

## Original Research Article

### CHANGES IN LIVER HISTOLGY, HEMATOLOGICAL PARAMETERS AND LIPID PROFILE OF CADMIUM-EXPOSED RATS TREATED WITH COMBINED LEAF EXTRACT OF *VERNONIA AMYGDALINA* AND *OCCIMUM GRATISSIMUM*

#### ABSTRACT

**Aim:** This study was aimed at **examining** the effect of mixture of *Ocimum gratissimum* extract and *Vernonia amygdalina* extract on lipid profile and hematological parameters of Wistar rats administered with cadmium.

**Methodology:** Sixteen female rats were divided into four groups and designated as follows: Group 1-Control (Normal Saline,) Group 2-(Cadmium 10mg/kg bwt), Group 3 (Normal saline, leaf extract 200kg/kg bwt) Group 4 (Cadmium 10mg/kg bwt and leaf extract 200kg/kg bwt) and were treated for 28 days (4 weeks).

**Results:** Exposure of rats to Cadmium alone (Group B) brought about significant increase in levels of Cholesterol, TAG and LDL with a decrease in HDL compared to control and rats maintained on plant extract alone (Group 3). However, treatment of Cd-exposed rats with mixture of the plant extracts (Group 4) significantly reduced Cholesterol, TAG and LDL with an increase in HDL compared to rats maintained on Cd alone (Group 2). The PCV and RBC of the rats administered with the cadmium showed a significant reduction when compared to the control group, whereas the WBC increased significantly ( $p < 0.05$ ). The group administered with the leaf extract and a combination of the leaf extract and cadmium recorded an increased in the (PCV), (RBC) level of the Wistar rat and a reduction in the (WBC) level when compared to the group administered with cadmium.

**Conclusion:** The findings of this study demonstrate that mixture of *Ocimum gratissimum* and *Vernonia amygdalina* extracts have lipid lowering properties that may be advantageous to those with the problem of cadmium toxicity. The extracts were also found to be efficient in decreasing cholesterol, **triacylglyceride**, and low density lipoprotein levels, indicating that they have **hypcholesterolaemic** properties.

**KEYWORDS:** *Vernonia amygdalina*; *Occimum gratissimum*; **haematology**, lipid profile.

## Introduction

Cadmium (Cd) is ubiquitous in nature and to great extent it is concentrated in the food chain due to its high soluble nature compared to other toxic heavy metals. It is not degradable, consequently it is easily transported from soil to plants which animals and humans largely depend on for survival <sup>[1]</sup>. Studies conducted by Vasey, [2] stated that there are two main ways for ionic form of Cd<sup>2+</sup> to get through the hepatocyte cell membrane: (a) binding with Fe<sup>2+</sup> and Zn<sup>2+</sup> transporters, or (b) through voltage-gated Ca<sup>2+</sup> channels. Protein-bound Cd usually binds to liver-produced metallothionein (MT) protein to form Cd metallothionein (Cd-MT) complex, which enters cells through receptor-mediated endocytosis and is then released from the Cd-MT as a Cd ion through the digestion of lysosome. Chronic Cd ion is stored in various tissues such as the liver, kidney, prostate and bone [3]. In the female liver, Cd absorption is 10%–20% higher than in males, and the female liver is more susceptible to Cd toxicity. This difference may be related to progesterone-activated receptor-dependent calcium channels, channels that are involved in the absorption and accumulation of Cd into the liver [4]. It was also documented in a study that the deposition of Cd in the liver can cause both liver injury and hepatotoxicity [5].

Extensive research on the molecular mechanism of Cd carcinogenesis has shown that chronic Cd exposure can induce oxidative stress interfere with gene expression [6], affect cell cycle regulation, inhibit cell apoptosis, induce inflammatory signaling, and promote genomic instability and mutation in key genes to promote tumorigenesis [7]. Indeed, high concentrations of Cd from acute exposure or low concentrations of chronic exposure are both linked to severe hepatotoxicity/liver injury that promotes liver diseases and HCC development.

