Phytochemical Analysis and Anti-Hyperlipidemic Potential of Ethanolic Extract of *Hunteriaumbellata* Seed

Morakinyo A. E.¹, Oladeji O. A.², Fajobi A. O.³, Adelowo J. M.⁴, Babarinde S.O.¹, Adepoju A. E.¹

¹(Department of Biochemistry, AdelekeUniversity, Ede,Nigeria)
²(Department of Food science and Technology, Ondo State University of Science and Technology, Ondo, Nigeria)

³(Department of Biochemistry, ObafemiAwolowo University, Ile-Ife, Nigeria)
⁴(Department of Basic Sciences, Adeleke University, Ede, Nigeria)

Abstract:

Background: Hyperlipidemia characterized by the abnormal elevation of plasma lipoprotein is the most prevalent indicator for cardiovascular diseases. This study screened, analyzed and accessed the anti-hyperlipidemic potentials of the phytochemicals and anti-nutrient compositions of ethanolic extracts of H. umbellata seeds with a view to utilizing the plants in the management of obesity and hyperlipidemia, and early stage renal dysfunction using high fat diet model in rats.

Materials and Methods: Thirty healthy albino rats were randomly divided into five equal groups: Group I received normal saline (2 ml/kg bwt); Group II received a high fat diet; Group III received Orlistat (50 mg/kg bwt); Group IV Orlistat (50 mg/kg bwt) + H. umbellata (200 mg/kg bwt); Group V received Orlistat (50 mg/kg bwt) + H. umbellata (400 mg/kg bwt); The hepatocellular injury markers (aspartate transaminase (AST) and alanine transaminase (ALT)), kidney injury markers (urea and creatinine) and lipid profile (total cholesterol (TC), triglycerides (TG), high density lipoprotein (HDL) were estimated using standard Randox diagnostic kits, Low Density Lipoprotein (LDL) concentration were calculated using Friedewald's equation

Results: The results revealed that H. umbellata seed has nutrients and anti-nutrients within permissible limits. It also revealed that HFD increased the body weight and the total trigycerides(TG), total cholesterol(TC) and low density cholesterol(L DL-c), post-treatment with 400 mg/kg H. umbellata seed significantly (p < 0.05)

decreased the concentrations of TG, TC and LDL when compared with the HFD group. Also post-treatment with 200 mg/kg plant extract significantly (p < 0.05) decreased urea and creatinine concentrations when compared to the HFD group and the Orlistat (standard drug) treated group.

Conclusion: These results revealed that the ethanolic extract of H.umbellataseed possesses safe to consume anti-nutrients and phytochemicals which elicited anti-hyperlipidemic potentials and normalizes the kidney injury markers at 400 and 200 mg/kg body weight respectively

Key Word: Hunteria umbellata; Hepatocellular injury; Kidney injury; Hyperlipidemic; Phytochemicals

Date of Submission: 23-06-2020 Date of Acceptance: 11-07-2020

I. Introduction

Abnormal and excessive fataccumulation isone of the major risk factors for a number of metabolic disorders like cardiovascular diseases, diabetes and cancer. Once considered high-income countries' problem, obesity and overweight is fast becoming a problem in urban areas of low-income countries mainly as a consequence of lifestyle changes¹. Pre-clinical studies have shown a direct correlation between a high fat diet and overweight/obesity among distinct animal models². An important aspect not often discussed is the relationship between the overweight and obesity caused byanhypercaloric and high fat diet and the development of kidney failure. Excess weight gain and obesity may cause renal changes, including glomerular hyperfiltration, sodium retention, enlargement of Bowman's space, increased glomerular cell proliferation, mesangial matrix expansion, inflammatory cell infiltration and tubuleinterstitial lesions³. These early renal alterations can progress to more intense and diffuse lesions in the kidneys, such as focal segmental glomerulosclerosis, proteinuria and tubule-interstitial lesions, which are observed in prolonged obesity⁴.

In addition, it hasbeen described that changes in lipoprotein metabolism and serum lipids occur with increasing renal dysfunction⁵. Moreover, the high levels of cholesterol andobesity have been associated with changes of glomerular structure during the final stages of renal disease⁶.

