

PERSISTENT ORGANIC POLLUTANTS IN LITHUANIA: ASSESSMENT OF AIR AND SOIL CONTAMINATION

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Persistent organic pollutants (POPs) such as polycyclic aromatic hydrocarbons (PAHs), polychlorobiphenyls (PCBs), and pesticides (hexachlorocyclohexanes (HCHs), dichlorodiphenyltrichlorometilmetanes (DDTs)) were investigated in the atmospheric air and soil at 5 sites of Lithuania during March–August, 2006. POPs concentration at different sampling sites varied in the range of 6.39–127.8 ng m⁻³, 0.017–0.440 ng m⁻³, 0.088–0.310 ng m⁻³, and 0.006–0.360 ng m⁻³ in the atmospheric air and in the range of 29.5–529.3 ng g⁻¹, 0.6–24.0 ng g⁻¹, 0.4–1.1 ng g⁻¹, and 0.3–7.7 ng g⁻¹ in soil for PAHs, PCBs, HCHs, and DDTs, respectively. The direct relationship between the POPs concentration level in the atmospheric air and soil at all sampling sites was observed. The concentrations of PAHs dominated over those of other POP groups in the atmospheric air and soil as well. The highest concentrations of almost all POP compounds were determined in the atmospheric air and soil in Vilnius city at the site with intensive traffic and commercial activity. Such wide range investigations of POPs in the natural environment components have been performed for the first time in Lithuania.

Keywords: PAHs, PCBs, HCHs, DDTs, concentration, passive samplers, atmospheric air, soil

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1. Introduction

Persistent organic pollutants (POPs) due to their wide distribution, ability to bioaccumulate in the fatty tissue, and carcinogen, mutagen, or endocrine disruption potential remain in the centre of researchers' attention [1]. Among different organic matters, POPs (polycyclic aromatic hydrocarbons (PAHs), polychlorinated biphenyls (PCBs), pesticides, polychlorinated dibenzop-dioxins, and dibenzofurans) are the most interesting from the point of view of their behaviour in the environment. Different classes of POPs are emitted from various sources: PAHs are mainly emitted in combustion processes from industrial or domestic sites and traffic [2], PCBs are spread via chemicals used in semiconductor industry and by waste disposal [3, 4], dibenzop-dioxins and dibenzofurans via operation of landfills and incinerations [5, 6], and pesticides are spread via dispersion to the soil [7, 8]. Furthermore, secondary sources of POPs include the spreading of sludge on the ground surface and remobilization of previously deposited compounds from soil and water bodies. The atmosphere often plays a key role in their transport within

the immediate vicinity of POP sources as well as over great distances [9, 10]. Atmospheric transport is also the main route for carrying POPs to the terrestrial and aquatic ecosystems [11, 12].

Research on POPs in the environment currently focuses on determination of uncertainties due to their ambient sources, atmospheric transport and fate, and air-surface exchange to clarify the complex factors controlling air concentrations. Measurements of the atmospheric air concentration at different sites are needed to assess contribution of sources, atmospheric processes, and the long-range atmospheric transport of potential toxic POPs.

High-volume samplers are usually used as conventional sampling techniques to determine POPs in the atmospheric air [6, 7, 13–15]. The relatively high cost of the equipment as well as the requirement for a pump and source of electricity are important disadvantages of this sampling method. A potential alternative is the use of passive samplers. These devices can be deployed at many sites at the same time, which offers a new approach to the large-scale investigation. As it provides information about long-term contamination of

the selected site, passive air sampling can be used as a screening method for semi-quantitative comparison of different sites having advantage of low sensitivity to accidental short-time changes in the concentration of pollutants. However, it has been demonstrated that passive air samplers are mainly available for study of vapour-phase POPs, they are applied to POP monitoring on the global and regional scale [16–18].

The aim of this study was to determine some polycyclic aromatic hydrocarbons, polychlorinated biphenyls, and pesticides by exposing passive samplers in the territory of Lithuania according to the program of the pilot study for development of the monitoring network in the Central and Eastern Europe (MONET CEEC), 2006 under the auspices of Research Centre for Environmental Chemistry and Ecotoxicology (RECETOX), Masaryk University, Brno, Czech Republic. Furthermore, this study was the first attempt to investigate the distribution of POPs such as PCBs and organic chlorinated pesticides in the environment of Lithuania.

