Use of Urinary PAH Metabolites to Assess PAH Exposure Intervention among Coke Oven Workers

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Abstract: Use of Urinary PAH Metabolites to Assess PAH Exposure Intervention among Coke Oven Workers: Soo-Hun Cho, et al. Department of Preventive Medicine, Seoul National University College of Medicine and Institute of Environmental Medicine, SNUMRC—To assess the effectiveness of protective skin coveralls in reducing skin contamination among coke-oven workers, 1-hydroxypyrene glucuronide (1-OHPG) was used as an internal dose marker of polycyclic aromatic hydrocarbon (PAH) exposure. Twenty coke-oven workers at a steel plant in South Korea provided their first morning void urine samples before beginning work, as well as postshift urine samples after working for five days with regular skin protection. Pre- and postshift urine samples from the same workers were collected after new skin coveralls made from Tyvek® had been worn during the week following regular skin protection. Pre-and postshift urine samples from the same workers were collected after new skin coveralls made from Tyvek® had been worn during the week following regular skin protection. Urine samples were quantitated for 1-OHPG by immunoaffinity purification and HPLC with a fluorescence detector. The benzene-soluble fraction (BSF) of coke oven emissions (COE) in personal air samples from the 20 workers was also quantitated. To obtain information relating to smoking, job history, dietary habits, drug use, past medical history, and the use of personal protective equipment, a self-administered questionnaire was used. The mean age of study participants was 39.7 yr and the average length of employment was 12 yr (11 months - 18 yr). In 20 workers, there was a statistically significant correlation between ambient COE and urinary 1-OHPG levels during the period of regular skin protection (r=0.50, p<0.05). The difference in 1-OHPG levels between post- and preshift urine samples using regular skin protection was higher than when the new skin coveralls were worn. Although this was not statistically significant, there was a statistically significant difference in 1-OHPG among topside workers (p<0.05). These results indicate that the introduction of the new skin coverall resulted in significant reductions of urinary PAH metabolites among workers exposed to higher levels of PAHs. The measurement of PAH metabolites in human urine appears to be ideally suited to biomonitoring in the workplace and testing the effectiveness of attempts to reduce PAH exposure.

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Key words: Urinary PAH metabolites, Coke-oven workers, Intervention

Increased risk of lung cancer is associated with employment in the coke production areas of the steel industry¹. The risk is believed to be due primarily to elevated air concentrations of polycyclic aromatic hydrocarbons (PAHs) found in this work environment²-⁴. A useful and direct approach to the assessment of human exposure and uptake of PAHs is to measure PAH metabolites excreted in urine⁵-¹⁰. The measurement of PAH metabolites in human urine provides a means of assessing an individual’s internal dose of PAHs¹¹. A rapid assay for quantitation of 1-hydroxypyrene-glucuronide (1-OHPG) in human urine has been developed¹². The assay has been applied in three validation studies of individuals with PAH exposure due to diet, occupation and/or smoking. Dietary exposure was examined in a controlled feeding study of charbroiled beef¹³. Occupational and smoking exposure was
investigated in a study of steel plant workers and on-site controls\textsuperscript{14}. Smoking exposure was also examined in a study of smokers of blond or black tobacco\textsuperscript{15}.

The development and institution of successful intervention to reduce PAH exposure levels in occupational settings requires evaluation of the methodologies involved, and a limited number of studies have examined the usefulness of urinary PAH metabolites for assessing the effect of such intervention\textsuperscript{16–18}. The measures involved included new workwear policy (e.g. new coveralls, shirt, trousers, underwear, socks, and boots) and improved skin protection measures (e.g. washing the hands and face before a break, taking a shower after work, putting on new gloves and laundered work clothes) including special skin coveralls made from Tyvek\textsuperscript{6}. The effect of such intervention, as assessed by the reduction of urinary 1-hydroxypyrene (1-OHP) levels, ranged from 35% to 50\%\textsuperscript{16–18}.

We recently conducted a study to evaluate several extra skin protective measures among 25 coke-oven workers at a steel plant in Korea\textsuperscript{19}. First morning void urine samples of the 25 workers were collected before work, and postshift urine samples were collected after five days’ work with regular skin protection. Pre- and postshift urine samples from the same workers were collected after several extra skin protective measures had been taken. These measures included: 1) encouraging washing the hands and face before each rest break; 2) the daily provision of clean work clothes; 3) the daily provision of new neck towels, gloves, and socks; 4) encouraging the use of clean underwear everyday; 5) the use of wristlets and skin protective creams. The results indicated that extra skin protective measures significantly reduced the excretion of PAH metabolites in coke oven workers by 31\%.

