Addition of Selenium to *Carica papaya* Linn Pulp Extract Enhances Dermal Wound Healing Activity

Nafiu B Abdulrazaq, Haris B Akram, Dinie Najwa Bero, Mohd Yusof bin Mohamad, Izzati Abd Malik and Mohammad Tariqur Rahman*

Department of Biomedical Science, Faculty of Science, International Islamic University Malaysia Jalan Istana, Bandar Indera Mahkota, Kuantan, Pahang 25200, Malaysia

*For correspondence: E-mail: tarique@iium.edu.my; Tel: +6095716 400 ext 2805; Fax: +6095716781

Received: 10 May 2012 Revised accepted: 3 January 2013

Abstract

**Purpose:** To investigate the combined effect of *Carica papaya* extract and mineral elements wound healing.

**Methods:** Phosphate-buffered saline (PBS) and water extracts of unripe *Carica papaya* papaya were analyzed for mineral element composition (Ca, calcium; Mg, magnesium; Na, sodium; Fe, iron; Mn, manganese; Zn, zinc; Ag, silver; and Se, selenium) using a microwave-assisted digestion procedure. Atomic absorption spectrophotometry was used for the analysis of the elements. Phytochemical analysis was conducted using standard protocols. Mice weighing 25 - 30 g were anesthetized, shaved and inflicted with wound at the dorsal region using a biopsy punch. The mice were treated topically twice daily with or without 5 mg/ml PBS or water extracts containing 100 or 200 µM Zn2+ or 1.0 or 0.5 µg/g Se2+. The standard (positive control) group was treated with solcoseryl and the negative control with deionised H2O. Wound area was monitored with a camera and evaluated by software.

**Results:** PBS extract formulated with Se2+. (0.5 or 1.0 µg/g) took approximately 9 days for complete wound healing while negative control took 14 days and other treatment groups least 10 days. PBS extract contained 9 % protein but not tannin while water extract contained 0.58 % protein and tannins. Calcium (137.98 mg/100 g extract) constituted the most abundant macro-element while Se (32.14 mg/100 g extract) was the most abundant microelement in C. papaya pulp.

**Conclusion:** Addition of selenium to C. papaya extracts augments the wound healing activity of the extracts.

**Keywords:** Papaya, Zn, Selenium, dermal wound healing

INTRODUCTION

*Carica papaya* (*L. Caricaceae*) is a lactiferous plant, and produces latex rich in the enzymes, papain (a potent wound healing enzyme) and chymopapain [1]. These enzymes have antimicrobial and antioxidant properties [2,]. C. papaya has also been used in many parts of Africa for the treatment of burns, prevention of burns infections and de-sloughing necrotic tissue in wounds [1]. Traditional uses of papaya latex for wound healing have been confirmed and attributed to its content [3]. It is expected that any chemicals and/or natural bioactive components having either or both antioxidant and cell proliferative effects might have an influence on the wound healing process. Therefore, we have added Se, a well-known antioxidant and Zn, a well-known cell proliferating element [4] to green papaya extract to determine if these added...
substances would enhance the wound healing properties of the extract.

Zinc (Zn) is an important element in the wound healing process as it is a cofactor for the synthesis of structural components and plays an essential regulatory role in cellular function [4]. Se functions within the active site of the antioxidant enzymes. Oral or topical Se supplementation resulted in less skin damage after test mice were irradiated with UV rays, as indicated by reduced inflammation and pigmentation, and lower incidence of skin cancer. A similar study in female human volunteers yielded encouraging results [5].

EXPERIMENTAL

Collection of papaya

The green (unripe) fruits of Carica papaya of Exotica variety were collected during the month of July from a farm at Jabatan Pertanian Tanjung Malim, Perak, Malaysia. The size of the fruits collected were in the range of 15 - 20 cm × 10 - 12 cm. The fruit was identified and authenticated by Dr Nurziana of Herbal Laboratory, Faculty of Pharmacy, International Islamic University Malaysia (IIUM). A voucher specimen (no. Eiium 34) was deposited at the herbarium of the Faculty of Pharmacy, IIUM for future reference.

