

Seasonal and Regional Contributors of 1-Hydroxypyrene among Children near a Steel Mill

Mi-Sun Lee,¹ Ki-Do Eum,¹ Kiyoung Lee,¹ Ho Kim,² and Domyung Paek¹

Departments of ¹Environmental Health and ²Biostatistics, Graduate School of Public Health and Institute of Health and Environment, Seoul National University, Seoul, Republic of Korea

Abstract

Urinary 1-hydroxypyrene (1-OHP) is a biomarker of exposure to polycyclic aromatic hydrocarbons (PAH). Effect of residence on children's PAH exposure was reported among children living near a polluted area. Instead of a snapshot assessment, however, a temporal history of exposure characteristics needs to be assessed in the studies of chronic disease development such as cancer. The urinary 1-OHP measurements were repeated to determine regional effect of ambient air pollution on 1-OHP levels over extended periods. Two sites were chosen: (a) one site located near the steel mill ("nearby" site) and (b) the other site located at a longer distance from the mill ("remote" site). Spot urinary 1-OHP levels were measured from 72 children for 3 consecutive days per month, repeated over 9-month period. Compared with remote site, the nearby site had

increased the urinary 1-OHP level by 62.3% [95% confidence interval (95% CI), 39.8-88.3%]. Other statistically significant factors that contributed to the level include sex [16.5% (95% CI, 1.2-34.1%) higher for girls than boys], consumption of charbroiled meat [16.5% (95% CI, 1.1-34.2%) higher], and an increase in PM₁₀ [10.1% (95% CI, 4.8-15.7%) higher for the interquartile range increment]. Controlling for covariates, the 1-OHP levels were increased in the summer and fall compared with winter. The magnitude of the effects of both seasons had diminished after adjusting for PM₁₀. This is the first report providing seasonal and regional contributors to environmental PAH exposure, assessed by urinary 1-OHP, with higher 1-OHP levels during summer when ambient pollution was also high. (Cancer Epidemiol Biomarkers Prev 2009;18(1):96-101)

Introduction

Polycyclic aromatic hydrocarbons (PAH) have raised much public concern because some congeners are listed as group 1 (i.e., "carcinogenic to humans") by IARC (1). Epidemiologic studies have shown increased lung cancer risks for employees exposed to PAHs (2-6) as well as for residents of nearby neighborhoods (7, 8). Various adverse health effects also have been reported among children residing near a known PAH source: decline of lung function (9), growth-retarded infant (10), and delay in child's neurodevelopment from prenatal exposure (11).

PAHs are released as a mixture into the environment in a variety of ways, and the exposure properties of these compounds vary depending on the source. Routes of human exposure to PAHs occur via inhalation, ingestion, and dermal contact because these pollutants are ubiquitous in living environments. Sources of PAHs are mainly cigarette smoke, wood-burning ovens and fireplaces, vehicle exhaust, and consumption of grilled and smoked foods (1). Occupational exposure generally occurs in the setting of coal-fired utilities, steel plants, and waste incineration facilities.

To provide information on the uptake of PAHs from various sources, pathways of exposure, and health risks, urinary 1-hydroxypyrene (1-OHP), as a metabolite of pyrene, has been widely used as the most relevant biomarker of PAH exposure in occupational and nonoccupational settings (12-14). Despite of its wide usage in biomonitoring studies for public health, essential questions about temporal variability should be addressed before applying urinary 1-OHP to the study on PAH-related chronic diseases such as cancer. Urinary 1-OHP levels reflect relatively recent exposures, especially within a few days before urine collection due to its short half-life (15). Therefore, information on the variation of long-term PAH exposure and its contributing factors are necessary for epidemiologic studies of chronic diseases and ensuing health policy. However, little information on the seasonal variation of PAH exposure and its contributing factors are available.

This study was designed to provide information on the seasonal and regional variation of PAH exposure among children living at two separate sites from a steel mill assessed by repeated urine samples over 9 months. The design can provide the insight for further understanding of the role of contributing factors on the long-term variation of biological PAH exposures.

Materials and Methods

Study Site and Subject. A total of 956 school children participated in biomonitoring part of the PAH exposure assessment study (16). Among the participants, 72 children agreed to take part in the repeated PAH

Received 7/7/08; revised 9/30/08; accepted 10/6/08.

Grant support: Local municipal government and Ministry of Environment, Republic of Korea.

