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Antiplasmodial and antiulcer activities of *Melanthera scandens*

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ABSTRACT

Objective: To evaluate the antimalarial and antiulcerogenic activities of leaf extract and fractions of *Melanthera scandens* (*M. scandens*). **Methods:** The crude leaf extract (37–111 mg/kg) and fractions (chloroform, ethylacetate and methanol; 78 mg/kg) of *M. scandens* were investigated for antiplasmodial activity against chloroquine-sensitive *Plasmodium berghei* infections in mice and for antiulcer activity against experimentally-induced ulcers. The antimalarial activity during early and established infections as well as prophylactic was investigated. Artesunate (5 mg/kg) and pyrimethamine (1.2 mg/kg) were used as positive controls. Thin films made from tail blood of each mouse were used to assess the level of parasitaemia of the mice. Antiulcer activity of the crude extract was also evaluated against indomethacin, ethanol and histamine induced ulcers. **Results:** The extract and its fractions dose-dependently reduced parasitaemia induced by chloroquine-sensitive *Plasmodium berghei* infection in prophylactic, suppressive and curative models in mice. These reductions were statistically significant ($P < 0.001$). They also improved the mean survival time (MST) from 9.28 to 17.73 days as compared with the control ($P < 0.01 - 0.001$). The activities of extract/fractions were incomparable to that of the standard drugs *i.e.* artesunate and pyrimethamine. On experimentally-induced ulcers, the extract inhibited indomethacin, ethanol and histamine induced ulcers. These inhibitions were statistically significant ($P < 0.001$) and in a dose-dependent fashion. **Conclusions:** The antiplasmodial and antiulcerogenic effects of this plant may in part be mediated through the chemical constituents of the plant.

1. Introduction

Melanthera scandens (*M. scandens*) (Schumach. & Thonn.) Roberty (Asteraceae) is a perennial herb up to 1 m to 4 m high, with branches quadrangular and scabrid. It distributes geographically in East, West and South Africa[1]. It is known as 'ayara edemerong' by the Ibibios of Akwa Ibom State of Nigeria. The leaves are traditionally used to treat various ailments such as stomach ulcer and sores in Gosomtwi-Atwimakwanwoma district of Ghana[2]. In Nigeria, the leaves are used to treat dysmenorrhoea, diabetes and malaria[3]. It is also used by the Bete people of Issia district of Cote d'Ivoire to treat malaria[4,5]. The antioxidant[6], *in vitro* antiplasmodial[4,5] and antidiabetic[7] activities have been reported. Triterpenoid saponins have been reported in the leaves[8]. In this study, we reported the *in vivo* antiplasmodial and antiulcer activities of *M. scandens* as the plant is traditionally used as remedy for malaria and ulcer.

2. Materials and Methods

2.1. Plant materials

The leaves of *M. scandens* were collected from a bush in Ukup in Ikono Local Government area of Akwa Ibom State. The leaves were identified and authenticated by Dr. Margaret Bassey, a taxonomist in the Department of Botany and Ecological Studies, University of Uyo, Uyo. A voucher specimen of the plant was deposited in the Herbarium of Department of Botany and Ecological Studies, University of Uyo, Uyo.

2.2. Extraction

The plant parts (leaves) were washed and shade-dried for two weeks. The dried leaves were further chopped into small pieces and reduced to powder. The powdered leaf was divided into two parts, one part (1.5 kg) was macerated in 97% ethanol for 72 h to give the crude ethanolic extract while the other part (1.5 kg) was successively and gradiently macerated for 72 h in each of these solvents *i.e.* chloroform, ethyl acetate and methanol to give the corresponding fractions of these solvents. The liquid filtrates were

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concentrated and evaporated to dryness in vacuo at 4 °C using rotary evaporator. The yield of each extract was determined; crude ethanolic extract (3.98%), chloroform fraction (0.50%), ethyl acetate fraction (0.30%) and methanolic fraction (0.78%). The dried crude extract and fractions were stored in a refrigerator at 4 °C until use for the proposed experiment.

2.3. Phytochemical screening

Phytochemical screening of the crude extract was carried out employing standard procedures^[9], to reveal the presence of chemical constituents such as alkaloids, flavonoids, tannins, terpenes, saponins, anthraquinones, reducing sugars, cardiac glycosides and others.

2.4. Animals

Swiss albino mice (17–24 g) and rats (120–145 g) of both sexes were used for these experiments. They were obtained from University of Uyo Animal House. The animals were housed in standard cages and were maintained on a standard pelleted feed (Guinea feed) and water was given *ad libitum*.

