Urinary mercury excretion and proteinuria in pathology laboratory staff

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ABSTRACT  The use of mercuric chloride as an histological fixative was associated with high environmental atmospheric concentrations of mercury vapour (up to 0.5 nmol/l) as well as mercury compounds (total Hg up to 1.0 nmol/l). Technicians exposed to this environment showed increased urinary mercury (median value 265 nmol/24h) and protein outputs (median value 117 mg protein/24h). Routine control measures, ventilation and careful handling of mercuric chloride solutions, reduced the level of atmospheric mercury vapour levels to within acceptable limits (threshold limit values 0.01 mg/m³ (0.05 nmol/l) alkyl compounds and 0.05 mg/m³ (0.25 nmol/l) for all forms except alkyl). This reduction was associated with the disappearance of trace proteinuria from the technicians’ urine. Contamination of histology laboratories by mercuric chloride should be minimised.

Corrosive sublimate (HgCl₂) was one of the first reagents to be used as a tissue preservative and fixative in histology (Blanchard, 1847). Braus (1896) appears to have been the first to use a mixture of formaldehyde and mercuric chloride. This mixture, generally known as formol-corrosive solution, was advocated as a routine fixative by Carleton (1926) and by Lendrum (1941). Its use is now promoted by the Association of Clinical Pathologists (1966). Other fixatives containing mercuric chloride, such as Zenker’s, Susa’s and Helly’s solutions, are also in common use.

Since 1947 histopathologists in Dundee have routinely used formol-corrosive solution (30g HgCl₂ in 100 ml commercial formalin diluted with water to 1 litre) for fixing surgical material. The routine procedure for post-mortem material, however, includes preliminary fixation in 10% formol-saline followed by prolonged immersion (secondary fixation) in 5% mercuric chloride (saturated corrosive).

This investigation was undertaken to assess whether or not there was mercury absorption in the laboratory staff.

Methods

POPULATION
Individuals were allocated to one or other of two groups depending on whether or not their laboratory used mercurial fixatives.

Group 1 (exposed)
The laboratory staff of three hospitals used formol-corrosive solution as a routine fixative. The staff consisted of 21 persons: 13 men and eight women, and they all handled, or worked for five to six days each week in a room where formol-corrosive solution was present. Their ages ranged from 17 to 50 years and their period of service in these laboratories varied from seven months to 20 years.

Group 2 (non-exposed)
The control group consisted of 21 histopathology laboratory workers from another hospital who were using only formalin for routine fixation. They did not use mercury or its compounds for any work, and were selected to match the first group for sex and by decade for age.

ENVIRONMENTAL ASSESSMENT
The room atmosphere of four laboratories within one hospital pathology department which traditionally uses mercuric chloride fixatives was investigated. These laboratories were:

Histology laboratory A
Mercuric chloride-containing solutions and corrosive-fixed tissue in specimen jars were constantly present; specimen cutting and fixation had always been undertaken in this room. No food or beverage
was taken in this crowded working space, which had no form of controlled ventilation system.

**Histology laboratory B**
This was used for microscopy; no mercury compounds were stored and corrosive-fixed tissues were rarely present.

(a) **Haematology laboratory H**
No mercury was used.

(b) **Haematology annex Hx**
This was adjacent to haematology laboratory H and had previously been used for manometric analysis and mercury cleaning. Liquid mercury was found in floor cracks and under parquet floor-blocks.

**PLAN OF INVESTIGATION**
During the initial phase, 24-hour urine collections were made from members of staff and measurements taken of mercury in air in histology laboratories (A and B) and in the haematology laboratories (H and Hx). Control measures were subsequently introduced in histology laboratory A. These consisted of:
1. Educating the staff about the potential hazard.
2. Using surgical gloves for handling fixed tissue specimens.
3. Improving exhaust ventilation with a minimum of 3·5 air changes an hour.

Twelve months after these control measures had been implemented in histology laboratory A, measurements of mercury in air were again taken in laboratories A, B, H, and Hx, and 24-hour collections of urine were obtained from nine members of staff then working in histology laboratory A.

**Mercury in laboratory air**
The total mercury present in the laboratory atmosphere was estimated by the method of Hanson et al. (1965). Atmospheric mercury vapour was measured by an ultraviolet absorption method using a Beckman model K 24 meter.

