Hematological effect of chronic administration of ethanolic extract of *Garcinia conruana* seed on rat

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**ABSTRACT**

Present work deals with the haemalogical effect of ethanolic extract of *Garcinia conruana* seed on wistar albino rats. 20 rats weighing 150-200g were divided into four groups of five rats per group, and were gavaged with the extract of *Garcinia conruana* seed daily for 21 days. Group 1 served as the control group and was fed with standard animal feed only, while group 2, 3 and 4 were the test groups. In addition to the standard feed, 100mg/kg, 200mg/kg and 300mg/kg of the extract was orally gavaged to the test groups 2, 3 and 4 respectively. The result showed increase in Hemoglobin concentration (Hb), Packed Cell Volume (PCV), Red Blood Cell (RBC), Mean Cell Hemoglobin (MCH) and Mean Cell Volume (MCV) but with a significant decrease in Mean Cell Hemoglobin Concentration (MCHC), total White Blood Cell (WBC) and differentials ($P<0.001$, $n=5$). This extract therefore is noted to have a mild erythropoietic effect but with a moderate leucopenia which is characterized by lymphocytosis but decrease in all other WBC lines.

**Keywords:** *Garcinia conruana*; Haematological Indices; RBC; WBC.

**INTRODUCTION**

*Garcinia conruana* (Family-Guttifera) seed is a brown nut–like seed, used in traditional medicine (Braide, et al., 1989), serves as an antidote to the effect of strophantus gratus, as a guinea worm remedy, Poison antidote, treatment of diarrhea, hepatitis, asthma, common cold, cough, dysmenorrhoea and anti-anemia (Holmes, 2001; Lewis, 1977; Dalziel, 1956). The consumption of this seed has increased globally since it’s used medically and traditionally. It is ingested in relatively large amount at various social gathering and so a number of studies have been done while pondering over the possible toxicity in humans vis–versa the pharmacotherapeutic effects of this plant product (Ahumibe, et al., 2009).

It has been shown that chronic ingestion causes histological changes in the liver, kidney and gastrointestinal tract of rats (Braide, et al., 1989). Its active phytochemical components have been shown to have a protective action against chemically-induced hemolysis in G6PD deficient human red blood cells, anti-inflammatory and antipyretic activity (Braide, 1990; Braide, 1993). It has
hypoglycaemic (anti-diabetic) activity, bronchodilatation in man and antispasmodic
effect on smooth muscle (Iwu, 1985; Iwu, et al., 1990; Braide, et al., 1989). In view of
the extensive use of G. conruana nut, with little or no documented fact on the possible
adverse effect on some physiochemical systems, it therefore becomes mandatory to
investigate into the potential effects of chronic administration of ethanolic extract of
Garcinia conruana on haematological indices in rats and to subsequently evaluate
whether it’s ethnopharmacological uses such as mentioned above could have side
effects such as destruction of blood cells which is common with the use of most
chemotherapeutic agents.

**MATERIALS AND METHODS**

**Collection and Preparation of plant extract:** Seeds of *Garcinia conruana* (*Guttiferae*)
were obtained from a local market (Itam Market), Akwa Ibom State, and were
authenticated by a Taxonomist of the department of Botany, University of Uyo. The
seeds were rinsed with water to remove debris, peeled, air-dried and pulverized by
grinding using a manual blender. The total weight of the *Garcinia* seed was 896g. The
powder was divided into two (448g each) and put into two different 200ml conical
flask containing ethanol (98.90%). The suspension was agitated with an electric
blender for about 10 minutes then allowed to store in the refrigerator (4°C). Twenty
four (24) hours later, it was filtered using glass funnel with cotton wool. The filtrate
was evaporated to dryness by heating in a water bath at 40°C. The extract weight was
57.46g (12.83% yields) of crude extract. This was reconstituted in distilled water to an
appropriate concentration, stored in refrigerator awaiting the administration. Lethality
studies put the LD50 of *Garcinia conruana* extract at about 450.24mg/kg. From here,
convenient doses were chosen to preclude the lethal range.

