Urinary hippuric acid and orthocresol excretion in man during experimental exposure to toluene

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ABSTRACT It is not known whether urinary excretion of hippuric acid (HA) or orthocresol (O-Cr) is to be preferred for the biological monitoring of workers with occupational exposure to toluene. To study this, 42 printing trade workers with more than 10 years' exposure to a mixture of organic solvents including toluene (0–20 ppm) and 43 control subjects matched by age, smoking habits, and living accomodation were investigated. Each matched pair was randomised to an experimental exposure of either 100 ppm or 0 ppm toluene for 6-5 hours under controlled conditions in an exposure chamber. Urinary excretion of HA and O-Cr was determined by high pressure liquid chromatography from samples obtained before exposure, during the first three hours, and during the last 3-5 hours of exposure. No difference in HA and O-Cr excretion was found between printing trade workers and controls. The median O-Cr excretion increased 29 times during exposure, whereas the HA excretion increased only five times. Thus only 3% of the O-Cr excretion originated from other sources than toluene whereas the corresponding value for HA was 19%. Standardisation of the concentrations of HA and O-Cr in relation to urinary creatinine reduced the relative variation by 29% and 56% respectively. This was not reduced further by expressing the excretions as average excretion rates based on total volume of urine collected. Background urinary O-Cr excretion was three to four times higher among smokers than non-smokers, probably due to the content of O-Cr in cigarettes. The O-Cr excretion in unexposed smokers was, however, 10 times lower that that of the non-smokers during the end of the experimental exposure to 100 ppm toluene.

The use of urinary metabolite excretion as an index of occupational exposure to organic solvents is suggested by the good correlation between these indices and biological changes. As a measure of exposure to toluene two urinary metabolites of toluene have received major interest in biological monitoring. Firstly, hippuric acid (HA) because most (80%) of the body burden of toluene is metabolised to this substance and, secondly, o-Cresol (O-Cr) because this is regarded as the only known specific metabolite of toluene. The lack of specificity of HA and the very small proportion of the retained toluene that is excreted as O-Cr (≤ 0-1%) raise the question which of these metabolites is to be preferred in the biological monitoring of people exposed to toluene.

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To investigate the excretion of toluene metabolites in man we have developed highly sensitive high performance liquid chromatographic (HPLC) methods to measure the urinary excretion of HA and O-Cr, and an experimental exposure to toluene was set up.

Material and methods

Eighty five people, 42 male printing trade workers and 43 controls aged 29-50, were included in the study. The printers were selected in collaboration with their trade union from six rotogravure and flexoprinting plants. Only those who had been employed for at least 10 years in printing plants and at least five years at the present plant were included. Workers with known heart, lung, liver, kidney, or any other disabling diseases, or with more than 15 days of absence from work during the past year were
excluded. Regular use of psychotropics, hypnotics, or analgesics was also used as an exclusion criterion.

The controls were selected from the municipal register of Aarhus, Denmark. A questionnaire was mailed to 300 people who matched the 42 printers, according to age, educational level (skilled or unskilled workers), and living accommodation. From the information gathered in this way the controls were selected as the closest match according to the same criteria as the printing trade workers, except that those with occupational exposure to organic solvents or other potentially harmful chemicals were excluded.

The printers had been exposed to a mixture of organic solvents including toluene. During the last years toluene had gradually been replaced by other solvents including ethylacetate and alcohols. Workroom air analyses had been performed on several occasions in all the plants during the past three years. The mean exposure level was 0-48 (range 0-1 to 1-1) of the sum of the concentrations divided by the respective maximal allowed time weighted average concentrations (TLV). The concentrations of toluene ranged from 0 to 20 ppm (parts per million). Everyone had been at work the day before the examination, giving an exposure free interval of 15–18 hours for printing trade workers.

The mean age for both printers and controls was 36-4 years. Body height and weight were 178 cm (range 162–192 cm) and 78 kg (range 58–105 kg) respectively. Sixteen of the printers were non-smokers, nine were moderate smokers (consumption below 11 cigarettes daily), and 17 were heavy smokers. The corresponding numbers in the control group were 16, seven, and 20.