DOI: 10.9790/3008-1504010109 www.iosrjournals.org 1 | Page

There is therefore the need to arrest the trend using relatively assessed plant-based drug with little or no side effects. *H. umbellata* which belongs to the family of apocynaceae, is a plant native to West Africa and reputed to be effective in the treatment and management of diverse ailments ranging from diabetes mellitus, gastric ulcers, liver disease, infections, hypertension and obesity. The plant has been reported to contain tannins, alkaloids, cardiac glycosides, saponins, flavonoids and anthraquinone, and possess antioxidant, anti-inflammatoryand aphrodisiac properties ^{7, 8}. Adeneyeet al., reported the effectiveness of *H. umbellata* in the management of obesity andhyperlipidemia in rats induced with triton and olive oil.

This study is designed to evaluate the effect of *H. umbellate*seed extract in rats fed with high fat diet with a view to ascertaining its effectiveness in the management of obesity and hyperlipidemia, and early stage, renal dysfunction using the high fat diet model in rats.

II. Material And Methods

A total 30 adult male Albino rats (both male and females) that weighed 120 – 130 g were obtained from the Animal House, Faculty of Pharmacy, ObafemiAwolowo University, Ile-Ife. The rats werehoused in the Animal House, Department of Biochemistry, Adeleke University, Ede, Osun stateand kept under standard conditions. The rats were acclimatized for 14 days; fed with standard chow and water *ad libitum*.

Study Design: Invitro and invivo study

Plant Materials: Dried seeds of *H.umbellata* were purchased from Ipetu-modu, Ife North Local Government, Osun state, Nigeria. The seeds were pulverized using a mechanical blender.

Preparation of Ethanolic Extract: The ethanolic extract of *H. umbellata* was prepared according to a modified method described by 16 Donald *et al.*, 2016. Typically, 500 g of pulverized plant material was defatted with n-Hexane (2000 ml) for 24 hours and was extracted with ethanol (1500 ml) for 48 hours. The suspension was stirred and filtered through a muslin cloth and Whatman No 1 filter paper. The residue was re-extracted five more times with ethanol until the filtrate was clear and became colourless. The filtrates were combined, allowed to settle, carefully decanted and re-filtered using WhatmanNo 1 filter paper. The combined filtrate was concentrated to dryness using a rotary evaporator. The extract was kept in the refrigerator for biochemical studies.

Study Duration:

Subjects &Treatment method: Thirty (3 male Albino rats was purchased and selected randomly for the purpose of this study. The animals were randomly allocated and orally gavagedas follows:

Group A(N = 6 rats) – standarddiet with distilled water (Control)

Group B (N = 6 rats) – High Fat Diet only

Group C(N = 6 rats)– High Fat Diet + 50 mg orlistat/kg bwt

Group D(N = 6 rats) - High Fat Diet + 200 mg extract/kg bwt

Group E(N = 6 rats) - High Fat Diet + 400 mg extract/kg bwt

Sacrificing and Samples Collection: The animals were sacrificed under mild anasthesia with diethyl ether, twenty four hours after the last treatment (oral administration of extract and orlistat). Blood was collected via cardiac puncture into bottles containing anticoagulant (trisodium citrate, 3.8% w/v) and mixed gently.Blood sample was centrifuged on Bench Centrifuge Model 90-2 (Searchtech Instrument England, UK.) at3000 rpm for 10 min. The plasma was carefully transferred into clean vial bottles and kept frozen at -4°C for further biochemical assays.Liver (1g) and kidney (1 g) was cut into bits and homogenized in a total of 10 ml with freshly prepared phosphate buffer (pH 6.8, 100 mM). The homogenate was centrifuged at 3000 rpm with Bench Centrifuge at 3000 rpm for 10 min. The supernatant was carefully transferred into clean vial bottles and kept frozen at -4°C for further biochemical assays.

Procedure methodology

The proximate analysis, mineral composition⁹ and phytochemical screening^{10,11} of the ethanolic extract of *H. umbellates*eeds were carried out. The quantification of tannin¹³, saponin¹⁴ and phytate¹⁵ in the seed ethanolic extract was also determined.

All lipid parameters were quantified on the blood plasma; Total cholesterol, High density lipoprotein-Cholesterol (HDL-c) and Triglyceride quantization was determined using standard kits from Randox Diagnostics[®]. Low density lipoprotein-Cholesterol (LDL-C) and very low density lipoprotein-Cholesterol (vLDL-C) was calculated using the Friedewald'sequation¹⁸.

