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# Neurotoxicity of polychlorinated biphenyls and related organohalogenes

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## Abstract

Halogenated organic compounds are pervasive in natural and built environments. Despite restrictions on the production of many of these compounds in most parts of the world through the Stockholm Convention on Persistent Organic Pollutants (POPs), many “legacy” compounds, including polychlorinated biphenyls (PCBs), are routinely detected in human tissues where they continue to pose significant health risks to highly exposed and susceptible populations. A major concern is developmental neurotoxicity, although impacts on neurodegenerative outcomes have also been noted. Here, we review human studies of prenatal and adult exposures to PCBs and describe the state of knowledge regarding outcomes across domains related to cognition (e.g., IQ, language, memory, learning), attention, behavioral regulation and executive function, and social behavior, including traits related to attention-deficit hyperactivity disorder (ADHD) and autism spectrum disorders (ASD). We also review current understanding of molecular mechanisms underpinning these associations, with a focus on dopaminergic neurotransmission, thyroid hormone disruption, calcium dyshomeostasis, and oxidative stress. Finally, we briefly consider contemporary sources of organohalogenes that may pose human health risks via mechanisms of neurotoxicity common to those ascribed to PCBs.

## Introduction

Epidemiologic [6, 12, 163] and preclinical [158, 192] studies have identified the brain as a vulnerable target of polychlorinated biphenyls (PCBs). This review focuses on the state of the science regarding the neurotoxicity of PCBs at epidemiologic, neuropathologic, and molecular levels. Much of the scientific literature describes the developmental neurotoxicity of PCBs, which are arguably among the most extensively

studied developmental neurotoxicants of the nonmetallic persistent organic pollutants (POPs). While our knowledge of how early or mid-life exposures to PCBs influence the aging brain is not as comprehensive, there is evidence linking PCBs to increased risk of neurodegenerative outcomes, and this literature is also addressed in this review.

PCBs are a chemically related class (chemotype) within a vast number of POPs of diverse anthropogenic origins. Common to all POPs is their chemical stability in the environment, and their resistance to metabolic degradation. These properties underlie the ubiquitous environmental distribution and bioaccumulation of POPs in human tissues, which have raised global concerns about adverse health consequences of widespread human exposures [36, 43, 179]. These concerns have led to restrictions on the production of POPs imposed by many governments through the Stockholm Convention on POPs signed in 2001 and appended in 2008 and 2014 [165, 166]. Despite these regulatory efforts, “legacy” PCBs persist in the environment and in human tissue, where they are still routinely detected, thus continuing to pose significant health risks, particularly among susceptible populations.

Chemically, PCBs are a complex mixture of isomers or congeners that differ significantly in their structure and modes of action. PCB congeners have been heuristically

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divided into two categories based on structure: PCBs with biphenyl rings substituted with zero chlorines in the *ortho* position that assume a coplanar biphenyl orientation in solution, and PCBs with one to four chlorines in the *ortho* position that assume increasing degrees of noncoplanar biphenyl orientation (Fig. 1). Of the 209 possible PCB congeners, 19 congeners are stable atropisomer or enantiomers, with chiral asymmetry about their biphenyl bond axes. PCBs can be hydroxylated or sulfonated metabolically, and each enantiomer and metabolite has potentially distinct interactions with biological targets, contributing to overall PCB neurotoxicity.

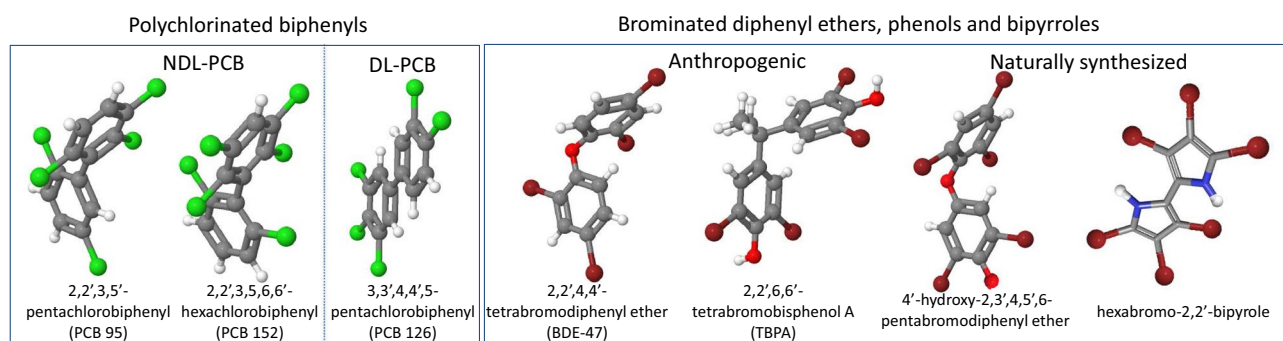
Coplanar PCBs have been shown to mimic dioxin in that they bind with relatively high affinity to the aryl hydrocarbon receptor (AhR), which regulates the transcription of a large group of dioxin-responsive genes [105, 156]. Although low-level exposures to dioxin and dioxin-like PCBs have been associated with a number of adverse outcomes in several organ systems, especially liver, skin, and immune function [11, 124, 203], and are probably carcinogenic [109], there is less evidence from epidemiologic and preclinical studies that they are associated with neurotoxic outcomes. Recent preclinical studies have demonstrated that dioxin alters neuronal migration [99] and ultrasonic vocalization [100] in mice, albeit at doses higher than those required for induction of other pathological endpoints. But whether dioxin-like PCBs have similar effects on these neurodevelopmental endpoints has yet to be determined. There is also emerging data linking dioxin-like compounds to increased risk of neurodegenerative disease, specifically amyotrophic lateral sclerosis (ALS) [4]. The most prevalent pathology of ALS is the accumulation of phosphorylated insoluble aggregates of transactive response DNA binding protein 43 (TDP-43) in neurons. Dioxin-like compounds were shown to significantly increase TDP-43 in induced pluripotent stem cells and mouse brains via AhR-dependent mechanism(s)

[4]. Whether environmentally relevant levels of dioxin-like PCBs similarly induce TDP-43 expression in neural tissues, and whether this effect translates into increased ALS risk remains to be determined.

In contrast, noncoplanar PCBs, also referred to as nondioxin-like PCBs (NDL PCBs) have little to no binding affinity for AhR. However, there is extensive scientific evidence linking NDL PCBs to neurotoxic outcomes [146, 180]. Further, NDL PCBs are predominant not only in contemporary environmental samples, but also in breast milk, serum, and adipose tissue from wildlife and humans [9, 146]. Given that NDL PCBs constitute the predominant congener subclass comprising contemporary human exposures, and that the weight of evidence implicates NDL PCBs in neurotoxic sequelae, this review largely focuses on the neurotoxicity of NDL PCBs.

While PCBs are among the most extensively studied POP chemotype, other major POPs have been shown to have neuropathogenic potential, including organochlorine insecticides (e.g., DDT, cyclodienes, hexachlorocyclohexanes) and, more recently, polybrominated diphenyl ether flame retardants (PBDEs) and newly discovered organohalogens produced as disinfectant byproducts [96, 146, 223]. Figure 1 compares PCBs with chemotypes that have emerged more recently as POPs of human health concern because of their potential as neuromodulators and neurotoxicants. Interestingly, many brominated organics from anthropogenic sources, especially those released as water disinfection byproducts, closely mimic organohalogens naturally produced in marine environments that function as primary ecological signaling molecules and secondary deterrence and defense toxins [1].

In this review, we present a critical evaluation of the state of the science regarding the neurotoxicity of PCBs from both a neurodevelopmental and neurodegenerative standpoint. We also review what is known about the underlying



**Fig. 1** Exemplary non-coplanar and coplanar polychlorinated biphenyls (PCBs) of anthropogenic origin. Coplanar PCBs are also referred to as dioxin-like (DL) PCBs because of their ability to binding the AhR with high affinity, a mechanism considered to contribute to can-

cer risk, whereas non-coplanar PCBs, which are referred to as nondioxin-like (NDL) PCBs, which possess one or more *ortho* chlorine substitution have negligible AhR activity and have been shown to be neurotoxic through other mechanisms

mechanisms of PCB neurotoxicity, including alterations in biogenic amines (dopamine in particular), endocrine disruption, altered calcium signaling in neurons, and oxidative stress. Our understanding about PCB neurotoxicity is likely to provide broad insights regarding the neurotoxic potential of a number of chemically related organohalogenes that are eliciting scientific and public attention as emerging concerns, and these chemically related organohalogenes are briefly discussed in the closing statements.

## Mechanisms of PCB neurotoxicity

While significant research effort has focused on understanding how PCBs interfere with neurodevelopment or promote neurodegeneration, the molecular mechanisms by which PCBs cause neurotoxicity, and whether all or only a subset of PCB congeners are neurotoxic, remain key questions in the field. Prevailing mechanistic hypotheses of PCB neurotoxicity include: (1) altered dopamine (DA) signaling; (2) disruption of thyroid hormone signaling; (3) perturbation of intracellular  $\text{Ca}^{2+}$  dynamics; and (4) oxidative stress. The mechanistic studies described below have been highlighted because the PCB exposure paradigms that elicited changes in these important biological pathways are relevant to the PCB levels detected in humans and wildlife as well as the congener profiles detected in contemporary environmental and biological samples [9, 67, 146]. Importantly, these mechanisms have also been implicated in the neurotoxicity of several contemporary POPs with NDL structures, including brominated diphenyl ethers (PBDEs) [34, 96] and bisphenols [222].