Preparation of papaya extract

The extract was prepared following the procedure of Anuar and colleagues [4] except that the pulp was used for wound healing studies while the peel and seed extracts were analysed only for mineral element composition. Briefly, the plant material was cut into small pieces and homogenized using either sterile phosphate-buffered saline (PBS) or sterile deionized water (dH₂O). One part by weight of the pulp was added to 3 parts of solvent, crushed in a clean blender and transferred to an incubator shaker at 37 °C for 8 h. The mixture was then centrifuged at 200 g for 30 min at 4 °C. The supernatant containing the extract was collected and freeze-dried to obtain the final extract. Either zinc (SYSTERM®, 100 or 200 mM) or selenium (Acros Organics, 0.5 or 1 µg per mg of extract) dissolved in dH₂O was added to either the water extract (WE) or PBS extract (PE). Both the weight of the pulp (without seeds) and that of the extract were recorded to determine yield.

Evaluation of mineral composition of C. papaya

This investigation was carried out on the elemental content of the seed, pulp and peel extracts of papaya. Both water and PBS extracts of Carica papaya seed, pulp and peel, prepared following the same method as explained in the section, “Preparation of papaya extract” above, were analysed to determine the concentrations of 8 mineral elements, namely: Ca, Mg, Na, Fe, Mn, Zn, Ag, and Se using microwave-assisted digestion using nitric acid-hydrogen peroxide procedure [6] for sample preparation and atomic absorption spectrophotometry (AAS) for analysis. Briefly, 0.3 g of the sample was taken in a 100 mL Teflon vessel and digested with 6 ml of HNO₃ and 1 ml H₂O₂ in microwave digestion system. The temperature program was set at 240 °C for 40 min with pressure of 60 bar applied. The resulting solution was cooled and diluted to 50 ml with deionized water. Analysis was carried out in triplicate.

Phytochemical analysis of the extract

Qualitative analysis of the phytochemical profile of both the PBS and water extracts was carried out to determine the presence or absence of proteins, alkaloids, terpenoids, steroids, anthocyanin, flavonoids, saponins, tannins, while protein content was quantitatively determined.

Protein analysis

Bradford method was used to quantify proteins in the PBS and water extracts [7]. Briefly, the extract (20 µL) was mixed together in a clear plastic 1 mL microcuvette with sodium chloride solution (80 µL) and Bradford reagent (1000 µL) and the absorbance was read at 595 nm wavelength. The resulting reading was compared to the standard solution of bovine serum albumin (Amresco™).

Test for alkaloid (Dragendorff’s test)

The extract (2 mL) was mixed with a few drops of concentrated hydrochloric acid and a few drops of Dragendorff’s reagent in a test tube. Colour change and precipitate formation (if any) were observed; a reddish brown precipitate indicated the presence of alkaloids [8].

Test for terpenoids and steroids (Libermann-Burchard test)

The extract (2 ml) was mixed with 5 drops of acetic anhydride in a test tube and stirred to to mix well. Thereafter, 2 drops of concentrated sulphuric acid were added to the mixture by slow dripping down the test tube wall. Colour change and precipitate formation (if any) were observed. Formation of a brown ring at the junction of two
layers with the upper layer turning green indicates the presence of steroids while formation of a deep red color indicates the presence of triterpenoids [9].

**Cyanidin test**

A piece of magnesium ribbon was dropped into a test tube containing 2 mL of extract, followed by a few drops of concentrated hydrochloric acid. The solution was allowed to mix and settle for 10 min. Colour change and precipitate formation (if any) were observed. A yellow precipitate indicates the presence of anthocyanin [9].

**Tannin test**

The extract (2 mL) was mixed with a few drops of iron (III) chloride solution in a test tube, and colour change and precipitate formation (if any) were observed. Brownish green to blue-black precipitate was indicative of the presence of tannins in the sample [10].

