Exposure of Remote Maritime Populations to Coplanar PCBs

by Éric Dewailly, John Jake Ryan, Claire Laliberté, Suzanne Bruneau, Jean-Philippe Weber, Suzanne Gingras, and Gaétan Carrier

Introduction

Fish consumption is a major contributor of organochlorinated compound (OCs) intake of humans. To minimize human exposure, regulations on fish contamination in Canada have been set at 2 mg/kg for polychlorinated biphenyls (PCBs) and 20 ng/kg for (2,3,7,8-tetrachlorodibenzo-p-dioxin) (TCDD). All of these foods for the general Canadian population, fish contains the highest levels of PCBs (1), and one of the highest for polychlorinated dibenzo-p-dioxins (PCDDs) and polychlorinated dibenzofurans (2). Because the daily intake of fish products in Canada by the general population is only moderate, i.e., about 12 g (3), the percentage daily intake of these contaminants from fish is not high. However, various groups and communities consume large quantities of fish (e.g., sportfishermen, commercial fishermen, and native peoples), and these individuals are potentially highly exposed to these contaminants.

In the Province of Québec, two remote populations are of special concern. About 6500 Inuit people of Arctic Québec are scattered in 14 communities along the 1500 km of seashore. Arctic Québec is located between 55° and 63° north latitude (Fig. 1).

The main source of proteins and lipids for these people arises from fish (Arctic char, cod) and sea mammal consumption (seal, beluga, walrus). The second population is 2000 km north east from Montréal on the north shore of the Gulf of the St. Lawrence River. Fishing is the main activity of these 6500 people (14 settlements) of Caucasian origin. These two populations are linked to the southern part of Canada only by plane. Because dietary habits of these populations are based principally on seafood, we decided to evaluate their biological exposure for different contaminants including organochlorines.

Organochlorines are part of a group of halogenated aromatic hydrocarbons (HAHs) and consist of chlorinated pesticides, PCBs, and by-products of industrial processes such as furans and dioxins. PCBs, PCDDs, and polychlorinated dibenzofurans (PCDFs) are probably the most toxic organochlorine compounds. Recent developments in experimental toxicology gave convincing evidence that these toxic halogenated aromatic hydrocarbons have common toxic properties due to a common receptor-mediated mechanism of action (4). Toxic equivalent factors (TEFs) were developed for PCDDs and PCDFs and more recently for PCBs (5). Compared with 2,3,7,8-TCDD toxicity, specific PCB congeners have TEF values ranging from 0.1 for non-ortho coplanar 3,3',4,4',5-,penta CB (IUPAC no. 126) to 0.00002 for di-ortho coplanar PCBs, which are the predominant congeners found in humans. In human tissues and fluids of nonoccupationally exposed populations, recent work shows that coplanar PCBs are present at higher levels than dioxins and furans and, based on the TEF concept, they account for the majority of possible toxic effects (6–8).
This paper presents data on coplanar PCB levels in the breast milk of the Inuit population of Arctic Québec and in the blood of fishermen of the Gulf of the St. Lawrence River in comparison with nonexposed populations. We also discuss the relative importance of these compounds with PCDD and PCDF levels.

**Populations and Methods**

**Populations and Sampling**

*Inuit Study.* Between July 1989 and July 1990, 224 live births occurred among the Inuit studied in Arctic Québec. One hundred nineteen babies were breast fed and 109 of these Inuit women provided us with a 60-mL milk sample collected within the first 3 days after delivery. Chlorinated pesticides and seven PCB congeners were quantitated in these 109 milk samples. Forty randomly selected individual subsamples were used for coplanar PCB and PCDD–PCDF determination analyses, 35 other subsamples were used for complementary measurement of the 118 and 170 (IUPAC nos.) congeners.