2. Experiment write-up

Sampling sites. Monthly sampling of POPs in the atmospheric air was performed at 5 chosen sites in the territory of Lithuania in the period of 21 March to 13 August 2006: two sampling sites were chosen in the western part (Preila, Plateliai) and three sampling sites in the eastern part of Lithuania (Vilnius, Aukštieji Paneriai, Rūgštelėškis) (Fig. 1). The selected sites comprised commercial and industrial part of the Vilnius city and its suburb Aukštieji Paneriai, while other sites represented the monitoring background stations. Detailed characteristics of the sampling sites are presented below:

- Preila (55°20'N; 21°00'E) is a background environmental pollution research station of the Institute of Physics (Lithuania) on the southeastern coast of the Baltic Sea. It is located on the Curonian Spit (70 km in length and 2–3 km in width) separating the Curonian Lagoon from the Baltic Sea. The main local sources of pollution are small villages situated along the spit and the traffic on the road Klaipėda–Kalininograd at a distance of 300 m.
- Plateliai (56°00'62"N; 21°52'28"E) is an integrated monitoring station located in Žemaitija national park, 50 km to the east from the Baltic Sea.
- Rūgštelėškis (55°26'26"N; 26°03'60"E) is an integrated monitoring station located in the Aukštaitija



Fig. 1. Geographical location of sampling sites.

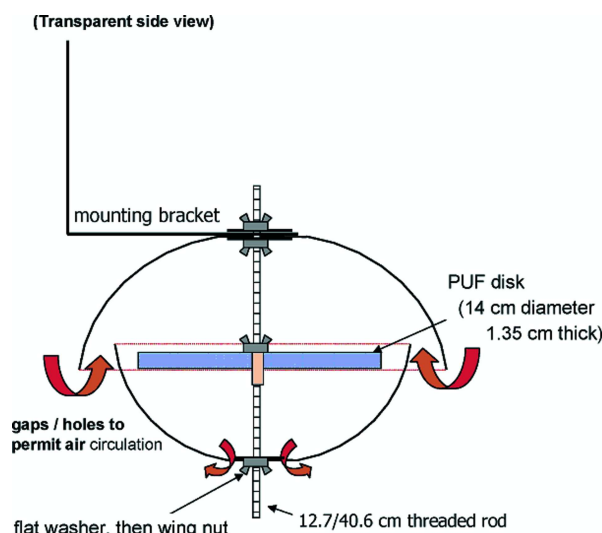


Fig. 2. Scheme of the passive air sampling device.

National Park where measurements of POPs according to EMEP project are performed.

- Aukštieji Paneriai sampling site (54°39'30"N; 25°14'16"E) is in the forested area, in the territory of the Institute of Physics in the southwestern part of Vilnius city. Experimental waste and local traffic can be a source of pollution.
- Vilnius sampling site (54°42'37"N; 25°20'40"E) is located in the commercial and industrial part of Vilnius city near the street with intensive traffic (Žemaitės St), among residential buildings.

Concentrations of POPs in the soil at air sampling sites were determined as well. The plot of about 10–15 m² area without vegetation cover in each location

was selected. From 5 to 10 subsamples (ca 0.3–0.5 kg) were taken at different random points from the soil surface layer (0–10 cm) of this area. Five combined samples of soil were collected at all air sampling sites at the end of air sampling campaign (August, 2006).

Sampling. For sampling of POPs in the atmospheric air, passive air samplers were used. They are composed of a foam disk (15 cm diameter, 1.5 cm thick, 0.03 g m^{-3} density, type N 3038, Gumotex Breclav, Czech Republic) placed in a stainless steel sampling chamber consisting of two domes (Fig. 2) [19]. This “flying saucer” design protects the foam disks from direct precipitation, sunlight, and coarse particle deposition.

Sampling chambers were prewashed and solvent-rinsed with acetone prior to installation. All filters were prewashed, cleaned (8-hour-extraction in acetone and 8-hour-extraction in dichloromethane), wrapped in two layers of aluminum foil, placed into zip-lock polyethylene bags, and kept in the freezer prior to deployment. Exposed filters were wrapped in two layers of aluminum foil, labelled, placed into zip-lock polyethylene bags, and transported in a cooler at 5°C to the laboratory where they were kept in the freezer at -18°C until the analysis. Air was allowed to flow over the sampling surface through a ~ 2.5 cm gap between two domes. The average sampling rate was estimated to be $3.5 \text{ m}^3/\text{day}$, which roughly corresponds to 100 m^3 for a 28-day sampling cycle. Exposure time of four to twelve weeks enables determination of many compounds from the POP group [20]. The technique of passive sampling and uptake of POPs by polyurethane foam is described in [16, 21] and it was shown that passive samplers with polyurethane foam can be used mainly for gas phase POPs and may collect only some part of fine particles [18].