### Evidence that PCBs alter DA neurotransmission

In vitro studies using pheochromocytoma cells (PC12) exposed to technical PCB mixtures, e.g., the Aroclors, provided the first evidence that PCBs significantly reduce cellular DA concentrations [174]. Subsequent structure–activity relationship studies revealed that *ortho*-substituted NDL PCBs, but not coplanar DL congeners, significantly depleted cellular DA [174, 190]. These in vitro observations were corroborated in vivo in *Macaca mulatta* (rhesus monkeys) orally exposed to Aroclor 1254 [168]. Analysis of striatal and substantia nigral concentrations of DA using high-performance liquid chromatography with electrochemical detection demonstrated significantly reduced nigrostriatal levels of DA in PCB-exposed subjects, an effect that persisted following termination of PCB exposure [169]. Studies using Aroclor 1016 (a mixture of lightly chlorinated NDL PCB congeners) in non-human primates also demonstrated significant reductions in nigrostriatal DA concentrations, providing additional verification of in vitro studies

demonstrating that NDL PCB congeners impair DA neurotransmission [170]. Similar observations were reported in adult male mice orally exposed to Aroclor 1254 for 4 weeks [111]. Experimental evidence suggests that DA depletion may result from reduced levels of tyrosine hydroxylase, the rate limiting enzyme in the formation of DA [28], or from inhibition of vesicular monoamine transporter (VMAT) or the plasma membrane dopamine transporter (DAT) [22, 120, 172]. However, evidence for direct molecular interactions of NDL PCBs with one or more of these targets remains weak.

### Evidence that PCBs alter thyroid hormone (TH) signaling

Maternal or infant hypothyroidism is linked to significantly impaired neurodevelopment, including intellectual disability [139, 205]. The importance of decreased circulating TH, primarily measured as changes in levels of thyroxine (T4) in serum, has been widely debated as the primary mechanism of PCB neurotoxicity since it was initially proposed [207]. Many studies, performed in a variety of species, including humans, have demonstrated a negative association between PCB exposures and serum T4 [19, 61, 62, 88, 224]. Developmental exposure of rats to Aroclor 1254 was shown to cause hypothyroxinemia (decreased serum T4) coincident with hearing loss, and supplementation with T4 throughout development attenuated PCB-induced motor and auditory deficits [61]. However, cognitive effects of developmental PCB exposure do not appear to be mediated by thyroid hormone-dependent mechanisms. For example, rats developmentally exposed to Aroclor 1254 at levels sufficient to reduce serum T4 levels had no deficits in learning and memory assessed in the T-maze or the Morris water maze [220]. Conversely, rats developmentally exposed to Aroclor 1254 at levels that did not significantly decrease serum T3 or T4 levels exhibited poorer performance in the Morris water maze [215]. Two additional studies of individual PCB congeners also failed to support a role for TH deficits in cognitive and behavioral deficits linked to PCBs [131, 162]. Consistent with these preclinical data are recently published results from a prospective birth cohort, the *Hokkaido study*, which included 222 mother–neonate pairs and was designed to determine whether there was a relationship between specific hydroxylated PCB metabolites (OH-PCBs) and maternal/neonate serum thyroxine levels [80]. Unexpectedly, and in direct opposition to the hypothyroidism theory of PCB developmental neurotoxicity, the Hokkaido study indicated that maternal and/or neonatal exposure to specific hydroxylated PCB metabolites (OH-PCBs) were positively correlated with free thyroxine levels [80]. Results from another recently published prospective study performed in Denmark indicated no significant association between the placental levels of 35 PCB congeners and placental T4 or T3, although

one PCB, PCB 81, was positively associated with reverse T3 (rT3) [117].

Further complicating the thyroid hormone disruption theory of PCB neurotoxicity, PCBs have been shown to have both antagonistic and agonistic effects on thyroid hormone receptors, depending on the congener and the dose [62, 131, 224]. Moreover, direct molecular interactions of PCBs with TH receptors have not been convincingly demonstrated. Recently Sethi et al. [173] used a novel stable in vitro reporter gene assay [52, 125] but failed to detect any thyroid hormone receptor activity for 12 most abundant PCB congeners mimicking the congener profile in maternal serum, either singly or as a mixture. These inconsistencies across both human and rodent studies raise significant questions regarding an association between PCB effects on serum thyroid hormone levels and adverse neurodevelopmental outcomes.

### Evidence that PCBs alter Ca<sup>2+</sup> channel function and Ca<sup>2+</sup>-dependent signaling

The precise spatial and temporal patterning of cytoplasmic Ca<sup>2+</sup> signals are critical for normal neurodevelopment, the establishment and refinement of neural networks, and the dynamic synaptic plasticity associated with many behaviors [7, 17, 103]. Diverse biochemical, biophysical and cellular approaches across many laboratories have consistently shown that NDL PCBs potentially alter intracellular Ca<sup>2+</sup> dynamics [146], and a number of studies have linked PCB-induced Ca<sup>2+</sup> dyshomeostasis to specific neurodevelopmental deficits [180]. An early hypothesis was that developmental exposures to NDL PCBs caused cognitive and behavioral deficits by disrupting Ca<sup>2+</sup>-dependent PKC signaling [101, 217]. This hypothesis derived in part from observations that PCB disruption of Ca<sup>2+</sup> fluctuations caused rapid redistribution of protein kinase C (PKC) isoforms in astrocytoma cells [191]. Further structure–activity relationship studies demonstrated that NDL PCBs [101, 217], but not DL PCBs [35], increased the translocation of PKC to the plasma membrane of cultured cerebellar neurons. In vivo studies demonstrated that developmental exposure to Aroclor 1254 changed the subcellular distribution of PKC isoforms in the brain in a complex region-specific manner [190]. However, whether influences on PKC signaling represent a primary mechanism driving the behavioral and cognitive impairments observed following developmental PCB exposure remains unanswered.

Subsequent studies focused on the primary molecular mechanism(s) responsible for PCB effects on intracellular Ca<sup>2+</sup> dynamics. In vitro studies using pharmacological blockade of specific Ca<sup>2+</sup> channels have shown that NDL PCBs increase extracellular Ca<sup>2+</sup> entry into cells through a number of mechanisms, including activation of L-type

voltage-sensitive Ca<sup>2+</sup> channels and NMDA receptors [78, 128]. However, such changes are elicited only at high PCB concentrations ( $\geq 10 \mu\text{M}$ ), which have been demonstrated to produce nonspecific changes in membrane fluidity [102]. NDL-PCBs have also been shown to facilitate the release of Ca<sup>2+</sup> from intracellular stores through sensitization of two genetically related Ca<sup>2+</sup> channels localized to the endoplasmic/sarcoplasmic reticulum (ER/SR): ryanodine receptors (RyR) [207–211] and inositol 1,4,5-trisphosphate receptors (IP<sub>3</sub>R) [77]. Of these, RyR sensitization is more sensitive to NDL PCBs, and this interaction has been shown to exhibit a consistent stringent structure–activity relationship, including stereoselectivity, as determined using biochemical, electrophysiological, cellular and in vivo approaches [47, 53, 54, 71, 72, 134, 146, 214]. Nanomolar PCB concentrations directly interact with RyR channels to stabilize RyR channels in their full open conformation [161], which sensitizes ER/SR Ca<sup>2+</sup> signals to physiological modulators including Ca<sup>2+</sup>, Mg<sup>2+</sup>, and ATP.

Sensitization of RyRs by NDL PCBs alters the fidelity of spontaneous Ca<sup>2+</sup> oscillations [200] and the neuroplasticity of hippocampal CA1 neurons [209] in vitro. Consistent with this observation, picomolar to nanomolar concentrations of NDL PCB 95, which is among the most RyR-active congeners, activate two Ca<sup>2+</sup>-dependent signaling pathways in cultured rat hippocampal neurons: (1) sequential activation of CaMKK, CaMKI $\alpha/\gamma$ , and MEK/ERK and CREB to increase transcription of Wnt2 [200]; and (2) CREB-mediated miR132 upregulation, which suppresses the translation of p250GAP [115]. In cultured rat hippocampal neurons, the former signaling pathway mediates PCB 95-induced dendritic growth [200], whereas the latter mediates PCB 95-induced synaptogenesis, which is evident as increased spine density and increased frequency of miniature excitatory post-synaptic currents [115]. Several lines of evidence indicate that PCB effects on RyR activity are causally linked to dendritic growth and spine formation: (1) RyR-active PCB congeners, such as PCB 95 and PCB 136, but not PCB congeners that lack activity at the RyR, such as PCB 66, enhance dendritic growth and promote spine formation in cultured hippocampal and cortical neurons; and (2) pharmacological blockade or siRNA knockdown of RyRs inhibit the dendrite and spine promoting activity of these NDL PCBs [115, 201]. An interesting observation from these studies is that PCB-induced dendritic growth exhibits a non-monotonic concentration–effect relationship, with effects observed in the pico- to nanomolar range but not at femto- or micromolar concentrations [201, 214]. Elucidating the mechanisms responsible for this non-monotonic concentration–effect relationship remains a critical data gap in the field.

