# Haematological Changes in Administration of *Chrysophyllum Albidu* Stem Extract to Wistar Rats

#### **Abstract:**

In medical practice, examination of blood for the presence of metabolites and other constituents is vital to the prediction of the physiological, nutritional and pathological status of any individual prior to prognosis. This study investigated the effect of crude n-hexane stem extract of Chrysophyllum Albidum on packed cell volume (PCV) and haemoglobin count in Albino Wistar rats. Twenty (20) Wistar rats of between 220-150g were procured and housed in well-ventilated animal House of  $28 \pm 2^{\circ}$ C, relative humidity 60-70%, for a 12hrs (light/dark cycle) duration of two weeks. They were then grouped into four (4) of five (5) rats per group. With group 1 rats fed with standard rat diet ad libitum (Control group), groups 2 -4 were respectively fed with 200mg/kg, 300mg/kg and 400mg/kg body weights of Chrysophyllum Albidum Extract. After 14 days of administration of test substance, rats were sacrificed, blood samples obtained and analysed for hematologic changes. Study found a statistically significant decrease in PCV and haemoglobin levels for group IV (high dose treated) when compared with other groups. Also, low dose treated group showed an increase in PCV values upon comparison. Study therefore (using analysis of variance, ANOVA at p < .05) established a dose-dependent change in most assayed haematological parameters. In conditions of abnormal spike in PCV levels, we recommend the use of Chrysophyllum Albidum extract.

**Keywords**: Chrysophyllum Albidum, Haematological parameters, PCV, haemoglobin count

# Introduction

With Blood being a major indicator of pathological status for sufferers of ailments and other conditions in disease states, Haematological variables are good indicators of the physiological status of animals<sup>1&2</sup>. Therefore, examination of blood gives an opportunity to investigate the presence of numerous metabolites and other constituents in humans, making it a vital and indispensable fluid in the estimation of their physiological, nutrition and pathological status<sup>3&4</sup>.

The adult human reportedly has about 5 to 6 liters (1 to 2 gallon) of blood which is pumped from the heart through a network of blood vessels collectively known as the cardio-vascular system. This (blood) accounts for roughly 7 to 8 percent of the total body weight. Infants and children have comparably lower volumes of blood that is proportionate to their

smaller size. The volume of blood in an individual is known to fluctuate. During dehydration for example, while running a marathon, blood volume decreases, and increases in circumstances like pregnancy, when the mother's blood needs to carry extra oxygen and nutrients to the baby<sup>3</sup>.

According to Olafedehan *et al.* (2010)<sup>5</sup> examining blood for its constituents can provide important information for the diagnosis and prognosis of diseases in humans. In disease conditions, blood constituents are known to change in relation to variations in an individual's health<sup>6</sup>. These changes are valuable in assessing their responses to treatments and several other physiological situations<sup>1</sup>. Afolabi *et al* (2010) once reported that changes in haematological parameters are often used to determine various status of the body and to determine stresses due to environmental, nutritional and/or pathological factors<sup>7</sup>.

Normalizing haematological variables and taming them to homeostatic levels in disease conditions remain debatable<sup>8</sup>. Partly due to its cost intensiveness, especially in developing societies where quality health care is an issue. Overtime, undesirable and adverse effects associated with the use of orthodox drugs has lead people to resort to the use of suitable herbs with minimal effect on hematological variables<sup>9&10</sup>. A great number of such herbs now serve traditional medical practitioners (Trado-Medics) in combating and ameliorating heamatological related ailments. One of such often alleged herbs of great importance is *C. Albidum*.

Chrysophyllum Albidum in South-western Nigeria is a fruit called "agbalumo" and popularly referred to as "udara" in South-eastern Nigeria. The plant often grows to a height of 36.5m though it may be smaller<sup>11</sup>; several other components of the tree including the roots and leaves are used for medicinal purposes<sup>11&12</sup>. The bark is used for the treatment of yellow fever and malaria, while the leaf is used as an emollient and for the treatment of skin eruption, stomach ache and diarrhoea<sup>8&13</sup>. The leaf has antiplatelet and hypoglycaemic properties<sup>14</sup>. The root bark has anti-fertility effects<sup>15</sup>. Stem bark extracts have antimicrobial effects and Antiplasmodial<sup>16</sup> effects. The seed cotyledon has been reported to possess Antihyperglycaemic and Hypo-lipidemic effects<sup>16</sup>.

Since little or no information is available about haematopoietic potentials or toxicity of *C. Albidum*, this study was therefore undertaken to unravel the haematological changes that accompany its administration to wistar rats. Specifically, study determined the effect of crude n-hexane stem extract of *C. Albidum* on pack cell volume (PCV) and haemoglobin counts in albino wistar rats.

