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5 ABSTRACT

6 **Aims:** To investigate the effect of food blends (plantain, soybean and ginger) on the blood  
7 glucose, lipid profile and haematological indices on *streptozotocin* induced diabetic rats

8 **Methodology:** A total of 35 rats of mean body weight 219.07g separated into 7 groups (5 per  
9 group) where induced by a single intraperitoneal (I.P) injection of *streptozotocin* (0.1g dissolved  
10 in 5ml of freshly prepared sodium citrate buffer 0.1M, pH 4.5) at a dose of 40 mg/kg body  
11 weight after fasting for 12 hours and fed with flours/blends. The flours were produced from plant  
12 materials for different treatments/blends (blend A=100% unripe plantain, B=80% unripe plantain,  
13 14% soybean, 6% ginger, C=70% unripe plantain, 26% soybean, 4% ginger, D= 60% unripe  
14 plantain, 38% soybean, 2% ginger, E= 50% unripe plantain, 50% soybean) and the  
15 phytochemicals and minerals content were determined. Blood glucose was determined at 5 days  
16 interval for 25 days. Diabetes was confirmed in rats with blood glucose concentrations  $\geq 200$   
17 mg/dl. After 25 days rats were anaesthetized with chloroform vapour and blood samples  
18 collected by cardiac puncture for haematology and lipid profile determination.

19 **Results:** The results showed that unripe plantain, soya beans and ginger in adequate  
20 proportion (C=70% unripe plantain, 26% soybean, 4% ginger or D= 60% unripe plantain, 38%  
21 soybean, 2% ginger) could help to reduce blood glucose, improve haematological parameters and  
22 lipid profile. Significant reduction was observed in the blood glucose level of rats fed blends C  
23 and D from 286 to 85mg/dl and 307 to 90mg/dl respectively at the end of experiment. These  
24 results also demonstrated that the inclusion of ginger at 6% causes rise in blood glucose  
25 level. Total cholesterol (TC) increased in all the blends. However, the lowest concentration of TC  
26 was observed in blends C and D. The highest packed cell volume (60%) and Haemoglobin  
27 (20g/dl) level observed in rats fed blend C was significantly higher than the normal control fed  
28 conventional feeds. The increase in packed cell volume (PCV) (50%) and Hb (17g/dl) in diabetic  
29 rats demonstrated that the formulated blend C was able to raise PCV and Hb above 50% and  
30 17g/dl (Normal control NC) respectively. Significant increase ( $P < 0.05$ ) in low density  
31 lipoprotein cholesterol (LDLc) was also observed in all the blends with blend C having the  
32 least (4.0mg/dl) close to NC (2.0mg/dl).

33 **Conclusion:** From the results it is evident that blend C will manage and improve the health  
34 status of diabetic patients.

35 **Key words:** Diabetes mellitus, streptozotocin (STZ), haematology, lipid profile, plant materials

36 **INTRODUCTION**

37  
38 Diabetes mellitus has become a major global problem in our world today. It is a common  
39 disorder associated with increased morbidity and mortality and can be defined as a group of  
40 metabolic diseases characterized by chronic hyperglycemia due to defective insulin secretion,  
41 insulin action, or both, resulting in impaired carbohydrate, lipid, and protein metabolism [1].

42 The combat against diabetes mellitus must be made a matter of top priority by all due to  
43 the continual increase in the global prevalence of this social ill. Globally the prevalence was  
44 estimated to increase in year 2000 to 2010 from 14.2 million to 17.5 million in North America,

45 15.6 million to 22.5 million in South America, 26.5 million to 32.9 million in Europe, 9.4 million  
46 to 14.1 million in Africa, 84.5 million to 132.2 million in Asia and 1.0 million to 1.3 million in  
47 Australia giving a total global increase in prevalence from 151 million people in 2000 to 221  
48 million people in 2010 [2]. This was projected to 324 million by 2025 by Zimmet *et al.* [3] and 366  
49 million 2030 [4]. In 2013, 382 million people had diabetes mellitus worldwide and this is  
50 expected to rise to 592 million by 2035 [5]

51 The increasing interest in herbal medicine for the treatment of diabetes and many  
52 prevailing diseases is not surprising. This may be attributed to the upsurge in cases of drug  
53 resistance, cost and several side effects associated with most orthodox medicines. The use of  
54 plant materials as spices, condiments and for medicinal purposes has therefore become more  
55 popular and as such more plants materials such as plantain and soybean that have low  
56 carbohydrate content with high mineral values are being exploited.

57 There is therefore no doubt that orthodox medicine itself appears to be strongly anchored  
58 on traditional medicine [6]. The fact that the tropics into which majority of Africa lies is host to  
59 about 2/3 of the world's flora and fauna means that a lot of medicinal plants can be found here  
60 for both curative and management of diseases [7].

