Brock, J. W., Ryan, L., Chen, Z., Herrick, R. F., Christiani, D. C., and Hauser, R. (2003). “Phthalate exposure and human semen parameters,” *Epidemiology* 14(3), 269-277. DOI: 10.1097/01.EDE.0000059950.11836.16
- Dykewicz, M. S., Patterson, R., Cugell, D. W., Harris, K. E., and Wu, A. F. (1991). “Serum IgE and IgG to formaldehyde human serum albumin: Lack of relation to gaseous formaldehyde exposure and symptoms,” *J. Allergy Clin. Immun.* 87(1, Part 1), 48-57. DOI: 10.1016/0091-6749(91)90212-7
- Eells, J. T., McMartin, K. E., Black, K., Virayotha, V., Tisdell, R. H., and Tephly, T. R. (1981). “Formaldehyde poisoning: Rapid metabolism to formic acid,” *JAMA* 246(11), 1237-1238. DOI: 10.1001/jama.1981.03320110049029
- El Mansouri, N. E., Pizzi, A., and Salvado, J. (2007). “Lignin-based polycondensation resins for wood adhesives,” *J. Appl. Polym. Sci.* 103(3), 1690-1699. DOI: 10.1002/app.25098

- Emery, J. A. (1986). "Formaldehyde release from wood panel products bonded with phenol formaldehyde adhesives," in: *Formaldehyde Release from Wood Products*, B. Meyer, B. A. K. Andrews, and R. M. Reinhardt (eds.), ACS Publications, Washington, D.C., pp. 26-39.
- European Commission (EC) Regulation 227/2012 (2017). "Commission Regulation (EU) No. 227/2017 of 9 February 2017 amending Annex XVII to Regulation (EC) No 1907/2006 of the European Parliament and of the Council concerning the Registration, Evaluation, Authorization and Restriction of Chemicals (REACH) as regards bis(pentabromophenyl)ether," European Union, Brussels, Belgium.
- European Commission (EC) Scientific Committee on Consumer Safety (SCCS) (2010) "Opinion on Boron compounds" European Union, Brussels, Belgium
- European Economic Community (EEC) Directive 2003/11/EC (2003). "Directive (EEC) 2003/11/EC of the European Parliament and of the Council of 6 February 2003 amending for the 24th time Council Directive 76/769/EEC relating to restrictions on the marketing and use of certain dangerous substances and preparations (pentabromodiphenyl ether and octabromodiphenyl ether)," European Union, Brussels, Belgium.
- Fong, J. P., Lee, F. J., Lu, I. S., Uang, S. N., and Lee, C. C. (2015). "Relationship between urinary concentrations of di(2-ethylhexyl) phthalate (DEHP) metabolites and reproductive hormones in polyvinyl chloride production workers," *Occup. Environ. Med.* 72(5), 346-353. DOI: 10.1136/oemed-2014-102532
- Frederiksen, H., Skakkebaek, N. E., and Andersson, A. M. (2007). "Metabolism of phthalates in humans," *Mol. Nutr. Food Res.* 51(7), 899-911. DOI: 10.1002/mnfr.200600243
- Frederiksen, M., Vorkamp, K., Mathiesen, L., Mose, T., and Knudsen, L. E. (2010). "Placental transfer of the polybrominated diphenyl ethers BDE-47, BDE-99 and BDE-209 in a human placenta perfusion system: An experimental study," *Environ. Health-Glob.* 9(1), 32-41. DOI: 10.1186/1476-069X-9-32
- Fromme, H., Hilger, B., Kopp, E., Miserok, M., and Völkel, W. (2014). "Polybrominated diphenyl ethers (PBDEs), hexabromocyclododecane (HBCD) and 'novel' brominated flame retardants in house dust in Germany," *Environment International* 64, 61-68. DOI: 10.1016/j.envint.2013.11.017
- Ganning, A. E., Brunk, U., and Dallner, G. (1984). "Phthalate esters and their effect on the liver," *Hepatology* 4(3), 541-547. DOI: 10.1002/hep.1840040331
- Ganning, A. E., Brunk, U., Edlund, C., Elhammer, A., and Dallner, G. (1987). "Effects of prolonged administration of phthalate ester on the liver," *Environ. Health Persp.* 73, 251-258. DOI: 10.1289/ehp.8773251
- Garner, A. A., O'Connor, B. C., Narad, M. E., Tamm, L., Simon, J., and Epstein, J. N. (2013). "The relationship between ADHD symptom dimensions, clinical correlates, and functional impairments," *J. Dev. Behav. Pediatr.* 34(7), 469-477. DOI: 10.1097/DBP.0b013e3182a39890
- Gascon, M., Vrijheid, M., Martínez, D., Forn, J., Grimalt, J. O., Torrent, M., and Sunyer, J. (2011). "Effects of pre and postnatal exposure to low levels of polybromodiphenyl ethers on neurodevelopment and thyroid hormone levels at 4 years of age," *Environ. Int.* 37(3), 605-611. DOI: 10.1016/j.envint.2010.12.005
- Gibson, E. A., Siegel, E. L., Eniola, F., Herbstman, J. B., and Factor-Litvak, P. (2018). "Effects of polybrominated diphenyl ethers on child cognitive, behavioral, and motor

- development,” *Int. J. Env. Res. Pub. He.* 15(8), 1636-1656. DOI: 10.3390/ijerph15081636
- Gunnarsen, L., Nielsen, P. A., and Wolkoff, P. (1993). “Design and characterization of the climpaq, chamber for laboratory investigations of materials, pollution and air quality,” *Proceedings of the 6th International Conference on Indoor Air Quality and Climate 2*, 507512.