Analysis. All samples were extracted with dichloromethane in a Büchi system B-811 automatic extractor. Surrogate recovery standards (d8-naphthalene, d10-phenanthrene, d12-perylene for PAH analysis, PCB 30 and PCB 185 for PCB analysis) were spiked on each filter prior to extraction. Terphenyl was used as internal standard for polyaromatic hydrocarbons (PAHs) and PCB 121 was used for polychlorinated biphenyl (PCB)/organochlorine pesticide (OCP) analyses. Volume of extracts after extraction was reduced under a gentle nitrogen stream at ambient temperature. Fractionation of PCB/OCP in extracts was performed on a silica gel column (a sulphuric acid modified silica gel). Samples were analysed using GC-ECD (HP 5890) supplied with

a Quadrex fused silica column for PCBs such as PCB28, PCB52, PCB101, PCB118, PCB153, PCB138, PCB180, and OCPs such as α -hexachlorocyclohexane (HCH), β -HCH, γ -HCH, δ -HCH, 1,1-dichloro-2,2-bis(*p*-chlorophenyl)ethylene (*p,p'*-DDE), 1,1-dichloro-2,2-bis (*p*-chloroprene) ethan (*p,p'*-DDD), 1,1,1-trichloro-2,2-bis (*p*-chloroprene) ethan (*p,p'*-DDT), *o,p'*-DDE, *o,p'*-DDD, hexachlorobenzene (HCB), and pentachlorobenzene (PeCB). Furthermore, 16 polycyclic aromatic hydrocarbons recommended by EPA (Environmental protection agency, USA) were determined in all samples using the GC-MS instrument (HP 6890 – HP 5972) supplied with a J&W scientific fused silica column DB-5MS. The laboratory blank test and the reference material were analysed with each set of ten samples. Limit of quantification was $0.2 \text{ ng filter}^{-1}$ ($=2 \text{ pg m}^{-3}$) [22].

Meteorological data such as ambient temperature, precipitation, humidity, and wind speed direction for all sampling sites during each period of investigation have been taken from the NOAA (National Oceanic and Atmospheric Administration) meteorological network [23].

3. Results and discussion

The ranges and averages of monthly concentrations of various POPs in the atmospheric air at five monitoring sites of Lithuania are presented in Table 1 and distribution of sum of concentrations of each group of determined pollutants at different sites is presented in Fig. 3. Data indicate that concentrations of various POPs are different in the territory of Lithuania with wide variation of some pollutants. The concentrations of PAHs profiles showed that 2–3-ringed (from naphthalene to anthracene) and 4-ringed (fluoranthene and pyrene) of PAH species were higher than those of 5- and 6-ringed PAHs, which, according to earlier studies, were mainly associated with particles in the atmospheric air [14]. A decreasing trend from March to August was observed for each of 16 PAHs at all sampling sites clearly indicating seasonality of pollution and suggesting generation of PAHs by fuel burning for domestic heating in the period of March–April. Furthermore, the concentrations of lighter PAHs determined in this study were very similar to those determined at the Preila background station in 1991 by using low volume air filtration, while the concentrations of particulate PAHs determined by the method of passive samplers were relatively low compared with those determined earlier at the background sites [24] and Vilnius city