*Fishermen Study.* One hundred eighty-five volunteers (fishermen and their spouses) from 7 of the 14 settlements were randomly selected within the local Fishermen Association list. Blood samples were collected in March 1990 for various analytes (organochlorines, heavy metals, biochemistry, biomarkers). For coplanar PCB and PCDD/PCDF determination, only those 10 individuals with the highest total PCB levels were chosen. Because a more complete evaluation is in progress, only preliminary results on coplanar PCBs are presented. Among the three
non-ortho coplanar PCBs only the penta CB (no. 126) and hexa CB (no. 169) have been analyzed.

**Laboratory Procedures**

2,3,7,8-Chloro-substituted congeners of PCDDs and PCDFs, coplanar PCBs 3,3′,4,4′-tetra CB, 3,3′,4,4′,5-penta CB, and 3,3′,4,4′,5,5′-hexa CB were determined by high-resolution mass spectrometry (HRMS). Milk samples, 25–40 g, were fortified with nine 
\(^{13}\)C-labeled PCDD and PCDF and three 
\(^{13}\)C-labeled PCB internal quantitation standards. These internal standards represented each of the PCDD and PCDF homologs and the three coplanar PCBs of interest (IUPAC nos. 77, 126, and 169). The milk samples were mixed with an aqueous solution of sodium oxalate, ethyl ether and ethanol, and then extracted with hexane. The hexane extracts were washed and then concentrated to constant weight, and the percent lipid was determined gravimetrically.

The lipid residue was diluted in hexane and was cleaned using a sulfuric acid-silica gel slurry followed by elution through a neutral/acid-silica gel chromatographic column. Subsequent clean-up steps included separation of the PCDDs/PCDFs and coplanar PCBs from interferences using neutral alumina 22 and Carbopack C/Celite columns. The eluent from the Carbopack C/Celite was concentrated to 5 \(\mu\)L. The final extracts were analyzed using a VG 7025S HRMS at a mass resolution of 10,000. Separation was achieved using a 60-m DB-5 column. Two ions characteristic of each PCDD and PCDF homolog, the coplanar PCBs, and the respective internal quantitation standards were monitored for each analysis. Identification of the PCDDs, PCDFs, and coplanar PCBs was based on retention time information and the comparison of the ratios of the characteristic ions with theoretical values.

For blood plasma analyses, isotope-labeled 
\(^{13}\)C-labeled PCDD, 
\(^{13}\)C-labeled PCDF, and 
\(^{13}\)C-labeled PCBs were added followed by ethanol and ammonium sulfate, and the mixture was extracted with hexane. After weighing the total hexane extracts to constant weight for the sample lipid content, the extract was defatted with \(\text{H}_2\text{SO}_4\), purified on columns of silicate, Florisil, and carbon, and measured by GC–MS according to Ryan et al. (9). The same extract was used to measure both the PCDDs/PCDFs and coplanar PCBs using the isotope dilution MS technique.

All data presented in Tables 1 and 2 are arithmetic means. Comparison data for the Inuit study are taken from a provincial survey (536 women) from which 16 pools of 6 milk samples each (96 milk samples) were constituted and analyzed at the same period by the same laboratory. In the fishery study we used comparison data from a pool of 10 plasma samples collected in 1988 from Red Cross donors from Ontario. For mono-ortho and di-ortho coplanar PCB comparison, we used new data from 59 blood samples collected in 1989 among firefighters in Québec.

Levels of PCBs and PCDDs/PCDFs were detected in all samples except for the 59 control blood samples, where mean values of PCB congeners were calculated only on plasma samples with levels above the detection limit (0.1 \(\mu\)g/L). TEFs used in Table 1 are international values for PCDDs and PCDFs (4) and those proposed by Safe for PCBs (5).

**Results and Discussion**

As shown in Table 1, levels of non-ortho coplanar PCBs range from 24.7 to 220.9 ng/kg for Inuit women, three times more than comparison levels except for the hexa CB (no.169), which is seven times higher in Inuit samples. These differences are also observed for mono-ortho and di-ortho coplanar PCBs with ratios ranging from 1/3 (no. 118) to 1/10 (nos. 153 and 180).

