Ancestry group	Ν	percentage (%)	N cases	N controls	Stratification <sup>a</sup>		
HIS	329	41.6%	148	181	PC1****; PC2***; PC5**		
CAU	273	34.5%	141	132	NS		
ASN	122	15.4%	68	54	PC9*		
SOUTH ASN	23	2.9%	10	13	NS		
AFR	21	2.7%	11	10	PC8*		
OTHERS	22	2.8%	10	12			

 Table S1. Self-reported ancestry of n=790 mothers

<sup>a</sup> Significant differences across ancestry components between case-control data tested with two-sample Mann-Whitney Wilcoxon test; \*\*\*\*p-value < 0.0005; \*\*\* P<0.001; \*\* P<0.01; \*P<0.05; . P<0.1

Abbreviations: CAU: Caucasians; ASN: Asians; HIS: Hispanics; SOUTH ASN: Indians; AFR: African-Americans; OTHERS: mixed ancestry, unknown ancestry, other ancestry (based on self-reported ancestry groups); PC: ancestry components

	age		multip	arity	educ	ation	count	try of birth	mont	h of birth	offspri	ng gender	ancestry
compound	dir	Р	dir	Р	dir	Р	dir	Р	dir	Р	dir	Р	P
Persistent Pestic	ides												
p,p'-DDE	+	***	-	***	-	**	+	***	+	**	+		***
T-NONA	+	***	-	***	-	NS	-	***	-	NS	-	NS	**
Polybrominated	dipher	nyl eth	ers an	d									
biphenyls													
BB-153	+	***	+	NS	+	***	-	***	+	**	-	NS	**
BDE-28	-	*	+	*	-	NS	-		-	*	+	NS	NS
BDE-47	-	***	+	*	+	NS	-	**	-	NS	+	NS	NS
BDE-99	-	***	+	***	-	NS	-		-	NS	+	NS	*
BDE-100	-	***	+	*	+	•	-	**	-		+	NS	NS
BDE-153	-	**	+	NS	-	•	-	***	-		-	NS	***
Sum PBDE	-	**	+	**	+	•	-	**	-	NS	+	NS	*
Polyclorinated bi	phenyls												
PCB28	+	NS	-	NS	-	*	+	NS	-	**	+	*	*
PCB99	+	***	-	***	+	**	-		+	NS	-	NS	***
PCB118	+	***	-	***	+	*	-	*	-	NS	+	NS	***
PCB153	+	***	-	***	+	***	+	NS	+		-	NS	***
PCB170	+	***	-	**	+	**	+	NS	+	NS	-	NS	***
PCB180	+	***	-	*	+	***	+	NS	+	NS	-	NS	***
PCB187	+	***	-	NS	+	***	+	NS	+	NS	-	NS	***
PCB194	+	***	-	NS	+	*	-	NS	+	*	-	NS	***
PCB199	+	***	-	NS	+	**	-	*	+	NS	-	NS	***
PCB138/158	+	***	-	***	+	**	+	NS	+	NS	-	NS	***
PCB196/203	+	***	-	NS	+	**	-	**	+	NS	-	NS	***
Sum PCB	+	***	-	•	+		+	NS	+	NS	+	NS	***

Table S2. Significance of sociodemographic confounding factors in linear regression models of 19 organohalogens and PCB and PBDE sums

Abbreviations: *dir*: direction; *P*: p-value; NS: not significant \*\*\*\*p-value < 0.0005; \*\*\* P<0.001; \*\* P<0.01; \*P<0.05; . P<0.1

 Table S3. Compressed maternal genome-wide summary-level results for each compound

**Table S4.** Compressed neonatal genome-wide summary-level results for each compound

Table S5. (	Quartile analysis in EMA	genotyped dataset	consistent with I	EMA full dat	taset showed in Ta	ble 3 in
Lyall et al.,	2016(Lyall <i>et al.</i> 2016).					