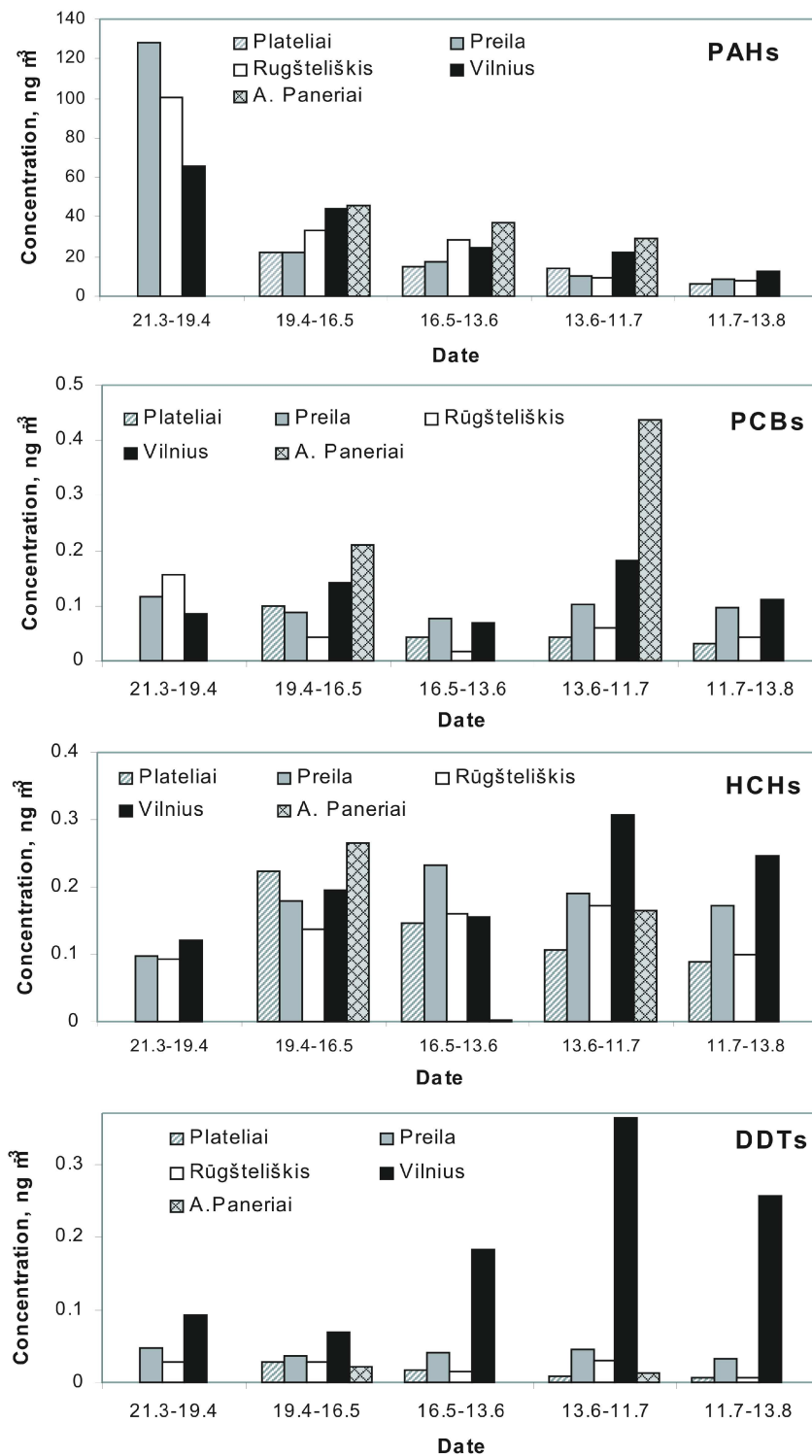


Fig. 3. Changes in sum of concentrations of each POP group in the atmospheric air at different sampling sites.

Table 1. Ranges and averages of POP concentrations in the atmospheric air (ng m^{-3}) at the sampling sites (5 sampling periods for each site, STD means standard deviation, DL is detection limit).

