Experimental Urology

Histologic alterations of rat kidneys perfused with a Euro-Collins diltiazem solution

Alterações histológicas ocorridas em rins de ratos perfundidos com solução de Euro-Collins associada à diltiazem

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ABSTRACT

Purpose: Analyse the histologic changes of rat kidneys perfused with isotonic saline solution (ISS), Euro-Collins solution (ECS) and Euro-Collins solution with diltiazem (ECSD).

Methods: Thirty-six Wistar rats were used divided equally, as follow: group A (ISS), group B (ECS) and group C (ECSD). Through a catheter placed into the abdominal aorta, a renal perfusion was performed using a solution according to the group to which the animal belonged. After the complete perfusion, bilateral nephrectomy was performed and the organs were preserved under hypothermia for five distinct periods of time. Glomerulus and tubule were evaluated through optical microscopy.

Results: Renal perfusion with ECS and ECSD proved effectiveness in the preservation of the organs up to 36 hours and an increase in the percentage of injured glomeruli was noticed only in the period of 48 hours.

Conclusions: The results showed that exists an association between the tubular injury and the glomeruli lesion degree; kidneys with a higher degree of tubular damage were related to severe glomerular lesion. Also, the addition of a calcium channel blocker, diltiazem, to the ECS for the renal perfusion does not decrease the percentage of glomerular lesion.

Key words: Diltiazem. Perfusion. Kidney. Rats.

RESUMO

Objetivo: Analisar as alterações histológicas nos rins de ratos perfundidos com solução salina isotônica (ISS), solução Euro-Collins (ECS) e solução Euro-Collins com diltiazem (ECSD).

Métodos: Foram divididos, de forma igual, 36 ratos Wistar, como se segue: grupo A (ISS), grupo B (ECS), grupo C (ECSD). Através de um cateter localizado na aorta abdominal, foi realizada a perfusão renal com a solução de acordo com o grupo ao qual o animal pertencia. Após a perfusão total, realizou-se nefrectomia bilateral com a preservação dos órgãos sob hipotermia por cinco períodos distintos de tempo. Glomérulos e túbulos foram avaliados por microscopia óptica.

Resultados: Tanto a perfusão renal com ECS quanto a com ECSD provaram sua efetividade na preservação dos órgãos até 36 horas e aumento da porcentagem de glomérulos injuriados foi notada apenas no período de 48 horas.

Conclusões: Os resultados mostraram haver uma correlação entre a injeção tubular e o grau de lesão glomerular; rins com um maior grau de dano tubular foram relacionados com lesão glomerular severa. Além disso, a adição de um bloqueador de canal de cálcio, diltiazem, à ECS para a perfusão renal não diminui a porcentagem de lesão glomerular.


Introduction

The goal of ideal renal preservation is to promote the necessary time for cross matching, receiver selection, transportation of graft and preparing the receiver for surgery as well as leading to early reestablishment of renal function as the delay would incur into considerable costs due to longer hospital stay and need for repeated dialysis.

Prolonged organ preservation has only become possible after the development of intracellular electrolyte composition solutions. Since then many changes in the composition of solutions have been performed. Aiming at prolonging renal graft preservation time without, however, altering the organ’s viability and function, studies have been performed by adding different substances to the perfusion solution such as oxygen, steroids, free radical scavengers and calcium channel blockers such as trifluoperazine, verapamil and diltiazem.

Considerable interest has been shown recently in the calcium function as possible irreversible cell damage mediator after ischemia period. Cell membrane ischemic injury is associated with cell membrane lipid peroxidation and calcium influx, which subsequently induces irreversible cell damage.
to the influx of calcium from the extracellular compartment into the intracellular compartment and as a consequence with the activation of calcium-dependent catabolic process. Based on previously mentioned data this study has been proposed in order to assess histological changes occurred in rat kidneys submitted to perfusion with isotonic saline solution (ISS), Euro-Collins solution (ECS) and Euro-Collins solution associated to diltiazem (ECSD), calcium channel blocker in different periods of preservation.