2.5. Determination of median lethal dose (LD₅₀)

LD₅₀ of the extract was determined using albino mice. The extract was administered intraperitoneally (i.p.) and the method of Miller and Tainter^[10] was adopted. This involved the administration of different doses of the extract (100 – 1 000 mg/kg) to groups of six mice each. The animals were observed for physical manifestation of signs of toxicity. The number of deaths in each group within 24 h was recorded.

2.6. Parasite inoculation

Each mouse used in the experiment was inoculated intraperitoneally with 0.2 mL of infected blood containing about 1×10^7 *Plasmodium berghei* (*P. berghei*) parasitized erythrocytes. The inoculum consisted of 5×10^7 *P. berghei* erythrocytes per mL. This was prepared by determining both the percentage parasitaemia and the erythrocytes count of the donor mouse and the blood was diluted with isotonic saline in proportions according to both determinations^[11].

2.7. Drug administration

The drugs (artesunate and pyrimethamine), extract and fractions used in the antiplasmodial study were orally administered with the aid of a stainless metallic feeding cannula.

2.8. Evaluation of antiplasmodial activity of the extract and fractions

2.8.1. Evaluation of suppressive activity of the extract and fractions (4–day test)

This test was used to evaluate the schizontocidal activity of the extract, fractions and artesunate against early *P. berghei* infection in mice. This was done as described by Antia *et al*^[12]. 48 mice were randomly divided into eight groups of six mice each. On the first day (D₀), 42 mice were infected with the parasite and randomly divided into various

groups. They were administered with the extract, fractions and artesunate. The mice in group 1 were administered with 37 mg/kg, the group 2 (74 mg/kg) and group 3 (111 mg/kg) of crude extract, groups 4, 5 and 6 were administered with 74 mg/kg of the chloroform, ethyl acetate and methanol fractions, respectively, while group 7 was administered with 5 mg/kg of artesunate (positive control), and 10 mL/kg of distilled water was given to group 8 (negative control) for four consecutive days (D₀–D₃) between 8 am and 9 am. On the fifth day (D₄), thin blood film was made from tail blood. The film was then stained with Leishman's stain to reveal parasitized erythrocytes out of 500 in a random field of the microscope. The average percentage suppression of parasitaemia was calculated in comparison with the controls as follows:

$$\frac{\text{Average \% parasitaemia in negative control} - \text{Average \% parasitaemia in positive control}}{\text{Average \% parasitaemia in negative control}}$$

2.8.2. Evaluation of prophylactic or repository activities of extract and fractions

The repository activity of the extract, fractions and pyrimethamine (daraprim) was assessed by using the method described by Okokon and Nwafor^[11]. The mice were randomly divided into seven groups of six mice each. Groups 1–3 were administered with 37, 74 and 111 mg/kg/day of the extract, respectively, while groups 4–7 were respectively given 74 mg/kg/day of the aqueous and chloroform fractions, 1.2 mg/kg/day of pyrimethamine (positive control) and 10 mL/kg of distilled water (negative control). Administration of the extract/fraction/drug continued for three consecutive days (D₀–D₂). On the fourth day (D₃) the mice were inoculated with *P. berghei*. The parasitaemia level was assessed by blood smears 72 h later.

2.8.3. Evaluation of curative activities of extract and fractions (Rane's test)

This was used to evaluate the schizontocidal activity of the extract, fractions and artesunate in established infection. This was done as described by Okokon and Nwafor^[11]. *P. berghei* was injected intraperitoneally into another 48 mice on the first day (D₀). 72 h later (D₃), the mice were divided randomly into eight groups of six mice each. Different doses of the extract (37, 74 and 111 mg/kg) were orally administered respectively to mice in groups 1–3. 74 mg/kg of chloroform, ethyl acetate and methanol fractions were administered to groups 4, 5 and 6, respectively, 5 mg/kg/day of artesunate to the group 7 (positive control) and group 8 was given 10 mL/kg of distilled water (negative control). The extract, fractions and drugs were administered once daily for 5 days. Leishman's stained thin smears were prepared from tail blood samples collected on each day of treatment to monitor parasitaemia level. The mean survival time (MST) of the mice in each treatment group was determined over a period of 29 days (D₀–D₂₈).

$$\frac{\text{No of days survived}}{\text{Total No of days (29)}} \times 100 = \text{MST}$$