- Hallgren, S., and Darnerud, P. O. (2002). “Polybrominated diphenyl ethers (PBDEs), polychlorinated biphenyls (PCBs) and chlorinated paraffins (CPs) in rats—Testing interactions and mechanisms for thyroid hormone effects,” *Toxicology* 177(2-3), 227-243. DOI: 10.1016/S0300-483X(02)00222-6
- Hansen, J., and Olsen, J. H. (1995). “Formaldehyde and cancer morbidity among male employees in Denmark,” *Cancer Cause. Control* 6(4), 354-360. DOI: 10.1007/BF00051411
- Harley, K. G., Berger, K., Rauch, S., Kogut, K., Henn, B. C., Calafat, A. M., Huen, K., Eskenazi, B., and Holland, N. (2017). “Association of prenatal urinary phthalate metabolite concentrations and childhood BMI and obesity,” *Pediatr. Res.* 82(3), 405-415. DOI: 10.1038/pr.2017.112
- Hauptmann, M., Stewart, P. A., Lubin, J. H., Freeman, L. E. B., Hornung, R. W., Herrick, R. F., Hoover, R. N., Fraumeni, Jr., J. F., Blair, A., and Hayes, R. B. (2009). “Mortality from lymphohematopoietic malignancies and brain cancer among embalmers exposed to formaldehyde,” *JNCI–J. Natl. Cancer I.* 101(24), 1696-1708. DOI: 10.1093/jnci/djp416
- Hauser, R., Meeker, J. D., Duty, S., Silva, M. J., and Calafat, A. M. (2006). “Altered semen quality in relation to urinary concentrations of phthalate monoester and oxidative metabolites,” *Epidemiology* 17(6), 682-691. DOI: 10.1097/01.ede.0000235996.89953.d7
- He, Y., Peng, L., Zhang, W., Liu, C., Yang, Q., Zheng, S., Bao, M., Huang, Y., and Wu, K. (2018). “Adipose tissue levels of polybrominated diphenyl ethers and breast cancer risk in Chinese women: A case-control study,” *Environ. Res.* 167, 160-168. DOI: 10.1016/j.envres.2018.07.009
- Herbstman, J. B., and Mall, J. K. (2014). “Developmental exposure to polybrominated diphenyl ethers and neurodevelopment,” *Current Environmental Health Reports* 1(2), 101-112. DOI: 10.1007/s40572-014-0010-3
- Hites, R. A. (2004). “Polybrominated diphenyl ethers in the environment and in people: A meta-analysis of concentrations,” *Environ. Sci. Technol.* 38(4), 945-956. DOI: 10.1021/es035082g
- Hoffman, K., Lorenzo, A., Butt, C. M., Hammel, S. C., Henderson, B. B., Roman, S. A., Scheri, R. P., Stapleton, H. M., and Sosa, J. A. (2017). “Exposure to flame retardant chemicals and occurrence and severity of papillary thyroid cancer: A case-control study,” *Environ. Int.* 107, 235-242. DOI: 10.1016/j.envint.2017.06.021
- Hogberg, H. T., Boufield, M., Ulker, O. C., Sa, R. C., Harris, G., Kleensang, A., Maertens, A., Pamies, D., Smirnova, L., Zhao, L., *et al.* (2016). “3D models and omics approaches to study developmental neurotoxicity,” *Toxicol. Lett.* 258(Supplement), S16. DOI: 10.1016/j.toxlet.2016.06.1172
- Holness, D. L., and Nethercott, J. R. (1989). “Health status of funeral service workers exposed to formaldehyde,” *Arch. Environ. Health* 44(4), 222-228. DOI: 10.1080/00039896.1989.9935887

- Hooper, K., and McDonald, T. A. (2000). "The PBDEs: An emerging environmental challenge and another reason for breast-milk monitoring programs," *Environ. Health Persp.* 108(5), 387-392. DOI: 10.1289/ehp.00108387
- Hubbe, M., Pizzi, A., Zhang, H., and Halis, R. (2018). "Critical links governing performance of self-binding and natural binders for hot-pressed reconstituted lignocellulosic board without added formaldehyde: A review," *BioResources* 13(1), 2049-2115. DOI: 10.15376/biores.13.1.Hubbe
- Ibhazehiebo, K., Iwasaki, T., Kimura-Kuroda, J., Miyazaki, W., Shimokawa, N., and Koibuchi, N. (2011). "Disruption of thyroid hormone receptor-mediated transcription and thyroid hormone-induced Purkinje cell dendrite arborization by polybrominated diphenyl ethers," *Environ. Health Persp.* 119(2), 168-175. DOI: 10.1289/ehp.1002065
- International Agency for Research on Cancer (IARC) (2004). *IARC Classifies Formaldehyde as Carcinogenic to Humans* [Press Release No. 153], Lyon, France, ([http://www.iarc.fr/ENG/Press\\_Releases/archives/pr153a.html](http://www.iarc.fr/ENG/Press_Releases/archives/pr153a.html)), Accessed 17 Dec 2018
- IARC Working Group on the Evaluation of Carcinogenic Risks to Humans (2006). "Formaldehyde, 2-butoxyethanol and 1-tert-butoxypropan-2-ol," *IARC Monog. Eval. Carcinog. Risk Hum* 88, 1-478.