**Test for saponin (Froth Test)**

The extract (0.5 g) was dissolved in 10 mL of distilled water. The test tube was closed with a stopper and shaken vigorously for 30 s. The test tube was allowed to stand in a vertical position and observed over 30 min. It was observed for the presence of “honey comb” froth above the surface of the liquid. Persistence of froth for over 30 min was indicative of the presence of saponin in the sample [9].

**Cardenolides (Keller Kiliani Test)**

The extract (0.5 g) was dissolved in 2 mL of glacial acetic acid containing one drop of ferric chloride solution and 1 mL of concentrated sulphuric acid was added. It was observed for the formation of a brown ring at the interface which is indicative of the presence of a deoxy sugar, a characteristic of cardenolides [8].

**Wound healing studies**

Female imprinting control region ICR mice (25 - 30 g, 7 - 8 weeks old, n = 7) were used to evaluate the effect of papaya extract alone and combined with Zn and Se on wound healing rate. The mice were purchased from Institute of Medical Research, Kuala Lumpur, Malaysia and housed under standard experimental conditions of 25 ± 2 °C, 12 h light/dark cycle and fed on normal pellet diet and tap water ad libitum. A uniform circular epidermal wound was made at the dorsal part of the mice using a 6 mm biopsy puncher. Prior to incision, the mice were anesthetized using intraperitoneal administration of 6 % Nembutal (Ceva Sante Animale™ 0.1 mL/10 g of body weight). The skin was shaved, swabbed with alcohol pad and allowed to dry. The PBS and water extracts of papaya pulp (~10 µL) were applied twice a day. All animal handling and experimental procedures were approved by the institutional animal ethics committee (no. IIUM/305/20/4/10) and carried out in accordance with the Guidelines for the Care and Use of Laboratory Animals of the Faculty of Medicine, International Islamic University Malaysia (IIUM).

Photos of the wounds from the first day of wound induction until the day of complete wound closure (CWC) were taken on alternative days to measure the rate of wound healing and the comparative wound healing efficiency of the extracts. A digital camera (Canon Powershot 5.0 MP) was used for the purpose. Photos were taken with the same settings of distance and aperture, and analyzed for wound surface area using Adobe ® CS3 Photoshop (Extended Version); the data were analyzed using SPSS for Windows version 16.0 software. Percent relative wound healing efficiency (RWHE) of the extracts was calculated to measure the rate of healing of the wound by the extracts compared with the time required for complete healing without any treatment (natural healing). RWHE (%) was calculated according to Eq 1.

\[
\text{RWHE} (%) = \left(\frac{T_N - T_E}{T_N}\right) \times 100
\]

where \(T_N\) = time required for natural wound healing i.e., CWC without any treatment and \(T_E\) the time (number of days) taken to attain CWC for the treated group.

In the current study, solcoseryl and a series of extracts were used: Negative control group was treated with sterile deionised water (NC); positive control group was treated with solcoseryl (PC) The extract-treated groups were PBS extract (PE) alone, water extract (WE) alone, PBS or water extract plus Zn\(^{2+}\) (100µM) or Se\(^{2+}\) (0.5µg/mg) (i.e., PEZ1 or WES1, respectively); and PBS or water extract plus Zn\(^{2+}\) (200µM) or Se\(^{2+}\) (1.0µg/mg) (i.e., PEZ2 or WES2, respectively).

**Statistical analysis**

Statistical analysis was performed using the software SPSS version 17, and the data expressed as mean ± SEM. One-way ANOVA was applied to the data followed by Tukey’s post-hoc test to assess mean differences between the groups at different time intervals. \(P\) value was set at \(p < 0.05.\)
RESULTS

Elemental composition of extracts

The results (Figure 1) showed that Ca is the most abundant mineral element in all the parts of papaya analyzed with the highest content in the aqueous extract of the peel. Similarly, Se constituted the most abundant essential microelement with the highest level in FPE. The highest concentrations of Na, Mn, Mg, Se, and Zn were found in FPE, SWE, FPE, FPE and FEW, respectively. Ag was not detected.