**Methods**

Thirty six Wistar rats have been used (Rattus norvegicus albinus, Rodentia mammalia) divided into three groups with twelve animals each. Group A: perfusion with ISS, group B: perfusion with ECS and group C: perfusion with ECSD (25mg/l). The rats have been submitted to inhalation anesthesia with commercial ether followed by xyphopubic median laparotomy with the exposure of the abdominal cavity. A dissection of the aorta above and below the renal vessels has been conducted after laparotomy. The aorta has been isolated above the celiac trunk being repaired at this level with 5-0 silk yarn. In the distal portion a ligation of the aorta and the vena cava has been performed right above its bifurcation to the iliac vessels. Aorta has been catheterized with catheter type Abocath® 24 introduced right above the iliac bifurcation. After checking of blood reflux in the catheter, meaning it was correctly positioned; renal blood flow has been interrupted by the traction of superior aortic repair. Immediately after a renal perfusion has been initiated using the solution according to the group the animal belonged to. Temperature of used solutions were always at 4ºC and perfusion pressure was of 100 cm H₂O for every group (A, B and C). During perfusion a change in color of the bowel and renal parenchyma has been observed. Such changes have been used as criteria for effective perfusion, being interrupted at the moment kidneys showed a homogeneous color or at least five minutes after beginning of perfusion. After bilateral nephrectomy kidneys have been stored in plastic package containing 30ml of perfusion solution (ISS, ECS, ECSD) at 4ºC and placed inside another plastic package containing 10ml of ISS and ice. After being properly identified, the package has been stored in thermal container with ice. The kidneys have been preserved and fixed for histological analysis in five different periods: 0 (right after perfusion was finished), 12, 24, 36 and 48 hours. The choosing of period of kidney has been done through draw.

**Histological study**

From 72 kidneys used in this experiment 12 have been analyzed from the histological point of view at period 0 and 15 kidneys in each one of the other periods. After draw, kidney was sectioned longitudinally on its anti-hilar edge for adequate fixation. The blades have been made with three cutting levels, with intervals of 5 micrometers between them stained with hematoxylin and eosin (HE). Two renal structures have been evaluated: glomeruli and tubules. Total number of glomeruli as well as damaged glomeruli have been accounted for. Glomeruli were considered normal (Figure 1) whenever capillary loops were open, with thin walls and whenever there was no content inside the Bowman’s capsule. On the other hand, glomeruli were considered as damaged (Figure 2) whenever they presented a contraction of glomerular tuft with approximation of structures, contents inside the Bowman’s capsule or vacuolization of endothelial cells. The degree of glomerular lesion has been characterized as: light (up to 25% of damaged glomeruli), moderate (26% to 50% of damaged glomeruli), sharp (51% to 75% of damaged glomeruli) and severe (above 75% of damaged glomeruli).

**FIGURE 1** - Photomicrography from kidney tissue showing two glomeruli (G) with usual capillary and mesangial cellularity (HE 400x)

**FIGURE 2** - Photomicrography from kidney tissue showing damaged glomeruli. Contraction of glomerular tuft (*). Mesangial disorganization and content amorphous eosinophilic (●) inside the Bowman’s capsule (HE 400x)

Evaluation of tubular damage has been carried out according to kidney’s compromising extension. Renal tubules have been considered normal whenever coated by a single layer of cuboid cells of dense eosinophilic cytoplasm and tubular lights were optically empty. Injured tubules presented cytoplasmic vacuolization in different degrees. Four different categories were used, according to Chart 1.
**Statistical analysis**

Thirteen rats were considered for the statistical analysis, randomly placed in 3 groups (10 rats each group), being both kidneys from each rat used. In total, 60 kidneys have been considered. Result evaluations have been done in 4 periods: 12 hours, 24 hours, 36 hours and 48 hours. For each one of the variables (weight of rats, time and volume of perfusion) null hypothesis of equal averages has been tested for the three groups considered (A, B and C), versus an alternative hypothesis in which at least one of the groups presented an average different from the others. For that matter Cochran test has been applied. In case there was significant difference between the groups “t as in student” tests was adopted for independent samples. For variable analysis (total number of glomeruli, percentual of damaged glomeruli and tubular damage) the Mann-Whitney non-parametrical test was adopted. For research of association between damaged glomeruli and tubular damage the Spearman’s rank correlation coefficient for evaluation of this association was estimated. In every test the adopted significance level was of 5%.

**Results**

From 36 rats used in the experiment 6 of them were used in period 0 (4 kidneys in each group: A, B and C). In each kidney histological analysis showed normal renal architecture, presenting normal glomeruli and absence of tubular damage in period 0. Therefore, such kidneys were not part of statistical analysis; however, they have been used as pattern for histological study in the remaining periods (Table 1).

As for rats’ weight (p=0.1786) and perfused solution volume (p=0.9288) in each rat, results showed that there was no significant difference between the three groups. As for perfusion time, kidneys perfused with ECS presented a bigger perfusion time than kidneys perfused with ISS (p=0.0000001) and ECSD (p=0.0014).

