

## **Original Research Article**

### **Biochemical Indices and Haematological Studies of Ethyl acetate Extract of *Persea americana* Leaf in Albino Rats**

#### **ABSTRACT**

The aim of this present study was to evaluate the effect of ethyl acetate extract of *P. americana* (avocado) on liver and kidney function, lipid profile as well as haematological parameters in albino rats. A total of 20 albino rats were used for this experiment and they were divided into four groups of 5 (A-D) rats each. Group A served as normal control, group B-D served as experimental groups administered with 100, 200 and 400mg/kg body weight of ethyl acetate extract of *Persea americana* leaf per day for 4 weeks respectively. This study was conducted in the Department of Biochemistry, Bayero University, Kano, in the month of May, 2018. Liver function test (colorimetric method), kidney indices and lipid profile (spectrophotometry method), and hematological examination (SYSMEX XE-2000) were analysed. The extract was found to significantly increase ( $P=.05$ ) the levels of serum ALT and albumin in the group administered with 400mg/kg of the extract, whereas total protein and globulin levels were significantly ( $P=.05$ ) decreased in all the treated groups. The extract significantly ( $P=.05$ ) decrease total cholesterol, triglyceride and low density lipoprotein in all the treated groups. Haematological parameters analysed were not significant affected in all the treated groups. In conclusion, ethyl acetate extract of *P. americana* may be safe and possesses hypolipidemic potentials but extremely high doses may not be advisable.

Keywords: *P. americana*, liver function test, kidney function test, lipid profile, haematological parameters

#### **1. INTRODUCTION**

Many medicinal plants used in ethnomedical practices in Nigeria are known or little known to scientific world [1]. Many important drugs used in medicine today are directly or indirectly derived from plants [2]. Plants and herbs have been a tremendous source of food and folk remedies for mankind and have served as starting material for the development of new synthetic drugs. They have the ability to synthesize a wide variety of chemical compounds which are used to perform biological functions and to defend against attack from predators such as insects, fungi, yeasts, bacteria, virus and other pathogens [3]. Chemical compounds in plants mediate their effect on the human body through processes identical to

those already well understood for the chemical compounds in conventional drugs [4, 5]. *Persea americana* belongs to the family *Lauraceae*, unflatteringly known in the past as alligator pear, midshipman's butter and vegetable butter, it is one of those plants currently used by indigenous persons for its nutritional value and to manage health problems. It is well known in many parts of the tropical world including Nigeria [6] and is commonly called avocado or pear. The fruit tree can attain a height up to 20m, with large spreading and flat topped crown. The plant is reported to possess antidiabetic, antihyperlipidemic potentials [2, 7], antioxidant [8, 9], cancer risk reduction [10], wound healing [11], hepatoprotection [12], analgesic and anti-inflammatory [13], anticonvulsant [14], vasorelaxant and blood pressure reducing [15, 16]. The use of traditional medicines as substitutes to orthodox medicines has been on the increase [17]. The reasons, which have given rise to this trend, include the cheapness, availability and accessibility of these natural medicines. Besides, there has been the erroneous belief that these medicines are free from adverse effects [18, 19]. On the other hand they have been rejected because many of the acclaimed medicinal values have not been scientifically evaluated and their safety profiles uncertain [19]. It is, therefore, pertinent that safety assessments should be conducted on natural products for which certain medicinal uses have been scientifically validated. It was recommended by the World Health Organization to evaluate herbal medicine, as orthodox medicines appear not to be safe [20]. It is, therefore, relevant that safety assessments should be conducted on ethyl acetate extract of *P. americana* leaf for which certain medicinal uses have been scientifically proven.

## **MATERIALS AND METHODS**

### **2.1 Sample Collection and Identification**

The leaf of *Persea americana* were collected from Jos, Plateau state of Nigeria. The plant part was authenticated by a Botanist at Plant Science Department, B.U.K. with accession number BUKHAN 0305 and was deposited in Bayero University, Kano herbarium for reference.

### **1.2 Extraction of Plant Material**

Soxhlet extraction method was used for the extraction of the plant materials. The sample (200g) was chopped into small pieces and then shade dried and ground into powdered form (120g). Ethyl acetate

was used as the extraction solvent and later concentrated in vacuo using rotary evaporator at 40°C resulting to a total extraction yield 35.5% (29.6g).

