Antidyslipidemic, Anti-Inflammatory, and Antioxidant Activities of Aqueous Leaf Extract of *Dioscoreophyllum cumminsii* (Stapf) Diels in High-Fat Diet-Fed Rats

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*Dioscoreophyllum cumminsii* (Stapf) Diels leaves are widely used in the treatment of diabetes, obesity, and cardiovascular related complications in Nigeria. This study investigates the anti-inflammatory and antiobesity effect of aqueous extract of *Dioscoreophyllum cumminsii* leaves in high-fat diet- (HFD-) induced obese rats. HFD-fed rats were given 100, 200, and 400 mg kg\(^{-1}\) body weight of aqueous extract of *Dioscoreophyllum cumminsii* leaves for 4 weeks starting from 9th week of HFD treatment. *D. cumminsii* leaves aqueous extract reversed HFD-mediated decrease in the activities of superoxide dismutase, catalase, glutathione peroxidase, glutathione reductase, and DNA fragmentation in rats liver was lowered. HFD-mediated alterations in serum total cholesterol, triacylglycerol, high-density lipoprotein cholesterol, low-density lipoprotein cholesterol, and very low-density lipoprotein cholesterol were significantly reversed by the extract. The treatment of HFD-fed rats reduced the levels of insulin, leptin, protein carbonyl, fragmented DNA, and tumour necrosis factor-α and interleukin- (IL-) 6 and IL- 8 and increased the adiponectin level. This study showed that aqueous extract of *Dioscoreophyllum cumminsii* leaves has potential antiobesity and anti-inflammatory effects through modulation of obesity-induced inflammation, oxidative stress, and obesity-related disorder in HFD-induced obese rats.

**1. Introduction**

Obesity, excessive visceral accumulation and distribution, is a risk factor in atherosclerosis, cancer, diabetes (type 2), dyslipidemia, and metabolic syndrome [1]. Indeed, its rising prevalence which continues to be a global challenge is associated with high-fat, caloric-dense diets, sedentary life styles, increased urbanization, and psychosocial stress (McLaren, 2007). Adiposity in high-fat consumption has been demonstrated in epidemiological and experimental studies [2, 3] and is associated with adipose tissue inflammation, endoplasmic reticulum stress of adipocyte, and necrosis-like cell death [4]. World Health Organization (WHO) estimated that there are more than 1.1 billion adults overweight in the world and about 115 million individuals suffering from obesity-related problems in low-income and middle-income populations [5]. Drugs including phentermine, fluoxetine, orlistat, sibutramine, and rimonabant are used for the treatment of obesity. However, associated side effects are not limited to nausea, dizziness, insomnia, diarrhea, dyspepsia, and constipation. Thus, there are growing demands for plant derived foods and compounds to manage and treat ailments such as metabolic syndrome, obesity, and diabetes [6]. Indeed, antidiabetic activity and capability of *Dioscoreophyllum cumminsii* leaves have been validated [7, 8]. In furtherance of these, the therapeutic importance of *Dioscoreophyllum cumminsii* leaves in obesity and associated complications was investigated.
Dioscoreophyllum cumminsii, a tropical rainforest vine, family Menispermaceae, is known as serendipity berry, (Omu-aja) Youba, and Okazi (Igbo) [9]. It is widely distributed in Guinea-Bissau, Sierra Leone, Liberia, Nigeria, Benin, and Congo. Monellin, a content of the fruit, is 3000 times sweeter than sugar [10]. Alkaloids, anthraquinones, cardiac glycosides, flavonoids, phlobatannins, saponins, and tannins are reported phytochemicals in Dioscoreophyllum cumminsii leaves [11]. Leaves of this plant are used in the treatment of diarrhea, dysentery, and uterine haemorrhages [11]. Recently, we reported in different models that magnoflorine, jatrohrrhizine, and columbamine are responsible for the antidiabetic and protective importance in metabolic syndrome model [7, 8]. As diabetes and metabolic syndrome could result from obesity, we evaluated the effect of aqueous leaf extract of Dioscoreophyllum cumminsii on HFD-induced dyslipidemia, inflammation, and oxidative stress.

2. Materials and Methods

2.1. Experimental Animals. Thirty-five male albino rats (Rattus norvegicus) of Wistar strain (141.24 ± 0.32 g) were obtained from the Animal House of Veterinary Physiology, Biochemistry and Pharmacology, University of Ibadan, Nigeria. Rats, kept in clean plastic cages, were placed in well-ventilated house conditions and supplied with food (Capefeed Ltd., Osogbo, Nigeria) and water ad libitum.