Generally, the properties of antioxidant of various plants have been applied in the treatment of different diseases especially in developing countries where they have been documented in traditional medicine [8]. A study had revealed about 80% of individuals with the use of traditional medicine in developing countries to meet up their primary health care needs [9]. The phytochemical evaluation of *Ocimum gratissimum* (OG) revealed the following bioactive compounds; flavonoids, triterpenes, alkaloids, citral, saponins, eugenol, and thymol etc <sup>9</sup>. Flavonoids compounds have been described with biological activities such as, anti-inflammatory, antitumor and antioxidant [10] activities. A study conducted by Udi, et al. [11] on the effects of aqueous extract of *Ocimum gratissimum* on the cerebellum of male wistar rats challenged by lead acetate revealed ameliorating qualities of *Ocimum gratissimum* as it reduce the degeneration of the purkinje cells. This also agree with the study on effects of Lead II Acetate induced PhysioMorphological Changes in Prefrontal Cortex of *Ocimum gratissimum* fed wistar Rats as *Ocimum grassimum* mitigate the effects of lead acetate on the Pyramidal cells of the prefrontal Cortex [12].

*Vernonia amygdalina*, popularly called bitter leaves belong to the family Asteraceae or Compositae, is consumed locally as food and for ethno-medicinal uses. The bitter taste of *Vernonia amygdalina* comes from the phytochemicals of the leaves, which include alkaloids, saponins, glycosides, and tannins [13].

Naringenin also restored the levels of antioxidant defense to normal levels and preserved the normal histological architecture of the liver tissue [14]. Similarly, a protective effect of alpha-tocopherol (vitamin E) on Cd toxicity in rat liver has been shown. Coadministration of vitamin E (300 mg/Kg/day for 3 weeks) decreased prooxidative state hepatic markers such as malondialdehyde (MDA) and peroxidase (POD) activities that are induced by Cd exposure and also increased superoxide dismutase (SOD) and catalase (CAT) activities, restored Ca levels, and improved liver architecture [15]. In addition, olive oil and colocynth oil prevented oxidative damage in Wistar rat livers induced by Cd. Cotreatment with olive oil or colocynth oil restored the antioxidant potential in plasma and liver and decreased MDA levels and transaminase activity [16]. Ferulic acid (FA) derivatives of curcumin also contribute to liver repair. **An interesting recent study demonstrated that supplementation of FA (50 mg/kg) significantly decreased Cd accumulation in rat liver and kidney tissues by elevating antioxidant enzyme expression and by decreasing the expression of hepatonephrotoxicity enzymes [17].** The aim of this study was therefore to examine the ameliorative effects of combined leaf extracts of *Ocimum gratissimum* and *Vernonia amygdalina* on the lipid profile, hematology and histomorphology of Cadmium induced liver of an adult Wistar rats.

## MATERIAL AND METHODS

### **MATERIALS (prefer to remove by author) no need**

#### **Experimental and laboratory Equipment/Apparatus**

The Materials used in this experimental study are as follow: Dissecting set, Wistar rats, Cadmium Chloride, animal cages, fresh leaves of *Ocimum gratissimum*, *Vernonia amygdalina*, digital weighing balance, water bottles, beakers, glass slides, 1ml syringe, Orogastric tubes, Graduated Measuring cylinder set, Molten paraffin wax, Rotary microtome, Bouin's fluid, Xylene, Ehrlich's Haematoxylin, Eosin, Spectrophotometer (UV/VIS. Model 752)-long/Tech. China, Centrifuge (Pro-PRP, ROTOFIX 32 Model DG-10-S)-US, Digital pH meter (NANBEL. L/W/H.) China, Digital Electronic weighing balance (RAD WAG. Model RS-R1) India, Oven (REMI. Model RDHO-50), Electric Blender (RICO. Medina, OH 44256), Soxhlet Extractor (PYREX, Model 1792) Jiangsu, China

#### **Methods**

##### **Extraction**

The leaves were destalked from the stem, sorted, washed with distilled water and air dried in a room temperature for 21 days (three weeks) milled to powder using (RICO blender Medina, OH 44256) and stored in an airtight beaker, 140g of the plant sample were extracted on a Soxhlet extractor using ethanol (96%) and water as the extraction solvents. The resulting eluate was dried at 40°C using (REMI Oven Model RDHO-50) and reconstituted in distilled water to appropriate concentrations.