Requests for reprints: Domyung Paek, Department of Environmental Health, Graduate School of Public Health and Institute of Health and Environment, Seoul National University, 28 Yeongseon-Dong, Jongno-Gu, Seoul 110-460, Republic of Korea. Phone: 82-2-740-8886; Fax: 82-2-743-8240. E-mail: paekdm@snu.ac.kr

Copyright © 2009 American Association for Cancer Research.

doi:10.1158/1055-9965.EPI-08-0612

biomarker assessment. This repeated prospective sampling was done from March to December of 2004, except for the school vacation period of January, February, and August. As the urine samples were collected during the attendance of school, study was unfeasible during the vacation. Usually, winter vacation is longer than the one in summer in Korea. Each month, a spot urine sample was collected every morning for 3 consecutive days, and this once-a-month, 3-d sampling was repeated for 9 mo. Each subject, then, could have up to 27 spot urine samples during the study period.

The study site was in the neighborhood of a steel mill located in the southern part of Korea. Two sites were chosen based on wind direction and distance from the steel plant: "nearby" site adjacent to the northern boundary of the plant complex and "remote" site, as a reference area, located on the same wind direction but about 6 km away from the plant. Participants of nearby site were consisted of 36 students from one elementary school ($n = 24$) and one junior high school ($n = 12$), whereas those of remote site were consisted of 36 students from another elementary school ($n = 24$) and another junior high school ($n = 12$). We hold meetings at each school to explain about the study objectives, sampling methods, and questionnaire guidelines. We distributed on the eve of the sampling day the guideline sheets, individually labeled questionnaires, and labeled sterile sampling tubes for boys and cups for girls. The exposure assessment questionnaires asked about activities and specific exposures such as diet with high PAH contents and indoor [secondhand smoke (SHS) exposure] and/or outdoor sources of exposure (transportation, time spent for the transportation, and location of outdoor play) during 1 d before the urine sampling. Children filled them out at home and, if needed, parents or guardians helped them. They were instructed to get the first spot urine samples next morning and bring back the samples together with the completed questionnaires to school for 3 consecutive days. We collected the urine samples and questionnaires each morning when the children attended school. Children were 7 to 15 y old with the mean age of 11.1 y. Thirty-two of 72 participants were females. All participants provided written informed consent before study participation.

Determination of Urinary 1-OHP. The first spot urine samples in the morning were collected in a sterile sampling tube. The collected urine samples were stored at freezer (-20°C) in school infirmaries less than 3 d, and then the frozen samples with packed ice were sent to the laboratory for the storage at deep freezer (-70°C) until the final analysis. The 1-OHP in urine was analyzed by a reverse-phase high-performance liquid chromatography method with enzymatic hydrolysis using β -glucuronidase/arylsulfatase (17) as previously described (16). Excitation and emission wavelengths were 242 and 388 nm, respectively. A reverse-phase C18 column (250 mm \times 4.6 mm, 5 μm , Supelco) was used for the segregation of 1-OHP. The detection limit of the method was 0.021 nmol 1-OHP/L urine. Urinary creatinine, as a marker of renal clearance, was measured according to the Jaffe method (18) and used for the correction of urinary concentration. The concentrations of 1-OHP were expressed as $\mu\text{mol}/\text{mol}$ creatinine.

Ambient Particulate Matter Assessment. Some previous studies used particulate pollutants as a proxy measure of ambient PAHs (16, 19). Recent study showed that >90% of the ambient PAHs measured at different regions of Korea were captured in PM_{10} fraction (20). Due to the lack of available data about ambient PAHs at the studied sites, we used the mean PM_{10} levels with a lag of 1 to 7 d before the urine sampling as the proxy indices of atmospheric PAH pollution. The Ministry of Environment of Republic of Korea is operating an ambient air monitoring station at the nearby site. The data on particulate matter <10 μm in aerodynamic diameter (PM_{10}) were measured every 5 min throughout the year using β -ray absorption method.³

Statistical Analysis. All the statistical analyses were done using the Statistical Analysis System software, version 9.1 (SAS Institute). Due to the skewed distribution, \log_{10} -transformed creatinine-adjusted 1-OHP concentrations were used to better approximate the normality assumption. To compare the differences in sex and age distributions between the two study groups, χ^2 test and Student's t test were adopted. The difference in the mean 1-OHP level between the two study groups was tested by Student's t test after log transformation. In addition, descriptive statistics were calculated, especially monthly mean, SD, geometric mean (GM), and geometric SD (GSD) of 1-OHP.

Two-day lag was selected in testing the effects of PM_{10} on 1-OHP levels based on graphical inspections of different lag periods and indicators of goodness of fit in the Proc Mixed models (Akaike information criterion, Bayesian information criterion, and corrected Akaike information criterion) as well as the reports of short half-lives of 1-OHP (14).

