Fertility Parameters of Male and Female Rats Simultaneously Treated with Aqueous Extract of *Eremomastax speciosa*

B. Nchegang1, P.A. Amang2, C. Mezui3, Z.E. Nkwengoua4, P.V. Tan5,*

1Department of Fisheries and Aquatic Ecosystems Management, Institute of Fisheries and Aquatic Sciences at Yaoundé, University of Douala, P.O. Box 2701 Douala – Cameroon.
2Department of Biological Sciences, Faculty of Science, University of Maroua, P.O. Box 814, Yaoundé, Cameroon.
3Department of Biological Sciences, Higher Teachers’ Training College, University of Yaoundé I, P.O Box 047: Yaoundé, Cameroon.
4Department of Organic Chemistry, Faculty of Science, University of Yaoundé I, P.O Box 84, Yaoundé, Cameroon.
5Department of Animal Biology and Physiology, Faculty of Science, University of Yaoundé I, P.O Box 812, Yaoundé, Cameroon.

**Article history:**
Received 25 June 2020
Accepted 17 July 2020
Available online 24 July 2020

**Keywords:**
*Eremomastax speciosa*
Reproduction
Fertility
Toxicity
Bat

**Abstract**

*Eremomastax speciosa* is widely used in ethnomedicine for various purposes, but the possible toxic effects on fertility parameters following sub-acute intake have not been studied. 36 immature male and 36 adult female rats were each divided into four groups of nine rats (Control, Extract 250, 500, 1000 mg/kg). Males received extract for 60 days and females for 14 days. From day 15, a fertility study was done by cohabiting one male with one female from each corresponding group for 10 days. In males, plant extract (250-1000 mg/kg) reduced testicular and spleen weights, increased epididymis weights (250-500 mg/kg; p<0.001) and increased spermatozoa density and motility (at 250 mg/kg). In females, the extract did not influence number of uterine resorptions and number of live offspring compared with controls. Red blood cell count significantly increased (p<0.01) in males and significantly decreased (p<0.05) in females compared with controls. Simultaneous treatment of male and female rats with the extract of *E. speciosa* does not negatively affect reproductive performance. *E. speciosa* may be safely used by reproducing couples at the dose of 250 mg/kg for the management of various disease conditions without the risk of reproductive toxicity.

1. Introduction

The use of medicinal plants as sources of medicine has gone on for several hundred years in many cultures of the world. In recent decades the acceptance and use of plant products as alternative therapy in place of synthetic modern medicines has witnessed a dramatic rise [1]. Top among the various reasons advanced for this turn of events is the debatable assertion that *natural* products are harmless, and that conventional medicines can be ineffective due to side effects and inefficient therapy [2]. A historic review of the use of minerals, plant and animal products as sources of drugs is provided by De Pasquale [3]. Although the WHO considers phytotherapy in its health programs, strict guidelines and basic procedures for the validation of drugs from plant origin in developing countries have been issued, and eastern countries, such as China and India, have developed well-established herbal medicines industry. In the same vein, Latin American countries have been investing huge resources in research programs on medicinal plants and the standardization and regulation of phytomedicines, following the example of European countries, such as France and Germany. In North America, the provision of information on efficacy and safety of phytomedicines products sold as “health foods” is a mandatory requirement for their registration by the FDA [2]. These measures and cautious approaches underline the importance of standards and safety considerations for the use of plant-derived products in health care systems. Thus, in addition to the systemic toxicity (acute, sub-acute, sub-chronic), and local toxicity tests, the WHO recommends that special toxicity tests for mutagenicity, and additional toxicity tests such as carcinoagenicity, teragenicity and reproduction studies be carried out especially when deviations from traditional use (new uses, new routes or more prolonged administration) are contemplated [4].