2.2. Plant Material and Authentication. Dioscoreophyllum cumminsii leaves were collected from Oja Titun, Ilorin, Nigeria. They were authenticated and deposited in the herbarium of Department of Plant Biology, University of Ilorin, Ilorin, Nigeria (UIH 001/1082).

2.3. Chemical Reagents and Assay Kits. Disodium salt, hexahydrate and guanidine hydrochloride, ultrapure water, 5,5-dithiobis-2-nitrobenzoic acid (DTNB), and trichloroacetic acid were purchased from Research Organics, 4353 East 49th Street, Cleveland, Ohio 44125; superoxide dismutase, glutathione peroxidase, glutathione reductase, glucose-6-phosphate dehydrogenase, catalase, total cholesterol, triglyceride, and HDL-cholesterol assay kits were purchased from Randox Laboratories Co., Antrim, UK. Adiponectin, insulin, and leptin (enzyme immunoassay kits) were products of Sigma-Aldrich Inc., St. Louis, USA. All other reagents used were products of Sigma-Aldrich Inc., St. Louis, USA.

2.4. Preparation of Plant Extract. Dioscoreophyllum cumminsii leaves were washed clean with distilled water, air-dried, and pulverized using domestic blender. Pulverized leaves (200 g) were extracted in distilled water (1 L) for 48 h, filtered, and concentrated on water bath. The extract yield (26.60 g) was reconstituted to 100, 200, and 400 mg/kg body weight doses. We reported magnoflorine (1.97 mg/g), jatrohrrhizine (1.35 mg/g), and columbamine (2.12 mg/g) as the antidiabetic and antidyldiabemic agents in aqueous leaf extract of Dioscoreophyllum cumminsii [7]. The extract was refrigerated all through the experimental period to avoid microbial contamination and maintain its composition.

2.5. Feed Composition and Formulation. HFD with composition presented in Table 1 was used for the study and formulated as described by Ajiboye et al. [12].

2.6. Animal Grouping and Treatments. Rats (35) were randomized into seven groups (A–G) of five rats each. All rats received HFD for 12 weeks except rats in groups A and C fed with control diet. In addition, rats in groups C–F were gavaged with 400, 100, 200, and 400 mg/kg BW of aqueous extract of D. cumminsii, respectively, for 4 weeks starting from 9th week of diet treatments. Group A rats, which served as control, were gavaged with distilled water (1 mL), while group G rats received 400 mg/kg BW metformin [8], reference drug, for 4 weeks starting from 9th week. This study was approved by Al-Hikmah University Ethical Committee on the use of laboratory animals (HUI/ECULA/014/009) and all treatments were done in accordance with the Guidelines of National Research Council's Guide for the Care and Use of Laboratory Animals [13].

2.7. Preparation of Serum and Tissue Homogenate. Rats were anaesthetized with diethyl ether and sacrificed 24 h after the last day of the experimental period. Blood collected from the jugular vein was allowed to clot for 15 min and centrifuged for 5 min at 500 g for serum collection. Liver was excised and homogenized in sucrose-Tris buffer (0.25 mol/L sucrose, 10 mmol/L Tris-HCl, pH 7.4).

2.8. Biochemical Assays

2.8.1. Blood Insulin, Adipokines, and Cytokines. Adiponectin, insulin, leptin, tumour necrosis factor-α, interleukin-6, and interleukin-8 were determined as described in manufacturer’s assay kit manual.

### Table I: Feed composition and formulation.

| Feed component | Control diet (g/kg) | High-fat diet (g/kg) |
|----------------|---------------------|---------------------|
| Corn starch    | 506                 | 396                 |
| Casilan 90*    | 250                 | 250                 |
| Lard           | 40                  | 140                 |
| Cholesterol    | 0                   | 10                  |
| Sucrose        | 100                 | 100                 |
| Rice husk      | 40                  | 40                  |
| DL-methionine  | 4                   | 4                   |
| Lysine         | 10                  | 10                  |
| Vitamin mix**  | 10                  | 10                  |
| Mineral mix*** | 40                  | 40                  |

