

Hypoglycaemic Activity and Biochemical Effects of *Cataranthusroseus* with Diabinese in Normal and Diabet IC Rats

Jones B.B¹, Tabe N. N¹, Ushie O.A² *

¹Department of Chemical Science, Cross River University of Technology Calabar, Nigeria

²Department of Chemical Science, Federal University Wukari Nigeria

***Corresponding Author:** Ushie O.A, Department of Chemical Science, Federal University Wukari Nigeria.

Abstract: The hypoglycaemic activity and biochemical effects of *cataranthusroseus* with diabinese in normal and diabetic rats were compared. Ethanolic extract of *C. roseus* were screened for their phytochemical constituents using standard methods. Biochemical effects were evaluated on alloxan induced diabetic rat (32) assigned into four study groups of eight rats each Group 1 (control) received diabinese (14.28mg/kg). *C roseus* (400mg/kg) all in 30% ethanol. At the end of 21 day administration period blood serum was obtained from sacrificed animals and biochemical indices of toxicity (lipid profile, aminotransferases activity, serum protein, urea and albumin).

1. INTRODUCTION

Diabetes mellitus is a chronic endocrinologic disorder characterized by high blood glucose levels arising from insufficient secretion of insulin by the pancreas or improper utilization by target cells. Diabetes is the most common endocrinologic disorder with widespread prevalence cutting across all populations groups (Report of the expert committee on Diagnosis and classification of diabetes mellitus, 1998). The insufficient secretion of insulin by pancreatic islet cells or defective insulin receptors. Leads to poor utilization of insulin by target cells culminating in raised fasting blood glucose levels (>8mmol/L) glycosylated haemoglobin (Hb A1C) greater than 6 – 9%. Glycosylated haemoglobin gives an indication of average blood glucose concentration over a 6 – 8 week period and is a biochemical test used for monitoring progression of diabetes or response to treatment. While the capillary or a venous glucose value gives the blood glucose levels. Besides the immediate perceptible metabolic dysfunction accompanying hyperglycemia. There are long – term complications of diabetes mellitus involving disorders of the eyes, kidneys, nerves and blood vessels.

The Canadian Diabetes Association (2003), reported three main types diabetes: Type 1 diabetes usually diagnosed in children and adolescence which occurs when the pancreas is unable to produce insulin necessary to meet the energy needs of tissues required in the body. About 10 percent of people with diabetes have type I diabetes. It is also called insulin dependent Diabetes. The remaining 90 percent of diabetes have type II diabetes that occurs when the pancreas does not produce enough insulin or body does not effectively use insulin that is produced and is developed in adulthood.

A third type, gestational diabetes which is a temporary condition, occurs during pregnancy. It affects 3.55 percent of all pregnancies and present an increased risk of developing diabetes for both mother and child in later life (Mayfield 1998; Canadian diabetes association 2003). According to Clavell (2002) report on Diabetes Management as documented by the Mayo foundation for Medical Education and Research in New York identified a fourth type of diabetes arising from illness or medications that can interfere with production of insulin or its action. Approximately 1-2% of all diagnosed cases belongs to this group and include inflammation of the pancreas, pancreatotomy, adrenal or pituitary disorders hydrocortisone treatments for another disease, some high blood pressure and Cholesterol lowering medications malnutrition and infection.

Diabetes is a syndrome of disturbed intermediary metabolism diabetes mellitus commonly concomitant with hypertension and other disease (Greff, 2000). It is therefore not a single disease entity but has been described to be a syndrome characterized by an absolute or relative lack of insulin

leading to persistent elevation of blood glucose as well as alteration in lipid and fat metabolism (spencer and cudworth 1989).A lot of specific secondary causes can lead to this syndrome. These include pancreatectomyheamachromatosis including iron over load of beta cells excessive cortisol production (cushing syndrome) excessive growth hormone production as in agromegally, insulin resistant syndrome occasioned in part by insulin receptor defects (Lernmarket *al*, 1981; Laakoet *al*, 1988) where the patients does not show these defects, he is said to have idiopathic diabetes (WHO, 1995).

2. MATERIALS AND METHODS

The ethanolic extracts of *Cataranthusroseus*were subjected to phytochemical screening to establish their phytochemical compositions. Ninety rats (Males and Females) of wistar strain weighing 140-245g obtained from the animal house. Department of Biochemistry, University of Calabar.Diabetes was induced on a pool of seventy rats by intraperitoneal injection of 150mg/kg body weight of alloxan monohydrate (Sigma, st Louis, Mo. USA). Using distilled water as the vehicle.The research work was organised, into phases. Phase 1 and phase 11. The first consisted of 24 diabetic and non-diabetic (normal) animals respectively assigned into 4 groups of six rats each. These were used for hypoglycaemic assessment only on the hand. The second phase consisted of purely diabetic animals (32 animals) also assigned into 4 groups of eight each and tread as in phase 1.

