

Original Scientific Paper

## SEASONAL VARIATIONS OF POLYCHLORINATED BIPHENYLS IN AMBIENT AIR IN ZAGREB, CROATIA

Snježana HERCEG ROMANIĆ and Blanka KRAUTHACKER

*Institute for Medical Research and Occupational Health, Zagreb, Croatia*

Received November 2001

Levels of polychlorinated biphenyls (PCBs) were monitored continuously in Zagreb between June 1999 and February 2000 and analysed qualitatively and quantitatively using high resolution gas chromatography (HRGC). Total PCBs were determined with respect to Aroclor 1260, which was used as the standard reference. The concentrations of PCBs ranged from 1.6 to 136  $\mu\text{g}/\text{m}^3$  and were higher in warmer seasons (temperatures above 10 °C; 5.1-136  $\mu\text{g}/\text{m}^3$ ) than in colder seasons (temperatures from -10 °C to about 10 °C; PCB range: 1.6-23.5  $\mu\text{g}/\text{m}^3$ ). The increase in PCBs concentration with temperature followed an exponential curve.

**KEY WORDS:** *levels in air, organochlorine compounds, persistent pollutants, POPs*

Polychlorinated biphenyls (PCBs) make a group of 209 synthetic aromatic compounds differing by the number and the position of chlorine atoms ( $\text{C}_{12}\text{H}_{10-n}\text{Cl}_n$ ;  $n$  is the number of chlorine atoms). They are non-polar lipophilic compounds, stable in the environment in either alkaline or acid conditions. Due to their physical and chemical properties they were used widely in open systems as additives to cement, paints, or copy paper, as well as in closed systems such as transformers, condensers, and vacuum pumps (1).

The atmosphere is an important environment compartment under direct influence of PCBs. Because of their persistence and toxic influence on environment and humans, PCBs have limited application today. They are still found in such sources as waste, certain transformers and condensers, and they also evaporate from natural sorbents (such as sediments and biota). In

ambient air, PCB compounds are present in the gas phase or are adsorbed on particulate matter. Higher chlorinated PCBs (with more than five chlorine atoms) are more lipophilic and therefore tend to adsorb on particles while PCBs with the lower chlorine content are mainly found in the gaseous phase. The evaporation from the sorbent's surface and long-range transboundary air transport of PCBs depend on the temperature and other meteorological conditions (2, 3). As PCB levels in ambient air tend to be higher at higher temperatures, seasonal variations are expected, with peaks in the summer (4).

The aim of this study was to monitor PCB levels in samples of ambient air collected in Zagreb, Croatia and establish their seasonal variations. One of the reasons for this study is that data on PCB levels in ambient air in Croatia are quite scarce (5), and seasonal monitoring had never been undertaken before.

## MATERIAL AND METHOD

Total PCBs in the analysed ambient air samples were determined with respect to the reference Aroclor 1260 mixture, which has the most similar pattern to that of PCB. Samples (about 1,000 m<sup>3</sup> air) were collected on polyurethane foam and quartz filters. From June 1999 to February 2000, 48 samples had been collected in the northern part of Zagreb around the clock. Data about air pressure and temperature were obtained from the Croatian Meteorological and Hydrological Service, and the volumes of sampled air were recalculated to standard temperature and air pressure. Polyurethane foam and quartz filters were extracted in Soxhlet (12 hrs, 4 cycles/h) with 5% diethyl ether in *n*-hexane. Extracts were evaporated to about 5 ml, cleaned with concentrated sulphuric acid and evaporated to dryness in a gentle stream of nitrogen. The samples were then analysed qualitatively and quantitatively using high-resolution gas chromatography with <sup>63</sup>Ni electron capture detector and a 60-m long capillary column with inner diameter of 0.25 mm and SPB-5 stationary phase of 0.25 µm film thickness (5).

