Topical application of silver-curcumin on wound healing in rabbits

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Abstract
Curcumin, a natural product obtained from the rhizomes of Curcuma longa, is used traditionally in prevention of wound infection. To determine the effects of silver and curcumin combination on surgical wound healing, a study was conducted in rabbits. A total of 32 surgical wounds were created in 16 rabbits in four groups. Information was recorded from Day 0 to Day 42 postoperatively. Swelling area of wound, elevation of suture line from the skin surface, width of sutured area and contraction length were recorded weekly. Treatment with curcumin, 1% silver sulfadiazine (1% SSD) and a mixture of the two resulted in swelling of 11.4 ± 0.1 mm, 11.1 ± 0.4 mm and 11.0 ± 0.1 mm, respectively. Swelling (12.2 ± 0.3mm) and elevation of sutured line was higher (3.4 ± 0.2 mm) in wounds of control group and lower (2.3 ± 0.1 mm) in wounds treated with a mixture of curcumin and 1% SSD. The scores of wound colonization were lowest (0.8) in 1% SSD group followed by mixture of curcumin and 1% SSD (1.0), curcumin (1.3) and tincture of benzoin (1.6) alone. Histopathologically reactive cells decreased markedly in wounds treated with combination of 1% SSD and curcumin at D3 with increased fibrous connective tissue. However, wounds treated with 1% SSD showed fewer reactive cells than curcumin group. Proliferation of fibrous connective tissue was highest in the silver treated wound, which indicates good wound healing process. Overall, wound healing was improved by the topical application of 1% SSD alone. Curcumin had positive effects on wound healing process but less than 1% SSD. These results indicate that 1% silver sulfadiazine is the best topical therapy for wound management in rabbits. (Bangl. vet. 2015. Vol. 32, No. 2, 55 – 64)

Introduction
Wounds are reported to be the most common surgical condition in animals. Healing of wounds is a complex biological event (Gillitzer et al., 2001) and extrinsic and intrinsic factors may result in complications (Hess et al., 2003). A wound may lead to serious consequences if not treated (Harpal and Kuldip, 1993; Mashhood et al., 2006). Typically, the steps in the wound repair process include inflammation, angiogenesis, development of granulation tissue, repair of connective and epithelial tissues, and ultimately remodelling (Midwood et al., 2004).

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Treatment of wounds is important to achieve the best functional and aesthetic results in a short time (Myers et al., 1980; Adams et al., 2003; Mamun, 2012). Topical antibacterial therapy may alter the wound environment by reduction of fibroplasia or promotion of epithelialization (Stashak, 1991; Mashhood et al., 2006). Silver nitrate, Nano crystalline silver, and some silver-containing dressings have anti-inflammatory effects and encourage neovascularization (Walker et al., 2007; Kortoes et al., 2010; Wilkinson et al., 2011).

Benefits of oral and topical use of curcumin (diferuloylmethane), a natural product obtained from the rhizomes of Curcuma longa, for the treatment of cutaneous wound has been reported (Sidhu et al., 1999). It has been reported that curcumin accelerates wound healing in cattle (Mamun, 2012). Curcumin improves re-epithelialization and migration of cells such as myofibroblasts, fibroblasts, and macrophages (Sidhu et al., 1999). Curcumin inhibits pain and inflammation by selectively inhibiting the arachidonic acid cascade (Gong et al., 2013). Curcumin acts as a scavenger of nitric oxide and inhibits cyclo-oxygenase (COX-2), a pro-inflammatory cytokines (Mamun, 2012). No research has been reported on open wound healing in rabbits using silver sulfadiazine and curcumin combination. The present study was designed to study the therapeutic effects of curcumin (Curcuma longa) and silver sulfadiazine on surgical wound in rabbits.

Materials and Methods

Study location
A series of studies were done on rabbits to find out the effect of silver and curcumin in healing of surgical wounds sutured with simple interrupted pattern using silk. The studies were conducted at the experimental shed of the Department of Medicine, Bangladesh Agricultural University, Mymensingh from January to May 2013.

Experimental animals
With the approval of the Department of Surgery and Obstetrics, Bangladesh Agricultural University, Mymensingh 16 apparently healthy rabbits were used. Body weight of the animals ranged from 1.8 to 2.0 Kg. The animals were kept under standard laboratory conditions and veterinary supervision with water and food ad libitum. Before the study the rabbits were kept in quarantine for three weeks.