**Urine collections**
Each subject provided a 24-hour urine specimen in heavy-metal-free polythene bottles. Five ml of concentrated hydrochloric acid were added to each bottle as a preservative and samples were stored at a temperature of 4°C.

**Urinary estimation of mercury**
A dithizone colorimetric method was used, modified from that of Kudsk (1964).

**Urinary protein**
The method was developed from those of Henry et al. (1957) and Kibrick (1958). The proteins were precipitated from a 20 ml aliquot of urine by addition of 5·0 ml of 25% trichloroacetic acid solution before spectrophotometric estimation using biuret reagent. A standard curve was made using crystallised bovine albumin.

**Urinary creatinine**
The alkaline picrate method of Bonsnes and Taussky (1945) was used. Creatinine output was calculated to check that a full 24-hour specimen had been received. All samples were found to be satisfactory by this rough check.

**Statistics**
Since the distribution of urinary output rates were not gaussian the significance of differences between the exposed and non-exposed groups in respect of mercury and protein output rates were calculated using the U test (Mann and Whitney, 1947).

**Results**
Levels of environmental mercury are given in Table 1. Measurements of total atmospheric mercury in histology laboratory A were up to five times higher than those in haematology laboratory H. Raised mercury vapour values, including some of 0·5 nmol/l, were detected in air sampled near the two sinks in histology laboratory A. One of these sinks is customarily used for block-taking, involving the handling of mercuric chloride-fixed specimens and it is conveniently near the ready-use supplies of fixative solution. Disused fixed-specimen storage jars are cleaned out with hot water in the second sink.

Histology laboratory B, selected as another control for laboratory A, gave zero readings for mercury vapour. The finding of traces of mercury vapour and 0·20 nmol/l of total mercury in laboratory H was unexpected in a control environment. Higher readings in the adjoining annex Hx also intended as a control led to the discovery of the source, mercury globules in gaps between and under the parquet floor-blocks.

The median urinary mercury excretion among 21 non-exposed technicians was 72 nmol/24h, range 0 to 157 (Fig. 1). By contrast, the median mercury excretion rate was 265 nmol/24h, range 119 to 442, in 21 similarly employed technicians who were exposed to mercury-containing fixatives. Thus the mercury excretion rate in the mercury-exposed personnel is four times greater than that in the control subjects. The difference is significant ($p < 0·001$). Similarly the amount of proteinuria in the mercury-exposed individuals as a group is at least twice that in the controls, using as an index either the output rate of protein expressed per 24 hours
Table 1. Atmospheric mercury levels in the laboratories

<table>
<thead>
<tr>
<th>Laboratory</th>
<th>Mercury vapour* (nmol/l)</th>
<th>Total mercury (nmol/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Initial phase†</td>
<td>Follow-up phase</td>
</tr>
<tr>
<td><strong>Histology laboratory A</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Specimen area, sink and cork cutting-board</td>
<td>0-15-0-50 0-10-0-25</td>
<td>0-05-0-125</td>
</tr>
<tr>
<td>Wash-up area and sink</td>
<td>0-05-0-20</td>
<td>0-05-0-10</td>
</tr>
<tr>
<td>Solution preparation area</td>
<td>0-05-0-15</td>
<td>0-05-0-10</td>
</tr>
<tr>
<td><strong>Histology laboratory B</strong> (No mercury or mercury compounds present)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Specimen area</td>
<td>&lt;0-025</td>
<td>&lt;0-025</td>
</tr>
<tr>
<td>Wash-up area and sink</td>
<td>0-05-0-20</td>
<td>0-05-0-10</td>
</tr>
<tr>
<td>Solution preparation area</td>
<td>0-05-0-20</td>
<td>0-05-0-10</td>
</tr>
<tr>
<td><strong>Haematology laboratories</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>(a) Main laboratory H</td>
<td>&lt;0-025</td>
<td>&lt;0-025</td>
</tr>
<tr>
<td>(b) Annex Hx</td>
<td>&lt;0-025-0-5</td>
<td>&lt;0-025-5-0</td>
</tr>
</tbody>
</table>

*Range of repeated meter readings
†And on different week-days
Mercury vapour meter detection limit was 0-005 mgHg/m³ (0-025 nmol/l)

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Fig. 1 Urinary mercury excretion rates in mercury-exposed and non-exposed technicians.

