



Original Research Article

The effects of aqueous leaf extract of *Mucuna pruriens* (agbala) on some selected biochemical indices of wister albino rats

B.Eze Obioma¹, I.Obeagu Emmanuel², P.C.Ugwu Okechukwu^{1,4} and J.C.Ifemeje³

¹Department of Biochemistry, University of Nigeria Nsukka, Enugu state Nigeria.

²Diagnostic Laboratory Unit, University Health Services, Michael Okpara University of Agriculture, Umudike, Abia State, Nigeria

³Department of Biochemistry, Anambra State University, Uli, Anambra State Nigeria

⁴Department of Biochemistry, Tansian University, Umunya, Anambra State Nigeria

*Corresponding author

ABSTRACT

Keywords

Mucuna pruriens;
Agbala Leaf;
Alanine amino-transferase;
Aspartate amino transferase;
Trace elements.

The effects of aqueous leaf extract of *Mucuna pruriens* on some selected biochemical parameters of aspartate amino transferase (ALT), alanine amino transferase (ALT) and some trace elements on wister albino rats were assayed. The results showed a significant ($p < 0.05$) increase in the liver enzymes of ALT and AST of the heated and raw extracts when compared with their corresponding controls. The iron and copper concentrations of the rats increased significantly ($p < 0.05$) in the raw and heated extract groups when compared with the controls. The zinc concentration increased significantly ($p < 0.05$) in the raw extract group but decreased significantly ($p < 0.05$) in the heated extract group when compared with the control. There was a progressive change in the relative body weights of the rats after 11 days of feeding. The results of this study showed that aqueous leaf extract of (Agbala) *Mucuna pruriens* are rich in iron and copper and could be dangerous to the liver as shown from the elevated levels of AST and ALT liver markers of the heated and raw extract groups.

Introduction

Plant extracts have been used in folk medicinal practices for the treatment of various ailments since antiquity. A medicinal plant as defined by the world health organization (WHO) is a plant which one or more parts of it contain substances that can be used for therapeutic purposes or which are precursors for the synthesis of useful drugs (Ogamba *et al.*,

2011). *Mucuna pruriens* is thought to have originated from India. It is one of the popular medicinal plants of India and it constitutes more than 200 indigenous drug formulations. All parts of *Mucuna pruriens* possess valuable medicinal properties.

In the eastern part of Nigeria *Mucuna*

pruriens popularly known as agbala leaf in Igbo populace is used as a blood tonic traditionally (Katzenshlager *et al.*, 2004; Akindele and Busavo, 2011 and Ogamba *et al.*, 2011). Among the natives of eastern part, the use of *Mucuna pruriens* (agbala leaves) extract is a very common remedy for the treatment of anemia. The fresh leaves are collected from the farm, garden or bush manually, the leaves are washed with clean water and then squeeze to remove the liquid content of the leaves (Katzenshlager *et al.*, 2004). This liquid extract is then boiled for about five minutes and is taken orally as blood tonic to boost blood production.

Aims and objectives of the research

The aim of this research was to determine the effects of aqueous leaf extract of *Mucuna pruriens* on some selected biochemical parameters of aspartate amino transferase (ALT), alanine amino transferase (ALT) and some trace elements on wister albino rats in order to determine the efficiency of this extract in African traditional medicine.

Materials and Methods

Preparation of the aqueous plant extract

The leaves of *Mucuna pruriens* were washed and weighed. 100g of the leaves were extracted with 100ml of distilled water with the aid of a manual (cheese cloth) sieve. 60ml of *Mucuna pruriens* was measured and heated for 5mins. The heated extract was allowed to cool at room temperature while the remaining 60ml (raw extract) was used like that. 60mls of raw and heated extracts were measured and given to the rats daily.

Plant materials

The leaves of *Mucuna pruriens* were collected from Umuoma and were authenticated by Mr. C.J Onyirioha of the Department of Biochemistry Anambra State University, Uli.