To examine the seasonal and regional variations of urinary 1-OHP levels and the effects of other contributing factors, we did multivariate mixed model analysis using log-transformed 1-OHP data. The intraindividual regression coefficients were considered to be random. Individual-level covariates, that is, demographic factors (age and gender), dietary source (charbroiled meat), indoor source [environmental tobacco smoke (ETS)], outdoor source (ambient PM_{10}), and season, were treated as fixed effects. Dummy variables were used to represent nominal variables, such as gender (boy versus girl), region (remote versus nearby), ETS, and diet. Seasons were categorized into spring (March to May), summer (June to July), fall (September to November), and winter (December) according to the classification by Korean Meteorological Administration.⁴ Spring, summer, and fall were included into the model using winter as the reference.

We estimated the percent change in creatinine-adjusted 1-OHP levels by each nominal variables as $(10^{\beta} - 1) \times 100\%$ with 95% confidence intervals (95% CI) of $[10^{(\beta \pm 1.96 \times \text{SE})} - 1] \times 100\%$ and for PM_{10} lag 2 as $(10^{\beta \times \text{IQR}} - 1) \times 100\%$ with 95% CI of $\{10^{[\text{IQR} \times (\beta \pm 1.96 \times \text{SE})]} - 1\} \times 100\%$, where β and SE are the estimated regression coefficient and its SE. We also estimated the percent change in 1-OHP levels associated with study region after stratifying by season.

³ <http://www.airkorea.or.kr>

⁴ <http://www.kma.go.kr/>

Table 1. General characteristics of study subjects

Variables	Overall, <i>n</i> (%)	Nearby site, <i>n</i> (%)	Remote site, <i>n</i> (%)	<i>P</i>
Sex				
Girl	32 (44)	17 (53)	15 (47)	0.27*
Boy	40 (56)	21 (53)	19 (47)	
Age, <i>y</i> (mean ± SD)	11.1 ± 2.5	11.2 ± 2.4	10.9 ± 2.7	0.64 [†]
1-OHP (μmol/mol creatinine) [‡]	0.050 ± 2.435	0.062 ± 2.112	0.040 ± 2.647	<0.0001 [†]

**P* value was calculated by χ^2 test.

[†]*P* values were calculated by Student's *t* test after log transformation.

[‡]GM ± GSD.

Results

Descriptive Analysis. Out of total 1,944 (72 subject × 3 days × 9 months) originally scheduled urine samples, 1,497 (77%) samples were actually collected. The number of urine samples finally available was 6 to 27 samples per subject. Twenty-two urine samples were excluded because of the errors in analytic procedure. The GM of urinary 1-OHP concentration for the total urine samples was 0.050 μmol/mol creatinine. When stratified by region, it was 0.062 μmol/mol creatinine for the nearby site and 0.040 μmol/mol creatinine for the remote site. The 1-OHP levels of nearby children were significantly higher than those of remote children (*P* < 0.0001), as shown in Table 1.

The monthly mean, SD, GM, and GSD of the creatinine-corrected 1-OHP levels by study sites are shown in Table 2. The overall monthly means of 1-OHP were higher for nearby site than remote site, but the absolute levels and differences varied by seasons. The difference between study sites was the largest during summer (GM = 0.087 μmol/mol creatinine for nearby site; GM = 0.036 μmol/mol creatinine for remote site).

Lag Effects of PM₁₀ as a Proxy Measure of Ambient PAHs. Monthly variation of creatinine-adjusted 1-OHP and ambient PM₁₀ levels are shown in Fig. 1. The monthly variation of 1-OHP followed a similar track of PM₁₀ variation. The ambient PM₁₀ levels were higher in June and July, and 1-OHP levels were also higher during the same periods.

The ambient monitoring data of PM₁₀ were used as a proxy measure of ambient PAHs, and mixed-effects

models incorporating PM₁₀ were analyzed with 1 to 7 days of lags before the urine sampling date. The daily mean PM₁₀ levels of 2, 3, and 7 days before survey (lag 2, 3, and 7) showed significantly positive associations with the creatinine-adjusted 1-OHP. Considering the distribution of different lag effects, the 2-day lag model was chosen as the most appropriate for evaluating the effect of air pollution on urinary 1-OHP level as shown in Fig. 2.