## RESULTS AND DISCUSSION

PCBs were present in all 48 analysed samples and their levels ranged from 1.6 to 136 pg/m<sup>3</sup>

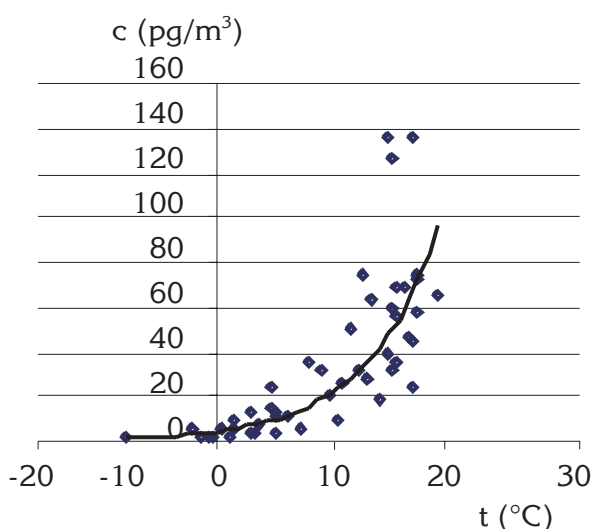


Figure 1 Levels of total PCBs (pg/m<sup>3</sup>) in ambient air samples collected in Zagreb in 1999/2000 in relation to air temperature (°C)

(Figure 1). It is evident from Figure 1 that PCB levels gradually increase with temperature. At temperatures below 0 °C concentrations are low (range: 1.6-2.7 pg/m<sup>3</sup>). Markedly higher levels were determined in samples collected when the average temperatures were above 10 °C. This increase with temperature is not linear, but follows an exponential equation:  $y=4.46 e^{0.126x}$ ; where *y* is PCB concentration and *x* is the temperature. This equation also includes concentrations from three samples which for unknown reason contained PCBs at considerably higher levels than other samples collected at the same temperature (Figure 1). This might be the result of long-range transport of PCBs through the air.

Table 1 Median (*M*) monthly concentrations of total PCBs in ambient air (pg/m<sup>3</sup>). Temperature (*t*) is the mean of average daily air temperatures over the sampling period in the specific month

Year	Month	t (°C)	M (pg/m <sup>3</sup> )
1999	June	19.7	66.0
	July	21.5	46.4
	August	19.8	55.9
	September	19.0	52.5
	October	11.5	23.2
	November	2.7	5.5
	December	1.3	4.3
2000	January	-1.4	2.3
	February	4.0	10.9

Table 1 gives a summary of all results; concentrations and temperatures are expressed as monthly median and mean values, respectively. PCB levels were two or more times higher in the months with average temperatures ranging between 19.0 °C and 21.5 °C than in the months when average temperatures ranged from -1.4 °C to 11.5 °C.

According to *Oehme and co-workers* (6), higher PCB levels in ambient air can be observed when ambient temperatures keep above 10 °C over a longer period, necessary for PCBs to evaporate from the sorbent's surface (6). Our results corroborate this finding. It seems that PCB levels in ambient air do not vary noticeably at low temperatures (7). Seasonal variations of PCB levels in ambient air were also determined in samples collected in Augsburg, Germany (8), as well as in Manchester, Cardiff, London, and Stevenage in the United Kingdom (9). According

to Kaupp and co-workers (8), PCB levels in ambient air vary because of temperature-dependent partition of PCBs between sorbents on the Earth's surface (soil, water, biota) and air (8).

PCBs can be determined as individual congeners or can be compared to commercial mixtures which always contain numerous PCB congeners. Commercial mixtures differ by their chlorine content, that is, by the number and type of congeners. Different abilities and performance of laboratories and the differences in the approach to PCB measurement, measurement in various biological matrices make the comparison of published results difficult. To facilitate their comparison, it is necessary to state whether PCBs were determined as individual congeners or in comparison to a commercial reference mixture, as well as to indicate its content and type. When PCBs are determined as individual congeners, concentrations can be expressed as levels of each congener, as a sum of all congeners analysed, or as a sum of congeners within a group of homologues. To express concentrations as the sum of all congeners means to list all congeners included in their sum.

Granier and co-workers (10) reported that total PCB levels in ambient air samples collected in Paris in 1986 determined with reference to mixtures Aroclor 1242 : Aroclor 1254 : Aroclor 1268 (1:1:1) were relatively high (range: 5.0-19.0 ng/m<sup>3</sup>), while in 1990 those levels were somewhat lower (range: 2.0-6.0 ng/m<sup>3</sup>) (10). PCB levels in samples of ambient air collected on eleven locations in the southern Sweden (expressed as sum of levels of 51 congeners) ranged from 7.0 to 983 pg/m<sup>3</sup> (4). If our results are compared to those cited above, it is evident that total PCB levels determined in samples of ambient air collected in Zagreb keep within the range reported in other European countries.

### Acknowledgement

This work was supported by the Ministry of Science and Technology of the Republic of Croatia. The author's wish to thank Elsa Reiner for her valuable suggestions and help.