*Casilan 90 (g100 g−1), energy (1572 kg 100 g−1), protein (90 g), carbohydrate (0.3 g), fat (1.0 g), fibre (trace), sodium (0.03 mg), and calcium (1400 mg). **Vitamin mix (per kg of diet): thiamine hydrochloride (6 mg), pyridoxine hydrochloride (7 mg), nicotinic acid (30 mg), calcium pantothenate (16 mg), folic acid (2 mg), biotin (0.2 mg), Cyanocobalamin (0.01 mg), retinol palmitate (4000 IU), cholecalciferol (100 IU), a-tocopherol acetate (50 IU), menadione (0.05 mg), and chole chloride (2 g). ***Mineral mix (g/kg−1): CoCl2·6H2O (0.001), CuSO4·5H2O (0.079), MnSO4·7H2O (0.178), KI (0.033), NaCl (3.573), ZnSO4·7H2O (1.60), CaSO4 (11.61), MgSO4·7H2O (2.292), K2HPO4 (10.559), and FeSO4·7H2O (1.075).
Catalase absorbance/min at 480 nm. The rate of autoxidation of epinephrine as determined by change in enzyme activity was defined as 50% inhibition of the rate monitored at 480 nm every 30 s for 150 s. A unit of prepared 0.3 mM epinephrine (0.3 mL). Increase in absorbance 2.5 mL carbonate buffer (0.05 M, pH 10.2), and freshly prepared liver homogenate (0.2 mL), rats was determined as described by Misra and Fridovich [17].

2.8.3. Antioxidant Enzymes and Oxidative Stress Biomarkers

Superoxide Dismutase. Superoxide dismutase in the liver of rats was determined as described by Misra and Fridovich [17]. The assay mixture consisted of liver homogenate (0.2 mL), 2.5 mL carbonate buffer (0.05 M, pH 10.2), and freshly prepared 0.3 mM epinephrine (0.3 mL). Increase in absorbance was monitored at 480 nm every 30 s for 150 s. A unit of enzyme activity was defined as 50% inhibition of the rate of autoxidation of epinephrine as determined by change in absorbance/min at 480 nm.

Catalase. Catalase activity was determined as described by Beers and Sizer [18]. The assay mixture consisted of 2 mL phosphate buffer and 30 mM H₂O₂ and liver homogenate (50 μL). Absorbance was read at 240 nm for 1 min and the activity was calculated using the extinction coefficient of H₂O₂ (43.6 M cm⁻¹).

Glutathione Peroxidase and Glutathione Reductase. Activities of glutathione peroxidase and glutathione reductase were determined as described in commercial kits (Randox Laboratories Ltd., Antrim, UK).

Reduced Glutathione (GSH). Glutathione content of liver was determined as described by Ellman [19]. Briefly, liver homogenate (1.0 mL) was mixed with 0.1 mL of 25% trichloroacetic acid (TCA). The mixture was centrifuged at 5,000 × g for 10 min to remove precipitate. Supernatant (0.1 mL) was mixed with 2 mL of 0.6 mM DTNB prepared in 0.2 M sodium phosphate buffer pH (8.0). Absorbance was read at 412 nm.

2.8.2. Lipid Profile. Serum TC, TAG, and HDLc were determined as described in commercial kits (Randox Laboratories Ltd., Antrim, UK). LDLc and VLDLc were calculated using the following expression:

\[
\text{LDLc} = 0.2 \times \text{TAG} \\
\text{LDLc} = \text{TC} - (\text{HDLc} + \text{VLDLc}).
\]  

Cardiac index (CI), atherogenic index, and coronary artery index were estimated as described by Kang et al. [14], Kayamori and Igarashi [15], and Ajiboye et al. [16], respectively.

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2.9. Statistical Analysis. All the data were expressed as the mean ± SEM of five replicates unless stated otherwise. Analysis of variance (ANOVA) followed by Tukey-Kramer test for difference between means was used to detect any significant difference between the treatment groups in this study. Statistical evaluation of data was performed with SPSS version 20.0. Differences were considered statistically significant at \( p < 0.05 \).
Parameters are expressed as mg/dL. Values with different alphabetical superscript are significantly different (p < 0.05). HDL cholesterol: high-density lipoprotein cholesterol; LDL cholesterol: low-density lipoprotein cholesterol; HFD: high-fat diet. Concentrations of lipid profile parameters are expressed as mg/dL. Values with different alphabetical superscript are significantly different (p < 0.05).

HFD-mediated alterations in these parameters. Indeed, the highest dose (400 mg/kg body weight) produced 85.11, 64.04, 51.60, 72.92, and 60.88% reversal of TC, TAG, VLDLc, LDLc, and HDLc, respectively, and compared significantly with reference drug, metformin (Table 3).

3.3. Antioxidant Enzymes. Activities of antioxidant enzymes, superoxide dismutase, catalase, glutathione peroxidase, glutathione reductase, and glucose 6-phosphate dehydrogenase, in the liver of HFD-fed rats decreased significantly (p < 0.05) when compared to control rats. This decrease was significantly reversed by aqueous leaf extract of Dioscoreophyllum cumminsii in dose dependent manner (Table 4), which compared significantly with metformin treated rats.