This study aimed to assess the effectiveness of reduced PAH exposure among coke-oven workers through the introduction of special skin coveralls made from Tyvek\textsuperscript{6}. To this end, the usefulness of urinary PAH metabolites, 1-OHP and 1-OHPG, as internal dose markers of PAH exposure was evaluated.

**Materials and Methods**

1) Study design and subjects

On the basis of their job title, 20 coke-oven workers were recruited from a steel plant in South Korea. Eight were topside workers, eight were oven side workers, and four were maintenance workers. The mean age of study participants was 39.7 yr and their length of employment at the plant averaged 12 yr (11 months - 18 yr). It was reported that at the time of the study, 83\% of participants were wearing protective respiratory equipment (Table 1). Seventy percent of subjects were current smokers and their average consumption was 12 pack-years and there were no significant differences in demographical characteristics among workers of different job titles.

First morning void urine samples were collected before work and postshift urine samples were collected after five days’ work with regular skin protection. Pre- and postshift urine samples of the same workers were collected after new skin coveralls made from Tyvek\textsuperscript{6} (DuPont, USA) had been worn during the week following regular skin protection. Clothing made from Tyvek\textsuperscript{6}, a spun-bonded olefin sheet structure of 100\% polyethylene, is known for its low permeability to particulates. Each worker had been provided with the skin coveralls everyday before they began to work. A self-administered questionnaire was used to obtain information about smoking habits, job history, major diseases, dietary habits, drug use, and the previous use of personal protective equipment. The design of the entire study is shown in Fig. 1.

2) Urinary 1-hydroxypyrene glucuronide (1-OHPG) measurement

Urine samples were quantitated for 1-OHPG by immunoaffinity purification and high performance liquid chromatography (HPLC) with a fluorescence detector, using a modification of the assay developed by Strickland \textit{et al.}\textsuperscript{12}. To hydrolyze acid-labile metabolites, 2 ml urine samples were treated with 0.1 N HCl (90°C). The hydrolyzed samples were loaded onto Sep-Pak C\textsubscript{18} cartridges (Waters), washed with 30\% methanol, and the relatively non-polar metabolites were eluted with 80\% methanol. Concentrated samples were loaded into immunoaffinity columns (IAC) containing CNBr-activated sepharose 4B coupled with monoclonal antibody

<table>
<thead>
<tr>
<th>Variables</th>
<th>Mean</th>
<th>SD</th>
</tr>
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<tbody>
<tr>
<td>Age (yr)</td>
<td>39.7</td>
<td>5.2</td>
</tr>
<tr>
<td>Body surface area (m\textsuperscript{2})</td>
<td>1.8</td>
<td>0.1</td>
</tr>
<tr>
<td>Duration of employment (months)</td>
<td>141.5</td>
<td>56.3</td>
</tr>
<tr>
<td>Smoking History (PY)</td>
<td>11.7</td>
<td>6.3</td>
</tr>
</tbody>
</table>

*Current smoker*: 70.0\%  
*Wearing Resp. PPE*: 83.3\%
which recognizes several PAH adducts and metabolites. Bound material was eluted with 40% methanol and eluated samples from the immunoaffinity columns were dried at 65°C in a vacuum and redissolved in 120 µl water. Each 20 µl of prepared samples was injected and analyzed by HPLC with a fluorescence detector (Waters gradient controller, Waters 515 pump, Waters 717 plus autosampler, Waters 474 scanning fluorescence detector and Autochro-Win chromatography data system, Young In, Korea). Gradient elution using methanol and water was carried out (10% methanol for 5 min, 10–100% methanol for 25 min and 100% methanol for 10 min) at a flow rate of 1 ml/min. The fluorescence detector was fixed as λex=347 nm and λem=381 nm.

3) Urinary 1-hydroxypyrene (1-OHP) measurement
Urinary 1-OHP concentration was measured by HPLC with a fluorescence detector. The method was slightly modified from that originally developed by Jongeneelen et al.

4) Coke-oven emission measurements
Benzene soluble fractions (BSF) of coke oven emissions (COE) in personal air samples from the 20 workers were quantitated by the NIOSH analytical method No. 5506. Table 2 shows COE concentrations in different workplaces at the plant during the period 1992–1995.

5) Statistical analysis
According to the Shapiro-Wilks W statistic, 1-OHPG levels observed in this study were log-normally distributed, and a log transformed 1-OHPG concentration (LOGOHPG) was used for their further analysis. The correlation between log transformed COE (LOGCOE) and urinary LOGOHPG was assessed by Pearson’s correlation analysis.