- Iversen, O. H. (1988). "Formaldehyde and skin tumorigenesis in SENCAR mice," *Environ. Int.* 14(1), 23-27. DOI: 10.1016/0160-4120(88)90373-X
- Jacobson, M. H., Barr, D. B., Marcus, M., Muir, A. B., Lyles, R. H., Howards, P. P., Pardo, L., and Darrow, L. A. (2016). "Serum polybrominated diphenyl ether concentrations and thyroid function in young children," *Environ. Res.* 149, 222-230. DOI: 10.1016/j.envres.2016.05.022
- Jin, X. B., Jiang, Z. H., Wen, X. W., Zhang, R., and Qin, D. C. (2017). "Flame retardant properties of laminated bamboo lumber treated with monoammonium phosphate (MAP) and boric acid/borax (SBX) compounds," *BioResources* 12(3), 5071-5085. DOI: 10.15376/biores.12.3.5071-5085
- Jones-Otazo, H. A., Clarke, J. P., Diamond, M. L., Archbold, J. A., Ferguson, G., Harner, T., Richardson, G. M., Ryan, J. J., and Wilford, B. (2005). "Is house dust the missing exposure pathway for PBDEs? An analysis of the urban fate and human exposure to PBDEs," *Environ. Sci. Technol.* 39(14), 5121-5130. DOI: 10.1021/es048267b
- Jyothi, A. N., Moorthy, S., and Vimala, B. (2007). "Physicochemical and functional properties of starch from two species of *Curcuma*," *Int. J. Food Prop.* 6(1), 135-145. DOI: 10.1081/JFP-120016630
- Kannerth, J., Hahn, G., and Gindl, W. (2009). "Feasibility of particle board production using bone glue," *Eur. J. Wood Wood Prod.* 67(2), 243-245. DOI: 10.1007/s00107-009-0307-3
- Kerns, W. D., Pavkov, K. L., Donofrio, D. J., Gralla, E. J., and Swenberg, J. A. (1983). "Carcinogenicity of formaldehyde in rats and mice after long-term inhalation exposure," *Cancer Res.* 43(9), 4382-4392. DOI:
- Khamgaonkar, M. B., and Fulare, M. B. (1991). "Pulmonary effects of formaldehyde exposure—An environmental-epidemiological study," *Indian J. Chest Dis.* 33(1), 9-13. DOI:
- Koch, H. M., and Angerer, J. (2007). "Di-iso-nonylphthalate (DINP) metabolites in human urine after a single oral dose of deuterium-labelled DINP," *Int. J. Hyg. Envir. Heal.* 210(1), 9-19. DOI: 10.1016/j.ijheh.2006.11.008

- Koch, H. M., Bolt, H. M., and Angerer, J. (2004). "Di(2-ethylhexyl)phthalate (DEHP) metabolites in human urine and serum after a single oral dose of deuterium-labelled DEHP," *Arch. Toxicol.* 78(3), 123-130. DOI: 10.1007/s00204-003-0522-3
- Koch, H. M., Bolt, H. M., Preuss, R., and Angerer, J. (2005). "New metabolites of di(2-ethylhexyl)phthalate (DEHP) in human urine and serum after single oral doses of deuterium-labelled DEHP," *Arch. Toxicol.* 79(7), 367-376. DOI: 10.1007/s00204-004-0642-4
- Koch, H. M., and Calafat, A. M. (2009). "Human body burdens of chemicals used in plastic manufacture," *Philos. T. Roy. Soc. B* 364(1526), 2063-2078. DOI: 10.1098/rstb.2008.0208
- Koch, H. M., Christensen, K. L., Harth, V., Lorber, M., and Brüning, T. (2012). "Di-n-butyl phthalate (DnBP) and diisobutyl phthalate (DiBP) metabolism in a human volunteer after single oral doses," *Arch. Toxicol.* 86(12), 1829-1839. DOI: 10.1007/s00204-012-0908-1
- Koch, H. M., Lorber, M., Christensen, K. L., Pälme, C., Koslitz, S., and Brüning, T. (2013). "Identifying sources of phthalate exposure with human biomonitoring: Results of a 48 h fasting study with urine collection and personal activity patterns," *Int. J. Hyg. Envir. Heal.* 216(6), 672-681. DOI: 10.1016/j.ijheh.2012.12.002
- Koch, H. M., Preuss, R., and Angerer, J. (2006). "Di(2-ethylhexyl)phthalate (DEHP): Human metabolism and internal exposure—An update and latest results," *Int. J. Androl.* 29(1), 155-165. DOI: 10.1111/j.1365-2605.2005.00607.x
- Köppel, C., Baudisch, H., Schneider, V., and Ibe, K. (1990). "Suicidal ingestion of formalin with fatal complications," *Intens. Care Med.* 16(3), 212-214. DOI: 10.1007/BF01724806
- Lacorte, S., and Ikononou, M. G. (2009). "Occurrence and congener specific profiles of polybrominated diphenyl ethers and their hydroxylated and methoxylated derivatives in breast milk from Catalonia," *Chemosphere* 74(3), 412-420. DOI: 10.1016/j.chemosphere.2008.09.050