# **Materials and Methods**

# **Scope of Study**

Study was an ex-vivo experiment, and was conducted with Wistar rats in the animal house of the Faculty of Basic Medical Sciences, Delta State University, Abraka, Delta State, Nigeria. Study was restricted to unveiling status of common haematological variables in administration of *C. Albidum* at different doses.

#### **Study Design**

Study was an experimental based, and animals [twenty (20) wistar rats] were purchased, acclimatized for two weeks, and grouped into four groups of 5Wistar rat per group as follow;

Group 1: Normal Control

Group 2: C. Albidum Extract, 200mg/kg

Group 3: C. Albidum Extract, 300mg/kg.

Group 4: C. Albidum Extract, 400mg/kg.

#### **Materials**

Electronic weighing balance, refrigerator, blender, heating mantle, centrifuge, meter rule, needle and syringe (2ml and 5ml), hand gloves, beaker, filter paper, tissue paper, cotton wool, distilled water, dissecting kits, crude n-hexane leaf extract of *C. Albidum*.

#### **Collection and Identification of Plant Sample**

The plant materials (*C. albidum*) were procured in the month of March 2015 from local market in Evbuobanosa Community, Edo State, Nigeria. The plant was authenticated at the Herbarium Section of the Department of Botany, Faculty of Science, Delta State University, Abraka, Nigeria.

#### **Preparation of Plant's extract**

Fresh leaves of *C. Albidum* were washed with tap water and oven-dried for 2days. Crisply dried leaves were powdered with the aid of a blender. 500g of powdered *C. Albidum* leaves were soaked in 1000ml of n-hexane. The mixture was stirred every 6hours for 72hours and then filtered with Whatman's filter paper. The filtrate was thereafter concentrated to

dryness with the aid of hot air oven set at  $40^{\circ}$ C. Final extract was weighed (9g) and used in the calculation of the percentage yield (0.02w/w %) using the relation below;

$$percentage\ yield = \frac{Final\ weight\ of\ extract}{weight\ of\ powdered\ extract} \times \frac{100}{1}$$

#### **Preparation of stock solution**

The crude plant extract was reconstituted in hydro-ethanol (ratio 1:9) solution to give the required doses of 200mg/kg, 300mg/kg and 400mg/kg body weight of the extract used in the study.

Plant extract: (Standard dose = 200mg/kg, 300mg/kg and 400mg/kg)

First, 2g of the leaf extract (of *C. albidum*) was weighed with electronic weigh balance and constituted in 100ml of hydro-ethanol solution. This gave a stock solution of 2000mg/100ml (20mg/ml) for low dose 200mg/kg. Next, 3g of the n-hexane leaf extract was then weighed with electronic weigh balance and constituted in 100ml of hydro-ethanol solution. This gave stock a solution of 3000mg/100ml (20mg/ml) for medium dose 300mg. lastly; 4g of the leaf extract was weighed with electronic weigh balance and constituted in 100ml of hydro-ethanol solution to give a stock solution of 4000mg/100ml (40mg/ml) for high dose 400mg/kg.

#### **Ethical Issues**

Ethical clearance was obtained from the Research and Ethics Committee of the Faculty of Basic Medical Sciences, College of Health Sciences, Delta State University, Abraka, Delta State. Animals were handled according to protocols approved by the institutional animal ethics committee (IAEC), as adopted by the Faculty of Basic Medical Sciences, Delta State University, Abrake, Nigeria.

#### **Procedure**

#### **Determination of Haemoglobin**

20ul (0.02ml) of capillary blood or well mixed venous blood was carefully collected (from rats) and dispensed into 2ml of the ammonia diluting fluid. The solution was read immediately at a stable colour for 6-8 hours. Next, the meter performance was checked by inserting the control statement glass provided in the cuvette aperture reading was then made

corresponding to the stated value of  $\pm$  5. Blood sample was then transferred to a clean 10mm light-path cuvette and read.

## Test method Packed Cell Volume (PVC)

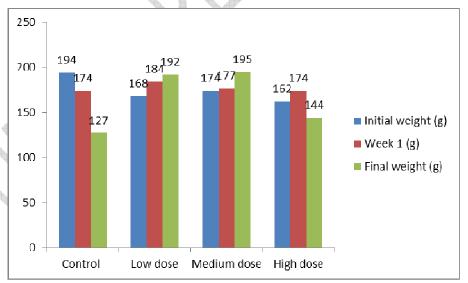
A heparinized capillary tube was filled up to three quarter with a well mixed EDTA anticoagulated blood (tested within 6hours of collection). Next, one end of the capillary tube was seal with a sealant material by heating sealing capillaries. The filler capillary in one of the numbered slots of the micro-haematocrit rotor was then carefully located to prevent breakage. Entire content was then centrifuged for 5minutes (RCF 12000-15000xg) and immediately read.