3.4. Oxidative Stress Biomarkers. Lipid peroxidation products, conjugated dienes, lipid hydroperoxides, and malondialdehyde, in the liver of HFD-fed rats increased significantly

| Group                          | Total cholesterol | Triglyceride | VLDL cholesterol | LDL cholesterol | HDL cholesterol |
|--------------------------------|-------------------|--------------|------------------|-----------------|-----------------|
| Control                        | 47.21 ± 1.28      | 52.33 ± 1.14 | 10.04 ± 0.69     | 3.63 ± 0.09     | 33.77 ± 1.26    |
| HFD                            | 66.21 ± 1.65      | 80.07 ± 1.86 | 15.68 ± 1.67     | 33.21 ± 0.12    | 15.06 ± 1.33    |
| 400 mg/kg body weight of extract| 43.11 ± 2.01      | 51.12 ± 2.70 | 9.76 ± 0.11      | 2.12 ± 0.24     | 34.29 ± 0.98    |
| HFD + 100 mg/kg body weight extract| 63.24 ± 1.46   | 76.34 ± 4.11 | 15.37 ± 0.43     | 30.04 ± 1.02    | 16.78 ± 1.54    |
| HFD + 200 mg/kg body weight extract| 58.67 ± 0.76    | 71.35 ± 3.19 | 14.99 ± 1.04     | 18.54 ± 0.92    | 23.11 ± 1.29    |
| HFD + 400 mg/kg body weight extract| 50.04 ± 0.48   | 62.35 ± 2.30 | 13.77 ± 0.87     | 11.64 ± 1.03    | 26.45 ± 0.89    |
| HFD + 400 mg/kg body weight metformin| 55.21 ± 0.92  | 64.76 ± 2.05 | 13.92 ± 0.92     | 17.65 ± 1.00    | 24.68 ± 0.57    |

Values are mean ± SEM of five determinations and are considered statistically significant at p < 0.05. VLDL cholesterol: very-low density lipoprotein cholesterol; HDL cholesterol: high-density lipoprotein cholesterol; LDL cholesterol: low-density lipoprotein cholesterol; HFD: high-fat diet. Concentrations of lipid profile parameters are expressed as mg/dL. Values with different alphabetical superscript are significantly different (p < 0.05).
**Table 4:** Specific activities of antioxidant enzymes in the liver of HFD-fed rats following the administration of aqueous extract of *D. cumminsii* leaves to high-fat diet-fed rats.

| Group | SOD (U/mg protein) | Catalase (U/mg protein) | GSH-Px (U/mg protein) | GSH-Red (U/mg protein) | Glucose-6-PD (U/mg protein) |
|-------|--------------------|-------------------------|-----------------------|------------------------|-----------------------------|
| Control | 44.03 ± 2.01<sup>d</sup> | 32.03 ± 0.16<sup>e</sup> | 78.21 ± 0.97<sup>d</sup> | 33.33 ± 1.41<sup>d</sup> | 23.71 ± 2.01<sup>d</sup> |
| HFD | 27.54 ± 3.12<sup>a</sup> | 14.21 ± 0.32<sup>a</sup> | 26.26 ± 1.34<sup>e</sup> | 13.18 ± 1.56<sup>e</sup> | 9.75 ± 1.98<sup>e</sup> |
| 400 mg/kg body weight of extract | 48.76 ± 1.11<sup>f</sup> | 38.23 ± 1.23<sup>f</sup> | 83.51 ± 1.26<sup>d</sup> | 34.10 ± 1.23<sup>d</sup> | 24.21 ± 1.65<sup>d</sup> |
| HFD + 100 mg/kg body weight extract | 33.24 ± 1.78<sup>b</sup> | 17.65 ± 2.19<sup>b</sup> | 42.08 ± 2.10<sup>b</sup> | 13.27 ± 1.34<sup>e</sup> | 13.45 ± 1.89<sup>b</sup> |
| HFD + 200 mg/kg body weight extract | 37.65 ± 0.99<sup>c</sup> | 23.15 ± 0.75<sup>c</sup> | 53.15 ± 2.31<sup>c</sup> | 17.37 ± 1.39<sup>c</sup> | 18.76 ± 3.41<sup>c</sup> |
| HFD + 400 mg/kg body weight extract | 40.19 ± 2.03<sup>c</sup> | 27.24 ± 1.14<sup>d</sup> | 67.22 ± 1.01<sup>d</sup> | 21.23 ± 0.64<sup>c</sup> | 7.02 ± 0.26<sup>c</sup> |
| HFD + 400 mg/kg body weight metformin | 41.07 ± 1.02<sup>c</sup> | 24.01 ± 1.23<sup>c</sup> | 72.92 ± 2.22<sup>c</sup> | 20.05 ± 0.68<sup>c</sup> | 8.22 ± 0.93<sup>c</sup> |

Values are mean ± SEM of five determinations and are considered statistically significant at *p* < 0.05. SOD: superoxide dismutase; GSH-Px: glutathione peroxidase; GSH-Red: glutathione reductase; Glc 6-PD: glucose 6-phosphate dehydrogenase; HFD: high-fat diet. Enzyme activities are expressed as nmol/min/mg protein. Values with different alphabetical superscript are significantly different (*p* < 0.05).

