Acceleration of Skin Wound Healing with Tragacanth (Astragalus) Preparation:  
An Experimental Pilot Study in Rats  
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Abstract- Gum tragacanth is a natural complex mixture of polysaccharides and alkaline minerals extracted from species of Astragalus plant, which is found widely in arid regions of the Middle East. In a pilot experimental study we examined the effects of its topical application on wound healing in ten albino adult male rats. Two similar parasagittal elliptical full-thickness wounds (control vs. test samples) were created on the dorsum of each animal. Test group samples were fully covered by a thin layer of gum tragacanth daily. The extent of wound healing was evaluated by planimetric analysis on multiple occasions during the 10-day study period. On the 7th day of the study, the percent of wound closure was significantly higher in gum tragacanth-treated specimens compared to the control samples (87%±2% vs. 70%±4%, P<0.001). The majority of wounds in the test group were completely closed by the 10th day of the study. The difference in wound healing index measured by histological examination on day 10 of the study was also statistically meaningful between the two groups (0.624±0.097 vs. 0.255±0.063, P<0.05). The results of this study clearly showed the useful effects of topical application of gum tragacanth in acceleration of skin wound contraction and healing. More studies are encouraged to identify the implicating agents and precisely understand the mechanism by which they exert their wound healing effects.  
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Keywords: Astragalus; Tragacanth; Skin; Wound healing  

Introduction  

Wound healing has historically been the focus of interest for many scientists. Since the pioneering research by Winter (1), the concept of wound dressings has been evolving from the traditional so-called passive products having a minimal role in the healing process to the functional more active materials which, through the interaction with the wounds they cover, establish and maintain an optimal environment for wound repair. An ideal wound dressing should protect the wound from bacterial infection, provide a moist and healing environment, and be safe and biocompatible (2).  

Various polysaccharide biopolymers such as chitosan (3,4), alginate (5) and glycosaminoglycans (6), having hydrogel-forming properties, have been shown to be effective in their application as wound dressing materials. In the present study, we aimed to assess the applicability of a different arabino-galactan-based natural material, namely tragacanth, as a dressing for wound occlusion and accelerator of the healing process. Gum tragacanth, an exudate of branches and roots used in the manufacturing of food pharmaceutical and industrial products because of its valuable hydrophilic and colloidal properties, is obtained from several tragacanth species of Astragalus occurring principally in the mountain regions of the Middle East (7). Tragacanth is a complex mixture of polysaccharides, which yields upon hydrolysis D-galacturonic acid, D-galactopyranose, L-fucose (6-deoxy-L-galactose), D-xylopyranose, L-arabinofuranose, and a very small amount of L-rhamnose. The acidic components are largely present as calcium, magnesium, and potassium salts. The gum also contains trace amounts of amino acids and their derivatives. It has a molecular weight of about 840,000 Dalton and an elongated shape of 4500Á by 19Á for a flake type of non-degraded gum. Tragacanth is considered to contain two primary constituents,
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tragacanthin and bassorin, which are both insoluble in alcohol and have high molecular weights. The minor component, tragacanthin, is a highly branched arabinoxylan and is soluble in water to give a colloidal hydrosol solution. It has a ring containing three molecules of galacturonic acid and one molecule of arabinose, with a side chain of two molecules of arabinose. The major component, bassorin, though insoluble in water, has the capacity to swell and form a gel. It is believed to contain polymethoxylated acids that yield tragacanthin upon demethoxylation. Gum tragacanth swells in cold water to give extremely viscous colloidal solutions (8-10). The wound-healing property of tragacanth has been evaluated in the current study using a rat model with full-thickness excisional skin wounds and histological examinations.

Materials and Methods

Tragacanth gel 5% (w/v) was prepared by dissolving powdered tragacanth gum extracted from Astragalus gummifer (Iranian commercial market) in deionized water. Sterilization was performed by tendalization.

Experimental design and procedure

The animal experiment was performed in accordance with “the Guide for the Care and Use of Laboratory Animals” published by the US National Institute of Health (NIH Publication No. 85-23, revised 1996). Ten albino rats weighing 250±50g (Wistar, male, 6-month-old, Inst. Pasteur Iran) were used in this study. After induction of anesthesia by intraperitoneal (ip) injection of ketamine (75 mg/kg) and xylazine (5 mg/kg), dorsal hair was clipped and the area was wiped with 70% (v/v) ethanol. The procedures were carried out on a warming blanket. Using a scalpel, two similar (15 mm) parasagittal full-thickness elliptical skin wounds, 3 cm away from each other, were created on the dorsum of each rat, thus each animal was its own control. Extreme caution was taken to avoid injuring the perforating vessels. About 250 µl of tragacanth gel was applied daily onto the right-side wound of each animal (test samples) for 10 days. The left-side control wounds received similar amounts of sterile deionized water. The procedures were carried out on a warming blanket. Using a scalpel, two similar (15 mm) parasagittal full-thickness elliptical skin wounds, 3 cm away from each other, were created on the dorsum of each rat, thus each animal was its own control. Extreme caution was taken to avoid injuring the perforating vessels. About 250 µl of tragacanth gel was applied daily onto the right-side wound of each animal (test samples) for 10 days. The left-side control wounds received similar amounts of sterile deionized water.