Phytochemical characteristics of the extracts

The extract yield of papaya fruit pulp was 6.70 ± 0.02 % (n = 3) for the PBS extract and 9.10 ± 0.10 % (n = 3). Mean protein content was 9.35 ± 0.09 and 0.58 ± 0.05 % for the PBS and water extracts, respectively. Alkaloids were present in both the PBS and water extracts while tannins were present only in the water extract. All the other phytochemicals analyzed were absent from both extracts (Table 1).

Effect of Se on papaya wound healing properties

In the early stage of treatment (Days 0 - 4), reduction of the initial wound area ranged from 30 to 70 % (Tables 2 and 3). By day 2, WE produced a significant (p < 0.05) reduction (~ 67 %) in wound area when compared to both NC (negative control) and PC (positive control) groups. Addition of Se²⁺ (0.5 µg/mg) to PBS extract (PE) did not significantly influence (p < 0.05) wound healing compared to positive control group. All other treatment groups showed significant reduction in wound area compared to NC but the reduction in wound areas were significantly (p < 0.05) lower compared to PC. At the end of the early stage of wound healing process (Day 4), all treated groups showed significant (p < 0.05) reduction in wound area compared to NC but the reduction was significantly (p < 0.05) lower than for PC except for the WE-treated group which showed no difference.

Table 1: Yield and phytochemical profiles of the extracts

<table>
<thead>
<tr>
<th>Extract</th>
<th>Alkaloid</th>
<th>Terpenoid</th>
<th>Steroid</th>
<th>Anthocyanin</th>
<th>Cardiolide</th>
<th>Saponin</th>
<th>Tannin</th>
<th>%Yield</th>
<th>% of protein</th>
</tr>
</thead>
<tbody>
<tr>
<td>PBS extract (PE)</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>6.7</td>
<td>9.35</td>
</tr>
<tr>
<td>Water extract (WE)</td>
<td>√</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>√</td>
<td>-</td>
<td>9.2</td>
<td>0.58</td>
</tr>
</tbody>
</table>
By day 6, addition of Zn²⁺ (100 µM) or Se²⁺ (0.5 µg/mg) to PE or WE did not significantly influence (p < 0.05) when compared to PC. Other treated groups produced significant reduction (p < 0.05) in wound area compared to NC, although the reductions were less pronounced than by PC. By day 8, all the treated groups showed no significant difference (p < 0.05) in wound area compared to PC except for Zn²⁺ (200 µM)/PE and PE-treated groups. A similar result was obtained at the end of day 10 except that only Zn²⁺ (200 µM)/PE showed significantly higher healing property (p < 0.05) than PC. Furthermore, only PE containing Se²⁺ (0.5 or 1 µg/mg) produced complete wound closure (Tables 2 and 3).

**Effect of addition of Se²⁺ to extract on wound healing**

When the wound was treated with PE containing Se, healing duration (CWC, ~9 days) was significantly shorter (p < 0.05) than for other groups. For the groups given either solcoseryl, WE/100 µM Zn or WE/0.5 µg/mg Se, CWC was 10 - 11 days while for the rest, wound healing was within 12 - 14 days. Compared to negative control group, all the treatment groups reduced significantly (p < 0.05) the duration for CWC. The most efficient healing (38 %), as determined RWHE, was observed in the PE/Se (0.5 µg/mg) (Table 2).

**DISCUSSION**

Papaya has been widely used for therapeutic purposes. The therapeutic effectiveness of green or raw papaya for medicinal purposes is perhaps related to its higher latex content compared with ripe papaya which has a reduced number of laticifer cells [11]. The latex of green papaya contains high amounts of protease group of enzymes namely, papain and chymopapain [1].

Some solvents have been used to extract various phytochemicals from papaya [10,11].