Experimental animals

The animals used for this study were both male and female wister albino rats with average weight of 45-66g. They were purchased from animal house of the Faculty of Pharmaceutical Sciences, University of Nigeria Nsukka. The animals were housed in locally fabricated cages in the animal house of Department of Biochemistry, Anambra State University, Uli for 4 weeks. They were allowed to acclimatize to the new environment for seven days before the commencement of the experiment. They were fed with animal feed water *ad libitum*.

Experimental design

Nine wister albino rats were used in this study. The rats were randomly divided into three groups made of three animals each as shown below. The animals were fed for three weeks. They were given extract to drink at their own will which served as their water.

Group A: served as the control and received only water and normal guinea feed.

Group B: 60ml of raw extracts

Group C: 60ml of heated extracts

Body weights

Initial and final body weights of the animals were recorded at the end of the treatments period (3 weeks), the animals

were sacrificed and blood sample collected.

Sample collection

Blood sample was collected from the rats fed with aqueous extracts using orbital technique. Blood sample was collected from the retro-bulbar plexus of the medial canthus of the eye to puncture the retro-bulbar plexus out flow of blood into bottle containing ethylene-diamine-tetra-acetic acid (EDTA). The sample was stored at 4°C before analysis.

Determination of liver markers

Liver markers of ALT and AST were assayed using Reitman and Frankel (1957) methods as outlined in the Randox kit.

Determination of trace elements

Iron, copper and Zinc concentrations of the experimental rats were assayed using AOAC, 1990 methods.

Results and Discussion

The data showed that ALT and AST concentrations of the rats increased ($p < 0.05$) significantly in both raw and heated extracts when compared to their respective controls (Table.1).

The results showed that copper and iron concentrations of the experimental rats also increased significantly ($p < 0.05$) in both copper and iron levels of raw and heated extract groups when compared with that of their respective control groups. But zinc decreased in heated extract group when compared with the control (Table.2).

There was a progressive change in the relative body weight of the rats after

eleven days of feeding. There was also, a relatively body weight change after twenty one days of feeding, but the change was not progressive like that of eleven days of feeding .

The results of this analysis carried out on the effects of aqueous leaf extract of *Mucuna pruriens* on some selected liver markers of AST, ALT and some trace elements on wister albino rats revealed an elevation in ALT, AST and some trace elements of copper and iron. These results showed that *Mucuna pruriens* raw and heated leaf extracts could be deleterious to the liver in high concentration and at the same time increase the production of blood.

Zinc is always in close association with proteins, it occurs wherever protein is and enhances protein function (Ezeokonkwo, 2004). Iron is an essential component of haemoglobin, therefore its high amounts in the extract indicates that the extract plays a role in the synthesis of blood.

The results in table 3 showed a progressive change in the relative body weight of the rats after 11 days of feeding. There was also a relatively body weight change after twenty one days of feeding, but not as progressive as that for 11 days of feeding. This shows that as the extract was serving as water to the rats, they did not actually drink it as water and they became dehydrated and did not gain much weight. Meanwhile, their nutritional status was also affected and they drank heated extract more than raw extract. Also, the heated extract rats gained more weights than that of the raw extract rats.

In conclusion, the results of this study suggest that the raw and heated extracts of *Mucuna pruriens* could be used in the

Table.1 The results of some selected biochemical parameters of rats fed with *Mucuna pruriens*

Sample	Aspartate amino transaminase (AST)	Alanine amino transaminase (ALT)
Raw extracts	41.67±0.015	52.65±0.0208
Heated extracts	41.35±0.015	51.35±0.305
Control	40.52±0.208	50.48±0.029

Table.2 The results of some selected trace elements of rats fed with *Mucuna pruriens*