	Odds of ASD relative to GP controls									
	Unadjusted OR (95% CI)	AOR (95% CI) $^{^{\wedge}}$	AOR (95% CI) <sup>#</sup>							
PCB138_158										
Q1	1	1	1							
Q2	0.97 (0.91, 1.03)	0.99 (0.93, 1.06)	1.00 (0.94, 1.06)							
Q3	0.99 (0.93, 1.05)	0.98 (0.92, 1.04)	0.97 (0.92, 1.04)							
Q4	1.14 (1.06, 1.21)	1.04 (0.99, 1.10)	1.03 (0.98,1.09)							
PCB153										
Q1	1	1	1							
Q2	0.95 (0.89, 1.01)	0.97 (0.91, 1.03)	0.98 (0.92, 1.03)							
Q3	0.97 (0.91, 1.03)	0.96 (0.90, 1.03)	0.96 (0.90, 1.02)							
Q4	1.17 (1.09, 1.25)	1.07 (1.02, 1.13)	1.06 (1.01, 1.12)							

Abbreviations: ASD Autism spectrum disorder; GP general population controls. Bold indicates statistical significance (p<0.05)

<sup>^</sup>Adjusted for matching factors (offspring gender, month and year of birth), maternal age (continuous), maternal race/ethnicity (Non-Hispanic White, Asian, Black/Pacific Islander/or other, Hispanic, or missing), maternal weight at time of sample collection (quartiles), parity (multi- vs primiparous), and maternal education (<high school, high school, college, graduate). Removal of race/ethnicity from these models did not materially alter estimates; maternal age and parity were the only significant covariates in models and adjustment for these factors only yielded AORs similar to fully adjusted estimates. Models adjusted only for matching factors yielded estimates nearly identical as unadjusted models.

<sup>#</sup>Adjusted for EMA genotyped dataset covariates (offspring gender, month of birth, maternal age, maternal education, country of birth, multiparity, PC1-10)

compound	SNP	CHR	BP	MAF	beta	SE	Р	locus
Persistent Pesticio	les							
p,p'-DDE	rs7259965	19	41521532	0.29	0.32	0.05	7.8x10 <sup>-11</sup>	CYP2B6
T-NONA	rs1181906	7	155977316	0.14	-0.17	0.04	5.7 x10 <sup>-6</sup>	lincRNA
Polybrominated di	phenyl ethers and bip	henyls						
BB-153	rs56385737	18	4414795	0.05	0.44	0.08	3.1x10 <sup>-7</sup>	DLGAP1
BDE-28	rs11789653	9	17993589	0.03	0.65	0.12	3.2x10 <sup>-8</sup>	ADAMTSL1 SH3GL2
BDE-47	rs2855812	6	31472720	0.21	-0.33	0.07	7.9x10 <sup>-7</sup>	MICB
BDE-99	rs7259965	19	41521532	0.29	0.32	0.06	2.1x10 <sup>-8</sup>	CYP2B6
BDE-100	rs2855812	6	31472720	0.21	-0.35	0.07	2.9x10 <sup>-7</sup>	MICB
BDE-153	rs17506613	8	38594656	0.48	-0.27	0.06	1.1x10 <sup>-6</sup>	TACC1
Sum PBDE	rs2855812	6	31472720	0.21	-0.29	0.06	1.8x10 <sup>-6</sup>	MICB
Polyclorinated bipl	henyls							
PCB28	rs11727994	4	91911417	0.35	-0.34	0.07	1.6x10 <sup>-6</sup>	TMSB4XP8
PCB99	rs1812081	2	134279837	0.40	0.13	0.03	4.8x10 <sup>-6</sup>	MGAT5
PCB118	rs8076755	17	13788655	0.02	-0.51	0.11	7.5x10 <sup>-6</sup>	HS3ST3A1 COX10
PCB153	rs76546227	15	33276215	0.02	-0.55	0.11	1.0x10 <sup>-6</sup>	FMN1
PCB170	rs10826182	10	60164820	0.32	0.15	0.03	2.1x10 <sup>-6</sup>	TFAM BICC1
PCB180	rs4946957	6	109641411	0.39	-0.15	0.03	1.2x10 <sup>-6</sup>	CCDC162P CEP57L1 CD164
PCB187	rs10500669	11	6328472	0.23	0.20	0.04	2.5x10 <sup>-7</sup>	PRKCDBP
PCB194	rs75107653	3	4181811	0.02	0.57	0.11	1.3x10 <sup>-7</sup>	SETMAR SUMF1
PCB199	rs4945829	6	109638828	0.43	-0.17	0.03	3.7x10 <sup>-7</sup>	CCDC162P CFP57L1

**Table S6.** Maternal top genetic loci genome-wide and suggestively associated with mid-gestational levels of organohalogens

