Evaluation of the Anti-ulcer Effect of Extra Virgin Avocado (Persea americana) Oil in Rats

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ABSTRACT

Peptic ulcer disease is a gastrointestinal disorder of high mortality and morbidity. Therapy of this disease condition with conventional drugs presents therapeutic challenge due to incidents of unwanted side effects, and the high cost of these medications. The use of products from natural sources will provide useful alternatives due to their minimal side effects, being inexpensive, and readily available, especially in resource-challenged nations of the continent of Africa and other developing economies. Oils derived from plants are increasingly being used in management of several disease conditions. This study aims at evaluating the antiulcer activity of extra virgin avocado oil in rodents using ethanol-induced, and indomethacin-induced ulcer models. Test group was given extra virgin avocado oil (1ml per 250g weight) for 7 days before induction of ulcer. Positive controls received omeprazole 30 mg/kg, while negative control animals were given distilled water (10ml/kg) for 7 days respectively, before induction of ulcer. The stomachs were excised, and their histopathological examinations carried out. Avocado oil significantly reduced the ulcer index in both models when compared to the negative control group, and histopathological findings corroborate that the oil ameliorates ulcerations in both models. The present study has demonstrated that extra virgin avocado oil possesses significant antiulcer activity.

Keywords: Avocado oil, extra-virgin, peptic ulcer disease, ulcer index.

I. INTRODUCTION

Peptic ulcer disease (PUD) is a disease of the gastrointestinal tract, which comprises both gastric and duodenal ulcers. It is hallmarked by imbalances between offensive (gastric acid, pepsin, and Helicobacter pylori) and defensive (prostaglandins, bicarbonate ions, mucin, growth factors, and nitric oxide) factors [1]-[3]. Other causes of peptic ulcer disease include long term and high dose of drugs such as non-steroidal anti-inflammatory drugs (NSAIDs), smoking, stress, alcohol consumption, age-related decline in prostaglandin levels, and diseases such as Zollinger-Ellison syndrome [4], [5]. Peptic ulcer disease is the most common gastrointestinal disorder with significant mortality and morbidity globally [6]. Synthetic drugs such as proton pump inhibitors, antacids (aluminum hydroxide and magnesium trisilicate), histamine-H₂-receptor blockers (cimetidine and ranitidine), proton pump inhibitors (omeprazole and lansoprazole), cytoprotective agents (sucralfate and the prostaglandin analogue misoprostol), muscarinic antagonists (pirenzepine), and antimicrobial agents (amoxicillin and clarithromycin) are commonly used for treatment of peptic ulcer disease [7], [8]. However, these drugs have numerous side effects such as diarrhoea, headache, constipation, arrhythmia, hypomagnesemia, hypersensitivity, impotence and gynecomastia [9], [10]. Additionally, some of these drugs are expensive [11]. Herbal products are considered better alternatives as they have fewer side effects and better compatibility with the human body [12].

Oils obtained from plants are gaining wide popularity for their therapeutic potentials [13]. Some oils reported to be effective for treatment of peptic ulcer disease include virgin coconut oil [14], olive oil [15], ginger, clove, castor oils [16], and basil oil [17]. Avocado (Persea americana) is a fruit that is native to Central America. It is grown mainly in warm
temperate and subtropical climates throughout the world. Hence, climate and country of origin can affect the fruit quality and therefore, the oil [18]. Extra-virgin avocado oil which is viscous, dark green in colour (due to chlorophylls and carotenoids contents), with a mild taste is obtained from the avocado fruit mostly by cold pressing, and without undergoing alterations in its nature by the addition of chemicals or subsequent processing [19]. Internationally, there are no defined parameters for avocado oil, and the values commonly used are those recommended for olive oil [18]. The quality standard for olive oil is documented in the Codex Alimentarius and the International Olive Oil Council [20]. Avocado oil was classified based on the extraction method and fruit quality by Woolfe and co-workers [21] as “Extra virgin,” which is that produced from high-quality fruit, extracted only with mechanical methods at temperature below 50 °C and without the use of chemical solvents. “Virgin” avocado oil, is produced using lower quality fruits i.e., having small areas of rot and physical alterations, extracted mechanically at temperature below 50 °C, and with no added chemical solvents. The third class is “Pure” avocado oil whose fruit quality is not important. It is bleached, deodorized, and contains natural fruits or herbs flavour. The fourth is “Mixed” avocado oil which combines avocado oil with olive, macadamia, and other oils [18], [21]. The major producers of avocado oil in the world are Mexico, New Zealand, the United States, South Africa, and Chile [22].

