Antinociceptive and Anti-Inflammatory Activities of Ethanol Extract of Bryophyllum Pinnatum Laboratory Animals

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Abstract: The ethanol extract of Bryophyllum pinnatum was investigated for possible anti-nociceptive and anti-inflammatory effects in mice and rats was carried out using acetic acid-induced abdominal contractions in mice and formalin-induced hind paw edema in rats. Three doses of the extract (50, 100, and 200mg/kg body weight) were used intraperitoneally for both anti-nociceptive and anti-inflammatory studies. There was a significant effect (P<0.05) for the anti-nociceptive study, the highest activity resides at the lowest dose 50mg/kg body weight of the extract, while for the anti-inflammatory study (P<0.05) the activity resides more at the dose of 100mg/kg. Preliminary phytochemical screening of the extract revealed the presences of alkaloid, triterpenoid, saponin, glycerin, flavonoid, glycerin and steroid. The intraperitoneal LD50 is 774 mg/kg body weight in mice. The results support the local use of the plant in painful and inflammatory conditions.

Key words: Bryophyllum pinnatum Anti-nociceptive; anti-inflammatory; Ethanol, Formalin, Acetic acid.

I. Introduction

Bryophyllum pinnatum was initially named kalanche. It belongs to the family, Crassulaceae. It has many common names such as air plant, life plant, resurrection plant, green love, Miracle leaf. It is widely naturalized in other parts of the world including tropical eastern Africa, Asia (e.g. Taiwan, Indonesia and New Guinea), New Zealand, south-eastern USA (i.e. Florida), the Caribbean, and the Pacific (i.e. the Galapagos Islands, Melanesia, Polynesia and Hawaii) (Biosecurity Queensland, 2011). The plant reproduces both from the seeds and vegetative from then bulbils at the margins[2].

The herbal remedies of Bryophyllum pinnatum have been seen in various culture and still serves as the main means of therapeutic medical treatment and it is also used for all set of respiratory conditions, kidney stones gastric ulcers, skin disorders, promote menstruation[2,3]. In Bagbahera, the indigenes use the fresh leaves juice to repel mosquitoes. They apply the juice on the infants body in order to protect them from mosquitoes bites[4]. Traditional healers in Northern part of Nigeria however claimed that the leaves and root serve effectively as an asthmatic remedy. It was claim to be useful for the treatment of other cardiovascular diseases[5].

This research was aimed at investigating the possible anti-nociceptive and anti-inflammatory activities of ethanol extract of Bryophyllum pinnatum in order to support or refute the claims by traditional herbalists.

II. Materials And Methods

2.1 Location of Study

The research was carried out in the Department of Human Physiology Faculty of Medicine, Ahmadu Bello University, Zaria, Nigeria.

2.2 Drugs

All chemicals and drugs used were obtained commercially and of analytical grade.

2.3 Collection of Plant Material

The plant Bryophyllum pinnatum was obtained from Department of plant science, Faculty of Agricultural science, Ahmadu Bello University, Zaria, Nigeria in the month of June 2012. It was identified at the herbarium unit of Biological Science Department, ABU, and Zaria, Nigeria by Mallam U. Galah, where a voucher specimen No.1838 has been deposited for future reference.

2.4 Extraction of Plant Material

The leaves were collected and dried under shade and ground into powder. The powder (500 g) was macerated in 30% of distilled water and 70% ethanol at room temperature for 24 hours. It was then filtered.
using a filtered paper (Whatman size No.1) and the filtrate evaporated to dryness in water bath at 37°C. A brownish residue weighing 30.5 g was obtained. This was kept in air tight bottle in a refrigerator until use.

2.5. Phytochemical Screening

Standard screening test [6] were employed in the screening the extract for the different constituents.

2.6. Animal Management

Adult Wistar rats of both sexes weighing between 200-250g and adult Swiss albino mice of both sexes weighing between 20-25g were used for the experiment. They were kept in well ventilated room and fed with standard grower mash. Water was given *ad libitum*. This research was carried out in Ahmadu Bello University, Zaria, Nigeria in accordance with the rules governing the use of laboratory animals as accepted internationally.

III. Experimental Procedures

3.1. Acute toxicity study: LD₅₀

This was conducted by using the method of [7]. In the first phase, mice were divided into 3 groups of three and treated with the ethanol extract of the plant at doses of 10, 100 and 1000mg kg body weight intraperitoneally (*i.p.*) They were then observed for 24 hours.

In the second phase, mice were divided into 4 groups of one mouse each and treated with the ethanol extract at doses of 200, 400, 600 and 800 mg kg body weight *i.p.* The LD₅₀ was calculated using the second phase.