The hepatocellular markers; Aspartate transaminase (AST) and Alanine transaminase (ALT), and kidney biomarkers (urea and creatinine) were determined using standard kits from Randox diagnostics[®].

Information about the type of orlistat was taken from the pharmacy database where it was purchased. Body weights of the rats were measured once a weekof the experiment, using digital mettler (Mettler Toledo Type BD6000, Mettler-Toledo GmbH, Greifensee, Switzerland). The weight difference to the initial weight per group was calculated

Statistical analysis

Datawere expressed as mean \pm SEM. Differences between the mean values of the control and treatment groups were determined by One-way Analysis of Variance withDunnettpost hoc test using the Graph Pad Prism 5. Significant difference was considered if p < 0.05.

III. Result

In Table no 1is the result of thephytochemical screening of the ethannolic extract of *H. umbellata* seed. Itrevealed that cardio glycoside was strongly positive, followed by flavonoids, terpenoids and saponin, that are moderately positive. While the tannins and steroids were slightly positive.

Table no 1: Photochemical screening of ethanolic extract of *H. umbellata*.

Phytochemicals	Status
Tannins	+
Saponin	++
Steroids	+
Terpenoid	++
Flavonoids	++
Cardiac glycoside	+++

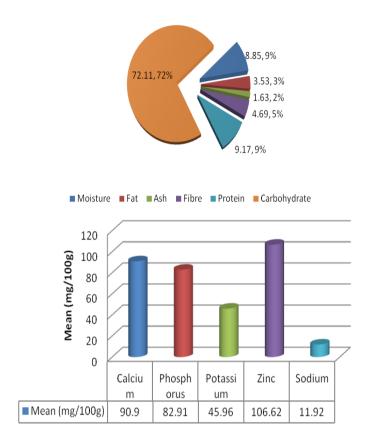
Proximate Analysis, Mineral and anti-nutrient Composition of H. umbellata

The Proximate Analysis and Mineral Composition of the *H. umbellata* indicated that the seed contain 72.11% of carbohydrate which is the highest followed by protein, moisture, fiber and fat. Ash is the least present. The mineral composition was also estimated as mg per 100 g of the seed. The result indicated zinc is the most abundant, followed by calcium, phosphorus, potassium, and with sodium as the least present mineral.

In Table no 2 is the result of the proximate analysis and mineral composition of the seed of *H.umbellata*. The carbohydrate, 72.11 \pm 0.80 %, Protein, 9.17 \pm 0.08 %, Moisture, 8.85 \pm 0.06 %, Fiber, 4.69 \pm 0.35 %, Fat, 3.53 \pm 0.44% and Ash 1.63 \pm 0.02%. The minerals include; Zinc 106.62 \pm 0.14 mg/100g, Calcium 90.9 \pm 0.08 mg/100g, Phosphorus 82.91 \pm 0.01 mg/100g, Potassium 45.96 \pm 0.02 mg/100g and Sodium 11.92 \pm 0.14 mg/100g

Table no 2:Proximate analysis and mineral composition of *H.umbellata seed*

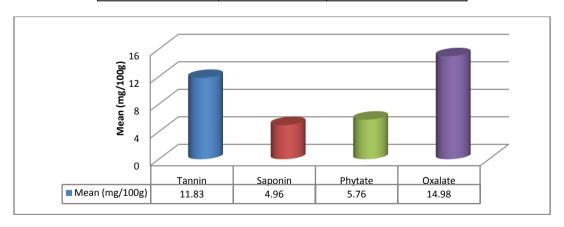
Proxima	te (%)	Mineral((mg/100g)
Moisture	8.85 ± 0.06	Calcium	90.9 ± 0.08
Fat	3.53 ± 0.44	Phosphorus	82.91 ± 0.01
Ash	1.63 ± 0.02	Potassium	45.96 ± 0.02
Fiber	4.69 ± 0.35	Zinc	106.62 ± 0.05
Protein	9.17 ± 0.08	Sodium	11.92 ± 0.14
Carbohydrate	72.11 ± 0.80		



In Table no 3 is the anti-nutrient composition of ethanolic seed extract of H.umbellata. Oxalate (16.98 \pm 0.44 mg/100g) is the most abundant anti-nutrient, with levels above the permissible limit. Phytate (5.67 \pm 0.11 mg/100g), saponin (4.96 \pm 0.07 mg/100g) and tannin (11.83 \pm 0.05 mg/100g) are well below the permissible limit.