								CD164
PCB138/158	rs76546227	15	33276215	0.02	-0.59	0.12	9.4x10 <sup>-7</sup>	FMN1
PCB196/203	rs75107653	3	4181811	0.02	0.57	0.11	8.6x10 <sup>-8</sup>	SETMAR SUMF1
PCB196/203	rs4945829	6	109638828	0.43	-0.16	0.03	1.1x10 <sup>-7</sup>	CCDC162P CEP57L1 CD164
Sum PCB	rs35560597	5	52678868	0.01	0.79	0.15	7.9x10 <sup>-8</sup>	FST NDUFS4

In each category the results are shown in alphanumeric order. The genome-wide significant associations are shown in **bold** and the suggestively significant associations are in *italics*. Genome-wide association threshold is  $P=5x10^{-8}$  and suggestive association threshold is  $P=5x10^{-7}$ . Abbreviations: CHR: chromosome; BP: position: SE: standard error

	chr3:rs75107653 - MAF=0.016					chr6:rs4945829 - MAF=0.43				chr11:rs10500669 - MAF=0.23			
compound	Ν	beta	SE	p-value	Ν	beta	SE	p-value	Ν	beta	SE	p-value	
PCB28	763	0.5	0.24	$3.7 \times 10^{-2}$	762	0.05	0.07	NS	760	-0.05	0.08	NS	
PCB118	714	0.24	0.11	$2.7 \times 10^{-2}$	713	-0.07	0.03	2.4x10 <sup>-2</sup>	711	0.04	0.03	NS	
PCB153	755	0.24	0.11	2.4x10 <sup>-2</sup>	754	-0.12	0.03	2.8x10 <sup>-5</sup>	752	0.12	0.03	$4.1 \times 10^{-4}$	
PCB170	727	0.39	0.11	1.9x10 <sup>-4</sup>	726	-0.13	0.03	1.2x10 <sup>-5</sup>	724	0.15	0.03	9.5x10 <sup>-6</sup>	
PCB180	759	0.41	0.11	1.9x10 <sup>-4</sup>	758	-0.14	0.03	3.6x10 <sup>-6</sup>	756	0.15	0.03	1.7x10 <sup>-5</sup>	
PCB187	714	0.38	0.12	1.6x10 <sup>-3</sup>	713	-0.16	0.03	2.6x10 <sup>-6</sup>	711	0.2	0.04	2.5x10 <sup>-7</sup>	
PCB194	712	0.57	0.11	1.3x10 <sup>-7</sup>	711	-0.15	0.03	6.5x10 <sup>-7</sup>	709	0.15	0.03	1.5x10 <sup>-6</sup>	
PCB199	715	0.56	0.12	1.6x10 <sup>-6</sup>	714	-0.17	0.03	3.7x10 <sup>-7</sup>	712	0.17	0.04	5.2x10 <sup>-6</sup>	
PCB138/158	752	0.18	0.11	NS	751	-0.12	0.03	9.1x10 <sup>-5</sup>	749	0.11	0.04	3.1x10 <sup>-3</sup>	
PCB196/203	715	0.57	0.11	8.6x10⁻ <sup>8</sup>	714	-0.16	0.03	1.1x10 <sup>-7</sup>	712	0.17	0.03	1.0x10 <sup>-6</sup>	
Sum PCB	706	0.4	0.12	4.8x10 <sup>-4</sup>	705	-0.04	0.03	NS	703	0.07	0.04	NS	

**Table S7**. Suggestively and nominally significant association between maternal chr3:rs75107653, chr6:rs4945829, chr11:rs10500669 and PCB congeners

In each category the results are shown in alphanumeric order. Suggestively significant associations are shown in *italics*. Suggestive association threshold is P=5x10<sup>-7</sup>. Abbreviations: MAF: minor allele frequency; SE: standard error; NS: not significant