### 1.3 Experimental animals

Experimental rats (80-100g) of same sex (males) were obtained from the Physiology department, Faculty of Basic Medical Sciences, Bayero University, Kano. The animals were kept in cages and clean drinking water provided *ad libitum* while they were fed with standard commercial pelleted feed (Vital Feed® Nigeria). The temperatures varied between 27-30°C and relative humidity of about 55%-60% with 12-h light-dark cycle and adequate ventilation maintained in the animal house. This study was conducted in the Department of Biochemistry, Bayero University, Kano, in the month of May, 2018. Ethical conditions governing the conducts of experiments with life animals as stipulated were strictly observed. Also, the experimental protocol was approved by the College of Health Science ethical committee with a reference number 0653.

### 2.4 Experimental Design

A total of 20 rats were used for this experiment and they were divided into four groups of 5 rats each.

Group 1- normal rats

Group 2- treated with ethyl acetate extract of *P. americana* leaf (100mg/kg body weight).

Group 3- treated with ethyl acetate extract of *P. americana* leaf (200mg/kg body weight).

Group 4- treated with ethyl acetate extract of *P. americana* leaf (400mg/kg body weight).

The sub-chronic toxicity study lasted for a period of four weeks. The various doses (100, 200 and 400mg/kg body weight.) were administered to group 2, 3 and 4 respectively of normal rats. At the end of 4 weeks treatment the rats were sacrificed and blood samples collected for liver function (Alanine amino transferase, Aspartate amino transferase, Alkaline phosphatase, Albumin, Total protein and Globulin), kidney function indices [creatinine, urea, electrolytes ( $\text{Na}^+$ ,  $\text{K}^+$ ,  $\text{Cl}^-$  and  $\text{HCO}_3^-$ )], lipid profile (Cholesterol, Triglyceride, High density lipoproteins, Low density lipoproteins and very low density lipoproteins) and hematological parameters (White blood cells, Red blood cells, Haemoglobin, Packed cell volume, Mean

corpuscular volume, Mean corpuscular haemoglobin, Mean corpuscular haemoglobin concentration and Platelets).

## 2.5 Assays

Liver function test were assayed using the methods of; serum alanine and aspartate amino transferase activity determination by Reitman and Frankel, [21], serum alkaline phosphatase activity determination by Rec, [22], serum albumin concentration determination by Grant, [23], serum total protein and serum globulin concentration determination by Tietz, [24]. Kidney function test were assayed using the methods of; serum urea concentration determination by Weatherburn, [25], serum creatinine concentration determination by Bartels and Bohmer, [26], serum potassium concentration determination by Henry, [27], serum sodium concentration determination by Kraut and Madias, [28], serum chloride concentration determination by White, [29] and serum bicarbonate concentration determination by Forrester et al. [30].

Lipid profile was analysed using the methods of; serum total cholesterol concentration determination by Lothar, [31], triglycerides concentration and serum HDL-cholesterol concentration determination by Jacobs et al. [32], serum LDL and VLDL-cholesterol concentration determination by Friedewald et al. [33].

Haematological parameters were analyzed using SYSMEX XE-200 (QBC Autoread Plus, UK).