Table no3: Anti-nutrients composition of Ethanolic seed extract of H. umbellata

	H.umbellata	Permissible limit (mg/100g)
	(mg/100g)	(Ndidiet al., 2014)
Tannin	11.83 ± 0.05	2000
Saponin	4.96 ± 0.07	50 – 200
Phytate	5.76 ± 0.11	250 – 500
Oxalate	14.98 ±0.44	0.3 - 0.5



Effect of H. umbellata on Body Weight Gain

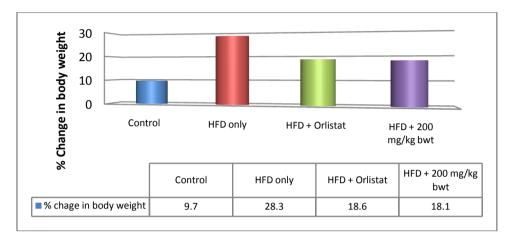
In Table no4is the result of percentage (%) change in body weight (g) on the ethanolic extract of *H. umbellata*. Therats fed with HFD showed the highest percentage change in body weight. The percentage change in body weight of rats fed with 200 mg/kg bwt and 400 mg/kg bwtwas reduced as compared to the standard Orlistat

DOI: 10.9790/3008-1504010109 www.iosrjournals.org 4 | Page

drug. The percentage change in weight of the animals were as follows: control group: 9.7 %, the HFD only: 28.3 %, HFD + Orlistat: 18.6 %, HFD + 200 mg/kg bwt: 18.1 % and HFD + 400 mg/kg bwt, 15.4%.

Table no4: Percentage (%) Change in body weight (g) on the ethanolic extract of *H.umbellata*

Groups	Initial weight (g)	Final weight (g)	Percentage Change	P value
Control	139.95 ± 5.58	153.60 ± 3.79	+ 9.7	< 0.01
HFD only	129.70 ± 7.38	166.40 ± 6.53	+ 28.3	< 0.01
HFD + Orlistat	132.5 ± 7.94	$157.2 \pm 8.92^*$	+ 18.60	< 0.01
HFD + 200 mg/kg bwt	128.00 ± 3.46	$151.20 \pm 4.35^*$	+ 18.1	< 0.01
HFD + 400 mg/kg bwt	136.90 ± 8.42	$157.90 \pm 8.66^*$	+ 15.4	< 0.01

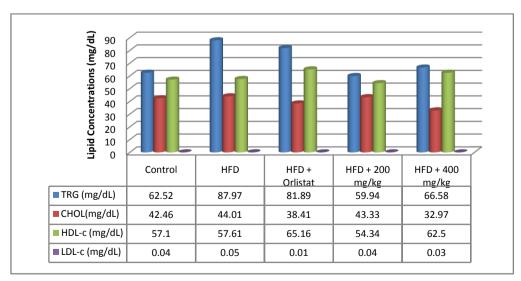


Effect of *H. umbellata* on Lipid profile:

In Table no 5is the resultof the effect of ethanolic seed extract of H. umbellata on plasma lipid profile.H. umbellata 400 mg/kg bwtsignificantly (p < 0.05) reduced the plasma triglyceride, cholesterol and LDL-c (66.58 ± 1.03 mg/dl, 32.97 ± 0.21 mg/dl, 0.03 ± 0.01 mg/dl) as compared to the HFD group87.97 ± 2.75 mg/dl,44.01 ± 0.43 mg/dl, 0.05 ± 0.00 mg/dl respectively) and the standard drug (orlistat group). Also, the plant extract at 400 mg/kg bwt significantly (p<0.05) increased the HDL-c (62.50 ± 4.05 mg/dl) as compared to the HFD group (57.61 ± 2.86 mg/dl).