POPs	Minimum	Maximum	Average	STD	Variance, %
PAHs					
Naphthalene	1.71	48.7	13.2	11.6	88.0
Acenaphthylene	0.04	3.26	0.78	0.95	120.8
Acenaphthene	0.11	2.42	0.58	0.61	104.8
Fluorene	0.32	21.3	3.46	5.15	148.9
Phenanthrene	1.07	43.4	9.73	10.6	108.9
Anthracene	0.02	4.86	0.43	1.02	235.7
Fluoranthene	0.41	12.6	2.27	2.62	115.4
Pyrene	0.22	4.16	1.28	0.93	72.8
Benz(a)anthracene	0.00	0.09	0.03	0.02	75.9
Chrysene	0.00	0.48	0.08	0.10	122.8
Benzo(b)fluoranthene	0.01	0.09	0.04	0.02	45.6
Benzo(k)fluoranthene	0.01	0.05	0.02	0.01	47.9
Benzo(a)pyrene	0.01	0.05	0.03	0.01	40.5
Indeno(1,2,3,c,d)pyrene	<DL	0.05	0.03	0.01	50.0
Dibenz(a,h)anthracene	<DL	<DL	<DL	<DL	–
Benzo(g,h,i)perylene	0.00	0.10	0.06	0.02	34.2
Sum of PAHs	6.4	127.8	32.1	30.6	95.3
PCBs					
PCB 28	0.008	0.265	0.039	0.054	138.7
PCB 52	0.006	0.087	0.027	0.019	70.6
PCB 101	<DL	0.036	0.016	0.009	66.6
PCB 118	<DL	0.024	0.013	0.006	48.0
PCB 153	<DL	0.022	0.010	0.006	62.0
PCB 138	<DL	0.013	0.006	0.003	42.3
PCB 180	<DL	0.029	0.006	0.007	101.1
Sum of PCBs	0.017	0.440	0.100	0.090	81.7
HCHs					
α -HCH	0.011	0.203	0.077	0.044	57.3
β -HCH	<DL	0.042	0.022	0.011	49.6
γ -HCH	0.027	0.147	0.071	0.029	41.3
δ -HCH	<DL	<DL	<DL	<DL	–
Sum of HCHs	0.088	0.310	0.166	0.060	36.2
DDTs					
<i>o,p'</i> -DDE	<DL	0.013	0.012	0.002	13.1
<i>p,p'</i> -DDE	0.006	0.151	0.034	0.036	103.3
<i>o,p'</i> -DDD	<DL	0.027	0.013	0.012	91.7
<i>p,p'</i> -DDD	0.000	0.039	0.010	0.012	118.8
<i>o,p'</i> -DDT	<DL	0.065	0.022	0.024	109.4
<i>p,p'</i> -DDT	<DL	0.083	0.023	0.027	117.8
Sum of DDTs	0.006	0.360	0.063	0.090	144.3

or its suburb as well [25]. It may be explained by the possibility of passive samplers to adsorb more vapour than particulate phase POPs. However, the concentrations determined for particulate PAHs were in the interval of monthly concentrations earlier determined at these sites. Long-term investigation of benzo(a)pyrene showed that its monthly concentration in the atmospheric air varied from 0.02 to 1.72 ng m^{-3} with high frequency of daily concentrations of 0.02–0.50 ng m^{-3} at the Preila background station in warm season (May–

September) and from 0.18 to 3.30 ng m^{-3} with high frequency of 0.20–1.00 ng m^{-3} daily concentration in cold season (October–April) during 1980–2002 [26].

The highest concentrations of PCBs were obtained at the background sites in springtime (March–April period), while at urban sites, Vilnius city and its suburb Aukštieji Paneriai, they were higher in summertime. The analysis of synoptic situation shows that in the period from 25 March to 19 April the territory of Lithuania was under the influence of polluted SW air masses

Table 2. Ranges and averages of POP concentrations in soil (ng g^{-1}) at all sampling sites (STD means standard deviation, DL is detection limit).