Model of Contributing Factors. Table 3 presents the estimated percent changes of 1-OHP for potential covariates. The urinary 1-OHP concentration increased by 17.6% (95% CI, 2.1-35.3%; *P* = 0.022) for girls compared with boys, by 17.6% (95% CI, 1.9-35.7%; *P* = 0.027) for charbroiled meat consumption, and by 62.1% (95% CI, 39.7-88.2%; *P* < 0.0001) for nearby children compared with remote children as shown in model I. Urinary 1-OHP concentrations in summer and fall were significantly higher than those of winter (model II): a 40.9% (95% CI, 20.6-64.7%; *P* < 0.0001) increase for summer and a 37.8% (95% CI, 18.6-60.2%; *P* < 0.0001) increase for fall compared with winter levels. The seasonal effects somewhat decreased after adjusting for south wind from a steel mill (model III): 30.2% (95% CI, 9.9-54.3%; *P* = 0.002) for summer and 34.0% (95% CI, 15.1-55.9%; *P* = 0.0002) for fall. The 1-OHP level increased by 10.1% (95% CI, 4.8-15.7%; *P* < 0.021) for the interquartile range (IQR) increment of PM₁₀ lag 2 (model IV). The magnitudes of the percent changes for both summer and fall also diminished after adjusting for PM₁₀ lag 2. Especially, the percent change by summer effect dropped to 20.4% (95% CI, 1.2-43.2%; *P* = 0.036).

Table 2. Means and SDs of creatinine-adjusted urinary 1-OHP levels by month and season

Season	Month	Nearby site					Remote site				
		<i>n</i>	Mean	SD	GM	GSD	<i>n</i>	Mean	SD	GM	GSD
Spring	March	105	0.061	0.038	0.053	1.716	102	0.056	0.042	0.044	2.285
	April	101	0.062	0.040	0.053	1.755	90	0.048	0.042	0.039	1.861
	May	93	0.070	0.073	0.047	2.654	78	0.052	0.065	0.035	2.403
	Total	299	0.064	0.052	0.051	2.035	270	0.052	0.050	0.039	2.186
Summer	June	84	0.107	0.075	0.081	2.344	83	0.065	0.081	0.036	3.580
	July	88	0.113	0.067	0.094	1.931	71	0.062	0.056	0.036	3.650
	Total	172	0.110	0.071	0.087	2.139	154	0.064	0.070	0.036	3.597
Fall	September	74	0.072	0.047	0.060	1.966	69	0.060	0.043	0.046	2.265
	October	70	0.087	0.075	0.071	1.770	71	0.082	0.097	0.061	1.966
	November	79	0.079	0.047	0.068	1.744	61	0.054	0.035	0.042	2.360
	Total	223	0.079	0.057	0.066	1.831	201	0.066	0.067	0.049	2.218
Winter	December	84	0.075	0.063	0.055	2.501	72	0.047	0.045	0.029	3.267

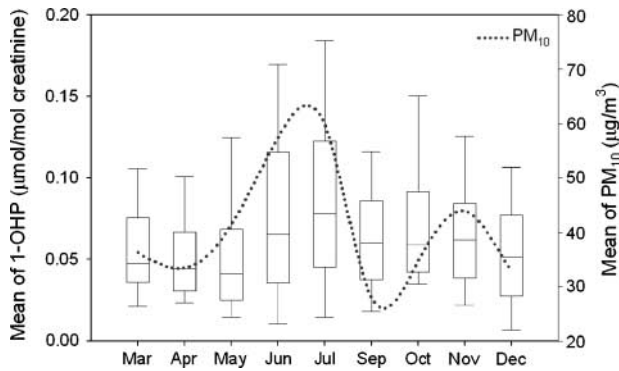


Figure 1. Distribution of creatinine-adjusted 1-OHP levels and mean levels of ambient PM₁₀ by month.

Discussion

To explore the chronic variation of environmental exposure to PAHs, biomonitoring with multiple measurements, at different time points of day, months, and seasons, can be adopted. Such a strategic design, to our knowledge, has not been yet reported in the literature. Repeated sampling design may have affected sample collection compliance. The collection rate was 77% of the total scheduled urine samples (1,944). During the overall study period, daily collection rate was 58% to 99% of the possible 72 urine samples. Although sampling rate had somewhat varied by season (88% for spring, 76% for summer, 65% for fall, and 72% for winter), there was no trend after the initial decline.

Our study focused primarily on environmental exposure to PAHs from a point industrial source of steel mill, whereas other studies concentrated on traffic and/or dietary exposures to PAHs (9, 21–24). Similar to our study, the spatial effect by different distances from a point source was reported by the study from Ukraine (12). The environmental exposure to PAHs at the steel mill site was compared with that in capital city, but only one cross-sectional measurement was taken without considering dietary PAH intake. Our study was to determine spatial differences in chronic exposure levels using repeated samples to account for the temporal variation.

