

## EFFECT OF FRACTION 2 OF *PORTULACA OLERACEA* ON PLASMA GONADOTROPHINS AND TESTOSTERONE LEVELS AND THEIR RECOVERIES IN MALE WISTAR RATS

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Article Received on  
13 Nov 2015,

Revised on 04 Dec 2015,  
Accepted on 24 Dec 2015

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### ABSTRACT

**Objective:** To investigate effect of chromatographic fraction 2 of *Portulaca oleracea* on plasma gonadotrophins and testosterone levels and their recoveries in male rats. **Methods:** Air-dried specimen of *Portulaca oleracea* was cold-extracted in methanol for 72 hours. The resulting methanol extract was then subjected to open column chromatography on silica gel for fractionation. Out of 5 fractions obtained, fraction 2 was then subjected to male rats' hormonal bioassays. Twenty male rats (120-150 g) were divided into control and fraction 2(1, 2, 3 mg/kg) treated groups (5 per group) for hormonal assays study. The animals were orally treated on daily basis for 50 days and allowed a recovery (withdrawal) period of 50 days after which hormonal assays were carried out at the end of dosing and recovery periods. Plasma LH, FSH and testosterone levels were

assayed using Enzyme – Linked Immunosorbent Assay (ELISA) technique. **Results:** Treatment of rats for 50 days with fraction 2(2 mg/kg, 3 mg/kg) caused significant ( $p < 0.05$ ) increase in LH levels relative to control. Fraction 2(1 mg/kg, 3 mg/kg) caused significant ( $p < 0.05$ ) increase in FSH levels relative to control. Fraction 2(1 mg/kg, 3 mg/kg) caused significant ( $p < 0.05$ ) decrease in testosterone levels relative to control. There was a significant ( $p < 0.05$ ) decrease in LH level of fraction 2(3 mg/kg) recovery group relative to control. **Conclusion:** It can therefore be concluded that fraction 2 of *Portulaca oleracea* probably

induced significant changes on plasma gonadotrophins and testosterone levels through the hypothalamic – pituitary – testicular axis which could result in sterility.

**KEYWORDS:** Fraction 2, Testosterone, Luteinizing hormone (LH), Follicle stimulating hormone (FSH), Rats.

## INTRODUCTION

*Portulaca oleracea* belongs to the family of Portulacaceae. It is a warm - climate annual herb and has cosmopolitan distribution. It is commonly called Purslane in English language and “Esan omode” or “Papasan” by the Yoruba language tribe of South - West Nigeria.<sup>[1]</sup>

It is used in Iranian folk medicine as a diuretic, vermifuge, antiscorbatic, antitussive, analgesic and gastroesophageal reflux.<sup>[2]</sup>

Pharmacologically, *Portulaca oleracea* extracts have been reported to decrease morphine dependence in mice.<sup>[3]</sup> Its extracts have been reported to have analgesic and anti-inflammatory effects.<sup>[4]</sup> The aqueous and methanol extracts of this plant have contractile effects on isolated intestinal smooth muscle in *in-vitro* preparations.<sup>[5]</sup> Its extracts have been reported to cause reduction in locomotor activity and an increase in the onset time of pentylenetetrazole (PTZ) – induced convulsion in rats.<sup>[6]</sup> Its crude extracts have also been reported to have beneficial effects on the hematological functions and blood chemistry of rats.<sup>[7]</sup>

Since the crude extracts of this plant have been reported to cause significant reductions in testosterone levels in male rats<sup>[8]</sup>, this study therefore aims at investigating the effect of chromatographic fraction 2 of *Portulaca oleracea* on plasma gonadotrophins and testosterone levels and their recoveries in male Wistar rats.

## MATERIALS AND METHODS

### Experimental Animals

Adult male rats weighing between 120 g and 150 g bred in the Pre-Clinical Animal House of the College of Medicine and Health Sciences, Afe Babalola University were used. They were housed under standard laboratory conditions and had free access to feed and water. They were acclimatized to laboratory conditions for two weeks before the commencement of the experiments. All experiments were carried out in compliance with the recommendations of Helsinki’s declaration on guiding principles on care and use of animals.