2.9. Evaluation of antiulcer activity

2.9.1. Indomethacin-induced ulcer

Male adult albino rats (150–170 g) were randomly divided into five groups of six rats each. The animals were starved from food and water for 24 h and 2 h, respectively before the commencement of experiment^[13]. They were treated as follows: group 1 (control) received only indomethacin (Sigma, 60 mg/kg p.o. dissolved in 5% Na₂CO₃); groups 2–4 were pretreated with *M. scandens* extract (37, 74 and 111 mg/kg p.o., respectively); group 5 received cimetidine (100 mg/kg p.o. dissolved in 50% Tween 80). 1 h later, groups 2–5 were administered with indomethacin. Animals were killed by cervical dislocation, 4 h after indomethacin administration. The stomachs were removed and opened along the greater curvature. The tissues were fixed with 10% formaldehyde in saline. Macroscopic examination was carried out with a hand lens and the presence of ulcer lesion was scored^[13]. Ulcer index (UI) and preventive ratio (PR) of each of the groups pretreated with extract were calculated using standard methods^[13,14].

2.9.2. Ethanol-induced gastric ulceration

The procedures were similar to that used in indomethacin-induced ulceration except that the negative control group (group 1) received only ethanol (2.5 mL/kg p.o.); groups 2–4 were pretreated with *M. scandens* extract (37, 74 and 111 mg/kg p.o., respectively); and positive control group (group 5) received propranolol (40 mg/kg p.o. dissolved in distilled water). Time of administration of ulcerogen and sacrifice, stomach processing and examination as well as ulcer scoring were similar to that used in indomethacin-induced ulceration.

2.9.3. Histamine-induced gastric ulceration in rats

The procedures were similar to that used in indomethacin-induced ulceration except that the negative control group (group 1) received only histamine acid phosphate (Sigma, 100 mg/kg i.p. dissolved in distilled water)^[15]; and positive control group (group 5) received cimetidine (100 mg/kg p.o. dissolved in 50% Tween 80); groups 2–4 were pretreated with *M. scandens* extract (37, 74 and 111 mg/kg p.o., respectively); group 5 received cimetidine (100 mg/kg p.o. dissolved in 50% Tween 80). Time of administration of ulcerogen (histamine acid phosphate, 100 mg/kg, i.p.), stomach processing and examination as well as ulcer scoring were similar to that used in indomethacin-induced ulceration except that the animals were killed by cervical dislocation 18 h after histamine administration.

2.10. Statistical analysis

Data were expressed as mean \pm standard error of the mean (SEM) and were analyzed statistically using one way ANOVA followed by Tukey–kramer multiple comparison test and values of $P < 0.01$ were considered significant.

3. Results

3.1. Phytochemical screening

The phytochemical screening of the ethanolic extract of the leaves of *M. scandens* revealed the presence of cardiac

glycosides, tannins, saponins, terpenes and flavonoids.

3.2. Acute toxicity

The median lethal dose (LD₅₀) was calculated to be (370.00 \pm 23.33 mg/kg). The physical signs of toxicity included excitation, paw licking, increased respiratory rate, decreased motor activity, gasping and coma which was followed by death.

3.3. Suppressive activity of ethanolic leaves extract and fractions of *M. scandens*

The crude extract demonstrated a dose-dependent chemosuppressive effect on the parasitaemia. These effects were statistically significant relative to the control ($P < 0.001$). The chloroform fraction exerted the highest activity that was comparable to that of the standard drug, artesunate. The chemoinhibitory percentages ranged from 37.60 to 68.83 (Table 1).

3.4. Repository activity of ethanolic leaves extract and fractions of *M. scandens*

The ethanolic crude leaves extract showed a dose-dependent chemosuppressive effect on the parasitaemia. These effects were statistically significant relative to the control ($P < 0.001$) with the chloroform fraction exerting the highest activity (Table 2).

Table 1

Suppressive activity of ethanolic leaves extract and fractions of *M. scandens* on *P. berghei* infection in mice (mean \pm SEM) ($n=6$).

Treatment	Dose (mg/kg)	Parasitaemia	% Chemosuppression
Control (normal saline)	10 mL/kg	25.66 \pm 2.84	–
<i>M. scandens</i> crude extract	37	16.00 \pm 1.40*	37.60
	74	13.00 \pm 2.53*	49.34
	111	10.00 \pm 2.10*	61.00
Methanol fraction	74	16.33 \pm 1.02*	62.46
Ethyl acetate fraction	74	11.33 \pm 2.30*	55.80
Chloroform fraction	74	8.00 \pm 2.10*	68.83
Artesunate	5	7.00 \pm 1.86*	72.75

* $P < 0.001$ as compared with the control.

Table 2

Repository/Prophylactic activity of ethanolic leaves extract and fractions of *M. scandens* on *P. berghei* infection in mice (mean \pm SEM) ($n=6$).

