Wound Healing Activity of the Ethanolic Extract of 
Onosma Bracteatum Wall

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Abstract
The wound healing activity of ethanolic extract of aerial parts of Onosma bracteatum evaluated on excision and incision model, in albino rats, in the form of an ointment with two concentrations (5% and 10% w/w ointment of bark extract in simple ointment base). Both concentrations of the ethanolic extract showed significant response in both the wound types tested when compared with the control group. Nitrofurazone ointment (0.2% w/w) used as standard.

I. Introduction
In tribal areas different crude drugs extract used to treat various skin disorders including wound. Wound healing process involve several steps, which involves coagulation, formation of granulation tissue, collagation and aquisation of wound strength. During the formation of new tissue, endothelial cells proliferate and form new blood vessels. Onosma bracteatum (OB), Wall (Family Boraginaceae, commonly known as Gaozaban, Gojihva) which has been reported to be used in the treatment of asthma and bronchitis. The drug is used as tonic, alterative, demulcent, diuretic and is considered cooling. It is useful as a spasmolytic. A decoction is used in the treatment of rheumatism, syphilis and leprosy. The plant is considered to be useful in relieving excessive thirst and restlessness in febrile excitement, and also to be useful in relieving functional palpitation of the heart, irritation of the bladder and stomach, and strangury.

Onosma bracteatum has been used in folk medicine for the treatment of skin diseases and wound. A survey of literature revealed that no systematic approach has been made to study the wound healing activity of this plant. Thus the present study was undertaken to assess the effect of this indigenous plant on different parameters related to wound healing in rats.

II. Materials and Methods
i. Plant material
The dried aerial parts of Onosma bracteatum was purchased from Dravid Herbs World, Pondicherry, India.
a. Incision wound
In incision wound model four groups (The group I was considered as control, the group II served as the reference standard and treated with 0.2%w/w Nitrofurazone ointment. The group III animals were treated with the 5%w/w ethanolic extract and the group IV animals were treated with 10%w/w ethanolic extract of Onosma bracteatum) of animals containing six in each group. Paravertebral incision of 6cm. long were made on either side of the vertebral column of the rat. Care was taken to see that incision was at least 1cm. lateral to vertebral column. The wounds were closed with interrupted sutures of 1cm. apart. The animals were caged individually. The sutures were removed on 8th post wounding day. The tensile strength of the wound was measured on 10th post wounding day².

b. Excision wound-In excision wound four groups (The group I was considered as control, the group II served as the reference standard and treated with 0.2%w/w Nitrofurazone ointment. The group III animals were treated with the 5%w/w ethanolic extract and the group IV animals were treated with 10%w/w ethanolic extract of Onosma bracteatum) of animals containing six in each group. A circular piece of full thickness (approx.500 mm²) was cut off from a predetermined area on the back of the rat. Wounds were traced on 1mm² graph paper on the day of wounding and subsequently on alternate days until healing was complete. Changes in wound area were calculated, giving an indication of the rate of wound contraction. Number of days required for falling of the eschar without any residual raw wound gave the period of epithelization⁵. The ointment of the fruit extract, standard drug and simple ointment was applied to the wound twice daily, until recovery to the respective groups of animals. Statistical analysis-The results are expressed as mean ±SE of six animals in each group. The data were evaluated by students t-test and the values of P≤0.001 were considered statistically significant.

III. RESULTS
It was observed that the wound healing contracting ability of the extract ointment in different concentrations was significantly greater than that of the control (i.e. simple ointment treated group). The 10% (w/w) extract ointment treated groups showed significant wound healing from the fourth day onwards, which was comparable to that of the standard drug, i.e. nitrofurazone ointment treated group of animals. The wound closure time was lesser, as well as the percentage of wound contraction was much more with the 10% w/w extract ointment treated group (18±1 days for 100% contraction which was almost similar to that of the nitrofurazone treated group). The 5%(w/w) extract ointment treated group of animals showed significant wound contraction from the eighth day onwards and achieved 100% with the wound closure time of 20±2 days.

In incision wound model the measurement of the effect of the extract and standard drug on the tensile strength is shown in Table II. The tensile strength of the 10% extract treated group and the nitrofurazone ointment treated group were comparable to each other. The 5% extract ointment treated group showed a lesser but significant increase in the tensile strength compared to the control group. Thus both concentrations of the extract as well as the standard drug showed a significant increase in tensile strength in the 10 days old wound. The results of the present study revealed that both concentration (5% and 10%w/w) of ethanolic extract of Onosma bracteatum fruit have significant wound healing activity in both incision as well as excision wound models.

IV. DISCUSSION
Wound healing is a fundamental response to tissue injury that results in restoration of tissue integrity. This is mainly achieved by the synthesis of the connective tissue matrix. Collagen is a major protein of the extracellular matrix and is the component that ultimately contributes to wound strength. Tannins promote the wound healing through several cellular mechanisms; chelation of the free radicals and reactive species of oxygen, promoting contraction of the wound and increasing the formation of capillary vessels and fibroblasts and including keratinocyte proliferation, but do not act on the differentiation towards cornified cells. Similar findings have been reported with the extracts of the plants containing tannins by earlier workers. However, our results revealed that tannins are one of the important phytoconstituents responsible for wound healing mainly due to their astringent and antimicrobial property. Hence, it can be inferred that the wound healing activity of the plant Onosma bracteatum is due to its high tannin content, which seems to be responsible for wound contraction and increased rate of epithelization.
Table 1: Effect of ethanolic extract of Onosma bracteatum on % wound closure of excision wound model

<table>
<thead>
<tr>
<th>Group</th>
<th>4th Day</th>
<th>8th Day</th>
<th>12th Day</th>
<th>16th Day</th>
<th>Period of epithelization in days</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>15.82±0.68</td>
<td>27.21±1.02</td>
<td>48.21±1.80</td>
<td>68.53±2.60</td>
<td>24</td>
</tr>
<tr>
<td>Nitrofurazone</td>
<td>35.28±0.15</td>
<td>76.80±0.19</td>
<td>89.81±0.58</td>
<td>97.11±0.048</td>
<td>18</td>
</tr>
<tr>
<td>Extract(10%)</td>
<td>31.44±1.01</td>
<td>72.24±1.24</td>
<td>81.39±2.36</td>
<td>90.84±2.10</td>
<td>18*</td>
</tr>
<tr>
<td>Extract(5%)</td>
<td>20.82±1.02</td>
<td>34.26±1.82</td>
<td>61.94±2.78</td>
<td>80.56±2.32</td>
<td>20</td>
</tr>
</tbody>
</table>

Values are mean ±SE, *P<0.001 vs control n=6 animals in each group.

Table 2: Effect of ethanolic extract of Onosma bracteatum on tensile strength of incision wound

<table>
<thead>
<tr>
<th>Treatment groups</th>
<th>Tensile strength(g)</th>
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</thead>
<tbody>
<tr>
<td>Control</td>
<td>310±4.6</td>
</tr>
<tr>
<td>Nitrofurazone</td>
<td>564±1.8*</td>
</tr>
<tr>
<td>Extract(10%)</td>
<td>522±4.2*</td>
</tr>
<tr>
<td>Extract(5%)</td>
<td>450±5.8*</td>
</tr>
</tbody>
</table>

Values are mean ±SE, *P<0.001 vs control n=6 animals in each group.

REFERENCES