**Table 5:** Levels of malondialdehyde, conjugated dienes, and lipid hydroperoxides in the liver of HFD-fed rats following the administration of aqueous extract of *D. cumminsii* leaves.

| Group | Malondialdehyde (nmol/mg protein) | Conjugated dienes (nmol/mg protein) | Lipid hydroperoxides (nmol/mg protein) |
|-------|----------------------------------|-------------------------------------|--------------------------------------|
| Control | 3.84 ± 0.37<sup>e</sup> | 23.89 ± 2.33<sup>b</sup> | 18.07 ± 0.62<sup>b</sup> |
| HFD | 27.21 ± 0.45<sup>d</sup> | 75.21 ± 3.07<sup>e</sup> | 64.62 ± 3.27<sup>e</sup> |
| 400 mg/kg body weight of extract | 3.62 ± 0.54<sup>e</sup> | 18.05 ± 0.29<sup>n</sup> | 10.12 ± 1.24<sup>e</sup> |
| HFD + 100 mg/kg body weight extract | 16.82 ± 1.79<sup>c</sup> | 38.24 ± 0.98<sup>c</sup> | 38.76 ± 0.43<sup>d</sup> |
| HFD + 200 mg/kg body weight extract | 12.05 ± 1.06<sup>c</sup> | 27.01 ± 1.34<sup>c</sup> | 33.21 ± 2.22<sup>c</sup> |
| HFD + 400 mg/kg body weight extract | 7.45 ± 1.92<sup>c</sup> | 27.23 ± 1.65<sup>c</sup> | 25.32 ± 1.86<sup>c</sup> |
| HFD + 400 mg/kg body weight metformin | 7.53 ± 1.63<sup>c</sup> | 30.61 ± 1.53<sup>d</sup> | 28.22 ± 0.37<sup>d</sup> |

Values are mean ± SEM of ten determinations and are considered statistically significant at *p* < 0.05. HFD: high-fat diet. Conjugated dienes, lipid hydroperoxides, malondialdehyde, and protein carbonyl are expressed as nmol/mg protein. Values with different alphabetical superscript are significantly different (*p* < 0.05).

by 608.59, 214.82, and 257.61% when compared to control rats. Administration of aqueous leaf extract of *D. cumminsii* significantly lowered HFD-mediated increase in levels of conjugated dienes, malondialdehyde, and lipid hydroperoxides when compared to the control rats (Table 5). Similar reduction was observed for HFD-fed rats treated with metformin. Also, protein carbonyl, product of protein oxidation, and fragmented DNA of HFD-fed rats were lowered by extract administration (Figures 4 and 5).

**4. Discussion**

Demands for health promoting/maintenance foods have led to increase in investigations into the bioactive constituents (phenolic acids, polyphenols, and micro- and macronutrients) conferring the medicinal properties [6]. Although studies have documented the usefulness of *D. cumminsii* leaves in the management of diabetes and high-fructose-induced metabolic syndrome, no study has evaluated the effect on HFD-induced obesity. This study thus presents the effect of aqueous leaf extract of *Dioscoreophyllum cumminsii* (Stapf) Diels in HFD-fed rats.

Leptin, adiponectin, and insulin are indicators of body mass fats and energy imbalance and are present in obesity [23, 24]. The increase in serum leptin and insulin of HFD-fed rats is in consonance with previous studies [25–27]. Reversal of HFD-mediated increase in leptin and insulin by the extract suggests inhibition of lipogenesis and stimulation of lipolysis and reduction of intracellular lipid levels in skeletal
muscle, liver, and pancreatic β-cells, leading to improved insulin sensitivity [24] and decreased lipid accumulation in adipocytes [28].

Previous studies have demonstrated decrease in adiponectin level in HFD-fed rats and have implicated its involvement in diseases presenting obesity [24]. This could be associated with insulin resistance and hyperinsulinemia [29]. The reversal of HFD-mediated decrease in adiponectin by aqueous leaf extract of Dioscoreophyllum cumminsii could have resulted from improved insulin sensitivity, as evident in this study, leading to decreased flow of free fatty acids and stimulating glucose utilization and fatty acid oxidation [30]. In addition, this may protect cardiovascular system and reduce incidence of myocardial infarction [29].