Differences in LOGOHPG between pre- and postshift urine samples with regular skin protection was higher than when the new skin coveralls were being worn (Fig. 3). Although the difference in 1-OHPG levels between post- and preshift urine samples using the new skin coverall (1.8 µg/g Cr) was lower than when regular clothing was worn (2.7 µg/g Cr), this difference was not statistically significant. Data analysis on the basis of job category showed that only among topside workers were 1-OHPG increments significantly lower when special skin coveralls were worn than during regular skin protection (p<0.05, by Wilcoxon’s rank sum test). Simple comparison of DLOGOHPG (Wilcoxon’s rank sum test) and ANCOVA adjustment for BMI was performed.

Results
There was a statistically significant correlation between concentrations of 1-OHP and 1-OHPG in the same urine samples (r=0.65, p<0.0001, Fig. 2). There was a statistically significant correlation between ambient COE and urinary 1-OHPG levels during the period of regular skin protection (r=0.50, p<0.05, data not shown).

The difference in 1-OHPG levels between post- and preshift urine samples with regular skin protection was higher than when the new skin coveralls were being worn (Fig. 3). Although the difference in 1-OHPG levels between post- and preshift urine samples using the new skin coverall (1.8 µg/g Cr) was lower than when regular clothing was worn (2.7 µg/g Cr), this difference was not statistically significant. Data analysis on the basis of job category showed that only among topside workers were 1-OHPG increments significantly lower when special skin coveralls were worn than during regular skin protection (p<0.05, by Wilcoxon’s rank sum test for log-transformed data; Table 3).

In order to assess the effect of special skin coveralls on urinary 1-OHPG levels, residuals were compared between the use of regular skin protection and the special skin coveralls. They were calculated using the regression model after adjusting COE levels and BMI (model: LOGOHPG=b0 + b1*(LOGCOE) + b2*BMI). Although urinary DLOGOHPG levels in all study participants were not significantly different for both simple and multivariate comparison (adjusting for LOGCOE and BMI), there were significant differences in topside workers between the regular and special coveralls periods for simple and

<table>
<thead>
<tr>
<th>Year</th>
<th>Top side</th>
<th>Coke ovens</th>
<th>Maintenance sites</th>
</tr>
</thead>
<tbody>
<tr>
<td>Apr, 95</td>
<td>0.08 (0.06–0.10)</td>
<td>0.03 (0.02–0.04)</td>
<td>0.06 (0.05–0.06)</td>
</tr>
<tr>
<td>Sept, 94</td>
<td>0.06 (0.05–0.08)</td>
<td>0.02 (N.D–0.06)</td>
<td>0.06 (N.D–0.09)</td>
</tr>
<tr>
<td>Apr, 94</td>
<td>0.08 (0.06–0.11)</td>
<td>0.04 (0.03–0.06)</td>
<td>0.05 (0.03–0.14)</td>
</tr>
<tr>
<td>Sept, 93</td>
<td>0.08 (0.05–0.15)</td>
<td>0.05 (0.02–0.07)</td>
<td>0.03 (0.01–0.06)</td>
</tr>
<tr>
<td>Apr, 93</td>
<td>0.08 (0.04–0.14)</td>
<td>0.03 (N.D–0.05)</td>
<td>0.04 (0.01–0.06)</td>
</tr>
<tr>
<td>Sept, 92</td>
<td>0.08 (0.01–0.17)</td>
<td>0.04 (0.01–0.13)</td>
<td>0.05 (N.D–0.14)</td>
</tr>
</tbody>
</table>

N.D=non-detectable
Table 3. COE and 1-OHPG differences according to job category

<table>
<thead>
<tr>
<th>GM (GSD)</th>
<th>Topside workers (n=8)</th>
<th>Others (n=12)</th>
</tr>
</thead>
<tbody>
<tr>
<td>COE (mg/m³)</td>
<td>1st pre: 0.053 (2.011)</td>
<td>0.027 (3.910)</td>
</tr>
<tr>
<td></td>
<td>1st post: 0.030 (2.883)</td>
<td>0.020 (2.655)</td>
</tr>
<tr>
<td></td>
<td>2nd pre: 0.041 (1.913)</td>
<td>0.039 (2.079)</td>
</tr>
<tr>
<td></td>
<td>2nd post: 0.045 (3.001)</td>
<td>0.029 (2.975)</td>
</tr>
<tr>
<td>increasing 1-OHPG₁ (µg/g Cr)</td>
<td>3.006 (0.002)</td>
<td>1.605 (0.004)</td>
</tr>
<tr>
<td>increasing 1-OHPG² (µg/g Cr)</td>
<td>2.203 (0.003)</td>
<td>1.591 (0.005)</td>
</tr>
</tbody>
</table>

₁During regular protection period.  ²After the use of special skin coveralls.
* p-value<0.05 (by Wilcoxon’s rank sum test; log-transformed data).