### Plant Material

Fresh specimens of *Portulaca oleracea* were collected from the Botanical Garden of the Forestry Research Institute of Nigeria, Jericho, Ibadan and was authenticated in the above named institute where a voucher specimen (No FHI 108334) was deposited.

### Extraction and Fractionation of *Portulaca oleracea*

About 3.2 kg of air-dried specimen of *Portulaca oleracea* was cold-extracted in methanol for 72 hours. The mixture was filtered using a wire-gauze and a sieve with tiny pores (0.25 mm) and concentrated at room temperature by exposing the extract for six days. The resulting solution was then placed in the oven at a reduced temperature (45 – 50°C).

The methanol extract was then pre-absorbed with silical gel and placed in the oven at a reduced temperature (45 – 50°C) overnight and then subjected to open column chromatography on silical gel (F<sub>254</sub>, 50 - 200 mesh, E. Merck) for fractionation. The solvents (mobile phases) were hexane (non-polar), ethylacetate (partially polar) and methanol (polar). The gradients of the mobile phases involved hexane with an increasing percentage of ethylacetate (hexane/ethylacetate mixture) and then ethylacetate with an increasing percentage of methanol (ethylacetate/methanol mixture) as shown below.

Hexane		Ethylacetate		Methanol
100% (50 ml)	:	0% (0 ml)		
90% (45 ml)	:	10% (5 ml)		
80% (40 ml)	:	20% (10 ml)		
70% (35 ml)	:	30% (15 ml)		
60% (30 ml)	:	40% (20 ml)		
50% (25 ml)	:	50% (25 ml)		
40% (20 ml)	:	60% (30 ml)		
30% (15 ml)	:	70% (35 ml)		
20% (10 ml)	:	80% (40 ml)		
10% (5 ml)	:	90% (45 ml)		
0% (0 ml)	:	100% (50 ml)	:	0% (0 ml)
		90% (45 ml)	:	10% (5 ml)
		80% (40 ml)	:	20% (10 ml)
		70% (35 ml)	:	30% (15 ml)
		60% (30 ml)	:	40% (20 ml)

50% (25 ml)	:	50% (25 ml)
40% (20 ml)	:	60% (30 ml)
30% (15 ml)	:	70% (35 ml)
20% (10 ml)	:	80% (40 ml)
10% (5 ml)	:	90% (45 ml)
0% (0 ml)	:	100% (50 ml)

Twenty-one fractions were obtained after the column chromatographic procedure.

### Thin Layer Chromatography (TLC)

The 21 fractions were spotted on pre-coated plates of silica gel GF<sub>254</sub> (20 x 20, 0.5 mm thick; E. Merck) using capillary tubes. The spotted TLC plates were developed in a tank that contained a mixture of ethylacetate/methanol (9:1) as the mobile phases.

The TLC plates were then examined under the ultraviolet (UV) light at a wavelength of 365 nm and the well - defined spots of the components were then revealed by the UV light. Fractions with similar relative fronts or retention or retardation factors ( $R_f$  value) were then pooled or bulked together, this then reduced the number of fractions to five (fractions 1, 2, 3, 4, 5).

$$R_f = \frac{\text{distance compound has moved from origin}}{\text{distance of solvent front from origin}}$$

Fraction 2 was then subjected to bioassay, *vis-à-vis*, its effect on hormonal profiles in male rats were evaluated.

### Acute Toxicity Test of Chromatographic Fraction

The acute toxicity test of chromatographic fraction 2 of *Portulaca oleracea* was evaluated in mice as described by.<sup>[9]</sup> Fifteen adult male mice weighing between 20 – 22 g were divided into five mice per group. Three doses of the fraction: 1 mg/kg, 5 mg/kg and 10 mg/kg were given orally to the animals. The control group mice (n=5) received 0.5 ml of distilled water. The animals were observed for seven days for behavioral changes and mortality.