Table no5:Effect of ethanolic seed extract of H. umbellata on plasma Lipid profile

Groups	TRG (mg/dL)	CHOL(mg/dL)	HDL-c (mg/dL)	LDL-c (mg/dL)
Control	62.52 ±0.87	42 46 ±0.48	57.10 ±4.22	0.04 ± 0.01
HFD	87.97 ±2.75	44.01 ±0.43	57.61 ±2.86	0.05 ± 0.00
HFD + Orlistat	81.89 ±0.76	38.41 ±0.46*	65.16 ±4.43*	$0.01 \pm 0.01^*$
HFD + 200 mg/kg bwt	59.94 ±2.35*	43.33 ±0.51	54.34 ±3.43*	0.04 ± 0.01
HFD + 400 mg/kg bwt	66.58 ±1.03*	32.97 ±0.21*	62.50 ±4.05*	$0.03 \pm 0.01^*$
p-value	p < 0.05	p < 0.05	p < 0.05	p < 0.05

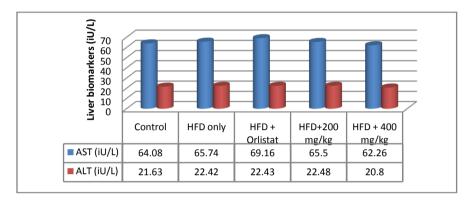


Effect of *H umbellata* on Hepatocellular injury

In Table no 6 is the resultof the effect of ethanolic extract of H. umbellateon hepatocellular injury. H. umbellateat 400 mg/kg bwt insignificantly (p > 0.05) reduced the AST and ALT (62.26 \pm 0.76iU/L and 20.80 \pm 0.42iU/L) as compared to the HFD group (65.74 \pm 1.95iU/L and 22.42 \pm 0.68iU/L respectively) and the standard drug (orlistat group).

Table no 6: Shows the effect of ethanolic extract of *H. umbellata*on hepatocellular injury

Groups	AST (iU/L)	ALT (iU/L)
Control	64.08 ± 1.50	21.63 ±0.46
HFD only	65.74 ± 1.95	22.42 ± 0.68
HFD + Orlistat	69.16 ± 1.38	22.43 ± 0.75
HFD+200mg/kg bwt	65.50 ± 3.00	22.48 ± 0.61
HFD + 400 mg/kg bwt	62.26 ± 0.76	20.80 ± 0.42
p-value	p > 0.05	p > 0.05

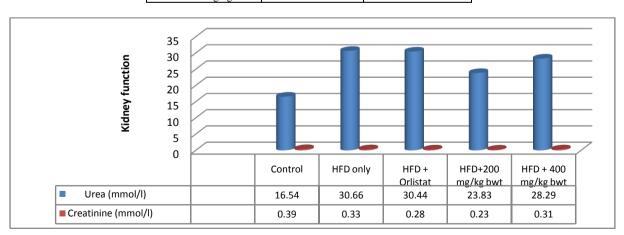


Effect of H umbellata on Kidney function

In Table no 6 is the effect of ethanolic extract of *H. umbellata*on kidney function. *H. umbellata*at 200 mg/kg bwt significantly (p < 0.05) reduced the Urea and Creatinine (23.83 \pm 2.71mmol/L and 0.23 \pm 0.03 mmol/L) as compared to the HFD group (30.66 \pm 2.08mmol/L and 0.33 \pm 0.03mmol/L respectively) and the standard drug (orlistat group).

Table no 6: Shows the effect of ethanolic extract of *H. umbellata*on Kidney function

Groups	Urea(mmol/l)	Creatinine(mmol/l)
Control	16.54 ± 2.08	0.39 ± 0.02
HFD only	30.66 ± 2.08	0.33 ± 0.03
HFD + Orlistat	30.44 ± 3.29	0.28 ± 0.02
HFD+200mg/kg bwt	$23.83 \pm 2.71^*$	$0.23 \pm 0.03^*$
HFD + 400 mg/kg bwt	$28.29 \pm 3.98^*$	0.31 ± 0.02



DOI: 10.9790/3008-1504010109

IV. Discussion

Phytochemicals are chemical compounds formed during plants normal metabolic processes; these chemicals are often referred to as secondary metabolites. The results of the present study revealed the presence of tannins, saponins, steroids, terpenoids, flavonoids and cardiac glycosides. Methanol and ethanol have been proved as effective solvents to extract phenolic compounds²⁰ which are found to be apparent in the ethanolicseed extract of *H. umbellata*(Table 2). Cardiac glycosidewhich was stronglyindicated inethanolic extract of *H. umbellates*eedis a class of natural products able to increase the cardiac contractile force in patients with congestive heart failure and cardiac arrhythmias²¹. Flavonoids and saponins have been established to possess antioxidant properties, reactive oxygen species scavenging potentials, and cell function modulation. These properties account for the large part of their pharmacological activity²².

