A simple method for the quantitative analysis of urinary trichloroethanol and trichloroacetic acid as an index of trichloroethylene exposure

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Ogata, M., Takatsuka, Y., and Tomokuni, K. (1970). Brit. J. Industr. Med., 27, 378-381. A simple method for the quantitative analysis of urinary trichloroethanol and trichloroacetic acid as an index of trichloroethylene exposure. A simple method of estimating trichloroethanol and trichloroacetic acid in the urine of workers exposed to trichloroethylene is described. The glucuronide of trichloroethanol was hydrolysed enzymatically to trichloroethanol by β-glucuronidase and the trichloroethanol released was allowed to react with pyridine and potassium hydroxide in that order, thereby avoiding decomposition of trichloroethanol with strong alkali. The colour which developed in 3-5 minutes at 100°C with pyridine was measured at 440 nm and 530 nm. This also allowed trichloroacetic acid to be determined. The results agreed well with those obtained by longer methods.

After a subject has been exposed to trichloroethylene, trichloroethanol and trichloroacetic acid are excreted in the urine (Butler, 1949) and this excretion has been correlated with the degree of exposure (Tanaka and Ikeda, 1968; Ogata and Tomokuni, 1969). As trichloroethanol and trichloroacetic acid excretion occur at different rates (Souček and Vlachová, 1960), and the excretion of the former is more closely related to exposure on the same day, whereas excretion of the latter is more closely related to exposure on a few previous days (Ogata and Tomokuni, 1969), both trichloroethanol and trichloroacetic acid must be estimated in the urine.

Previous methods required four hours’ oxidation of trichloroethanol by chromium trioxide, and involved a difficult neutralization step before Fujiwara’s colour reaction was carried out (Seto and Schultz, 1956; Weitbrecht, 1965; Tanaka and Ikeda, 1968).

We describe here a shorter method based on the work of Marshall and Owens (1954), who hydrolysed urinary trichloroethanol glucuronide enzymically before estimating trichloroethanol, and on the modification made by Friedman and Cooper (1958) of Fujiwara’s reaction for trichloroethanol. They did not test the determination of urinary trichloroethanol in the form of glucuronide. The method described here allows the simultaneous estimation of trichloroethanol and trichloroacetic acid.

**Materials and methods**

Urine specimens were obtained from volunteers exposed for seven hours to 170 ppm of trichloroethylene and from workers exposed for three hours to 50-150 ppm of trichloroethylene at a motor car component factory.

**Analytical procedure**

To 1 ml of urine were added 1 ml of 0.15 M phosphate buffer (pH 6.0) and 1 ml of phosphate buffer containing 500 units (Fishman, Springer, and Brunetti, 1948) of β-glucuronidase (Sigma Chem. Co.), and the mixture was incubated at 37.5°C for 40 or more minutes. A 1-ml portion was taken and 5-0 ml of pyridine was added,
followed by 2.0 ml of 10 m potassium hydroxide. The mixture was agitated with a mechanical shaker, heated in a boiling water bath for exactly 3.5 minutes and immediately transferred to an ice bath for 3 to 5 minutes. Then 3 ml of the pyridine layer was transferred to 0.5 ml of water, and the mixture was diluted three times with pyridine/water (1/1). The optical densities of the solution at 440 nm and 530 nm were read immediately in a Beckman DU spectrophotometer. When a number of samples were to be determined, the solutions were kept in an ice bath until they were assayed. Standard solutions were obtained by using aqueous solutions of trichloroethanol and trichloroacetic acid instead of urine in the above method.

All analyses were carried out by an autorecording spectrophotometer (Beckman DU).

As shown in Fig. 1, absorption at 440 nm was almost wholly due to trichloroethanol and that at 530 nm was almost entirely due to trichloroacetic acid.

The calculation of the results depends on the separation of the optical densities measured at 440 and 530 nm, $U_{440}$ and $U_{530}$, into their component parts, $U_{440E}$ etc., the optical densities (OD) due to each component at each wave length. Thus we define:

\[
\begin{align*}
U_{440} &= \text{OD at 440 nm from urine} \\
U_{530} &= \text{OD at 530 nm from urine} \\
U_{440E} \text{ and } U_{530E} &= \text{OD in urine at 440 and 530 nm} \\
U_{440A} \text{ and } U_{530A} &= \text{OD in urine at 440 and 530 nm due to trichloroacetic acid.}
\end{align*}
\]

By definition:

\[
\begin{align*}
U_{440} &= U_{440E} + U_{440A} \quad (1) \\
U_{530} &= U_{530E} + U_{530A} \quad (2).
\end{align*}
\]

Experimentally, from Fig. 1:

\[
\begin{align*}
U_{440E} &= 0.1U_{440} \quad (3) \\
U_{530E} &= 0.06U_{530A} \quad (4).
\end{align*}
\]

Hence, substituting (4) in (1) and (3) in (2), we obtain the simultaneous equations:

\[
\begin{align*}
U_{440} &= U_{440E} + 0.06U_{530A} \quad (5) \\
U_{530} &= 0.1U_{440E} + U_{530A} \quad (6).
\end{align*}
\]

Whence $U_{440E} = (16.67U_{440} - U_{530})/16.57 \ldots (7)$

$U_{530A}$ can be found by substituting the result from equation (7) in equation (6).

