Hippuric acid, phenol, and trichloroacetic acid levels in the urine of Japanese subjects with no known exposure to organic solvents

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Ikeda, M., and Ohtsui, H. (1969). Brit. J. indus. Med., 26, 162-164. Hippuric acid, phenol, and trichloroacetic acid levels in the urine of Japanese subjects with no known exposure to organic solvents. Urine samples from 36 male and 30 female university students and 31 male factory workers with no known exposure to industrial organic solvents were analysed for hippuric acid, phenol, and trichloroacetic acid, which are the major metabolites of toluene, benzene, and trichloroethylene respectively. The normal levels were less than 1.4 g./l. for hippuric acid, less than 80 mg./l. for phenol, and less than 1 mg./l. for trichloroacetic acid. No evidence was obtained to suggest that correction for urine concentration with either specific gravity or creatinine concentration minimizes individual variation of metabolite levels.

Recently a number of papers have been reporting the increased levels of metabolites in the urine from workers exposed to various organic solvents (Souček and Vlachová, 1960; Walkley, Pagnotto, and Elkins, 1961; Pagnotto, Elkins, Brugsch, and Walkley, 1961; Ogata, Sugiyama and Moriyasu, 1962; Elkins, Compron, and Pagnotto, 1963; Salmowa, Piotrowski, and Neuhorn, 1963; Ikeda and Kita, 1964; Van Haaften and Sie, 1965; Rainsford and Lloyd Davies, 1965; Docter and Zielhuis, 1967; Pagnotto and Lieberman, 1967; Ogata and Nagao, 1968; Ogata, Nagao, and Tomokuni, 1968; Tanaka and Ikeda, 1968). Because some of these metabolites can originate non-industrially, for example from food (Quick, 1931), it was important to determine the levels of metabolites in the urine of non-exposed Japanese subjects. Hippuric acid, phenol, and trichloroacetic acid were chosen because their parent substances (toluene, benzene, and trichloroethylene) are among the most important organic solvents.

Materials and methods

Urine samples
Sampling was conducted in April. Second morning urines were sampled from university students around 10 a.m. Workers in three factories were asked to urinate at 1 p.m. and the urines voided later, around 3 p.m., were sampled. Unless otherwise specified, the subjects had no known exposure to industrial organic solvents for at least one week prior to the sampling.

Chemical and statistical analyses

Hippuric acid Hippuric acid was determined essentially by Ogata et al.'s modification (1962) of the method of Gaffney, Schreier, DiFerrante, and Altman (1954). Fresh urine, 0.01 to 0.05 ml., was spotted on Toyo No. 525 paper (roughly equivalent to Whatman No. 3 mm. paper) along a 2.5 cm. line and air-dried. The paper was subjected to ascending chromatography with n-butanol:acetic acid: water (4:1:1, by vol.). When the solvent had risen 20 cm., the paper was air-dried, sprayed with 4% dimethyldimethylbenzaldehyde in acetic anhydride (saturated with sodium acetate), and heated in an oven at 130°C. for 3 minutes. An area of 6 × 6 cm., which included the orange spot, was taken, and the dye was extracted into 6 ml. of methanol. Extinction was measured at 460 m.μ.

Phenol Determination of phenol was as described elsewhere (Ikeda, 1964). Phenol in the acid-hydrolysed
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The results and discussion

The results are summarized in the Table. The mean specific gravity of the urine samples was closer to the values reported by Rainsford and Lloyd Davies (1965) and Van Haaften and Sie (1965) than to that reported by Elkins and Pagnotto (1965).

Practically no Fujiwara-reaction-positive substances were observed in the normal urine samples. Oxidation of the urine to convert trichloroethanol to trichloroacetic acid (Tanaka and Ikeda, 1968) did not give significant increases in Fujiwara-reaction-positive substances. Thus, the concentrations of trichloroacetic acid in the Table can be considered to represent the concentrations of total trichloro-compounds as well. These low concentrations were in sharp contrast to those found in the urine from factory workers exposed to trichloroethylene (120 to 250 p.p.m., 8 hours a day), where the levels of trichloroacetic acid and trichloroethanol are as high as 80 mg./l. and 130 mg./l. respectively (Tanaka and Ikeda, 1968).