Avocado oil has gained wide application in human nutrition, food industry, and cosmetics. The lipid content which consists mainly of monounsaturated fatty acids, has cardiovascular system benefits and anti-inflammatory potentials [18], [23]. Avocado oil contains more than 60% of monounsaturated fatty acids, a characteristic similar to that of olive oil, hazelnut, and macadamia nut. When compared with olive oil, avocado oil possesses a higher proportion of saturated fatty acids (16.4%), mostly palmitic acid (15.7%), a lower proportion of monounsaturated fatty acids (67.8%), most of which is oleic acid (60.3%), and a higher proportion of polyunsaturated fatty acids (15.2%), most of which is linoleic acid (13.7%) [24]. Avocado oil has been reported as having a higher polyunsaturated fatty acid/saturated fatty acid, and higher omega-6/omega-3 ratios than olive oil [25]. The presence of bioactive compounds such as tocopherols, tocotrienols, phytosterols, carotenoids, and polyphenols in avocado oil makes it of particular interest for research, targeted at the prevention and management of several disease conditions [18]. This study aims at the evaluation of antulcer activity of Avocado fruit (Persea americana) oil using rodent models.

II. MATERIALS AND METHODS

A. Preparation of Extra Virgin Avocado Oil

Ten (10) mature, high-quality avocado fruits were bought from Uyo market, in Uyo Local Government Area of Akwa Ibom State, Nigeria. They were washed and kept until soft. The pulp was removed, mashed and spread on a large surface open container kept under room temperature. During this period, the pulpy mash was turned occasionally for faster drying. After about 4 days, droplets of the oil were seen on the surface (this shows its ready for extraction). The oil was squeezed from the paste using sieve cloth and then filtered to remove unwanted particles.

B. Experimental Animals

A total of 36 rats of both sexes, weighing (128-221 g) were obtained from the animal house of the Department of Pharmacology and Toxicology, Faculty of Pharmacy, University of Uyo. They were maintained on standard animal pellets and water ad libitum, housed in cages to acclimatize to the animal house, and maintained under standard conditions (25-28 °C) with 12h dark/12h light cycles.

C. Experimental Design

The experimental rats were divided into three groups consisting of six rats per group. Rats of each group were orally pre-treated as follows:

- **Group 1 (Negative control):** Given distilled water (10ml/kg) for 7 days
- **Group 2 (Positive control):** Given Omeprazole (30mg/kg) for 7 days
- **Group 3 (Treatment group):** Given Extra Virgin Avocado oil (1ml per 250g body weight) for 7 days [26]. Animals were fasted for 24 hours into the 8th day before the ulcer induction. Ethanol and Indomethacin models were used.

D. Ethanol-Induced Ulcer Model

Rats in each group were pre-treated for 7 days and fasted for 24h into the 8th day. On the eighth day, ulcer was induced through the administration of ethanol (2 ml/kg) 1 hour after administration of distilled water (group 1), omeprazole (group 2) and avocado oil (group 3), respectively. The animals were kept further for 4 hours for the ethanol to take effect. After the 4 hours, animals were euthanized by cervical dislocation under chloroform anaesthesia. The stomachs were excised and gently rinsed with normal saline, then inflated and with 1% formalin solution (10 ml), and immersed in the same solution to fix the outer layer of the stomach [1], [27]. After about 10 minutes, each stomach was opened along the greater curvature, rinsed with normal saline to remove gastric contents, and examined by using a 10x magnifier lens to assess the formation of ulcers, then scored by using the Kulkarni method (0 = no ulcer, 0.5 = red coloration, 1 = spot ulcers, 2 = deep ulcers, and 3 = perforations) [28]. This procedure was performed by an expert in identification of ulcer types.

