Urinary cytology in experimental toxic renal injury

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Aspirin causes the desquamation of certain epithelial cells and this effect can be demonstrated both in the stomach and, in particular, in the kidney (Harvald, Valdorf-Hansen, and Nielsen, 1960; Harvald and Clausen, 1960; Scott, Denman, and Dorling, 1963; Prescott, 1965, 1966). In the case of the kidney the nephron is lined by a variety of different epithelial cells and it is not clear which of these types of cell is excreted in response to aspirin. In view of the development of analgesic nephritis in a proportion of people who habitually take combinations of analgesics which include aspirin, it is of some interest to localize the site of this desquamative action of aspirin on the kidney. This paper is an account of a morphological study of the cells excreted after giving a series of nephrotoxins of known action to rats. The cells excreted were compared both with their site of origin and with the cells excreted in response to aspirin.

Material and methods

Young adult male rats of the Sprague-Dawley strain were used. During the experiments these were kept in individual stainless steel metabolism cages. They were fed on Diet 41B (Oxoid Ltd.) and fresh water *ad libitum*. After the collection of baseline specimens of urine, groups of six rats were given the different compounds by the doses and routes shown in the Table. The controls were injected with the solvent alone, *i.e.* physiological saline, by the appropriate routes. Specimens of urine were collected approximately 24 and 48 hours after giving the injections. 48 hours after the injections had been given the animals were killed. Previous work has shown that after the administration of nephrotoxins, the increase of cell excretion usually occurs within the first 2 days (Davies and Kennedy, 1967a,b; Davies, Kennedy, and Saluja, 1968; Davies, Kennedy, and Roberts, 1969).

The urine samples were collected over a period of time sufficient to give a volume of 1 to 2 ml. of fresh urine. The samples were mixed and films were made using a Shandon Cytocentrifuge running at a speed of 1,500 r.p.m. for 5 minutes. The films were then immediately fixed in absolute alcohol, rehydrated, and stained with haematoxylin and eosin. Specimens containing large numbers of cells were diluted as necessary. When the cells were required for staining by histochemical methods, these were carried out immediately after preparation of films by centrifugation.

At necropsy a thin slice of each kidney was fixed in formol-corrosive and dehydrated for the preparation of conventional paraffin sections. In addition, frozen sections of unfixed pieces of kidney were cut for staining by the same histochemical methods applied to the cells. Staining for alkaline phosphatase was carried out using the calcium-cobalt technique and succinic dehydrogenase using nitro BT according to the methods of Pearse (1960). The viability of the cells was tested by assessing their permeability to dilute eosin. Measurements of cells were made with an eyepiece micrometer.

### Table: Doses and routes used for the administration of the different nephrotoxins

<table>
<thead>
<tr>
<th>Compound</th>
<th>Solution used (mg./ml.)</th>
<th>Route</th>
<th>Dose per 100 g. body weight</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mercuric chloride</td>
<td>1</td>
<td>Intramuscular</td>
<td>2 mg.</td>
</tr>
<tr>
<td>Uranyl nitrate</td>
<td>30</td>
<td>Subcutaneous</td>
<td>30 mg.</td>
</tr>
<tr>
<td>dl-serine</td>
<td>40</td>
<td>Intraperitoneal</td>
<td>80 mg.</td>
</tr>
<tr>
<td>Potassium dichromate</td>
<td>10</td>
<td>Subcutaneous</td>
<td>4 mg.</td>
</tr>
<tr>
<td>Hexamethyamine bromide</td>
<td>4</td>
<td>Intramuscular</td>
<td>4 mg.</td>
</tr>
<tr>
<td>Ethyleneimine*</td>
<td>1 per cent. v/v</td>
<td>Intramuscular</td>
<td>0.002 ml.</td>
</tr>
<tr>
<td>Calcium aspirin</td>
<td>10</td>
<td>Gastric tube</td>
<td>20 mg.</td>
</tr>
</tbody>
</table>

**Abbreviation:** © Abbott Laboratories, Aldrich Chemical Co. Inc., Milwaukee, Wisconsin.

Results

None of the controls showed any change in the very scanty number of cells they excreted; the rats given the toxins behaved as follows:

**Mercuric chloride**

The cells excreted in response to mercuric chloride were all large with mean diameters in the order of

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15 x 10μ. In shape they were oval, pear-shaped, or polygonal and had rather indistinct borders (Fig. 1).

The cytoplasm was granular and brighty eosinophilic and some cells had faintly staining or pyknotic nuclei about 4μ in diameter. These cells were very similar to the necrotic cells seen in sections of the renal cortex where there was extensive necrosis of the straight segments of the proximal convoluted tubules (Fig. 2). In the urine most of the cells had lost their nuclei and they were permeable to dilute eosin indicating that they were not viable.