Nutritional and proximate analyses of plants are used to evaluate their nutritional importance as they are used as food by animals as well as being employed by humans for medicinal purposes²³. The importance of mineral elements in health maintenance and protectionagainst various diseases is well documented. The result of this study revealed that *H. umbellata* contained an appreciable amount of calcium, phosphorus, zinc and sodium. Zinc serves as biocatalyst, functions as a co-factor of many enzymes in the synthesis of biomolecules. Potassium and sodiumplay a key role in the maintenance of electrical potential, healthy centralnervous system; strong teeth and bone formation. Phosphorus helps in development of reproductive and immune systemsamong other functions²⁴. Calcium is essential for bone and teeth formation and development, blood clotting and for normal functioning of heart, nervous system and muscles. Calcium deficiency can lead to ricket, osteomalacia and tooth decay²⁵. The seed of *H. umbellata* is a good source of carbohydrate for energy and a relative source of mineral (ash), protein and fat. The relatively low moisture content of *H. umbellate* indicates a possible stable plant shelf life²⁶.

In this present study, *H. umbellate*anti-nutrient contents (tannin, oxalate, phytate and saponin) were all safely below the permissible limits. Saponins have been shown to lower cholesterol concentration by competing against it for absorption in the body²⁵. Dietary oxalate has been known to complex with calcium, magnesium, and iron leading to the formation of insoluble oxalate salts resulting in oxalate stones and interferes with utilization of the minerals. Phytic acid interferes with the bioavailability of nutrients in diet and inhibits several proteolytic enzymes and amylases²⁰. Phytic acid is one of few chelating therapies for uranium removal. Phytic acid has some mineral binding properties. It may prevent colon cancer by reducing oxidative stress in the lumen of the intestinal tract^{25,26}. Tannins bind to proline-rich proteins and interfere with protein synthesis and enzyme function²⁶. Obesity, characterized by increased adipose tissue results from both increased fat cell number and increased fat size, develops from excessive energy intake and reduced energy expenditure⁶.

In the present study, the rats fed HFD for three weeks showed overweight as indicated by the significant increase in weight (Table 5). This is closely related to the significantly development of dyslipidemia. After treatment with *H. umbellata* extract, the body weight of the animals was significantly decreased. The result suggested that treatment of *H. umbellate*might have anti-hyperlipidemiceffect in HFD-induced obese rats. Ibeh*et al.*, reported that the exposure to extracts of abeere seed did not change significantly the body weights of affected rabbit, which suggests no adverse effect on metabolic activities of the rabbits treated with the seed extract²⁷.

Lipids are found in the blood and are stored in tissues; they play an important role in the body. Lipid disorders, such as high cholesterol, may lead to life-threatening illnesses, such as coronary artery disease (CAD), heart attack, or stroke⁶. Lipid profile describes the level/status of different lipids: triglycerides, totalcholesterol, high-densitylipoproteincholesterol and low-densitylipoproteincholesterol in the blood. It serves as a medical screening tool for abnormalities in lipids⁶. In this present study, (Table 6) *H. umbellate* at 200 mg/kg significantlyattenuated the effect of the HFD on the level of triglyceride and HDL-C when compared to the standard drug (orlistat). The *H. umbellata* at 400 mg/kg significantly decreased the concentrations of cholesterol and LDL-c levels when compared to the HFD only fed rats. This revealed the anti-hyperlipidemic effect of ethanolic extract of *H. umbellata*. Akpabio and Ikpe (2013) reported that phyticacidhas been implicated as having a significant effect on reducing plasma cholesterol and levels of triglyceride, this effect is thought to berelated to the ability of phytic acid to bind to zinc and thuslower the ratio of plasma zinc tocopper which is known to dispose humans to cardiovascular disease²⁶. Uzoekwe andMohammedreported that the hypocholesterolemic effect of saponins²⁵.

The liver plays a central role in the maintenance of metabolic homeostasis²⁸. The membranes of liver cells can become permeable when damaged, allowing for escape of intracellular enzymes into the bloodstream. The major intracellular enzymes are alanine aminotransferase (ALT) and aspartate aminotransferase (AST)²⁸. In this present study, there was no significant difference in the activities of liver indices (AST and ALT) of HFD fed rats when compared to the treated group (Table 7). This indicated that the HFD does not have effect on the liver. This was probably due to short term duration of the study. Several studies have reported increase in

activities of AST and ALT. Ibeh*et al.*, reported that the effect of abeere seed extracts on selected enzymes showed an enhancement in the activities of aspartate transaminase (AST) and alanine transaminase (ALT)²⁷.