Sample	Zinc (mg/l)	Copper (mg/l)	Iron (mg/l)
Raw extracts	6.67±0.005	1.21±0.015	112.95±0.029
Heated extracts	4.42±0.028	0.99±0.015	81.32±0.028
Control	5.73±0.028	0.52±0.028	76.23±0.028

Table.3 The results of the mean body weights (g) of rats fed with *Mucuna pruriens*

Sample	Mean body weight in (g) before feeding	Mean body weight in (g) after eleven days of feeding	Mean body weight in (g) after twenty one days of feeding.
Raw extracts	53.85	70.48	78.45
Heated extracts	50.62	71.32	81.82
Control	63.53	96.99	120.44

synthesis of blood when consumed because of its high contents of iron, copper and zinc. The results also showed that heated and raw leaf extracts of *Mucuna pruriens* could be deleterious to the liver in high concentration.

References

Akindede A.J. and Busavo F.L. 2011. Effects of the hydroethanolic extract of *Mucuna pruriens* on haematological profile in normal and haloperidol treated rats. Nigeria Quarterly Journal of Hospital Medicine, 212: 8-93.
 Amin, K.M.Y; Khan, M.N and Ziller – Rehman, S. 1996. Sexual function improving effect of *Mucuna pruriens* in sexually normal male rats .3rd ed.

Fitoterapia publication. Pp53-58.
 Association of Official Analytical Chemists AOAC 1990, Official Methods of Analysis, 15th Edition, pp. 220-224, Washington DC.
 Butler, E. and Waalen, J. 2006. The definition of an Aemia :Blood heamoglobin . 107:1747-1750.
 Bolarin, D.M. 2006. Hemoglobin: clinical chemistry 2nd edition. Spectrum books limited Ibadan. Pp62-80
 Corey , C. 2007 . Hemolytic Anemia : Sidney kimmed comprehensive cancer center Hopkinds, Baltimore 15:441-449.
 Churchill , E, Echeobi , O. and Nnelii , R. 2007 . Implication of nutrition. Journal of medical sciences 5:242-244.

- Crook , M.A. 2006 Biochemical test for liver disease : Clinical chemistry and metabolic medicine . 7th ed. Edward Arnold publishers ltd . London .
- Elmer, H.C1997. Better living through agricultural sciences: Agricultural science fundamental and application. 2nd ed. Delmera, a division of Thoson learning , inc , united states of America Pp.30, 116.
- Essig ,M.G. and Poore , R. 2008http:1 www.health link : bc .caik base|Topic\modest|Iw 2033\describe .Htm Retrived on 25|05|09|.
- Ezeanyika , L.U.s 2004 . Biochemistry for Beginners . Basic :Aspect of Nutritional Biochemistry 1st ed. Published by Great AP express Publishers limited .Pp 113 -131.
- Ezeokonkwo, C.A. 2004 Biochemistry for Beginners : Elements Relevant in Biology 1st ed. Publishers limited .Pp30-45.
- Dacie, J.V. and Lewis, S.M. 2000. Practical Haematology. 9th Edition Churchill Livingstone.
- Gasket , H. Derry , S . More , R.A2008. Prevalance of aneamia in older persons : Journal of BMC Geriatrics 8 2318-36,8-21.
- Giuliano F . Allard , J Dopamine and male sexual function . 2001.Eur urol 40:601-608.
- Grund, S.2007. Permicious anaemia : journal of new Britain General hospital .15:98-110.
- Health stout, N. 2009. Definition of anaemia . http: www. Health stout :Com|ency|68|112|main-html.Retrieved on 25|05|09.
- Hund, K. 2008. Prevention of anemia htt:ll www.Cureresearch .com|a|anaemia|prevalence.Htm.Retrieved on 24|08|09.
- Human serum albutin .2008. Wikipedis.org :|wiki|human serum albutin . Retrieved on09|09|09
- Human albumin. 2007. Hutchison encyclopedias :www.Right health .com||pediaetrics Retrieved 09|09|09
- Ifon ,E.T.1997. The nutrient composition of some Nigerian leafy green vegetables and physiological availability of their iron contents :Ph.d. Thesis ,Department of Biochemistry, University of Ibadan Nigeria . Pp 335-445.
- Islam ,M.S. Lucky ,N.S and Siddi ,M.S.I.2004 variation hematological Parameter :International journal of Poultry Science 2:144-147.
- Juhn , G.Eltz, D.R. 2006.Iron deficiency anaemia :journal of varimed Health care Network Baltimore 15:98-110.
- Katzenschlager ,R. Evans, A and Manson , A 2004. Mucuna Pruriens in parkinsons disease .A double blind clinical and Pharmacological study :Journal of neural Neurosury psychiatry. 75:6672-1677.
- Kavach, S .2001.http://.www.Herbal provider .com /mucuna htm http:Wikipedia org |wiki| mucuna - pruriens.
- Kimberly, 12|03|09.serum total protein :en. Wikipedia-org |wiki | serum -total protein Retrieved on 20|09|09.
- Kingsley , R,S Shelly. J and James, E.B. 2003. Flowering plants and civilization :Introduction to plant biology .9th ed.marger ,J kemp. Mc Grew-Hill companies. Pp 138-140,470-472.
- Lumeij J.T., de Bruijne J.J., Kwant, M.M. 1990. Comparism of different methods of measuring protein and albumin in Pigeon Sera. Avian Pathol 192: 61-255.
- Matsui, W.2006. B12 deficiency anaemia : Journal of verimed Health care Network Baltimore .10:216-220.
- Nancy,J.N. 2008. *Mucuna pruriens* information from :NPGS/GRN:www. ars-grin .gov|http: www.arsgrin