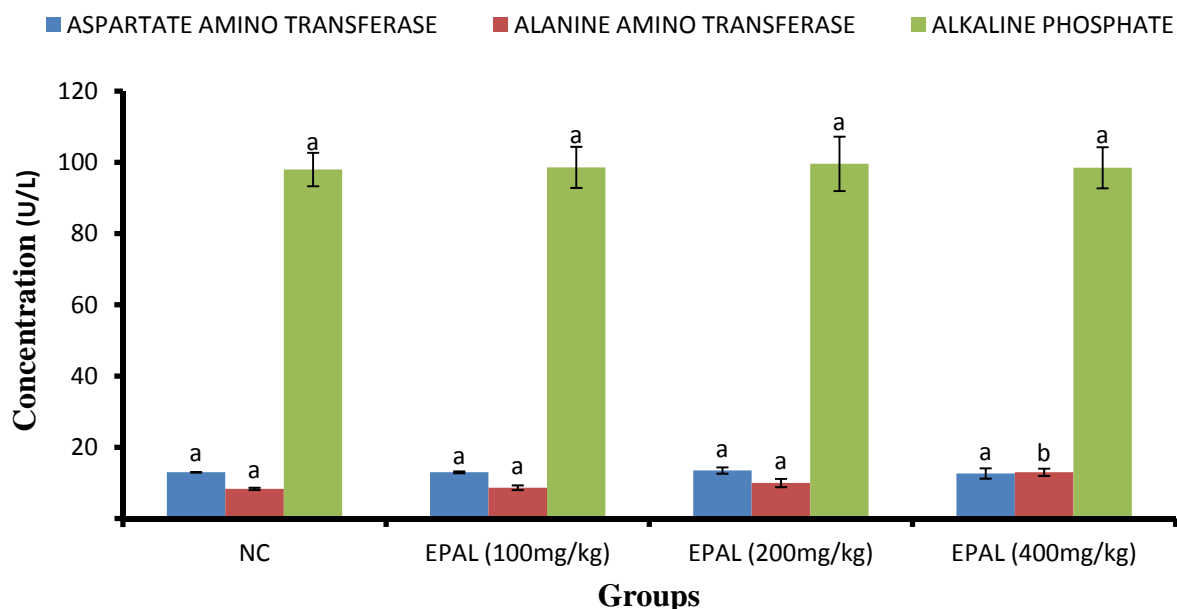
## 2.5 Statistical Analysis

Statistical package for the social sciences (SPSS) version 17 software was used for all calculations and statistical analysis. Analyses were performed using student t-test at 95% confidence level with P=.05 being significant. Results were presented as mean  $\pm$  standard deviation.

### 3 RESULTS

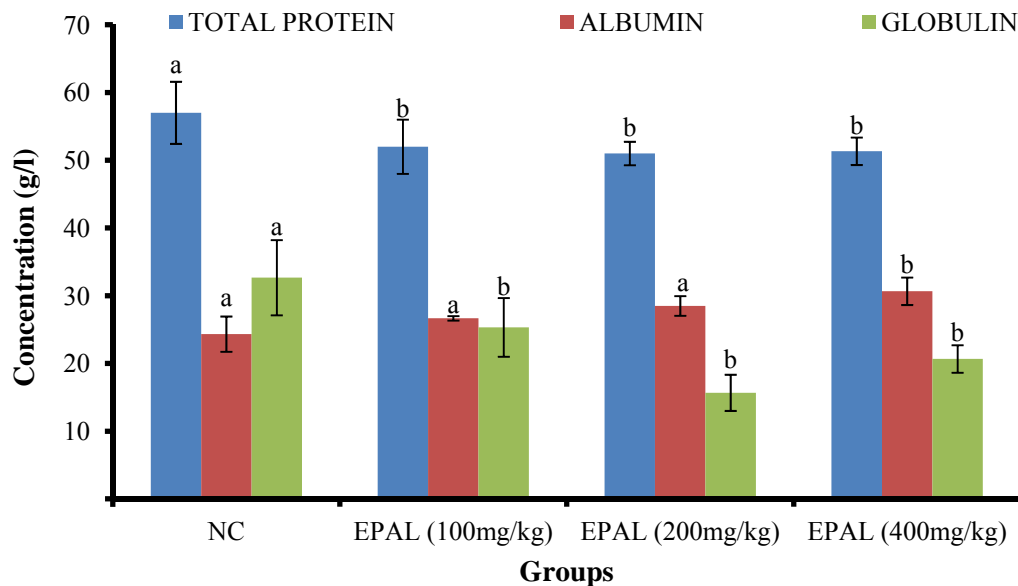
#### 3.1 Liver Function Test

Result of sub-chronic toxicity study showed no significant difference ( $P=0.05$ ) in AST and ALP activities in all the treated groups when compared with normal control. However, significant increase ( $P=0.05$ ) was observed in serum level of ALT of the entire treated group when compared with normal control (Figure 1). There was a significant decrease ( $P=0.05$ ) in serum total protein and globulin in all the treated groups compared with the normal control (Figure 2). The level of serum albumin was found to increase significantly ( $P=0.05$ ) in all the treated groups compared with normal control.