#### **Statistical Analysis**

Study presented results as mean  $\pm$  standard error mean (SEM) n=5. Raw data were analysed by post Hoc LSD alpha t-test for multiple comparison, using statistical package for social science (SPSS-20) p-value less than .05 were considered statistically significant.

## Results

Figure 1: Effect of C. Albidum extract on body weights of Wistar rat.



n=5, p < .05: significant as determined by post Hoc LSD alpha t-test for multiple comparison. Statistically increased significant difference was seen in group weights upon comparison with control group

Figure 2: Changes in Packed Cell Volume of *C. Albidum* extract administration to Wistar rat.

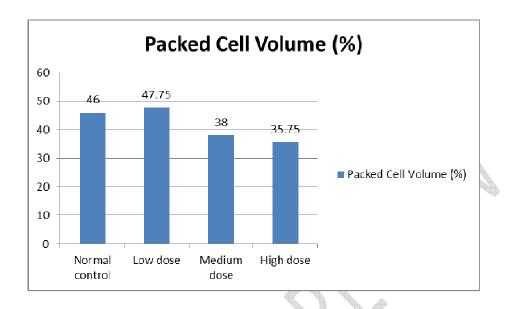


Figure 3: Changes in Haemoglobin count of *Chrysophyllum Albidum* extract administration to Wistar rat.

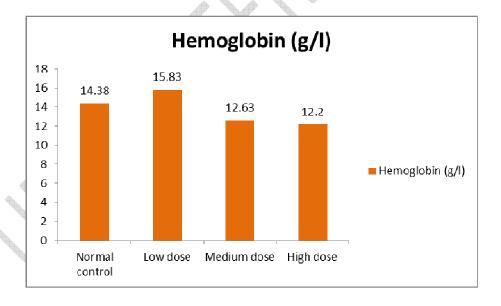
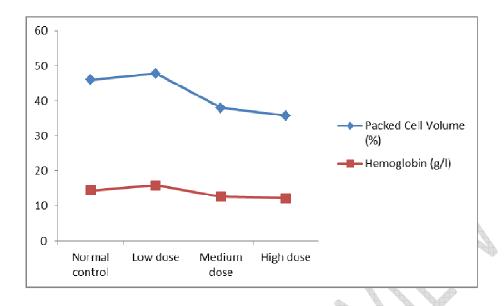


Figure 4: Relationship between PCV and Haemoglobin count in *Chrysophyllum Albidum* extract administration to Wistar rat.



## Discussion

Sometimes referred to as "the river of life", blood as a vital fluid in humans is reportedly built of 45% of three major types of cells; red blood cells (Erythrocytes), white blood cells (Leukocytes), and platelets (Thrombocytes), the remaining 55% of blood is composed of a liquid known as plasma. Several factors have been implicated to affect the ratio of the volume of packed red cells to that of total blood volume, which is the haematocrit (38–52% for males and 37–47% for females)<sup>18</sup>. To this point, current study was designed to investigate the effect of crude n-hexane stem extract of *C. Albidum* has on body weight, packed cell volume (PCV) and haemoglobin count, using Albino Wistar rats as experimental model.

From the study, a statistically significant increase was seen in the body weight of rats treated with *Chrysophyllum albidum*. This could be as a result of tannins present in the extract as tannins have been implicated to stimulate increase in body mass. For example, a recent study by Marcus *et al.*, 2003 revealed that tannins present in small quantities in medicinal plants are potent in increasing body mass<sup>19</sup>. Also, a phytochemical analysis of *C. Albidum*<sup>16</sup> has shown the presence of small quantities of tannins (among other things) as one of its active component. This could be responsible for the increased body weight observed in this study.