The amounts of both hippuric acid and phenol in the normal urine were measurable. The levels of hippuric acid were slightly higher than those reported by Ogata et al. (1962). The difference might be due to the number of specimens examined. The phenol levels were roughly in agreement with or somewhat higher than those reported from various countries (Rainsford and Lloyd Davies, 1965; Walkley et al., 1961; Van Haaften and Sie, 1965).

TABLE

LEVELS OF HIPPURIC ACID, PHENOL, AND TRICHLOROACETIC ACID IN NORMAL URINE SAMPLES

<table>
<thead>
<tr>
<th></th>
<th>Students</th>
<th></th>
<th>Factory workers</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Male</td>
<td>Female</td>
<td>Male</td>
<td>Female</td>
</tr>
<tr>
<td>No. of samples examined</td>
<td>36</td>
<td>30</td>
<td>31</td>
<td>1-022</td>
</tr>
<tr>
<td>Mean specific gravity</td>
<td>1:019</td>
<td>0-89 (0:24-2:93, 28%)</td>
<td>1:56 (0:87-2:79, 11%)</td>
<td></td>
</tr>
<tr>
<td>Creatinine (g./l.)</td>
<td>1:32</td>
<td>(0:48-3:68, 20%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hippuric acid</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Uncorrected (mg./l.)</td>
<td>301</td>
<td>398 (110-1431, 11%)</td>
<td>350 (112-1084, 10%)</td>
<td></td>
</tr>
<tr>
<td>Corrected with</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Specific gravity (mg./l.)</td>
<td>290</td>
<td>570 (153-2125, 10%)</td>
<td>291 (93-909, 10%)</td>
<td></td>
</tr>
<tr>
<td>Creatinine (mg./g. creatinine)</td>
<td>229</td>
<td>449 (163-1242, 8%)</td>
<td>235 (78-711, 10%)</td>
<td></td>
</tr>
<tr>
<td>Phenol</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Uncorrected (mg./l.)</td>
<td>26:1</td>
<td>25:2 (8:4-75:5, 17%)</td>
<td>22:8 (8:9-58:3, 15%)</td>
<td></td>
</tr>
<tr>
<td>Corrected with</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Specific gravity (mg./l.)</td>
<td>23:3</td>
<td>34:8 (9:8-123:8, 18%)</td>
<td>18:2 (8:0-41:5, 14%)</td>
<td></td>
</tr>
<tr>
<td>Creatinine (mg./g. creatinine)</td>
<td>18:9</td>
<td>28:5 (11:3-71:7, 14%)</td>
<td>14:5 (7:1-29:9, 14%)</td>
<td></td>
</tr>
<tr>
<td>Trichloroacetic acid</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Uncorrected (mg./l.)</td>
<td>0-0-90</td>
<td>0-0-60</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Corrected with</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Specific gravity (mg./l.)</td>
<td>0-2-00</td>
<td>0-1-03</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Creatinine (mg./g. creatinine)</td>
<td>0-0-93</td>
<td>0-0-71</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

1Adjusted to a specific gravity of 1-016.

2Metabolite concentrations divided by creatinine concentrations.

Numbers in the table are geometric means. Fiducial limit (P = 0.05) ranges and coefficients of variation of log1. [metabolite] are given in parentheses. In the case of trichloroacetic acid, minimum and maximum values observed are shown. In urine samples from 23 males and 21 females no Fujiwara reaction-positive substance was observed.
In order to evaluate the significance of corrections for urine density, metabolite concentrations were corrected to a constant specific gravity of 1.016. Metabolite concentrations were also expressed relative to creatinine concentrations. Contrary to expectation that the corrected values would be less variable than the uncorrected values (i.e., the observed values), it was found that the corrected values were not always less variable than the uncorrected ones (see Table). It appears that correction for urine concentration either by correction to constant specific gravity (1.016) or to constant creatinine concentration does not lessen variation of normal urine samples.

The possible correlation between the specific gravity of urine and its creatinine concentration was also examined on 254 urine samples from male factory workers exposed to toluene or trichloroethylene or not exposed. The calculated regression line of the scatter diagram was:

\[ y = 0.0585x + 0.2068 \]

where \( y \) = creatinine concentration (g./l.), and \( x = \) (specific gravity - 1.000) \( \times \) 1,000. Although the regression line passed quite close to the origin, the correlation coefficient (r) was only 0.63, indicating a rather poor correlation between specific gravity and creatinine concentration. Thus, correction to constant specific gravity may not replace correction to constant creatinine concentration, nor vice versa.

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References


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