Ulcer index and percentage of ulcer inhibition were determined as follows:

\[
\text{Ulcer index (UI)} = U_N + U_S + U_P \times 10^{-3}
\]

where \(U_N\) = average number of ulcers per animal, \(U_S\) = average of severity score, and \(U_P\) = percentage of animals with ulcers.

\[
\text{Ulcer inhibition (\%)} = \frac{(\text{UI control} - \text{UI test})}{\text{UI control}} \times 100
\]

E. Indomethacin-Induced Ulcer Model

Rats in each group were pre-treated for 7 days and fasted for 24 h into the 8th day. On the eighth day, ulcer was induced...
according to the method of Nwafor and co-workers with modifications [29] via the administration of indomethacin (100 mg/kg) 1 hour after administration of distilled water, indomethacin and avocado oil respectively. The animals were kept further for 4 hours for the indomethacin to take effect. After the 4 hours, the animals were euthanized by cervical dislocation under chloroform anaesthesia for macroscopical examination carried out with a hand lens and scored for the presence of lesions.

F. Histopathology

The stomach tissues which were preserved in 10% formalin pending histopathological studies were dehydrated serially through progressive concentrations of alcohol and cleared using xylene. After clearing, the tissues were embedded in paraffin wax and thin sections of about 5µm were made using the microtome. Each section was mounted on a clean glass slide and stained with Haematoxylin and Eosin. Later, a mounting medium (Canada balsam) was dropped on each tissue section and a cover slip placed on it and allowed to dry [30]. They were examined with a light microscope, and photomicrographs were captured using a Moticam Images plus 2.0 (Motic China Group Ltd.) digital Camera attached to the microscope.

G. Ethical Issues

The procedures were performed according to the guidelines on the use of animals and approved by the Institutional Animal Ethical Committee of the Faculty of Pharmacy, University of Uyo, Nigeria (Ethical Approval No: FPharm/EC/010).

H. Statistical Analysis

Results were expressed as mean ± standard error of mean (SEM). Statistical analysis was carried out using SPSS version 23 statistical software. Statistical significance was determined using one-way Analysis of Variance (ANOVA) followed by Tukey’s post hoc test. Values of p<0.05 were considered to be significant.

III. RESULTS AND DISCUSSIONS

A. Effect of Persia Americana Oil on Ethanol-Induced Ulceration in Rats

The effect of avocado oil on ethanol-induced gastric ulceration is as shown in Table I. Avocado oil pre-treated group significantly (p<0.05) prevented ulcer formation compared to that of negative control. The standard drug, omeprazole also caused reduction in ulcer index compared to the negative control but the variation is not statistically significant(p>0.05). In addition, avocado oil showed better protection against ulcer with a higher percentage inhibition of 39.2% when compared to that of the standard drug, omeprazole with 17.5 % ulcer inhibition. Induction of ulcer by ethanol is widely used in antulcer animal model. Ethanol metabolism in the body releases superoxide anion, and hydroperoxyl free radicals, which are known to be part of the mechanisms of acute and chronic ulceration of the gastric mucosa [31].

B. Effect of Persia Americana Oil on Indomethacin-Induced Ulceration in Rats

The effect of Avocado oil on indomethacin-induced gastric ulceration is as shown in Table II. Rats pre-treated with Avocado oil showed significant (p<0.05) reduction in ulcer index compared to those in the normal and control groups, and have percentage ulcer inhibition of 38. However, this anti-ulcer potential of avocado oil was lower than that of the standard drug, omeprazole which apart from significantly (p<0.05) reducing the ulcer index relative to the control, also has a higher percentage inhibition (59 %). Indomethacin has been shown to exhibit greater gastric damage in rodents when compared to other NSAIDs [31]. Hence, it has become preferable for inducing ulcers [32]. It causes damage to stomach tissue by increasing gastric acid, pepsin activity, and enhances lipid peroxidation and oxidative stress via production of free radicals in mucus [33].

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C. Histopathology of the Stomach

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TABLE I: EFFECT OF VIRGIN AVOCADO OIL ON ETHANOL-INDUCED ULCERATION IN RATS

<table>
<thead>
<tr>
<th>Group</th>
<th>Treatment</th>
<th>Ulcer Index</th>
<th>Percentage Inhibition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal Control</td>
<td>Distilled Water-10 ml/kg</td>
<td>3.83±0.31</td>
<td>-</td>
</tr>
<tr>
<td>Positive Control</td>
<td>Omeprazole-30 mg/kg</td>
<td>3.16±0.31</td>
<td>17.5</td>
</tr>
<tr>
<td>Avocado Oil</td>
<td>1 ml per 250 g body weight</td>
<td>2.33 ± 0.42*</td>
<td>39.2</td>
</tr>
</tbody>
</table>

Values are expressed as mean ±SEM. Significant at p<0.05 when compared to control (n=6).

