

Title	Demographic, behavioral, dietary, and socioeconomic characteristics related to persistent organic pollutants and mercury levels in pregnant women in Japan
Author(s)	Miyashita, Chihiro; Sasaki, Seiko; Saijo, Yasuaki; Okada, Emiko; Kobayashi, Sumitaka; Baba, Toshiaki; Kajiwara, Jumboku; Todaka, Takashi; Iwasaki, Yusuke; Nakazawa, Hiroyuki; Hachiya, Noriyuki; Yasutake, Akira; Murata, Katsuyuki; Kishi, Reiko
Citation	Chemosphere, 133, 13-21 https://doi.org/10.1016/j.chemosphere.2015.02.062
Issue Date	2015-08
Doc URL	http://hdl.handle.net/2115/87376
Rights	© 2015. This manuscript version is made available under the CC-BY-NC-ND 4.0 license https://creativecommons.org/licenses/by-nc-nd/4.0/
Rights(URL)	https://creativecommons.org/licenses/by-nc-nd/4.0/
Туре	article (author version)
File Information	44_Chemosphere.pdf



1 Demographic, behavioral, dietary, and socioeconomic characteristics related to 2 persistent organic pollutants and mercury levels in pregnant women in Japan.

3

4 Chihiro Miyashita^a, Seiko Sasaki^b, Yasuaki Saijo^c, Emiko Okada^b, Sumitaka Kobayashi^b,

5 Toshiaki Baba^b, Jumboku Kajiwara^d, Takashi Todaka^e, Yusuke Iwasaki^f, Hiroyuki

6 Nakazawa^f, Noriyuki Hachiya^g, Akira Yasutake^h, Katsuyuki Murataⁱ, Reiko Kishi^a*

 $\overline{7}$

^aCenter for Environmental and Health Sciences, Hokkaido University, North 12 West 7 8 9 Kita-ku, Sapporo, 060-0812, Japan; ^bDepartment of Public Health Sciences, Hokkaido University Graduate School of Medicine, North 15 West 7 Kita-ku, Sapporo 060-8638, 10 11 Japan; ^cDepartment of Health Sciences, Asahikawa Medical University, Midorigaoka-Higashi 2-1-1-1, Asahikawa 078-8510, Japan; ^dFukuoka Institute of 12Health and Environmental Sciences, Mukaizano 39, Dazaifu 818-0135, Japan; 13 ^eDepartment of Dermatology, Graduate School of Medical Sciences, Kyushu University, 14Maidashi 3-1-1, Higashi-ku, Fukuoka 812-8582, Japan; ^fDepartment of Analytical 15Chemistry, Faculty of Pharmaceutical Sciences, Hoshi University, 2-4-41 Ebara, 16 Shinagawa-ku, Tokyo 142-8501, Japan; ^gDepartment of Epidemiology, National 17Institute for Minamata Disease, 4058-18 Hama, Kumamoto 867-0008, Japan; ^hGraduate 18 School of Science and Technology, Kumamoto University, 2-39-1 Kurokami, 19 Kumamoto 860-8555, Japan; Department of Environmental Health Sciences, Akita 2021University, Graduate School of Medicine, Akita 010-8543, Japan

22

23 *Corresponding author:

24 Reiko Kishi

25 Center for Environmental and Health Sciences, Hokkaido University, North 12 West 7

26 Kita-ku, Sapporo 060-0812, Japan

27 Tel: +81-11-706-4746, Fax: +81-(0)11-706-4725.

- 28 E-mail: rkishi@med.hokudai.ac.jp
- 29

30 Author e-mail addresses:

31 Chihiro Miyashita: miyasita@med.hokudai.ac.jp

- 32 Seiko Sasaki: <u>sasakis@med.hokudai.ac.jp</u>
- 33 Yasuaki Saijo: <u>y-saijo@asahikawa-med.ac.jp</u>
- 34 Emiko Okada: <u>ekat_oka@yahoo.co.jp</u>
- 35 Sumitaka Kobayashi: <u>sukobayashi@cehs.hokudai.ac.jp</u>
- 36 Toshiaki Baba: <u>baba.toshiaki@gmail.com</u>

- 37 Jumboku Kajiwara: <u>kajiwara@fihes.pref.fukuoka.jp</u>
- 38 Takashi Todaka: todaka@dream.ocn.ne.jp
- 39 Yusuke Iwasaki: <u>iwasaki@hoshi.ac.jp</u>
- 40 Hiroyuki Nakazawa: <u>hironakazawa@jcom.home.ne.jp</u>
- 41 Noriyuki Hachiya: <u>hachiya@nimd.go.jp</u>
- 42 Akira Yasutake: <u>nimdyasutake@yahoo.co.jp</u>
- 43 Katsuyuki Murata: winestem@med.akita-u.ac.jp
- 44

45 Abstract

Persistent organic pollutants and mercury are known environmental chemicals 46 47that have been found to be ubiquitous in not only the environment but also in humans, including women of reproductive age. The purpose of this study was to evaluate the 4849 association between personal lifestyle characteristics and environmental chemical levels 50during the perinatal period in the general Japanese population. This study targeted 322 pregnant women enrolled in the Hokkaido Study on Environment and Children's Health. 5152Each participant completed a self-administered questionnaire and a food-frequency questionnaire to obtain relevant information on parental demographic, behavioral, 5354dietary, and socioeconomic characteristics. In total, 58 non-dioxin-like polychlorinated 55biphenyls, 17 dibenzo-p-dioxins and -dibenzofuran, and 12 dioxin-like polychlorinated biphenyls congeners, perfluorooctane sulfonate, perfluorooctanoic acid, and mercury 56were measured in maternal samples taken during the perinatal period. Linear regression 57models were constructed against potential related factors for each chemical 58concentration. Most concentrations of environmental chemicals were correlated with the 5960 presence of other environmental chemicals, especially in the case of non-dioxin-like polychlorinated biphenyls and, polychlorinated dibenzo-p-dioxins and -dibezofurans 61 62 and dioxin-like polychlorinated biphenyls which had similar exposure sources and persistence in the body. Maternal smoking and alcohol habits, fish and beef intake and 63 household income were significantly associated with concentrations of environmental 64 65 chemicals. These results suggest that different lifestyle patterns relate to varying exposure to environmental chemicals. 66