In a similar vein, current study, upon careful analysis observed a statistically significant increase in Haemoglobin and PCV levels within the duration of treatment with *C. Albidum* n-hexane leave extract at low, medium and high doses. Studied groups showed possible haematinic potentials of this plant upon administration. Before now, Adewoye *et al.*,

2011 had shown that *C. Albidum* contains tannins, flavonoids, saponins, alkaloids, anthraquinone and cardenolides. Some species of plants known to contain similar phytochemical constituents as *C. Albidum* have been reported to pose anti-anaemic effects<sup>20-22</sup> on experimental animals. It is likely that these phytochemical constituents might be responsible for the anti-anaemic properties observed in this study. The increase in the blood indices was progressive, giving a notable effect on the 14<sup>th</sup> day our treatment. Under normal condition, the body generates new red blood cells (RBCs) to replace the lost red cells and this process takes a much longer time<sup>14</sup>. The quick attainment of normal RBC count could well be an indication of accelerated erythropoiesis occurring as a result of treatment with *C. Albidum*. This result suggests that *C. albidum* leaf extract might have directly stimulated increased production of red blood cell precursors thereby increasing blood components.

A close look at figure 1 shows a huge dose-dependent increment in weight with the administration of *C. albidum* extract. Here, a statistically significant difference in weight was seen among various groups upon comparison with control. This however proved insignificant in final weight gain for high dose administration of test extract. The exact reason for this is yet to be fully understood. However, active ingredient in *C. albidum* may be implicated. This finding concurs with that of Adisa (2000); who posited that at high dose, *C. albidum* has a weight lowering effect on animals<sup>13</sup>.

Again, Flavonoids have been suggested to be a possible factor responsible for the increased erythrocyte count in Wistar rat<sup>23&24</sup>. It might well be that the flavonoid content in C. albidum was responsible for the erythropoietic ability observed in this study (as seen for high PCV in figure 2).

# **Importance of Study**

In effect, this study will supply necessary information regarding the effect of medicinal plant (*Chrysophyllum albidum*) body weight, as well as selected haematological variables (PCV and Hb count). Finding of this study will also help patient with anaemia to make necessary adjustment in therapy.

#### **Conclusion and Recommendations**

In this study, the crude n-hexane leave extract of *C. albidum* has caused a great deal of increase in blood volume for treated rats. Thus, it might be possible that *C. albidum* has the ability to balance between the rate of destruction and production of blood cells as evident by the increased PCV and Hb counts from this study. This increased haematological parameters observed in Wistar rat treated with *C. albidum* could be a result of the flavonoid present in the plant, while the tannin content might have been responsible for the increased body weight of the animals. *C. albidum* may therefore be said to be a good plant source for haematinics, antiplatelets and drug development.

# References

- 1. Khan, T. A and Zafar, F. (2005). Haematological Study in Response to Varying Doses of Estrogen in Broiler Chicken. *International Journal of Poultry Science*, 4(10), 748-751.
- 2. Martini, Frederic et al. (2007). Anatomy and Physiology. Rex Bookstore, Inc. p. 643, ISBN 9789712348075.
- 3. Aderemi, F. A. (2004). Effects of replacement of wheat bran with cassava root sieviate supplemented or unsupplemented with enzyme on the haematology and serum biochemistry of pullet chicks. *Tropical Journal of Animal Science*, 7, 147-153.
- 4. Doyle, D. (2006). William Hewson (1739-74). The father of haematology. *British Journal of Haematology*, 133(4), 375-381.
- 5. Olafedehan, C. O., Obun, A. M., Yusuf, M. K., Adewumi, O. O., Oladefedehan, A. O., Awofolaji, A. O., &Adeniji, A. A. (2010). Effects of residual cyanide in processed cassava peal meals on haematological and biochemical indices of growing rabbits (p.212). Proceedings of 35th Annual Conference of Nigerian Society for Animal Production.
- 6. Togun, V. A., Oseni, B. S. A., Ogundipe, J. A., Arewa, T. R. (2007). Effects of chronic lead administration on the haematological parameters of rabbits a preliminary study (p. 341). Proceedings of the 41st Conferences of the Agricultural Society of Nigeria.
- 7. Afolabi K. D., Akinsoyinu, A. O., Olajide, R., &Akinleye, S. B. (2010). Haematological parameters of the Nigerian local grower chickens fed varying dietary levels of palm kernel cake (p.247). Proceedings of 35th Annual Conference of Nigerian Society for Animal Production.
- 8. Idowu, T.O., Iwalewa, E.O., Aderogba, M.A., Akinpelu, B.A., Ogundaini, A.O. (2006). Antinociceptive, Anti inflammatory and Antioxidant activities of *Eleagnine*: an alkaloid isolated from seed cotyledon of *C. albidum. Journal of Biological Science*, 6 (6): 1029 1034.
- 9. Isaac, L. J., Abah, G., Akpan, B and Ekaette, I. U. (2013). Haematological properties of different breeds and sexes of rabbits(p.24-27). Proceedings of the 18th Annual Conference of Animal Science Association of Nigeria.