- gov|cgibin|npgs|htm|taxon.PI? 24652.
Retrieved on 18|6|09.
- Ogamba J.O., Eze N.A., Ogamba S.E., and Chilaka K.C.2010. *Tropical Journal of Medical Research*. 142: Pp1-6.
- Oxford Advanced learners Dictionary. 2001.6th edition Oxford University press Pp 672 and 887.
- Oxford Concise Medical Dictionary .2003.6th edition. Oxford Publishers New York. Pp475 and476
- Pearson, D.A. 1969. *Chemical analysis of food* ,7th ed. Churchill living stone New York .Pp27-32.
- Penninx, B.W. Pahor, M. 2004. Anaemia is associated disability :*Journal of Jam Geriater Soc*.52.719-724.
- Purves, W.K. David, S. Gordon, H.O 2004. *Life: The Science of Biology*. 7th edition Sunderland. Mass:sinader Association Pp954.
- Reitman, S. and Frankel, S. 1957. *In vitro* determination of glutamic-pyruvic transaminase in serum. *American Journal of Clinical Pathology*, **28**:56.
- Roper ,N 2000.*Pocket medical Dictionary* :Churchill Living Stone 14th edition . Icn New York.Pp42,56,78,107,281.
- Solomon,B.M.V.2002 *Reproduction in flowering plants: Biology*. 3rd .Martins brother publisher Ltd. Pp 747-748.
- Tessmer , K.2009. Total protein : [www.Healthopedia.com/total protien](http://www.Healthopedia.com/total_protien). Retrieved on 20|90|09.
- Wikipedia ,2009. Alanine Transamines : [http://en : Wikikipedia. Org/Wik/ Alanine transaminase](http://en.wikipedia.org/wiki/Alanine_transaminase).Retrieved on June 2009.
- William,B.Van.V. 2007.*Aspartate transaminase*: [http://www.medindia.net blood test BiochemistryAspartate transamines](http://www.medindia.net/blood_test/Biochemistry/Aspartate_transamines).Retrieved on 06|20|2009.
- William, M. 2006. Idiopathic autoimmune hemolytic Anaemia: *Journal of verimed Health care Network Baltimore*. 15:741-449.