Preparation of 1% silver cream and turmeric paste
Silver cream was prepared with 1% silver sulfadiazine (SSD) and 99% paraben (Burnsile® Beximco Pharmaceuticals, Bangladesh). Fresh herbal pastes were prepared from turmeric root (Curcuma longa). The fresh turmeric roots were purchased from KR Shopping centre, Bangladesh Agricultural University campus, Mymensingh. The roots were cleaned with water. These were ground in a mortar to prepare cream. A combination was made by mixing (1:1) 1% SSD cream and turmeric paste.
**Experimental design**

A total of 32 surgical wounds were made on the skin of 16 rabbits with two in each. Rabbits were divided into four groups with four animals in each group (Table 1).

**Group-T:** Fresh turmeric paste was applied locally to surgical wounds daily. These animals were maintained so as to avoid interference with granulation tissue formation.

**Group-S:** Silver sulfadiazine 1% cream was applied to surgical wounds in each animal daily. The treatment schedule was as in Group-T.

**Group-ST:** A mixture of silver sulfadiazine 1% cream and turmeric was applied locally daily to surgical wounds.

**Group-C:** This group was kept as control. In this group tincture of benzoin was applied in surgical wounds in rabbits.

<table>
<thead>
<tr>
<th>Group</th>
<th>Materials for treatment</th>
<th>Form of materials</th>
<th>No. of animals</th>
<th>No. of wounds</th>
</tr>
</thead>
<tbody>
<tr>
<td>T</td>
<td>Turmeric</td>
<td>Paste</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>S</td>
<td>1% SSD</td>
<td>Cream</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>ST</td>
<td>Turmeric and 1% SSD</td>
<td>Paste</td>
<td>4</td>
<td>8</td>
</tr>
<tr>
<td>C</td>
<td>Control (Tincture of benzoin)</td>
<td>Solution</td>
<td>4</td>
<td>8</td>
</tr>
</tbody>
</table>

SSD = silver sulfadiazine

All wounds were closed with simple interrupted suture using silk thread. Follow-up information was obtained from day of surgical operation up to Day 42 after surgery. Swellings, elevation of sutured line from the skin surface, width of sutured area were recorded. Elevation of sutured line was recorded up to 7 days of surgery. Width of sutured area was measured at Day 0 (D₀), Day 3 (D₃), Day 7 (D₇), Day 14 (D₁₄), Day 21 (D₂₁) to determine wound contraction length. Tissue samples were collected from all treatment groups at D₁, D₃ and D₇ for histopathological study.

**Anaesthesia**

The rabbits were anaesthetized with single intramuscular injections of 6 mg/kg xylazine hydrochloride (Xylaxon®, 23.2 mg/mL, Indian Immunological Ltd., India) and 40 mg/kg ketamine hydrochloride (G-Ketamine®, Gonosasthya, Bangladesh, 50 mg/mL ketamine hydrochloride USP).

**Surgical wound**

The thigh region of the rabbits was shaved and cleaned with 10% Povidone Iodine solution. Surgical wound of 1 cm length and 0.5 cm depth was made. Exudates were
removed by dry gauze. Wound was closed with simple interrupted suture using silk thread. All sutures were placed 8 mm apart. Distance between suturing needle placement and border of cutting edge was 5 mm.

**Observation of morphological changes**
Slide calipers was used to measure swelling area (mm), elevation of suture line (mm), wound contraction and width of sutured area of wound (mm). Swelling was observed up to three days after operation because swelling started decreasing gradually from Day 3 (D3). Elevation of sutured line was recorded until 7 days after surgery. Width of sutured area was measured from the day of surgical intervention Day 0 (D0), Day 3 (D3), Day 7 (D7), Day 14 (D14), Day 21 (D21).

**Assessment of wound colonization**
Wound colonization was assessed according to exudation, purulent efflux, efflux odour, erythema and oedema. A score range from 0 and 3 was presented here:

- 0: No finding of colonization
- 1: Mild exudation and odour
- 2: Erythema, moderate purulent efflux, exudation and odour
- 3: Severe exudation, purulent efflux, odour, oedema and erythema.