POPs	Minimum	Maximum	Average	STD	Variance, %
PAHs					
Naphthalene	1.6	10.4	5.2	3.2	61.9
Acenaphthylene	0.3	3.8	1.2	1.5	123.2
Acenaphthene	0.2	1.8	0.6	0.7	108.4
Fluorene	0.3	3.0	1.2	1.0	83.5
Phenanthrene	3.5	49.5	16.0	22.3	138.8
Anthracene	0.2	9.0	2.4	3.7	156.5
Fluoranthene	3.5	70.0	20.6	27.8	135.3
Pyrene	2.4	61.3	17.9	24.5	136.8
Benz(a)anthracene	0.9	38.4	10.7	15.7	146.1
Chrysene	1.9	40.3	11.9	16.0	134.4
Benzo(b)fluoranthene	2.4	74.2	21.1	29.9	142.0
Benzo(k)fluoranthene	1.3	22.3	7.1	8.6	121.1
Benzo(a)pyrene	2.1	49.8	14.7	19.9	134.9
Indeno(1,2,3,c,d)pyrene	2.9	44.8	14.8	17.0	115.2
Dibenz(a,h)anthracene	0.2	5.2	1.5	2.1	136.4
Benzo(g,h,i)perylene	2.4	45.5	13.9	17.8	127.9
Sum of PAHs	29.5	529.3	158.5	209.1	131.9
PCBs					
PCB 28	<DL	0.20	0.15	0.07	47.1
PCB 52	0.10	0.50	0.20	0.17	86.6
PCB 101	0.10	2.60	0.68	1.08	158.2
PCB 118	0.10	1.50	0.46	0.61	131.9
PCB 153	0.20	9.40	2.20	4.03	183.1
PCB 138	0.10	4.30	1.04	1.82	175.4
PCB 180	0.10	5.50	1.60	2.61	162.8
Sum of PCBs	0.60	24.0	5.94	10.1	170.3
HCHs					
α -HCH	0.10	0.50	0.32	0.16	51.4
β -HCH	0.10	0.30	0.24	0.09	37.3
γ -HCH	0.10	0.40	0.27	0.15	63.2
δ -HCH	<DL	<DL	<DL	<DL	–
Sum of HCHs	0.40	1.10	0.80	0.29	36.4
DDTs					
<i>o,p'</i> -DDE	<DL	<DL	<DL	<DL	–
<i>p,p'</i> -DDE	0.20	2.40	0.76	0.92	121.5
<i>o,p'</i> -DDD	<DL	0.80	0.45	0.49	110.0
<i>p,p'</i> -DDD	0.10	3.60	0.80	1.57	195.7
<i>o,p'</i> -DDT	<DL	<DL	<DL	<DL	–
<i>p,p'</i> -DDT	<DL	<DL	<DL	<DL	–
Sum of DDTs	0.30	7.70	1.92	3.23	168.5

from industrial regions of Europe [27], while at the urban sites the local sources were dominating and concentrations of PCBs might increase with an increase of ambient temperature. The increase of PCB concentrations with an increase of ambient temperature in the majority of PCB studies was observed earlier [9, 28]. Furthermore, the level of concentration increased going from PCBs with higher degree of chlorination (PCB 180) to PCB congeners with lower degree of chlorination (PCB 28) which can be related to increasing volatility of these pollutants. The lowest concentrations

of PCBs at urban sites were determined in cold period of March–April, while the highest concentrations of PCBs with large amount of PCB 28 were measured at highest ambient temperature (average temperature varied from 19.2 to 20.6 °C at different sampling sites) at all sampling sites during 13 June – 11 July. The sum of concentrations of PCB 28, 52, 101, 118, 153, 138, and 180 congeners varied between 0.017 ng m^{-3} in Rūgštelėškis and 0.155 ng m^{-3} in Preila with the highest value of 0.440 ng m^{-3} in Vilnius suburb. Concentrations of PCBs at background Lithuanian sites were

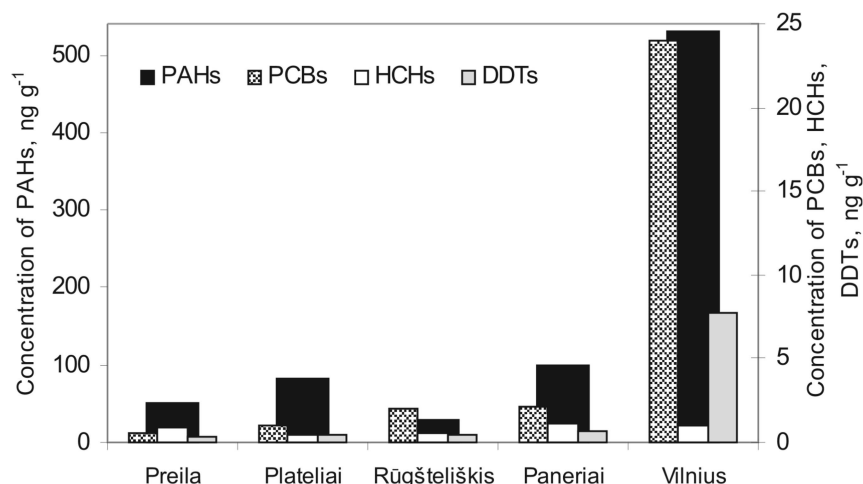


Fig. 4. Variation of concentrations of various POPs in soil at different sites.

similar to those determined in UK, Wraymires rural site (from 12 to 230 pg m^{-3}) in 1972–1992 [29].

