OBJECTIVE: To determine bleeding time using Moringa oleifera leaf extract versus saline control in an experimental epistaxis model.

METHODS:
Design: Randomized controlled trial
Setting: Tertiary Government Training Hospital
Participants: Ten adult male New Zealand White rabbits were acclimatized for 1 week in a standard environment. One-centimeter long, full-thickness mucosal wounds in the junction of the nasal floor and anterior part of the septum were treated randomly with topical Moringa oleifera extract or colored isotonic saline control in either right or left nasal cavity, one site at a time. The duration of bleeding – time bleeding started to time bleeding stopped -- was recorded in seconds. Data was subjected to a t-test for paired samples.

RESULTS: The mean bleeding time for wounds treated with Moringa extract was 53 seconds (range 38-70 secs), versus 159 seconds (range 100-218 secs) for controls. The bleeding time in the former was significantly shorter than in the latter (p = .000019, t-stat = 8.139), with a mean difference of 106 seconds between the two groups.

CONCLUSION: Moringa oleifera leaf extract was associated with significantly shorter bleeding time than saline control in this experimental epistaxis model and may be worth investigating further as a hemostatic agent for epistaxis.

KEYWORDS: epistaxis, malunggay, Moringa oleifera extract, topical hemostatic agent

bleeding time using Moringa Oleifera (Malunggay) Leaf Extract versus Saline Control in a Rabbit Epistaxis Model: A Randomized Controlled Trial

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dislodged nasal packing. Thus, investigators have continued to look for alternative methods to treat this common problem.  

*Moringa oleifera,* also known as *malungay,* *mulangay,* horseradish tree, drumstick tree, or Ben oil tree, is a perennial softwood tree with low timber quality. It has long been known for its traditional medicinal as well as its industrial uses, and has been attributed with antibiotic and anti-inflammatory properties. Studies revealed that topical application of its extract promotes wound healing. However, its hemostatic effect on epistaxis has yet to be established. Using the keywords “Moringa,” “hemostatic” and “epistaxis,” we found no published study regarding the hemostatic effect of *Moringa* on epistaxis in a search of PubMed, the Cochrane Library, Science Direct, Philippine E-Journals, and the journals Philippine Journal of Health Research and Development, Cochrane Library, Science Direct, Philippine E-Journals, and the journals Philippine Journal of Health Research and Development.

In order to explore its potential hemostatic effects, the objective of this study is to determine the bleeding time using *Moringa oleifera* leaf extract versus saline control in an experimental epistaxis model.

**METHODS**

**Preparation of Moringa Extract and Control**

A modification of extract preparation yielding 13% aqueous extract was utilized. Fresh, green *Moringa oleifera* leaves procured from the local market were authenticated by a local biologist. The leaves were chopped and air-dried under shade for 72 hrs. Powdered leaves were obtained by crushing the dried leaves using a mortar and pestle. One hundred grams of powdered leaves was soaked in 500 ml of distilled water and left standing for 48 hours at 30°C, then filtered using Whatman No. 1 filter paper to obtain *Moringa* extract.

A liter of isotonic saline solution with 20 ml of green food coloring (McCormick, McCormick Philippines Inc., Novaliches, Quezon City) made up control solution with similar appearance to the *Moringa* extract.

The *Moringa* extract and control solutions were separately put in opaque specimen bottles with 20 ml solution per bottle. The bottles were labeled Solution A and Solution B by the principal investigator, with Solution A (*Moringa* extract) and Solution B (control) unknown to the blinded examiner, a licensed veterinarian. The solutions in each bottle were only used once.

**Animal Experiment**

Ten adult male New Zealand White rabbits with a mean weight of 1.26 kg (range, 1.1 – 1.5 kg) were purchased from a certified animal distributor and acclimatized for 1 week in a standard environment. They were housed in standard cages of 1m² area each, in a room with a constant temperature of 22°C ± 4°C and a 12-hour light/dark cycle, fed twice daily with 200g generic rabbit growing pellets, and unlimited access to tap water. The study was approved by the Institutional Animal Care and Use Committee (IACUC): <IACUC Protocol Number M003>. The experiment was conducted in the clinic of one licensed veterinarian, who performed all the procedures with one assistant. Both veterinarian and assistant were blinded.