##### **Experimental Design (Add ethical permission) for animal ethics check list provide by author**

This was an experimental study involving sixteen (16) Wistar rats which were purchased from the animal house of the Faculty of Basic Medical Sciences, Delta State University, Abraka weighing between 120-200g were divided at random into four (4) groups of 4 animals each of female Wistar rats thus: Group A – control fed with normal feed and water alongside with normal **salin**. Group B – Fed with Normal feed and water alongside with administration of 1mg of cadmium (14 days to study the direct effect of Cadmium). Group C – Fed with Normal feed and water alongside with administration of 1mg of Extract (14 days to study the direct effect of *Ocimum gratissimum* and *Vernonia amygdalina* extracts). Group D – Fed with Normal feed,

water and 1mg of cadmium along side with 1mg of the extract was administered (14 days to study the ameliorating effects of the extracts)

### **Blood sample and organ collection**

At the end of the treatment period the experimental rats were sacrificed by cervical dislocation. Blood samples were collected by cardiac puncture and dispensed into heparinized bottles for clinical chemistry analysis and EDTA (Ethylenediaminetetra acetic acid) container for haematological tests centrifuged at 5000g for 10 minute. Sera and supernatants collected and were stored frozen until used for analysis.

### **Assessment of Biochemical Parameters**

#### **LIPID PROFILE (It prefer to rewrite in a brief, no need to the details below)**

##### **Determination of Serum Total Cholesterol Using Randox Assay Kit**

###### **Principle.**

Cholesterol was determined after enzymatic hydrolysis and oxidation based on the understanding that quinoneimine (an indicator) is formed when hydrogen peroxide and 4-aminoantipyrine reacts in the presence of phenol and peroxidase.

###### **Procedure:**

Exactly 10 $\mu$ l each of distilled water, sample, and standard (CAL) were pipetted into 'blank', 'Test', and 'Standard' test tubes respectively. Thereafter 1000  $\mu$ l of Reagent was added to all test tubes. They were mixed, incubated for 5 minutes at 37 $^{\circ}$ C and the absorbance of the sample ( $A_{\text{sample}}$ ) and standard ( $A_{\text{standard}}$ ) were read against the reagent blank within 60 minutes at 500nm.

##### **Determination of low density lipoprotein Cholesterol (LDP-C) Using Randox Assay Kit**

###### **Principle:**

Low density lipoproteins (LDL) in the sample precipitate with polyvinyl sulphate. Their concentration is calculated from the difference between the serum total cholesterol and the cholesterol in the super after centrifugation, the cholesterol is spectrophotometrically measured by means of the coupled reactions described below.

###### **Procedure**

0.2  $\mu$ l sample and reagent (A) was pipetted into centrifuge tubes mixed thoroughly and stand for 15minutes at room temperature, was centrifuge at a minimum of 4000 r.p.m. for 15minutes. Supernatant was carefully collected. After which the reagent (cholesterol kit) to room temperature, then pipette into a test tubes

## **Determination of High density lipoprotein Cholesterol (HDP-C) Using Randox Assay Kit**

### **Principle:**

Low density lipoprotein (LDL and VLDL) and chylomicron fractions are precipitated quantitatively by the addition of phosphotungstic acid in the presence of magnesium ions. After centrifugation, the cholesterol concentration in the High density lipoprotein (HDL) fraction, which remains in the supernatant, is determined.