- Lamaming, J., Sulaiman, O., Sugimoto, T., Hashim, R., Said, N., and Sato, M. (2013). "Influence of chemical components of oil palm on properties of binderless particleboard," *BioResources* 8(3), 3358-3371. DOI: 10.15376/biores.8.3.3358-3371
- Lee, A. T., Shah, J. J., Li, L., Cheng, Y., Moore, P. K., and Khanna, S. (2008). "A nociceptive-intensity-dependent role for hydrogen sulphide in the formalin model of persistent inflammatory pain," *Neuroscience* 152(1), 89-96. DOI: 10.1016/j.neuroscience.2007.11.052
- Leng, G., Koch, H. M., Gries, W., Schütze, A., Langsch, A., Brüning, T., and Otter, R. (2014). "Urinary metabolite excretion after oral dosage of bis(2-propylheptyl) phthalate (DPHP) to five male volunteers—Characterization of suitable biomarkers for human biomonitoring," *Toxicol. Lett.* 231(2), 282-288. DOI: 10.1016/j.toxlet.2014.06.035
- Luce, D., Leclerc, A., Bégin, D., Demers, P. A., Gérin, M., Orłowski, E., Kogevinas, M., Belli, S., Bugel, I., Bolm-Audorff, U., et al. (2002). "Sino nasal cancer and occupational exposures: A pooled analysis of 12 case control studies," *Cancer Cause. Control* 13(2), 147-157. DOI: 10.1023/A:1014350004255
- Lunder, S., Hovander, L., Athanassiadis, I., and Bergman, A. (2010). "Significantly higher polybrominated diphenyl ether levels in young U.S. children than in their mothers," *Environ. Sci. Technol.* 44(13), 5256-5262. DOI: 10.1021/es1009357

- Marceaux, J. C., Dilks, L. S., and Hixson, S. (2008). "Neuropsychological effects of formaldehyde use," *J. Psychoactive Drugs* 40(2), 207-210. DOI: 10.1080/02791072.2008.10400632
- Martin, O. V., Evans, R. M., Faust, M., and Kortenkamp, A. (2017). "A human mixture risk assessment for neurodevelopmental toxicity associated with polybrominated diphenyl ethers used as flame retardants," *Environ. Health Persp.* 125(8), Article ID 87016. DOI: 10.1289/EHP826
- Meeker, J. D., Calafat, A. M., and Hauser, R. (2007). "Di(2-ethylhexyl) phthalate metabolites may alter thyroid hormone levels in men," *Environ. Health Persp.* 115(7), 1029-1034. DOI: 10.1289/ehp.9852
- Meeker, J. D., and Ferguson, K. K. (2011). "Relationship between urinary phthalate and bisphenol A concentrations and serum thyroid measures in U.S. adults and adolescents from the National Health and Nutrition Examination Survey (NHANES) 2007-2008," *Environ. Health Persp.* 119(10), 1396-1402. DOI: 10.1289/ehp.1103582
- Migneault, S., Koubaa, A., Nadji, H., Riedl, B., Zhang, S. Y., and Deng, J. (2011). "Binderless fiberboard made from primary and secondary pulp and paper sludge," *Wood Fiber Sci.* 43(2), 180-193. DOI: 10.1007/s00107-009-0379-0
- Milner, H. R., and Woodart, A. C. (2016). "Sustainability of engineered wood products," in: *Sustainability of Construction Materials* (Second Edition), J. M. Khatib (ed.), Woodhead Publishing, pp: 159-180. DOI: 10.1016/B978-0-08-100370-1.00008-1
- Morgan, K. T., Jiang, X. Z., Starr, T. B., and Kerns, W. D. (1986). "More precise localization of nasal tumors associated with chronic exposure of F-344 rats to formaldehyde gas," *Toxicol. Appl. Pharm.* 82(2), 264-271. DOI: 10.1016/0041-008X(86)90201-2
- Moubarik, A., Allal, A., Pizzi, A., Charrier, F., and Charrier, B. (2010). "Characterization of a formaldehyde-free cornstarch-tannin wood adhesive for interior plywood," *Eur. J. Wood Wood Prod.* 68(4), 427-433. DOI: 10.1007/s00107-009-0379-0
- Müller, A. K., Nielsen, E., and Ladefoged, O. (2003). *Human Exposure to Selected Phthalates in Denmark* (Report No. 15), The Danish Veterinary and Food Administration, Søborg, Denmark.
- Müller, C., Kües, U., Schöpfer, C., and Kharazipour, A. (2007). "Natural binders," in: *Wood Production, Wood Technology, and Biotechnological Impacts*, U. Kües (ed.), Universitätsverlag Göttingen, Göttingen, Lower Saxony, Germany.
- Nasir, M., Gupta, A., Beg, M. D. H., Chua, G. K., Jawaid, M., Kumar, A., and Khan, T. A. (2013). "Fabricating eco-friendly binderless fiberboard from laccase-treated rubber wood fiber," *BioResources* 8(3), 3599-3608. DOI: 10.15376/biores.8.3.3599-3608
- National Toxicology Program (2011). *12<sup>th</sup> Report on Carcinogens* (Report No. 12), U.S. Department of Health and Human Services, Research Triangle Park, NC, USA.