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**CHART 1 – Tubular damage degree**

<table>
<thead>
<tr>
<th>Degree</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>+</td>
<td>Only a faint cortical band presented tubular vacuolization</td>
</tr>
<tr>
<td>++</td>
<td>Besides cortical compromising, some medullar tubules have also been compromised</td>
</tr>
<tr>
<td>+++</td>
<td>Cortical and medullar region compromised but still presenting preserved areas</td>
</tr>
<tr>
<td>++++</td>
<td>Diffuse cortical and medullar vacuolization, with no preserved areas</td>
</tr>
</tbody>
</table>

**TABLE 1 – Statistical comparison of total glomeruli counting and damaged glomeruli among groups a, b and c in the analyzed periods**

<table>
<thead>
<tr>
<th>PERIOD</th>
<th>COMPARED GROUPS</th>
<th>COUNTING GLOMERULI (p value)</th>
<th>DAMAGED GLOMERULI (p value)</th>
</tr>
</thead>
<tbody>
<tr>
<td>12</td>
<td>A x B</td>
<td>0.1150</td>
<td>0.0723</td>
</tr>
<tr>
<td></td>
<td>A x C</td>
<td>0.1666</td>
<td>0.0947</td>
</tr>
<tr>
<td></td>
<td>B x C</td>
<td>0.9166</td>
<td>0.4101</td>
</tr>
<tr>
<td>24</td>
<td>A x B</td>
<td>0.0361 *</td>
<td>0.0088 *</td>
</tr>
<tr>
<td></td>
<td>A x C</td>
<td>0.0361 *</td>
<td>0.0439 *</td>
</tr>
<tr>
<td></td>
<td>B x C</td>
<td>0.6015</td>
<td>0.2448</td>
</tr>
<tr>
<td>36</td>
<td>A x B</td>
<td>0.0283 *</td>
<td>0.0088 *</td>
</tr>
<tr>
<td></td>
<td>A x C</td>
<td>0.2506</td>
<td>0.0723</td>
</tr>
<tr>
<td></td>
<td>B x C</td>
<td>0.6752</td>
<td>0.5296</td>
</tr>
<tr>
<td>48</td>
<td>A x B</td>
<td>0.5284</td>
<td>0.0088 *</td>
</tr>
<tr>
<td></td>
<td>A x C</td>
<td>0.0749</td>
<td>0.3321</td>
</tr>
<tr>
<td></td>
<td>B x C</td>
<td>0.4620</td>
<td>0.3458</td>
</tr>
</tbody>
</table>

* Comparisons in wich were a rejection of the null hypothesis at significance level of 5%.

A: Kidneys perfused with SSI.
B: Kidneys perfused with SEC.
C: Kidneys perfused with SECD.

There was no significant difference in total number of glomeruli in the different periods (p=0.05). There was a significant difference, however, in the percentual of damaged glomeruli in each group between the periods 12x24 (p=0.0088), 12x36 (p=0.0088) and 12x48 (p=0.0088) hours in group A, between periods 12x48 (p=0.0465) hours in group B and between periods 12x48 (p=0.0232) and 24x48 (p=0.0439) hours in group C (Table 2).

**TABLE 2 – Frequencies observed in comparison of degree of glomerular injury and tubular damage**

<table>
<thead>
<tr>
<th>Glomeruli</th>
<th>Tubular Damage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Light lesion</td>
<td>+   ++  +++  ++++  Total</td>
</tr>
<tr>
<td>12</td>
<td>12   12   12   0   36</td>
</tr>
<tr>
<td>Moderate lesion</td>
<td>1    0    0    1    2</td>
</tr>
<tr>
<td>Sharp lesion</td>
<td>0    0    4    4    8</td>
</tr>
<tr>
<td>Severe lesion</td>
<td>0    0    3    11   14</td>
</tr>
<tr>
<td>Total</td>
<td>13   12   19   16   60</td>
</tr>
</tbody>
</table>
Discussion

The choice of rat as an experimental animal in this study was due to ease in obtaining and handling such animal and the accordance of literature which states rats’ isolated kidney as a doable and reliable method for evaluation of organ preservation. The choice for diltiazem in this work was based on the action mechanism of this blocker in the calcium channel. Diltiazem enhances blood renal flow through direct vasodilatation of glomeruli afferent arterioles. It also prevents calcium ions from entering intracellular space, consequently interrupting vicious cycle initiated by ischemia which leads to cellular damage. Besides, this substance enhances urinary volume and it is also natriuretic, as it inhibits sodium reabsorption by the distal tubules. That fact itself enhances glomerular filtering index due to the bigger renal perfusion produced by the increase of cardiac debit.

Perfused kidneys with ECS presented bigger perfusion time than kidneys perfused with ISS (p=0.000001) and ECSD (p=0.0014). On the Friarte and collaborators study, performed in 1981, there has been a difference in perfusion time between kidneys perfused with the same solution (Collins and Sacks-Kaufman solutions). Difference in perfusion time found on this work may be due to the different compositions of solutions used in renal perfusion, when a given solution demands a bigger time to perfuse the same volume.