The pattern of distribution of the three non-ortho coplanar PCBs in Inuit mothers’ milk is different from that of Caucasian...
women. Congener 169 is the most prominent in Inuit milk, whereas congener 126 predominates in control samples. For PCDDs and PCDFs (data not shown), differences are not so great: 13.3 ng/kg TEQs for Caucasian women and 19.1 ng/kg TEQs for Inuit women.

The relative contributions for toxicity of PCBs and PCDDs/PCDFs are presented in Figure 2. For di-ortho, mono-ortho, non-ortho coplanar PCBs and PCDDs/PCDFs, TEQs are, respectively, 17.3, 58.7, 32.2, and 19.1 ng/kg for Inuit women and 2.1, 17.4, 9.8, and 13.3 for Caucasian women.

Total PCBs represent 70 and 85% of total TEQs for Caucasians and Inuits, respectively. In the fisherman study, levels of coplanar PCBs were 30 times more than those of controls (Table 2). However, coplanar PCBs were determined only on those 10 fishermen with the highest total PCB blood levels. Levels of the sum of the 10 main PCB congeners was 46.4 μg/L of plasma in this highly exposed group compared with 12.3 μg/L in the overall fishing population studied. If we assume that this ratio (1 to 4) also holds true for coplanar PCBs, then we can estimate that plasma levels of non-ortho coplanar PCBs are probably near 400 and 250 ng/kg lipids for coplanars 126 and 169 respectively, 8 to 10 times more than in southern populations. For the highly exposed fishermen, TEQs of 204 ng/kg are due to the two non-ortho coplanar PCBs measured. The main contributors for total TEQs is the mono-ortho coplanar 118 with a TEQ of 568 ng/kg (60% of total PCB TEQs).

**Conclusion**

Inuit people from Arctic Québec and fishermen from the lower north shore of the St. Lawrence River depend on fish consumption for their subsistence. Because of their extremely high daily intake of seafood (300 g for Inuit, 140 g for fishermen) and despite the relatively low concentrations of contaminants in fish from these remote areas (J1), it appears that not only consumers of highly contaminated fish but also remote maritime communities obtain elevated doses.

In these two maritime populations, as in general populations, the main contributors to total TEQs are PCBs, particularly 2,3',4,4',5 pent-CB (no.118), which is responsible for more than 60% of the total toxicity. This finding indicates the necessity for further experimental studies to confirm TEFs for non- and mono-ortho coplanar PCBs. In these two regions, epidemiologic studies are now in progress and the recent knowledge of the relative toxicity of coplanar PCBs, PCDDs, and PCDFs will be useful to better characterize biological exposure. However, it is not clear whether all the toxic effects linked to these PCB exposures are mediated by a TCDD-like mechanism. The principal effects reported for OCs in children exposed during their fetal life affect the central nervous system (J1) and seem not to be mediated by Ah receptors (J2). This example demonstrates the present limitations of using TEQs in exposure assessment.

This study was made possible through the following agencies and their financial assistance: Ministry of Health (Québec), Ministry of Environment (Québec), Health and Welfare Canada and Hydro-Québec. We thank Kathy E. Boggess of Midwest Research Institute for analysis of coplanar PCBs in milk samples and in the firefighters' blood samples, Evelyne Pelletier and Liliane Ferron of Québec Toxicology Center for specific PCB congeners GC analyses offishermen blood samples and Inuit milk samples, and Lise Côté for technical support. For the plasma analyses, we are grateful to L. Panopio for sample cleanup and to W.-F. Sun and M. Boyle for MS determination.