Treatment	Dose (mg/kg)	Parasitaemia	% Chemosuppression
Control (normal saline)	10 mL/kg	21.56 \pm 1.01	–
<i>M. scandens</i> crude extract	37	14.41 \pm 0.32*	33.16
	74	12.88 \pm 0.40*	40.25
	111	10.40 \pm 1.01*	51.76
Methanol fraction	74	12.15 \pm 1.04	43.64
Ethyl acetate fraction	74	13.54 \pm 1.04	37.19
Chloroform fraction	74	11.20 \pm 0.06*	48.05
Pyrimethamine	1.2	8.00 \pm 0.42*	62.89

* $P < 0.001$ as compared with the control.

3.5. Antiplasmodial effect of ethanolic leaf extract and fractions of *M. scandens* on established infection

The extract showed a dose-dependent schizonticidal effect on the parasitaemia. These effects were statistically significant relative to the control ($P<0.001$) (Figure 1 and 2). Though both the extract and its fractions showed a significant dose-dependent MST on established infection ($P<0.001$), the ethyl acetate fraction showed greater protective effect with a much longer MST than other fractions and crude extract but lower than that of the standard drug, artesunate (Table 3).

3.6. Indomethacin-induced gastric ulceration

The extract pretreatment on indomethacin induced gastric ulceration showed a dose-dependent reduction in ulcer indices in pretreated groups relative to control. The reduction was statistically significant ($P<0.001$) compared with control (Table 4). The effect was incomparable to that of the standard drug, cimetidine.

Table 3

MST of mice receiving the various doses of ethanolic leaves extract and fractions of *M. scandens* during established *P. berghei* infections in mice (mean±SEM) ($n=6$).

Treatment	Dose (mg/kg)	MST (days)
Control (normal saline)	10 mL/kg	9.28±0.80
<i>M. scandens</i> crude extract	37	10.68±0.40
	74	12.70±0.25*
	111	17.01±0.40*
Chloroform fraction	74	12.38±0.44*
Ethyl acetate fraction	74	17.73±2.33*
Methanol fraction	74	12.00±0.73*
Artesunate	5	29.83±0.12*

* $P<0.001$ as compared with the control.

3.7. Ethanol-induced gastric ulceration

The extract significantly protected rats from ethanol-induced ulcer (Table 5). There was a significant ($P<0.001$) dose-dependent reduction in the ulcer indices relative to control. The effect was comparable to that of control.

Table 4

Effect of *M. scandens* extract on indomethacin-induced ulcer (mean±SEM) ($n=6$).

Treatment	Dose (mg/kg)	Ulcer indices	Preventive ratio
Control (indomethacin)	60	18.78±1.22	–
<i>M. scandens</i> extract	37	7.46±1.42*	60.27
	74	5.20±0.34*	72.31
	111	4.08±1.38*	78.27
Cimetidine	100	1.25±0.40*	93.34

* $P<0.001$ as compared with the control.

Table 5

Effect of *M. scandens* extract on ethanol-induced ulcer (mean±SEM) ($n=6$).

Treatment	Dose (mg/kg)	Ulcer indices	Preventive ratio
Control (ethanol)	–	5.26±0.55	–
<i>M. scandens</i> extract	37	2.45±0.25*	53.42
	74	2.45±0.17*	53.42
	111	1.32±0.86*	74.90
Propranolol	40	1.41±0.42*	73.19

* $P<0.001$ as compared with the control.

3.8. Histamine-induced ulceration

Administration of the extract significantly ($P<0.001$) reduced histamine-induced gastric ulceration in a dose-dependent fashion compared with control (Table 6).

Table 6

Effect of *M. scandens* extract on histamine-induced ulceration in rats (mean±SEM) ($n=6$).

Treatment	Dose (mg/kg)	Ulcer index	Preventive ratio
Control (histamine)	100	15.33±0.45	–
<i>M. scandens</i> extract	37	4.50±0.15*	70.64
	74	3.64±0.46*	76.25
	111	2.78±1.37*	81.86
Cimetidine	100	1.21±0.24*	92.10

* $P<0.001$ as compared with the control.