*Eremomastax speciosa* (Hochst.) Cufod. (Acanthaceae) is widespread from West Africa through Central African Republic and N. Congo-Kinshasa to S. Sudan and S.W. Ethiopia, Madagascar. The plant is widely distributed in tropical Africa and is the only species of the genus *Eremomastax* (syn. *Paulownia helmia* (Lindau), and *Ruellia* (S. Moore)) [5]. The robust, polymorphous shrub is commonly referred to in Cameroon as ‘blood plant’ due to its reputed use in the treatment of cases of anemia. Its widely-claimed anti-anemic activity has been experimentally demonstrated by workers Okoko et al. [6] who also showed anti-microbial actions against pure clinical cultures of *Staphylococcus aureus*, *E. coli, Candida albicans* and *Aspergillus niger*. It is also used in Cameroonian ethnomedicine for the treatment of various stomach complaints and information from traditional practitioners suggested that it possesses antiallergic effects. The leaf aqueous extract has antimicrobial activity [7]. The Douala peoples of Cameroon employ *E. Speciosa* variously for malaria, kidney pain, scabies, anemia, diabetes, and nerves pain [8]. The plant has been cited [9] for the treatment of menstrual pains, gonorrhea, appendicitis and dry burns; as an anti-poison, and to increase and purify blood in the mount Cameroon region. Comparative phytochemical screening of *E. speciosa* revealed the presence of flavonoids, alkaldoids, phenols, tannins, terpenes and saponins [10]. *E. speciosa* has also been cited for its local use in the treatment of male infertility in the west region of Cameroon [11], for the treatment of male infertility among the *Ifo Nkri* People of Akwa Ibon State, Nigeria [12], as well as for the treatment of irregular menstruation by the Agaumba-Bamumbu peoples of the Lebialem highlands in the South West Region of Cameroon [13]. The medicinal importance has also been documented in the Cameroononian and Ghanaian Government/WHO-sponsored ethnobotanical surveys [14, 15]. The antioxidant activity of the extracts includes cytoprotection [16], anti-cholesterin and antiishtaminergic actions, as well as healing actions against chronic gastric ulcers [17]. Since the extract of *E. speciosa* is widely used to treat many conditions, it is important that possible secondary effects related to reproductive physiology are taken into consideration during prescription. In the present experiment we studied the effects of the aqueous extract of *E. speciosa* on the reproductive performance and fertility parameters of male and female rats that were given the extract simultaneously.

*Corresponding Author: verdzekowsno@yahoo.com* (P.V. Tan Paul)

https://doi.org/10.30799/jnpr.087.20060103
2455-0299 / JACS Directory ©2020. All Rights Reserved

2. Experimental Methods

2.1 Experimental Animals

Male immature (35 - 40 days) and female adult (75 days) albino Wistar rats (70-75 g and 110-125 g, respectively) raised in the animal house of the Animal Physiology Laboratory, Faculty of Science, University of Yaoundé 1, were used. The animals were maintained in a well-ventilated room with a 12:12 hour light/dark cycle at room temperature. They were fed a standard laboratory diet (supplied by SPC Ltd, Bafoussam, Cameroon) and given tap water ad libitum. Prior authorization for the use of laboratory animals in this study was obtained from the Cameroon National Ethics Committee (Reg. No. FWA-IRB0001954). The use, handling and care of animals were done in adherence to the European Convention (Strasbourg, 18.III.1986) for the protection of vertebrate animals used for experimental and other purposes (ETS-123), with particular attention to Part III, articles 7; 8 and 9.

2.2 Plant Material and Preparation of the Aqueous Extract

The plant material (Eremomastax speciosa) (Fig. 1) was collected from the outskirts of Yaoundé between August and September 2014 and identified at the National Herbarium, Yaoundé, where voucher specimen No. HNC/136984 has been deposited. The aqueous extract of E. speciosa was prepared by infusing 560 g of the ground powder in 5 liters of boiled water for 15 minutes. After filtration through Whatman filter paper No. 3, the filtrate was evaporated at 40 °C using a Raven convection air oven (Jencons-PLS, UK). The brownish solid obtained (10% yield) was stored at 4 °C. The crude extract dissolved readily in distilled water which was used as vehicle in the subsequent experiments.