**Figure 1: Serum Levels of Aspartate Amino Transferase, Alanine Amino Transferase and Alkaline Phosphatase of Rats Administered with Ethyl acetate Extract of *P. americana* leaf (100, 200 and 400mg/kg b.w.) for 4 Weeks**

Results are presented as Mean  $\pm$  SD,  $n=5$ . Values with different superscripts (a or b) are significantly different ( $P<0.05$ ) with respect to normal control, NC= Normal control, EPAL= Ethyl acetate extract of *P. americana* leaf

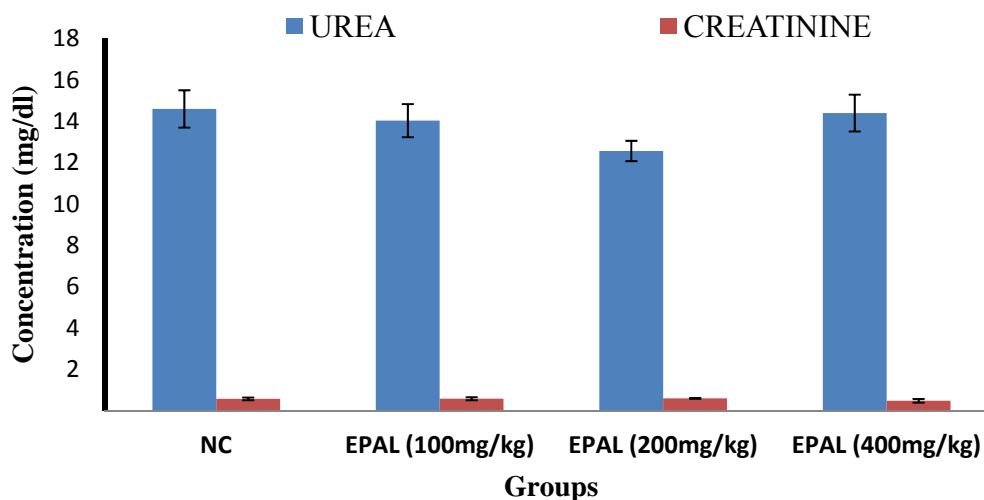


**Figure 2: Serum levels of Total Protein, Albumin and Globulin of Rats Administered with Ethyl acetate Extract of *P. americana* Leaf (100, 200 and 400mg/kg b.w.) for 4 Weeks**

Results are presented as Mean  $\pm$  SD, n=5. Values with the different superscripts (a or b) are significantly different ( $P < 0.05$ ) with respect to normal control, NC=Normal control, EPAL=Ethyl acetate extract of *P. americana* leaf