3.2. Test for Anti-Nociceptive Study

The Acetic acid induced writhing test in mice as described by [8] was employed. Swiss albino mice were divided into 4 groups of 5 mice each. The first group was given 10 ml/kg of Normal saline *i.p.* and served as control, groups 2, 3 and 4 received 50, 100 and 200 mg of extract per kg of body weight *i.p.* respectively. Thirty minutes later, mice in all the groups were treated with Acetic acid (0.6%/v/v, 1 ml per 100g body weight *i.p.*). Five minutes after Acetic acid injection mice were placed in individual cage and the number of abdominal contractions was counted for each mouse for a period of 10 mins. Percentage inhibition of writhing was calculated using the formula

\[
\text{Inhibition} \ (%) = \frac{\text{Mean number of writhings(control)} - \text{Mean No. of writhings(test)}}{\text{Mean number of writhing (control)}} \times 100
\]

3.3 Tests for Anti-Inflammatory Study

The method used for this test was similar to that described by [9] and modified by [10]. Adult wistar rats were dived into four groups of five rats each. Group 1 received distilled water as control group , group 2 3 and 4 received 50,100 and 200mg/kg extract Thirty minutes after treatment they were administered 50µl of 2.5%  ', solution of formalin, subcutaneously under the subplanter surface of the left hind paw. Oedema was assessed in terms of the difference in linear diameter at the injected paw and its diameter 1, 2, 3, 4 and 5 hours interval after formalin injection.[11]. The degrees of inflammation and percentage inhibition of oedema were evaluated by measuring the paw diameter using vernier caliper [12]

\[
\text{Inhibition} \ (%) = \frac{\text{Mean paw diameter (control)} - \text{Mean paw diameter (treated)}}{\text{Mean paw diameter (control)}} \times 100
\]

3.4 Statistical analysis

Results were expressed as mean ± standard error of mean (SEM). The data were statistically analyzed using the one-way ANOVA to determine whether results in a particular group were significantly different from those in the corresponding control groups. Results were statistically significant when p values is less than 0.05 (P<0.05) as described by [13].

IV. Result And Discussion

4.1 Phytochemical Analysis

The freshly prepared extracts were subjected to preliminary phytochemical screening test for various constituents, which revealed the presence of alkaloids, triterpenoid, saponins, glycerin, flavonoids, glycerin and steroids.
4.2 Acute Toxicity Study (LD<sub>50</sub>)

The sign of toxicity were first noticed after 2-3 hours of extract administration. There was decreased locomotor activity and decrease in sensitive to touch and jerking. Also there was decrease feed intake, tachypnoea and prostrations after 8 hours of extract administration.

The median lethal dose (LD<sub>50</sub>) in mice was calculated to be 774 mg/kg body weight.

4.3 Anti-Nociceptive Study

The extract at doses of 50, 100 and 200mg/kg showed anti-nociceptive when compared to control (Fig 1). The extract decrease the number of acetic acid-induced abdominal constrictions in mice and the values were found to be significant (P<0.05). The activity resides more at 50mg/kg with percentage inhibition of 61.3%

![Fig 1: Anti-Nociceptive Study](image)

4.4 Anti-Inflammatory Study

The result of this experiment shown in Fig 2, showed the extract caused inhibition of formalin-induced oedema over a period of 1, 2, 3, 4 and 5 hours respectively. The effect were not dose-dependent at the doses tested (50,100 and 200mg/kg). The peak inhibitory effects after 5 hours were observed with the dose of 100mg/kg. (34.4%) Result were found to be statistically significant (P<0.05 ) in all the three doses when compared to control group.

![Fig 2: Anti-Inflammatory Study](image)

V. Discussion

The ethanol extract of Bryophyllum pinnatum was found to have shown a significant (P<0.05) anti-nociceptive effect at all the doses tested. Although, the inhibitory effect on the acetic-induced writhings in mice was not dose-dependent, the percentage inhibition at a dose of 50 mg/kg body weight of extract was found to be highest (67.1 %). (Figure 1 ) The abdominal constriction method used in evaluating the activity anti nociceptive according to [8,14] was a very sensitive one and can detect antinociceptive effect of substances at a dose that cannot be detected by other methods, such as the tail- flick test [15] Abdominal constriction responses were found to partly involve local peritoneal receptors[16]. The method has been associated with prostanoids in general, e.g increased levels of PGE<sub>2</sub> and PGF<sub>2α</sub> in peritoneal fluids[17,18] as well as lipoxygenase products by some researchers [19,20] Therefore, the result of the acetic acid-induced writhing strongly suggest that mechanism of action of this extract may be linked partly to lipoxygenases and / or cyclo-oxygenases. The activity demonstrated by the extract might be due to the presence of flavonoids and tannins that were present in the extract. This was supported by other workers who found that flavonoids and tannins were found to have antinociceptive and /or anti-inflammatory activities [21]
The significant (P< 0.05) anti-inflammatory activity exhibited by the extract at all the doses used (50, 100, and 200mg/kg body weight ) against oedema induced by formalin in rats compared to the control group the activity reside more at the dose 100mg/kg body weight with percentage inhibition of 34.4% after 5 hours of extract administered. (Figure 2) The plant might serve as a useful source of anti-inflammatory agent. This anti-inflammatory effect of the extract observed might be due to the presence of flavonoids, in the plant. This was supported by other workers, who found that flavonoids inhibit phosphodiesterases which are involved in cell activation, and their effect depend upon the biosynthesis of protein cytokines that mediate adhesion of circulating leucocytes to the sites of injuries. This was supported by [22] that flavonoids and tannins were found to have anti-nociceptive and/or anti-inflammatory activities.

In conclusion, this study has shown that the ethanol extract of *Bryophyllum pinnatum* does possess significant anti-nociceptive and anti-inflammatory effects in the laboratory animal at the doses tested. The results support the traditional use of this plant in some painful and inflammatory conditions and also suggest the presence of biologically active principles which may worth further investigation and elucidation.

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**References**