Fig. 1 Photograph of Eremomastax speciosa

2.3 Experimental Protocol

The protocol employed was the segment 1 study procedure (administration of test substance prior to and in the early stages of pregnancy) recommended for reproductive and development toxicity [4]. Thirty-six immature male rats were randomly divided into four groups of nine rats each. Group I served as the control and received distilled water (2.5 mL/kg of body weight) while groups II, III and IV were treated orally with 250, 500 and 1000 mg/kg of body weight of the aqueous extract of E. speciosa, respectively, for 60 days (the duration of treatment depends on the duration of spermatogenesis which is 54 days in rats). Thirty-six adult female rats were also randomly divided into four similar groups of nine rats each and given the extract (2; 250, 500, 1000 mg/kg of body weight) orally for 14 days (beginning 47 days from the start of male rats’ treatment). At the end of the treatment period (60 days for the males and 14 days for the females) a fertility study was done by cohabiting one male with one female from each corresponding group for 10 days. The females were examined regularly and the presence of a vaginal plug was taken as a positive indicator for mating and considered the first day of pregnancy. Extract administration for the males continued until successful copulation and administration for the females continued during mating and after successful copulation until the beginning of organogenesis. During the experimental period, mortality was recorded, general signs were noted and body weights and food intake were measured. After successful copulation, the males were sacrificed and autopsied. The testis, ovaries or uterine and kidneys were removed and weighed, relative organ weights (ROW) were calculated as:

\[
\text{ROW} = \frac{\text{absolute organ weight (g)}}{\text{body weight on the day of sacrifice (g)}} \times 100, \text{ and samples were preserved in 10% neutral buffered formaldehyde solution for subsequent histological analysis.}
\]

2.3.1 Fertility Parameters

The females were autopsied on the 12th day of gestation and examined for the number of ovarian corpora lutea, number of successful pregnancies and number of uterine resorptions. Non-gravid uteri were subjected to ammonium sulphide staining for confirmation of non-pregnant status and number of corpora lutea and resorptions were noted. In addition, gross examinations of the organs and tissues were carried out. The left cauda epididymis of each male rat was split into 3 mm pieces in a petri dish containing 10 mL of NaCl (0.9%). The solution was incubated at 37 °C for 10 min, and an aliquot observed under a light microscope (10x mag.) for sperm motility. Sperm counts (millions/mL of suspension) were made using a haemocytometer. A separate aliquot of sperm was stained in 1% eosin and a glass slide smear examined for spermatozoa morphological deformities.

2.3.2 Haematology and Organ Biochemistry

The animals were weighed and sacrificed using an overdose of ether. Each rat was opened up surgically and blood samples were drawn by cardiac puncture. Blood was collected into tubes with and without ethylenediamine tetra-acetic acid (EDTA) as anticoagulant. Two millilitres of blood were collected into tubes with EDTA. Red blood cell count (RBC), white blood cell count (WBC), differential leucocyte count (lymphocyte, monocyte, granulocyte), platelets (PLT), haemoglobin (HGB), hematocrit (HCT), mean corpuscular haemoglobin concentration (MCHC), mean corpuscular volume (MCV), distribution index of red blood cells (RDW), standard deviation of red blood cell distribution index (RDW-S), mean platelet volume (MPV), procalcitonin (PCT), and platelet distribution index (PDW) were determined using an automatic analyzer (Hospitex Diagnostics Hema Screen 18).

2.3.2.1 Hematological Parameters

Hematological analyses were performed on total blood collected in tubes with EDTA. Red blood cell count (RBC), white blood cell count (WBC), differential leucocyte count (lymphocyte, monocyte, granulocyte), platelets (PLT), haemoglobin (HGB), hematocrit (HCT), mean corpuscular haemoglobin concentration (MCHC), mean corpuscular volume (MCV), distribution index of red blood cells (RDW), standard deviation of red blood cell distribution index (RDW-S), mean platelet volume (MPV), procalcitonin (PCT), and platelet distribution index (PDW) were determined using an automatic analyzer (Hospitex Diagnostics Hema Screen 18).

2.3.2.2 Dosage of Serum Cholesterol, Proteins and Transaminases

Biochemical analyses were performed on serum obtained after centrifugation of total blood without anticoagulant at 2000 rpm for 15 min. The analysis of transaminase activities (ASAT, ALAT) and total protein and total cholesterol, were estimated in serum using Commercial kits (Fortress and GCM). Enzymatic activities and atherogenic index were calculated as described by various authors [18-20].