These observations suggest that PCB effects on dendritic growth contribute to PCB-induced behavioral deficits. In

support of this possibility, developmental exposure of rats to Aroclor 1254 at 1, but not 6, mg kg<sup>-1</sup> in the maternal diet was found to cause learning and memory deficits in the Morris water maze, and this dose of Aroclor 1254 also modulated RyR activity and dendritic arborization in brain regions that subserve Morris water maze behavior [215]. Several interesting findings emerged from this study. First, developmental PCB exposure promoted dendritic growth in cerebellar Purkinje cells and neocortical pyramidal neurons among untrained animals but attenuated or reversed experience-dependent dendritic growth in these brain regions among Morris water maze-trained littermates. Second, deficits in learning and memory were only seen in the 1 mg kg<sup>-1</sup> group, and the most robust changes in dendritic arborization were also observed in this dose group, replicating the non-monotonic dose–response relationship observed in the *in vitro* studies. Third, this study demonstrated that the behavioral deficits correlated better with changes in RyR expression and activity than with changes in serum levels of thyroid hormone or sex hormones. In a separate study, a similar non-monotonic dose–response relationship was observed for dendritic arborization of CA1 pyramidal neurons in the hippocampus of rats exposed to PCB 95 in the maternal diet throughout gestation and lactation [201]. Developmental exposure to PCB 95 has also been shown to dramatically alter the tonotopy of the primary auditory cortex (A1) [91], presumably by tipping the balance of excitatory–inhibitory currents within A1 towards excitation during critical periods of its development [91, 97, 98, 209]. While these data strongly support the hypothesis that PCB effects on RyR activity contribute to behavioral deficits via modulation of calcium-dependent signaling pathways that regulate neuroplasticity, experimental evidence demonstrating that the molecular and cellular effects are causally linked to behavioral deficits are lacking. Testing this hypothesis using RyR knockout animals is not feasible because of embryo lethality, but alternative approaches using knock-in animals expressing human gain-of-function mutations that enhance RyR sensitization by organohalogen [44, 219] may increase confidence in causal inferences.

Although the current data support a primary mechanistic contribution of RyRs to the developmental neurotoxicity of NDL PCBs, their possible role in the induction and progression of neurodegenerative diseases, such as Parkinson's disease (PD), is less clear. Although Ca<sup>2+</sup> dyshomeostasis induced by ER stress is an important molecular mechanism of selective loss of dopaminergic (DA) neurons in PD, an etiological role for RyR channels has only recently received experimental attention. Using a 6-hydroxydopamine (6-OHDA)-induced *in vitro* SN4741 cell model of PD, Huang et al. [75] showed that 6-OHDA significantly increased cytoplasmic Ca<sup>2+</sup> levels, an effect that could be blocked by either an ER stress inhibitor or a specific RyR

blocker. The authors further demonstrated that 6-OHDA reduced the spike number and rheobase of DA neurons in acute brain slices, effects that were also reversed by RyR blockade. Furthermore, novel 6-aminoquinoxaline derivatives appear to confer neuroprotective effects on DA neurons in cellular animal models of PD that are at least partially mediated through interactions with RyRs [110]. Interestingly, blocking Ca<sup>2+</sup> influx did not change PCB effects on intracellular dopamine levels in a catecholaminergic cell line [87], suggesting that PCBs may be more likely affecting dopamine homeostasis by mediating dysfunction of ER Ca<sup>2+</sup> stores. Clearly this area needs more attention before inferences can be made about convergent mechanisms linking PCB neurotoxicity across life stages.

### Evidence of PCB-induced oxidative stress

Oxidative stress occurs when the production of reactive oxygen species (ROS) and reactive nitrogen species (RNS) exceeds the antioxidant capacity of the system [8, 92, 137]. This imbalance can cause oxidative and nitrative damage to macromolecules, which in turn can cause cell damage or even cell death. Numerous *in vitro* studies have demonstrated that PCBs cause oxidative stress in neurons [35, 74, 121]. In cultured cerebellar granule neurons, exposure to the NDL-congeners, PCB 4 or 153, or to Aroclor 1254, but not to DL PCBs, increased cellular ROS levels, which caused concentration-dependent cell death [35]. Similarly, Aroclor 1254 or the NDL PCB 47, but not the DL PCB 77, triggered neuronal cell apoptosis in primary hippocampal neurons that was blocked by co-exposure to the antioxidant alpha-tocopherol [74], suggesting that NDL PCBs induced apoptosis via increased ROS. Interestingly, this same study demonstrated that apoptosis induced by PCB 47 was also prevented by pharmacologic blockade of RyRs [74]. Whether and how RyRs and ROS interact to mediate PCB-induced apoptosis has yet to be determined. One possibility is that ROS is a consequence of RyR activation. As discussed above, NDL PCBs stabilize RyRs in the open conductance state, which increases release of Ca<sup>2+</sup> from the ER [146]. Elevated intracellular Ca<sup>2+</sup> can trigger apoptosis either by directly activating caspases in neuronal cells and/or by increasing Ca<sup>2+</sup> flux into mitochondria, thereby increasing ROS production, which triggers release of cytochrome c from mitochondria with subsequent activation of caspases [113]. Alternatively, ROS can act as signaling molecules that initiate apoptosis via targeted interactions with specific cellular components, one of which is the RyR [23, 144, 146]. ROS have been shown to interact directly with hyperreactive cysteine residues on the RyR to enhance its open probability [45, 136, 188, 213]. This suggests an alternative model of PCB-induced apoptotic signaling in which NDL PCBs activate RyRs

indirectly as a consequence of PCB-induced increase in ROS [74]. There is precedence for this proposed mechanism in that ROS generated by quinone have been shown to increase the calcium conductance of RyR from rabbit sarcoplasmic reticulum membranes. In fact, RyR channels are tightly regulated by changes in the ER/SR membrane redox potential [45, 213] and are highly sensitive to modification by redox active environmental chemicals including naphthoquinones and benzo[a]pyrene-7,8-dione [46, 145], providing a possible mechanistic link between  $\text{Ca}^{2+}$  dysregulation and oxidative stress. These may not be mutually exclusive models in that PCBs may independently influence both RyR activation and ROS generation, with each effect augmenting the other in a positive feedback mechanism.

There is conflicting evidence as to whether PCBs cause oxidative stress in the brain *in vivo*. Oral exposure of adult male mice to a relatively high dose (25 mg/kg/day) of Aroclor 1254 for 4 weeks was associated with increases in superoxide dismutase (SOD), heme oxygenase (HO-1), lipid peroxides, and protein carbonyl levels in the striatum and cerebellum [111]. In rats exposed throughout gestation and lactation to environmentally relevant low doses of Aroclor 1254 in the maternal diet (0.1 or 1.0 mg/kg/day), multiple brain regions showed increased expression of biomarkers of oxidative stress [216]. In contrast, a study of mice exposed to a mixture of six NDL PCBs via lactation failed to find any evidence of oxidative damage in the brain [39]. The discrepancy between these studies may be due to differences in species, timing and length of PCB exposure, dose and composition of PCB mixtures to which animals were exposed, as well as the use of different biomarkers to measure oxidative stress. Collectively, the available studies suggest that oxidative stress may contribute to the developmental neurotoxicity and neurodegenerative toxicity associated with NDL PCBs, potentially via RyR-dependent mechanisms.

## The neuropathologic effects of PCBs

We performed a literature search using PubMed (searched December 2018) to identify human and animal studies of neuropathologic effects of PCBs. We used the following search terms: “polychlorinated biphenyls”, “PCBs but not printed circuit boards”, “neuropathology”, “neurophysiology”, and “neuroimaging”. Our search yielded seven unique articles, of which three were reviews [42, 118, 129, 150, 152, 171, 182]. Inspection of the references cited in these three reviews yielded an additional two articles describing the neuropathology of PCBs. These articles, as well as other relevant publications describing PCB-induced effects

on neuronal cytoarchitecture *in vivo* are discussed in this section.

## The neuropathology of PCB developmental neurotoxicity

Currently, there is a very limited literature regarding pathologic effects of PCBs in the developing human brain. A pilot study that conducted magnetic resonance imaging (MRI) scans on 60 children age 4 years found stronger associations of PCBs with response inhibition among participants with smaller splenium size [182]. This indicates that children with suboptimal development of the splenium could be particularly vulnerable to exposure to PCBs, increasing their risk for poorer response inhibition, a core trait of attention-deficit hyperactivity disorder (ADHD). Another pilot study administered functional magnetic resonance imaging (fMRI) to 12 adolescent boys from a Faroese birth cohort assembled in 1986–1987 [204]. Boys with high exposure to both PCBs and methylmercury (MeHg) showed more activation across brain regions during visual and motor tasks than boys with lower mixed exposures to PCBs and MeHg. In a separate cohort of Inuit from Nunavik (Northern Québec) [40], cord blood concentrations of methylmercury or lead, but not PCBs, were associated with pattern-reversal visual evoked potentials (VEPs) recorded at the occipital cortex. Prenatal exposure to lead was associated with a delay of the N150 latency at most contrast levels. This study suggests that heavy metal exposure, in particular during the gestational period, but not PCB exposure, can impair the development of visual processing. Whether early-life exposures negatively impact healthy brain aging later in life [167] remains unanswered.

