A Study on the Effects of Modulation of Intracellular Calcium on Excisional Wound Healing in Rabbit

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ABSTRACT

An in vitro study on the role of intracellular calcium ions in healing of excisional wound in rabbit was undertaken. We employed two drugs namely, glibenclamide and nitroglycerin that are topically applied in vivo to modulate the activity of intracellular calcium. Our model consisted of a 15 × 15 mm excisional wound. Seven groups of New Zealand rabbits were used. The first three groups served as untreated, Vaseline- and lubricating jelly vehicle-treated as controls. The remaining groups received topically 0.5 g of nitroglycerine (2 % in Vaseline base) or glibenclamide (1, 2 and 4% in lubricating jelly) on the day of excision and continued for 11 days. Using wound surface area measurement, complemented with measurement of the breaking strength and histological assessment, the results showed that inhibition of intracellular calcium ion had a favorable effect on wound healing. The mean wound half-lives and breaking strength were significant and concentration dependently reduced in glibenclamide-treated group. In contrast, in nitroglycerin-treated group the rate of wound healing and breaking strength were increased relative to untreated control and Vaseline treated groups. Histological findings revealed more organized collagen fibers and angiogenesis in nitroglycerin-treated wounds. The present study demonstrated that intracellular calcium ion has an important role on the overall process of wound healing. This information may be utilized in further studies to assess its role in healing of human wounds.

Keywords: Excisional wound, Nitroglycerin, Glibenclamide, Wound surface area, Tensile strength

INTRODUCTION

Accumulating evidences from biomedical research implicate calcium ion as a key intracellular signal that modulates the expression of cellular function and gene expression in a variety of systems including the developing ones [1]. As early as 1983, Chapman [2] demonstrated changes in calcium gradient following chemical or physical stimulation that correlated with activation of calcium channels across the cell membrane. Cell culture studies on keratinocytes and fibroblasts demonstrated the capacity of local calcium to modulate cellular proliferation, modification, maturation and the creation of epidermal lipid function through signal transduction and gene expression [3]. Such functional capabilities implicate calcium ion as a central role in epidermal regeneration and reconstruction in wound healing [4].

In the wound site, calcium concentration consistently changes with the biological events of the healing process [5]. Experimental studies have shown that calcium channel blockers, verapamil and nifedipine to abolish terminal differentiation of cultured keratinocytes [6]. On the other hand, calcium concentration in the extracellular media was found to be inversely proportional to cellular motility [7]. Furthermore, excessive calcium concentration in the wound environment induced reduction in proliferation and chemotactic responses and is believed to lead to delay in healing [8].

Topical application of verapamil to chronic wounds in experimental animals resulted in improved wound contraction and healing [9]. In contrast, diltiazem was not found useful in treatment of fibrous tissue formation in conjunctival wounds in rabbit [10]. All these observations are likely to have important clinical implications, but the available evidence is equivocal.

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Grzesiak and Pierschbacher in 1995 [11], suggested that the level of extracellular Mg$^{++}$ and Ca$^{++}$ could have a distinct impact on the adhesion and migratory activities of many cell types. The experimental models support the notion that management of calcium is essential for successful epidermal reconstruction [12], the benefits and possible implications under in vivo conditions of wound healing has received little attention. Therefore, we used an in vivo setting to assess the role of intracellular calcium on the healing of an experimentally induced excisional wound. For this purpose, we topically applied two drugs known to modulate the intracellular calcium level namely: glibenclamide, an antidiabetic drug known to increase the intracellular calcium ion via blocking ATP-dependent potassium channel (K$_{ATP}$) [13], and nitroglycerin, a potent coronary vasodilator that increases cGMP level and inhibits the intracellular calcium activity [14].

**MATERIALS AND METHODS**

**Animals and treatment.** Thirty-five white New Zealand rabbits of an average weight of 2.5 Kg (range 2.3 to 2.8 Kg) were utilized in this study, and divided randomly into seven groups of five animals each. The groups were allocated into the following categories: untreated control, vehicle 1 control (lubricating jell) (Abazar Darman Company, Tehran), vehicle 2 control (Vaseline) (Tehran Chemi Company), nitroglycerin (Berenguer-Infale, Germany) (2 %, in Vaseline base), and 1, 2 and 4 % glibenclamide (Gifted by Chemidaru Pharmaceutical Company, Tehran) in lubricating jell. The vehicles and treatment groups received 0.5 g of topical treatment daily commenced on the day of wounding and continued for 11 days post injury.