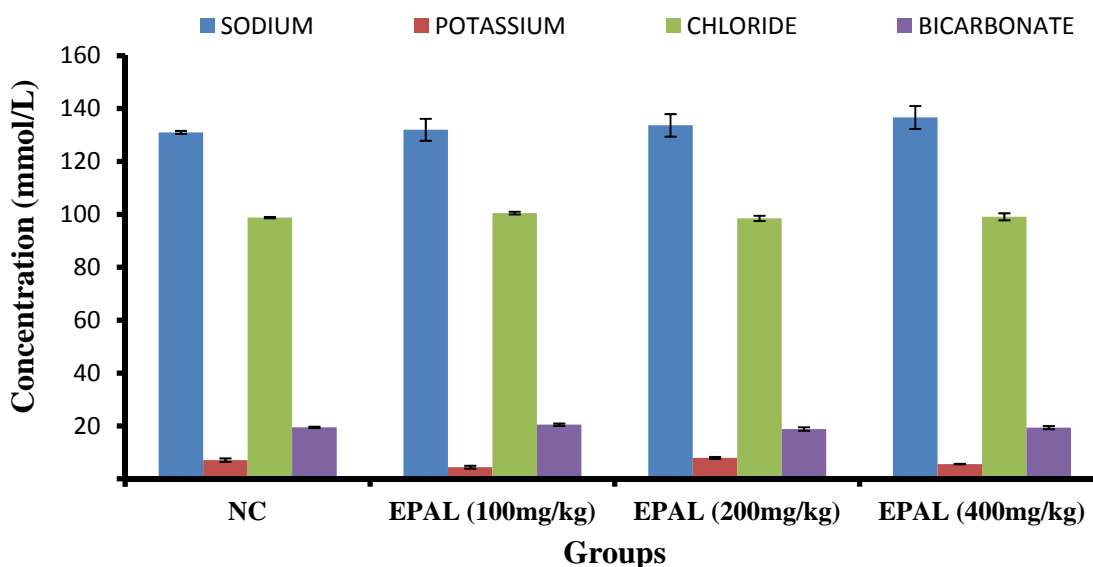
### 3.2 Kidney Function Test

Kidney function tests carried out include serum urea, creatinine, sodium, potassium, chloride and bicarbonate (Figures 3 and 4). No significant difference ( $P = 0.05$ ) was observed in serum urea, creatinine, sodium, chloride and bicarbonate in all the groups when compared with normal control group. Serum Potassium level was found to decrease significantly in group 2 (100mg/kg b.w.) When compared with normal and groups 3 and 4 (200 and 400mg/kg b.w.).



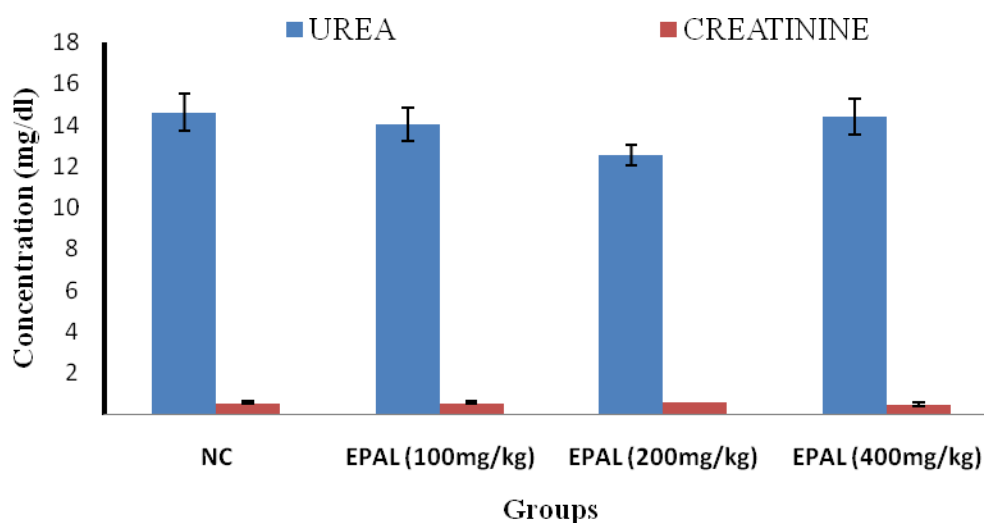
**Figure 3: Serum Levels of Urea and Creatinine Levels of Rats Administered with Ethyl acetate Extract of *P. americana* Leaf (100, 200 and 400mg/kg b.w.) for a Period of 4 Weeks**

Results are presented as Mean  $\pm$  SD, n=5. Values with the different superscripts are significantly different ( $P < 0.05$ ) with respect to normal control, NC= Normal control, EPAL= Ethyl acetate extract of *P. americana*