All animals were intramuscularly sedated using 0.02 mg/kg Acepromazine (Labistress® 5 mg/mL, Labiana Life Sciences, Barcelona, Spain) and 11 mg/kg Ketamine chloride (Ceva Ketamine 100 mg/mL, Ceva Santé Animale, Sydney, Australia). One-centimeter long full-thickness longitudinal mucosal wounds, not advancing to the cartilage, were made using Blade #11 (Kai Medical, Kai Industries Co. Ltd., Seki City, Japan), at the junction of the nasal floor and anterior part of the septum of the right and left nasal cavity, one site at a time. (Figure 1) The wounds were randomized to be treated with either Solution A or B on either side.

Solutions were topically applied with cotton pledgets soaked in either solution as soon as bleeding started, and pressed gently on the wounds for 1 second, every 30 seconds (re-soaking the cotton pledget in solution for every application), until the bleeding stopped. The duration of bleeding – time bleeding started to time bleeding stopped-- was recorded in seconds by the assistant.

Post-operative care consisted of application of topical Clotrimazole 10 mg, gentamicin sulfate 1 mg, betamethasone dipropionate 500 mcg, per gram cream (Polyderm 3, Lloyd Laboratories Inc., Malolos, Bulacan) on the wounds, right after the procedure. The rabbits were observed and monitored after 1 hour and 24 hours after conclusion of the procedure for any adverse effects or complications such as recurrence of bleeding, swelling, or mucosal irritation, and donated to a local animal habitat 3 days after the procedure.
Sample Size Computation

Sample size was computed using an online sample size calculator (Select Statistical Services Ltd., Exeter, Devon, UK) to compare 2 means (http://select-statistics.co.uk/sample_size_calculation_two_means). Assuming a 95% confidence level with a power of 80 to detect a mean difference of 30 seconds with a variance of 500 seconds, 10 samples each were needed for the Moringa and control groups.

Statistical Analysis

Data was subjected to the Wilk-Shapiro test for normality (W statistic, 0.842), with computed W statistic of 0.889 (normal distribution). Data was then analyzed manually using the t-test for paired samples.

RESULTS

Ten adult male New Zealand white rabbits with ages ranging from 16 – 20 weeks completed the study. None of the rabbits died or acquired any disease during the time of acclimatization. The weight of the rabbits during the study period ranged from 1.1 – 1.5 kg.

The mean bleeding time for wounds administered Moringa extract was 53 seconds (range, 38 – 70 secs) compared to 159 seconds (range, 100 – 218 secs) for controls. The bleeding time in the former (mean, 53.3 secs) was significantly shorter than in the latter (mean, 159.3), with a mean difference of 106 seconds between the two groups (p = .000019, t-stat = 8.139). No adverse effects or complications were noted.

DISCUSSION

Mucosal healing undergoes a process that follows the stages of fibroplasia, angiogenesis, and reepithelialization. Topical application of Moringa oleifera extract has been shown to promote wound healing. Application of the extract on wounds increases its tensile strength through rapid re-epithelialization and collagen formation. It also increases fibroblast proliferation.

Moringa extract has also been known to act on the blood coagulation cascade. The presence of proteolytic activity is one of the important characteristic features of Moringa oleifera. In a study by Satish et al., both extracts from the leaves and roots of the plant showed proteolytic activity in a dose-dependent manner with the leaf extract exhibiting a significantly higher activity. The study suggested that both extracts exhibited procoagulant activity and reduced recalcification time in clot formation by activation of factors involved in the blood coagulation cascade or by precipitation of the co-factors. Both extracts also exhibited fibrinogenolytic and fibrinolytic activities.

Whether procoagulant activity and reduced recalcification time in clot formation or fibrinogenolytic and fibrinolytic properties of Moringa extract were involved in reducing bleeding time in this study were not established, and can at best be inferred. However, the significant difference in bleeding time between treatment and controls makes a compelling argument to pursue the potential hemostatic effects of Moringa for epistaxis.

Further studies are recommended to determine the histopathologic effects of Moringa extract on nasal mucosa; its toxic dose and approximate effective dose; the effect of tonicity, pH and temperature settings of Moringa extract in the coagulation process; the hemostatic effect of Moringa extract versus such usual topical agents as oxymetazoline and epinephrine; sterilization techniques in the preparation of the extract; before actual clinical trials.

Our study has established that Moringa oleifera leaf extract was associated with significantly shorter bleeding time than saline control in this experimental epistaxis model, and may thus be worth investigating further as a hemostatic agent for epistaxis.

REFERENCES