### **Procedure;**

200 µl of sample was Pipetted into centrifuged tubes then 500 µl was used to dilute the precipitant, after which it was mixed thoroughly and allowed for 10 minutes at room temperature. Then it was centrifuged at 4,000 rpm for 10 minutes at 12,000 rpm. Then the clear supernatant was carefully removed using a pipette.

## **Determination of Serum Triglycerides using Manual Method**

### **Principle:**

Triglycerides from serum or plasma are extracted with isopropanol, extracted triglycerides are saponified with an alkali, sodium methylate, forming glycerol and fatty acids.

Triglyceride + Sodium methylate  $\longrightarrow$  Glycerol + fatty acids

Glycerol is oxidized with sodium periodate to produce formaldehyde,

Glycerol + Sodium periodate  $\longrightarrow$  formaldehyde

Formaldehyde is reacted with acetylacetone in the presence of ammonium ions to produce a yellow compound (diacetyldihydrolutidine) which is measured colorimetrically.

### **Procedure:**

Three glass stoppered tubes were set up which contain the test, standard and blank, the tubes were mixed for 30 seconds in a vortex mixer, then the tubes were stand at a room temperature for 10 minutes for clear separation of two layers. There after the top solvent layer was used for further assay.

## **Determination of the red blood percentage using a centrifugation technique.**

**Principle (It prefer to rewrite in a brief, no need to the details below)**

The packed cell volume is that proportion of whole blood occupied by red cells, expressed as a ratio (litre/litre). Anticoagulated blood in a glass capillary of specified length, bore size, and wall thickness is centrifuged in a microhaematocrit centrifuge at RCF 12000-15000xg for 3-5 minutes to obtain constant packing of the red cells. A small amount of plasma remains trapped between the packed red cells. The packed cell volume value is read from the scale of a microhaematocrit reader or calculated by dividing the height of the red cell column by the height of the total column of blood.

**Procedure:**

The capillary tube was filled with sample (blood). After which it was cleaned with a cottonwood. The tip was sealed with plasticine to avoid spillage of blood while spinning. Thereafter Capillary tube was placed on the hematocrit centrifuge, then was allowed to spin for 5min at 12,000rpm. After which the capillary tube was brought out and was placed on the hematocrit reader at the eye level, then the reading was taken.

**Determination of the Red blood cell using (Manual Method)**

**Principle: (It prefer to rewrite in a brief, no need to the details below)**

Whole blood is diluted appropriately using an isotonic diluent to avoid lysis of red cells.

The number of red cells in a known volume and of known dilution is counted using a counting chamber.

**Determination of the White blood cell using (Manual Method)**

**Principle: (It prefer to rewrite in a brief, no need to the details below)**

Whole blood is diluted appropriately using a diluent which haemolyses red cells. Leaving all the nucleated cells intact. The number of white cells in known volume and known dilution are counted using a counting chamber.

**Analysis of Data.**

Data were expressed as mean  $\pm$  standard error (SEM) the mean values between the groups were compared by using analysis of variance (ANOVA) and least significance test (LSD) produced using the software (SPSS). The results were considered significant at  $p < 0.05$  level.

## RESULTS

### Effect of combined leaf extract of *Vernonia amygdalina* and *Ocimum gratissimum* on lipid profile on cadmium induced Wistar Rats

The effect of combined leaf extract of *Vernonia amygdalina* and *Ocimum gratissimum* on lipid profile on cadmium induced wistar rats is presented in Table 1. The effect of *Vernonia amygdalina* and *Ocimum gratissimum* on lipid profile on cadmium induced Wistar Rats showed There was a significant increase in the level of serum cholesterol, triglyceride and LDL levels in group administered with only cadmium when compared with the control group ( $p < 0.05$ ), while those administered with only the plant extract and a combination of the plant extract and cadmium showed significant decrease in level of cholesterol, triglyceride, high density lipoprotein, and low density lipoprotein when compared to the group administered with cadmium only. High density lipoprotein (HDL) level of group administered with cadmium only recorded a significant decrease when compared to those of the control group ( $p < 0.05$ )

The administration of the extract to Cd-exposed rats brought about significant reduction in cholesterol, TAG and LDL compared to rats maintained on Cd alone. (Table 1) below.