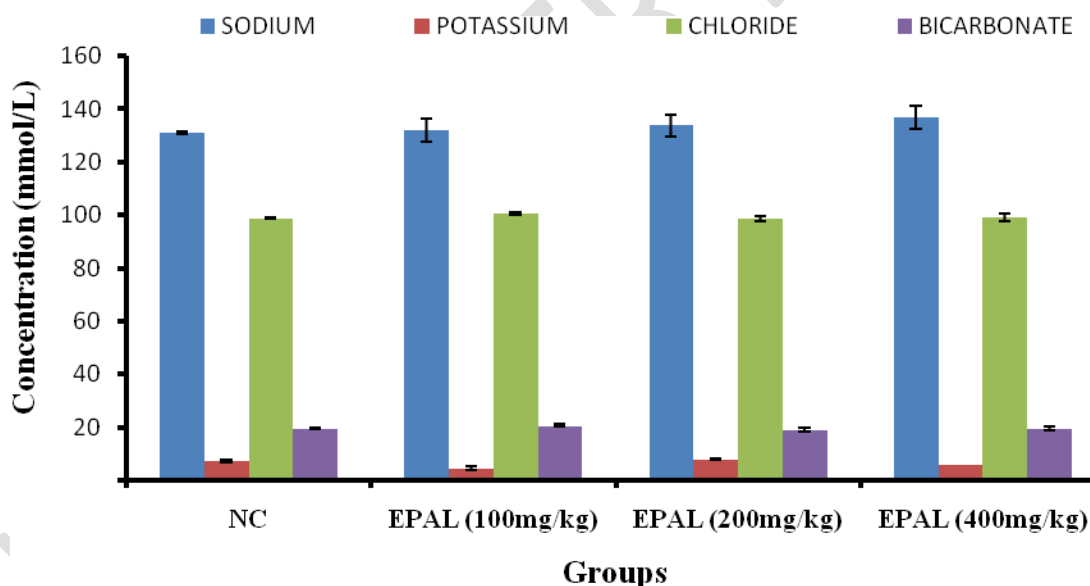
**Figure 4: Serum Levels of Sodium, Potassium, Chloride and Bicarbonate of Rats Administered with Ethyl acetate Extract of *P. americana* Leaf (100, 200 and 400mg/kg b.w.) For 4 Weeks**

Results are presented as Mean  $\pm$  SD, n=5. Values with the different superscripts (a or b) are significantly different ( $P < 0.05$ ) with respect to normal control, NC= Normal control, EPAL= Ethyl acetate extract of *P. americana*



**Figure 3: Serum Levels of Urea and Creatinine Levels of Rats Administered with Ethyl acetate Extract of *P. americana* Leaf (100, 200 and 400mg/kg b.w.) for a Period of 4 Weeks**

Results are presented as Mean  $\pm$  SD, n=5. Values with the different superscripts are significantly different (P < 0.05) with respect to normal control, NC= Normal control, EPAL= Ethyl acetate extract of *P. americana*



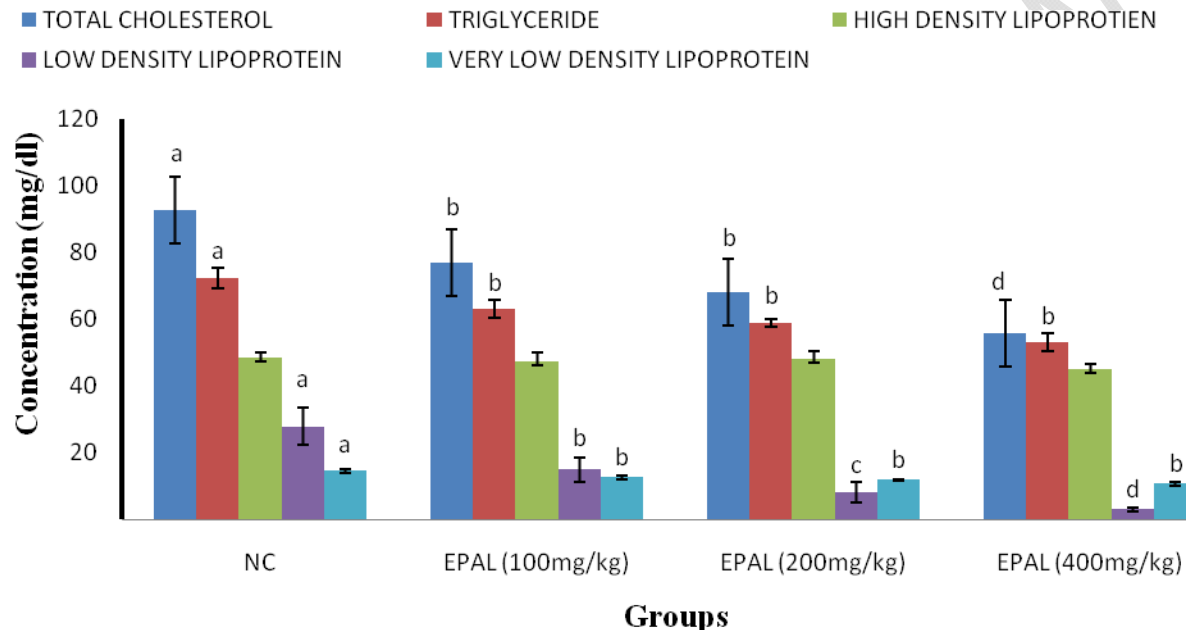
**Figure 4: Serum Levels of Sodium, Potassium, Chloride and Bicarbonate of Rats Administered with Ethyl acetate Extract of *P. americana* Leaf (100, 200 and 400mg/kg b.w.) For 4 Weeks**