67

68 Keywords: persistent organic pollutants, mercury, pregnant women, biomarkers, POPs

69

70 1. Introduction

Polychlorinated biphenyls (PCBs), polychlorinated dibenzo-p-dioxins and 7172-dibezofurans (PCDDs/PCDFs), perfluorooctane sulfonate (PFOS), perfluorooctanoic 73acid (PFOA)-categorized as persistent organic pollutants-and mercury (Hg) are 74known environmental chemicals that have been detected ubiquitously in animal samples 75and the environment. Exposure to environmental chemicals during prenatal and neonatal periods, which are considered windows of vulnerability for fetuses, may cause 76 77various toxicities including carcinogenicity, teratogenicity, endocrine, immune, and 78reproductive disruption, and neurobehavioral effects (Clarkson and Magos, 2006; Olsen 79et al., 2009; Todaka et al., 2010; Wigle et al., 2008).

80 Epidemiological studies of Asian, European, and US populations have revealed 81 that environmental chemical levels measured in maternal samples were associated with demographic, behavioral, dietary, and socioeconomic characteristics. Fish and seafood 82 are the main dietary sources of PCB and PCDDs/PCDFs exposure in Japan, Taiwan, 83 Nordic countries, and Italy (Arisawa et al., 2011); whereas, meat products, dairy 84 85 products and fish are the main dietary sources in the US, The Netherlands, and Germany (Larsen, 2006). Potential exposure sources of PFOS and PFOA were reported to be fish 86 and marine mammals, red meat, animal fat, tap (drinking) water, and household dust in 87 Spain, Norway, and Denmark (D'Hollander et al., 2010; Haug et al., 2011). Many 88 reports to date have also found fish/seafood consumption responsible for 89 90 bio-accumulated methylmercury in humans (Clarkson and Magos, 2006; Kim et al., 2008; Ramon et al., 2008). Consequently, it is plausible that the presence of exposure 91 sources and their contribution to whole body burden levels of environmental chemicals 9293 would vary according to the specific characteristics of populations in different countries or regions (Glynn et al., 2007; Halldorsson et al., 2008; Kim et al., 2008; Ramon et al., 94952008; Sonneborn et al., 2008; Brauner et al., 2011; Ibarluzea et al., 2011).

The elimination rate of toxic substances as a reflection of internal metabolism 96 97 is an effective way to detect body burden levels of environmental chemicals..Tobacco smoking and alcohol habits are considered behavioral factors related to altered 98 elimination rates of environmental chemicals. For example, tobacco smoking induces 99 100increased expression of dioxin-metabolizing enzymes, such as cytochrome P450 (CYP) 1A2, leading to enhanced elimination of PCDDs/PCDFs and dioxin-like PCBs 101(DL-PCBs) (Milbrath et al., 2009). Animal and human studies have also demonstrated 102 that fluorinated organic compounds can regulate CYP enzymes (Ishibashi et al., 2008; 103Narimatsu et al., 2011). 104

105

To date, limited epidemiological studies have been conducted in Japan among

pregnant women with no history of accidental poisoning in Japan. Some studies found 106 107 that PCBs and PCDDs/PCDFs in maternal samples increased with maternal age, alanine aminotransferase levels and alcohol intake, as well as decreased with maternal history 108 of delivery and smoking (Arisawa et al., 2011; Tajimi et al., 2005; Nakamura et al., 109 110 2008). However, no study has assessed maternal smoking and alcohol habits during the 111 pre-pregnancy periods, which is considered an important period because chemicals that have a long half-life could be influenced by lifestyle factors during the entire perinatal 112113period. There have also been no current studies to evaluate associations between 114 background exposure levels of environmental chemicals even though certain chemical 115levels could be correlated with the presence of other chemicals in the human body. This 116 information could help in estimating the magnitude of body burden levels after 117 exposure to various chemicals.

Thus, the purpose of this study was to evaluate associations between concentrations of individual chemicals including non-dioxin-like PCBs (NDL-PCBs), PCDDs/PCDFs and DL-PCBs, PFOS, PFOA and Hg and the potential factors responsible for their varied elimination rates and exposure sources in the general Japanese population.

123

124 2. Materials and Methods

125

126 2.1 Study population

We enrolled 514 Japanese women at 23-35 weeks gestation who were visiting 127128the Sapporo Toho Hospital to take part in the Hokkaido Study on Environment and 129Children's Health Study (Kishi et al., 2011) between July 2002 and September 2005 130 (Supplementary Figure 1). In their last trimester, the subjects filled out a 131self-administered questionnaire regarding the following parental information: tobacco 132smoking and alcohol habits during pre- and post-pregnancy; frequency of food consumption during pregnancy of items such as shoreline fish (e.g., saury, Pacific 133134herring, mackerel), pelagic fish (e.g., tuna, bonito, salmon), beef, pork, chicken, milk, 135and eggs; education level; and household income. Estimated intake value for alcohol 136 (g/day) was calculated from a modified self-administered questionnaire about frequency 137 and type of alcohol consumption (Washino et al., 2009).

From enrollment to delivery, 10 subjects dropped out because of intrauterine growth retardation (2), hospital transfer (1), or voluntary withdrawal (7). The medical records for the remaining 504 mother–newborn pairs were used to obtain data on maternal height and weight before pregnancy. To obtain information on maternal fish

intake throughout pregnancy, we contacted subjects within 5 days of delivery. They 142completed part of a food-frequency questionnaire (FFQ) and provided information 143about intake frequency and portion size for 28 fish and seafood items and their 144estimated total fish intake (g/day) was calculated as previously described (Yasutake et 145146 al., 2003) (Supplementary Table 1). We were not able to contact 74 subjects because of 147poor health conditions immediately after delivery. Subjects also provided a sample of their hair for Hg measurements and information on their past history of having their hair 148149permed. This study was conducted with written informed consent from all subjects and 150was approved by the institutional ethics board for epidemiological studies at the 151Hokkaido University Graduate School of Medicine.