### Experimental Design

Twenty animals were randomly divided into four groups with each group consisting of five rats. The four groups were subjected to the following oral daily treatments for 50 days.

Group I rats received 1 mg/kg of fraction 2.

Group II rats received 2 mg/kg of fraction 2.

Group III rats receive 3 mg/kg of fraction 2.

Group IV rats received 0.5 ml of distilled water as the control group.

### Collection of Blood Samples

Twenty four hours (day 51) after the last dosing of the four groups and also twenty four hours after the last day of the 50 days recovery period (day 101), blood samples were collected from all the animals through the medial cantus into EDTA bottles for the determination of plasma gonadotrophins and testosterone levels using ELISA technique with Fortress Kit.

### Statistical Analysis

The mean and standard error of mean (S.E.M.) were calculated for all values. Comparisons between the control and the treated groups were done using one-way analysis of variance (ANOVA) with Duncan's Multiple Range Test. Differences were considered statistically significant at  $p < 0.05$ .

### RESULTS

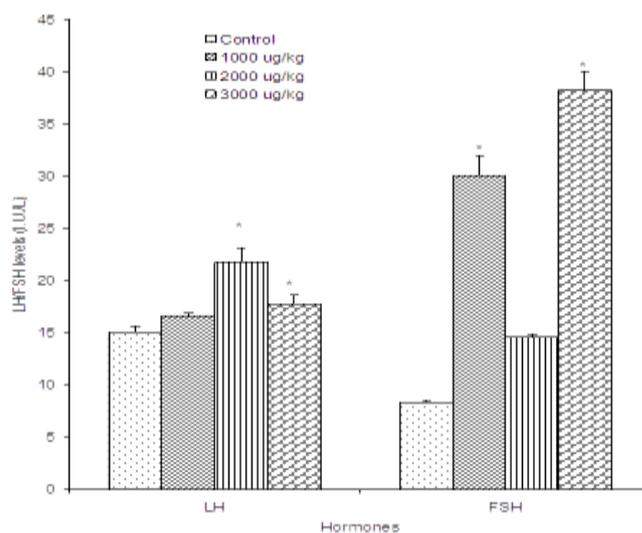
No mortality or changes in behavior were observed in all the treated and control groups of rats.

Treatment of rats for 50 days with fraction 2 (2 mg/kg, 3 mg/kg) caused significant ( $p < 0.05$ ) increase in LH levels relative to the control, while fraction 2 (1 mg/kg) caused insignificant ( $p > 0.05$ ) change in LH level relative to the control. Treatment of rats for 50 days with fraction 2 (1 mg/kg, 3 mg/kg) caused significant ( $p < 0.05$ ) increase in FSH levels relative to the control, while fraction 2 (2 mg/kg) caused insignificant ( $p > 0.05$ ) change in FSH level relative to the control (Fig. 1). Treatment of rats for 50 days with fraction 2 (1 mg/kg, 3 mg/kg) caused significant ( $p < 0.05$ ) reductions in testosterone levels relative to the control, while fraction 2 (2 mg/kg) caused insignificant ( $p > 0.05$ ) change in testosterone level relative to the control (Fig. 2).