Several recent studies among children have shown that diet is the most distinct contributor to 1-OHP levels

(21, 22). Among the dietary food items, the ingestion of charbroiled meat is one of the main sources of PAH exposure (1, 25). Whereas other Korean study reported no relationship (19), significant contributions of dietary intake to the 1-OHP level were found in this study based on dietary questionnaire after adjusting for other intra-individual variation.

Several studies examined the effect of gender on children's 1-OHP concentrations (12, 15, 23, 26). The gender effect was inconsistent. Several studies reported higher 1-OHP levels for girls (12, 16), whereas others reported higher levels for boys (15, 26). This study found significantly higher level of 1-OHP for girls when creatinine-adjusted log-transformed values were analyzed (estimate = 0.087; $P = 0.031$), whereas the difference was not statistically significant with creatinine-unadjusted values (estimate = 0.004; $P = 0.941$). Boys, especially those in junior high schools, are bigger and more masculine than girls and, therefore, excrete more creatinine in urine, resulting in the smaller 1-OHP values when adjusted with creatinine. In this way, the differences in body mass might have contributed to this gender-related difference in creatinine-adjusted log-transformed values. In addition, as we did not take account of the amount of intake, the difference in the frequency and amount of charbroiled foods between boys and girls may also have influenced this gender-related difference.

SHS from household members can be an important contributor to PAH exposure. Dutch studies showed the contribution of SHS in nonoccupationally exposed male adults (27) and children (28), whereas several studies (26, 29) did not. Ukraine study showed an effect of SHS among total groups of children but not within unexposed and exposed group (12). In our results, exposure to SHS did not contribute to 1-OHP levels. Inaccurate estimation of the SHS based on questionnaire, and the amounts and conditions of SHS in South Korea, especially housing structures, ventilations under different weather conditions, and the change of parent's smoking behavior at home, may have influenced the results.

Because the ambient PAH data were not available at the study site, we were not able to assess the direct contribution of ambient PAHs to 1-OHP levels. Ambient PM₁₀ was used as a proxy measure of ambient PAH exposure. More than 70% of PAHs are adsorbed onto suspended particles at ambient temperatures (20). Lighter PAHs with two to three benzene rings are mostly present in gas phase, whereas those with four or more

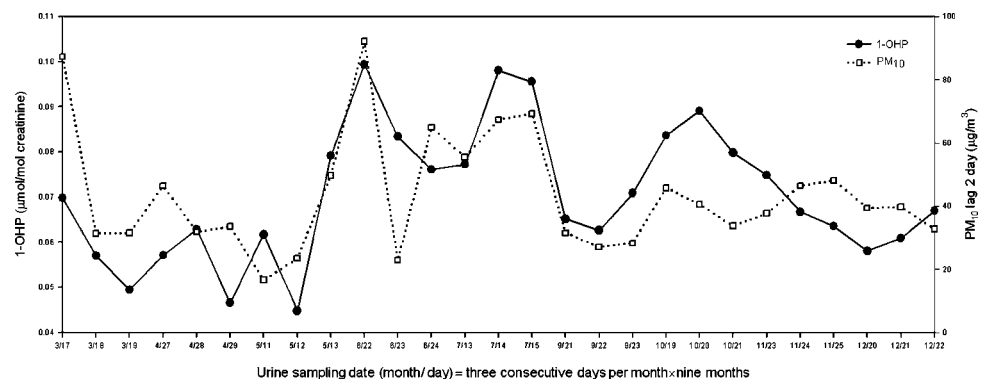


Figure 2. Mean levels of urinary 1-OHP and PM₁₀ lag 2 day during study periods.

Table 3. Estimated percent changes (95% CIs) in 1-OHP ($\mu\text{mol/mol}$ creatinine) levels and four-random effects models for contributing factors

		Estimated percent change (95% CI)			
		Model I	Model II	Model III	Model IV
Age		-1.7 (-4.5 to 1.3)	-1.4 (-4.3 to 1.5)	-1.5 (-4.4 to 1.4)	-1.6 (-4.4 to 1.4)
Gender	Boy	Reference	Reference	Reference	Reference
	Girl	17.6 (2.1-35.3)*	16.8 (1.5-34.4)*	16.3 (1.1-33.9)*	16.5 (1.2-34.1)*
ETS	No	Reference	Reference	Reference	Reference
	Yes	-8.8 (-20.5 to 4.7)	-8.1 (-19.9 to 5.4)	-7.8 (-19.6 to 5.8)	-8.0 (-19.8 to 5.5)
Food	No	Reference	Reference	Reference	Reference
	Yes	17.6 (1.9-35.7)*	18.8 (3.1-37.0)*	18.4 (2.7-88.5)*	16.5 (1.1-34.2)*
Region	Remote	Reference	Reference	Reference	Reference
	Nearby	62.5 (40.2-88.5) [†]	62.1 (39.7-88.2) [†]	62.4 (40.0-88.5) [†]	62.3 (39.8-88.3) [†]
Season	Spring		8.7 (-6.0 to 25.8)	6.3 (-8.2 to 23.0)	6.8 (-7.7 to 23.5)
	Summer		40.9 (20.6-64.7) [†]	30.2 (9.9-54.3)*	20.4 (1.2-43.2)*
	Fall		37.8 (18.6-60.2) [†]	34.0 (15.1-55.9) [†]	36.1 (16.9-58.3) [†]
	Winter		Reference	Reference	Reference
South wind (%)				28.5 (3.8-59.1)*	9.3 (-13.0-37.4)
PM ₁₀ lag 2 [‡]					10.1 (4.8-15.7) [‡]

