Amikacin Induces Renal Morphohistological Alterations in Wistar Rats

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The aim of this experimental study was to investigate the morphohistological effects of amikacin in rats. Male Wistar rats aged 8-12 weeks and weighing 200-250g were placed in a temperature (23±2°C) controlled room in which a 12/12h light: dark cycle was maintained. Twenty-five rats were randomly assigned to five experimental groups of five animals each that received daily treatment with 15 mg/kg of amikacin intraperitoneally. The control group was treated with 0.9% saline for 28 days. Groups I, II, III and IV were injected intraperitoneally with amikacin for 7, 14, 21 and 28 days, respectively. Renal tissues slides were stained with PAS reagent and examined light microscopy. The morphology of the PAS stained structures of the renal glomeruli was evaluated in the different experimental groups using scores. The renal morphohistology and the morphologic analysis of PAS staining were analysed statistically with the Kruskal-Wallis Test and Dunn’s Multiple Comparisons Test (5% statistical significance). The results show microscopically that kidneys in the groups treated with amikacin for 3 and 4 weeks showed marked cellular necrosis (p<0.0001), tubular obstruction, and basal membrane interruption. This result clearly shows that amikacin administration induces renal morphohistological alterations.

Key-words: amikacin; aminoglycosides; nephrotoxicity; tubular injury.

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INTRODUCTION

Due to its prominent role in major excretory pathways, the kidney is particularly sensitive to toxicity from antimicrobials drugs, especially the aminoglycosides. Storage of these drugs in the renal cortex with their effect on renal cells has consequences for renal function, and when re-absorbed by renal tubules induces nephrotoxicity (1,2).

Aminoglycosides are natural or semi-synthetic antibiotics with a heterocyclic structure formed by two or more aminosugars linked by glycoside bonds to an aminocyclitol ring. Aminoglycoside antibiotics are clinically used for treatment of Gram-positive and Gram-negative infections by inhibiting bacterial protein synthesis. However, they show clinical side effects with long-term administration, such as nephrotoxicity and ototoxicity (3). It is considered that aminoglycoside nephrotoxicity is caused by accumulation of aminoglycosides in proximal tubular cells, therefore cellular membrane structure and some transport functions would be affected (4).

Amikacin has become an antibiotic used especially for the early treatment of bacterial Gram-negative infections in hospitals, since bacterial resistance to gentamicin and tobramycin has become a significant problem. Adverse effects are ototoxicity and nephrotoxicity (5). Thus, the purpose of the present article is to investigate morphohistological kidney alterations due to chronic treatment with amikacin, by means of PAS staining.

MATERIALS AND METHODS

Male Wistar rats 8-12 weeks old, weighing 200-250g, were placed in a temperature (23±2°C) controlled room in which a 12/12h light: dark cycle was maintained. Twenty-five rats were randomly assigned to five experimental groups of five animals each that received daily amikacin treatment, 15mg/kg, intraperitoneally. The control group was treated with 0.9% saline for 28 days. Groups I, II, III and IV were injected intraperitoneally with amikacin for 7, 14, 21 and 28 days, respectively. After treatment, the animals in all groups were sacrificed and the kidneys were quickly removed, decapsulated and divided longitudinally into two equally sized pieces. One piece was placed in formaldehyde solution to be embedded in paraffin and 5 µm-thick sections were cut for histological examination by light microscopy.
Histological evaluation

**Histology**

Renal tissue slides were stained with PAS reagent and examined under light microscopy. The PAS reaction stains carbohydrates and carbohydrate-rich macromolecules. It is used to visualize the basal membrane that underlies epithelia. Three coded slides from each kidney were examined in a blinded manner.

**Qualitative and quantitative evaluation of morphology**

The morphology of the PAS-stained structures of the renal glomeruli was evaluated in different experimental groups using scores (Table 1) to classify the histological alterations with respect to the state of the basal membrane of the glomerular capillaries, and the continuity of the basal membrane of the renal glomeruli and parietal cells. The proximal and distal tubules were evaluated for the morphology of the revestment cells (necrosis). Kidney sample sections were analyzed with a microscope (with camera) connected to a microcomputer.

**Statistical analysis**

The renal morphohistology and the morphologic analysis for PAS staining were analysed statistically with the Kruskal-Wallis Test and Dunn’s Multiple Comparisons Test (5% statistical significance).

**RESULTS**

Figure 1 shows the results of renal morphohistological alterations found after histochemical PAS staining to the different groups treated with amikacin. Scores to histochemical alterations were classified according to table 1.