Then $C_E = U_{440E} \times S_E/U_{440SE}$, where $C_E$ and $S_E$ are the concentrations of trichloroethanol in urine and in the aqueous standard, and $U_{440E}$ is the OD given by the standard.

An exactly analogous equation:

\[ C_A = U_{530A} \times S_A/U_{530SA} \]

gives $C_A$ for trichloroacetic acid.

**Results**

The absorption spectra of the colours developed with pyridine from trichloroethanol and trichloroacetic acid in urine are shown in Figure 1. The maxima for trichloroethanol were at 440 nm and 370 nm and those for trichloroacetic acid were at 530 nm and 370 nm respectively.

![FIG. 1. Absorption spectra of the colours given by trichloroethanol and trichloroacetic acid.](http://oem.bmj.com/)

The percentage of the glucuronide of trichloroethanol hydrolysed by $\beta$-glucuronidase at 37.5°C is shown in Figure 2. The trichloroethanol released reached its maximum 40 minutes after incubation, apparently according to first order kinetics. Forty minutes was taken as sufficient time for the completion of enzymic hydrolysis. The same urines were analysed for trichloroethanol by the chromium trioxide oxidation method of Tanaka and Ikeda (1968). It was found that 97% of the trichloroethanol glucuronide was hydrolysed by the enzymic reaction, and that urinary trichloroacetic acid was stable in the procedure.

![FIG. 2. Rate of hydrolysis of trichloroethanol glucuronide by $\beta$-glucuronidase at 37.5°.](http://oem.bmj.com/)
A small amount of trichloroethanol was found in the urine before enzymatic hydrolysis, perhaps caused by some exposure to trichloroethylene. Monochloroacetic acid did not show a colour reaction under the conditions of the test.

Recoveries of trichloroethanol added to urine ranged from 90 to 102% and those of trichloroacetic acid from 93 to 96% (Table). The variations probably resulted from differences in the efficiencies of extraction and not from incomplete colour formation. In the case of trichloroethanol, hydrolysis of the glucuronide by β-glucuronidase was not negligible.

### Table

**Recovery of Added Trichloroethanol (TCE-OH) and Trichloroacetic Acid (TCA) from Normal Urine**

<table>
<thead>
<tr>
<th>TCE-OH added (μg)</th>
<th>Recovery %</th>
</tr>
</thead>
<tbody>
<tr>
<td>50</td>
<td>90</td>
</tr>
<tr>
<td>100</td>
<td>92</td>
</tr>
<tr>
<td>150</td>
<td>102</td>
</tr>
<tr>
<td>200</td>
<td>102</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>TCA added (μg)</th>
<th>Recovery %</th>
</tr>
</thead>
<tbody>
<tr>
<td>50</td>
<td>95</td>
</tr>
<tr>
<td>100</td>
<td>93</td>
</tr>
<tr>
<td>150</td>
<td>95</td>
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<tr>
<td>200</td>
<td>96</td>
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</tbody>
</table>

A comparison was made of the concentrations of trichloroethanol and trichloroacetic acid found by the oxidation method (Tanaka and Ikeda, 1968) and by our method. Urine specimens were taken during and after exposure to trichloroethylene. The relationship between the values obtained by the two methods can be described by regression lines.

For trichloroethanol:

\[ y_1 = 1.03x_1 + 0.53, \quad r = 0.986 \]

where \( x_1 \) represents the values obtained by the oxidation method and \( y_1 \) those obtained by our direct colorimetric method (Fig. 3). There was good agreement between the results, though our method gave slightly higher values than the oxidation method.

For trichloroacetic acid:

\[ y_2 = 1.08x_2 - 5.06, \quad r = 0.996 \]

where \( x_2 \) represents the values obtained by the oxidation method and \( y_2 \) those obtained by our method (Fig. 4).

That the absorption at 440 nm was mostly due to trichloroethanol was shown by measuring the absorption spectrum of some specimens of urine obtained from a volunteer exposed to 170 ppm of trichloroethylene for eight hours (Fig. 5). The large absorption peak at 440 nm appeared after treatment with β-glucuronidase. There was also slightly increased absorption at 530 nm, also due to the trichloroethanol released.
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Discussion

The quantitative determination of trichloroethanol and trichloroacetic acid in urine can be conducted very simply by the method described and the values obtained coincide with those obtained by the much longer oxidation method. Trichloroethanol was released from its glucuronide by β-glucuronidase. The quantitative determination of trichloroethanol by the Fujiwara reaction could not succeed because trichloroethanol is decomposed by strong alkali; but, by adding pyridine first, as described by Friedman and Cooper (1958), trichloroethanol was estimated successfully by a modified Fujiwara reaction. Friedman and Cooper (1958) attempted to measure chloral hydrate, trichloroacetic acid, and trichloroethanol simultaneously, this involving, therefore, a very complicated calculation. But, in urine excreted from workers exposed to trichloroethylene, only trichloroacetic acid and trichloroethanol were present, and their concentrations were easily calculated.

If the concentration of trichloroacetic acid is much lower than that of trichloroethanol, the former can be estimated by the pyridine-alkali method without prior treatment with β-glucuronidase.

References


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