152

153 2.2 Experimental and exposure assessment

154A 40-mL blood sample was taken from the maternal peripheral vein in the last trimester, except in those subjects with pregnancy-related anemia, from whom blood 155samples were taken immediately after delivery. All blood samples were stored at -80°C. 156157NDL-PCBs (Supplementary Table 2) and, PCDDs/PCDFs and DL-PCBs levels (Supplementary Table 3) in maternal blood were detected by high-resolution gas 158chromatography/high-resolution mass spectrometry equipped with a solvent-cut 159large-volume injection system at the Fukuoka Institute of Health and Environmental 160 Sciences as previously described (Iida and Todaka, 2003; Todaka et al., 2003; Todaka et 161162al., 2008). NDL-PCBs and, PCDDs/PCDFs and DL-PCBs levels were adjusted by total lipid content (pg/g lipid)(Todaka et al., 2003). Toxic equivalent (TEQ) values were 163 164 calculated by multiplying the concentration of each individual congener of 165PCDDs/PCDFs and DL-PCBs by its specific toxic equivalency factor value (Van den 166 Berg et al., 2006). PFOS and PFOA levels in maternal serum were detected by 167 column-switching liquid chromatography-tandem mass spectrometry at Hoshi University as previously described (Inoue et al., 2004a; Inoue et al., 2004b; Nakata et al., 168 169 2005a; Nakata et al., 2005b). Values below the detection limit were assigned as 50% of 170the detection limit. The remaining samples were not analyzed owing to unavailable or 171insufficient sample volumes for measurement. Total Hg levels were detected in the 1-cm 172hair fiber closest to the scalp (0.7-1.2 mg) by the oxygen combustion-gold 173amalgamation method using an MD-1 atomic absorption detector (Nippon Institute, Co., 174Ltd., Osaka) at the National Institute for Minamata Disease as previously described (Yasutake et al., 2003). Total hair Hg concentration is a convenient biomarker for 175methylmercury exposure because >90% of total hair Hg is methylmercury, which 176 177covalently binds to cysteines in hair proteins (Clarkson and Magos, 2006). Finally, 58

NDL-PCBs, 12 DL-PCBs and 17 PCDDs/PCDFs congeners—were detected in 426
blood samples. PFOS and PFOA, and total Hg were detected in 447 sera samples and
430 hair samples, respectively.

- 181
- 182 2.3 Statistical analysis

In total, 322 subjects that had complete data about concentration of 183environmental chemicals and personal characteristics were included in the statistical 184 185analyses. Subjects were divided into four categories for each of maternal age, BMI, blood sampling period, and fish intake during pregnancy as shown in Table 1. 186 187 Spearman's rank test was used to determine correlations between concentrations of 188 environmental chemicals. The Mann-Whitney U-test and Kruskal-Wallis test were used 189 to evaluate simple associations between subject characteristics and the concentrations of 190 each environmental chemical. Linear regression analyses were performed to evaluate associations between concentrations of environmental chemicals and subject 191192characteristics. Because of skewed distributions in these concentrations, 193 log₁₀-transformed values were used for linear regression analysis. Linear regression models were constructed for explanatory variables that had previously been reported as 194195related to concentrations of environmental chemicals or that were significantly 196 associated with these concentrations by bivariate analysis in this study. Backward 197 stepwise regression was used to eliminate those variables with a p-value >0.1.

198 Subgroup analyses were performed to confirm significant associations between maternal smoking history and alcohol consumption during pregnancy and 199 200 concentrations of each environmental chemical. Duration of maternal smoking (years) 201was used as a continuous explanatory variable in subgroup analyses of subjects with a 202 history of smoking. Alcohol intake levels (g/day), after categorization into four groups 203 according to their quartile distribution, were used in subgroup analyses of a subjects' 204 alcohol consumption during pregnancy. Presence of NDL-PCBs and, PCDDs/PCDFs 205and DL-PCBs congeners were examined in subgroup analyses among alcohol drinkers 206 during pregnancy. Statistical significance was defined as p < 0.05. Statistical analyses 207 were performed using SPSS for Windows version 19.0J (SPSS, Inc., USA).

208

209 3. Results

Parental characteristics based on the self-administered questionnaire and the FFQ are shown in Table 1. Approximately half (52.2%) of the mothers had a history of tobacco smoking; 17.1% of mothers smoked during the pregnancy. There was a history of alcohol consumption in 73.6% of the mothers, and 30.1% of the mothers consumed

alcohol during pregnancy. Median concentrations of NDL-PCBs, PCDDs/PCDFs and 214DL-PCBs, PFOS, PFOA, and hair Hg were 95.1 ng/mL lipid, 13.8 TEQ pg/g lipid, 5.00 215ng/mL, 1.30 ng/mL, and 1.39 µg/g, respectively (Table 2). Table 3 shows the 216217correlations between concentrations of individual environmental chemicals. The 218strongest correlation was found between NDL-PCBs and, PCDDs/PCDFs and DL-PCBs 219(r = 0.80, p < 0.01). In univariate analyses of maternal levels of environmental chemicals in relation to maternal characteristics, levels of environmental chemicals were 220221significantly associated with maternal age at delivery, parity, blood sampling period, 222education level, smoking and alcohol habits, fish intake, frequency of food consumption, 223and annual household income (p < 0.05; Table 4).

224The linear regression models in Figure 1 show the potential relationship between various factors and maternal blood concentrations of NDL-PCBs and, 225PCDDs/PCDFs and DL-PCBs, PFOS, and PFOA, and total hair Hg. Significant positive 226associations with each log₁₀-transformed concentration of environmental chemicals 227228were observed for maternal age, maternal alcohol consumption during pregnancy, fish 229intake, pelagic fish intake, beef intake, and household income (Supplementary Table 4). Significant negative associations with each log10-transformed concentration of 230231environmental chemicals were observed for multiparous subjects, smoking history, and blood sampling period (Supplementary Table 4). In the subgroup analyses of the 168 232subjects with a history of smoking, the duration of tobacco smoking was inversely 233234associated with log₁₀-transformed PFOS values [Figure 2; -0.08 (-0.15, -0.02)]. With additional adjustment for maternal age, statistical significance remained. In the 235236subgroup analysis of the 97 subjects who reported drinking alcohol during their pregnancy, there was no significant association between NDL-PCBs and, 237238PCDDs/PCDFs and DL-PCBs and their congeners with alcohol intake for any quartile 239as well as across quartiles (Table 5; Supplementary Table 5).