Elevated levels of TC, TG, VLDLc, and LDLc with concomitant reduction in HDLc characterize the dyslipidemic changes reported for HFD [12, 31]. Indeed, TC, TG, VLDLc, LDLc, and HDLc indicate disordered lipid metabolism and predisposition to cardiovascular disease [12, 31, 32]. These alterations could predispose the risk of developing atherosclerosis and cardiovascular diseases [33], while reduction in HDL cholesterol could intensify the development of atherosclerosis and cardiovascular diseases. Indeed, studies have demonstrated the importance of aqueous leaf extract of Dioscoreophyllum cumminsii in the regulation of dyslipidemia in diabetic and metabolic HFD-fed rats [7, 8]. Thus, the reversal of HFD-fed rats mediated alterations in lipid profile by aqueous leaf extract of D. cumminsii suggests antidyslipidemic activity of the extract.

Oxidative stress associated with consumption of HFD results from overwhelmed antioxidant enzymes, which act in concerted manner to detoxify reactive oxygen species [34]. The decreased antioxidant enzymes observed in this study have been documented in HFD-fed rats [12, 32, 35, 36]. Reversal of HFD-mediated decrease in these enzymes suggests antioxidant activity of the extract, although, in a different animal model, antioxidant activities of D. cumminsii have been reported [7, 8].

Lipid peroxidation, protein oxidation, and DNA fragmentation are consequential effects of overwhelmed antioxidant defense system. Elevated levels of lipid peroxidation products, CD, LH, and MDA, in this study are in accordance with previous studies [12, 32, 37, 38]. This may lead to disorganization and functional loss of membrane [39]. Similar increased protein carbonyl and fragmented DNA, associated with HFD consumption [40–43], indicate oxidative stressed rats. The capability of D. cumminsii to reverse the increase in oxidative stress biomarkers further provided the antioxidant capability of the extract.

5. Conclusion

Arising from the data obtained from this study, it is evident from the reversal of HFD-mediated alterations in proinflammatory cytokines, metabolic hormones, and antioxidant enzymes that aqueous leaf extract of Dioscoreophyllum cumminsii leaves possesses antioxidants, antidyslipidemic, and anti-inflammatory properties.

Conflicts of Interest

The authors declare no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

References

[1] X.-M. T. Nguyen, J. Lane, B. R. Smith, and N. T. Nguyen, "Changes in inflammatory biomarkers across weight classes in a representative US population: A link between obesity and inflammation," Journal of Gastrointestinal Surgery, vol. 13, no. 7, pp. 1205–1212, 2009.
[2] W. P. T. James, "The epidemiology of obesity: the size of the problem," Journal of Internal Medicine, vol. 263, no. 4, pp. 336–352, 2008.
[3] M. Korbonits, Obesity and Metabolism, KARGER, Basel, 2008.
[4] S. Cinti, G. Mitchell, G. Barbatelli et al., "Adipocyte death defines macrophage localization and function in adipose tissue of obese mice and humans," Journal of Lipid Research, vol. 46, no. 11, pp. 2347–2355, 2005.
[5] World Health Organization, Controlling the Global Obesity Epidemic, vol. 7, World Health Organization, 2015.
[6] H. O. B. Oloyede, T. O. Ajiboye, Y. O. Komolafe, and A. K. Salau, "Polyphenolic extract of Blighia sapida arilli prevents N-nitrosodiethylamine-mediated oxidative onslaught on microsomal protein, lipid and DNA," Food Bioscience, vol. 1, pp. 48–56, 2013.
[7] H. O. B. Oloyede, T. O. Bello, T. O. Ajiboye, and M. O. Salauw, "Antidiabetic and antidiyslipidemic activities of aqueous leaf extract of Dioscoreophyllum cumminsii (Stapf) Diels in alloxa-indueiated diabetic rats," Journal of Ethnopharmacology, vol. 166, pp. 313–322, 2015.
[8] T. O. Ajiboye, H. Aliyu, M. A. Tanimu, R. M. Muhammad, and O. B. Ibitoye, "Dioscoreophyllum cumminsii (Stapf) Diels leaves halt high-fructose induced metabolic syndrome: Hyperglycemia, insulin resistance, inflammation and oxidative stress," Journal of Ethnopharmacology, vol. 192, pp. 471–479, 2016.
[9] J. A. Morris and R. H. Cagan, “Purification of monellin, the sweet principle of Dioscoreophyllum cuminensis,” BBA - General Subjects, vol. 261, no. 1, pp. 114–122, 1972.