**Table 1: Effect of combined leaf extract of *Vernonia amygdalina* and *Ocimum gratissimum* on lipid profile on cadmium induced Wistar Rats**

Groups	Lipid profile			
	Cholesterol	Triglyceride	HDL	LDL
Group 1	25.73±0.70 <sup>a</sup>	23.50±0.96 <sup>a</sup>	3.29±0.17 <sup>a</sup>	19.02±0.34 <sup>a</sup>
Group 2	44.59±3.59 <sup>bc</sup>	28.99±0.98 <sup>bc</sup>	2.15±0.36 <sup>bc</sup>	37.71±2.88 <sup>bc</sup>
Group 3	19.67±0.33 <sup>ade</sup>	17.66±0.34 <sup>ade</sup>	5.43±0.11 <sup>ade</sup>	14.51±0.15 <sup>ade</sup>
Group 4	33.34±0.70 <sup>adf</sup>	25.49±0.68 <sup>adf</sup>	3.08±0.04 <sup>adf</sup>	21.80±0.83 <sup>adf</sup>

Values are Mean±standard Error Mean (SEM). Values with different superscript are statistically different at ( $p < 0.05$ ). superscript (a,b) compares Group 2, Group 3 and Group 4 to Group 1 ((1<sup>st</sup> letters) along the column. Superscript (c,d) compares Group 3 and Group 4 to Group 2 (2<sup>nd</sup> letters) along the column while Superscript (e,f) compares Group 4 to Group 3 along the column.

**Effect of combined leaf extract of *Vernonia amygdalina* and *Ocimum gratissimum* on Hematological Parameters on cadmium induced Wistar Rats**

The effect of combined leaf extract of *Vernonia amygdalina* and *Ocimum gratissimum* on Hematological Parameters on cadmium induced Wistar Rats showed a significantly reduction in the packed cell volume (PCV) and red blood cell (RBC) level of the rats when compared to the control group, while the white blood cell (WBC) level increased ( $p < 0.05$ ). The group administered with the extract and a combination of the extract and cadmium recorded an increase in the packed cell volume (PCV) and red blood cell (RBC) level of the wistar rats and a reduction in the white blood cell (WBC) level when compared to the group administered with cadmium below.

**Table 2: Effect of combined leaf extract of *Vernonia amygdalina* and *Ocimum gratissimum* on lipid profile on cadmium induced Wistar Rats**

Groups	Hematological parameters		
	PCV	RBC	WBC
<b>Group 1</b>	31.33 ± 1.20 <sup>a</sup>	5.00±0.42 <sup>a</sup>	4833.33 ± 88.19 <sup>a</sup>
<b>Group 2</b>	20.67±1.20 <sup>b,c</sup>	3.97±0.09 <sup>b,c</sup>	7766.67 ± 284.80 <sup>b,c</sup>
<b>Group 3</b>	34.00±0.58 <sup>a,d,e</sup>	5.33±0.09 <sup>a,d,e</sup>	4566.67 ± 66.67 <sup>a,d,e</sup>
<b>Group 4</b>	29.33±0.88 <sup>a,d,f</sup>	4.47±0.15 <sup>a,d,f</sup>	5033.33±120.19 <sup>a,d,f</sup>

Values are Mean ± Standard Error Mean (SEM). Values with different superscript are statistically different at ( $p < 0.05$ ). Superscript (a,b) compares Group 2, Group 3, (1<sup>st</sup> letters) along the column. Superscript (c,d) compares Group 3 and Group 4 to Group 2 (2<sup>nd</sup> letters) along the column while superscript (e,f) compares Group 4 to Group 3 along the column.