2.3.2.3 Dosage of Tissue Proteins and Cholesterol

Biochemical analyses were performed on homogenates obtained after homogenization and centrifugation of tissue samples (testes, ovaries or uterus) at 2000 rpm for 20 min. and stored at −20 °C. The analysis of tissue protein and total cholesterol in homogenates were done using commercial kits (Fortress and GCM). Atherogenic index was calculated as described by authors Gornal et al. and Youmbissi et al. [19, 20].

2.3.3 Histological Processing

The procedure described by Akpantah et al. [21] was essentially followed. The organs (testes and ovaries initially preserved in 10% neutral buffered formaldehyde solution) were transversely cut in slabs of 0.5 cm thick and fixed in Bouin solution for a day after which it was transferred to 70% alcohol for dehydration. The tissues were passed through 90% alcohol and chloroform for different durations before they were transferred into two changes of molten paraffin wax for 20 minutes each in an oven at 57 °C. Serial sections were cut using a rotary microtome at 5 microns. Slides were prepared from these tissues. The slides where dewaxed and passed through absolute alcohol (two changes); 70% alcohol and then to water for 5 minutes. The slides where then stained with haematoxylin.

2.3.4 Statistical Analysis

The results were analyzed using the one-way ANOVA followed by the Student-Newman-Keuls posttest for comparison of treatment means. P values ≤0.05 were considered significant. Values in tables are given as mean ± standard error for the mean (SEM).
3. Results and Discussion

3.1 Effect of Extract on Body Weight Gain and Relative Organ Weights

The results showed that the rats that received the aqueous extract of \textit{E. speciosa} (250 - 1000 mg/kg) had significantly higher body weights at the end of the experimental period compared with the controls, even though food intake reduced progressively with time for all the treatments (Figs. 2 and 3).

3.2 Effect of Extract on Body Weight Gain and Relative Organ Weights

3.3 Effect of Extract on Haematological Parameters

Tables 1 and 2 show the effects of the extract on vital and reproductive organ weights of females and males respectively. In the male rats treated with the extract for 60 days, there was a significant dose-dependent increase in the relative weights of spleens and epididymis, and a significant dose-dependent reduction of testicular weights. In the female rats treated for 20 days (14 days and up to the beginning of organogenesis), there was a significant dose-dependent drop in relative weights of ovaries, with a drop up to 50-60% decrease at the highest dose of extract compared with the controls. Relative weights of spleen increased while liver weights decreased significantly (p<0.01) up to the 500 mg/kg dose.

Table 3 Sperm characteristics of rats treated (60 days) with the aqueous extract of \textit{E. speciosa}.

<table>
<thead>
<tr>
<th>Treatment group (mg/kg)</th>
<th>Motility (%)</th>
<th>Sperm counts (million/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>82.4±7.6</td>
<td>8.98±0.5</td>
</tr>
<tr>
<td>250</td>
<td>80.0±7.9</td>
<td>10.25±2.1</td>
</tr>
<tr>
<td>500</td>
<td>78.6±7.2</td>
<td>11.75±2.8</td>
</tr>
<tr>
<td>1000</td>
<td>75.7±6.6</td>
<td>12.66±1.1</td>
</tr>
</tbody>
</table>

Values are expressed as Mean ± SEM of 9 rats per group. Statistically significant relative to the controls, p<0.05.

Table 4 Effects of aqueous extract of \textit{E. speciosa} on fertility parameters of female rats.

<table>
<thead>
<tr>
<th>Fertility parameters</th>
<th>Dose (mg/kg of bw)</th>
<th>Number of implantations</th>
<th>Number of fetuses (day 12)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
<td>6.00±0.26</td>
<td>6.55±1.40</td>
</tr>
<tr>
<td></td>
<td>250</td>
<td>4.11±1.34</td>
<td>5.00±1.01</td>
</tr>
<tr>
<td></td>
<td>500</td>
<td>1.33±1.00</td>
<td>3.33±0.64</td>
</tr>
<tr>
<td></td>
<td>1000</td>
<td>1.89±0.77</td>
<td>2.44±0.68</td>
</tr>
</tbody>
</table>

Values are expressed as Mean ± SEM of 9 rats per group. Statistically significant relative to the controls, p<0.05.