240

241 4. Discussion

242

243 4.1 Correlations between concentrations of environmental chemicals

NDL-PCBs and, PCDDs/PCDFs and DL-PCBs are reported to have high lipophilicities and the biological half-life of most their congeners ranges from a few years to approximately 20 years (Todaka et al., 2010). Perfluoroalkyl acids (PFAAs) are reported to distribute mainly in blood serum and the liver as a result of protein fraction binding (Karrman et al., 2010), and the half-lives of PFOS and PFOA are estimated to be 3.8 and 5.4 years, respectively (Olsen et al., 2009). Methylmercury binds to

hemoglobin in the blood, and its half-life is estimated to be 2 months (Clarkson and 250Magos, 2006). The degrees of correlations between environmental chemicals could 251252correspond to differences in their exposure sources as well as their individual 253pharmacokinetics. Almost all of the environmental chemicals, with the exception of 254relationships between PFOS and PCBs, PFOA and PCBs, and Hg and PFOA, were 255significantly correlated, implying that concentrations of certain environmental chemicals could be used to estimate the magnitude of exposure among the general 256257population in Japan to other environmental chemicals, especially those from similar 258exposure sources and with similar persistence in the body.

259

260 4.2 Tobacco smoking

261Maternal smoking history was significantly related to a decline in 262concentrations of PCDDs/PCDFs and DL-PCBs, and PFOS in this study. A previous study reported that tobacco smoking lead to a decrease in PCDDs/PCDFs and DL-PCBs 263264levels because of increased expression of dioxin-metabolizing enzymes after activation 265of the aryl hydrocarbon receptor (Milbrath et al., 2009). PFAAs are known to activate peroxisome proliferator-activator receptor (PPAR), and a study in wild animals 266267suggested the possibility that a signaling pathway exists between receptor PPAa and CYP that promotes elimination of PFAAs from the body after PFAA exposure (Ishibashi 268et al., 2008). Previous epidemiological studies reported inconsistent relationships 269270between PFAA levels and smoking status among pregnant women when categorized by history or current smoking status (Halldorsson et al., 2008; Halldorsson et al., 2012; 271272Jain, 2013; Ode et al., 2013). In a Swedish study, maternal cotinine levels among current smokers were not associated with PFOA and PFOS plasma levels; in fact, PFOA 273274and PFOS plasma levels were significantly lower than those of subjects who had never 275smoked (Ode et al., 2013). Ode discussed that these results could reflect differences in 276lifestyle patterns between smokers and non-smokers that were associated with sources 277of PFOA and PFOS exposure or an enhanced elimination rate of these environmental 278chemicals in smokers. Our study is the first study to indicate an inverse association 279between PFOS concentrations and the duration of tobacco smoking using a linear 280regression model adjusted for confounding factors. This result supports previous studies 281and suggests that a smoking habit may lead to enhanced elimination rate of not only 282PCDDs/PCDFs and DL-PCBs but also PFOS through activation of PPAα and CYP.

283

284 4.3 Alcohol consumption

285

Our results showed that mothers who drank alcohol during pregnancy had

higher blood concentrations of NDL-PCBs and, PCDDs/PCDFs and DL-PCBs. In a 286previous Japanese study that included women of reproductive age, two possible 287288explanations for this positive association between PCDDs/PCDFs and DL-PCBs and alcohol consumption were proposed. The first is that alcohol intake likely affects 289290 hepatic drug-metabolizing enzymes, which could result in slowed elimination of these 291environmental chemicals. The second is that alcohol intake may indicate a greater likelihood of rich, fatty food consumption, which may result in increased 292293PCDDs/PCDFs and DL-PCBs levels (Arisawa et al., 2011). In this study, no significant associations were found across alcohol-intake quartiles and concentrations of 294295NDL-PCBs and, PCDDs/PCDFs and DL-PCBs, and congeners among alcohol drinkers 296during pregnancy. However, concentrations NDL-PCBs and, PCDDs/PCDFs and DL-PCBs in maternal drinkers were higher than those of women who did not drink 297during pregnancy. Because of their long half-lives, NDL-PCBs and, PCDDs/PCDFs and 298DL-PCBs could be influenced by drinking in the pre-pregnancy period as well as that of 299300 pregnancy, on the assumption that maternal alcohol intake could affect the elimination 301 rate of these chemicals. However, in this study, history of alcohol consumption had no association with NDL-PCBs and, PCDDs/PCDFs and DL-PCBs levels. Therefore, 302303 maternal alcohol consumption may reflect subsequent lifestyle patterns during 304 pregnancy that increase concentrations of NDL-PCBs and, PCDDs/PCDFs and 305 DL-PCBs, rather than indicating an effect on hepatic drug-metabolizing enzymes.

306

307 4.4 Food intake

308 We found that meat, especially beef intake, may be an important exposure 309 source of NDL-PCBs in Japan, similar to that in the US and the Europe (Larsen, 2006). 310 In a Japanese food market study, meat provided the second highest contribution to total 311daily dietary intake of PCDDs/PCDFs and DL-PCBs (Sasamoto et al., 2006). Our 312results indicated that fish/seafood, especially pelagic fish, may be an important exposure 313 source for Hg. This is supported by another study that showed that large predatory fish 314 were the largest contributor to total hair Hg among pregnant women in Japan 315(Yaginuma-Sakurai et al., 2009).

316

317 4.5 Other related factors

In agreement with previous studies, a history of parity was associated with decreasing concentrations of NDL-PCBs and, PCDDs/PCDFs and DL-PCBs, PFOS and PFOA, suggesting that reproductive events could play a role in elimination of environmental chemicals from the maternal body (Milbrath et al., 2009; Olsen et al., 322 2009). PFOS and PFOA were inversely associated with gestational age at the time of blood sampling, possibly due to the dilutional effect of plasma volume expansion, 323 especially after the last trimester (Glynn et al., 2007). NDL-PCBs and, PCDDs/PCDFs 324325and DL-PCBs increased with maternal age, which could be explained by previous 326 reports that maternal age might be a good marker for the estimated duration of exposure 327 to chemicals with long half-lives (Milbrath et al., 2009). Hg in hair increased with household income in our study, which is supported by a previous report indicating that 328 329 high socioeconomic status is related to increased fish consumption, dental amalgams 330 and vaccines, which are all associated with increased exposure to Hg (Tyrrell et al., 3312013).