The earliest neuropathologic studies of developmental exposures to PCBs in preclinical models utilized very high PCB levels intended to mimic the exposures associated with the accidental mass poisonings due to ingestion of PCB-contaminated cooking oil in Japan in 1968 and Taiwan in 1979 [155]. In one of these studies [29], pregnant CD-1 mice were dosed orally with the DL PCB 77 at 32 mg/kg/day on days 10 through 16 of gestation. This exposure paradigm caused significant maternal toxicity as evident by significantly smaller litter sizes (60% of control) and higher neonatal mortality (40% at postnatal day 20). In approximately 55% of surviving offspring from PCB-exposed dams, a permanent motor disturbance, referred to as “spinning syndrome”, developed at weaning. Spinning syndrome was characterized by repetitive jerking of the head, hyperkinesia, and swift stereotyped circular or spinning movement, described as “chasing one’s tail”.

From over 100 spinners resulting from 60 PCB-exposed litters, 24 spinners ranging in age from 4 to 34 weeks along with age-matched controls and 12 littermates who did not

develop spinning syndrome (nonspinners) were subjected to pathologic studies [29]. Interestingly, no detectable neuronal changes, gliotic foci or demyelinating lesions were detected in the CNS of either PCB-exposed spinners or nonspinners by light microscopy. However, cylindrical CNS peninsulas (CCPs) that projected into the cranial and spinal nerve roots were noted in all spinners and some nonspinners, but none of the control subjects. The CCPs were characterized by a cylindrical column of nerve fibers, myelinated or nonmyelinated, and/or glial bundles in varying proportions, which were well-circumscribed and delineated from the neighboring PNS-type nerve fibers. The diameter and length of CCPs varied considerably up to 100  $\mu\text{m}$  in diameter and 5 mm in length. They were seen predominantly in the ventral roots, and to a lesser extent in the dorsal spinal roots, and most abundantly at lumbar levels. Large, apparently displaced, motor neurons were often detected within CCPs. Electron microscopy revealed that the CCPs in both the ventral and dorsal spinal roots were ensheathed in a glycocalyx layer that was indistinguishable from the basement membrane; and while collagen fibrils invested the outer surface of the CCPs, collagen fibrils were not observed within the CCPs. The myelinated fibers within the CCPs were distinguished by the lack of an outer basement membrane, the presence of oligodendroglia and shorter periodicity of myelin lamellae, all characteristics typical of CNS myelin. Signs of myelin sheath degeneration in the absence of macrophage infiltration were very common in CCPs, but not in the PNS-type myelinated fibers surrounding the CCP. All the cellular elements of the CNS (neurons, astroglial and oligodendroglia) were observed within CCPs. Some of the heterotopic neurons appeared normal with well-preserved terminal boutons detected along the cell surfaces. Except for frequent distal displacement in the ventral white matter, ventral horn motor neurons showed no ultrastructural changes suggestive of degenerative process in the perikaryon; however, along the cell surfaces, the nerve terminals appeared dense, amorphous and shrunken.

The authors noted that the glial outgrowth observed in the CCPs in the spinner mice was similar to that reported in pig spinal roots following transection and anastomosis, but concluded that mechanical injury was unlikely to be the cause of CCP development in the spinner mice [29]. Rather, the neuropathology induced by developmental exposure to very high levels of PCB 77 appeared remarkably similar to that observed in Werdnig–Hoffman disease. The authors concluded that PCB 77 interfered with the organization rather than the differentiation of neurons and glia since no histopathologic abnormalities could be detected in the CNS of mice exposed developmentally to PCB 77. Specifically, the authors conjectured that PCB 77 selectively interfered with synaptogenesis of inhibitory dopaminergic terminals, and that PCB exposure may be relevant to the high incidence of

aberrant motor development in hyperactive boys [29]. This hypothesis derived in part from pharmacologic observations in this study that, like hyperactivity in children, amphetamine was effective in attenuating spinning behavior in the spinner mice [29].

A later histopathologic study of Sprague–Dawley rats exposed to a more moderate dose of PCBs similarly concluded that PCBs did not cause overt damage to the developing brain, but rather PCBs interfered with the organization of neural networks [150]. In this study, pregnant Sprague–Dawley rats were fed chow containing Aroclor 1254 at 125 ppm throughout gestation and lactation. Upon weaning, offspring were maintained on chow containing 125 ppm Aroclor 1254 until euthanized for histological processing at 16, 30 and 60 days of age. Using Timm's silver sulfide staining to visualize the hippocampal mossy fibers, the authors demonstrated that continuous exposure to Aroclor 1254 from conception significantly reduced the relative size of II-P mossy fibers in 16-, 30- and 60-day old rats. In contrast, developmental exposure to Aroclor 1254 had no effect on the size of the hilar or suprapyramidal mossy fibers or on cortical thickness. The authors speculated that PCBs decreased the growth of mossy fibers as a result of increased glutamate or delayed myelination secondary to PCB-induced hypothyroidism or as a result of altered  $\text{Ca}^{2+}$  homeostasis [150]. However, none of these postulated mechanisms explains the selective effect of Aroclor 1254 on a subpopulation of granule cells that give rise to the II-P mossy fibers.

Subsequent preclinical studies of neuropathologic changes induced by developmental exposures to more environmentally relevant doses of PCBs further support the hypothesis that PCB-induced functional deficits reflect subtle organizational and/or functional defects of key neural networks of the brain [57, 167, 180]. Classic histopathologic analyses of brains from weanling rats exposed to Aroclor 1254 at 1 or 6 mg/kg/day in the maternal diet throughout gestation and lactation did not detect overt structural damage in the brain [215]. However, Golgi analyses of rat CA1 hippocampal pyramidal neurons and cerebellar Purkinje cells indicated that exposure to Aroclor 1254 at 6 mg/kg/day in the maternal diet throughout gestation and lactation caused a pronounced age-related increase in dendritic growth. Thus, even though dendritic lengths were significantly attenuated in PCB-exposed animals at postnatal day 22, by postnatal day 60, dendritic growth was comparable to or exceeded that observed in vehicle controls [114]. A subsequent study of postnatal day 31 rats [215] demonstrated that developmental Aroclor 1254 exposure increased dendritic arborization of cerebellar Purkinje cells and neocortical pyramidal neurons in animals not trained in a Morris water maze but caused dose-dependent dendritic retraction in these same neuronal cell types in littermates trained in the Morris water maze. These dendritic effects were significantly



more pronounced in the animals exposed to Aroclor 1254 at 1 mg/kg/day compared to 6 mg/kg/day in the maternal diet, and the lower dose, but not the higher dose of Aroclor 1254 elicited deficits in spatial learning and memory in the Morris water maze [215]. Interestingly, postmortem examinations of brains from Autism Spectrum Disorder (ASD) patients have revealed increased neuropil, which comprised dendrites, non-myelinated axons, synapses, vasculature and glial cell processes, in cortical regions [193]. Morphometric analyses of pyramidal neurons in various cortical regions altered in ASD, has revealed increased dendritic complexity, including increased dendritic spines [76, 79], on projection neurons. While studies from the 1990s indicated less complex dendritic arborization in the hippocampus of subjects with ASD [193], more recent studies of genetic mouse models of ASD have identified increased dendritic complexity of hippocampal neurons [27, 85, 90].

Another pathologic feature associated with ASD is an imbalance of excitatory to inhibitory neural circuits [60]. Kenet et al. [91] reported that rats developmentally exposed to NDL PCB 95 at 6 mg/kg/day in the maternal diet during gestation and lactation exhibited profoundly abnormal development of the primary auditory cortex (A1) without impairment of hearing sensitivity and brainstem auditory responses of pups. A1 was irregularly shaped and marked by internal nonresponsive zones, with grossly abnormal or reversed topographic organization in about half of the exposed pups. Changes in A1 network organization occurred concomitantly with changes in the balance of neuronal inhibition to excitation for A1 neurons, with significant alterations in the critical period plasticity that underlies normal postnatal auditory system development [91].

Histologic studies in preclinical models of PCB developmental neurotoxicity have also detected increased apoptosis in the brains of rat pups exposed to Aroclor 1254 at 0.1 or 1.0 mg/kg/day in the maternal diet throughout gestation and lactation [216]. TUNEL staining of sections from the cortex, hippocampus and cerebellum revealed significantly increased apoptosis in all three brain regions, with the most pronounced effect in the cerebellum, at postnatal day 1 but not postnatal day 21 [216]. These findings were confirmed by assays of caspase 3 activity. Histologic evidence of increased apoptosis coincided spatiotemporally with biochemical evidence of oxidative stress, specifically increased levels of 4-HNE and 3-NT as detected by western blotting [216].

Collectively, the available neuropathologic data suggest that neither overtly toxic levels of PCBs nor more environmentally relevant levels of PCBs cause major structural changes in the developing brain. Rather, developmental exposures to PCBs interfere with the organization of neural networks, evident by CCPs in spinal roots in response to very high PCB levels, or by changes in axonal and dendritic

morphology or altered patterns of apoptosis elicited by more environmentally relevant PCB exposures.