3.3 Effect of Extract on Haematological Parameters

Tables 5 and 6 show the effects of the extract on haematological parameters. Extract treatment of males for 60 days resulted in a significant increase in lymphocytes, red blood cells, haematocrit, haemoglobin, blood platelets and mean corpuscular haemoglobin concentration. Maximum effects of extract were obtained with the 500 mg/kg dose although effects at 1000 mg/kg were still higher than in controls (Table 5). The only significant effects of the extract on female rats for 20 days were the drop in red blood cell counts and haematocrit for all the doses compared with the controls. As in the male rats, the 500 mg/kg dose had more significant effect compared with the 1000 mg/kg dose (Table 6).

Table 5 Effects of aqueous extract of \textit{E. speciosa} on haematological parameters in male rats.

<table>
<thead>
<tr>
<th>Haematological parameters</th>
<th>Treatment group (mg/kg)</th>
<th>White Blood Cells</th>
<th>Lymphocyte</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
<td>5.18±0.75</td>
<td>0.15±0.02</td>
</tr>
<tr>
<td></td>
<td>250</td>
<td>4.92±1.03</td>
<td>0.19±0.02</td>
</tr>
<tr>
<td></td>
<td>500</td>
<td>6.81±1.12</td>
<td>0.29±0.02</td>
</tr>
<tr>
<td></td>
<td>1000</td>
<td>5.68±0.85</td>
<td>0.39±0.01</td>
</tr>
</tbody>
</table>

Values are expressed as Mean ± SEM of 9 rats per group. Statistically significant relative to the controls, p<0.05.

Table 6 Effects of aqueous extract of \textit{E. speciosa} on haematological parameters in female rats.

<table>
<thead>
<tr>
<th>Haematological parameters</th>
<th>Treatment group (mg/kg)</th>
<th>Red Blood Cells</th>
<th>Haematocrit</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0</td>
<td>5.98±0.45</td>
<td>0.38±0.02</td>
</tr>
<tr>
<td></td>
<td>250</td>
<td>5.66±0.75</td>
<td>0.32±0.01</td>
</tr>
<tr>
<td></td>
<td>500</td>
<td>6.54±1.01</td>
<td>0.35±0.01</td>
</tr>
<tr>
<td></td>
<td>1000</td>
<td>7.78±0.77</td>
<td>0.43±0.02</td>
</tr>
</tbody>
</table>

Values are expressed as Mean ± SEM of 9 rats per group. Statistically significant relative to the controls, p<0.05.
Effects of aqueous extract of *E. speciosa* on haematological parameters in female rats

<table>
<thead>
<tr>
<th>Haematological parameters</th>
<th>Treatment groups (mg/kg)</th>
<th>0</th>
<th>250</th>
<th>500</th>
<th>1000</th>
</tr>
</thead>
<tbody>
<tr>
<td>RBC (x10^6/μL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>9.82±2.11</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lymphocytes (x10^3/μL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5.92±1.51</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mid cells total (x10^3/μL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20.0±0.55</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Granulocytes (x10^3/μL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>2.49±1.04</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Red blood cells (x10^6/μL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5.92±1.51</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Haemoglobin (g/dL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>13.9±0.25</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>RDW (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>12.9±0.25</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>MPV (fL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>9.82±2.11</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PLT (x10^9/L)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>5.92±1.51</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Values are expressed as Mean ± SEM of 9 rats per group. Statistically significant relative to the controls, *p<0.05; **p<0.01; ***p<0.001; RDW: distribution index of red blood cells; MCH: mean red blood cell hemoglobin content; MCHC: mean corpuscular concentration in hemoglobin; RDW-SD: standard deviation of red blood cell distribution index; MPV: mean platelet volume; PCT: procalcitonin; PDW: platelet distribution index; PLT: blood platelets.

3.4.3 Effect of *E. speciosa* Extract on Transaminase Activity (u/L)

*E. speciosa* aqueous extract induced changes in the activity of ALAT and ASAT. However, at a dose of 500 mg/kg, the extract caused a significant decrease (p<0.05) ASAT (42.69 ± 4.36) in males compared to control (56, 93 ± 2.94). Similarly, ASAT decreased in females that received the extract at doses of 250, 500 and 1000 mg/kg and ASAT decreased in males at 250 mg/kg compared to the corresponding controls. On the contrary, ALAT increased (p > 0.05) in treated females and males treated at doses of 500 and 1000 mg/kg compared to the corresponding control groups (Table 9).