Normal and damaged glomeruli counting has been carried out in a comparative way and aiming at evaluating kidney preservation. This method is little regarded in literature. In the present study and average of 95.85 glomeruli were counted in each kidney, varying from 66 to 126 glomeruli. In order to keep an average of counted glomeruli the same level of histological cut has been chosen, making sure the same renal area was assessed in every case. Results showed statistically significant difference in the total number of counted glomeruli between groups ISS x ECS and ISS x ECSD in the 24-hour period and between groups ISS x ECS in the 36-hour period (p<0.05). There was no significant difference, however, between the different periods in each group (p<0.05), confirming that glomeruli counting was similar in the evaluated groups.

Evaluation of glomerular damage is another factor used for analysis under the histological point of view, changes and preservation occurred in graft after perfusion. In the present study, glomerular damage present in the different groups and periods have been evaluated. When groups were compared there was no significant difference in the percentual of damaged glomeruli between groups A, B and C in the period of 12 hours, showing that the three solutions obtained similar results in the renal preservation during this period. Such data is according to the Fisher and collaborators work, in 1967, which demonstrated little glomerular lesion in kidneys perfused with Ringer lactate solution and preserved under simple hypothermia for up to 12 hours. Results, however, showed that there has been significant difference in the percentual of damaged glomeruli between ISS X ECS groups in the periods of 24, 36 and 48 hours and between ISS x ECSD groups in the period of 24 hours (p<0.05). Data showed that ECS, being in intracellular solution, has been more effective in the preservation of glomerular damage than ISS, which is extracellular.

As for comparisons between ECS x ECSD groups there hasn’t been significant differences between the percentual of damaged glomeruli in any of the evaluated periods (p<0.05), showing that the addition of calcium channel blocker to the Euro-Collins solution has not been effective in the preservation of glomerular damage. As for the percentual of damaged glomeruli the present study evaluated different periods for each group. Perfused kidneys with ECS showed significant difference when compared to periods of 12x48 and 24x48 hours. Data suggests that ECS, associated to diltiazem, provides protection against glomerular damage for a period up to 36 hours.

Tubular damage is another important and observed factor in histological changes occurred in preserved kidneys. A tubular damage evaluation has been carried out during this study as well as the association to the degree of glomerular damage. Obtained results showed direct correlation between tubular damage and the degree of glomerular damage. Eleven out of fourteen kidneys which presented severe glomerular damage have also presented tubular damage. However, none of the kidneys presenting light glomerular damage showed severe tubular damage. The result was similar to the one obtained by Hill and collaborators in 1976, which performed renal biopsy one hour after transplantation and observed that kidneys with a higher degree of tubular alteration correlated to a more intense glomerular damage, which proves the direct correlation between the degree of tubular alteration and glomerular damage.

Literature on the subject is controversial when it comes to the results obtained by calcium channel blockers in renal preservation. Blank et al. evaluated the effects of hypothermia and verapamil in rats’ kidneys submitted to sixty minutes of ischemia, showing that isolated hypothermia provides a better protection and that verapamil has not obtained protective effects against hot and cold ischemia. Such results have been confirmed by findings of electronic microscopy.

Nakamoto et al. has evaluated the effects of verapamil in renal preservation after 24 hours of hypothermic perfusion compared to simple perfusion using Euro-Collins solution. Authors showed that verapamil has not only enhanced kidney preservation but also improved their function after reperfusion. The fact that difference in the percentual of glomerular damage has not been found between groups ECS x ECSD in this study may be explained by the renal damage caused by tissue reperfusion. Lesions due to ischemia/tissue reperfusion have been described in the myocardium, brain, intestine, liver and kidney. Besides, tissue reperfusion may lead to cellular damage when tissue reoxygenating.

Cellular injury derived from ischemia/renal reperfusion may lead to a series of events: depletion of ATP reserves, free radicals formation derived from oxygen, cellular acidosis, increase in calcium cellular ions influx, lytic actions of some enzymes (proteases, phospholipases) and cellular apoptosis. The ischemic/reperfusion phenomena also involves the production of inflammatory mediators, activation of endothelial cells and leukocitary activation.

Therefore, protective action of calcium channel blockers may happen when beginning tissue reperfusion. Anaïse et al. has used calcium channel blocker (verapamil) during kidney perfusion in dogs and submitted to reperfusion after preservation.
for 48 and 72 hours. This substance showed favorable effect in the renal blood flow, decreasing the effects caused by tissue reperfusion.

Conclusions

Based on the results in this work it was concluded that the addition of diltiazem to the Euro-Collins solution has not decreased the percentual of damaged glomeruli when compared to the Euro-Collins solution itself in the periods of 12, 24, 36 and 48 hours. Is has also been observed that perfusion with Euro-Collins solution and Euro-Collins solution with addition of diltiazem are effective in the preservation of renal histology for a period up to 36 hours.

References


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