### Table 2: Effect of topical treatment of wound with PBS extract of *C. papaya* containing Zn²⁺ and Se²⁺.

<table>
<thead>
<tr>
<th>Day</th>
<th>NC</th>
<th>PC</th>
<th>PE</th>
<th>PEZ1</th>
<th>PEZ2</th>
<th>PES1</th>
<th>PES2</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>100±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
</tr>
<tr>
<td>2</td>
<td>17.86±8.51*</td>
<td>51.52±3.42*</td>
<td>42.20±2.71*</td>
<td>41.97±3.92*</td>
<td>20.29±5.42*</td>
<td>51.55±2.70*</td>
<td>44.18±6.24*</td>
</tr>
<tr>
<td>4</td>
<td>26.71±10.46*</td>
<td>73.77±0.89*</td>
<td>69.93±0.69*</td>
<td>61.91±4.88*</td>
<td>46.76±3.94*</td>
<td>66.58±3.33*</td>
<td>59.23±6.18*</td>
</tr>
<tr>
<td>6</td>
<td>39.78±12.12*</td>
<td>85.45±2.23*</td>
<td>78.87±5.06*</td>
<td>86.13±2.59*</td>
<td>78.91±5.37*</td>
<td>82.66±2.51*</td>
<td>78.76±4.43*</td>
</tr>
<tr>
<td>8</td>
<td>75.17±7.23*</td>
<td>97.04±0.94*</td>
<td>93.03±5.67*</td>
<td>97.31±0.83*</td>
<td>92.90±3.14*</td>
<td>98.62±1.22*</td>
<td>98.99±0.63*</td>
</tr>
<tr>
<td>10</td>
<td>88.97±3.74*</td>
<td>99.51±0.32</td>
<td>96.70±2.77*</td>
<td>99.77±0.09*</td>
<td>95.20±3.92*</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
</tr>
<tr>
<td>12</td>
<td>94.35±2.69*</td>
<td>100±0.00</td>
<td>98.70±1.30*</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
</tr>
<tr>
<td>14</td>
<td>97.89±1.23</td>
<td>100±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
</tr>
<tr>
<td>16</td>
<td>100±0.00</td>
<td>100±0.00</td>
<td>100±0.00</td>
<td>100±0.00</td>
<td>100±0.00</td>
<td>100±0.00</td>
<td>100±0.00</td>
</tr>
</tbody>
</table>

**Table 3: Effect of topical treatment of wound with *C. papaya* water extract containing Zn²⁺ and Se²⁺.

<table>
<thead>
<tr>
<th>Day</th>
<th>NC</th>
<th>PC</th>
<th>WE</th>
<th>WEZ1</th>
<th>WES1</th>
<th>WES2</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>100±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
<td>0.00±0.00</td>
</tr>
<tr>
<td>2</td>
<td>17.86±8.51*</td>
<td>51.52±3.42*</td>
<td>67.37±8.81*</td>
<td>42.43±6.22*</td>
<td>29.64±7.43*</td>
<td>36.39±5.36*</td>
</tr>
<tr>
<td>4</td>
<td>26.71±10.46*</td>
<td>73.77±0.89</td>
<td>71.23±1.89</td>
<td>66.11±4.23*</td>
<td>55.97±3.92*</td>
<td>54.10±4.12*</td>
</tr>
<tr>
<td>6</td>
<td>39.78±12.12*</td>
<td>85.45±2.23*</td>
<td>81.50±3.86*</td>
<td>87.30±2.62*</td>
<td>75.26±4.35*</td>
<td>84.90±2.83*</td>
</tr>
<tr>
<td>8</td>
<td>75.17±7.23*</td>
<td>97.04±0.94*</td>
<td>97.00±0.42</td>
<td>98.10±0.78</td>
<td>94.29±3.26*</td>
<td>97.30±0.77</td>
</tr>
<tr>
<td>10</td>
<td>88.97±3.74*</td>
<td>99.51±0.32</td>
<td>98.47±2.23*</td>
<td>99.87±0.09*</td>
<td>98.62±0.34*</td>
<td>99.53±3.32*</td>
</tr>
<tr>
<td>12</td>
<td>94.35±2.69*</td>
<td>100±0.00</td>
<td>99.17±0.56*</td>
<td>100.00±0.00</td>
<td>99.47±0.25*</td>
<td>100.00±0.00</td>
</tr>
<tr>
<td>14</td>
<td>97.89±1.23</td>
<td>100±0.00</td>
<td>99.80±0.20</td>
<td>100.00±0.00</td>
<td>99.90±0.10</td>
<td>100.00±0.00</td>
</tr>
<tr>
<td>16</td>
<td>100±0.00</td>
<td>100±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
<td>100.00±0.00</td>
</tr>
</tbody>
</table>

