Influence of *Aspilia pluriseta* Schweinf (Asteraceae) on the healing of dermal excision wounds (mouse model) and skin sensitization activity (Guinea pig model)

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**Background:** The skin is highly predisposed to injury because of its direct contact with the environment. The aim of treating wounds is to both hasten healing, and to minimise the occurrence of perturbations of the healing process. Many plants traditionally used to treat wounds have been proven to support the healing process using scientific models. *Aspilia pluriseta* has been used by a number of communities in East and Southern Africa to treat wounds.

**Objectives:** This study aimed at testing ethnomedical claims of wound healing activity of *A. pluriseta* using preclinical models.

**Methods:** Aerial parts of the plant were ground and incorporated into an ointment base (10% and 20% w/w) to evaluate the influence of the plant on the healing of acute excision wounds in mice compared to Silverex Cream® and Simple Ointment (B.P.). The 20% ointment was tested for skin sensitization in guinea pigs.

**Results:** The effects of the plant-based ointments on wound contraction and gross epithelialisation time were less than significantly different from the controls (p≥0.05), but histopathologic examination revealed remarkable epithelialisation and collagen deposition in the wounds treated with these ointments. The 20% *A. pluriseta*-based ointment induced moderate allergic contact dermatitis.

**Key words:** *Aspilia pluriseta*, wound healing, skin sensitization, excision wound model

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1. **Introduction**

A wound is an injury that results from the disruption of the normal tissue anatomy and function. It may result in a break of the epithelium with or without loss of underlying connective tissue. Wounds may result from physical, chemical, microbial agents or immunological mechanisms (Raina et al, 2008). Wound healing is the process through which the wound tissue is restored as closely as possible to its pre-injury state and is the result of interaction between cells, blood and connective tissue components directed by growth factors and cytokines. It is characterized by three major phases, inflammation, proliferation and remodelling which are distinct but overlap in time (Singer and Clark 1999). Cutaneous wounds primarily heal by three mechanisms: connective tissue matrix deposition, contraction and epithelialisation (Diegelmann and...
The primary objective of this study was informed by the need for pharmacological and toxicological testing of traditional remedies before recommending them for therapy (Niles Gutha 2010; Midiwo et al, 2001). We evaluated the effects of the ointment containing powdered aerial parts of *A. pluriseta* on selected processes of excision wound healing and dermal safety, specifically allergy induction.

2. Materials and Methods

2.1 Plant material

*Aspilia pluriseta* aerial parts (stems, leaves and flower heads) were collected in Ruiru, Kiambu County in the month of January, 2012. The identity of the plant was confirmed by the East Africa Herbarium where a voucher specimen was deposited (Voucher no: Kuria/Asp/001). The plant material was cleaned with tap water and then dried in the shade for 10 days, then ground using an electric mill.

2.2 Preparation of the ointments

The powdered *A. pluriseta* was triturated into an ointment base (Simple Ointment B.P) to form a 10% and 20% (w/w) *A. pluriseta* ointment for testing.

2.3 Excision wound healing assay

Four month old, male, Swiss Albino mice were purchased from the Kenya Agricultural Research Institute (KARI) Trypanosomiasis Research Centre. The animals were allowed 7 days to acclimatize to conditions at the animal house in the Department of Public Health, Pharmacology and Toxicology, University of Nairobi. They were fed standard pelleted feed (Unga Mice Pencils, Unga Feeds) and water *ad libitum*.

