Conjugated serum bile acid concentrations in workers exposed to low doses of toluene and xylene

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Recent studies of non-halogenated cyclic carbon solvents have shown little evidence of hepatotoxicity, although these negative findings might be explained as a consequence of the incapacity of the existing screening procedures to detect liver changes at an early stage. In fact, the uncertainties regarding the evaluation of liver function have been underlined and the need for simple, non-invasive tests of liver function has been emphasised. Recently, a relatively new test of hepatic function, the measurement of serum bile acids, has been studied in workers occupationally exposed to a variety of chemicals (styrene, vinyl chloride, and solvent mixtures).

During exposure to toluene and xylene at concentrations below the current TLV-ACGIH, we have studied the behaviour of serum bile acids in relation to environmental concentrations of toluene and xylene and to indices of internal dose (toluene and xylene in blood and urine) to evaluate the existence of the relation between exposure and effect.

Materials and methods

A group of 25 workers (mean age 42:3 years, age range 27–50) at a chemical factory producing varnishes and exposed mostly to toluene and xylene (mean exposure duration 10 years, range 5–15) were selected from among 30 subjects using the following criteria:

(1) exposure to solvents for at least two years;
(2) daily ethanol consumption less than 50 g;
(3) no history of hepatic disease;
(4) no drug taken in the previous month.

The survey was performed on a Friday after four days at work. The environmental concentration of solvents in the breathing zone of each subject was measured by means of a personal passive sampler. Samples were analysed according to the methods previously described. Table 1 shows that there was exposure to toluene and xylene (o, m, p-isomers) and to isobutyl acetate and methyl isobutyl ketone at low levels.

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Table 1 Time weighted individual value (Cw) of solvent exposure

<table>
<thead>
<tr>
<th></th>
<th>Median value (mg/m³)</th>
<th>Geometric SD (μmol/mol/m³)</th>
<th>TLV-ACGIH (mg/m³)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Toluene</td>
<td>21.5</td>
<td>223</td>
<td>292</td>
</tr>
<tr>
<td>Xylene</td>
<td>27.8</td>
<td>261</td>
<td>490</td>
</tr>
<tr>
<td>Isobutyl acetate</td>
<td>12.4</td>
<td>106</td>
<td>710</td>
</tr>
<tr>
<td>Methyl isobutyl ketone</td>
<td>15.8</td>
<td>157</td>
<td>205</td>
</tr>
</tbody>
</table>

Urine was collected from the subjects at 0800 (immediately before the beginning of work) and at 1200 (at the end of the half-shift) to measure toluene and xylene concentrations. At 1200 a 10 ml blood sample was taken from the antecubital vein from subjects who had been fasting from 0600 to assess toluene, xylene, and serum bile acid concentrations. Blood samples were also taken at 1200 from 25 controls who had been fasting from 0600 (mean age 44; age range 23–53) chosen from among unskilled workers using the above criteria but with no exposure to solvents or hepatotoxic xenobiotics.

Serum was separated within 20 minutes and kept at −20°C before analysis. Fasting levels of serum conjugated cholic acid (S-CCA) and serum conjugated chenodeoxycholic acid (S-CCDCA) were determined by an immunoenzymatic method (CCA and CCDCA Immunosystem, Sibar, Italy) in which unconjugated cholic and chenodeoxycholic acids had cross reactivities toward the CCA and CCDCA antisera of 5% and 7% respectively. The ratio of S-CCA to S-CCDCA was calculated.

From the fasting values of the 25 controls the upper limits of the reference range were expressed as the 90th percentile (1.4 nm/ml for S-CCA and 2.4 nm/ml for S-CCDCA). Statistical evaluation of the results was performed by the Wilcoxon rank sum test for unpaired samples and by χ². Correlations between serum bile acids and solvents in air (C1), in blood (Cg) in urine
with increased concentrations of serum bile acids. The present observations show that exposure to toluene and xylene produces a significant increase in the S-CDA/S-CCDCA ratio.

These observations might be explained by a change in hepatocyte function and, in particular, in one of the various steps involved in bile acid transport. Since the hepatic uptake of bile acids is a sodium dependent carrier mediated process, coupled to the sodium potassium pump (Na/K-ATPase), xenobiotics themselves or their metabolites, or both, may be responsible for an impairment of this mechanism (or simply competition with it). Even though this hypothesis seems attractive, the lack of any relation between the individual doses (environmental, urinary, and blood toluene and xylene) and serum bile acid concentrations remains to be explained.

The measurement of serum bile acid concentrations seems to be helpful in detecting early changes of hepatic function, even though it needs further validation before being used as an index of effect.

### References