Results are presented as Mean  $\pm$  SD, n=5. Values with the different superscripts (a or b) are significantly different (P < 0.05) with respect to normal control, NC= Normal control, EPAL= Ethyl acetate extract of *P. americana*



### 3.3 Lipid Profile

The levels of serum total cholesterol, triglyceride and low density lipoprotein in the treated groups were found to decrease significantly ( $P=0.05$ ) when compared with normal control in a dose dependent pattern within the treated groups. No significant difference was observed ( $P=0.05$ ) in serum high density lipoprotein and very low density lipoprotein levels (Figure 5).



**Figure 5: Levels of Serum Total Cholesterol, Triglyceride, High Density Lipoprotein, Low Density Lipoprotein and Very Low Density Lipoprotein of Rats Administered with Ethyl acetate Extract of *P. americana* Leaf (100, 200 and 400mg/kg b.w.) for 4 Week**

Results are presented as Mean  $\pm$  SD,  $n=5$ . Values with the different superscripts (a or b) are significantly different ( $P<0.05$ ) with respect to normal control, NC= Normal control, EPAL= Ethyl acetate extract of *P. americana* leaf

### 3.4 Haematological Parameters

There was no significant difference ( $P=0.05$ ) between the normal control and the tested groups in all the haematological parameters assessed. These include, white blood cell, red blood cell, haemoglobin, packed cell volume, mean corpuscular volume, mean corpuscular hemoglobin, mean corpuscular hemoglobin concentration and platelets (Table 1).

**Table 1: Haematological Parameters of Rats administered with Ethyl acetate Extract of *P. americana* leaf (100, 200 and 400mg/kg b.w.) for 4 weeks**

GROUPS	RBC( $10^3/\mu\text{l}$ )		WBC( $10^6/\mu\text{l}$ )		Hb(g/dl)		PCV(%)		MCV(FL)		MCHC(g/dl)		MCH(pg)		PLT( $10^3/\mu\text{l}$ )	
Normal Control	6.45	$\pm .10^a$	8.23	$\pm .20^b$	7.60	$\pm .36^c$	36.67	$\pm 1.86^d$	55.30	$\pm .06^e$	21.80	$\pm .23^f$	11.23	$\pm .35^g$	346.67	$\pm 3.22^h$
EE(100mg/kg)	6.47	$\pm .14^a$	8.73	$\pm .23^b$	6.97	$\pm .27^c$	37.67	$\pm 1.86^d$	58.07	$\pm .54^e$	22.93	$\pm .17^f$	11.83	$\pm .19^g$	336.67	$\pm 3.93^h$
EE (200mg/kg)	6.73	$\pm .04^a$	8.53	$\pm .35^b$	8.03	$\pm .35^c$	36.67	$\pm .88^d$	55.20	$\pm .47^e$	23.33	$\pm .98^f$	11.40	$\pm .42^g$	349.00	$\pm 8.50^h$
EE (400mg/kg)	6.63	$\pm .22^a$	8.77	$\pm .09^b$	7.47	$\pm .35^c$	37.67	$\pm 2.40^d$	53.00	$\pm .78^e$	22.23	$\pm .73^f$	11.60	$\pm .40^g$	357.33	$\pm 4.10^h$