- 332
- 333 4.6 Strengths and limitations

334 This study provides useful information on associations between demographic, behavioral, dietary, and socioeconomic characteristics and background concentrations of 335336 individual chemicals including NDL-PCBs and, PCDDs/PCDFs and DL-PCBs, PFOS, 337 PFOA and Hg during the perinatal period by liner regression models. These characteristics may also influence the level of fetal exposure to environmental 338 339 chemicals through effects on maternal exposure levels. However, we did not collect data 340 during maternal breast-feeding despite indications that this is an important determinant in the body burden of environmental chemicals (Milbrath et al., 2009). Further studies 341342 are also needed to evaluate etiological mechanisms of maternal smoking on the elimination rate of PFOS mediated by PPAa and CYP activation. 343

In conclusion, most concentrations of individual NDL-PCBs, PCDDs/PCDFs 344 345and DL-PCBs, PFOS, PFOA and Hg were correlated, especially the association between 346 NDL-PCBs and, PCDDs/PCDFs and DL-PCBs, which had similar exposure sources 347 and persistence in the body. PCDDs/PCDFs and DL-PCBs and PFOS decreased with maternal smoking history. NDL-PCBs and, PCDDs/PCDFs and DL-PCBs increased 348 349 with maternal alcohol consumption during pregnancy. Total hair Hg increased with 350 household income. Beef and fish/seafood intake may be important exposure sources of 351NDL-PCBs. These results may reflect various lifestyle patterns associated with 352exposure sources and elimination rates of these environmental chemicals.

 $\frac{353}{354}$

355 **Conflicts of Interest**

The authors declare they have no competing financial interests. This study was supported by a Grant-in-Aid for Scientific Research from the Japanese Ministry of

- Health, Labour and Welfare, and from the Japanese Ministry of Education, Culture, Sports, Science & Technology. The funding sources did not play a role in the study design; the collection, analysis and interpretation of the data; the writing of the report; or the decision to submit the article for publication.
- 362

363 Acknowledgements

- We would like to express our gratitude to all study participants and staff members at Sapporo Toho Hospital for their generous collaboration.
- 366

367 **References**

Arisawa, K., Uemura, H., Hiyoshi, M., Kitayama, A., Takami, H., Sawachika, F.,
Nishioka, Y., Hasegawa, M., Tanto, M., Satoh, H., Shima, M., Sumiyoshi, Y., Morinaga,
K., Kodama, K., Suzuki, T., Nagai, M., 2011. Dietary patterns and blood levels of
PCDDs, PCDFs, and dioxin-like PCBs in 1656 Japanese individuals. Chemosphere 82,
656-662.

- 373
- Brauner, E.V., Raaschou-Nielsen, O., Gaudreau, E., LeBlanc, A., Tjonneland, A.,
 Overvad, K., Sorensen, M., 2011. Predictors of polychlorinated biphenyl concentrations
 in adipose tissue in a general Danish population. Environ Sci Technol 45, 679-685.
- 377

380

D'Hollander, W., de Voogt, P., De Coen, W., Bervoets, L., 2010. Perfluorinated
substances in human food and other sources of human exposure. Rev Environ Contam
Toxicol 208, 179-215.

384

Glynn, A., Aune, M., Darnerud, P.O., Cnattingius, S., Bjerselius, R., Becker, W., Lignell,
S., 2007. Determinants of serum concentrations of organochlorine compounds in
Swedish pregnant women: a cross-sectional study. Environ Health 6, 2.

388

Halldorsson, T.I., Fei, C., Olsen, J., Lipworth, L., McLaughlin, J.K., Olsen, S.F., 2008.
Dietary predictors of perfluorinated chemicals: a study from the Danish National Birth
Cohort. Environ Sci Technol 42, 8971-8977.

392

Halldorsson, T.I., Rytter, D., Haug, L.S., Bech, B.H., Danielsen, I., Becher, G.,
Henriksen, T.B., Olsen, S.F., 2012. Prenatal exposure to perfluorooctanoate and risk of
overweight at 20 years of age: a prospective cohort study. Environ Health Perspect 120,
668-673.

397

Haug, L.S., Huber, S., Becher, G., Thomsen, C., 2011. Characterisation of human
exposure pathways to perfluorinated — Comparing exposure estimates with biomarkers
of exposure. Environ Int 37, 687-693.

401

402 Ibarluzea, J., Alvarez-Pedrerol, M., Guxens, M., Marina, L.S., Basterrechea, M.,

<sup>Clarkson, T.W., Magos, L., 2006. The toxicology of mercury and its chemical
compounds. Crit Rev Toxicol 36, 609-662.</sup>

- Lertxundi, A., Etxeandia, A., Goni, F., Vioque, J., Ballester, F., Sunyer, J., 2011.
 Sociodemographic, reproductive and dietary predictors of organochlorine compounds
 levels in pregnant women in Spain. Chemosphere 82, 114-120.
- 406
- 407 Iida, T., Todaka, T., 2003. Measurement of dioxins in human blood: improvement of408 analytical method. Ind Health 41, 197-204.
- 409
- Inoue, K., Okada, F., Ito, R., Kato, S., Sasaki, S., Nakajima, S., Uno, A., Saijo, Y., Sata,
 F., Yoshimura, Y., Kishi, R., Nakazawa, H., 2004a. Perfluorooctane sulfonate (PFOS)
 and related perfluorinated compounds in human maternal and cord blood samples:
 assessment of PFOS exposure in a susceptible population during pregnancy. Environ
 Health Perspect 112, 1204-1207.
- 415
- Inoue, K., Okada, F., Ito, R., Kawaguchi, M., Okanouchi, N., Nakazawa, H., 2004b. 416 417Determination of perfluorooctane sulfonate, perfluorooctanoate and perfluorooctane 418 sulfonylamide in human plasma by column-switching liquid chromatography-electrospray mass spectrometry coupled with solid-phase extraction. J 419420 Chromatogr B Analyt Technol Biomed Life Sci 810, 49-56.
- 421

Ishibashi, H., Iwata, H., Kim, E.Y., Tao, L., Kannan, K., Tanabe, S., Batoev, V.B., Petrov,
E.A., 2008. Contamination and effects of perfluorochemicals in Baikal seal (Pusa
sibirica). 2. Molecular characterization, expression level, and transcriptional activation
of peroxisome proliferator-activated receptor alpha. Environ Sci Technol 42,
2302-2308.