Fig. 2 Urinary protein excretion rates in mercury-exposed and non-exposed technicians.

Table 2. Urinary mercury and protein in exposed and non-exposed technicians

<table>
<thead>
<tr>
<th>Phase</th>
<th>Population</th>
<th>Mercury output (nmol/24h)</th>
<th>Protein output (mg/24h)</th>
<th>Protein output (mg protein/nmol creatinine)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>Group 1 (exposed) 21 subjects</td>
<td>119-443 266</td>
<td>35-1029 117</td>
<td>2-7-576 115</td>
</tr>
<tr>
<td></td>
<td>Group 2 (non-exposed) 21 subjects</td>
<td>0-157 72</td>
<td>12-119 48</td>
<td>1-9-17-3 3-8</td>
</tr>
<tr>
<td>Follow-up*</td>
<td>Group 1 (exposed) 9 subjects</td>
<td>0-70 40</td>
<td>16-59 35</td>
<td>1-8-7-5 4-0</td>
</tr>
</tbody>
</table>

*12 months after control measures were implemented
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(Fig. 2), or, an indirect measure in which urinary protein is expressed per gram of accompanying creatinine output. The difference in the amount of proteinuria shown by the exposed and non-exposed groups is also significant (p < 0.01). These results are summarised in Table 2.

In the follow-up phase, mercury vapour levels were not changed materially in either of the histology laboratories but total mercury levels in histology laboratory A were lower (Table 1). Urinary mercury and protein outputs of technicians working with formaldehyde, fixatives, but using control measures (Table 2), were comparable with those of non-exposed technicians (Group 2).

An attempt was made to remove metallic mercury found on the floor of the haematology annex (Hx), and then the parquet-blocks were covered with linoleum. Spillages were first cleared up by mechanical means, that is by vacuum probe, and the affected areas were then treated with a wash composed of equal parts of slaked lime and flowers of sulphur mixed with water to form a paste. Twenty-four hours later the wash was removed with clean water and the surfaces allowed to dry.

Discussion

ENVIRONMENTAL MERCURY

The demonstration of significant amounts of mercury in the atmosphere of one representative laboratory (A), in which technicians used mercuric fixatives, raises the question of a potential mercury hazard. The presence of mercury in the haematology annex was unexpected and was found to be a legacy from an earlier use of this room for mercury cleaning and manometric analysis. Other control work-spaces gave negligible values for mercury.

There were several sources of contamination including bench spillage of corrosive sublimate solution, floor spillage (in this case on to wooden parquet-blocks), saturated filter papers lying exposed in the laboratory atmosphere, as well as the possibility of contamination of clothing worn by laboratory staff. Local transient high concentrations of mercury vapour were obtained by warming the copper handles of test-tube brushes found in sink areas. When hot water was allowed to run through the sinks to waste, consistent mercury vapour values above 0.5 nmol/l were obtained.

The threshold limit value for non-alkyl mercury absorbed through the skin has been reduced to 0.25 nmol/l from 0.5 nmol/l and for alkyl compounds to 0.05 nmol/l (American Conference of Governmental Industrial Hygienists, 1975; Department of Employment, 1973).

It is noteworthy that mercuric chloride has a vapour pressure of the same order as that of metallic mercury. The values are $3.1 \times 10^{-4}$ torr for HgCl$_2$ and $18 \times 10^{-4}$ torr for Hg at 25°C and under one atmosphere (extrapolated from data of Johnson, 1911). The maximum concentration of mercuric chloride in air would occur only in a closed space and lesser values would be found in open laboratories depending on air movement, temperature, and proximity to the mercury source. Formaldehyde which is commonly associated with mercuric chloride in fixatives can partly reduce mercuric to mercurous chloride. Reduction to free mercury has not been found. Solid mercuric chloride sublimes at room temperature and a likely source of mercury atmosphere contamination in the histology laboratory is evaporation of spilled solutions.

Several common metals interact with mercuric chloride releasing mercury or forming amalgams. These metals include copper, zinc, lead, and aluminium which are likely to be present as part of the plumbing, sinks, and the related metal fittings of histology laboratories. This effect may in part account for the higher concentrations of mercury found in air near the sinks in this investigation.