*Negative control group was treated with sterile deionised water (NC); positive control group was treated with solcoseryl (PC); extract-treated groups are PBS extract (PE) alone, water extract (WE) alone, PBS extract plus Zn²⁺ (100µM) or Se²⁺ (0.5µg/mg) (i.e., PEZ1 or WEZ1, respectively); and PBS extract plus Zn²⁺ (200µM) or Se²⁺ (1.0µg/mg) (PEZ2 or WES2, respectively). Values are expressed as mean ± SEM (n = 7); *significantly different compared to positive control rats (p < 0.05); #significantly different compared to negative control rats (p < 0.05).*
PBS and water were used as solvents in the present research and has also been used in the earlier study [3], because of its isotonic or physiological properties which aid in the extraction of protein-like bioactive compounds. The extraction temperature (37°C) was selected for its importance as a physiological temperature and to avoid any possible denaturation of bioactive proteins in the extract which might be important for the wound healing. Tannins were detected in the extract prepared in the current study but neither saponins nor cardenolides were detected which were reportedly present in a previous study [12]. A probable reason for this difference could be different method of extraction, in which the aforementioned study used sun-dried raw papaya flesh while we used papaya flesh without drying. Another possible reason is the difference in papaya varieties used; the previous study used the Hawaiian Solo variety whereas we used the Exotica variety.

The mineral element composition of *C. papaya* has been analyzed in some studies but the results vary widely was and this has been attributed to differences in analytical methods, preparation and storage methods, varieties, place of cultivation, and parts of the fruit analyzed [13]. This, therefore, justifies the present study.

Mineral nutrients play an important role in wound healing. Fibroblasts and keratinocytes require optimal calcium signalling for cellular migration and differentiation leading to re-epithelization during the proliferative phase of wound healing [14,15]. Se, the most abundant micronutrient in *C. papaya*, is an essential cofactor for endogenous antioxidant enzymes which can reduce oxidative tissue damage resulting from neutrophil infiltration and microbial invasion of wounds [16].

During the early phase of wound healing (up to day 2), the water extract without Zn or Se exhibited the highest reduction in initial wound area; this effect was even higher than that obtained with the commercially available wound healing agent tested (solcoseryl) which served as positive control. However, the mean time required for complete wound closure (CWC) by the the water extract is longer than that of solcoseryl. This indicates that the contents of the water extract might have contributed significantly during the early phase of wound healing but not in the latter phases such as tissue regeneration and re-modelling. Notably, the protein content in the water extract was significantly lower than that in the PBS extract. Moreover, the water extract contains tannins, unlike the PBS extract. On the whole, the PBS extract (PE) required less time for the CWC than the water extract. This may be attributed to the higher protein content of PE which might have contributed to higher tissue regeneration, and hence, faster wound closure.

Addition of Se to PE has resulted in the shortest time to achieve CWC. Thus, both the higher protein content and Se might have contributed to the accelerated wound healing. Zn has been reported to aid the wound healing cascade, especially with regards to collagen and protein formation [5]. Inclusion of Zn in the water extract resulted in a shorter time to attain CWC than the water extract without the element. This may be causally linked to the cell proliferating effect of Zn and lower protein content of the water extract. Zn participates in the regulation of cell proliferation in several ways. It is essential to enzyme systems that influence cell division and proliferation [17]. Indeed, the exact mechanism and dynamics of collagen formation due to Zn added to extracts are yet to be determined. However, the addition of Zn to PE, which in itself has high protein content, did not cause further reduction in the time to achieve CWC. This is