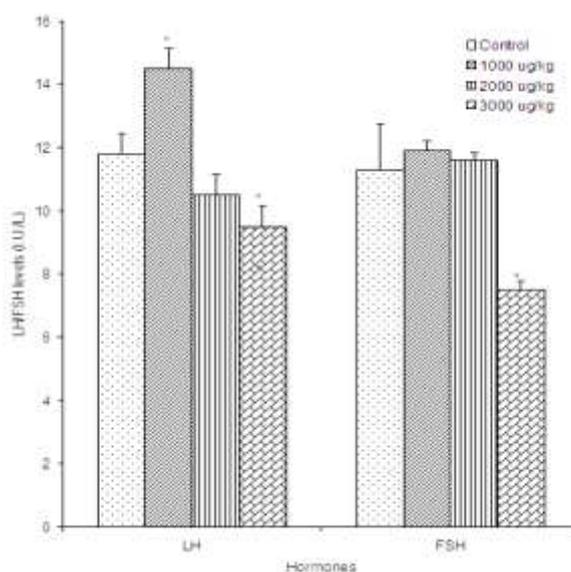
There were significant ( $p < 0.05$ ) increase and decrease respectively in LH levels of 1 mg/kg and 3 mg/kg of fraction 2 recovery groups relative to the control. There was insignificant

( $p > 0.05$ ) change in LH level of fraction 2 (2 mg/kg) recovery group relative to the control. There were insignificant ( $p > 0.05$ ) changes in FSH levels of fraction 2 (1 mg/kg, 2 mg/kg) recovery groups relative to the control, but there was a significant ( $p < 0.05$ ) decrease in FSH level of fraction 2 (3 mg/kg) recovery group relative to the control (Fig. 3).

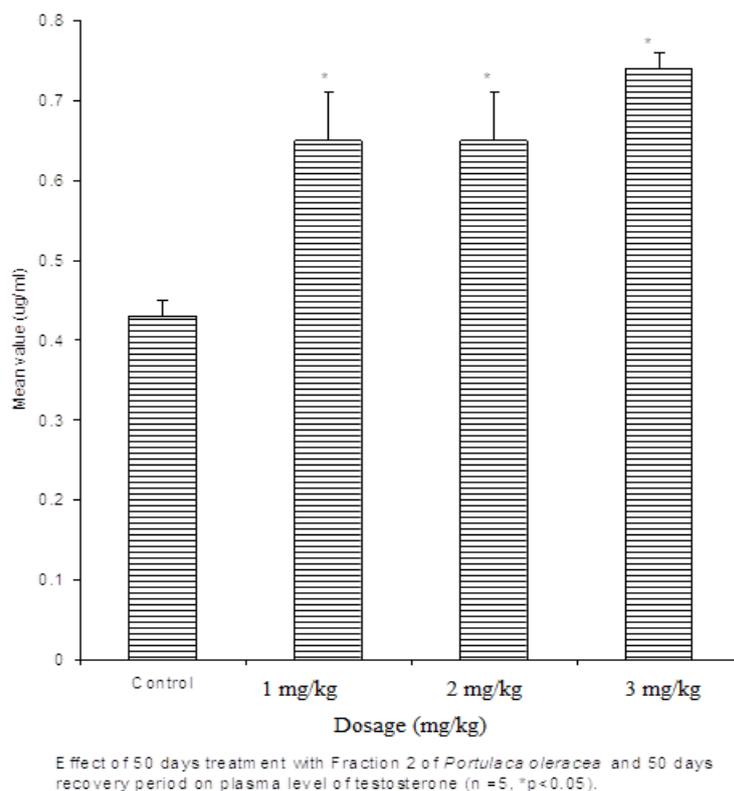
There were significant ( $p < 0.05$ ) increases in testosterone levels of fraction 2 (1 mg/kg, 2 mg/kg, 3 mg/kg) recovery groups relative to the control (Fig. 4).