The function of the kidneys is to remove wastes and excess fluid from the blood. Ureaandcreatinine are good markers that indicate regular functioning of the kidney. Urea is a waste product that is created by protein metabolism and excreted in the urine. Creatinine is a waste product of muscle energy metabolism and is produced at a constant rate that is proportional to the muscle mass of an individual. The body does not recycle it, so the quantity filtered by the kidneys in a given amount of time excreted with the urine, making creatinine clearance a specific measure of the kidney function⁴. In this present study, there was significant increase it the concentration of creatinine and urea in the HFD fed animals when compared to the control group (Table 8). However, there was significant reduction when treated with the extract at 200 mg/kg bwt. There were no significant changes when treated with the extract at 400 mg/kg bwt and orlistat at 50 mg/kg bwt. The results obtained evealed that consumption of HFD might play a role in the development of kidney impairment associated with obesity and *H. umbellate*extract exhibits attenuation in the impairment, indicated with the improved plasma urea and creatinine levels in HFD-induced obese rats. This is in constrast to the result of Hall*et al.*, who reported an insignificant increase in the levels of creatinine and urea⁴.

V. Conclusion

In conclusion, the results of this study revealed that *H. umbellata* seed has secondary metabolites and anti-nutrients within permissible limitswhich normalized the lipid profile and renal injury markers of high fat diet-induced obese rats. The seed could be a good source of bioactive substances useful in the treatment and management of dyslipidemia and related disorders. Moreover, the level of plant's secondary metabolites indicated that the *H. umbellata* may serve a medicinal and therapeutic function.

References

- [1]. Joshi PK., Chawla R A, Mistry D H,Pithawala M A. Obesity- An excuse or a blame. Eastern Journal of Medicine. 2016; 21(2): 56-63
- [2]. Dalboge LS, Pedersen PJ, Hansen G, Fabricius K, Hansen HB, Jelsing J. A hamster model of diet-induced obesity for preclinical evaluation of anti-obesity, anti-diabetic and lipid modulating agents. PLoS One. 2015; 10(8):e0135634.
- [3]. Tang J. Yan H,Zhuang S. Inflammation and oxidative stress in obesity-related glomerulopathy.Int. J.Nephrol. 2012: doi:10.1155/2012/608397.
- [4]. Hall JE, Henegar JR, Dwyer TM, Liu J, Da Silva AA, Kuo JJ, Tallam L. Is obesity a major cause of chronic renal disease? Adv.Ren. Replace Ther. 2004; 11: 41-54.
- [5]. Choudhury D, Tuncel M, Levi M. Disorders of lipid metabolism and chronic kidney disease in the elderly. SeminNephrol. 2009; 29(6):610-620
- [6]. Sasatomi Y, Tada M, Uesugi N, Hisano S, Takebayashi S. Obesity associated with hypertension or hyperlipidemia accelerates renal damage. Pathobiology. 2001; 69(2):113-118
- [7]. Adeneye AA, Adeyemi OO,Agbaje EO. Anti-obesity and antihyperlipidaemic effect of *Hunteriaumbellata* seed extract in experimental hyperlipidemia. Journal of Ethnopharmacology. 2010^a;130: 307-314.
- [8]. Oboh G, Adebayo AA,Ademosun AO, Erection-stimulating, anti-diabetic and antioxidant properties of HunteriaumbellateandCylicodiscusgabunensiswater extractable: phytochemicals. Journal of Complementary and Integrative Medicine. 2017: 20160164.
- [9]. Association of Official Analytical Chemists (AOAC). Official methods of Analysis of AOAC international.17th ed. U.S.A. Gatherburg, 2000
- [10]. Oyedapo OO, Sab FC,Olagunju JA. Bioactivity of fresh leaves of Lantana camara. Biochemical Letters.1999; 59, 175-183.
- [11]. TreaseGE, Evans WC. Pharmacognosy.15th Ed.LondonSaunders Publishers. 2002; pp. 42-393.
- [12]. Sofowora EA. (2006). Medicinal Plants and Traditional Medicine in Africa. Second Edition. Ibadan. Spectrum Books Limited, Nigeria.
- [13]. Price M, Van Scoyoc S, Butler LG. A critical evaluation of the vanillin reactions as an assay for tannin in sorghum grain. Journal of Agriculture and Food Chemistry. 1978; 26: 1214.
- [14]. Brunner JH. Direct spectrophotometric determination of saponin. Annalsof Chemistry. 1984; (34): 1314-1326
- [15]. Reddy NR, Balakrishman CV,SalunkheDK..Phytate phosphorus and mineral changes during germination and cooking of blackgram (Phaseolus mango L.) seeds.Journal of Food Science.1978; 43: 540-542.
- [16]. Fashakin JB, Ilori MO, Olarewaju I. Cost and Quality Optimization of a complementary diet from plant protein and corn flour using a computer aided Linear Programming Model. Nigerian Food Journal. 1991; 9: 123-127
- [17]. Aletor O.Nutrient composition of soya bean meal and soy protein isolate and their effects on the growth performance and haematological parameters of albino rats. FUTA Journal of Research in Sciences.2013; 2: 259-268
- [18]. Friedewald WT, Levy RI, Fredrickson DS.Estimation of the concentration of low density lipoproteins cholesterol in plasma without the use of propagative centrifuge. Journal of Clinical Chemistry. 1972; 18: 499.
- [19]. Chattodhyaypa RR,Bandyopadhyay. Effects of Azadirachiaindicaleaf extract on serum lipid profile changes in normal and streptozotocin induced diabetic rats, African Journal of Biomedicinal Research. 2005; 8: 101 104.
- [20]. Siddhuraju P, Mohan PS,Becker K.Studies on the antioxidant activity of Indian Laburnum (Cassia fistula L): a preliminary assessment of crude extracts from stem bark leaves, flowers and fruit pulp. Food Chem. 2002; 79:61-67.
- [21]. Hauptman PJ, Garg R, Kelly RA. Cardiac glycosides in the next millennium. Prog. Cardiovasc. Dis. 1999; 41: 247 254.
- [22]. Olabinri BM, Adepoju EA, Zainab AA, Ahmed AA.Phytochemical profiling of phytoconstituents of grape, Jatrophacurcas and Neem (Azadirachtaindica) extracts. J. PharmacognosyPhytother. 2014; Vol. 6(2), pp. 17-23
- [23]. Hussain J, Khan A, Rehman N, Zainullah KF, Hussain ST, Shinwari ZK. Proximate and Nutritional Investigations of Selected Medicinal Plants Species of Pakistan. Pakistan Journal of Nutritional. 2009; 8(5): 620-324.