**Table 4: Mean time (days) for complete wound healing**

<table>
<thead>
<tr>
<th>Treatment/group</th>
<th>Time (days)</th>
<th>RWHE (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Negative control (NC)</td>
<td>14.3 ± 0.7</td>
<td>-</td>
</tr>
<tr>
<td>Solcoseryl (PC)</td>
<td>10.6 ± 0.4</td>
<td>26</td>
</tr>
<tr>
<td>PBS extract (PE)</td>
<td>12.0 ± 1.2</td>
<td>16</td>
</tr>
<tr>
<td>Water extract (WE)</td>
<td>14.0 ± 1.2</td>
<td>2</td>
</tr>
<tr>
<td>PE + Zn 100 mM</td>
<td>11.1 ± 0.6</td>
<td>22</td>
</tr>
<tr>
<td>PE + Zn 200 mM</td>
<td>12.0 ± 0.1</td>
<td>16</td>
</tr>
<tr>
<td>PE + 0.5µg Se/mg of extract</td>
<td>8.9 ± 0.4</td>
<td>*16, *18</td>
</tr>
<tr>
<td>PE + 1.0 µg Se/mg of extract</td>
<td>9.1 ± 0.4</td>
<td>*16, *18</td>
</tr>
<tr>
<td>WE + Zn 100 mM</td>
<td>10.6 ± 0.3</td>
<td>26</td>
</tr>
<tr>
<td>WE + Zn 200 mM</td>
<td>12.9 ± 0.7</td>
<td>10</td>
</tr>
<tr>
<td>WE + 0.5 µg Se/mg of extract</td>
<td>10.6 ± 0.6</td>
<td>36</td>
</tr>
<tr>
<td>WE + 1.0 µg Se/mg of extract</td>
<td>11.7 ± 0.3</td>
<td>18</td>
</tr>
</tbody>
</table>

**Note:** RWHE = relative wound healing efficiency; values are expressed as mean ± SEM (n = 7); *,**, *** significantly shorter compared to NC, PC, PE and WE, respectively. P-value was set at p < 0.05.
perhaps due to the fact that proliferative effect, either by Zn or any protein in the extract, might have reached a plateau. Additional proliferative effect might require other physiological responses such as antioxidant activity.

The addition of Se was designed to take advantage of its antioxidant properties which aid in the reduction of free radicals. This phenomenon is important when we consider that Se is an important component in the enzyme, glutathione peroxidase. The enzyme debridging mechanisms of papain and chymopapain when combined with the protein synthetic functions of Zn might have resulted in the zinc-containing extracts producing faster wound healing in the late phase. It is noted that none of the mice treated with papaya extracts resulted in either inflammation or pus formation. However, the mice treated with solcoseryl showed pus formation while the mice treated with sterile deionized water (NC) showed inflammation.

Generally, for all groups, the wounds were observed to still be ‘fresh’ on the first day, i.e., pinkish and moist, but turned to a darker colour (forming a scab) and seemed to be drier as the days went by. This was recognized as granulation tissue covered with epithelial cells. The phytochemical profiles of the extracts indicate that water as extracting solvent resulted in the extraction of tannins. Tannins are of special importance in the regulation of vascular endothelia growth factor (VEGF) expression [18] which in turn is responsible for angiogenesis and important for wound healing. The regulation of these growth factors is especially important in the proliferative stage of wound healing which probably explains why the water extract containing 100 μM of Zn resulted in faster wound healing than PBS extract.

The re-modeling phase is the phase in which the collagen replacement occurs. Collagen formation is important as it increases the local tensile strength of the newly-formed skin. This is where it is noted that the zinc/extract treatment groups picked up speed as it has been stated that Zn is responsible for collagen and protein synthesis [5]. Although the Se/extract groups showed faster wound healing overall, they reached this phase slower than the Zn/extract groups but managed to close this gap.

CONCLUSION

Green papaya fruit extract reduced the healing time of dermal wounds in mice. The addition of Se, in contrast to Zn addition, decreased healing time. Thus, incorporation of Se in green papaya fruit extract should be considered a potential therapeutic intervention for wound healing.

REFERENCES

17. MacDonald RS. The role of zinc in growth and cell proliferation. J Nutr 2000; 130(14): 1505S–1508S.