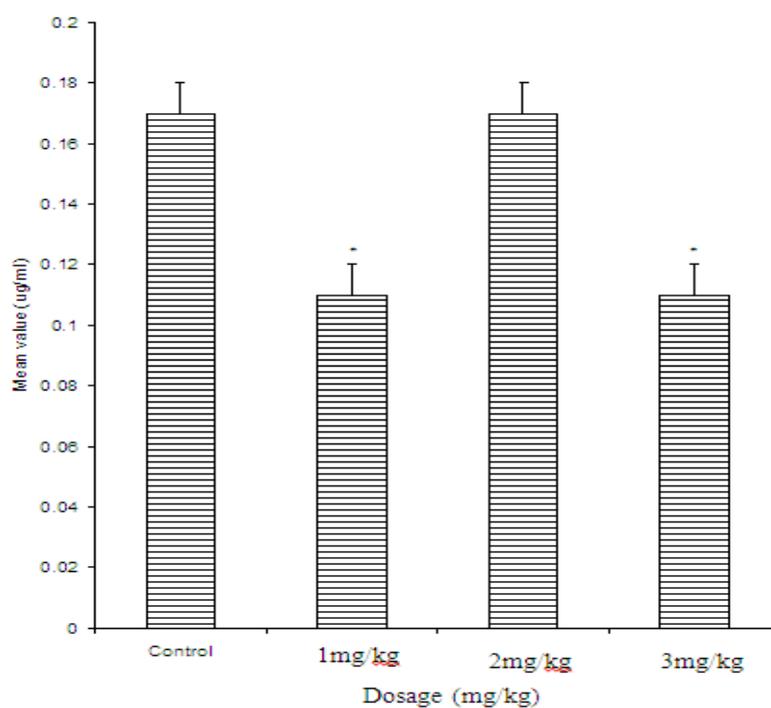
**Fig. 1: Effect of 50 days treatment of rats with fraction 2 of *Portulacaoleracea* on plasma LH and FSH levels (n=5, \* $p < 0.05$ ).**



**Fig. 2: Effect of 50 days recovery period from 50 days treatment of rats with fraction 2 of *Portulaca oleracea* on plasma LH and FSH levels (n=5, \* $p < 0.05$ ).**



**Fig. 3:** Effect of 50 days treatment with fraction 2 of *Portulaca oleracea* on plasma testosterone levels (n=5, \*p<0.05).



**Fig. 4:** Effect of 50 days recovery period from 50 days treatment of rats with fraction 2 of *Portulaca oleracea* on plasma testosterone levels (n=5, \*p<0.05).

## DISCUSSION

It was observed that the highest dose of fraction 2 caused no mortality or behavioral change in all the treated animals, which probably indicates that the fraction has a wide safety margin.

The fraction induced significant increases in LH levels which could be due to the indirect action of this fraction on the anterior pituitary gland thereby causing the stimulation of LH release by the adenohypophysis. Contrary result was reported by<sup>[10]</sup> in *Caricapryl* – 99 extract treated rats. However, there was still significant increase in LH levels after the recovery period which could be due to slow renal clearance of this fraction.

The fraction induced significant increases in FSH levels which could be due to the indirect action of this fraction on the anterior pituitary, thereby causing the stimulation of FSH release by the adenohypophysis. Similar result was reported by<sup>[11]</sup> in *Ruta chalepensis* extract treated rats. However, there was a significant reduction in FSH level after withdrawal (recovery) of treatment with the fraction which could be due to the negative feedback effect on the anterior pituitary by the hormone called inhibin which is secreted by the Sertoli cells. This hormone (inhibin) has a strong direct effect on the anterior pituitary gland to inhibit the secretion of FSH and possibly a slight effect on the hypothalamus to inhibit the secretion of GnRH.<sup>[12]</sup>

The fraction caused significant reductions in testosterone levels. Similar results were reported in *Carica papaya* and *Azadirachta indica* extracts treated rats.<sup>[13,14]</sup> This decreases in testosterone levels could indicate that fraction 2 inhibit the mechanism intervening in the process of the hormone synthesis in the Leydig cells. However, there were significant increases in testosterone levels after withdrawal of treatment (recovery) with the fraction which probably indicates that the effect of this fraction is reversible.

## CONCLUSION

It can therefore be concluded that chromatographic fraction 2 of *Portulaca oleracea* induced significant changes on plasma gonadotrophins and testosterone levels through the hypothalamic – pituitary – testicular axis which could result in sterility.

## CONFLICT OF INTEREST

We vehemently declare that there is no conflict of interests in this research work.

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