[10] G. E. Inglett and J. F. May, “Serendipity berries—source of a new intense sweetener,” Journal of Food Science, vol. 34, no. 5, pp. 408–411, 1969.

[11] B. Oliver-Bever, Medicinal Plants in Tropical West Africa, Cambridge University Press, 1986.

[12] T. O. Ajiboye, S. A. Akinpelu, H. F. Muritela et al., “Trichosanthes cucumerina fruit extenuates dyslipidemia, protein oxidation, lipid peroxidation and DNA fragmentation in the liver of high-fat diet-fed rats,” Journal of Food Biochemistry, vol. 38, no. 5, pp. 480–490, 2014.

[13] Committee, Guide for the Care and Use of Laboratory Animals, 8th edition, 2011.

[14] M. J. Kang, E. K. Lee, and S. S. Lee, “Effects of two P/S ratios with same peroxidizability index value and antioxidants supplementation on serum lipid concentration and hepatic enzyme activities of rats,” Clinica Chimica Acta, vol. 350, no. 1-2, pp. 79–87, 2004.

[15] F. Kayamori and K. Igarashi, “Effects of dietary nasunin on the serum cholesterol level in rats,” Bioscience, Biotechnology, and Biochemistry, vol. 58, no. 3, pp. 570–571, 1994.

[16] T. O. Ajiboye, A. A. Hussaini, B. Y. Nafiu, and O. B. Ibiboie, “Aqueous seed extract of Hunteria umbellata (K. Schum.) Hallier f. (Apocynaceae) palliates hyperglycemia, insulin resistance, dyslipidemia, inflammation and oxidative stress in high-fructose diet-induced metabolic syndrome in rats,” Journal of Ethnopharmacology, vol. 198, pp. 184–193, 2017.

[17] H. P. Misra and I. Fridovich, “The role of superoxide anion in the autoxidation of epinephrine and a simple assay for superoxide dismutase,” The Journal of Biological Chemistry, vol. 247, no. 10, pp. 3170–3175, 1972.

[18] R. F. Beers and I. W. Sizer, “A spectrophotometric method for measuring the breakdown of hydrogen peroxide by catalase,” The Journal of Biological Chemistry, vol. 195, pp. 133–140, 1952.

[19] G. L. Ellman, “Tissue sulfhydryl groups,” Archives of Biochemistry and Biophysics, vol. 82, no. 1, pp. 70–77, 1959.

[20] Reilly C. A. and Aust S. D., “Measurement of lipid peroxidation,” in Current Protocols in Toxicology, chapter 2, unit 2.4, 2001.

[21] R. L. Levine, D. Garland, C. N. Oliver et al., Oxygen Radicals in Biological Systems Part B: Oxygen Radicals and Antioxidants, Elsevier, 1990.

[22] Burton, K. “A study of the conditions and mechanism of the diphenylamine reaction for the colorimetric estimation of deoxyribonucleic acid,” Biochemical Journal, vol. 62, no. 2, pp. 315–323, 1956.

[23] U. Meier and A. M. Gressner, “Endocrine regulation of energy metabolism: review of pathobiochemical and clinical chemical aspects of leptin, ghrelin, adiponectin, and resistin,” Clinical Chemistry, vol. 50, no. 9, pp. 1511–1523, 2004.

[24] A. Fernández-Sánchez, E. Madrigal-Santillán, M. Bautista et al., “Inflammation, oxidative stress, and obesity,” International Journal of Molecular Sciences, vol. 12, no. 5, pp. 3117–3132, 2011.

[25] Y. Matsu, U. Tomaru, A. Miyoshi et al., “Overexpression of TNF-α converting enzyme promotes adipose tissue inflammation and fibrosis induced by high fat diet,” Experimental and Molecular Pathology, vol. 97, no. 3, pp. 354–358, 2014.

[26] A. Caimari, F. Puiggròs, M. Suárez et al., “The intake of a hazelnut skin extract improves the plasma lipid profile and reduces the lathiocholic/deoxycholic bile acid faecal ratio, a risk factor for colon cancer, in hamsters fed a high-fat diet,” Food Chemistry, vol. 167, pp. 138–144, 2015.

[27] P. J. Scarpace and Y. Zhang, “Leptin resistance: a predisposing factor for diet-induced obesity,” American Journal of Physiology-Regulatory, Integrative and Comparative Physiology, vol. 296, no. 3, pp. R493–R500, 2009.