Phytochemical Analysis and Anti-Hyperlipidemic Potential of Ethanolic Extract of ...

- [24]. Mohammed MI, Sharif N. Mineral composition of some leafy vegetables consumed in Kano, Nigeria, Nig. J. Basic Appl. Sci. 2011; 19(2): 208-212
- [25]. Uzoekwe NM, Mohammed JJ.Phytochemical, proximate and mineral contents of leaves and bark of Ficuscapensis. Journal of Applied Science and Environmental Management. 2015; 19(4): 633-637.
- [26]. Akpabio UD, Ikpe EE. Proximate composition and nutrient analysis of Aneilemaa equinoctial eleaves. Asian Journal of Plant Science and Research. 2013; 3(2): 55-61.
- [27]. IbehIN, IduMD, Ejimadu IM.Toxicological assessment of 'abeere' seed *hunteriaumbellate* mk. Schum (apocyanceae)BIOCIÊNCIAS, Porto Alegre. 2007; v15:04 07
- [28]. Olufunmilayo LA, Johnson M, Adeleke AP, Modupe AS.Hypoglycemic Effects of the Methanolic Seed Extract of Hunteriaumbellate(Abeere) and Its Effect on Liver, Hematological and Oxidative Stress ParametersinAlloxan-Induced Diabetic Male Albino RatsInt. J. Curr. Res. Biosci. Plant Biol. 2015; 2(6): 27-34

Morakinyo A. E, et. al. "Phytochemical Analysis and Anti-Hyperlipidemic Potential of Ethanolic Extract of *Hunteriaumbellata* Seed." *IOSR Journal of Pharmacy and Biological Sciences (IOSR-JPBS)*, 15(4), (2020): pp. 01-09.

DOI: 10.9790/3008-1504010109 www.iosrjournals.org 9 | Page