### The neuropathology of PCB-induced neurodegeneration

A [<sup>123</sup>Iodine] 2β-carbomethoxy-3β-[4-iodophenyl] tropane] single-photon emission computed tomography ([<sup>123</sup>I] β-CIT SPECT) imaging study of former capacitor workers with documented PCB exposures found that relative to non-exposed controls, individuals with occupational exposure to PCBs had significantly decreased striatal dopamine terminal densities [171], suggesting loss of dopaminergic neurons. Interestingly, this association was sex-specific. After statistical correction for a number of confounders, including age, only women show a significant inverse relationship between lipid-adjusted total serum PCB concentrations and β-CIT densities in the caudate, putamen or combined caudate and putamen, despite the fact that neither serum PCB levels nor age was statistically different between men and women. In a separate cohort of women participating in the Nun Study who had no clinical signs or symptoms of PD at death, elevated levels of NDL PCBs 138, 153, and 180 were observed in brain tissue that exhibited moderate nigral depigmentation compared to subjects with mild or no depigmentation [68].

Preclinical studies of neuropathologic changes in adult subjects exposed to PCBs at levels approximating the high end of occupational exposures are consistent with the limited neuropathologic findings in humans. As discussed above, studies in non-human primates demonstrated that long-term exposure to PCBs significantly reduced basal ganglia DA concentrations [168, 169]. To determine whether subchronic PCB exposure would cause cell loss in midbrain DAergic systems, ventral midbrain neurons were quantified by stereologic analyses at 2 weeks after a month-long exposure of adult male mice to Aroclor 1254 at 6, 12 and 25 mg/kg/day [111]. Moderate to high doses of Aroclor 1254 elicited death of both tyrosine hydroxylase (TH) immunopositive and TH immunonegative neurons in the ventral tegmental area. In the substantia nigra, the low and moderate doses of Aroclor 1254 significantly reduced only TH immunopositive cells, while the highest dose significantly decreased both TH immunopositive and TH immunonegative cells. The loss of DAergic and non-DAergic neurons correlated with increased levels of oxidative stress biomarkers in these same brain regions [111]. However, PCB exposure caused persistent increases in locomotor activity that correlated with dose-dependent increases in DA levels in the ventral midbrain. The authors interpreted these paradoxical data as suggesting the existence of compensatory mechanisms to maintain DA homeostasis in the face of decreased numbers of DAergic neurons [111].

Collectively, the limited literature available regarding the neuropathologic effects of PCBs on the aging brain suggest that PCBs promote the loss of DAergic and, at high exposure levels, non-DAergic neurons, within brain regions critically involved in motor function. Whether PCBs promote neuronal cell loss in other regions of the adult brain remains an open question.

## Epidemiological evidence of PCB neurotoxicity

### PCB developmental neurotoxicity

Accidental mass poisonings due to ingestion of PCB-contaminated cooking oil in Japan in 1968 and Taiwan in 1979 [155], resulted in a number of clinical morbidities, including adverse neurodevelopment in response to prenatal exposure [25, 26]. These incidents were the basis for classifying PCBs and other organochlorine compounds as suspected developmental neurotoxicants. The epidemiologic literature on lower-level PCB exposure, mostly via diet (e.g., fish, meat and dairy) [107, 119], and neurodevelopment is among the richest in environmental neuroepidemiology. Results of these studies, while generally supportive of an adverse impact of PCBs on the developing brain, have been mixed, with differences based on methodologic considerations, such as study design (prospective vs. cross-sectional), window of exposure (e.g., prenatal vs. postnatal), neurobehavioral domain (e.g., language, memory, executive function, inhibitory control) and age of neuropsychological assessment. PCBs that bioaccumulate in maternal tissues efficiently cross the placenta during pregnancy and are transferred postnatally through breast milk. This makes them a particular concern for the developing central nervous system, which undergoes rapid development during gestation and early life. Here, we review a subset of the available literature on PCBs and neurodevelopment with the aim of focusing on studies with more statistical power and methodologic rigor. To this end, we reviewed the state of the literature to date on prospective epidemiologic studies of prenatal PCB exposure measured in biospecimens collected during pregnancy and neurodevelopment across cognitive and behavioral endpoints in children age 3 years or older.

### Selection of studies for review

We performed a literature search using PubMed (searched June 2018) to identify human studies of prenatal exposure to PCBs and neurodevelopmental outcomes across domains related to cognition (e.g., IQ, language, memory, learning), attention, behavioral regulation and executive function, and social behavior, including traits related to attention-deficit

hyperactivity disorder (ADHD) and autism spectrum disorders (ASD). We used the following search terms: “polychlorinated biphenyls”, “PCBs”, “organochlorines”, “prenatal”, “pregnancy”, “neurodevelopment”, “attention”, “cognition”, “behavior”, “impulsivity”, “hyperactivity”, “attention deficit hyperactivity disorder”, and “autism”. We also inspected references of previously published review papers on PCBs and neurodevelopment [5, 41, 163, 207] to confirm that we identified all papers that met our criteria (detailed below). We limited our search to papers published in English-language journals after 1990.

For this review, we included a select body of literature to identify studies more robust in methodologic rigor and study power. Our inclusion criteria were: (1) studies that included analysis sample sizes of 100 or more participants, to focus on studies with more statistical precision; (2) prospective cohort studies that measured PCB exposure in the prenatal period, as neurodevelopment is generally recognized to be most sensitive to environmental insults during this period [63]; (3) studies that measured PCB exposure in biospecimens, primarily serum or plasma from blood collected from the mother during pregnancy or at delivery, including from the umbilical cord and placenta; and (4) studies of children 3 years of age or older, as neurodevelopmental outcomes can be measured more reliably in older children.

### Review findings

We identified 29 papers that met our inclusion criteria. These papers represented 14 different prospective cohort studies. Geographically, these studies were diverse and included countries in North America (Canada, Greenland and the USA), Europe (Belgium, Denmark and the Faroe Islands, Germany, the Netherlands, Spain, and Ukraine), and Asia (Japan). Studies were also diverse with respect to cohort birth year and, therefore, pregnancy PCB concentrations, which have been slowly decreasing since PCBs were banned. Studies most often presented findings for the sum of PCB congeners, though the number of congeners included and the limit of detection for these individual congeners varied widely.

### Prenatal PCB exposure and child cognition

We identified 12 papers from 9 different cohort studies that examined associations of PCBs with cognitive function among children ranging from ages 3 to 11 years. As shown in Table 1, most studies conducted tests of general intelligence (most commonly the Wechsler Intelligence Scale for Children, Kaufman Assessment Battery for Children and McCarthy Scales of Children’s Abilities), though a number also looked at specific cognitive domains, such as language and memory. Among these 12 papers, most (8)

**Table 1** Prenatal PCBs and cognitive function

Location	References	Sample size	PCB exposure matrix (congeners)	Median/mean concentrations	Outcome measure	Age at assessment (years)	Study results
Denmark (Faroe Islands)	Grandjean et al. [64]	435	Cord blood ΣPCBs (118, 138, 153, 170, 180)	Mean = 1.88 µg/L wet	NES-2 finger tapping, hand-eye coordination; BVMG; WISC-r digit span, similarities, block design; CVLT; BNT	7	↓ Language development
Japan (Tohoku)	Tatsuta et al. [189]	387	Cord blood ΣPCBs	Mean = 46.5 ng/g lipid	K-ABC	3	↓ Sequential and mental processing scores with highly chlorinated PCBs
Netherlands (Rotterdam and Groningen)	Patandin et al. [142]	395	Maternal plasma in last month of pregnancy ( <i>n</i> = 415) and cord ( <i>n</i> = 373) ΣPCBs (118, 138, 153, 180)	Median = 2.04 µg/L in maternal plasma and 0.38 µg/L in cord wet	K-ABC ( <i>n</i> = 395), RDLS ( <i>n</i> = 193)	3	↓ Overall cognitive and processing on K-ABC, ↓ verbal comprehension on RDLS
Spain (Menorca)	Vreugdenhil et al. [198]	372	Maternal plasma in last month of pregnancy and cord ΣPCBs (118, 138, 153, 180)	Median = 2.04 µg/L in maternal plasma and 0.38 µg/L in cord wet	MCSA	6	∅ (though possible effect modification by parental/home characteristics)
	Forns et al. [51]	405	Cord serum ΣPCBs (118, 138, 153, 180)	Median = 0.71 ng/ml wet	MCSA	11	↓ Executive function, verbal function, and visuospatial abilities, especially for PCB 153
United States (Collaborative Perinatal Project: 12 states)	Gray et al. [66]	894	Maternal serum ΣPCBs (28, 52, 74, 105, 118, 138, 153, 170, 180, 194, 203)	Median = 2.85 ug/L wet	WISC, WRAT-R	7	∅
United States (Cincinnati, Ohio)	Zhang et al. [221]	239	Maternal serum ΣPCBs (118, 138, 153, 158, 180)	Median = 31.3 ng/g lipid	WJ-III (5 years), WRAT-4 (8 years), WISV-IV (8 years)	5–8	∅
United States (Michigan)	Jacobson et al. [84]	226	Cord serum ΣPCBs (70% non-detects)	Mean = 0.4 ng/mL serum	MCSA*, Sternberg (memory), Kagan's (visual discrimination), vigilance task	4	↓ Visual discrimination, ↑ errors in short-term memory, ∅ sustained attention
	Jacobson et al. [81]	212	Cord serum ΣPCBs (118, 138, 153, 180)	Mean = 3 ng/ml serum	WISC-R, WRAT-R, WRMT-R	11	↓ Full scale, ↓ verbal IQ, ↓ freedom from distractibility, ↓ word/reading comprehension
United States (New Bedford, Massachusetts)	Orenstein et al. [140]	393	Cord serum ΣPCBs	Median = 0.3 ng/g serum	WRAML	8	∅