URINARY MERCURY EXCRETION

The technicians forming the non-exposed group showed a range of mercury excretion rates in urine which is similar to the normal range for the method used: 0 to 141 nmol/24h, median value 52 in 33 normal subjects (Taylor et al., 1969). The detection limit for this method is 20 nmol/l. Earlier authors quote comparable normal mercury values: 0 to 150 nmol/l (Nobel and Nobel, 1958; Kudsk, 1964) and less than 100 to 125 nmol/l (Goldwater, 1964).

Technicians who were exposed to mercury compounds had as a group appreciably higher urinary mercury output than the paired control subjects or other normal subjects. In 19 of 21 exposed subjects the output of mercury exceeded 150 nmol/l with fairly wide variability. Large day-to-day variability in and between exposed individuals in the output of mercury has been noted before (Buckell et al., 1946; Goldwater, 1964).

URINARY PROTEIN EXCRETION

As a group the exposed workers had an increased rate of excretion of protein in the urine compared with the group of matched controls who were all within the expected upper limit of adult normality for the method (Taylor et al., 1969). Although method dependent (Jorgensen, 1967), the upper limit of normal urinary protein excretion is between 100 and 150 mg/24h, mean value between 40 and 80 mg/24h (Relman and Levinsky, 1971). Paraproteinuria (globulins rather than albumin) in $1 \text{torr} \approx 1 \text{mmHg} \approx 133 \text{N/m}^2$
particular is not readily detected by Albustix-reagent strips (Rennie and Keen, 1967) but can be measured colorimetrically.

In nine of the 21 technicians in Group 1 the daily protein output rate exceeded 180 mg. This finding, accompanying a raised level of mercury excretion suggests that the repeated handling and/or inhalation of mercury and its bichloride salt, may cause an increase in the rate of protein excretion. There was a large variation in the protein excretion rate within the exposed group and a poor correlation between mercury and protein output rates for individual subjects, which suggest that a simple dose and effect relationship does not apply. This variability may well reflect differences in the extent of exposure from day to day, as well as underlying differences between technicians in their susceptibility to the effects of mercury once absorbed. No attempt was made to measure the duration of exposure time in technicians.

Overt clinical evidence of mercury poisoning was not found in the pathology technicians during the course of this study. The clinical significance of the minimal proteinuria found in the histopathology laboratory staff is not known. Comparative quantitative studies of trace proteinuria present in other populations exposed to industrial processes with environmental mercury concentrations at or above the threshold limit values would be valuable. In industrial situations proteinuria and urinary mercury excretion values (West and Lim, 1968) greater by a factor of 10 than those found in this study are currently accepted and are assumed to carry minimal health risk. The possibility remains however that trace proteinuria is a manifestation of mercury toxicity, albeit minimal. The occurrence of heavy proteinuria (more than 2g/24h) would warrant the removal of the subject from exposure to mercury and its compounds. In the case of the trace proteinuria associated with minimally increased urinary excretion of mercury as found here accompanying routine histopathological work, the long-term clinical significance is unknown.

**Mercury Absorption Control Measures**

Mercury vapour from manometric apparatus is a recognised risk in hospital laboratories (Williams et al., 1968). The current data indicate that the mercuric chloride used in histology also represents a potential hazard. To reduce contamination of skin and absorption by inhalation, laboratory staff should avoid spillage, work benches and floors should have impermeable surfaces without open joints, efficient ventilation should be provided, and the wearing of impermeable gloves is advisable. Simple control measures applied in histology labora-

tory A were effective in reducing the total atmospheric mercury to acceptable levels. Concurrently with this environmental change the urinary output rates of mercury and of protein also returned to the normal range.

The fortuitous observation of high levels of total mercury and mercury vapour concentration in the haematology annex led to the discovery of long-standing contamination of that room by metallic mercury. This finding, together with the accompaniments of the use of formol-corrosive tissue fixative already described, re-emphasizes that mercury and its compounds are an occupational hazard of laboratory technicians (Rose et al., 1972; Lancet, 1975).

The use of mercury-containing fixatives calls for special precautions. Not the least of these is the safe, pollution-avoiding disposal of the waste fixative as recommended by Porter (1972).

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**References**


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Mann, H. B., and Whitney, D. R. (1947). On a test of whether one or two random variables is stochastically larger than the other. Annals of Mathematical Statistics, 18, 50-60.


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