[28] P. Brahmanaidu, H. Nemani, B. Mergia, S. K. Mehra, S. Potana, and S. Ramgopalrao, “Mitigating efficacy of piperine in the physiological derangements of high fat diet induced obesity in Sprague Dawley rats,” Chemico-Biological Interactions, vol. 221, pp. 42–51, 2014.

[29] A. Psilonapaniagoti, H. Papadaki, E. F. Kranjioti, T. K. Axandrides, and J. N. Varakis, “Expression of adiponectin and adiponectin receptors in human pituitary gland and brain,” Neuroendocrinology, vol. 89, no. 1, pp. 38–47, 2009.

[30] G. Lastra, C. M. Manrique, and M. R. Hayden, “The role of beta-cell dysfunction in the cardiometabolic syndrome,” Journal of the CardioMetabolic Syndrome, vol. 1, no. 1, pp. 41–46, 2006.

[31] Oloyede H. O. B., Ajiboye T. O., and Salau A. K., “Potential roles of garlic and ginger in the management of metabolic syndrome,” Transaction of the Nigerian Society of Biochemistry and Molecular Biology, vol. 1, pp. 1–8, 2015.

[32] T. O. Ajiboye, G. A. Iliasu, A. O. Adeleye et al., “A fermented sorghum/millet-based beverage, Obiolor, extenuates high-fat diet-induced dyslipidemia and redox imbalance in the livers of rats,” Journal of the Science of Food and Agriculture, vol. 96, no. 3, pp. 791–797, 2016.

[33] Q. Chen, S. E. Reis, C. Kammerer et al., “Association of anti-oxidized LDLI and candidate genes with severity of coronary stenosis in the Women’s Ischemia Syndrome Evaluation study,” Journal of Lipid Research, vol. 52, no. 4, pp. 801–807, 2011.

[34] S. Muthulakshmi and R. Saravanan, “Protective effects of azealic acid against high-fat diet-induced oxidative stress in liver, kidney and heart of C57BL/6J mice,” Molecular and Cellular Biochemistry, vol. 377, no. 1-2, pp. 23–33, 2013.

[35] R.-L. Yang, W. Li, Y.-H. Shi, and G.-W. Le, “Lipoic acid prevents high-fat diet-induced dyslipidemia and oxidative stress: A microarray analysis,” Nutrition Journal, vol. 24, no. 6, pp. 582–588, 2008.

[36] R. D. Malheiros, V. M. B. Moraes, A. Collin, G. P. J. Janssens, E. Decuyper, and J. Buysse, “Dietary macronutrients, endocrine functioning and intermediary metabolism in broiler chickens: Pair wise substitutions between protein, fat and carbohydrate,” Nutrition Research, vol. 23, no. 4, pp. 567–578, 2003.

[37] M. Ming, L. Guanhua, Y. Zhanhai, C. Guang, and Z. Xuan, “Effect of the Lycium barbarum polysaccharides administration on blood lipid metabolism and oxidative stress of mice fed high-fat diet in vivo,” Food Chemistry, vol. 113, no. 4, pp. 872–877, 2009.

[38] J. Ling, B. Wei, G. Lv, H. Li, and S. Li, “Anti-hyperlipidaemic and antioxidiant effects of turmeric oil in hyperlipidemic rats,” Food Chemistry, vol. 130, no. 2, pp. 229–235, 2012.

[39] E. Niki, “Lipid peroxidation: physiological levels and dual biological effects,” Free Radical Biology & Medicine, vol. 47, no. 5, pp. 469–484, 2009.

[40] N. Matsuazawa-Nagata, T. Takamura, H. Ando et al., “Increased oxidative stress precedes the onset of high-fat diet-induced insulin resistance and obesity,” Metabolism - Clinical and Experimental, vol. 57, no. 8, pp. 1071–1077, 2008.

[41] S. A. Noeman, H. E. Hamooda, and A. A. Baalash, “Biochemical study of oxidative stress markers in the liver, kidney and heart.”
of high fat diet induced obesity in rats,” *Diabetology & Metabolic Syndrome*, vol. 3, article 17, 2011.

[42] L. V. Yuzefovych, S. P. Ledoux, G. L. Wilson, and L. I. Rachek, "Mitochondrial DNA damage via augmented oxidative stress regulates endoplasmic reticulum stress and autophagy: crosstalk, links and signaling," *PLoS ONE*, vol. 8, no. 12, Article ID e83349, 2013.

[43] C. Bonnard, A. Durand, S. Peyrol et al., "Mitochondrial dysfunction results from oxidative stress in the skeletal muscle of diet-induced insulin-resistant mice," *The Journal of Clinical Investigation*, vol. 118, no. 2, pp. 789–800, 2008.