- Navarrete, P., Mansouri, H. R., Pizzi, A., Tapin-Lingua, S., Benjelloun-Mlayah, B., Pasch, H., and Rigolet, S. (2010). "Wood panel adhesives from low molecular mass lignin and tannin without synthetic resins," *J. Adhes. Sci. Technol.* 24(8-10), 1597-1610. DOI: 10.1163/016942410X500972
- Net, S., Sempéré, R., Delmont, A., Paluselli, A., and Ouddane, B. (2015). "Occurrence, fate, behavior and ecotoxicological state of phthalates in different environmental matrices," *Environ. Sci. Technol.* 49(7), 4019-4035. DOI: 10.1021/es505233b
- Nikvash, N., Kharazipour, A., and Euring, M. (2012). "Effects of wheat protein as a biological binder in the manufacture of particleboards using a mixture of canola,

- hemp, bagasse, and commercial wood,” *Forest Prod. J.* 62(1), 49-57. DOI: 10.13073/FPJ-D-11-00102.1
- Noor, M. A. B. M., Mohammad, A. M. D., Islam, M. N., and Mehat, N. A. (2000). “Physico-chemical properties of oil palm trunk starch,” *Starch-Stärke* 51(8-9), 293-301. DOI: 10.1002/(SICI)1521-379X(199909)51:8/9<293::AID-STAR293>3.0.CO;2-F
- Özdemir, F., Serin, Z. O., and Tutuş, A. (2017). “Investigation of the effect of some fire-retardant chemicals and mineral materials used in surface coating on combustion performance of particleboard,” *BioResources* 12(4), 8862-8869. DOI: 10.15376/biores.12.4.8862-8869
- Parry, E., Zota, A. R., Park, J. S., and Woodruff, T. J. (2018). “Polybrominated diphenyl ethers (PBDEs) and hydroxylated PBDE metabolites (OH-PBDEs): A six-year temporal trend in Northern California pregnant women,” *Chemosphere* 195, 777-783. DOI: 10.1016/j.chemosphere.2017.12.065
- Patel, K. G., Bhatt, H. V., and Choudhury, A. R. (2003). “Alteration in thyroid after formaldehyde (HCHO) treatment in rats,” *Ind. Health* 41(3), 295-297. DOI: 10.2486/indhealth.41.295
- Pedersen, G. A., Jensen, L. K., Fankhauser, A., Biedermann, S., Petersen, J. H., and Fabech, B. (2008). “Migration of epoxidized soybean oil (ESBO) and phthalates from twist closures into food and enforcement of the overall migration limit,” *Food Addit. Contam. A* 25(4), 503-510. DOI: 10.1080/02652030701519088
- Pitten, F. A., Kramer, A., Herrmann, K., Bremer, J., and Koch, S. (2000). “Formaldehyde neurotoxicity in animal experiments,” *Pathol. Res. Pract.* 196(3), 193-218. DOI: 10.1016/S0344-0338(00)80100-4
- Pizzi, A. (1983). “Tannin-based wood adhesives,” in: *Wood Adhesives: Chemistry and Technology*, A. Pizzi (ed.), Marcel Dekker, New York City, NY, USA, pp. 364.
- Pizzi, A. (2006). “Developments in biobased adhesives for wood bonding: Opportunities and issues,” *J. Adhes. Sci. Technol.* 20(8), 829-846. DOI: 10.1163/156856106777638635
- Pizzi, A. (2014). “Natural adhesives, binders, and matrices for wood and fiber composites: Chemistry and technology,” in: *Research Developments in Wood Engineering and Technology*, A. Aguilera and J. P. Davim (eds.), IGI Global, Hershey, PA, USA, pp. 131-181.
- Porterfield, S. P., and Hendrich, C. E. (1993). “The role of thyroid hormones in prenatal and neonatal neurological development—Current perspectives,” *Endocr. Rev.* 14(1), 94-106. DOI: 10.1210/edrv-14-1-94
- Qiu, X. H., Bigsby, R. M., and Hites, R. A. (2009). “Hydroxylated metabolites of polybrominated diphenyl ethers in human blood samples from the United States,” *Environ. Health Persp.* 117(1), 93-98. DOI: 10.1289/ehp.11660
- Responsible Purchasing Network (RPN) (2013). “Green purchasing best practices: Office and dorm furniture,” ([http://www.responsiblepurchasing.org/purchasing\\_guides/furniture/naspo\\_rpn\\_furniture\\_purchasing\\_guide.pdf](http://www.responsiblepurchasing.org/purchasing_guides/furniture/naspo_rpn_furniture_purchasing_guide.pdf)), Accessed 4 January 2019.
