The Effect of Time on Apoptosis Changes Following Ischemia-Reperfusion in Isolated Heart of Rats

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**Abstract**

**Background and objective:** To study the time-dependent effects of ischemia – reperfusion on myocardium cells. Methodology: Male SD rats (270-330gr) were randomly grouped in four groups of 10 in an experimental study. After anesthesia using pentobarbital sodium (50-60 IP-kg/mg) the rats’ hearts in the treatment groups were isolated immediately and connected to langendorff apparatus with krebs solution at 37°C and a fixed pressure. During stabilization, 30m ischemia and 60, 90, and 120min reperfusion was carried out on the hearts. Hearts of the control group remained intact. Immunhistochemistry of apoptotic cells was performed using TUNEL POD Kit and positive TUNEL myocardium cells of each group, in five microscopic view fields, was counted. The results were represented as mean ± SD. **Findings:** Apoptotic cell counts in control group was 1±0.4, and this figure in the treatment groups T/60min, T/90min, and T/120min was 2.0±0.5, 3.0±0.3, and 6.0±0.3 respectively. Comparison between the treatment groups and control group was carried out using one-way ANOVA and the results showed no significant difference between the groups T/90min and T/120min and the groups T/60 and control (p>0.05). However, the differences between the groups T/120min and T/60min (p<0.001) and groups T/120min and T/90min (p<0.01) was significant. **Conclusion:** The study showed that ischemia-reperfusion time can be effective on apoptotic changes of myocardium cells of heart.

**Keywrds:** Ischemia –reperfusion- apoptosis- infraction

**Introduction**

Necrosis and apoptosis are two separate pathways of cell death in myocardium cells. In fact, there is a correlation between ischemia (I) and reperfusion (R) and biological and morphological specifications of cardiocytes under necrosis and apoptosis are very similar. Necrosis is usually referred to as random or pathologic cell death that is coincident with severe cell inflammation and split, cytoplasm protein coagulation and denaturation, breakdown and destruction of cell internal organelles and severe inflammatory response. On the contrary, apoptosis is a programmed and genetically controlled cell death. The main morphological features of apoptosis are cellular leakage, chromatin condensation, formation of cytoplasm blebbing and apoptotic objects that happen without losing integrity of cell membrane and inflammatory response. Although necrosis and apoptosis mechanisms are different, there are common features between them in pathological conditions [1-4]. Several studies have shown that ischemia-reperfusion induces apoptosis of myocardia cells in vivo; still, there is an ongoing debate if apoptosis starts with I or R. Gutlib et al. found that the traces of touchstone of apoptosis – i.e. nucleosome fragments of DNA- were found in rabbits’ ischemic myocardium 30min after infraction and 4hrs after reperfusion; however, this was not found in ischemic myocardium. Based on their surveys, the authors proposed that apoptosis only happens with reperfusion [5].

A brief review of the pathology literature indicated that abnormal apoptosis appears in many disorders and diseases. There are a few diseases known in which apoptosis plays...
no role and a major part of pathogens of diseases can be explained by taking into account the role of apoptosis. Studies have shown that short and limited periods of in vitro ischemia-reperfusion in the right kidney induced apoptosis, while prolonged ischemia induced necrosis. Gutli et al. demonstrated that ischemia-reperfusion induced apoptosis in myocytes of the rat and elaborated on the role of pH and ATPase in apoptosis following reperfusion. Piot et al. showed that ischemia-reperfusion in the rat’s heart could be a sign of apoptosis and then they found the evidences after a long reperfusion of coronary blood flow. Although, they managed to explain apoptosis changes following ischemia-reperfusion, they failed to determine how much apoptosis was induced after each period of trivial preconditioned ischemia [5, 6]. Leski studied the probable effects of ischemia-reperfusion in the case of coronary veins angioplasty and reported that increase of reperfusion time was effective on the post-surgery side-effects [5, 7]. Heart infraction pathology is featured with several factors and conditions, and gradual loss of myocardium is one of the top pathologic issues. Over the years, we have been provided with a great deal of information about the role of post-infraction and ischemia-reperfusion apoptosis and that the both phenomena highlight importance of apoptosis. In light of this, the possibility to limit myocardium cell death through preventing apoptosis is potentially important in treating heart infarction and adopting an optimum length of time of ischemia-reperfusion [6, 8]. Based on the findings of previous studies, the present one is an attempt to survey the role of time in ischemia-reperfusion of myocardium tissues of rats’ heart.

Materials and Methods

Through an experimental study, male rats (SD; 250-300 gr) were randomly categorized in four groups of ten (one control and three treatment groups). The subjects were anesthetized using pentobarbital sodium (50-60 IP-kg/mg) and immediately the hearts were isolated and connected to langendorff apparatus with krebs solution containing carbogen gas at 37ºC and fixed pressure. The subjects in the treatment groups received standard krebs solution during stabilization, 30mins limited ischemia, and 60, 90, and 120min reperfusion. The groups were labeled as T/60min, T/90min, and T/120min based on ischemia-reperfusion time. The subjects in the control groups did not have ischemia-reperfusion. The hearts of the treatment group, after the reperfusion, were covered by fixative and kept in autotechnicon filled with formalin, ethanol, xylene, and paraffin. Afterward, apoptosis detecting method (TUNEL assay) was implemented by sectioning the specimens. Apoptotic cells number in five random views of each specimen was counted under optical microscope. Data analyses were carried out using One-way ANOVA and Bartlett’s test and the results were reported as mean score ± standard deviation.