Treatment was administered twice a day by gastric intubation in a 12 hour cycle 7.00am and 7.00pm everyday for 14days. Body weight of the animal monitored after every two days throughout the administration period using a beam balance. Twelve hours after last feeding and administration (overnight fast) the animals were anesthetized under chloroform vapour then dissected. Whole blood was collected from the heart by cardiac puncture and divided into two fractions. Serum triglyceride level was estimated with the use of Dialab laboratory kit. The method employs enzymatic hydrolysis of triglyceride in the sample with lipoprotein lipase (LPL).In Hayems and Turks fluids a 1:200 dilution of blood was made by taking 20 microlitres of blood in a glass tube (75x12mm). The tube sealed with rubber band was tilted through an eagle of 120 and rotated for 2 minutes. With a Pasteur pipette. the country chamber (New improved Neubauer country chamber) was filled with the diluted blood sample, ensuring that no blood spilled into surrounding moat, The cells were allowed to settle by leaving the chamber undisturbed for 2 minutes, after which the cells were counted under low magnification (4mm dry objective and x 10 eye piece). The country involved cells in the four squares at the corners of the chamber of WBC as well the cells in the centre square alongside those in the squares at the corners for the RBC.

2.1 Statistical

Data obtained was expressed as meant standard deviation and analysed the Analysis of variance (ANOVA) or student's t-test where applicable. Values $P < 0.05$ were regard as significant in comparison to appropriate controls.

3. RESULTS

Phytochemical composition of *Catharanthusroseus* extract shows the ethanolicleaf extracts of *C roseus* contain various phytochemical components which differ in their relative abundance. They include cardiac glycosides, sapouins, tannins and anthraquinones which showed slight presence.Effect of treatment on blood glucose concentration of rats show comparative effect of ethanoic extracts of *Cataranthusroseus* and diabinese treatment on serum glucose of diabetes rats. The treatment with these drugs reduced significantly ($P < 0.05$) serum glucose levels of normal non-diabetic animals (81.32 ± 5.32 , and 78.64 ± 6.47 for diabinese, and *Cataranthus* respective) and diabetic rats (163.05 ± 3.75 , and 105.61 ± 3.62 respectively) for diabeinese and *Cataranthusroseus* relative to their respective controls (107.88 and 247 ± 4.83 for normal and diabetic rats).

Table1. Comparative effect on fasting blood glucose (mg/dl) following a 21day administration of diabinese*Cataranthusroseu* extracts on normal and diabetic rats.

Treatment groups rat	Normal-Diabetic rats	Diabetic rat
Control (30% ethanol	107.88 ± 7.38	247.25 ± 4.83
Diabinese treated (14.29mg/kg)	$*81.31 \pm 5.32$	$*163.05 \pm 3.75$
C roseus treated (400mg/kg)	$*78.64 \pm 6.47$	$*105.61 \pm 3.63$

Values of expressed as mean \pm SD, = 6

*= Statistically significant ($P < 0.05$) compared with controls values without asterisks are not significant.

3.1 Effect of Treatment on Serum Lipid Fraction of Diabetic Rats

The comparative effect of ethanolic extracts of *Cataranthusroseus* and diabinese on serum lipid fractions including Total cholesterol, Triglycerides (TG) and High density lipoprotein have been shown in table 3. Treatments with diabinese produce decreases in Total Cholesterol and Triglycerides but an increase in High density lipoprotein of these changes only that of Triglycerides was significant (88.16 ± 25.41) compared to the control (142.19 ± 40.13). Treatment with diabinese however, caused significant decrease ($P < 0.05$) in High density lipoprotein level (33.99 ± 8.25) and a decrease in total cholesterol (63.62 ± 16.11) and triglycerides (86.16 ± 18.92) compared to their respective controls. The decrease was only significant in total cholesterol ($P < 0.05$) and not total cholesterol.

Table 2. Comparative effect of diabinese, ethanolic extracts of *C. roseus* on serum lipids of diabetic rats treated for 21 days.

Treatment Groups	Total Cholesterol (mg/dl)	Triglycerides (mg/dl)	High density Lipoprotein (mg/dl)
Diabetic Control	69.98 ± 9.50	142.19 ± 40.13	43.10 ± 4.00
Diabetic Diabinese treated	63.62 ± 16.11	*88.16 ± 18.92	39.99 ± 8.25
Diabetic <i>C. roseus</i> treated	60.08 ± 8.22	*86.59 ± 22.76	**60.05 ± 7.87

Values represented mean ± SD n = 8

*= Statistically significant ($P < 0.05$) compared with the control

**= Statistically significant ($P < 0.01$) compared with the control are not significant.

Table 3. Comparative effect of a 21 day treatment with diabinese ethanolic extract of *C. roseus* on serum levels of total protein, albumin and urea of diabetic rats

Treatment Group	Total Protein (g/dl)	Albumin (g/dl)	Urea (mg/dl)
Diabetic control	7.16 ± 0.52	3.43 ± 0.14	7.30 ± 6.67
Diabetic diabinese treated	*8.41 ± 0.98	3.39 ± 0.34	**12.63 ± 0.29
Diabetic <i>Cataranthusroseus</i> treated	**8.31 ± 0.18	*3.33 ± 0.30	**25.17 ± 7.31