Clinical assessment

The wound sites were digitally photographed on days 0, 3, 7, and 10 after the initial procedure. The length and area of each wound was measured. As well, the percent of wound closure was calculated according to the following equation: % wound closure = [(S₀-Sₓ)/S₀] × 100, in which S₀ and Sₓ are wound areas on the day of wound creation and the day of evaluation respectively. Planimetric calculations were made using Image J software (version 1.40 for Windows, NIH, Bethesda, USA).

Histological examination

After the animals were sacrificed, 3×5 cm dorsal skin patches including the wounds were excised and fixed overnight in 10% (v/v) phosphate-buffered formaldehyde solution. The tissues were washed with PBS, embedded in paraffin, and sectioned in 10 µm increments. The sections were made perpendicular to the surface of the wounds. The sections were then positioned on a slide, and stained with hematoxylin and eosin (H&E) reagent. To analyze the degree of wound healing, central portions of the wounds were viewed under 40× magnification. The area of granulation tissue (consisting of fibroblasts and matrix deposits, inflammatory cells and blood vessels) was quantitatively evaluated by a color image analyzing software (E200, Nikon, Tokyo, Japan). Moreover, the area of new epithelium over the wound bed was measured in a similar way and the index of wound healing was calculated according to the following formula: wound healing index = epidermis area/ (granulation tissue area + epidermis area).

Statistical analysis

All data are expressed as mean±standard error (SE) of mean. Differences between groups were determined by paired Student’s t-test. P-values less than 0.05 were considered statistically significant. Computations were performed using the SPSS statistical software (version 15.0 for Windows, SPSS Inc., Chicago, USA).

Results

Macroscopic observations

Results of macroscopic evaluations including wound lengths, wound areas and the percentage of wound closure are presented in table 1. As also seen in figure 1, in contrast with the control wounds, tragacanth-treated wounds healed far more rapidly.
Table 1. The wound length (mm), the wound areas (mm²), and the percents of wound closure in control and tragacanth-treated groups 3, 7, and 10 days after creating the wounds.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Day</th>
<th>Control</th>
<th>Tragacanth</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wound length (mm)</td>
<td>3</td>
<td>13.4±0.2</td>
<td>12.8±0.1</td>
<td>&gt;0.05</td>
</tr>
<tr>
<td></td>
<td>7</td>
<td>10.0±0.8</td>
<td>7.4±0.6</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>5.6±1.2</td>
<td>1.4±0.6</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Wound area (mm²)</td>
<td>3</td>
<td>26.0±2.1</td>
<td>21.4±1.5</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>7</td>
<td>11.4±1.8</td>
<td>5.4±0.9</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>7.2±2.3</td>
<td>0.6±0.3</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>Wound closure (%)</td>
<td>3</td>
<td>32±5</td>
<td>45±4</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>7</td>
<td>70±4</td>
<td>87±2</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>83±5</td>
<td>99±1</td>
<td>&lt;0.05</td>
</tr>
</tbody>
</table>

Statistically significant differences in wound contraction and closure between the two groups were observed upon day 3 of the study (P<0.001). Signs of wound infection were noted in neither of the groups. Moreover, there were no evident signs of an inflammatory reaction to tragacanth in treated wounds.

Microscopic observations

On histologic examination of the specimens on day 10, improved epithelialization, extracellular matrix deposition and tissue remodeling were evident in tragacanth-treated wounds in contrast to the extensive inflammation and granulation tissue formation noted in control group samples (Figure 2). No significant infiltration of inflammatory cells to the wound areas was observed in the tragacanth-treated wounds.

The differences in epithelialized tissue area and wound healing index between the two groups were statistically significant with p values less than 0.01 and 0.05 respectively (Table 2).

Table 2. The area of repaired epithelium (μm²), granulation tissue (μm²), and the wound healing index in control and tragacanth-treated groups 10 days after creating the wounds.

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control</th>
<th>Tragacanth</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Repaired epithelium</td>
<td>27262±3427</td>
<td>61362±10587</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Granulation tissue</td>
<td>131667±3504</td>
<td>99029±39422</td>
<td>0.551</td>
</tr>
<tr>
<td>Wound healing index</td>
<td>0.255±0.063</td>
<td>0.624±0.097</td>
<td>&lt;0.05</td>
</tr>
</tbody>
</table>

Figure 1. Acceleration of wound closure and healing using tragacanth (right-side wounds) compared to control samples (left-side wounds) in two animals on day 10 after the initial wounding.
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Figure 2. Photomicrograph of control (A1, 4X; A2, 10X) and tragacanth-treated wounds (B1, 4X; B2, 10X) on day 10. Note the near complete tissue epithelialization (black arrows) and mature scar formation (transparent arrows) in tragacanth-treated wounds compared to the vast tissue granulation (transparent arrowheads) and vessel dilation (black arrowheads) in the control group.