**URANYL NITRATE**

This also causes necrosis of the proximal convoluted tubules (Smith, 1951) and the histological appearances of the kidneys resembled those of the rats given mercuric chloride. In addition, there was some evidence of action on the collecting tubules, the cells of which had basophilic cytoplasm with large nuclei and a number of mitotic figures.

In the urinary sediment (Fig. 3), the main type of cell excreted was large, pear-shaped, and granular. These cells measured 20 x 12μ in diameter and in all respects resembled the cells excreted in response to mercuric chloride. In addition, some smaller round or oval cells 12-10μ in diameter were found; these had a round nucleus and smooth non-granular
cytoplasm. These cells were often present in large clumps and were most frequent on the first day of the experiment. They have some resemblance to the cells excreted by animals given ethyleneimine (vide infra).

**DL-SERINE**

The histological changes induced by dl-serine were almost identical with those produced by mercuric chloride. Areas of necrosis occurred in the proximal convoluted tubules in the inner part of the cortex. Large numbers of cells were found in the urinary deposit at both 24 and 48 hours. These cells also were large and had diameters between 15 and 20μ. The cells were oval or rounded with granular eosinophilic cytoplasm and they had a general appearance similar to that of the cells excreted after mercuric chloride had been given. Few cells had nuclei but, like the sediment seen in uranyl nitrate poisoning, small less granular nucleated cells were also present.

**POTASSIUM DICROMATE**

The most notable feature in these animals was the disparity between the small number of intact cells seen in the deposit and the severity of the renal damage. The main component of the deposit was amorphous eosinophilic debris in which only a few cells were recognizable (Fig. 4). Some of these cells measured about 12 x 10μ and were, therefore, slightly smaller than those excreted after giving mercuric chloride. Their margins were ill defined and some of them contained well-preserved nuclei between 5 and 7μ in diameter. A few larger cells 17 x 15μ were also seen. Very few recognizable cells were present 24 hours after giving the injections but moderate numbers were present at 48 hours. The presence of so much debris in the deposit was explained by the appearance of the kidney (Fig. 5).

**FIG. 4** Few recognizable cells can be seen in the urinary sediment after potassium dichromate had been given: the sediment consists almost entirely of amorphous debris (c.f. Fig. 5). Haematoxylin and eosin. × 440.

**FIG. 5** Renal cortex of a rat which had been injected with potassium dichromate 48 hours previously. There is extensive necrosis of the tubules but the necrotic cells have disintegrated in situ so that the affected tubules contain amorphous debris only. Haematoxylin and eosin. × 440.

In the outer part of the cortex there was necrosis affecting proximal convoluted tubules. In these tubules the cells had disintegrated in situ leaving an amorphous granular deposit similar to that seen in the cytological preparations. This was in complete contrast to the action of mercuric chloride where the necrotic cells remained distinct (Fig. 2).

**HEXADIMETHRINE BROMIDE**

After this compound had been given the urine con-
After the administration of hexadimethine bromide, moderate numbers of medium-sized cells with dark round nuclei were found in the urine. Haematoxylin and eosin. x 440.

Renal cortex of a rat injected with hexadimethrine bromide. A necrotic distal tubule can be seen in the centre of the field; cells resembling those in Fig. 6 are present in this tubule. Haematoxylin and eosin. x 440.

tained moderate numbers of brightly eosinophilic medium-sized cells, some of which had prominent dark nuclei (Fig. 6). The necrosis induced by hexadimethrine bromide occurs in the ascending limbs of Henle's loops and in the distal convoluted tubules in the cortex. The similarity between the urinary cells and the damaged cells in the renal cortex can be seen by comparing Figs 6 and 7. These cells had an average size of 10 × 7μ and were usually oval or pear-shaped with slightly granular cytoplasm.

This compound causes a highly selective necrosis of the renal papilla (Davies, 1969). The deposit contained well-preserved cells which were characteristically found in clumps or sheets (Fig. 8). The cells had abundant palely-staining cytoplasm and, where they occurred in a sheet, their outlines were angular. Single cells were more rounded and often had a single large vacuole in the cytoplasm; this appearance

Typical sheet of well-preserved cells in urine of a rat given ethyleneimine. Haematoxylin and eosin. x 440.
is most marked in wet preparations stained by the method of Prescott and Brodie (1964). The nuclei were almost invariably present and varied in appearance from a faintly granular chromatin pattern to a rather coarse lumpy arrangement. Cells similar to these were found in the necrotic papilla (Fig. 9, overleaf), but some of the sheets of cells may have been derived from the pelvic surface of the papilla or from the epithelium of the renal pelvis itself.