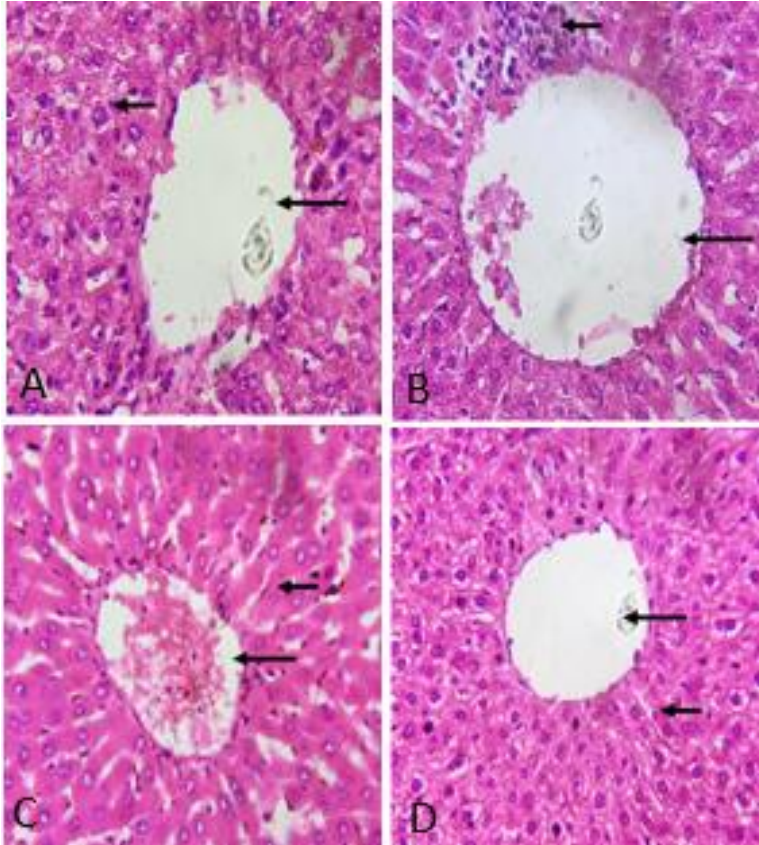


Fig.1 : Histopathological slides

Liver centriole appears normal, large and distinct (long arrow). The hepatocytes also reveal normal nucleus (short arrow).

Liver centriole appears large with thickened wall (long arrow). The hepatocytes also reveal slightly pyknotic nucleus with focal pool of mononuclear infiltrates (short arrow).

Liver centriole appears large and distinct surrounded by mild mononuclear cells (long arrow). The hepatocytes also reveal slightly vacuolated nucleus with mild fatty changes (short arrow).

Liver centriole appears large and distinct surrounded (long arrow). The hepatocytes also reveal pyknotic nucleus (short arrow).

## Discussion

The study clearly shows that the administration of the combined extract of *Ocimum gratissimum* and *Vernonia amygdalina* decreased the cholesterol, triacylglycerol and low density lipoprotein (LDL) level of the experimental animal significantly ( $p < 0.05$ ) when compared to the control while the level of high density lipoprotein (HDL) increased significantly. Whereas, Wistar rat induced with Cadmium and combined plant extract displayed an increased in the lipid profile of the Wistar rat with the exception of the high density lipoprotein (HDL) which showed a significant reduction ( $p < 0.05$ ). Adaramoye *et al.* [18] also reported a decrease in the cholesterol, glyceride, low density lipoprotein (LDL) level of Wistar rat administered with different dose of *Vernonia amygdalina* and an increase in the high density lipoprotein (HDL) level when compared to those of the control, this is in agreement with the findings of this present study.

Lipid disorders play a key role in the pathogenesis and progression of atherosclerosis and cardiovascular disease, according to research, and environmental factors play a role in these disorders as well [19]. The use of medicinal plants for therapeutic purposes are increasingly becoming prevalent in modern society as alternatives to synthetic medicines. Various parts of *Ocimum gratissimum*, including the leaves, stems, and roots, are used pharmaceutically in folklore medicine to treat various ailments and diseases such as diabetes, hypertension, pile, rheumatism, and others [20].