Implementation of TUNEL assay

1. After removing paraffin and hydrating, the prepared sections were exposed to proteinase K enzyme and rinsed by PBS solution after incubation (30min, 37ºC).
2. Tissue sections were exposed to TUNEL reaction mixture (50µl, 60min, 37ºC) and then rinsed by PBS solution.
3. After incubation with Converter-POD (50µl, 30min, 37ºC) the sections were rinsed by PBS and exposed to DAB for 20min at 25ºC.
4. The sections were rinsing by PBS solution and stained by blue toluidine [1, 9].

Results

Microscopic examinations showed an increase in condensation of chromatin and fragmentation of cell nucleuses and development of chromatin crescent in some samples. Due to cell DNA fragmentation that was

<table>
<thead>
<tr>
<th>Groups</th>
<th>mean score</th>
<th>Mean ± SD</th>
<th>Mean ± SE</th>
</tr>
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<tbody>
<tr>
<td>T/60min</td>
<td>2.4</td>
<td>2.4±1.1</td>
<td>2.4±0.5</td>
</tr>
<tr>
<td>T/90min</td>
<td>3.8</td>
<td>3.8±0.8</td>
<td>3.8±0.3</td>
</tr>
<tr>
<td>T/120min</td>
<td>6.0</td>
<td>6±0.707</td>
<td>6±0.316</td>
</tr>
<tr>
<td>Control</td>
<td>1.6</td>
<td>1.6±80</td>
<td>1.6±0.4</td>
</tr>
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</table>

Table 1. Mean Score, SD, and SE of Number of Apoptotic Cells in Treatment and Control Groups

Figure 1. Sectional Photomicrograph of Myocardium Tissue of The Rats in the Treatment Group After 120min Reperfusion. Positive TUNEL cells are spotted by arrows (TUNEL staining, magnification:100X)
Discussion

In terms of pathophysiology of cardiomyocytes apoptosis, several pro-apoptotic factors play a role in inducing apoptosis and tissue ischemia-reperfusion is one them that uses several pathway to induce cell apoptosis. Following blood reperfusion, an increase in cell calcium cytosol level happens that results in increase of penetrability potential of mitochondria membrane. Consequently, a large amounts of calcium enter mitochondria and then cytochrome C (Apaf-2) along with Apaf-13, dATP, and Apaf-3 trigger a cascade of caspases enzymes that results in apoptotic death of the cell. Moreover, high Ca++ content in cytoplasm of cell following tissue reperfusion activates endonuclease enzyme and fragmentation of DNA strands with factor of 180-200 base pairs, TUNEL staining with changes in cellular death including: apoptosis, several pro-apoptotic factors play a role in inducing apoptosis and tissue ischemia-reperfusion was significant (p<0.01) (Table 1).

The difference between T/90min and T/120min groups was significant (p<0.001) and between T/120min and T/90min and T/60 and control group was not significant (p>0.05). However, the difference between T/120min and T/60min groups (p<0.001) and between T/120min and T/90min groups was significant (p<0.01) (Table 1).

Acknowledgement

The authors wish to thank the Research Department and Mr. Gholamreza Danesh Pazhouhan who helped us as pathology expert.

References

7. Harper IS BJ, Chacon E, Reece JM, Herman B, Lemasters JJ. Inhibition of Na+/H+ exchange preserves viability, restores mechanical function, and prevents the pH paradox peak value after 60min reperfusion, which results in an increase of penetration potential of mitochondria membrane and facilitation of discharge of cytochrome C and AIF. However, AIF volume is higher than cytochrome C, which is due to the fact that the former is bound to anion phosphor-lipid in intra-membrane space and the latter is not bound in inter-membrane space; this means that AIF can easily cross the membrane [4, 11].

2. Expression level of the triggering gens and inducers of apoptosis like BAX, that play a role in formation of mitochondria membrane channels and discharge of cytochrome C, increases in long-term reperfusion and in presence of BAX active protein [12].

3. Kim et al. reported that activity level of caspase 3, which is a highly effective factor on apoptosis, increased after 2hrs reperfusion. Thereby, more changes at T=120min probably is due presence of adequate volume of caspase enzymes, and caspase 3 in particular [10-12].

At any rate, knowing about the role of time and its effect on tissue reperfusion paves the path for future works on reducing apoptotic changes in tissue reperfusion compounds. Using anti-apoptotic agents along with tissue reperfusion, and heart/liver transplant operation in particular, at proper moment can be highly effective on controlling tissue damages. Given the results and that number of apoptotic cells increases over time, prescribing anti-apoptotic medicines before tissue reperfusion or at early stages of reperfusion is recommended [2, 3, 6].

Figure 2. Sectional Photomicrograph of Myocardium Tissue of the Rats in Control Group (TUNEL Staining, Magnification:100X)