**Wounding procedure.** A 15 × 15-mm full thickness excisional wound was made on the lower flank of each animal according to the model proposed by Cross et al. [15]. The animals were held in standard crouching position, and the lower right flank were clipped with scissors and wet shaved with a scalpel blade. The area to be excisioned was outlined using a 15 × 15 mm template and then was locally anaesthetized. After established local anesthesia, by a subcutaneous injection of 1-2 ml of 2% lidocaine solution, a full-thickness wound was made by excising the skin within the confines of the template square down to the level of lose subcutaneous fascia.

**Wound measurement.** While the animals were held in standard crouching position, five wound boundaries for each wound were traced by a fine-tipped permanent marker onto an acetate transparency. This procedure was performed daily starting from day 0, the day of wounding and continued for 11 days post wounding. The areas were quantified by placing the transparency film over graph paper and counting the squares. The areas for each day after wounding were converted into percentage of wound surface area relative to day 0, which was considered 100%. The degree of effectiveness of drug treatments on the overall process of wound healing was quantified and compared by extrapolation of wound half-lives [16]. Furthermore, the area-under-the-curve (AUC) for all treatments relative to vehicle control was calculated [17].

**Histological observations.** At the end of the wound measurement period on day 11, the wound area together with 0.5 cm of intact skin from the outer boundary was removed. Half of the samples were selected for histological assessment. For this purpose, the skin samples were placed in 10 % formalin in normal saline. The tissues were processed by normal standard procedures of embedding in paraffin, sectioning, dehydration, and staining with haematoxylin and eosin, and finally mounted on a microscopic slide. The following parameters were assessed microscopically: the apparent number of polymorphonuclear leucocytes and fibroblastic-shaped cells, the relative extent of development new vascular beds (or angiogenesis) and the organization of collagen fibers were compared in different treatment groups.

**Measurement of wound tensile strength.** The remaining half of the removed wound samples was used for measurement of the tensile strength. For this purpose, a simple manual method was employed, by which the tissue was hanged by cotton threads vertical to the wound surface and fixed at one end to a metal bar and the other end to a 1 g aluminum envelope. With an aid of a large fixed magnifying glass and drop-wise addition of tap water, the point at which detachment of the wound surface from the skin or a break within the body of the wound tissue was considered the end point for breaking. The envelope was removed and weighed which reflected the force necessary for breaking the wound tissue.

**Statistical analysis.** The data are expressed as mean ± standard error of means (s.e.m.). Wound
RESULTS

The general observation from measurement of wound surface area showed a progressive reduction in the wound surface area with time in all groups. The data collected are illustrated in Figure 1, and revealed the mean half-lives were significantly different in untreated-control as compared to both glibenclamide and nitroglycerin-treated groups. Glibenclamide produced a dose-dependent delay in the wound surface area compared to untreated and vehicle 2 (lubricating jell) ($P<0.01$). Wound half-lives were 6.5, 8.9, and 10.3 days for 1, 2 and 4 % glibenclamide-treated groups respectively (Fig. 1). The half-life for lubricating jell (vehicle) treated group was 6 days and similar to untreated control group ($P>0.05$). In comparison, the nitroglycerin-treated group showed an acceleration in the rate of healing producing a reduction of wound half-life of 3.52 days, statistically significance compared to both the untreated control ($P<0.01$), and vehicle (Vaseline) treated groups ($P<0.05$).

For vaseline-treated group, the wound half-life was 5.1 days, in contrast to 6 days for untreated control group ($P<0.05$). Calculation of the AUC for all treatment groups relative untreated control group are summarized in Table 1. Where the values obtained reflected the relative extent of the rate of healing on the overall process of wound healing for each treatment relative to untreated control, and a negative or a positive sign represents the direction of this effects, i.e. decrease or increase respectively (17). These data were correlated with the times for 50% reduction in wound surface area measurements (Fig. 1).

**Histological observations.** Haematoxylin and eosin staining of 4 µm thick paraffin embedded sections, taken from the glibenclamide-treated wounds, revealed delayed epithelialization and a dose-dependent increase in the inflammatory polymorphonuclear leukocytes and dis-organized collagen fibers (Figs. 2e and f) compared to untreated control (Fig. 2a).