The excision model was used to provide information on wound contraction, epithelialisation and histologic data. The backs of the mice were shaved using an electric clipper. The shaved area was disinfected using povidone iodine. A square area, approximately $100\text{mm}^2$ was marked on the dorsum using a stencil. Full thickness excision wounds were created using thumb forceps and scissors along the edges of the marked areas (Yadav et al, 2014). The excision wounds were created with the mice under halothane anaesthesia. The mice were then randomly divided into four groups each having eight members. The first two groups were treated with the 10% and 20% *A. pluriseta* ointment. The third group was treated with the vehicle ointment (Simple Ointment B.P) and acted as the negative control group. The fourth group was treated with a standard wound drug, Silverex Cream® (Ranbaxy) containing 1% silver sulfadiazine. The wounds were left open (Sünantar et al, 2011) and treated once daily for 21 days. Before the application of the respective treatment, the wounds were cleaned with antiseptic solution. The wounds were observed daily over the test duration. Any hair growing over the wound area was trimmed to facilitate observation. The time it took for each wound to be fully covered by an epithelium (time when the scab fell off without leaving a raw area) was recorded for each animal. The wounds margins were traced onto tracing paper for area determination after wound creation. Wound areas were estimated by the use of 1mm$^2$ graph paper(Yadav et al, 2014). Wound contraction was tracked by determining the wound area every three days until the wounds were fully re-epithelialised grossly. Wound contraction was calculated as percent reduction of wound area relative to the initial wound area on the day of wounding (initial wound area = 100%).
On the 7th and 14th days, a representative animal from each group was sacrificed and a biopsy of the wound harvested for histology. On the 21st day, all the animals were euthanized using halothane and histopathology samples were harvested from all the animals. The harvested samples were fixed in 10% formalin, processed routinely and blocked with paraffin wax. Serial sections, 5µm thick were cut with a microtome and then stained with Hematoxylin and Eosin and Masson’s Trichrome (Nisbet et al, 2010). The stained sections were examined using a light microscope. The selected histologic parameters (neutrophil population, macrophage population, fibroblast population, collagen deposition, vascularisation and epithelialisation) were evaluated qualitatively and scored into three categories, +, ++ and ++++, representing increasing magnitude of each respective parameter.

2.4 Skin sensitization assay

This was carried out along OECD Test Guideline 406 (Buehler Non-adjuvant test method) (OECD 1992) with modifications on the number of animals and the use of an occlusive patch. 3 female adult (3 months old) albino guinea pigs were used. A 4cm² square area was marked on the right posterior flank which was earlier shaved, about 1cm from the dorsal midline and a film of the 20% A. pluriseta ointment was applied on it. This application was repeated on the 7th and 14th days and constituted the induction phase of the trial. On the 28th day, the left side, equally shaved and marked was used for challenge phase. Six hours after the application of the ointment on the new site, the residual ointment was wiped off using moist cotton wool. 24 hours later, the site was observed for a skin reaction, and the lesions scored. Challenge exposure was repeated again on the left side on the 35th day to confirm the observations made on after the first challenge exposure.

2.5 Data analysis

Means (±SEM) were computed for measures of wound contraction and epithelialisation times. The means for the four treatment groups were compared with each other using one-way Analysis of Variance (ANOVA). P values ≤0.05 were considered significant. The histology and skin sensitization data were not subjected to statistical analysis.

2.6 Ethical considerations

The animals used in these assays were handled humanely at all times in accordance with animal care and use guidelines of the University of Nairobi and other applicable guidelines and regulations.

3. Results

3.1 Epithelialisation time

The 20% A. pluriseta treated group took less time than the other groups to fully epithelialise as shown in Figure 1. The difference in the mean epithelialisation time was however not statistically significant.

![Figure 1](image)

*Figure 1*: Graph showing the epithelialisation time (mean number of days ± SEM (standard error of mean)) for the different treatments from left to right: 10% A pluriseta ointment, 20% A pluriseta ointment, Simple Ointment (negative control) and Silverex Cream (reference drug).
3.2 Wound contraction

The 20% *A. pluriseta* ointment showed marginally higher contraction (assessed as percent wound area reduction) towards the end of the first week and during the second week of healing. However, the difference in wound contraction rates between the test and the control groups was not statistically significant. All the wounds were fully closed (100% wound area reduction) on day 21. The wound contraction results are shown in Figure 2.