NOTE: Coefficients are expressed as percent changes in 1-OHP-associated contributing factors: model I included age (years), gender, ETS, and charbroiled meat consumption (yes/no); model II included model I plus season; model III included model II plus south wind; and model IV included model III plus PM₁₀ lag 2.

* $P < 0.05$.

[†] $P < 0.001$.

[‡] Percent change (95% CI) based on IQR (75-25%) of PM₁₀ lag 2.

rings, including carcinogenic PAHs such as benzo(a) pyrene, are mainly found in particle phase (30). The majority of PAHs exist predominantly on ultrafine particles of $<0.1 \mu\text{m}$ in diameter in Los Angeles (31) or fine particles of 0.1 to $2.5 \mu\text{m}$ in diameter in China (32). More than 60% of ambient PAHs were in particles with diameter of $<3 \mu\text{m}$ and $>90\%$ of PAHs were in particles with diameter of $<9 \mu\text{m}$ at studied sites in Korea (20). The concentration of airborne pyrene in urban fine particles was the highest during summer (32) and 43% of total 16 PAHs, the highest value among the measurements, was pyrene during summer in Ulsan, Korea (33).

Over the study period, ambient PM₁₀ had ranged from 30.29 to 88.66 $\mu\text{g}/\text{m}^3$ (mean, 51.32 $\mu\text{g}/\text{m}^3$) during the 1 week before urine collection. Except for September, October, and December, the mean values of PM₁₀ exceeded the air quality standard of the annual mean of Korean Ministry of Environment (50 $\mu\text{g}/\text{m}^3$) and United Kingdom (40 $\mu\text{g}/\text{m}^3$). In a Ukraine study (12), much higher annual level of particulate (200 $\mu\text{g}/\text{m}^3$) was reported, and even the level in their control area (100 $\mu\text{g}/\text{m}^3$) was higher than in our study. Very high level of ambient pollution may have increased the level of 1-OHP among Ukraine children, but beyond this ecological association, no further temporal or spatial variation was examined. In this study, short-term contribution of ambient PM₁₀ could be captured by the repeated measurements over 3 days.

In addition to the short-term variation, we observed significant seasonal variations in 1-OHP levels among children, especially higher during summer and fall compared with winter. Very few studies reported the seasonal variations in biomonitoring. Higher levels of 1-OHP during winter were found among occupational employees (34, 35). One study found the seasonal differences only among nonsmoking employees (34), whereas the other study attributed the seasonality to the variation in sources such as residential heating and

higher traffic density during winter time in urban area (35). In our study, seasonal variation could be explained largely by ambient PM₁₀ levels and possibly by wind directions (models II, III, and IV in Table 3). Differences in indoor residence time and outdoor activities between different seasons might have also contributed to this seasonal variation.

This is also consistent with our results on the contribution of regional effects for different seasons (Fig. 3). Regional effect by season was calculated as the percent change in creatinine-adjusted 1-OHP levels by region as $(10^\beta - 1) \times 100\%$ with 95% CI of $[10^{(\beta \pm 1.96 \times \text{SE})} - 1] \times 100\%$, where β and SE are the estimated regression coefficient and its SE in a

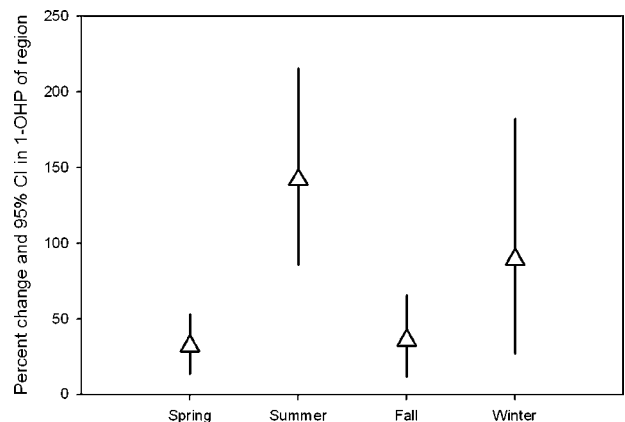


Figure 3. Estimated percent change in creatinine-adjusted 1-OHP levels for region by season. The covariates adjusted for age, sex, ETS, and charbroiled meat consumption. Each triangle and bar represents the percent change and 95% lower and upper confidence intervals.