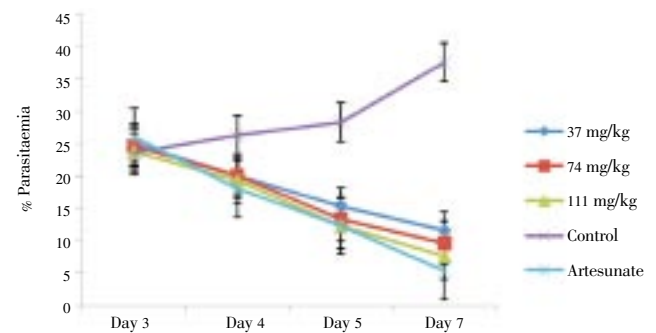


Figure 1. Curative effect of crude leaf extract of *M. scandens* on established infection in mice.

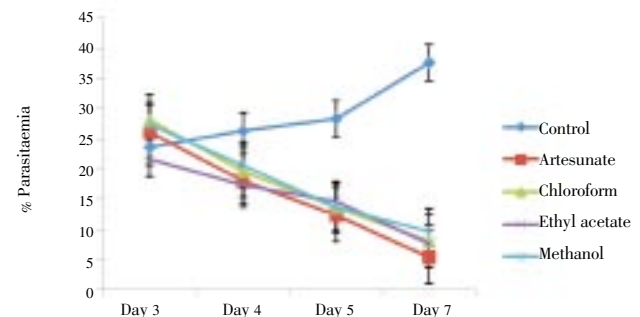


Figure 2. Curative effect of leaf fractions of *M. scandens* on established infection in mice.

4. Discussion

M. scandens leaves are traditionally used to treat various ailments such as stomach ulcer and sores[2], dysmenorrhoea[3] and malaria[4,5]. These prompted the need to evaluate the antiplasmodial and antiulcer potentials of the crude extract, fractions of the leaves of *M. scandens*.

In this work, LD₅₀ was determined to be (370.00±23.33 mg/kg) and the extract was relatively safe[16].

The antiplasmodial properties of the extract and its fractions were investigated using standard models. It was found that both the extract and its fractions significantly reduced the parasitaemia in prophylactic, suppressive and curative models in a dose-dependent fashion. Some secondary metabolites of plants have been reported to have antiplasmodial activity. Among these metabolites are alkaloids, flavonoids and triterpenoids such as limonoids and quassinoids[16]. These compounds (alkaloids, flavonoids

and triterpenoids) present in this plant extract may in part have contributed to the plasmocidal activity of this extract and therefore explained the mechanism of antiplasmodial effect of the extract and its fractions.

The antiulcer activity of the leaf extract was evaluated using indomethacin, ethanol and histamine-induced ulcer models. Indomethacin is known to cause ulcer especially in an empty stomach and mostly on the glandular (mucosal) part of the stomach by inhibiting prostaglandin synthetase through the cyclooxygenase pathway^[13]. Prostaglandins function to protect the stomach from injury by stimulating the secretion of bicarbonate and mucus, maintaining mucosal blood flow and regulating mucosal turn over and repair^[17]. Suppression of prostaglandins synthesis by indomethacin results in increased susceptibility of the stomach to mucosal injury and gastroduodenal ulceration. The extract was observed to significantly reduce mucosal damage in the indomethacin-induced ulcer model, suggesting the possible extract mobilization and involvement of prostaglandin in the anti-ulcer effect of the extract. Administration of ethanol has been reported to cause disturbances in gastric secretion, damage to the mucosa, alterations in the permeability, gastric mucus depletion and free radical production^[13]. It was observed in this study that the extract significantly reduced ethanol-induced ulcer. This may be due to cytoprotective effect of the extract via antioxidant effects. Ethanol is also reported to cause gastric mucosal damage by stimulating the formation of leukotriene C₄ (LTC₄)^[18]. The gastroprotective effect of the extract may in part be due to the suppression, by the extract of lipoxygenase activity. Histamine is known to enhance gastric acid secretion thereby leading to ulceration. Inhibition of histamine-induced ulcer by the extract may be due to its suppression of histamine-induced vasospastic effect and gastric secretion. The leaf extract has been found to contain flavonoids, terpenes, saponins, alkaloids and cardiac glycosides among others. Flavonoids have been reported to protect the gastric mucosa from damage by increasing the mucosal prostaglandin content, inhibiting histamine secretion from mast cells, inhibition of histidine decarboxylase and free radical scavenging ability^[13,19]. This activity corroborates the antioxidant activity of the leaf of *M. scandens* earlier reported by Adeseguna^[6] and maybe attributable to its antiulcer potential.

The results of this study demonstrated that *M. scandens* possesses considerable antiplasmodial and antiulcer activities. These confirm its use to treat malaria and ulcer in folkloric medicine. Therefore, it would be interesting if the active principle is isolated, identified and characterised.

Conflict of interest statement

We declare that we have no conflict of interest.

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