- Ritchie, I. M., and Lehnen, R. G. (1987). “Formaldehyde-related health complaints of residents living in mobile and conventional homes,” *Am. J. Public Health* 77(3), 323-328. DOI: 10.2105/AJPH.77.3.323
- Robledo, C. A., Yeung, E., Mendola, P., Sundaram, R., Maisog, J., Sweeney, A. M., Barr, D. B., and Louis, G. M. (2015). “Preconception maternal and paternal exposure



- to persistent organic pollutants and birth size: The LIFE study,” *Environ. Health Persp.* 123(1), 88-94. DOI: 10.1289/ehp.1308016
- Rusch, G. M., Clary, J. J., Rinehart, W. E., and Bolte, H. F. (1983). “A 26-week inhalation toxicity study with formaldehyde in the monkey, rat, and hamster,” *Toxicol. Appl. Pharm.* 68(3), 329-343. DOI: 10.1016/0041-008X(83)90276-4
- Rydén, A., Nestor, G., Jakobsson, K., and Marsh, G. (2012). “Synthesis and tentative identification of novel polybrominated diphenyl ether metabolites in human blood,” *Chemosphere* 88(10), 1227-1234. DOI: 10.1016/j.chemosphere.2012.03.076
- Salthammer, T., Mentese, S., and Marutzky, R. (2010). “Formaldehyde in the indoor environment,” *Chem. Rev.* 110(4), 2536-2572. DOI: 10.1021/cr800399g
- Samandar, E., Silva, M. J., Reidy, J. A., Needham, L. L., and Calafat, A. M. (2009). “Temporal stability of eight phthalate metabolites and their glucuronide conjugates in human urine,” *Environ. Res.* 109(5), 641-646. DOI: 10.1016/j.envres.2009.02.004
- Sathyanarayana, S., Calafat, A. M., Liu, F., and Swan, S. H. (2008). “Maternal and infant urinary phthalate metabolite concentrations: Are they related?,” *Environ. Res.* 108(3), 413-418. DOI: 10.1016/j.envres.2008.07.002
- Sharma, N. K., Verma, C. S., Charier, V. M., and Prasad, R. (2013). “Eco-friendly flame-retardant treatments for cellulosic green building materials,” *Indoor Built Environ.* 24(3), 422-432. DOI: 10.1177/1420326X13516655
- Silva, M. J., Barr, D. B., Reidy, J. A., Kato, K., Malek, N. A., Hodge, C. C., Hurtz, D., Calafat, A. M., Needham, L. L., and Brock, J. W. (2003). “Glucuronidation patterns of common urinary and serum monoester phthalate metabolites,” *Arch. Toxicol.* 77(10), 561-567. DOI: 10.1007/s00204-003-0486-3
- Sjödin, A., Jones, R. S., Caudill, S. P., Wong, L. Y., Turner, W. E., and Calafat, A. M. (2014). “Polybrominated diphenyl ethers, polychlorinated biphenyls, and persistent pesticides in serum from the National Health and Nutrition Examination Survey: 2003-2008,” *Environ. Sci. Technol.* 48(1), 753-760. DOI: 10.1012/es4037836
- Soffritti, M., Belpoggi, F., Lambertin, L., Lauriola, M., Padovani, M., and Maltoni, C. (2002). “Results of long-term experimental studies on the carcinogenicity of formaldehyde and acetaldehyde in rats,” *Ann. NY. Acad. Sci.* 982(1), 87-105. DOI: 10.1111/j.1749-6632.2002.tb04926.x
- Song, W., Zhu, M., Lin, W., and Zhang, S. (2018). “Determining optimum material mixture ratio and hot-pressing parameters for new hybrid fiber-reinforced composites: Modeling and optimization by response surface methodology,” *BioResources* 13(2), 4202-4223.
- Song, W., Zhu, M., and Zhang, S. (2018). “Comparison of the properties of fiberboard composites with bamboo green, wood, or their combination as the fibrous raw material,” *BioResources* 13(2), 3315-3334.
- Song, W., Wei, W., Wang, D., and Zhang, S. (2017). “Preparation and properties of new plywood composites made from surface modified veneers and polyvinyl chloride films,” *BioResources* 12(4), 8320-8339.
- Song, W., Cao, Y., Wang, D., Hou, G., Shen, Z., and Zhang, S. (2015). “An investigation on formaldehyde emission characteristics of wood building materials in Chinese standard tests: Product emission levels, measurement uncertainties, and data correlations between various tests,” *PloS One* 10(12), e0144374. DOI:10.1371/journal.pone.0144374
- Sonnier, R., Taguet, A., Ferry, L., and Lopez-Cuesta, J. M. (2018). *Towards Bio-based Flame Retardant Polymers*, Springer International Publishing, New York, NY, USA.

- Stoker, T. E., Laws, S. C., Crofton, K. M., Hedge, J. M., Ferrell, J. M., and Cooper, R. L. (2004). "Assessment of DE-71, a commercial polybrominated diphenyl ether (PBDE) mixture, in the EDSP male and female pubertal protocols," *Toxicol. Sci.* 78(1), 144-155. DOI: 10.1093/toxsci/kfh029
- Sulaiman, N. S., Hashim, R., Amini, M. H. M., Sulaiman, O., and Hiziroglu, S. (2013). "Evaluation of the properties of particleboard made using oil palm starch modified with epichlorohydrin," *BioResources* 8(1), 283-301. DOI: 10.15376/biores.8.1.283-301
- Takahashi, M., Hasegawa, R., Furukawa, F., Toyoda, K., Sato, H., and Hayashi, Y. (1986). "Effects of ethanol, potassium metabisulfite, formaldehyde and hydrogen peroxide on gastric carcinogenesis in rats after initiation with N-methyl-N'-nitro-N-nitrosoguanidine," *Jpn. J. Cancer Res.* 77(2), 118-124. DOI: 10.20772/cancersci1985.77.2\_118
- Terzi, E., Kartal, S., Pişkin, S., Stark, N., Kantürk Figen, A., and White, R. (2018). "Colemanite: A fire retardant candidate for wood plastic composites," *BioResources* 13(1), 1491-1509. DOI: 10.15376/biores.13.1.1491-1509
- Testa, C., Nuti, F., Hayek, J., De Felice, C., Chelli, M., Rovero, P., Latini, G., and Papini, A. M. (2012). "Di-(2-ethylhexyl) phthalate and autism spectrum disorders," *ASN Neuro.* 4(4), 223-229. DOI: 10.1042/AN20120015
- The Lowell Center for Sustainable Production (2011). "Phthalates and their alternatives: The health and environmental concerns," *University of Massachusetts Lowell*, (<https://www.sustainableproduction.org/downloads/PhthalateAlternatives-January2011.pdf>), Accessed 4 January 2019.