**Table 1 - Scores of renal histological alterations in Wistar rats.**

<table>
<thead>
<tr>
<th>Scores</th>
<th>Histological characteristics</th>
<th>1 (unaltered)</th>
<th>2 (slight)</th>
<th>3 (moderate)</th>
<th>4 (severe)</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>Alterations of parietal cells; Bowman’s capsule</td>
<td>Unaltered characteristics</td>
<td>Cells parietal alterations morphological</td>
<td>Alterations of parietal cells</td>
<td>Marked alterations of parietal cells</td>
</tr>
<tr>
<td>B</td>
<td>Basal membrane</td>
<td>Unaltered characteristics</td>
<td>Discreet interruption basal membrane</td>
<td>Interruption basal membrane 1/2</td>
<td>Intense interruption basal membrane</td>
</tr>
<tr>
<td>C</td>
<td>Tubule proximal cells</td>
<td>Unaltered characteristics</td>
<td>Integral brush-border membrane; level desquamation cells</td>
<td>Integral brush-border membrane; tubular necrosis</td>
<td>Complete interruption basal membrane; necrosis</td>
</tr>
</tbody>
</table>

Histological examination of the kidneys from animals in the control group revealed, as expected, entirely normal histological features, as illustrated in figure 1. However, there was tubular necrosis in kidneys from animals in the amikacin treated group, initially in group 2 and particularly in group 4, as illustrated in figures 2 and 3, respectively.
The lumen of the tubules was filled with degenerate and desquamated epithelial desquamated epithelial cells. Furthermore interruptions in the basal membrane are present.

DISCUSSION

Most of the aminoglycoside injected into the body is excreted into the urine without being metabolized. However, the remainder of the injected dose accumulates selectively and abundantly in the renal cortex. Aminoglycoside taken up by renal proximal tubular cells remains there for an extended period, leading to renal damage, such as structural changes and functional impairment of the plasma membrane, mitochondria and lysosome (6). Consequently, there has been great interest in the mechanism by which aminoglycosides are absorbed into the renal proximal tubular cells across the brush-border membrane. So far, results have indicated that aminoglycosides are taken up by receptor-mediated endocytosis following the binding of aminoglycosides to the brush-border membrane. There is also evidence that aminoglycosides directly disturb glomerular physiology, reducing SNGFR by lowering both the Kf and the afferent glomerular arteriolar output (7).

Perazella, (8) showed experimental data suggesting that both vascular, glomerular and tubular targets are involved in drug-induced nephrotoxicity, as a result of mechanisms that disrupt normal cellular structures and functions (mitochondria, membrane integrity, etc.), induce renal injury through intratubular obstruction (crystal deposition), and promote cellular swelling and tubular luminal occlusion (through osmotic effects). These effects were observed in our work, as seen in group 2 treated with amikacin, where we observed discrete basal membrane interruptions.

Studies of nephrotoxicity involving animal models have found that once-daily administration of gentamicin or tobramycin was significantly less toxic than more frequent (i.e., twice or three times daily) dosages or continuous infusion (9-12). Animal studies have further elucidated the probable mechanism involved in aminoglycoside nephrotoxicity, which involves an absorptive influx at proximal convoluted tubule cells and is mediated by a low-affinity, high-capacity mechanism that can be saturable, linear, or a mixture of the two (13-14). The exact mechanism is not known, but speculation has led primarily to the following hypotheses: a lysosomal mechanism, with aminoglycosides causing an accumulation of myeloid bodies within lysosomes and thereby inhibiting lysosomal phospholipases with a subsequent decrease in sphingomyelinase activity, or an extralysosomal mechanism, with inhibition of Na, K-ATPase and phospholipase C, leading to unopposed angiotensin II activity, or a combination of these mechanisms. In these experiments amikacin was administered once-daily, therefore significantly less toxic effects were observed than when traditional dosing regimens are applied.
Aminoglycoside-induced nephrotoxicity is characterized by tubular necrosis, basal membrane disruption, mesangial cell contraction, proliferation and apoptosis, indicated by a decrease in glomerular filtration and alteration in intraglomerular dynamics (15).

Our results clearly show that adverse effects observed in rats induced by chronic treatment with amikacin occur mainly in group 4, where morphological alterations were marked. Our results suggesting that amikacin is a potent factor are consistent with the clinical literature regarding nephrotoxicity. Our morphohistological findings include cellular necrosis, tubular obstruction, and basal membrane interruption.

REFERENCES


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