- Iwuji, T. C and Herbert, U. (2012). Haematological and serum biochemical characteristics of rabbit bucks fed diets containing garcimiola kola seed meal (p.87-89). Proceedings of 37th Annual Conference of Nigerian Society for Animal Production.
- 11. Bada, S.O. (1997). Preliminary information on the ecology of *Chrysophyllumalbidum*done in the West and Central Africa; In proceedings of a National Workshop on the potentials of Star Apple in Nigeria (Denton, O.A., Ladipo, A.O., Adetoro, M.A., Sarumi, M.B.), pp. 16 25.
- 12. Adewusi, H.A. (1997). The African Star Apple *Chrysophyllum albidum* Indigenous Knowledge from Ibadan, Southwestern Nigeria. In: Proceedings of a National Workshop on the Potentials of the Star Apple in Nigeria (Eds.), pp. 25-33.
- 13. Adisa, S.A. (2003). Vitamin C, Protein and Mineral contents of African Apple (*Chrysophyllum albidum*) In: proceedings of the 18th annual conference of MIST (eds) Garba SA, Ijagbone IF, Iyagba A O, IyamuA O, Kilani AS, Ufaruna N, pp. 141-146.
- 14. Agbor, A.G., Odetola, A.A. (2001). Haematological studies of *Parguetinanigrescenson* haemorhagicanaemic rats. *African Journal Medicine and Science*, 30: 105 109.
- 15. Onyeka, C.A., Aligwekwe, A.U., Olawuyi, T.S., Nwakama, E.A., Kalu E.C., Oyeyemi, A.W (2012a). Antifertility Effects of Ethanolic Root Bark Extract of *Chrysophyllumalbidum*in Male Albino Rats. *International Journal of Applied Research in Natural Products*, 5 (1): 12-17.
- Adewoye, E.O., Salami, A.T., Lawal, T.O., Adeniyi, B.A. (2011). The Antimicrobial and Kill Kinetics of *Chrysophyllumalbidum*stem bark extract. *European Journal of Scientific Research*, Vol 56 No 3 pp. 434 444.
- 17. Olorunnisola, D.S., Amao, I.S., Ehigie, D.O and Ajayi, Z.A.F (2008). Anti-hyperglycaemic and hypolipidemic effect of ethanolic extract of *Chrysophyllum albidum* seed cotyledon in alloxan induced-diabetic rats. *Research Journal of Applied Science*, 3: 123-127.
- 18. Iwuji, T. C and Herbert, U. (2012). Haematological and serum biochemical characteristics of rabbit bucks fed diets containing garcimiola kola seed meal (p.87-89). Proceedings of 37th Annual Conference of Nigerian Society for Animal Production.
- 19. Marcus, C., Karin, L., Jain, G., Matthias, L., Jorns, F., Tilman, G., Jurgen, S. (2003). Captive roe deer (*Capreoluscapreolus*) select for low amount of tannic acid but not quebracho: flunctuation of preference and potential benefit. *Biochemistry and Molecular Biology*, volume 136, issue 2: pg. 369 -382.
- 20. Amusa, N.A., Ashaye, O.A., Oladapo, M.O. (2003). Biodeterioration of the African Star apple (*Chrysophyllumalbidum*) in storage and the effect on its food value. *African Journal of Biotechnology*, 2: 56 57.
- 21. Chia, S., Nagurney, J.T., Brown, D.F.M., Raffel, O.C., *et al.* (2009). Association of leucocyte and neutrophil counts with infarct size, left ventricular function and outcome after percutaneous coronary intervention for ST- elevation myocardial infarction. *American Journal of Cardiology*, 103: 333–337.

- 22. Dina, O. A., Adedapo, A. A., Oyinloye, O.P., Saba, A.B. (2006). Effect of *Telfairiaoccidentalis*ex tract on experimentally induced anaemia in domestic rabbits. *African Journal of Biomedical Research*, 3: 181-183.
- 23. Dhakar, R., Katare, Y.K., Patcl, U.K., Pawar, R.K. (2012). In vivo assessment of bioactivity of *Trichosanthesdioica* Roxb for the management of haemolyticanaemia. *International Journal of Pharmacology and Technology Research*, 4 (2).
- 24. Esomonu, U.G., El Taalu, A.B., Anuka, J.A., Ndodo, N.D., Salim, M.A., Atiku, M.K. (2005). Effect of ingestion of ethanol extract of *Garcina Kola* seed on erythrocyte in wistar rats. *Nigerian Journal of Physiological Science*, 20 (1-2): pg. 30 -32.