Discussion

The plant *Astragalus* is employed in folk medicine as a mild laxative. According to traditional beliefs of some peoples in the Middle East and North Africa, it is implied that the leaves contain useful elements in preparation of wound healing lotions. Tragacanth, the dried viscid exudation from the trunk and branches of *Astragalus*, is the source of a pharmaceutical suspending agent, so-called gum tragacanth (9). Tragacanth gum is a highly branched, heterogeneous hydrophilic carbohydrate polymer containing galacturonic acid, galactose, arabinose, xylose, etc. It is a complex slightly acidic polysaccharide bounded with small proportions of protein and trace amounts of starch and cellulosic material present. It also contains a few mineral elements including calcium, magnesium, and potassium (10,11).

Gum tragacanth is a widely used natural emulsifier and thickener employed in food, drug and allied health industries (12-14). It has been shown to decrease plasma LDL and cholesterol in hyperlipidemic patients (15). Consumption of tragacanth is considered to be safe as no toxic effects have been reported, even with large dietary intakes of it in human experiments (16). Moreover, no evidence of carcinogenicity has been found in animal studies involving tragacanth gum (17,18). Strobel *et al.* investigated the immunogenicity of various types of polysaccharide gums including tragacanth and concluded that proper processing of these substances can significantly reduce the immune response, although a complete systemic immune reaction could not be utterly averted (19). Based on a quality control assessment performed by Moghbel *et al.*, the tragacanth gum available in the traditional Iranian market, which was also used in our study, meets the regulatory requirements of USP23 with acceptable very low amounts of arsenic, lead and heavy metals (20).

In the present study, about 90% wound closure in the tragacanth-treated group occurred within 7 days, and all of the tragacanth-treated wounds were completely or almost completely closed by the 10th day of the study. Moreover, the wound healing index, incorporating two factors of granulation tissue formation and epithelial regeneration, was also notably higher in this group (Tables 1 and 2). It could be therefore inferred that tragacanth components accelerate the transition from the inflammation and tissue granulation phases of the wound healing process and enhance mature scar formation and extracellular matrix remodeling which boosts the eventual wound contracture and closure. The findings of our study are consistent with similar wound healing experiments employing various materials including tragacanth (20) and chitosan (3,4,21,22). Moghbel *et al.* studied the effect of topical tragacanth application on excisional wounds in rabbits and concluded that this treatment could significantly shorten the period of wound healing and closure compared to eucerin (control) and phenytoin-treated groups; however, they did not establish any implicating mechanisms (20). Based on a series of studies by Ishihara et al. on chitosan, another arabinogalactan containing polysaccharide biopolymer, about 90% wound closure was achieved within 8 days in chitosan hydrogel-treated wounds, whereas similar amount of wound closure was seen only after 10 days in control specimens (3,4).

Several mechanisms encompassing all major phases of the wound healing process (including acute inflammation, granulation tissue formation, and tissue remodeling) have been proposed for the wound healing effects of chitosan and other arabinogalactan-based substances in different studies, which may be also involved in the effect observed in the present experiment. Chitosan has been shown to increase infiltration of inflammatory cells including...
polymorphonuclears (PMNs) (22,23). It also exerts chemotaxis of PMNs and macrophages via complement C5a activation (24). Another study has shown that it mediates stimulation of macrophage function (25). Chitosan also potentiates basic fibroblast growth factor (FGF-2) activity by protecting it from inactivation by the interaction between FGF-2 and chitosan polysaccharide constituents (26). Ishihara et al. proposed that physical forces initiated by adsorption of various protein molecules from the wound surface into the chitosan preparation may cause wound contraction. Chitosan also stimulates migration and proliferation of dermal fibroblasts. The positively charged chitosan molecules adsorb some substances involved in cell proliferation and migration, such as growth factors and cytokines, from blood plasma or wound exudate. The adsorbed substances subsequently could induce angiogenesis, fibrosis, and epithelialization (4). Arabinogalactans possess a potent dose-dependent complement fixation activity and lead to T cell-independent induction of B cell proliferation (27,28). Arabinogalactan polysaccharides also stimulate the proliferation and enhance the viability of skin fibroblasts (29). The mineral constituents of gum tragacanth including calcium, magnesium, and potassium may also play a substantial role in the wound healing process. Sequential events in wound repair require a conducive environment within the wound bed and a balanced pool of metal ions among which these elements are important. It has been demonstrated by Lansdown et al. that as wounds heal, local concentrations of calcium and magnesium change according to the phase in the wound healing cascade and associated biochemical events (30). Calcium has an established role in the normal homeostasis of mammalian skin and serves as a modulator in keratinocyte proliferation and maturation. Furthermore, it may be required in epidermal cell migration and regeneration (31). Magnesium might also by contributory to increased motility of fibroblasts and keratinocytes (32).

The results of our study explicitly reveal the efficacy of gum tragacanth as an effective, facile to prepare, and yet safe, biocompatible, cheap and readily available agent for wound dressing. Hence, tragacanth-containing formulations could be considered as efficacious preparations in chronic and slow-healing wounds such as diabetic foot ulcers, bed sores, and ischemic wounds; however, further studies are required to investigate its exact mechanisms of action in the healing process and verify its efficiency in clinical settings.

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