3.3 Histology

In the first week of healing, the wounds that were treated with the *A. pluriseta* ointments had intensive granulation as opposed to the negative control group which had thinly deposited granulation tissue. Also, the...
A. pluriseta ointment and the standard drug treated wounds had more mature fibroblasts than the negative control group. On Day 14 of healing, all the wounds except those treated with the 20% A. pluriseta ointment had haemorrhage. The A. pluriseta ointment treated wounds had more collagen, comparable to the standard drug treated wounds and more than the negative control group. At the end of the experiment on the 21st day, the A. pluriseta ointment treated wounds had fully epithelialised. They also had abundant collagen and a diminished number of blood capillaries. The histology results are summarised in Table 1 and Figure 3.

### 3.4 Skin sensitization

The 20% A. pluriseta ointment induced moderate allergic reaction after the challenge exposure on the 28th day which was indicated by a confluent erythema on the skin area used for challenge exposure. The erythema persisted for approximately 48 hours and then resolved on its own. This exposure was repeated 7 days later (35 days after the first application) to confirm the results obtained, and similar observations were made.

### Table 1: Summary of the histological observations

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group</th>
<th>Day 7</th>
<th>Day 14</th>
<th>Day 21</th>
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<tr>
<td>Neutrophils</td>
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<td>+++</td>
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<tr>
<td></td>
<td>20% A pluriseta</td>
<td>+++</td>
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<tr>
<td></td>
<td>Simple ointment</td>
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<td></td>
<td>Silverex® Cream</td>
<td>++</td>
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<tr>
<td>Macrophages</td>
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<td></td>
<td>Simple ointment</td>
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<td></td>
<td>Silverex® Cream</td>
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<tr>
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<tr>
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### 4. Discussion

Aspilia pluriseta at 10% and 20% concentration in simple ointment was observed to have an effect on the contraction, granulation and epithelialisation of excision wounds in a mouse model. It also induced allergic dermatitis in guinea pigs as evidenced by development of erythema on repeated application. The observed effects on the process of wound healing are in agreement with the observations made in similar
studies on the wound healing activity of plants (Yadav et al, 2014). The wound healing activity is similar to that reported by Okoli, Akah, and Okoli (2007) who reported enhancement of epithelialisation of excision wounds by the methanol extract of Aspilia africana, a member of the same genus as the test plant, A. pluriseta. The extent to which Aspilia pluriseta promoted wound contraction and gross epithelialisation was less than significantly different from the controls, as opposed to the measures of the same parameters in like studies. This observation may be explained by the fact that in this study, whole plant material was used as opposed to aqueous and organic solvent extracts from plant tissue in other studies. The allergy inducing activity is in agreement with the findings of other studies, that plants in family Asteraceae (formerly Compositae) readily induce allergic contact dermatitis due to sesquiterpene lactones in these plants’ tissues (Jovanović and Poljacci 2003).

The observed activity of Aspilia pluriseta on wounds is in agreement with other studies on the effect of medicinal plants in the healing of wounds. The wound healing enhancement that was observed in this project is similar to that seen in other experiments on in vivo support of wound healing by plant extracts (Kumar et al, 2007).

The findings of this study point to a multi-pronged enhancement of healing by A. pluriseta ointments. The plant material supported early polymorphonuclear inflammation, while at the same time allowing the recruitment of wound macrophages. It also induced or supported an early intensive granulation with numerous and more mature fibroblasts. Moreover, the plant-based ointments seemed to enhance collagen synthesis and deposition. Sections from groups treated with the ointments were more fibrotic than those from the control groups. Also, the A. pluriseta ointments promoted more epidermal remodelling.

5. Conclusion

In view of these observations, we confirm the wound healing activity of A. pluriseta reported ethnomedicinally. Further research is needed to determine the exact mechanism of action of the plant in the healing process of wounds. Development of a drug for treating acute wounds from the plant should be explored. The allergenic potential of the plant should be borne in mind during use of the plant or products derived from it.

Conflict of Interest declaration

The authors declare no conflict of interest.

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References