Table 1 (continued)

Location	References	Sample size	PCB exposure matrix (congeners)	Median/mean concentrations	Outcome measure	Age at assessment (years)	Study results
United States (Oswego, New York)	Stewart et al. [185]	293	Cord blood ΣPCBs	Median = 0.52 ng/g wet	MSCA	3, 4	↓ Executive function, verbal function, and visuospatial abilities at 3 years but not 4 years ↓ Full scale and verbal IQ
	Stewart et al. [184]	189	Placental ΣPCBs	Median = 1.5 ng/g wet	WISC-III	9	

*BVT* Boston naming test, *BV/MG* Bender visual motor gestalt, *CVLT* California verbal learning test, *K-ABC* Kaufman assessment battery for children, *MSCA* McCarthy Scales of Children's Abilities, *RLDS* Reynell Language Development Scales, *WISC-R* Wechsler Intelligence Scale for Children-Revised, *WISC-III* Wechsler Intelligence Scale for Children-Third Edition, *WISC-IV* Wechsler Intelligence Scale for Children-Fourth Edition, *WJ-III* Woodcock-Johnson III tests of achievement, *WRAT-R* wide range achievement test-revised, *WRAT-4* wide range achievement test-fourth edition, *WRAML* wide range assessment of memory and learning, *WRMT-R* Woodcock reading mastery tests-revised

found associations of PCBs with at least one aspect of cognition, including poorer scores on overall intelligence [51, 81, 142, 184, 185], language [64], processing speed [84], short-term memory [187], and reading comprehension [81]. Four papers report null effects of PCBs on all reported measures of cognition [66, 140, 198, 220]. In addition, two studies which found associations of PCBs with IQ at age 3 years [142, 185], found that these associations attenuated to the null at slightly older ages (ages 4 and 6 years) [185, 198]; though, for one of the studies, associations re-emerged when children were older (age 9) [184].

### Prenatal PCB exposure and child attention, behavioral regulation and social behavior

Table 2 summarizes results of 17 studies from 11 different cohorts that examined associations of prenatal PCBs with child attention, behavioral regulation and social behavior among children ranging mostly from ages 3–12 years (one study included participants up to age 22 years [187]). The majority of studies (10 of the 17) reported PCB-related associations with traits related to ADHD, including problems with impulse control [14, 82, 160, 182, 183, 186], hyperactivity [157, 160], and attention [51, 64, 82, 132, 159, 160, 195]. However, one of these studies, which reported more omission errors (indicating poorer attention) on a continuous performance test, also reported the opposite result when looking at parent-reported behaviors, with PCBs related to fewer parent-reported ADHD-related behaviors [132]. Four studies reported null associations of PCBs with behavioral problems, including ADHD diagnosis [13, 177, 187, 221]. Only two studies examined prenatal PCBs in relation to social behavior and autistic traits, with one study reporting that PCBs were associated with fewer autistic traits [135] and another study reporting mixed associations with autistic traits, depending on the specific PCB congener [16].

The epidemiologic literature on PCBs and neurodevelopment is extensive and continues to grow. We present a focused summary based on studies we determined to be the most rigorous in study design and statistical precision. Despite these narrow inclusion criteria, we found a large number of studies for this review (29 papers representing 14 prospective cohort studies). While the results of these studies were mixed, overall our review showed that most studies reported that prenatal PCBs were related to poorer cognitive function and more behavior problems. We chose to focus our review on prospective birth cohort studies with PCB exposure measured during the prenatal period. Studies have shown that gestation may be the most sensitive period of neurodevelopment [63], when the central nervous system is rapidly developing. However, neurodevelopment continues through childhood into early adulthood and is also vulnerable to chemical exposures. Early infancy, in particular,

**Table 2** Prenatal PCBs and child attention, behavioral regulation and social behavior

Location	References	Sample size	PCB exposure matrix (congeners)	Median/mean concentrations	Outcome measure	Age at assessment (years)	Study results
Belgium (Flanders)	Stoen et al. [177]	270	Cord blood $\Sigma$ PCBs (138, 153, 180)	Median = 74.5 ng/g lipid	SDQ	7–8	Ø
Canada (Nunavik)	Boucher et al. [12]	279	Cord blood PCB 153	Median = 93.6 ug/kg lipid	CBCL (teacher report form)	8–14	Ø
	Boucher et al. [13]	191	Cord blood PCB 153	Median = 93.3 ug/kg lipid	Go/No-go	9–12	↓ Error monitoring (aspect of behavioral regulation)
Denmark (Aarhus)	Strøm et al. [187]	876	Maternal serum $\Sigma$ PCBs (118, 138, 153, 154, 170, 180)	Median = 3.4 pmol/ml	Clinical diagnosis/prescribed medication for ADHD or depression	Up to 22	Ø
Denmark (Faroe Islands)	Grandjean et al. [64]	435	Cord blood $\Sigma$ PCBs (118, 138, 153, 170, 180)	Mean = 1.88 ug/L wet	CPT (NES2)	7	↑ (Slower) reaction time only in the presence of high mercury exposure
Germany (Duisburg)	Neugebauer et al. [132]	116	Maternal serum (23 weeks) $\Sigma$ PCBs (138, 153, 180)	Median = 0.16 lipid	KITAP (8 years) and FBB-ADHS (9 years)	8–9	↑ Omission errors for attention, but protective of parent-reported ADHD behavior
Greenland and Ukraine (Kharkiv)	Nowack et al. [135]	100	Maternal serum (28–42 weeks) $\Sigma$ PCBs	Median = 6.85 wet	SRS	10	↓ Autistic traits
	Rosenquist et al. [157]		Maternal serum (25 weeks gestation in Greenland, 23 weeks in Ukraine) PCB 153	Median = 107 (Greenland) and 27 (Ukraine) ng/g lipid	SDQ	5, 9	↑ Conduct problems and hyperactivity
Spain (Menorca)	Forns et al. [51]	393	Cord serum $\Sigma$ PCBs (118, 138, 153, 180)	Median = 0.54 ng/ml serum	Conners' CPT	11	Slower response time
United States (Cincinnati, Ohio)	Zhang et al. [221]	239	Maternal serum $\Sigma$ PCBs (118, 138, 153, 158, 180)	Median = 31.3 ng/g lipid	BASC-2	5–8	Ø
	Braun et al. [16]	175	Maternal serum PCBs (25 different congeners, including 138 and 153)	Median 138/158 = 7.7 and 153 = 11 ng/g lipid	SRS	4–5	Inconsistent associations with autistic traits across PCB congeners. 138/158 associated with ↑ autistic traits and 153 associated with ↓ autistic traits, though very imprecise

Table 2 (continued)

Location	References	Sample size	PCB exposure matrix (congeners)	Median/mean concentrations	Outcome measure	Age at assessment (years)	Study results
United States (Michigan)	Jacobson et al. [181]	154 (age 4 years), 148 (age 11 years)	Cord serum $\Sigma$ PCBs (70% non-detects)	Mean = 2.7 ng/mL serum	CPT, WCST, SWCT	4, 11	↑ Impulsivity, ↓ concentration, verbal, visual and auditory working memory
United States (New Bedford, Massachusetts)	Sagiv et al. [160]	573	Cord serum $\Sigma$ PCBs (118, 138, 153, 180)	Median = 0.19 ng/g serum	CRS-T	8	↑ ADHD-related behaviors (attention, impulsivity/hyperactivity)
	Sagiv et al. [159]	578 (CPT), 584 (WISC)	Cord serum $\Sigma$ PCBs (118, 138, 153, 180)	Median = 0.19 ng/g serum	CPT, WISC-III processing speed, freedom from distractibility	8	↑ Errors of omission (inattention) and slower processing speed among boys only
United States (Oswego, New York)	Stewart et al. [185]	189	Cord blood $\Sigma$ PCBs	Median = 0.52 ng/g wet	CPT (Catch the Cat)	4	↓ Response inhibition (more commission errors)
	Stewart et al. [183]	182 (age 8 years), 183 (age 9 years)	Cord blood $\Sigma$ PCBs	Categorized as ND, low, med and high	CPT (NES2 8 years), extended CPT (9 years)	8 and 9	↓ Response inhibition
	Stewart et al. [186]	167	Cord blood $\Sigma$ PCBs	Mean = 0.96 ng/g wet	DRL	9	↓ Response inhibition (shorter interresponse times and reinforcements earned)

BASC-2 behavioral assessment system for children, second edition, *CBCL* child behavior checklist, *CPT* Continuous Performance Test, *CRS-T* Conners Rating Scale for Teachers, *DRL* differential reinforcement of low rates task, *FBF-ADHS* parent rating scale for attention-deficit hyperactivity disorder, *KITAP* computerized test battery for attentional performance for children, *NES-2* neuropsychological examination system-2, *SRS* social responsiveness scale, *SDQ* Strengths and Difficulties Questionnaire, *SRS* Social Responsiveness Scale, *SWCT* stroop color-word test, *WCST* Wisconsin card sort test, *WISC-III* Wechsler Intelligence Scale for Children-Third Edition

may be a sensitive window of exposure due to high levels of PCBs in breastmilk. In addition, PCBs in dust from caulking materials and light ballasts used in schools and other buildings constructed before 1977 may be a significant source of exposure among children [20, 66]. Studies of associations of PCBs from breast milk with neurodevelopment show mixed findings, with many studies reporting null associations [5, 82, 83, 149, 194]. In addition, disentangling the impact of prenatal and postnatal exposure on neurodevelopment, particularly when it comes from breast milk, which may be highly correlated with transplacental exposure, presents a challenge.