	chr9:rs7	2692916 - N	1AF=0.05		chr18:rs72913475 - MAF=0.08				
compound	Ν	beta	SE	p-value	Ν	beta	SE	p-value	
Persistent Pesticides									
T-NONA	644	0.05	0.07	NS	664	0.15	0.05	5.5x10 <sup>-3</sup>	
Polybrominated diphenyl e	thers and b	iphenyls							
BB-153	635	-0.15	0.09	NS	655	0.16	0.07	2.3x10 <sup>-2</sup>	
BDE-28	629	-0.10	0.08	NS	649	0.26	0.07	1.2x10 <sup>-4</sup>	
BDE-47	686	-0.20	0.12	NS	708	0.51	0.10	2.9x10 <sup>-7</sup>	
BDE-99	666	-0.16	0.12	NS	688	0.33	0.10	7.2x10 <sup>-4</sup>	
BDE-100	675	-0.27	0.13	3.6x10 <sup>-2</sup>	697	0.55	0.10	4.6x10 <sup>-8</sup>	
BDE153	674	-0.29	0.12	2.0x10 <sup>-2</sup>	696	0.44	0.10	7.1x10 <sup>-6</sup>	
Sum PBDE	640	-0.26	0.11	1.8x10 <sup>-2</sup>	660	0.35	0.09	1.8x10 <sup>-4</sup>	
Polyclorinated biphenyls									
PCB99	642	-0.22	0.06	3.0x10 <sup>-4</sup>	709	0.10	0.05	4.3x10 <sup>-2</sup>	
PCB118	644	-0.20	0.07	2.9x10 <sup>-3</sup>	664	0.09	0.06	NS	
PCB153	681	-0.30	0.07	1.0x10 <sup>-5</sup>	703	0.13	0.05	2.0x10 <sup>-2</sup>	
PCB170	654	-0.32	0.07	1.7x10 <sup>-6</sup>	675	0.12	0.55	2.9x10 <sup>-2</sup>	
PCB180	683	-0.34	0.07	7.6x10 <sup>-7</sup>	705	0.12	0.55	3.6x10 <sup>-2</sup>	
PCB187	644	-0.42	0.08	2.8x10 <sup>-8</sup>	664	0.11	0.06	NS	
PCB194	642	-0.31	0.07	7.0x10 <sup>-6</sup>	662	0.10	0.06	NS	
PCB199	645	-0.36	0.07	1.3x10 <sup>-6</sup>	665	0.09	0.06	NS	
PCB138/158	676	-0.26	0.07	2.8x10 <sup>-4</sup>	698	0.11	0.06	NS	
PCB196/203	645	-0.33	0.07	1.5x10 <sup>-6</sup>	665	0.12	0.05	3.5x10 <sup>-2</sup>	
Sum PCB	637	-0.26	0.07	4.9x10 <sup>-4</sup>	657	0.08	0.06	NS	

**Table S8.** Genome wide and nominally significant association in fetal chr9:rs72692916 and chr18:rs72913475 across organohalogens

In each category the results are shown in alphanumeric order. The genome-wide significant associations are shown in **bold** and the suggestively significant associations are in *italics*. Genome-wide association threshold is *P*=5x10<sup>-8</sup> and suggestive association threshold is *P*=5x10<sup>-7</sup>. Abbreviations: MAF: minor allele frequency; SE: standard error; NS: not significant

compound	SNP	CHR	BP	MAF	beta	SE	p-value	locus
PCB199	rs1400087	1	106,108,925	0.07	0.33	0.07	7.6x10 <sup>-7</sup>	lincRNA
PCB180	rs1400087	1	106,108,925	0.08	0.32	0.06	1.1x10 <sup>-7</sup>	lincRNA
PCB199	rs6660284	1	106,115,714	0.08	0.34	0.07	7.2x10 <sup>-7</sup>	lincRNA
PCB180	rs6660284	1	106,115,714	0.08	0.33	0.06	1.2x10 <sup>-7</sup>	lincRNA
BDE-100	rs829661	2	30,726,691	0.19	0.37	0.07	2.3x10 <sup>-7</sup>	LCLAT1
BB-153	rs114147886	3	17,854,598	0.02	0.82	0.15	7.6x10 <sup>-8</sup>	near TBC1D5
BDE-28	rs76882575	3	179,782,262	0.04	0.49	0.10	8.1x10 <sup>-7</sup>	near <i>PEX5L</i>
PCB118	rs11755413	6	125,846,604	0.10	0.28	0.05	2.8x10 <sup>-7</sup>	between <i>HEY2</i> and <i>HDDC2</i>
BDE-47	rs1034591	7	114,443,038	0.43	-0.27	0.05	6.3x10 <sup>-7</sup>	between <i>FOXP2</i> and <i>MDFIC</i>
PCB187	rs72692916	9	8,446,518	0.05	-0.42	0.08	2.8x10 <sup>-8</sup>	PTPRD
BDE-28	rs72896141	11	5,845,481	0.05	0.40	0.08	5.6x10 <sup>-7</sup>	TRIM5
BB-153	rs763311	12	12,796,067	0.01	0.89	0.17	1.6x10 <sup>-7</sup>	CREBL2
PCB138/158	rs61989804	14	65,697,140	0.05	-0.38	0.07	1.3x10 <sup>-7</sup>	pseudogenes
BDE-153	rs8054028	16	73,159,970	0.17	-0.39	0.07	2.1x10 <sup>-7</sup>	near c16orf47
BDE-99	rs117896058	18	8,168,060	0.01	1.33	0.26	5.4x10 <sup>-7</sup>	PTPRM
BDE-100	rs72913475	18	44,170,156	0.08	0.55	0.10	4.6x10 <sup>-8</sup>	LOXHD1