Nonetheless, some unfavorable effects of *Vernonia amygdalina* have been documented, indicating that the saponin fractions of the extract have a hemolytic effect [21]. Wistar rats' body weight was significantly reduced after prolonged feeding with *Vernonia amygdalina* leaves, according to Asuquo *et al.*, [22]. This corroborate with this present study where Wistar rat administered with the combined extract of *Ocimum gratissimum* and *Vernonia amygdalina* displayed a significant decrease in the weight of the rat when compared to those of the control group and groups administered with cadmium. Cholesterol is an important chemical involved in a variety of cellular processes, including membrane fluidity, vitamin D generation on the skin's surface, hormone manufacturing, and even aiding cell connections in the brain<sup>23</sup>. It is critical that the body's cells receive an adequate quantity of cholesterol.

When cholesterol levels in the blood increase, unfortunately, it can have negative implications. Cholesterol, in particular, has gained prominence for its role in the development of **atherosclerosis**, the leading cause of mortality in affluent countries around the world [23]. The therapeutic effects of plant foods have been the focus of many significant dietary research because great efforts have been made to reduce the risk of cardiovascular illnesses through cholesterol management [24, 23].

Reduced erythrocyte survival is one of the adverse implications of this membrane lipid peroxidation [25]. The results obtained from this study, it is apparent that the RBC parameters measured decrease significantly which is an indication of reduced and abnormal erythropoiesis.

This observation is consistent with the reports of Ekweogu *et al.* [26] but differ from the reports of Akpaso *et al.*, [13]. Administration of combined extract of *Ocimum gratissimum* and *Vernonia amygdalina* to cadmium induced rats appreciably improved the levels of red blood cell (RBC) and its indices ( $p < 0.05$ ). This suggests that some phytoconstituents present in the extract can stimulate the formation or secretion of erythropoietin which stimulates the stem cells in the bone marrow to produce RBC [27].

The white blood cell (WBC) acts as a scavenger, removing foreign chemicals from the body. The number of white blood cell (WBC) is known to increase in response to a hazardous environment as a body defense mechanism [28]. Changes in white blood cell (WBC) have been linked to cardiovascular problems [29]. Coronary artery disease, insulin resistance, Type 2 diabetes, stroke, and diabetes-induced macro and microangiopathy have all been linked to leukocytosis [30]. AGEs, oxidative stress, angiotensin II, and pro-inflammatory cytokines have all been shown to activate leucocytes [30]. The result of this study showed a significant ( $P < 0.05$ ) increase in WBC of cadmium induced Wistar rats which became reduced significantly ( $P < 0.05$ ) on combined extract of *Ocimum gratissimum* and *Vernonia amygdalina* treatment. This may have been as a result of the ability of the extract to ameliorate AGEs production and reduce oxidative stress within the blood cells. This finding is in agreement with earlier reports [30].

### **Limitations of Study**

The study was limited to unavailability equipment to conduct a smooth research, it also suffered several challenges such as finance, and time.

### **Conclusion**

The findings of this study demonstrated that *Ocimum gratissimum* and *Vernonia amygdalina* extracts have lipid lowering properties that may be advantageous to patients at risk of cardiovascular disease. The extracts were also found to be efficient in decreasing cholesterol, triacylglyceride, and LDL levels, indicating that they have hypocholesterolaemic properties. The study also shows that the combined plant extract can enhance hematological parameters in Wistar rats who have been exposed to Cadmium. Finally, the study's findings revealed that the combined extracts diet preparation had greater anti lipidemic and hematological properties, suggesting that it could be advantageous to persons at high risk of cardiovascular disease.

### **Recommendation**

Owing to the finding of this study it is recommended that the combined plant extract of *Ocimum gratissimum* and *Vernonia amygdalina* should be explored for its adjuvant therapy for the management of diabetes mellitus treatment.

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