**Collection and maintenance of animals:** Twenty adult albino rats of both sexes
weighing 150-200g were randomly divided into 4 groups of five rats per group. Group
1 served as the control, and was fed with standard animal feed (Bendel Feed and Flour
Mill Ltd, Benin) only, while groups 2, 3 and 4 were the test groups and in addition to
the standard animal feed they were also gavages with 100mg/kg, 200mg/kg and
300mg/kg ethanol extract of *Garcinia conruana* respectively.

All animals were weighed before treatment commenced and were housed in a
standard wooden cage with wooden shavings as their beddings, kept and maintained in
the animal house unit of the department of Pharmacy and Toxicology, Faculty of
Pharmacy, University of Uyo. They were allowed free access to water ad libitum, good
light and maintained in room temperature. After 21 days of extract administration, all
the experimental rats were sacrificed by a blow on the head, dislocating the neck and
the whole blood was obtained by Cardiac Puncture into EDTA bottles for blood
analysis of some haematological indices. This research was carried out in University
of Uyo, according to the rules in Nigeria (Revised Helsinki Declaration, 2008)
governing the use of laboratory animals as acceptable internationally.

**Determination of Hematological Indices:** The determination of the hematological
indices was done within two hours of sample collection using SYSMEX (KX 21)
Automated Hematologic Analyser in the hematology unit of the University of Uyo
Teaching Hospital.

**Statistical analysis:** This was carried out using windows Statistical Package for Social
Sciences (SPSS) version 13. One way analysis of variance was adopted for
comparison, and the results were subject to post hoc test using least square deviation
(LSD). The data were expressed as mean ± standard error. \( P < 0.05 \) were considered significant.

**RESULT**

Table 1 and 2 shows chronic administration of *Garcinia conruana* extract for 21 days produced various effects on the hematological parameters of wistar albino rats. There were increases in PCV, Hb, RBC, MCH, MCV and lymphocyte count, while MCHC, total WBC and differentials were decreased. These were significantly different from control at \( P < 0.01 \).

**DISCUSSION**

The mechanism leading to the increase in RBC parameters is probably mediated by the anti-oxidant property of *Garcinia conruana* seed extract which has been variously demonstrated by other researchers. This gives the extract the hemopoietic, protective and stimulating potential. Previous research has shown that, prophylactic and therapeutic oral administration of anti-oxidant supplement significantly increased cells of hemopoietic origin in animals exposed to potentially lethal dose of radiation (Chris, et al., 2008). Similarly, anti-oxidants such as Vit C, Vit E, succinate and alpha-lipoic have been used to abolish various forms of chemically and metabolically induced oxidative stress to which human hemopoietic cell lines is exposed to (Chris, et al., 2008). The direct link between anti-oxidant activity and hemopoietic boosting effect was also demonstrated by the observation that ascorbic acid supplementation, through its action as a free radical scavenger, increased significantly the hemoglobin levels of children suffering from sickle cell anemia (Jaja, et al., 2002). In a similar way, the active phytochemical component of *G. conruana* seed extract has been shown to have a protective action against chemically-induced hemolysis in G6PD deficient human red blood cells (Braide, 1990).

Two teams of researchers at different times and locations have also demonstrated unequivocal increases in RBC parameters in their work on the effect of the *G. conruana* seed extract on rats and rabbits respectively, as was observed in this study (Esomonu, et al., 2005; Uniqwe, et al., 2009). We also observed that while the
erythropoietic modulated hemocyte lines showed increases in response to administration of the extract, the white blood cells lines rather decreased with exception of lymphocytes. This observation suggests that G. conruana extract effect relating to increase in RBC lines may be effected through an influence on the stimulant cytokine erythropoietin (Ahumibe, et al., 2009). The selective increase in lymphocytes count in this study was similarly observed by other researchers. Adedeji et al., 2006 in their work with pullet chicks observed that, though no significant increase occurred in other white cell lines, lymphocytes specifically proliferated. Ahumibe et al., 2009 also made similar observation in their study of the effect of the extract on erythrocyte membrane integrity and selected hematological indices in male Albino rats. This probably explains the antimicrobial effect of G. conruana extract, in view of the major role that lymphocytes assume in the immune mechanism of the body in both man and animals.

CONCLUSION

Garcinia conruana seed extract through its anti-oxidant property has hematopoietic, stimulating, enhancing and protective effects. Its use as antianemia, antimicrobial and immune booster is therefore supported.

REFERENCES