EXPERIMENTAL EXPOSURE
Each matched pair was randomised to exposure to either 100 or 0 ppm toluene during 6-5 hours in a controlled exposure chamber. Before the experiment everyone gave informed consent and underwent a general medical examination before entering the chamber. The exposure was carried out under the continuous supervision of a physician.

EXPOSURE CHAMBER
The exposure took place in a climatic chamber with a floor area of 29 m² and a volume of 83 m³. The walls were of stainless steel, and all the equipment of the chamber was made either of steel or teflon to minimise adsorption of toluene or other vapours.

The chamber was equipped with a recirculation system with a capacity of 1400 m³h⁻¹. Conditioned particle and charcoal filtered outdoor air was supplied with a rate of 380 m³h⁻¹ corresponding to an air renewal of 4-6 changes h⁻¹. The temperature in the room was 22·8 ± 0·5°C (mean ± SD) and the relative humidity was 40·2 ± 1·2%.

Toluene vapour (Merck Uvasol art 8331 (99-7%)), 20% saturated, was led to the fresh air supply. The concentration of toluene in the chamber air was continuously monitored by a flame ionisation detector (Bendix model 8401), which was calibrated daily with a standard gas containing 106 ppm toluene in synthetic air (de Linde, Germany). The concentration of toluene was continually adjusted by changing the degree of saturation of the inlet vapour.

During 6-5 hours of experimental exposure to toluene the concentration was 102 ± 2 ppm (382 ± 7 mg/m³), and during the control experiment the concentration of hydrocarbons was below 0-1 ppm.

Three or four people were exposed at a time. The physical activity was moderate. They were usually seated or working lightly in a standing position.

URINE SAMPLING AND ANALYSIS
Sampling of the total volume of urine was carried out in three consecutive periods before and during exposure. The time of the last micturition before entering the chamber was registered, and the urine was collected before exposure (period -2.5-0 h), after three hours of exposure (period 0–3 h), and at the end of exposure (period 3–6.5 h). The exact time of micturition and the volume of urine were registered, and samples were taken for analysis.

The concentration of hippuric acid (HA) and o-Cresol (O-Cr) was determined by a HPLC technique described elsewhere.³ Detection limits were 0-005 mg × ml⁻¹ for HA and 0-005 μg × ml⁻¹ for O-Cr. Concentration of creatinine in urine was determined by the Jaffé method.⁴

STATISTICS
The statistical comparisons were carried out using analysis of variance techniques (repeated measurement design) after logarithmic transformation. The covariances between the urinary excretion of metabolites and the personal variables age, body height and weight, and consumption of alcohol and tobacco were calculated and the statistical model was adjusted for the influence of the factors correlating significantly to the excretion rates.

The three samples in the four groups were compared using a standard analysis of variance. To investigate the time course of the excretion the differences between the first and the second, and the first and the third samples were analysed using a two dimensional analysis of variance. The calculations were made by the computer system Genstat.⁵ Probability values of less than 5% were considered statistically significant.
Results

The urinary excretion of the two metabolites through the experiment is shown in the figure. The median and quartiles are depicted for each of the

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Geometric mean excretion rate of hippuric acid and β-cresol in non-smokers, moderate smokers (≤ 10 cigarettes per day), and heavy smokers. Difference between non-smokers and each of the groups of smokers is indicated by an asterisk (*p ≤ 0.05, **p ≤ 0.01)
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This may partly be explained by the “noise” from other sources of benzoic acid.

The lack of difference in urinary excretion of HA and O-Cr between printers and controls indicates an equal metabolism of toluene in the two groups. Thus the self stimulating effect of toluene on its own metabolism found in animals exposed to 300 ppm could not be found among printers in the present study. This was probably due to the much lower exposure to toluene in this group, and the fact that the concomitant exposure to other solvents does not alter the metabolism significantly.

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