multivariate mixed model analysis. Overall, the regional effect was the largest during the summer (Fig. 3) when the relative contribution of PM₁₀ to the 1-OHP was also the largest compared with other seasons (models II and IV in Table 3). These results suggest that when seasonal variation in the exposure level is suspected, the assessment of chronic exposure to the ambient PAHs should cover at least two different periods, including seasons of high and low exposures.

Disclosure of Potential Conflicts of Interest

No potential conflicts of interest were disclosed.

Acknowledgments

The costs of publication of this article were defrayed in part by the payment of page charges. This article must therefore be hereby marked *advertisement* in accordance with 18 U.S.C. Section 1734 solely to indicate this fact.

References

- IARC. Overall evaluations of carcinogenicity. IARC Monogr Eval Carcinog Risks Hum 1-42. Supplement No. 7. Lyon (France): IARC; 1987.
- Ahn YS, Park RM, Stayner L, Kang SK, Jang JK. Cancer morbidity in iron and steel workers in Korea. *Am J Ind Med* 2006;49:647-57.
- Austin H, Delzell E, Lally C, Rotimi C, Oestestad K. A case-control study of lung cancer at a foundry and two engine plants. *Am J Ind Med* 1997;31:414-21.
- Rodriguez V, Tardon A, Kogevinas M, et al. Lung cancer risk in iron and steel foundry workers: a nested case control study in Asturias, Spain. *Am J Ind Med* 2000;38:644-50.
- Xu Z, Brown LM, Pan GW, et al. Cancer risks among iron and steel workers in Anshan, China. Part II: case-control studies of lung and stomach cancer. *Am J Ind Med* 1996;30:7-15.
- Armstrong B, Tremblay C, Baris D, Theriault G. Lung cancer mortality and polynuclear aromatic hydrocarbons: a case-cohort study of aluminum production workers in Arvida, Quebec, Canada. *Am J Epidemiol* 1994;139:250-62.
- Pless-Mulloli T, Phillimore P, Moffatt S, et al. Lung cancer, proximity to industry, and poverty in northeast England. *Environ Health Perspect* 1998;106:189-96.
- Bhopal RS, Phillimore P, Moffatt S, Foy C. Is living near a coking works harmful to health? A study of industrial air pollution. *J Epidemiol Community Health* 1994;48:237-47.
- Northridge ME, Yankura J, Kinney PL, et al. Diesel exhaust exposure among adolescents in Harlem: a community-driven study. *Am J Public Health* 1999;89:998-1002.
- Dejmek J, Solansky I, Benes I, Lenicek J, Sram RJ. The impact of polycyclic aromatic hydrocarbons and fine particles on pregnancy outcome. *Environ Health Perspect* 2000;108:1159-64.
- Perera FP, Rauh V, Whyatt RM, et al. Effect of prenatal exposure to airborne polycyclic aromatic hydrocarbons on neurodevelopment in the first 3 years of life among inner-city children. *Environ Health Perspect* 2006;114:1287-92.
- Mucha AP, Hryhorczuk D, Serdyuk A, et al. Urinary 1-hydroxypyrene as a biomarker of PAH exposure in 3-year-old Ukrainian children. *Environ Health Perspect* 2006;114:603-9.
- Seidel A, Dahmann D, Krekler H, Jacob J. Biomonitoring of polycyclic aromatic compounds in the urine of mining workers occupationally exposed to diesel exhaust. *Int J Hyg Environ Health* 2002;204:333-8.
- Viau C, Vyskocil A, Martel L. Background urinary 1-hydroxypyrene levels in non-occupationally exposed individuals in the Province of Quebec, Canada, and comparison with its excretion in workers exposed to PAH mixtures. *Sci Total Environ* 1995;163:191-4.
- Jongeneelen FJ. Biological monitoring of environmental exposure to polycyclic aromatic hydrocarbons; 1-hydroxypyrene in urine of people. *Toxicol Lett* 1994;72:205-11.
- Lee MS, Eum KD, Zoh KD, Kim TS, Pak YS, Paek D. 1-Hydroxypyrene as a biomarker of PAH exposure among subjects living in two separate regions from a steel mill. *Int Arch Occup Environ Health* 2007;80:671-8.