Table 9 Effect of *E. speciosa* extract on transaminase activities (U/L) in male and female albino rats

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Male</th>
<th>Female</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum</td>
<td>ALAT</td>
<td>ASAT</td>
</tr>
<tr>
<td>0</td>
<td>56.93±2.86</td>
<td>61.91±5.17</td>
</tr>
<tr>
<td>250</td>
<td>27.99±4.38</td>
<td>36.26±5.68</td>
</tr>
<tr>
<td>500</td>
<td>24.96±5.68</td>
<td>35.74±3.33</td>
</tr>
<tr>
<td>1000</td>
<td>20.25±3.54</td>
<td>24.90±4.7</td>
</tr>
</tbody>
</table>

Values are expressed as mean ± SEM, n=9.

3.4.4 Effect of *E. speciosa* Extract on Histology of the Testes and Ovaries

Fig 4 shows the histological parameters in the male rats treated with the extract. The testes (A) show interstitial edema (Oed) with a high spermatocyte density (250 mg/kg of bw). The animals that received the high dose extract (500-1000 mg/kg of bw); have the less developed seminal epithelium (Es).

In control and treated rats, we observed growing follicles (F) in the cortical zone of the ovary (B). In the treated rats, there is noticeable desquamation (dq) of the cells and the presence of follicular atresia (Af) (Fig. 5).

3.4.4 Effects of the extract on testicular and ovarian histology of male rats (x 200)

Oral treatment of the rats with *E. speciosa* extract revealed no evident signs of toxicity in male rats. There were no significant changes in daily body weights, liver, kidney and reproductive organ weights. Significant declines in organ and body weights usually correlate to the impairment of reproduction [22]. The body weights of the male rats increased consistently during the 60 days of extract administration. In addition, no toxicity-related behavioral signs were observed. However, rats given the 500 and 1000 mg/kg of extract showed signs of diarrhea earlier in the experiment but quickly recovered during the succeeding days. Similar findings were recorded during the assessment of reproductive toxicity in

https://doi.org/10.30799/jipps.087.20060103

female rats. Fetal morphology was not adversely affected, indicating that *E. speciosa* extract might not affect fetal development if administered before conception.

Female reproduction is functionally controlled by normal estrogen levels which usually peak during the estrous phase of the reproductive cycle. In the female rats, both ovarian and uterine cholesterol and protein levels increased with administration of *E. speciosa* extract, while serum levels decreased especially at the 250 mg/kg dose. The high affinity of the uterus and ovary for cholesterol and proteins during the reproductive cycle would be responsible for the observed drop in serum concentrations.

This suggests that *E. speciosa* extract may have estrogenic effects in the ovary. The extract had no significant effects on male testicular and serum cholesterol and total protein levels except for the observed serum cholesterol spike observed for the 500 mg/kg of bw dose. The significant drop in ovarian weights at the highest dose of extract indicates toxic effects that could be due to extract-induced degeneration of the follicular walls. Ovarian histology shows follicular atresia.

Following treatment of female rats for 14 days, the mean number for corpora lutea increased and the number of resorptions decreased. In addition, copulation and fertility rates increased. These changes suggest that *E. speciosa* extract may have beneficial effects on female reproductive function.

At the doses employed, there were no adverse effects on the mean weights of liver, kidney, heart, lungs and spleen which were consistent with previous studies. Also, there were no treatment-related abnormalities in the body weights of the female rats throughout the period of treatment. Body weight is well known to play an important role in the regulation of gonadotrophin secretion and plays a crucial role for regular cyclic function [23, 24].