427

Jain, R.B., 2013. Effect of pregnancy on the levels of selected perfluoroalkyl
compounds for females aged 17-39 years: data from National Health and Nutrition
Examination Survey 2003-2008. J Toxicol Environ Health A 76, 409-421.

431

Karrman, A., Domingo, J.L., Llebaria, X., Nadal, M., Bigas, E., van Bavel, B.,
Lindstrom, G., 2010. Biomonitoring perfluorinated compounds in Catalonia, Spain:
concentrations and trends in human liver and milk samples. Environ Sci Pollut Res Int
17, 750-758.

436

Kim, S.A., Jeon, C.K., Paek, D.M., 2008. Hair mercury concentrations of children and
mothers in Korea: implication for exposure and evaluation. Sci Total Environ 402,

439 36-42.

440

Kishi, R., Sasaki, S., Yoshioka, E., Yuasa, M., Sata, F., Saijo, Y., Kurahashi, N., Tamaki,
J., Endo, T., Sengoku, K., Nonomura, K., Minakami, H., 2011. Cohort profile: the
Hokkaido study on environment and children's health in Japan. Int J Epidemiol 40,
611-618.

445

Lackmann, G.M., Angerer, J., Tollner, U., 2000. Parental smoking and neonatal serum
levels of polychlorinated biphenyls and hexachlorobenzene. Pediatr Res 47, 598-601.

448

Larsen, J.C., 2006. Risk assessments of polychlorinated dibenzo- p-dioxins,
polychlorinated dibenzofurans, and dioxin-like polychlorinated biphenyls in food. Mol
Nutr Food Res 50, 885-896.

452

Milbrath, M.O., Wenger, Y., Chang, C.W., Emond, C., Garabrant, D., Gillespie, B.W.,
Jolliet, O., 2009. Apparent half-lives of dioxins, furans, and polychlorinated biphenyls
as a function of age, body fat, smoking status, and breast-feeding. Environ Health
Perspect 117, 417-425.

457

Nakamura, T., Nakai, K., Matsumura, T., Suzuki, S., Saito, Y., Satoh, H., 2008.
Determination of dioxins and polychlorinated biphenyls in breast milk, maternal blood
and cord blood from residents of Tohoku, Japan. Sci Total Environ 394, 39-51.

461

Nakata, H., Nakata, A., Kawaguchi, M., Iwasaki, Y., Ito, R., Saito, K., Nakazawa, H.,
2005a. Development of an analytical method for perfluorochemicals in human plasma
and blood by liquid chromatography-tandem mass spectrometry coupled with
solid-phase extraction using a column-switching technique. Organohalogen Compounds
64, 219-221.

467

Nakata, H., Nakata, A., Okada, F., Ito, R., Inoue, K., Saito, K., Nakazawa, H., 2005b.
Development of online solid-phase extraction-HPLC/MS/MS method for the
determination of perfluorochemicals in human plasma. Bunseki Kagaku 54, 877-884.

471

472 Narimatsu, S., Nakanishi, R., Hanioka, N., Saito, K., Kataoka, H., 2011.
473 Characterization of inhibitory effects of perfluorooctane sulfonate on human hepatic
474 cytochrome P450 isoenzymes: focusing on CYP2A6. Chem Biol Interact 194, 120-126.

475

Ode, A., Rylander, L., Lindh, C.H., Kallen, K., Jonsson, B.A., Gustafsson, P., Olofsson,
P., Ivarsson, S.A., Rignell-Hydbom, A., 2013. Determinants of maternal and fetal
exposure and temporal trends of perfluorinated compounds. Environ Sci Pollut Res Int
20, 7970-7978.

480

Olsen, G.W., Butenhoff, J.L., Zobel, L.R., 2009. Perfluoroalkyl chemicals and human
fetal development: an epidemiologic review with clinical and toxicological perspectives.
Reprod Toxicol 27, 212-230.

484

Ramon, R., Murcia, M., Ballester, F., Rebagliato, M., Lacasana, M., Vioque, J., Llop, S.,
Amurrio, A., Aguinagalde, X., Marco, A., Leon, G., Ibarluzea, J., Ribas-Fito, N., 2008.
Prenatal exposure to mercury in a prospective mother-infant cohort study in a
Mediterranean area, Valencia, Spain. Sci Total Environ 392, 69-78.

489

Sasamoto, T., Ushio, F., Kikutani, N., Saitoh, Y., Yamaki, Y., Hashimoto, T., Horii, S.,
Nakagawa, J., Ibe, A., 2006. Estimation of 1999-2004 dietary daily intake of PCDDs,
PCDFs and dioxin-like PCBs by a total diet study in metropolitan Tokyo, Japan.
Chemosphere 64, 634-641.

494

Sonneborn, D., Park, H.Y., Petrik, J., Kocan, A., Palkovicova, L., Trnovec, T., Nguyen,
D., Hertz-Picciotto, I., 2008. Prenatal polychlorinated biphenyl exposures in eastern
Slovakia modify effects of social factors on birthweight. Paediatric and perinatal
epidemiology 22, 202-213.

499

Tajimi, M., Uehara, R., Watanabe, M., Oki, I., Ojima, T., Nakamura, Y., 2005.
Relationship of PCDD/F and Co-PCB concentrations in breast milk with infant
birthweights in Tokyo, Japan. Chemosphere 61, 383-388.