- Thrasher, J. D., and Kilburn, K. H. (2001). "Embryo toxicity and teratogenicity of formaldehyde," *Arch. Environ. Health* 56(4), 300-311. DOI: 10.1080/00039890109604460
- Thrasher, J. D., Wojdani, A., Cheung, G., and Heuser, G. (1987). "Evidence for formaldehyde antibodies and altered cellular immunity in subjects exposed to formaldehyde in mobile homes," *Arch. Environ. Health* 42(6), 347-350. DOI: 10.1080/00039896.1987.9934357
- Thuvander, A., and Darnerud, P. O. (1999). "Effects of polybrominated diphenyl ether (PBDE) and polychlorinated biphenyl (PCB) on some immunological parameters after oral exposure in rats and mice," *Toxicol. Environ. Chem.* 70(1-2), 229-242. DOI: 10.1080/02772249909358751
- Trasande, L., Spanier, A. J., Sathyanarayana, S., Attina, T. M., and Blustein, J. (2013). "Urinary phthalates and increased insulin resistance in adolescents," *Pediatrics* 132(3), e646-e655. DOI: 10.1542/peds.2012-4022
- United States Consumer Product Safety Commission (US CPSC) (2008). "Consumer Product Safety Improvement Act (CPSIA)," CPSC, (<https://www.cpsc.gov/Regulations-Laws--Standards/Statutes/The-Consumer-Product-Safety-Improvement-Act>), Accessed 15 December 2018
- US Department of Health and Human Services, Public Health Service, ATSDR (1999). *Toxicological Profile for Formaldehyde*, ATSDR Atlanta, GA, USA.
- US Department of Health and Human Services, Public Health Service, ATSDR (2017). *Toxicological Profile for Polybrominated Diphenyl Ethers (PBDEs)*, ATSDR Atlanta, GA, USA.
- US Department of Health and Human Services, Public Health Service, ATSDR (2010). *Toxicological Profile for Boron*, ATSDR Atlanta, GA, USA.

- US Environmental Protection Agency (EPA) (1998). *Toxicological Review of Methylene Diphenyl Diisocyanate (MDI)*, U.S. EPA, Washington D.C., USA.
- US EPA (2012). *Phthalates Action Plan*, U.S. EPA, Washington D.C., USA.
- US EPA (2013). *Polybrominated Diphenyl Ethers (PBDEs) Significant New Use Rules (SNUR)*, U.S. EPA, Washington D.C., USA.
- US EPA (2014). *An Alternatives Assessment for the Flame Retardant Decabromodiphenyl Ether (DecaBDE)*, U.S. EPA, Washington D.C., USA.
- US EPA (2017). *Technical Fact Sheet – Polybrominated Diphenyl Ethers (PBDEs)*, U.S. EPA, Washington D.C., USA.
- Upton, K., Sathyanarayana, S., De Roos, A. J., Thompson, M. L., Scholes, D., Dills, R., and Holt, V. L. (2013). “Phthalates and risk of endometriosis,” *Environ. Res.* 126, 91-97. DOI: 10.1016/j.envres.2013.07.003
- Vandi, L. J., Chan, C., Werker, A., Richardson, D., Laycock, B., and Pratt, S. (2018). “Wood-PHA composites: Mapping opportunities,” *Polymers* 10(7), 751-766. DOI: 10.3390/polym10070751
- Van Vliet, E. D., Reitano, E. M., Chhabra, J. S., Bergen, G. P., and Whyatt, R. M. (2011). “A review of alternatives to di (2-ethylhexyl) phthalate-containing medical devices in the neonatal intensive care unit,” *J. Perinatol.* 31(8), 551-560. DOI: 10.1038/jp.2010.208
- Vargová, M., Wagnerová, J., Lísková, A., Jakubovský, J., Gajdová, M., Stolcová, E., Kubová, J., Tulinská, J., and Stenclová, R. (1993). “Subacute immunotoxicity study of formaldehyde in male rats,” *Drug Chem. Toxicol.* 16(3), 255-275. DOI: 10.3109/01480549309081819
- Vieira, M. G. A., Da Silva, M. A., Dos Santos, L. O., and Beppu, M. M. (2011). “Natural-based plasticizers and biopolymer films: A review,” *Eur. Polym. J.* 47(3), 254-263. DOI: 10.1016/j.eurpolymj.2010.12.011
- Vonderheide, A. P., Mueller, K. E., Meija, J., and Welsh, G. (2008). “Polybrominated diphenyl ethers: Causes for concern and knowledge gaps regarding environmental distribution, fate and toxicity,” *Sci. Total Environ.* 400(1-3), 425-436. DOI: 10.1016/j.scitotenv.2008.05.003
- Vuong, A. M., Braun, J. M., Yolton, K., Xie, C., Webster, G. M., Sjödin, A., Dietrich, K. N., Lanphear, B. P., and Chen, A. (2017a). “Prenatal and postnatal polybrominated diphenyl ether exposure and visual spatial abilities in children,” *Environ. Res.* 153, 83-92. DOI: 10.1016/j.envres.2016.11.020
- Vuong, A. M., Yolton, K., Poston, K. L., Xie, C., Webster, G. M., Sjödin, A., Braun, J. M., Dietrich, K. N., Lanphear, B. P., and Chen, A. (2017b). “Prenatal and postnatal polybrominated diphenyl ether (PBDE) exposure and measures of inattention and impulsivity in children,” *Neurotoxicol. Teratol.* 64, 20-28. DOI: 10.1016/j.ntt.2017.09.001