Fig. 2. Correlation between urinary 1-OHP and 1-OHPG levels.

Fig. 3. Concentrations of urinary 1-OHPG.
A: preshift urine samples during regular clothing.  B: postshift urine samples during regular clothing (p-value<0.05, by Wilcoxon’s rank sum test between post- and preshift urine samples during regular skin protection).  C: preshift urine samples during special skin coveralls.  D: postshift urine samples during special skin coveralls.

Discussion
In this study, we observed significantly reduced urinary 1-hydroxypyrene glucuronide (1-OHPG) excretion among topside workers exposed to higher levels of coke oven emissions (COE). Among all subjects, 1-OHPG increments during the use of special skin coveralls were lower than during the use of regular skin protection, though the significance disappears when residuals, calculated using the regression model of log-transformed multivariate comparisons (adjusting for LOGCOE and BMI, p<0.05) (Fig. 4).
1-OHPG (LOGOHPG) as a dependent variable, COE and body mass index (BMI) - which have shown significant association with LOGOHPG - as independent variables, were compared between the use of regular skin protection and the use of the special skin coveralls. The findings observed in this study have various possible explanations: 1) due to small sample size, there was not enough statistical power to build a robust regression model; 2) in this model, most 1-OHPG variation was not explained by COE, BMI, or the use of special skin coveralls, and incomplete adjustment of other confounders and individual differences in pyrene metabolism might thus affect 1-OHPG levels; 3) PAHs might penetrate the skin through uncovered areas; and 4) among these workers, skin absorption might not be the major route of exposure.

Since Van Rooij et al. first reported that in ten creosote-exposed workers, special skin coveralls reduced urinary 1-OHP excretion by about 63%, several studies assessed the effectiveness of intervention measures aimed at protecting workers from exposure to PAHs, and the subsequent excretion of PAH metabolites. Quinlan et al. reported that among ten coal liquefaction workers, new workwear policy (new coverall, shirt, trousers, underwear, socks, and boots) significantly reduced 1-OHP excretion and deposition of PAHs on the skin. Van Rooij et al. indicated that among 13 coke oven workers, simple hygienic skin protective measures (new gloves and laundered working clothes, washing both hands and face before each break) led to a 37% reduction in 1-OHP excretion.

By measuring urinary PAH metabolites as markers of effectiveness, these studies show the significant effect of skin exposure reduction and also report that skin contamination is the main determinant of internal doses of PAHs exposure. Several aspects of these reports need to be clarified, however, the first being inaccurate assessment of dermal and respiratory intake of PAHs. Because PAHs levels on a skin pad might not accurately reflect the exposure levels of natural skin, and also because the absorption rate constant used in the study was obtained from only four individuals, the extent to which PAHs entered via the skin might be overestimated. Second, because of variations in labor practices and the composition of PAHs in different job categories, the extent of reductions in skin exposure is affected, and intake varied according to the type of industry and occupation. This is born out by the findings of this study: a significant effect was seen only in topside workers. Finally, the rapidity and simplicity of the assay for 1-OHPG provides practical advantages in assessing the intervention effectiveness. Since 1-OHPG is approximately three- to five-fold more fluorescent than 1-OHP, it provides a more sensitive biomarker for assessing exposure to pyrene. However, concentration of urinary 1-OHP after deconjugation is correlated with 1-OHPG concentration in human urine, indicating that either assay can be used to determine human exposure to pyrene. The results of comparison of three different assays for 1-OHPG indicated that a simple and rapid method (fluorescence quantitation after immuno-affinity purification) shows reliable and comparable results to the assays using HPLC. This research clearly demonstrates, however, the usefulness of urinary PAH metabolites as markers for the effectiveness of exposure reduction programs in
PAH exposed workers.

In conclusion, although the present study suffers from several limitations, including the absence of skin exposure monitoring, the use of a surrogate exposure indicator of specific PAH measurements, and small sample size, its findings indicate that the measurement of PAH metabolites in human urine appears to be ideally suited to biomonitoring in the workplace, and testing the intervention effectiveness of attempts to reduce PAH exposure.

References