503

Todaka, T., Hirakawa, H., Kajiwara, J., Hori, T., Tobiishi, K., Yasutake, D., Onozuka, D.,
Sasaki, S., Miyashita, C., Yoshioka, E., Yuasa, M., Kishi, R., Iida, T., Furue, M., 2010.
Relationship between the concentrations of polychlorinated dibenzo-p-dioxins,
polychlorinated dibenzofurans, and polychlorinated biphenyls in maternal blood and
those in breast milk. Chemosphere 78, 185-192.

509

510 Todaka, T., Hirakawa, H., Tobiihi, K., Iida, T., 2003. New protocol of dioxins analysis

- 511 in human blood. Fukuoka Igaku Zasshi 94, 148-157.
- 512

Todaka, T., Hori, T., Hirakawa, H., Kajiwara, J., Yasutake, D., Onozuka, D., Iida, T.,
Furue, M., 2008. Congener-specific analysis of non-dioxin-like polychlorinated
biphenyls in blood collected from 127 elderly residents in Nakagawa Town, Fukuoka
Prefecture, Japan. Chemosphere 73, 865-872.

517

Tyrrell, J., Melzer, D., Henley, W., Galloway, T.S., Osborne, N.J., 2013. Associations
between socioeconomic status and environmental toxicant concentrations in adults in
the USA: NHANES 2001-2010. Environ Int 59, 328-335.

521

Van den Berg, M., Birnbaum, L.S., Denison, M., De Vito, M., Farland, W., Feeley, M.,
Fiedler, H., Hakansson, H., Hanberg, A., Haws, L., Rose, M., Safe, S., Schrenk, D.,
Tohyama, C., Tritscher, A., Tuomisto, J., Tysklind, M., Walker, N., Peterson, R.E., 2006.
The 2005 World Health Organization reevaluation of human and mammalian toxic
equivalency factors for dioxins and dioxin-like compounds. Toxicol Sci 93, 223-241.

527

Washino, N., Saijo, Y., Sasaki, S., Kato, S., Ban, S., Konishi, K., Ito, R., Nakata, A.,
Iwasaki, Y., Saito, K., Nakazawa, H., Kishi, R., 2009. Correlations between prenatal
exposure to perfluorinated chemicals and reduced fetal growth. Environ Health Perspect
117, 660-667.

532

Wigle, D.T., Arbuckle, T.E., Turner, M.C., Berube, A., Yang, Q.Y., Liu, S.L., Krewski,
D., 2008. Epidemiologic evidence of relationships between reproductive and child
health outcomes and environmental chemical contaminants. J Toxicol Env Heal B 11,
373-517.

537

Yaginuma-Sakurai, K., Shimada, M., Ohba, T., Nakai, K., Suzuki, K., Kurokawa, N.,
Kameo, S., Satoh, H., 2009. Assessment of exposure to methylmercury in pregnant
Japanese women by FFQ. Public Health Nutr 12, 2352-2358.

541

Yasutake, A., Matsumoto, M., Yamaguchi, M., Hachiya, N., 2003. Current hair mercury
levels in Japanese: survey in five districts. Tohoku J Exp Med 199, 161-169.

		$Mean \pm SD$	n (%)
Maternal characteristics			
Age at delivery (years)		30.63 ± 4.70	
BMI before pregnancy (kg/m ²)		21.12 ± 3.21	
Parity	≥1		170 (52.8
Blood sampling period	<28 weeks		19 (5.9)
	28 to <36 weeks		144 (44.7
	≥36 weeks		70 (21.2)
	After delivery		99 (30.7)
Education level (years)	>12		191 (59.3
Tobacco smoking history	Yes		168 (52.2
Tobacco smoking during pregnancy	Smoker		55 (17.1)
Alcohol consumption history	Yes		237 (73.6
Alcohol consumption during pregnancy	Drinker		97 (30.1)
Alcohol intake (g/day) during pregnancy		$0.00 (0.00, 0.46)^{a}$	
Fish intake (g/day) during pregnancy		50.00 (30.00, 50.00) ^a	
Frequency of food consumption during pregnancy			
Shoreline fish	≥once/week		155 (48.1
Pelagic fish	≥once/week		178 (55.3
Beef	≥once/week		86 (26.7)
Eggs	≥once/week		322 (100)
Milk	≥once/week		285 (88.5
Paternal characteristics			
Tobacco smoking history	Yes		279 (86.7
Tobacco smoking during their partner's pregnancy	Smoker		225 (69.9
Annual household income (million yen)	≥5		110 (34.2

2 BMI: body mass index

3

^aMedian (minimum, maximum)

1	Table 2.	Concentrations	of environme	ntal chemicals	s in maternal sar	nples ((n = 322))
T	1 aoite 2.	Concentrations	of chivinonine	mai enemieais	5 m maternar sa	iipies (II 522)	£

			Р	ercenti	le	
	Geometric mean	Minimum	25th	50th	75th	Maximum
NL-PCBs (ng/g lipid)	94.0	16.0	66.0	95.1	130	445
DL-PCBs and PCDDs/PCDFs (TEQ pg/g lipid) ^a	13.5	3.17	9.86	13.8	18.3	42.9
PFOS (ng/mL)	4.78	1.30	3.20	5.00	6.98	14.7
PFOA (ng/mL)	1.20	0.25	0.80	1.30	1.80	5.30
Hair Hg (µg/g) ^b	1.35	0.24	0.96	1.39	1.89	7.55

2 ^aTEQs were calculated from the individual congener toxic equivalency factor values (Van den Berg et al., 2006).