- Vuong, A. M., Yolton, K., Xie, C., Webster, G. M., Sjödin, A., Braun, J. M., Dietrich, K. N., Lanphear, B. P., and Chen, A. (2017c). “Childhood polybrominated diphenyl ether (PBDE) exposure and neurobehavior in children at 8 years,” *Environ. Res.* 158, 677-684. DOI: 10.1016/j.envres.2017.07.028
- Wang, C., Lin, Z., Dong, Q., Lin, Z., Lin, K., Wang, J., Huang, J., Huang, X., He, Y., Huang, C., *et al.* (2012). “Polybrominated diphenyl ethers (PBDEs) in human serum from Southeast China,” *Ecotox. Environ. Safe.* 78, 206-211. DOI: 10.1016/j.ecoenv.2011.11.016

- Wantke, F., Focke, M., Hemmer, W., Bracun, R., Wolf-Abdolvahab, S., Götz, M., Tschabitscher, M., Gann, M., and Tappler, P. (2000). "Exposure to formaldehyde and phenol during an anatomy dissecting course: Sensitizing potency of formaldehyde in medical students," *Allergy* 55(1), 84-87. DOI: 10.1034/j.1398-9995.2000.00307.x
- Weschler, C. J. (2009). "Changes in indoor pollutants since the 1950s," *Atmos. Environ.* 43(1), 153-169. DOI: 10.1016/j.atmosenv.2008.09.044
- Williams, G. R. (2008). "Neurodevelopmental and neurophysiological actions of thyroid hormone," *J. Neuroendocrinol.* 20(6), 784-794. DOI: 10.1111/j.1365-2826.2008.01733.x
- Wilmer, J. W., Woutersen, R. A., Appelman, L. M., Leeman, W. R., and Feron, V. J. (1987). "Subacute (4-week) inhalation toxicity study of formaldehyde in male rats: 8-hour intermittent versus 8-hour continuous exposures," *J. Appl. Toxicol.* 7(1), 15-16. DOI: 10.1002/jat.255007104
- Wilmer, J. W., Woutersen, R. A., Appelman, L. M., Leeman, W. R., and Feron, V. J. (1989). "Subchronic (13-week) inhalation toxicity study of formaldehyde in male rats: 8-hour intermittent versus 8-hour continuous exposures," *Toxicol. Lett.* 47(3), 287-293. DOI: 10.1016/0378-4274(89)90147-1
- Wolkoff, P. (1996). "An emission cell for measurement of volatile organic compounds emitted from building materials for indoor use - the field and laboratory emission cell FLEC," *Gefahrstoffe - Reinhaltung der Luft.* 56, 151-157.
- World Health Organization (WHO) (2000). *Air Quality Guidelines for Europe* (No. 91), WHO Regional Publications, Copenhagen, Denmark.
- Xu, X., Liu, J., Zeng, X., Lu, F., Chen, A., and Huo, X. (2014). "Elevated serum polybrominated diphenyl ethers and alteration of thyroid hormones in children from Guiyu, China," *PLOS ONE* 9(11), e113699. DOI: 10.1371/journal.pone.0113699
- Yu, L., Cai, J., Li, H., Lu, F., Qin, D., and Fei, B. (2017). "Effects of boric acid and/or borax treatments on the fire resistance of bamboo filament," *BioResources* 12(3), 5296-5307. DOI: 10.15376/biores.12.3.5296-5307
- Yu, Y. X., Pang, Y. P., Li, C., Li, J. L., Zhang, X. Y., Yu, Z. Q., Feng, J. L., Wu, M. H., Sheng, G. Y., and Fu, J. M. (2012). "Concentrations and seasonal variations of polybrominated diphenyl ethers (PBDEs) in in- and out-house dust and human daily intake via dust ingestion corrected with bio-accessibility of PBDEs," *Environ. Int.* 42, 124-131. DOI: 10.1016/j.envint.2011.05.012
- Zhang, B., Xu, T., Huang, G., Yin, D., Zhang, Q., and Yang, X. (2018). "Neurobehavioral effects of two metabolites of BDE-47 (6-OH-BDE-47 and 6-MeO-BDE-47) on zebrafish larvae," *Chemosphere* 200, 30-35. DOI: 10.1016/j.chemosphere.2018.02.064
- Zhang, D. H., Zhang, A. J., and Xue, L. X. (2015). "A review of preparation of binderless fiberboards and its self-bonding mechanism," *Wood Sci. Technol.* 49(4), 661-679. DOI: 10.1007/s00226-015-0728-6
- Zhou, T., Taylor, M. M., DeVito, M. J., and Crofton, K. M. (2002). "Developmental exposure to brominated diphenyl ethers results in thyroid hormone disruption," *Toxicol. Sci.* 66(1), 105-116. DOI: 10.1093/toxsci/66.1.105

Zoeller, R. T., and Rovet, J. (2004). "Timing of thyroid hormone action in the developing brain: Clinical observations and experimental findings," *J. Neuroendocrinol.* 16(10), 809-818. DOI: 10.1111/j.1365-2826.2004.01243.x

Article submitted: February 19, 2019; Peer review completed: May 3, 2019; Revised version received: May 19, 2019; Accepted: May 25, 2019; Published: June 3, 2019.  
DOI: 10.15376/biores.14.3.Ulker