Each assessment was performed by the same blinded worker who was unaware of the scores of rabbits determined before and during the treatment.

**Assessment of wound healing**
This study was continued for six weeks after the formation of surgical wound. Each wound was clinically observed and digital photos were taken with a camera every three days during the following six weeks. Wounds were considered to have healed when visible epithelium covered the wound and cicatrisation and pigmentation was found. The days to healing were recorded for all the animals and the mean time was calculated for each group.

**Histopathological assessment**
Biopsies (1.5 cm × 1 cm) were collected from the wound areas of each animal on the 1st, 3rd and 7th days after wounding using standard surgical procedure. The wound tissue contained dermis and epidermis. The samples were fixed in 10% buffered neutral formalin solution more than seven days for histopathology. Histopathological slides were prepared and the whole procedure was performed by the Department of Pathology, Bangladesh Agricultural University, Mymensingh.

**Statistical analysis**
The data obtained in the present study were analysed with SPSS statistics 17.0 software. Probability P<0.05 was considered statistically significant.
Results and Discussion

At the beginning of the study, all the surgical wounds were similar. At the end of six weeks, no unhealed wounds were present.

Morphological changes

Swelling of the wound edges was observed in all four groups (Table 2). Treatment with curcumin (group T), silver (group S) and mixture of both (group ST) resulted in the swelling area 11.4 ± 0.1 mm, 11.1 ± 0.4 mm and 11.0 ± 0.1 mm respectively; there was no significant difference (Table 2). The swelling (12.2 ± 0.3 mm) was significantly (P<0.05) higher in group C (Control group). Elevation of sutured line was higher (3.4 ± 0.2 mm) in group C and lower (2.2 ± 0.1 mm) in group ST (Table 3). Higher swollen area and elevated suture line indicate more inflammation in control group. The contraction length varied insignificantly (P>.05) among the groups (Table 2).

<table>
<thead>
<tr>
<th>Groups</th>
<th>Swelling of suturing area (mm)</th>
<th>Elevation of sutured line (mm)</th>
<th>Average contraction length (mm/week)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group S</td>
<td>11.1 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.3 ± 0.11&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.5 ± 0.2</td>
</tr>
<tr>
<td>Group T</td>
<td>11.4 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.4 ± 0.21&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.6 ± 0.2</td>
</tr>
<tr>
<td>Group ST</td>
<td>11.0 ± 0.1&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.2 ± 0.10&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.4 ± 0.1</td>
</tr>
<tr>
<td>Group C</td>
<td>12.2 ± 0.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.4 ± 0.23&lt;sup&gt;b&lt;/sup&gt;</td>
<td>1.8 ± 0.1</td>
</tr>
</tbody>
</table>

Table 2. The effect of 1% silver sulfadiazine-curcumin on wound healing in rabbits

<sup>a, b</sup> indicate significant (P<0.05) difference in groups; Mean ± SEM

Wound colonization

The scores of wound colonization were lowest in group S (0.8) followed by group ST (1.0), T (1.3) and C (1.6). This reveals that odour and exudation from wounds of silver-treated group were minimum due to least microbial changes. These differences were significant only at weeks 1, 2 and 3 (P<0.05) (Graph 1).

Days to healing

No significant differences (P>0.05) on the healing time were observed among the groups (Table 4). The mean days to complete healing ranged from 30 days for group S to 38 days for group C with a mean for all groups of 34 days (Graph 2).

Histopathological changes

Infiltration of reactive cells including neutrophils, macrophages and lymphocytes were present in all samples, which indicated inflammation. Additionally, tissue debris and haemorrhage were present and blood vessels were congested in the wounds of control group (Group C) at D<sub>1</sub>. Reactive cells decreased markedly in wounds treated with combination of 1% silver sulfadiazine and curcumin at D<sub>3</sub>. Fibrous connective tissue was observed on this sample at D<sub>1</sub> (Fig. 2d). Wounds treated with 1% SSD showed fewer reactive cells than those of curcumin group collected at D<sub>3</sub> (Fig. 2b and 2c). Notably, the proliferation of fibrous connective tissue was the highest in the silver-
treated wound (Fig. 2c). Continuous formation of keratin layer was observed in group S (Fig. 3a) at D7. In case of group ST, healing was progressed (Fig. 3c) at D7. A comparatively thick keratinized layer was in group T (Fig. 3b). However, the edges of the wound sample of control group at D7 did not come in apposition although; keratin layer was formed along the edges of the incised tissue (Fig. 3d). It may be due to lack of joining the two edges and consequently retarded healing of the wound in this group.