<table>
<thead>
<tr>
<th></th>
<th>Vaseline</th>
<th>Jell</th>
<th>Ntg</th>
<th>Glb 4%</th>
<th>Glb 2%</th>
<th>Glb 1%</th>
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<tr>
<td>A.U.C.</td>
<td>+325.8</td>
<td>-1.03</td>
<td>+785.9</td>
<td>-1772</td>
<td>-992.9</td>
<td>-237.2</td>
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| Jell (lubricating jell vehicle); Ntg (Nitroglycerin 2 % in vaseline base); Glb (Glibenclamide in lubricating jell). The plus and minus signs indicate an increase or decrease rate of wound surface area reduction relative to the control, respectively. |
Table 2. Summary of the means ± s.e.m. of the tensile breaking strengths of the excisional wound granulation tissues taken from different treatment groups of rats.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Mean ± sem</th>
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<tbody>
<tr>
<td>Control</td>
<td>16.7 ± 1.2</td>
</tr>
<tr>
<td>Vaseline</td>
<td>18.5 ± 1.4</td>
</tr>
<tr>
<td>Jell</td>
<td>16.6 ± 1.2</td>
</tr>
<tr>
<td>Ntg</td>
<td>35.1* ± 2</td>
</tr>
<tr>
<td>Glb 4%</td>
<td>6.98* ± 0.9</td>
</tr>
<tr>
<td>Glb 2%</td>
<td>11 ± 0.7</td>
</tr>
<tr>
<td>Glb 1%</td>
<td>14.5 ± 1.3</td>
</tr>
</tbody>
</table>

*P<0.001 relative to no treatment control group, ANOVA followed by Dunnet’s test. Jell (lubricating jelly vehicle); Ntg (Nitroglycerin 2% in vaseline base); Glb (Glibenclamide in lubricating jelly).

Fig. 2. Photomicrographic illustrations of haemotoxylline and oesin stained rabbit skin samples taken from (a), (100 ×) normal skin; (b), (100 ×) and (c), (400 ×) untreated control; (d), (400 ×): nitroglycerin-treated and (e) and (f), (400 ×): 4% glibenclamide-treated wound granulation tissues 11 days post wounding. A, angiogenesis; C, collagen fibres; E, epidermis; F, fibroblasts; K, keratin layer; N, neutrophil; O, oedema; R, red-blood cell and S, scab.
**Wound tensile strength measurements.** The tensile strengths measurements were found to be closely correlated with both the histological and wound surface area measurements. The greatest tensile strength measured was for the nitroglycerin-treated group, while there was a dose dependent and significant reduction in the tensile strength in the glibenclamide-treated group (Table 2).

**DISCUSSION**

This study demonstrated that modulation of the intracellular calcium ions causes marked changes in the overall process of wound healing. Glibenclamide, a drug known to increase the level of intracellular calcium concentration via inhibition of ATP-dependent potassium channels (K<sub>ATP</sub>) [13], produced adverse effects on the wound healing. In contrast, nitroglycerin a drug known to act via activation of cGMP, an intracellular intermediate that inhibits the calcium activity [14], was found to favorably promote wound healing. All these observations suggest a paradoxical role for calcium ions or the inhibition of intracellular calcium activity [13]. Findings showed a correlated increase in the blood glucose level (results not shown). This suggests that the effects produced by glibenclamide were solely local. Glibenclamide was found to produce reduction in both wound breaking strength and slowed the rate of closure of the wound surface area, suggestive to adverse effects on wound healing process. In addition, the histological observations showed correspondingly disorganized collagen and increased polymorphonuclear leucocytes, indicating an abnormality in collagen synthesis and persistent inflammation at the site of the wound. Whether healing of wounds in diabetic patients, who are under glibenclamide treatment, is also adversely affected warrants further clinical or experimental assessment.

Additional pharmacological effects, other than K<sub>ATP</sub> channel blockage and via still unknown mechanism attributed to glibenclamide, were reported [20, 21]. Glibenclamide was found to cause depolarization effect on the cardiac muscles during the ventricular fibrillation [20]. It also produced relaxant effect on isolated rat aorta. Although, we did not measure changes in the intracellular calcium, it seems the calcium level
increases since all the parameters considered in this study were opposite to those observed for nitroglycerin, a drug known to produce inhibition of intracellular calcium activity. However, the exact cells affected by glibenclamide cannot be allocated from these findings, and deserve further studies.

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REFERENCES