The course of concentrations of different HCH isomers was similar at all sampling sites with higher concentrations from mid-April to August. Concentration of γ -HCH (the main component of lindane) was very similar to the α -HCH pattern, but sometimes it was lower because γ -HCH may be transformed to α -HCH due to photochemical reactions in the atmosphere [7]. In our study the sum of HCH concentrations varied from 0.088 ng m^{-3} in Plateliai to 0.310 ng m^{-3} in Vilnius. The highest sum of HCHs concentrations was observed from May to June with a decrease in August at all sampling sites. The level of α -HCH concentration was relatively uniform at the background sites, while its level in Vilnius city was different.

Relatively high concentrations of DDTs, from 0.122 to 0.307 ng m^{-3} , were determined in Vilnius city, but they ranged from 0.006 to 0.048 ng m^{-3} at the rural sites. The highest concentrations of pesticides (HCHs and DDTs) were observed in the period of their usage (April–June) and in the period of high atmospheric temperature during 13 June – 11 July. The lowest pollution by pesticides was determined at Plateliai and Rūgštelėškis integrated monitoring stations. Wide distribution of pesticides may be explained by their intensive application on local and global scales in the past and dispersion of these compounds over long distances through the atmosphere due to their relatively high volatility. As a result of their persistence, organochlorine residues were found in the air and precipitation after some decades [8, 9].

Distribution of different POPs in the soil depends on their deposition processes which are highly controlled by the physico-chemical properties of com-

pounds (physical state, solubility, etc.) in the atmospheric air and meteorological conditions (ambient temperature, precipitation, etc.) [30]. Deposition of some heavier PCBs and high molecular weight PAHs is primarily caused by their insolubility. It was found that dry deposition of PAHs was larger than that of PCBs [14]. Particle deposition seems to be the major pathway of PAHs from the atmosphere to the ground surface, while the largest deposition of PCBs and HCHs, due to their high water solubility, occurs by washout with precipitation. Furthermore, higher solubility of γ -HCH than of α -HCH may affect the ratio α -HCH/ γ -HCH in soil and air samples as well.

In our study, 5 combined samples of soil were analysed for the same POPs as in the atmospheric air of the sampling sites. Concentrations of 16 PAHs ranged from 29.5 to 99.1 ng g^{-1} at rural sites and with the highest value of 529.3 ng g^{-1} in the soil of Vilnius city (Table 2 and Fig. 4). Contrary to the atmospheric air, phenanthrene, fluoranthene, and pyrene together with other high molecular weight particulate PAHs (5–6 rings) were dominating in soil. The dominance of particulate PAHs in the soil indicates that exchange of gaseous PAHs between the atmosphere and soil seems to be insignificant. High concentrations of PAHs in comparison with other POPs show that hydrophobic POPs compounds mainly tend to accumulate in the surface layer of soil.

The sum of PCB concentrations in the soil varied from 0.6 to 2.1 ng g^{-1} at background sites with the highest value of 24.0 ng g^{-1} at the site in the centre of Vilnius city. The concentrations of PCBs determined during this investigation were several times lower than those determined in England, ranging between 10 and 670 $\mu\text{g kg}^{-1}$ (median 30 $\mu\text{g kg}^{-1}$) [11].

The concentrations of HCHs and DDTs were found to be low and uniform in the soil of all rural areas, while concentrations of DDTs were exclusively elevated in the soil of Vilnius investigation site. The concentration of pesticides in Vilnius city was determined to be 1.0 ng g^{-1} and 7.7 ng g^{-1} for HCHs and DDTs, respectively. It is evident that the concentrations of all investigated POPs in the soil were by an order of magnitude higher in the urban soil. Furthermore, a relatively low variation of data showed that more uniform concentrations of all POPs were determined for HCHs at all investigated sites, which indicated stability of pollution sources compared with other POPs. The data presented in Figs. 3 and 4 showed a direct relationship of POP distribution between soil and the atmospheric air at all sampling sites. Consequently, the atmospheric air and soil were the cleanest at rural sampling sites (Plateliai, Rūgšteliškis, and Preila), except the period of March–April when the influence of heating season on the increase of PAH concentration level in the atmospheric air at the Preila background station was evident.