### REFERENCES

1. Danse IR, Jaeger RJ, Kava R, Kroger M, London WM, Lu FC, et al. Position paper of the American council on science and health: public health concerns about environmental polychlorinated biphenyls (PCBs). *Ecotoxicol Environ Safety* 1997;38:71-84.
2. Ballschmiter K, Wittlinger R. Interhemispheric exchange of hexachlorocyclohexanes, hexachlorobenzene, polychlorobiphenyls, and 1,1,1-trichloro-2,2-bis(p-chlorophenyl)ethane in the lower troposphere. *Environ Sci Technol* 1991;25:1103-11.
3. Ballschmiter K. Persistent, ecotoxic, and bioaccumulative compounds and their possible environmental effects. *Pure Appl Chem* 1996;68:1771-80.
4. Backe C, Larsson P, Okla L. Polychlorinated biphenyls in the air of southern Sweden - spatial and temporal variation. *Atmos Environ* 2000;34:1481-6.
5. Herceg S. Određivanje organoklorovih spojeva u lišću drveća i zraku. [Determination of organochlorine compounds in tree leaves and the air, in Croatian] [M. Sc. thesis]. Zagreb: Faculty of Science and Mathematics, University of Zagreb; 1999.
6. Oehme M, Haugen JE, Schlabach M. Seasonal changes and relations between levels of organochlorines in Arctic ambient air: First result of an all-year-round monitoring program at Ny-Alesund, Svalbard, Norway. *Environ Sci Technol* 1996;30:2294-304.
7. Stern GA, Halsall CJ, Barrie LA, Muir DCG, Fellin P, Rosenberg B, Rovinsky FYA, Kononov EYA, Pastuhov B. Polychlorinated biphenyls in Arctic air. 1. Temporal and spatial trends: 1992 - 1994. *Environ Sci Technol* 1997;31:3619-28.
8. Kaupp H, Dörr G, Hippelen M, McLahlan MS, Hutzinger O. Baseline contamination assessment for a new resource recovery in Germany. Part IV: Atmospheric concentrations of polychlorinated biphenyls and hexachlorobenzene. *Chemosphere* 1996;32:2029-42.
9. Halsall CJ, Lee RMG, Coleman PJ, Burnett V, Harding-Jones P, Jones KC. PCBs in U.K. air. *Environ Sci Technol* 1995;29:2368-76.
10. Granier LK, Chevreuil M. Behaviour and spatial and temporal variations of polychlorinated biphenyls and lindane in the urban atmosphere of the Paris area, France. *Atmos Environ* 1997;31:3787-802.

**Sažetak****SEZONSKE VARIJACIJE POLIKLORIRANIH BIFENILA U ZRAKU GRADA ZAGREBA**

Kontinuirano su praćene razine polikloriranih bifenila (PCB) u zraku skupljenom u sjevernome dijelu Zagreba tijekom devetomjesečnog razdoblja (lipanj 1999. - veljača 2000.). Kvalitativna i kvantitativna analiza provedena je plinskom kromatografijom visokog razlučivanja. Određivani su ukupni PCB-i prema smjesi Aroclor 1260. Koncentracije PCB-a bile su u rasponu 1,6 - 136 pg/m<sup>3</sup>. U razdoblju s višim temperaturama (iznad 10 °C) izmjerene su više koncentracije PCB-a (raspon: 25,1 pg/m<sup>3</sup> - 136 pg/m<sup>3</sup>) nego u razdoblju s nižim temperaturama kada je raspon temperatura bio od -10 °C do +10 °C (raspon: 1,6 pg/m<sup>3</sup> - 23,5 pg/m<sup>3</sup>). Porast razina ukupnih PCB-a s temperaturom zraka nije linearan, nego slijedi eksponencijalnu krivulju. Razine ukupnih PCB-a u uzorcima zraka skupljenim u Zagrebu unutar su raspona razina izmjerenih u uzorcima zraka iz drugih europskih zemalja.

**KLJUČNE RIJEČI:** *organoklorovi spojevi, perzistentna zagađivala, POP, razine u zraku*

**REQUESTS FOR REPRINTS:**

Snježana Herceg Romanić, M. Sc.  
Institute for Medical Research and Occupational Health  
P. O. Box 291, HR-10001 Zagreb, Croatia  
E-mail: [Snjezana.Herceg@imi.hr](mailto:Snjezana.Herceg@imi.hr)