Graph 1. Progress of colonized wounds of four groups in six consecutive weeks

Table 3. Time required for healing of wounds treated with silver, curcumin, a combination of silver and curcumin, and tincture of benzoin

<table>
<thead>
<tr>
<th>No. of animals</th>
<th>Healing time (days)</th>
<th></th>
<th></th>
<th></th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Silver</td>
<td>Silver + Curcumin</td>
<td>Curcumin</td>
<td>Tincture of benzoin</td>
</tr>
<tr>
<td>1.</td>
<td>29</td>
<td>42</td>
<td>42</td>
<td>38</td>
</tr>
<tr>
<td>2.</td>
<td>32</td>
<td>32</td>
<td>38</td>
<td>36</td>
</tr>
<tr>
<td>3.</td>
<td>28</td>
<td>28</td>
<td>32</td>
<td>38</td>
</tr>
<tr>
<td>4.</td>
<td>31</td>
<td>30</td>
<td>35</td>
<td>40</td>
</tr>
<tr>
<td>Average</td>
<td>30.0 ± 0.1</td>
<td>33.0 ± 0.1</td>
<td>36.7 ± 0.2</td>
<td>38.0 ± 0.2</td>
</tr>
</tbody>
</table>

P>0.05 comparing the groups

Graph 2. Mean days to wound healing
Fig. 1. Presence of reactive cells (arrows) beneath the keratinized tissues of epidermis of wounds of control group (a), group S (b), group T (c) and group ST (d) at D1 after treatment. Congested blood vessel is seen (dark circle) in control group (a).

Fig. 2. Presence of reactive cells (dark color) and newly formed fibrous tissue (whitish area) in wounds of group C (a), group T (b), group S (c) and group ST (d) after treatment.
Cicatrisation is an intricate process in which the skin repairs itself after injury (Nguyen et al., 2009). The application of topical medications is one means of keeping a wound free from contamination. These preparations allow high local antimicrobial efficacy while avoiding systemic toxicity and are most effective in the earlier stages of healing prior to a solid granulation bed (Brandt et al., 2012). The increased swelling and width of sutured area following treatments were noted up to Day 3 (D3) postoperatively. The record of the width of wound area at Day 7 (D7) and Day 14 (D14) postoperatively has been done to understand wound contraction process morphologically. There was no significant variation in wounds of all groups in term of diminishing contraction length per week. This result supports the hypothesis that wound contraction depends on the myofibroblast located at the periphery of the wound, its connection to components of the extra cellular matrix and myofibroblast proliferation (Rohrich, 1990). Among four groups, higher swelled area and elevation of suture line were observed in wounds of control group (treated with tincture of benzoin) in comparison to that of other three groups. This result hypothesizes that tincture of benzoin cause marked tissue reaction in wounds.

In this study 1% SSD cream produced comparatively better results in wound healing as a topical medication. It produced wound healing in 30 days, 33 days treated with mixture of 1% SSD and curcumin, 36 days with curcumin and 38 days in control group. Additionally, the scores of wound colonization were the lowest (2.2 in 1st
week) in silver treated animals (group S). Silver sulfadiazine increases the rate of re-
epithelialization in acute and chronic wounds and had good effects in wound healing
(Kontoes et al., 2010; Choi et al., 2013). It may be suggested that, 1% silver sulfadiazine
may be used as an effective topical medication for treating wounds in rabbits.

In this study, no negative effect from either 1% SSD or curcumin paste was observed.
It is believed that either the turmeric or the silver can be used clinically in wound
healing. However, in silver-treated wounds, there was early subsidence of
inflammation, better control of infection and quicker wound healing compared to the
other three groups. It is recommended that 1% SSD may be a practical choice for
wound care. It also relieves pain, makes the wound sterile and is available in the local
markets. Further investigation is necessary to determine the relative safety of these
products on the healing wound. Once that is done, the relative value of the products
can be determined.

Conclusions

Wound care is improved and simplified by the topical use of 1% silver sulfadiazine.

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