Results are presented as Mean  $\pm$  SD, n=5. Values with the different superscripts in the same column are significantly different ( $p < 0.05$ ) with respect to normal control, EE= Ethyl acetate Extract of *P. americana* leaf

## 4 DISCUSSION

Toxicity studies are essential tools in assessing the bioaccumulative effects of xenobiotics in biological systems. Assessment of liver and kidney function is very important in evaluating toxicity of modern and traditional medicines since these organs play major roles in metabolism of xenobiotics in the body. Elevated activities of liver enzymes are often diagnostic of underlying cellular injuries [34, 35]. From the biochemical result of this study, AST and ALP activities were not affected in all the groups administered with the extract. Serum concentration of ALT was found to increase in the group treated with 400mg/kg, and this can be as a result of increase in hepatocytes proliferation or regeneration [36]. Chemicals often cause subclinical injury to the liver, which manifests only as abnormal liver enzyme tests. Certain medicinal agents, when taken in overdoses and sometimes even when introduced within therapeutic ranges may injure the organ [37]. Studies conducted by [38] showed that a large intake of tannins may cause liver damage. Reduction in total protein and globulin are indications of diminished synthetic function of the liver or might be due to impaired hepatocellular function [39, 40, 41, 42]. The increased level of albumin observed in this study depends on a number of factors such as the nutritional status, catabolism, hormonal factors, urinary and gastrointestinal losses [42]. Ethyl acetate extract of *P. americana* was found to contain appreciable amount of saponins and tannins [8, 43] which can be related to the observed increase in ALT activity. Saponins may act by damaging cell membranes causing leakage of cellular materials, ultimately leading to cell death [44, 45].

Kidney function was evaluated by means of serum urea, creatinine and blood electrolyte concentrations. Urea is formed in the liver and is mainly excreted by the kidney, it is useful in evaluating kidney function in conjunction with creatinine which originates from the muscle and is filtered by the kidney. The serum urea and creatinine levels of treated groups in this study were not significantly changed. Increased blood urea and creatinine is a good indicator of compromised kidney function. These results suggest that the extract may not have altered the kidney function. In a similar study by [40] serum creatinine, urea and electrolytes such as sodium and calcium were not affected after *P. americana* extract treatment.

In this study, the administration of graded doses of EPAL significantly reduced serum levels of TC, TG, and LDL in treated rats. Saponins are known anti-nutritional factors, which lower cholesterol by binding

with cholesterol in the intestinal lumen, preventing its absorption, and/or by binding with bile acids, causing a reduction in the enterohepatic circulation of bile acids and increase its faecal excretion [46, 47]. Increased bile acid excretion is offset by enhanced bile acid synthesis from cholesterol in the liver and consequent lowering of the plasma cholesterol; hence saponins have been reported to have hypocholesterolaemic effect [48]. Thus, the observed hypolipidemic effect of EPAL may be linked to the synergistic actions of phytochemicals like saponins and polyphenolic compounds contained in the plant extract.

The haematopoietic system is one of the most sensitive targets of toxic compounds and is an important index of physiological and pathological status in man and animals as it investigate the extent of damage to the blood [37]. It provides vital information regarding the status of bone marrow activity and intravascular effects such as haemolysis and anaemia [49]. Assessment of the haematological indices showed that the extract did not cause any significant effect on WBC, RBC, PCV, Hb, PLT, MCV, MCH and MCHC. The results on the haematological parameters indicate normal haemopoiesis and absence of anaemia confirming the non-toxic nature of the extract.

## CONCLUSION

While the medicinal importance of the *P. americana* is increasingly becoming of scientific interest, the consumption of high dosage should be discourage, as oral administration of ethyl acetate extract of *P. americana* leaf results in an increase in the level of ALT in group administered with 400mg/kg. The extract was found to possessed hypocholesterolemic activity with no significant effect in kidney function and haematological parameters tested.

## ETHICAL APPROVAL

All authors hereby declare that; principle of laboratory animals care (NHI publication number 829 revised 1985) were followed, as well as all experiment have been examined and approved by the appropriate ethic committee.

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Figure 6: Image of *P. americana* (avocado) tree.