- Jongeneelen FJ, Anzion RB, Henderson PT. Determination of hydroxylated metabolites of polycyclic aromatic hydrocarbons in urine. *J Chromatogr* 1987;413:227-32.
- Jaffe M. Creatinine Jaffe endpoints methods with deproteinization. *Z Physiol Chem* 1986;10:391-400.
- Kang JW, Cho SH, Kim H, Lee CH. Correlation of urinary 1-hydroxypyrene and 2-naphthol with total suspended particulates in ambient air in municipal middle-school students in Korea. *Arch Environ Health* 2002;57:377-82.
- Bae SY, Yi SM, Kim YP. Temporal and spatial variations of the particle size distribution of PAHs and their dry deposition fluxes in Korea. *Atmos Environ* 2002;36:5491-500.
- Fiala Z, Vyskocil A, Krajak V, et al. Environmental exposure of small children to polycyclic aromatic hydrocarbons. *Int Arch Occup Environ Health* 2001;74:411-20.
- Vyskocil A, Fiala Z, Chenier VV, et al. Assessment of multipathway exposure of small children to PAH. *Environ Toxicol Pharmacol* 2000;8:111-8.
- Kanoh T, Fukuda M, Onozuka H, Kinouchi T, Ohnishi Y. Urinary 1-hydroxypyrene as a marker of exposure to polycyclic aromatic hydrocarbons in environment. *Environ Res* 1993;62:230-41.
- Zhao ZH, Quan WY, Tian DH. Urinary 1-hydroxypyrene as an indicator of human exposure to ambient polycyclic aromatic hydrocarbons in a coal-burning environment. *Sci Total Environ* 1990;92:145-54.
- Kang DH, Rothman N, Poirier MC, et al. Interindividual differences in the concentration of 1-hydroxypyrene-glucuronide in urine and polycyclic aromatic hydrocarbon-DNA adducts in peripheral white blood cells after charbroiled beef consumption. *Carcinogenesis* 1995;16:1079-85.
- Siwinska E, Mielzynska D, Smolik E, Bubak A, Kwapulinski J. Evaluation of intra- and interindividual variation of urinary 1-hydroxypyrene, a biomarker of exposure to polycyclic aromatic hydrocarbons. *Sci Total Environ* 1998;217:175-83.
- Van Rooij JG, Veeger MM, Bodelier-Bade MM, Scheepers PT, Jongeneelen FJ. Smoking and dietary intake of polycyclic aromatic hydrocarbons as sources of interindividual variability in the baseline excretion of 1-hydroxypyrene in urine. *Int Arch Occup Environ Health* 1994;66:55-65.
- van Wijnen JH, Slob R, Jongmans-Liedekerken G, van de Weerdt RH, Woudenberg F. Exposure to polycyclic aromatic hydrocarbons among Dutch children. *Environ Health Perspect* 1996;104:530-4.
- Scherer G, Conze C, Tricker AR, Adlkofer F. Uptake of tobacco smoke constituents on exposure to environmental tobacco smoke (ETS). *Clin Investig* 1992;70:352-67.
- Park SS, Kim YJ, Kang CH. Atmospheric polycyclic aromatic hydrocarbons in Seoul, Korea. *Atmos Environ* 2002;36:2917-24.
- Li N, Sioutas C, Cho A, et al. Ultrafine particulate pollutants induce oxidative stress and mitochondrial damage. *Environ Health Perspect* 2003;111:455-60.
- Zhou JB, Wang TG, Huang YB, Mao T, Zhong NN. Size distribution of polycyclic aromatic hydrocarbons in urban and suburban sites of Beijing, China. *Chemosphere* 2005;61:792-9.
- Lee BK, Lee CB. Development of an improved dry and wet deposition collector and the atmospheric deposition of PAHs onto Ulsan Bay, Korea. *Atmos Environ* 2004;38:863-71.
- Pastorelli R, Guanci M, Restano J, et al. Seasonal effect on airborne pyrene, urinary 1-hydroxypyrene, and benzo(a)pyrene diol epoxide-hemoglobin adducts in the general population. *Cancer Epidemiol Biomarkers Prev* 1999;8:561-5.
- Perico A, Gottardi M, Boddi V, Bavazzano P, Lanciotti E. Assessment of exposure to polycyclic aromatic hydrocarbons in police in Florence, Italy, through personal air sampling and biological monitoring of the urinary metabolite 1-hydroxypyrene. *Arch Environ Health* 2001;56:506-12.