Values represent mean ± SD n = 8

* = Statistically significant ($P < 0.05$) compared with control

**= Statistically significant ($P < 0.01$) compared with control

Treatment effect on aminotransferase activity of diabetic rats. Aminotransferase activities were determined and the results serum AST activity decreased in treatment (101.00 ± 10.42 and 102.60 ± 8.23 for diabetic diabinese treated and diabetic *Cataranthusroseus*). Compared to the control (110.13 ± 14.26). The decreases were however, non significant ($P > 0.05$). Serum ALT activity in serum also decreased upon treatment (14.62 ± 2.90 , 15.50 ± 3.60 and 13.13 ± 1.31 , diabetic diabinese treated). The decrease in ALT was only significant ($P < 0.05$) in the group treated with *C. roseus*. The presence of these substances may be responsible for their anti hyperglycemia action. Win/eman (1989) had earlier in his report indicated that plants endowed with flavonoids, glycosides and phytosterols are likely to possess both hypoglycaemia and antihyperglycemic action. Serum lipid fraction information available from pathophysiology reveals that it is associated with lipoprotein abnormalities due to defect in insulin, which is an antihyperglycemic and lipogenic hormone. As a consequence diabetes has increased risk for development of premature atherosclerosis due to the increased HDL levels (Bierman, 1992).

In this study total cholesterol and triglycerides raised in diabetic controls were reduced significantly following treatment with the herbs and diabinese. Nimeribo-Uadia (2003) and Uhegbu and Ogbuehi (2004). The saponins are thought to bind cholesterol to bile acids. Consequently lowering cholesterol in plasma (Nimeribo-Uadia, 2003). However, the effects of *C. roseus* and to a lesser extent diabetes are more beneficial with respect to HDL whereas *C. roseus* increased significantly the levels of HDL.

4. CONCLUSION

Some of the indices assayed after 21 days administration period indicates that the herbs could protect the diabetic animals better than the standard drug. The herbs also proved more efficacious in their hypoglycemic action. *Cataranthusroseus* are very efficacious, in its hypoglycemic and antihypoglycemic action than the standard drug. *C.roseus* also proved to possess the ability to reverse hyponatremia seemingly induced by diabinese treatment. *C. roseus* could protect the rats against damage on the hematopoietic tissue i.e. leucopenia caused by diabinese treatment. *Cataranthusroseus* might give a better and holistic protection as it is the only in the study with a vantage and positive impact on HDL cholesterol (good cholesterol).

REFERENCES

- [1] Canadian Diabetes Association (2003). Clinical practice guidelines for prevention and management of diabetes in Canada. Ontario: Canadian Diabetes Association.
- [2] Banley, C. V. & Day, C. (1989). Traditional treatment for diabetes
- [3] Spencer K. M & Cud Worth, A. G. (1989). Diabetes in epidemiological.
- [4] Lernmark, A. Hagglof. D & Freedman. W. (1981). A prospective analysis of antibodies reacting with pancreatic islet cells in Insulin dependent diabetic children. *Diabetologia*. 20.471- 474.
- [5] Clavell. M. C. (2002). Managing diabetes in Mayo, I. (ed). Mayo Foundation For medical education and research (2nd edition) (pp.5-20). New York: Kensington press.
- [6] Nimenibo-Uadia, R. (2003). Effect of Vernonia amygdalina in alloxan-induced diabetes albino rats. *Journal of Medical Laboratory Science*. 12(1) 25-31.
- [7] Thompson. L. U. (1993) Potential health benefits and problems associated with antinutrients in food *Research International*. 26, 137-149.
- [8] World Health Organization.(1985). World health organization study group technical report on diabetes mellitus. Geneva. World Health Organization.
- [9] Greoff, O. R. W. (2000). The Management of diabetes mellitus. *GeneeSkunde: The Medicine Journal*. October 2000. Retrieved on October 5, 2005 from <http://www.medpharm.co.2a/safp/2000/oct/mellitus.html>.
- [10] Wineleman. M. (1989). Ethnobotanical treatment of diabetes. *Baja medical Anthropology II*
- [11] Bierman, E.L. (1992). Artherosclerosis in diabetes. *Artherosclerosis and thrombosis* 12, 647 – 656.
- [12] Nimenibo – Uadia, R. (2003). Effect of vernonia amygdalina in alloxan – induced diabetes albino rats. *Journal of medical laboratory Science*. 12 (1).
- [13] Uhegbu, F.O & Ogbuehi, J. (2004). Effect of aqueous extract (crude) of leaves of veronnia amygdaline. Del on blood glucose serum albumin and cholesterol level in diabetic albino rats. *Global Journal of pure and Applied Science*, 10, 189 – 194.
- [14] Nimoibouadia (2003) argued that the saponins are likely to cause blood glucose reduction via their effect on plasma lipids. Furthermore, McCune and Jolius (2002) in a survey of medicinal plants (35) of North America revealed that the phytochemicals of these plants support a lifestyle historically low in the incidence of diabetes in Canada (North America).

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