We also chose to limit our review to studies of slightly older children (age 3 years or older). A sizable literature reports association of PCBs with neurodevelopment under the age of 3 using tests such as the Bayley Scales of Infant Development [15, 33, 58, 59, 104, 141, 206]; however, neuropsychological measures of infants and toddlers tend to be noisy, in part due to difficulty in achieving a desired state for testing in younger children [178]. Measured endpoints are therefore more likely to be reliable among older children. In addition, as demonstrated by three studies included in this review [142, 185, 198], PCB-related decrements in neurodevelopment may be transient, where decrements in cognitive or behavioral function found in early childhood recede in later childhood due to compensation by other brain regions or more stable measures of neurodevelopment at older ages.

The main limitation of any review article is publication bias, and this review is no exception. Pressures in the scientific and editorial processes promotes publication of positive results over null findings, which can skew the balance when trying to represent the weight of the evidence. Our stringent inclusion criteria may have offset this slightly, by filtering out smaller, methodologically less rigorous studies that are at greater risk for producing spurious findings. Yet, no review is immune to publication bias, and it is likely that a number of studies that would have met our inclusion criteria, but found null associations of PCBs and neurodevelopment, remain unpublished. Findings of this review paper should therefore be interpreted with this significant caveat in mind.

Another limitation of this review is that we did not evaluate associations with neurodevelopment for specific PCB congeners, or mechanism-based classes of congeners. This limitation is underscored by emerging studies that suggest it is specific RyR-active NDL PCBs and related organohalogenes that may be associated with increased risk of heritable neurodevelopmental disorders, including ASDs [21, 63, 65, 70, 176, 180], and may be at least in part mediated by cellular  $\text{Ca}^{2+}$  dyshomeostasis, mitochondrial dysfunction and alterations in gene-specific patterns of DNA methylation [38, 116, 126, 130, 146, 212]. The majority of studies included in this review used a sum of PCB congeners, which may be dominated by specific individual congeners

in the mix, such as PCB 153. Correlations across individual congeners also tend to be high, which makes disentangling individual congeners difficult due to collinearity. In addition, the mechanism for PCB neurotoxicity remains unclear and may be different across congeners and cognitive/behavioral domains, making a review of congener-specific associations challenging.

### Summary and priorities for future epidemiologic study of PCBs and neurodevelopment

In summary, we found that among larger studies with more methodologic rigor, most showed that PCBs were associated with poorer cognitive and behavioral development. Though PCB levels have been steadily decreasing since their ban in the 1970s, their persistence in the environment and in human tissue and continued exposure in older building materials make PCBs a continued public health concern. In addition, as analytic methods improve, measurement of congeners at lower limits of detection is possible. This may make it possible to examine outcomes in relation to lower-level congeners that could not be previously quantified, such as lower chlorinated congeners, which may also be related to neurodevelopment [133]. In addition, assessment of neurodevelopment continues to improve, in particular neuroimaging modalities, which have not yet been explored in relation to PCBs. This presents the opportunity to examine PCB-related associations with different, and potentially more sensitive, neurodevelopmental endpoints, including neuroimaging [73]. Finally, this review may inform the potential neurodevelopmental impact of more contemporary and emerging chemicals with structural and toxicological properties similar to PCBs.

### Epidemiology of occupational exposures to PCBs

In contrast to a wealth of epidemiological data showing associations of perinatal PCB exposures with poorer neurodevelopmental outcomes, there are considerably fewer published studies that have examined PCB exposures in relation to impaired brain aging. Negative consequences have been reported in adults exposed via consumption of PCB-contaminated foods. Subtle cognitive and behavioral changes have been reported in recreational fishermen from the Great Lakes [164] and in adults who consumed PCB-contaminated fish from the upper Hudson River [49, 50]. Similarly, humans who consumed high amounts of whale meat, which typically has very high PCB levels, were found to be at increased risk for PD [148]. With respect to the latter, it should be noted that a prospective study reported no association between PCB exposure history and serum levels of PCBs before PD diagnosis [202].

A number of studies have examined the effects of occupational PCB exposures on adult neurologic function. Several studies provide suggestive evidence that occupational PCB exposures exert subtle influences on adult behavior and well-being. For example, PCB exposures during adulthood were associated with moderately lower well-being scores for teachers who worked in PCB-contaminated school buildings [143]. In another study, PCB-exposed firefighters exhibited higher rates of depressive disorders compared with respective control groups [94]. Five additional studies have provided evidence of a positive correlation between PCB body burden and depressive symptoms in older PCB-exposed individuals [48, 49, 55, 56, 151]. Results from one of these studies, a cross-sectional study, indicated a significant negative influence of NDL PCB exposure on urinary levels of homovanillic acid, which is a metabolite of dopamine [151].

There have also been indications of an association between PCBs and increased risk of Parkinson's disease (PD). An early study of the caudate nucleus obtained post mortem from PD patients and from controls not diagnosed with PD showed significantly higher concentrations of NDL PCB 153 in the PD tissue [30]. More recently, a study of postmortem brain tissues from control patients and those diagnosed with PD disease obtained from the Emory University Brain Bank showed NDL PCB congeners 153 and 180 were significantly elevated in the brains of PD patients. When stratified by sex, the female PD group demonstrated significantly elevated concentrations of total PCBs and specifically congeners NDL PCB 138, 153, and 180 compared to controls, whereas PCB concentrations in males were not significantly different between control and PD groups [68]. Epidemiologic studies have similarly noted sex differences in neurodegenerative outcomes related to PCB exposures. For example, Steenland and colleagues [181] demonstrated that female electrical capacitor workers were more likely to die from either Parkinson's disease (PD) or amyotrophic lateral sclerosis than their male counterparts. Seegal and colleagues [171] reported that occupational exposure to PCBs significantly decreased striatal dopamine terminal densities only in women. In a separate cohort of women participating in the Nun Study who had no clinical signs or symptoms of PD at death, elevated levels of the same three PCB congeners were observed in brain tissue that exhibited moderate nigral depigmentation compared to subjects with mild or no depigmentation [68]. The reasons for these sexually dimorphic findings remain unexplained. One potential explanation, particularly for older women who were either peri- or post-menopausal during occupational exposure, is decreased neuroprotection because of reduced levels of circulating estrogen [197]. Another possible explanation is that PCBs are lipophilic [93], and women typically have higher PCB body burdens due to a higher body fat content compared to men [10].

The limited human epidemiological data available regarding associations between PCB exposure and frank neuropathological changes associated with Parkinson's disease is mixed with studies supporting [30, 68, 148] and not supporting [181, 202] an association between occupational PCB exposure with Parkinson's disease. It should be emphasized that the available occupational literature is limited by a paucity of specific PCB biomarkers and exposures likely included other organohalogens. Nevertheless, the environmental persistence of PCBs and evidence of continued human exposure to PCBs, combined with robust animal and human evidence of PCB neurotoxicity, provide strong justification for further longitudinal studies of early life exposures and later neurodegenerative pathology in aging. Importantly, future epidemiological studies need to incorporate in their sampling and exposure assessments the more recent literature that clearly indicates that not all PCB congeners are equivalent in their neurotoxic potential. As reviewed above, there is strong experimental evidence that NDL PCB congeners are significantly more neuroactive than DL congeners. Among the NDL PCBs there appears to be a stringent structure–activity relationship that includes a significant degree of stereoselectivity among the 19 chiral PCBs. Thus, using the sum of the PCBs or a small number of sentinel PCBs, such as PCB 153, to reflect exposure may limit or underestimate associations with neurotoxicity throughout the lifespan.