The results of the histopathological studies on the inner lining of the excised stomachs of the experimental animals, are shown in Fig. 1. A–C displays a representative image of each of the groups of the alcohol-induced ulcer model, while D–F shows the results of the indomethacin-induced ulcer model.
Results from observation of stomach of the various groups established severe ulceration induced by both alcohol and indomethacin (control groups). However, avocado treatment tends to ameliorate the ulcerations in both models, especially in the alcohol-induced model when compared to the control group. Indomethacin tends to be less gastro-toxic as compared to alcohol. Omeprazole, the standard drug also attenuates the toxicity induced by both alcohol and indomethacin, especially in the indomethacin model when compared to the negative control group. The antiulcer potential of avocado oil may be attributable to the bioactive constituents present in avocado fruit. Reference [35] reported the presence of alkaloids, saponins, unsaturated steroids and triterpenoids (Leucoanthocyanins), fats and oils in the ethanolic extract of P. americana. Tocopherols have also been identified in the avocado acetone extract [36]. The oil retains most of the bioactive substances and carotenoids present in the fruit [37].

IV. Conclusion

The present study showed that extra virgin avocado oil has significant anti-ulcer activity. However, further research work should be carried out to elucidate the exact mechanism(s) of action. Since the oil was administered in its natural form without any adulteration, it could be used as a nutritional recipe for ulcer patients (for example, as salad dressing, for stir frying etc.), and as potential candidate for preparation of natural remedies used in mitigating peptic ulcer disease in humans.

Conflict of Interest

Authors declare that they do not have any conflict of interest.

Fig. 1. (A)-Histological photomicrograph of the Avocado treated rat stomach alteration induced with alcohol (AV-A, group 3) stained with H&E techniques showing an abnormal digestive tissue with the gastric pits having wide area of hemorrhagic blood vessel (arrow), gastric glands (Gg) and infiltrating inflammatory cells within the stomach endometrium at magnification (x100). Inference: Moderately ulcerated; (B)-Photomicrograph of the longitudinal section of Control rat stomach alteration induced with alcohol (C-A) showing the histo-structure of an abnormal digestive tissue with the gastric pits having wide areas of hemorrhagic blood vessel (arrow), degenerating glandular epithelial cells (hp), wide infiltration of inflammatory cells (arrowhead) within the stomach endometrium and eroding endothelial lining cells at magnification (x100). Inference: Severely ulcerated; (C)- Photomicrograph of the longitudinal section of the Omeprazole treated rat stomach alteration induced with Alcohol (OM-A) showing the histo-architecture of an abnormal digestive tissue with the gastric pits having mild area of hemorrhagic blood vessel (arrow), hyperplastic glandular epithelial cells (hp) and few infiltrating inflammatory cells (arrowhead) within the stomach endometrium at magnification (x100). Inference: Mildly ulcerated. (D)-Photomicrograph of the longitudinal section of the Avocado treated rat stomach alteration induced with Indomethacin (AV-I) showing the histo-architecture of an abnormal digestive tissue with the gastric pits having scanty areas of hemorrhagic blood vessels (arrow) and infiltrating inflammatory cells within the stomach endometrium at magnification (x100). Inference: Mildly ulcerated; (E)- Photomicrograph of the longitudinal section of the Omeprazole treated rat stomach alteration induced with Indomethacin (OM-I) showing the histo-architecture of an abnormal digestive tissue with the gastric pits having wide area of hemorrhagic blood vessels (arrow), atrophying glandular epithelial cells (At) and infiltrating inflammatory (arrowhead) cells within the stomach endometrium at magnification (x100). Inference: Moderately ulcerated; (F)-Photomicrograph of the longitudinal section of the negative control rat stomach alteration induced with indomethacin (C-I) showing the histo-structure of an abnormal digestive tissue with the gastric pits having wide areas of hemorrhagic blood vessel (arrow), degenerating glandular epithelial cells, wide infiltration of inflammatory cells (arrowhead) within the stomach endometrium and eroding endothelial lining cells at magnification (x100). Inference: Severely ulcerated.

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