4. Conclusions

The sum of PAH, PCB, HCH, and DDT concentrations at 5 sampling sites in Lithuania varied in the range of $6.4\text{--}127.8 \text{ ng m}^{-3}$, $0.017\text{--}0.440 \text{ ng m}^{-3}$, $0.088\text{--}0.310 \text{ ng m}^{-3}$, and $0.006\text{--}0.360 \text{ ng m}^{-3}$ in the atmospheric air and in the range of $29.5\text{--}529.3 \text{ ng g}^{-1}$, $0.6\text{--}24.0 \text{ ng g}^{-1}$, $0.4\text{--}1.1 \text{ ng g}^{-1}$, and $0.3\text{--}7.7 \text{ ng g}^{-1}$ in the soil, respectively. A direct relationship between POP level in the atmospheric air and soil at all sampling sites was determined. PAH concentrations dominated over the other POP groups in the atmospheric air and soil as well. The highest concentrations almost of all POP compounds were determined in the atmospheric air and soil in Vilnius city at the site with intensive traffic and commercial activity. The concentrations of PCBs, especially of their lower congeners (PCB 28), in the atmospheric air were higher in summertime during the period of higher ambient temperature. Contrary to other POP groups, concentrations of PAHs indicated a decreasing trend from spring months to the end of summertime. Comparison of the concentrations of benzo(a)pyrene determined by using passive samplers and low volume air filtration showed that concentrations of this particulate PAH were lower but they were in the interval of monthly variation determined earlier at the Preila background station by using the method of low volume air filtration.

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PATVARŪS ORGANINIAI TERŠALAI LIETUVOJE: TARŠOS ORE IR DIRVOJE ĮVERTINIMASA. Milukaitė^a, J. Klánová^b, I. Holoubek^b, I. Rimšelytė^a, K. Kvietkus^a^a Fizikos institutas, Vilnius, Lietuva^b Masaryko Universitetas, Brno, Čekija**Santrauka**

Patvarių organinių teršalų (POT), tame tarpe policiklinių aromatinių angliavandenilių (PAA), polichlorbifenilų (PCB) ir pesticidų (heksachlorcikloheksanų (HCH), dichlordifeniltrichlormetilmetanų (DDT)), tyrimai atlikti 2006 m. kovo–rugpjūčio mėnesiais atmosferos ore ir dirvoje penkiose Lietuvos vietovėse. POT atmosferos ore tirti panaudojant pasyvius sorbentus, kurie pastaruoju metu yra plačiai taikomi atmosferos užterštumo tyrimams regioniniu ir globaliu mastu. Tyrimų rezultatai parodė, kad 16-kos PAA junginių, 7-ių PCB, 4-ių HCH ir 6-ių DDT suminės koncentracijos ore Lietuvos teritorijoje kito $6,4\text{--}127,8\text{ ng m}^{-3}$, $0,017\text{--}0,440\text{ ng m}^{-3}$, $0,088\text{--}0,310\text{ ng m}^{-3}$, $0,006\text{--}0,360\text{ ng m}^{-3}$ ribose, o dirvoje $29,5\text{--}529,3\text{ ng g}^{-1}$, $0,6\text{--}24,0\text{ ng g}^{-1}$, $0,4\text{--}1,1\text{ ng g}^{-1}$, $0,3\text{--}7,7\text{ ng g}^{-1}$ ribose. Nustatyta tiesioginė priklausomybė tarp koncentracijų POT atmosferos ore ir dirvoje. Tarp tirtų POT grupių

policiklinių aromatinių angliavandenilių koncentracijos buvo didžiausios ir atmosferos ore, ir dirvoje. Beveik visų POT didžiausios koncentracijos buvo Vilniaus mieste, o mažiausios – Rūgšteliškio stebėsenos stotyje. Polichlorbifenilų, ypač PCB28, koncentracijos buvo didesnės vasarą, esant aukštesnei oro temperatūrai. Skirtingai nuo kitų POT, policiklinių aromatinių angliavandenilių koncentracija ore mažėjo nuo šildymo sezono iki vasaros pradžios. Vidutinės benz(a)pireno koncentracijos atmosferos ore vertės, nustatytos naudojant pasyvius sorbentus, buvo mažesnės už jų vidutinės vertes, nustatytas imant oro bandinius mažu filtracijos greičiu. Tai rodo, kad pasyvių sorbentų metodas labiau tinka nustatant dujines negu aerosolines POT koncentracijas. Tokie plataus spektro patvarių organinių teršalų tyrimai gamtinės aplinkos sanduose Lietuvoje atlikti pirmą kartą.