3 ^b>90% methylmercury

1 Table 3. Correlation coefficients between individual environmental chemicals (n = 322)

	PCDDs/PCDFs	PFOS	PFOA	Hair Hg
	and DL-PCBs	1105	ITOA	Hall Hg
NDL-PCBs	0.80**	0.07	0.10	0.38**
PCDDs/PCDFs		0.24**	0.14*	0.30**
and DL-PCBs		0.24	0.14	0.30
PFOS			0.25**	0.12*
PFOA				0.03

2 *p <0.05, **p <0.01 by Spearman's rank correlation

			PCDDs/PCDFs			
Characteristics		NDL-PCBs	and DL-PCBs	PFOS	PFOA	Hair Hg
Characteristics		(ng/g lipid)	(TEQ pg/g	(ng/mL)	(ng/mL)	$(\mu g/g)$
			lipd)			
Age at delivery (years)	<25	60.78**	9.80**	5.0	1.4	1.4
	25 to <30	86.0	13.7	5.3	1.2	1.4
	30 to <35	101.3	14.4	5.0	1.4	1.3
	≥35	136.4	16.9	4.2	1.2	1.7
BMI at delivery (kg/m ²)	<18.5	89.7	13.7	5.4	1.4	1.3
	18.5 to <25	97.2	13.9	5.0	1.3	1.4
	25 to <30	108.4	13.4	4.4	1.4	1.2
	≥30	77.3	12.8	4.3	1.2	1.1
Parity	0	101.0	14.6**	5.50**	1.50**	1.4
	≥1	90.8	13.3	4.6	1.0	1.4
Timing of blood sampling	<28 weeks	114.6	16.7	6.4**	1.8**	
	28 to <36 weeks	108.6	13.9	5.6	1.5	
	≥36 weeks	102.1	13.8	4.6	1.2	
	After delivery	108.2	13.5	3.8	1.2	
Education level (years)	≤12	89.4	12.8	4.8	1.3	1.4
	>12	99.2	14.1	5.3	1.4	1.4
Tobacco smoking history	No	101.0	15.2**	5.30**	1.4	1.5
	Yes	87.8	12.6	4.7	1.2	1.3
Tobacco smoking during pregnancy	Nonsmoker	96.9	14.1	5.0	1.3	1.4
	Smoker	84.8	12.2	4.8	1.2	1.4
Alcohol consumption history	No	86.0	13.3	4.8	1.2	1.3
	Yes	101.0	14.0	5.0	1.4	1.4
Alcohol consumption during pregnancy	Non-drinker	92.1	13.8	5.0	1.3	1.4
	Drinker	101.0	13.8	5.1	1.3	1.4
Alcohol intake (g/day) during pregnancy	Quartile 1 (<0.73)	0.1	0.0	0.0	0.0	0.1
	Quartile 2 (0.73 to <1.52)	96.0	12.3	4.1	1.2	1.3
	Quartile 3 (1.52 to <3.52)	117.3	13.7	5.6	1.2	1.6
	Quartile 4 (≥3.52)	100.4	15.7	6.2	1.4	1.4
Fish intake (g/day) during pregnancy	Quartile 1 (<25)	101.5*	14.0	4.6	1.3	1.5**
	Quartile 2 (25 to <38.75)	84.8	13.0	5.3	1.4	1.3
	Quartile 3 (38.75 to <50)	101.3	13.8	4.9	1.4	1.4
	Quartile 4 (≥50)	104.1	14.3	5.0	1.2	1.7
Frequency of food consumption during pregnancy						
Shoreline fish	<once td="" week<=""><td>87.2</td><td>13.3</td><td>4.7</td><td>1.2</td><td>1.28*</td></once>	87.2	13.3	4.7	1.2	1.28*
	≥once/week	101.0	14.5	5.3	1.3	1.5

1 Table 4. Maternal environmental chemical levels in relation to characteristics (n = 322)

Pelagic fish	<once th="" week<=""><th>94.8</th><th>13.4</th><th>4.7</th><th>1.3</th><th>1.25**</th></once>	94.8	13.4	4.7	1.3	1.25**
	≥once/week	95.6	14.2	5.4	1.3	1.5
Beef	<once td="" week<=""><td>95.0</td><td>13.7</td><td>5.0</td><td>1.3</td><td>1.34*</td></once>	95.0	13.7	5.0	1.3	1.34*
	≥once/week	100.0	14.1	5.2	1.3	1.5
Egg	<once td="" week<=""><td>92.6</td><td>13.6</td><td>4.10**</td><td>1.3</td><td>1.28**</td></once>	92.6	13.6	4.10**	1.3	1.28**
	≥once/week	95.1	13.8	5.0	1.3	1.4
Milk	<once td="" week<=""><td>66.2</td><td>11.0*</td><td>4.3</td><td>1.2</td><td>1.3</td></once>	66.2	11.0*	4.3	1.2	1.3
	\geq once/week	99.2	14.1	5.0	1.3	1.4
Paternal characteristics						
Tobacco smoking history	No	97.5	15.1	5.4	1.3	1.3
	Yes	94.7	13.7	5.0	1.3	1.4
Tobacco smoking during their partner's pregnancy	Non-smoker	102.0	15.1*	5.4	1.4	1.4
	Smoker	92.5	13.1	4.8	1.3	1.4
Annual household income (million yen)	<5	91.8**	13.3**	4.7	1.2	1.27*
	≥5	113.0	15.8	5.5	1.5	1.5

1 Values shown are medians. BMI: body-mass index

 $2 \qquad *p < 0.05, **p < 0.01$ by the Mann–Whitney U-test and Kruskal–Wallis test

3

- 1 Table 5. Partial regression coefficients (95%CI) for environmental chemical concentrations from mothers
- 2 who consumed alcohol during pregnancy (n = 97)

Quartiles by alcohol intake	NDL-PCBs		PCDDs/PCDFs and DL-PCBs		
(n, range in g/day)	B (95%CI) p for trend ^a		B (95% CI)	p for trend ^a	
Quartile 1 (n = 26, <0.73)	reference	0.802	reference	0.404	
Quartile 2 (n = 27, 0.73–1.52)	0.07 (-0.05, 0.18)		0.06 (-0.03, 0.16)		
Quartile 3 (n = 24, 1.52–3.52)	0.07 (-0.05, 0.18) 0.08 (-0.		0.08 (-0.02, 0.17)		
Quartile 4 (n = 20, \ge 3.52)	0.00 (-0.12, 0.13)		0.04 (-0.07, 0.14)		

3 B: partial regression coefficient provides the expected change in the log₁₀-transformed environmental chemical

4 concentrations between quartiles in the regression linear model, adjusted for maternal age, parity, smoking history,

5 fish intake, shoreline fish intake, and beef intake.

6 ^aQuartiles are represented as ordinal variables.

 $\mathbf{7}$