The significant decrease in the testicular weights of male rats treated with the extract for 60 days is not readily explainable. The results of testicular histology corroborate those of their relative weight. In the extract-treated rats, interstitial edema and a small thickness of seminiferous epithelium were observed. Extract treatment resulted in increased sperm count and motility at the dose of 250 mg/kg. This effect may occur through increased activity of lactate dehydrogenase activity which has been reported to be related to sperm counts and motility [25] as a result of its involvement in the energy-supply metabolic processes. The extract may therefore contain phytochemicals which may improve spermato genesis at the level of the germinal epithelium. Increases in sperm motility caused by chemical agents had earlier been reported to be due to their ability to permeate the blood-testis diffusion barrier [26]. At higher doses, there was a drop in sperm motility which could be accounted for by the observed dose-dependent increase in the number of deformed spermatozoa. There were no significant decreases in testicular cholesterol levels (precursor of testosterone) to support the possible implication of reduced androgenic effects by the extract. Testosterone is required for the growth, development and maintenance of male reproductive organs [27] and in association with follicle stimulating hormone (FSH), acts on the seminiferous tubules to initiate and maintain spermatogenesis [28]. However, these potential adverse effects of extract at high doses did not have negative effects on male sexual performance and fertility rates.

In the present study, there were significant increases in erythrocytes values in the male rats given the extract of *Eremomastax* for 60 days, suggesting that the extract would confer good respiratory capability to the animals during sexual function. The corresponding significant increase in the hematocrit of extract-treated animals confirms the widely reputed use of the extract in the treatment of cases of anaemia [6]. The mean values of RBC, WBC, and hematocrit in rats given the extract were within the normal physiological ranges of 5.79 - 7.14 x 10^6 μL, 5.5 - 12.1 x 10^6 μL and 32 - 49%, respectively, reported by CRL (1982) [26] for young and adult male rats.

On the contrary, the extract may have negative effects on hematological status in females in which erythrocyte and hematocrit values dropped significantly. The mean values of RBC, WBC, and hematocrit of pregnant extract-treated rats were not within the normal physiological ranges of 6.2 - 7.6 x 10^6 μL, 4.39 - 9.73 x 10^6 μL and 34 - 42%, respectively, reported by CRL (1982) [29] for young and adult female rats. The results suggest that the animals could have suffered from leucocytosis, which may result from intoxications including those produced by metabolic disturbances [30].

Hepato-renal toxicity was studied by measurement of some biochemical parameters. ALAT is a cytosolic enzyme secreted in the liver cells when it is released into the bloodstream when liver cell necrosis occurs [31]. It is a specific enzyme in the liver, making it an important and very sensitive indicator of hepatotoxicity [32]. ASAT is also an indicator of hepatocyte destruction even if, in addition to the liver, it is also found in the heart, skeletal muscle, lungs and kidneys [31]. ALAT and ASAT levels rise rapidly when the liver is damaged for various reasons including hepatic cell necrosis, hepatitis, cirrhosis and liver toxicity of certain drugs [33]. In the present study, ALAT levels tended to increase in female and male rats (at doses above 250 mg/kg), and ASAT levels tended to decrease in the females but the differences were not statistically significant compared with the controls. These results suggest that the extract may not have hepatoprotective effects at high doses in spite of the presence of flavonoids [17], molecules with known hepatoprotective activity [34].

Previous studies on the acute and subacute toxicity of *E. speciosa* aqueous extract led to the recommendation of the 250 mg/kg dose for therapeutic use since higher doses [800-1600 mg/kg] caused histopathological signs in the lungs (diffuse alveolar damage), kidneys (tubular cell necrosis), reduced glomerular space, scattered inflammation and liver (vascular congestion, biliary stasis) [35]. The results of the present study are in line with the earlier recommendation.

4. Conclusion

Administration of *E. speciosa* aqueous extract in male rats for 60 days caused a decrease in relative testes weights, and increased epididymal weights. These effects were accompanied by a significant increase in the density and motility of spermatozoa. In the female rats, high doses of extract resulted in decreased ovarian weights and reduced counts of some haematological parameters. These effects had no observable negative influence on the reproductive performance of the rats. The extract of *E. speciosa* can be safely used by reproducing couples (prior to and in the early stages of pregnancy) at the dose of 250 mg/kg for the management of various disease conditions without the risk of reproductive toxicity.

Acknowledgements

This work was carried out with the contributing support of the authors. Authors B. Nchegang and P.V. Tan designed the study and wrote the protocol, did the literature analysis, and finalized the drafts for structural sections, and statistical analysis. Authors B. Nchegang, Z.E. Nkwengoua and P.V. Tan did the phytochemical study of the plant. Authors B. Nchegang and C. Mezui managed the biochemical analysis. Author B. Nchegang wrote the first draft, and author P.V. Tan supervised the study.

References