## Are PCBs the tip of the NDL-POP iceberg?

### Anthropogenic and naturally synthesized polybrominated diphenyl ether (PBDEs)

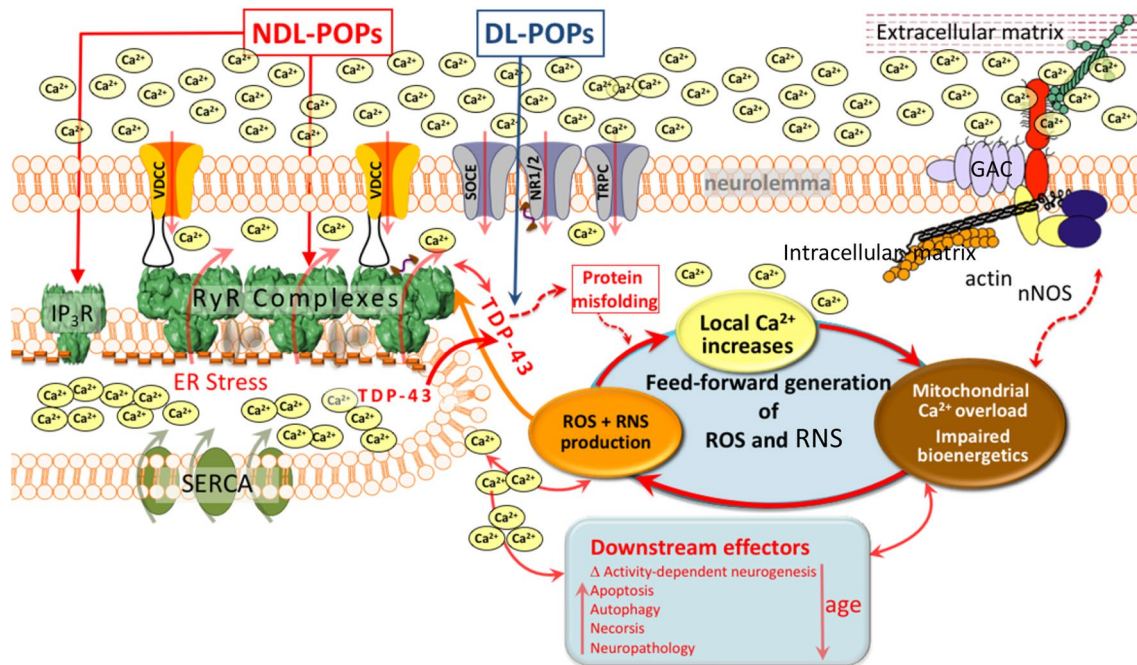
PBDE flame retardants share several common chemical properties with NDL PCBs, including a noncoplanar orientation of the phenyl rings (Fig. 1). Halogen and hydroxy substitutions at *ortho*-, *meta*- and *para*-positions exhibit similar structure–activity profiles toward triggering  $\text{Ca}^{2+}$  release from microsomal vesicles [96] as have been reported for NDL-PCBs [47, 53, 54, 71, 72, 147]. PBDEs were shown to cause intracellular  $\text{Ca}^{2+}$  dyshomeostasis in vitro by altering both ionotropic glutamate receptors [32] and RyR channels [94], and perturbations in the functional coupling of these channels has been linked to degenerative neuropathology [152]. Like PCBs, PBDEs alter TH signaling [34], and increase oxidative stress [31, 130]. Several studies have indicated that the neurotoxic activities of NDL PCBs, PBDEs and their hydroxylated metabolites are additive, and that a dose addition model for estimating risk may be appropriate [34, 71]. However, unlike NDL PCBs, which selectively alter dendritic complexity, neuronal-glia co-cultures from rat hippocampus exposed to BDE-47 or BDE-49 exhibited delayed neuronal polarization resulting in significant inhibition of axonal outgrowth during



the first few days in vitro, but no later influences on dendritic complexity [24]. Interestingly, the axon inhibitory effects of these PBDEs were blocked by pharmacological antagonism of RyR or siRNA knockdown of RyR2, indicating common molecular mechanisms affecting two distinct developmental processes of neuronal connectivity. The explanation for the differential cellular effects of NDL PCBs and PBDEs despite a common molecular mechanism of action remains unknown but may possibly reflect differential effects on distinct RyR isoforms that are spatially segregated [7, 112]. A recent systematic review regarding the effects of developmental exposure to POPs on intelligence or ADHD and attention-related behavioral conditions in humans concluded that there is sufficient evidence supporting an association between developmental PBDE exposure and reduced IQ [108], but a similar association was not observed for PCBs [221].

Finally, contemporary sources of organohalogenes produced as disinfection byproducts [95, 106, 218] and biosynthesized by marine organisms [1] are receiving considerable attention as emerging health risks because of their abundance, persistence, and potential to mimic natural organohalogenes produced by bacteria that serve signaling and/or toxicological functions in marine environments.

Recently 34 organohalogenes from anthropogenic and marine sources, including hydroxylated PBDEs, bromopyrroles and bromobipyrroles (Fig. 1), and bromoindoles were tested to identify compounds active towards RyRs. Several of these compounds were found to be highly active in not only sensitizing RyR channel activation, but also inhibiting microsomal sarcoplasmic/endoplasmic reticulum (SR/ER)  $\text{Ca}^{2+}$  ATPase (SERCA) [223]. The temporal and spatial properties of SR/ER  $\text{Ca}^{2+}$  signals not only depend on the density and activities of RyRs and  $\text{IP}_3\text{R}$  channels, but also the activity of ATP-dependent driven SERCA pumps, which are necessary for accumulating luminal  $\text{Ca}^{2+}$  and setting the ionic force that drives  $\text{Ca}^{2+}$  signal properties. Several halopyrroles formed as disinfectant byproducts that are detected in waste and drinking water treatment streams have recently been shown to exert cytotoxic, genotoxic and carcinogenic activity in mammalian cell culture models [153, 154, 218]. Whether they have neurotoxic effects as predicted by their mechanistic similarity to NDL PCBs remains to be determined. However, the risk of developmental neurotoxicity is plausible given recent evidence that organohalogenes of bio-synthetic and anthropogenic origins accumulate in marine mammals and human breast milk [3, 175, 196].



**Fig. 2** Heuristic hypothesis of modeling known mechanisms by which nondioxin-like (NDL) and dioxin-like (DL) persistent organic pollutants (POPs) influence mechanisms converging through their respective activities on ryanodine receptor (RyR) calcium channel complexes and expression of TDP-43. Expression of TDP-43 is significantly upregulated by a variety of DL environmental chemicals [4] and TDP-43 mediated neurotoxicity proceeds by  $\text{Ca}^{2+}$  dyshomeostasis and ER stress [37, 89] via mechanisms requiring RyR channel activity [2]. Such a model could serve as a basis for improving our

understanding of adverse outcome pathways that contribute to risk for neurodevelopmental and neurodegenerative disorders, and the relationships between them. *GAP* glycoprotein anchoring complex, *IP<sub>3</sub>R* inositol 1,4,5-trisphosphate receptor, *nNOS* neuronal nitric oxide synthase, *ROS* reactive oxygen species, *RNS* reactive nitrogen species, *SOCE* store-operated calcium entry channel, *SERCA* sarcoplasmic/endoplasmic reticulum calcium ATPase, *TDP-43* transactive response DNA binding protein 43, *TRPC* canonical transient receptor protein, *VDCC* voltage-dependent calcium channel

## Concluding remarks

There is compelling evidence from animal research and human epidemiological studies that PCBs are developmental neurotoxicants that are associated with a number of early cognitive and motor deficits later in life. Although less well studied, the influence of PCB exposure on the risk, onset and severity of late-onset neurodegenerative disorders are suggestive. It is important to note that the diversity of PCB chemistry (degree and position of chlorination, and their major hydroxylated and sulfonated metabolites) make the design of animal and epidemiological studies more challenging, as few studies include analysis of the major DL- and NDL-congeners and metabolites present in human tissues for their biochemical and/or behavioral outcomes, which can vary significantly by population demographics, age, sex and specific life-stages of exposure.

Experiments designed to assess the consequences of long-term exposures to PCBs and related environmental organohalogenes on aging-related neuropathology present unique set of challenges. Aging-related declines in cognitive capacity, clinical dementia, and the major neurodegenerative disorders such as Parkinson's disease (PD), Alzheimer's disease (AD) and Amyotrophic Lateral Sclerosis (ALS) have both genetic and environmental components that contribute to temporal onset, progression and severity [69, 122, 199]. Gene by environment interactions have also been implicated in several neurodevelopmental disorders [70, 113, 176]. Future research should encourage approaches that test convergent biologically plausible mechanisms. For example, the transactive response DNA binding protein 43 (TDP-43) has been implicated in the etiology of several aging-related neurodegenerative disorders, including PD [138], AD [127], ALS [86], frontotemporal dementia [18], and idiopathic late-onset dementia [123].

RyR dysfunction has been implicated in several aging disorders and may provide a focal point for more detailed studies of chronic exposures to NDL organohalogenes and neuropathological markers. ALS may serve as a proof of principle for studying convergent mechanisms. Expression of TDP-43 is significantly upregulated by a variety of DL environmental chemicals [4] and TDP-43 mediated neurotoxicity proceeds by  $\text{Ca}^{2+}$  dyshomeostasis and ER stress [37, 89] via mechanisms requiring RyR channel activity [2]. Considering the widespread distribution of dioxin-like and NDL environmental chemicals, TD-43 and RyR (Fig. 2) could serve as a basis for improving our understanding of adverse outcome pathways that contribute to risk for neurodevelopmental and neurodegenerative disorders. A better understanding of the relationships between the neurodevelopmental impairments

highlighted in this review and their potential impact on neuropathological sequelae later in life are needed.

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