**CALCIUM ASPIRIN**

After aspirin had been given, no necrosis was found in the kidneys but a certain amount of eosinophilic debris was found in the lumina of the convoluted tubules. The most prominent change in the urine was the presence of large pale cells with abundant cytoplasm. These cells had mean diameters of 15 × 20μ and varied in form, being oval, polygonal, or pear-shaped; nuclei were usually absent. The cytoplasm was faintly granular and in most respects the cells resembled those which were excreted after mercuric chloride, uranyl nitrate, or dl-serine had been given (Fig. 10). In addition, a few smaller cells were present. These resembled the smaller variety of cell seen after uranyl nitrate had been given.

**Discussion**

These results show that the administration of a number of well-known nephrotoxins is associated with changes in the urinary deposit and that the morphology of the cells excreted depends on the agent given and its site of action. In general, the nephrotoxins which are known to act on the proximal tubule, i.e. mercuric chloride, uranyl nitrate, and dl-
serine, caused the excretion of large eosinophilic granular cells 15μ in diameter or greater. Allowing for the mechanical difference in preparing sections and cytological specimens, these cells clearly resembled those seen in the necrotic tubules in the kidney. While it is tempting to conclude that the excretion of cells of this type is a characteristic finding in necrosis of the proximal tubules, it must be pointed out that potassium dichromate, which also acts on the proximal tubule although on a different segment, produced a rather different cytological picture. We have not yet carried out quantitative studies with potassium dichromate, but it appears that relatively few recognizable cells are excreted after the administration of this agent, and the most prominent feature of both the urinary deposit and the histological sections was the presence of eosinophilic debris. This supports the conclusion, drawn previously, that the number of cells excreted after the kidney has been damaged by a toxin depends not only on the degree of damage but also on the site and mode of action of the damaging agent (Davies and others, 1969).

In the case of the two compounds which act more distally, hexadimethrine bromide and ethyleneimine, the cells were smaller and more frequently retained their nuclei. The origins of these cells were also demonstrable histologically but, when ethyleneimine was given, the whole papilla became necrotic so that the cells might have arisen from a variety of sites, including the pelvic epithelium, the ducts of Bellini, and Henle’s loops. However, both these toxins caused the excretion of cells which were different from those excreted after giving substances which act only on the proximal convoluted tubules.

Aspirin caused the excretion of large cells which were comparable in size with those excreted after the injection of mercuric chloride, uranyl nitrate, or dl-serine. Their cytoplasm was less granular than that seen after giving the three proximal poisons, but some cells did show alkaline phosphatase activity so that it seems probable that many of the cells excreted in response to aspirin are of proximal origin under the conditions used in this experiment; in the rat very large increases of excretion of proximal cells can occur without irreversible damage to the kidney (Davies and Kennedy, 1967b). While no necrosis was demonstrated in the sections in this experiment, it is relevant that necrosis has been found histologically after large doses of sodium salicylate were given to rats (Robinson, Nichols, and Taitz, 1967) and that the affected part of the nephron was the proximal convoluted tubule. Scott and others (1963) concluded that the ‘aspirin cells’ originated along a large part of the nephron, including the proximal convoluted tubule, and the unstained cells shown in their Fig. 6 closely resemble the cells we have found after giving mercuric chloride to rats. In our experiments there were also some smaller cells excreted after giving aspirin and these may have had a more distal origin. This is in agreement with the conclusion of Scott and his colleagues that aspirin may have a general irritative effect on a long length of the nephron.

If the main site of action of aspirin is the proximal convoluted tubule, the increase of cell excretion which it produces may not necessarily be relevant to the problem of analgesic nephritis. The primary lesion in analgesic nephritis is probably situated in the medulla, the cortical changes being secondary to the medullary necrosis which eventually occurs (Sanerkin, 1966; Kincaid-Smith, 1967).

**Summary**

The cytology of the urinary sediment of rats given different nephrotoxins has been correlated with the histological changes observed in the kidney. Mercuric chloride, uranyl nitrate, and dl-serine all led to the excretion of large eosinophilic granular cells caused by necrosis of the proximal convoluted tubules. The responses to potassium dichromate, hexadimethryrine bromide, and ethyleneimine are described. Aspirin in the dose used did not cause histologically detectable necrosis, but the cells were of at least two types. The most abundant cell was of a type similar to that excreted after the administration of toxins which act on the proximal convoluted tubule. It is concluded that much of the desquamative action of aspirin affects the proximal convoluted tubules, but that other part of the nephron may also be involved.

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