**REFERENCES**

1. Mes, J., Newsome, W. H., and Conacher, H. B. S. Levels of specific polychlorinated biphenyls in fatty foods from five Canadian cities between 1986-1988. Food. Add. Contam. 8: 351-361 (1991).
2. Ryan, J. J., Panopio, L., Weber, D. F., and Conacher, H. B. S. PCDDs/PCDFs in 22 categories of food collected from six Canadian cities between 1985 and 1988. In: Organohalogen Compounds, Vol. 1. Dioxin, 90/EPRI-Seminar. (O. Hutzinger and H. Fiedler, Eds.) Ecoinforma Press, Bayreuth, Germany, 1990 pp. 497-500.
3. Conacher, H. B. S., Graham, A., Newsome, W. H., Graham, G. F., and Verdier, P. The Health Protection Branch total diet program and overview. Can. Inst. Food Sci. Technol. J. 22: 322–326 (1989).
4. NATO/CCMS. Pilot Study on International Information Exchange on Dioxins and Related Compounds. Scientific Basis for the Development of the International Toxicity Equivalency Factor (I-TEF). Method of Risk Assessment for Complex Mixtures of Dioxins and related compounds. North Atlantic Treaty Organization Committee on the Challenges of Modern Society. Report no. 176. Bayreuth, Germany, 1989.
5. Safe, S. Polychlorinated biphenyl, dibenzo-p-dioxins, dibenzofurans and related compounds: environmental and mechanistic considerations which support the development of Toxic Equivalency Factors. CRC Crit. Rev. Toxicol. 21: 51–88 (1990).
6. Norén, K., and Lundén, A. Trend studies of polychlorinated biphenyls, Dibenzo-p-dioxins and Dibenzo furans in human milk. In: Organohalogen Compounds, Vol. 1. Dioxin, 90/EPRI-Seminar. (O. Hutzinger and H. Fiedler, Eds.) Ecoinforma Press, Bayreuth, Germany, 1990, pp. 263–266.
7. Devaillie, E., Weber, J. P., Girgas, S., and Laliberté, C. Coplanar PCBs in human milk in the Province of Québec, Canada. Are they more toxic than dioxin for breast fed infants? Bull. Environ. Contam. Toxicol. 47: 491–498 (1991).
8. Patterson, D. G., Jr., Todd, G. D., Turner, W. E., Isaacs, S. G., and Needham, L. L. Levels of non-ortho-substituted polychlorinated biphenyls, dibenzo-p-dioxins, and dibenzofurans in human serum and adipose tissue. In: Organohalogen Compounds, Vol. 4. Miscellaneous Papers and index. (O. Hutzinger and H. Fiedler, Eds.), Ecoinforma Press, Bayreuth, Germany, 1990, pp. 133–136.

9. Ryan, J. J., Panopio, L., Lewis, D. A., and Weber, D. F. Polychlorinated dibenzo-p-dioxins and polychlorinated dibenzofurans in cows' milk packaged in plastic-coated bleached paperboard containers. J. Agr. Food Chem. 39: 218–223 (1991).

10. Muir, D. C. G., Wagemann, R., Hargrave, B. T., Thomas, D. J., Peakall, D. B., and Norstrom, R. J. Arctic marine ecosystem contamination. Sci. Total Environ. 122: 75–134 (1992).

11. Jacobson, J. L., Jacobson, S. W., and Humphrey, H. E. B. Effects of in utero exposure to polychlorinated biphenyls and related contaminants on cognitive functioning in young children. J. Pediatr. 116: 1:38–45 (1990).

12. Seegal, R. F., Bush, B., and Shain, W. Neurotoxicology of PCBs: a novel structure/activity relationship. In: Organohalogen Compounds, Vol. 1. Dioxin 90/EPRI Seminar. (O. Hutzinger and H. Fiedler, Eds.), Ecoinforma Press, Bayreuth, Germany, 1990, pp. 157–160.

13. Carrier, G., Tremblay, C. G., Groulx, S., and Dewailly, E. Polychlorobiphenyl (PCB), polychlorodibenzodioxin (PCDD), and polychlorodibenzofuran (PCDF) exposure of firefighters involved in the PCB fire at St-Basile-le-Grand, Québec, Canada. In: Organohalogen Compounds, Vol. 3. Dioxin 90/EPRI Seminar. (O. Hutzinger and H. Fiedler, Eds.), Ecoinforma Press, Bayreuth, Germany, 1990, pp. 361–368.