**Table S9.** Fetal genetic loci genome-wide, suggestively and nominally associated with mid-gestational levels of organohalogens

In each category the results are shown in alphanumeric order. The genome-wide significant associations are shown in **bold** and the suggestively significant associations are in *italics*. Genome-wide association threshold is  $P=5x10^{-8}$  and suggestive association threshold is  $P=5x10^{-7}$ . Abbreviations: CHR: chromosome; BP: position: SE: standard erro

## **Figure Legends**

**Figure S1.** Multi-dimensional scaling plot of EMA (a) maternal and (b) newborn cohort, and (c) non-overlapping Caucasian, Asian and Hispanic EMA maternal groups. Each individual is represented with a dot and the distance between two individuals represents the genetic distance between them and summarizes the ancestry differences. The X-axis represents the first ancestry coordinate that explains 57.6% and 52.7% of the total genetic variance of the (a) maternal and (b) fetal cohort, respectively. The Y-axis represents the second ancestry coordinate that explains 25.3% and 25.5% of the total genetic variance of the (a) maternal and (b) fetal cohort, respectively. The Y-axis represents the second ancestry coordinate that explains 25.3% and 25.5% of the total genetic variance of the (a) maternal and (b) fetal cohort, respectively; (c) Clustering of 90% of EMA mothers in three non-overlapping ethnic groups. Abbreviations: White: Caucasians; ASN: Asians; HIS: Hispanics; SOUTH ASN: Indians; AFR: African-Americans; OTHERS: mixed ancestry, unknown ancestry, other ancestry (based on self-reported ancestry groups); not compared: individuals not used in analyses comparing ancestry groups.

Figure S2. Residual PBDE congener serum levels by ASD outcome after covariate adjustment.

Boxplot of residual levels of BDE-100 (a), BDE-153 (b) and Sum PBDE (c) in mothers of ASD controls (white) and ASD cases (light grey) obtained with a linear regression after adjustment for sociodemographic covariates. Two-sample Mann-Whitney Wilcoxon test: \* P < 0.05; \*\* P < 0.01.

**Figure S3.** Linear non-parametric correlation coefficients for 19 organohalogens. Correlation coefficients across organhalogens (total sum of PBDEs and PCBs were excluded) calculated with Spearman's test. From weakest to strongest, positive correlation coefficients are represented in a red gradient; negative correlation coefficients are displayed in a blue gradient.

**Figure S4**. Linkage disequilibrium regional genomic plots of maternal suggestively associated SNPs with PCBs and BB-153 serum levels. (a) rs35560597 associated with sum PCB maps near *FST* and *NDUFS4* on chromosome 5q11.2 ( $\beta$ =0.79, SE=0.15, *P*=7.9x10<sup>-</sup> <sup>8</sup>), (b) rs75107653 on chromosome 3p26.1 (PCB196/203,  $\beta$ =0.57, SE=0.11, *P*=8.6x10<sup>-8</sup>) maps near *SETMAR* and *SUMF1*, (c) rs4945829 on chromosome 6q21 (PCB196/203,  $\beta$ =-0.16, SE=0.03, *P*=1.1x10<sup>-7</sup>) maps to *CCDC162P* and near *CD164*, (d) rs10500669 on chromosome 11p15.4 (PCB187,  $\beta$ =0.20, SE=0.04, *P*=2.5x10<sup>-7</sup>) is located near *PRKCDBP* and (e) rs56385737 on chromosome18p11.31 (BB-153, beta=0.44, SE=0.08, *P*=3.1x10<sup>-7</sup>) maps to the *DLGAP1* gene The X-axis represents the genomic position; the Y-axis shows the negative logarithm of the observed association p-value for each tested SNP and the recombination rate. Plotted with Locuzoom tool(Pruim *et al.*2011;).



Figure S1



Figure S2



Figure S3



Figure S4