61 Plantain (*M. paradisiaca*) is a staple food crop in West Africa where its starchy fruits are  
62 generally cooked or fried before consumption. During unripe plantain ripening, the starch is  
63 changed to reducing sugars and sucrose. The medicinal value of plants have assumed a more  
64 important dimension in the past decades owing largely to the discovery that their extracts contain  
65 not only minerals but also a diverse array of secondary metabolites with antioxidant potentials  
66 [8,9]. These antioxidants have been implicated in the therapeutic effects of several plants and  
67 vegetables that are used in traditional medicine [10, 11]. Plantain contains a high fiber content,  
68 and thus is capable of lowering cholesterol and helps to relieve constipation and hence  
69 prevention of colon cancer. Besides, its high potassium content is found to be useful in the  
70 prevention of rising blood pressure and muscle cramp [12]. Various parts of the plant such as the  
71 leaves, root, fruit stalk, bract and fruit have been used for medicinal and domestic purposes.

72 Soybean is known as the "Golden bean" or the super legume of the twentieth century,  
73 because it contains a good proportion of oil more than 20 percent. Soybean is also categorized as  
74 oilseed, represents an excellent source of unsaturated fatty acids, high quality proteins and fiber.  
75 Soybean contains very small amount of saturated fatty acid but do not contain any Trans fatty  
76 acid. Both omega-6 and omega-3 fatty acids such as linoleic acid (56 % of total fat) and alpha  
77 linolenic acid (7-8 % of total fat) are present in soybean. Cooked Soybeans are rich in iron,  
78 phosphorous, magnesium, vitamin B2 (riboflavin) and folate. Kadamet *et al.* [13] stated that  
79 legumes have been known as "a poor man's meat". They supply protein, complex carbohydrates,  
80 fiber and essential vitamins and minerals to the diet, which are low in fat and sodium and contain  
81 no cholesterol.

82 Spices are food adjunct commonly added to food to improve the sensory properties but  
83 many spices have been observed to exert medicinal effects. Some spices which have been  
84 reported to exert hypoglycemic effect both in laboratory animals and human subjects are:  
85 Fenugreek seeds (*Trigonella foenum*), garlic (*Allium sativum*), Onion (*Allium cepa*), turmeric  
86 (*Curcuma longa*), cumin seeds (*Curminum cuminum*), ginger (*Zingiber officinale*), mustard  
87 (*Brassica nigra*), curry leaves (*Murrayakoenigi*) and coriander (*Coriandumsativum*)  
88 [14].

89 Ginger is a perennial plant with narrow, bright green, grass-like leaves. It is cultivated in  
90 the tropics for its edible rhizomes and has been found to be useful for both culinary and medicinal  
91 purposes [15, 16]. Fresh ginger contains 80.9% moisture, 2.3% protein, 0.9% fat, 1.2% minerals,  
92 2.4% fiber and 12.3% carbohydrates. The minerals presented in ginger are iron, calcium and  
93 phosphorous. It also contains vitamins such as thiamine, riboflavin, niacin and vitamin C. The  
94 composition varies with the type, variety, agronomic conditions, curing methods, drying and  
95 storage conditions [17].

96 Several studies have reported the hypoglycemic effect of different forms of ginger in both  
97 animals and human subjects. Among the fairly recent reports are: Arablouet *al.*[18]; Mozaffari-  
98 Khosravi *et al.*[19] and Mahluji *et al.*[20] used ginger powder in Type 2 diabetic patients; Son *et*  
99 *al.*[21] used 6-gingerol isolated from ginger in obese diabetic mice; Sukalingam *et al.*[22] used 6-  
100 gingerol in STZ-induced diabetic rats; Abdulrazaq *et al.* [23] used aqueous ginger extract STZ-  
101 induced diabetic rats; while Jafri *et al.* [24] used aqueous extract in alloxan-induced diabetic rats.  
102 Very limited studies have reported the hypoglycemic effect of ginger juice while there is abject  
103 scarcity of scientific findings on hypoglycemic effect of cooked ginger extract, which is highly  
104 needed since the spice is mostly consumed in cooked forms in various cuisines. Hence, the  
105 objective of this study is to determine the effect of food blends (plantain, soybean and ginger) on  
106 the blood glucose, lipid profile and haematological indices on streptozotocin induced diabetic rats

## 107 MATERIALS AND METHODS

### 108 Materials

109 Unripe plantain and ginger roots were bought from Jattu market in Auchi, Edo State;  
110 defatted soy bean flour (Variety TGX 1448-2E) was purchased from Benin City in Edo State.  
111 Streptozotocin (STZ) Sigma NO SO130 was a product of Sigma-Aldrich chemical company,  
112 UK. Every other chemical used were bought from Promise laboratory in Ekpoma, Edo State.

### 113 Processing of plantain flour:

114 Fresh unripe plantain was peeled, sliced using slicer and dried in an oven at 60°C for  
115 48 hours. Dried sample was ground into powder (plantain flour).

### 116 Processing of soybean to defatted flour:

117 Soybean seeds were cleaned and sorted manually to remove dirt, leaves and stones. The  
118 clean soybean seeds were coarsely milled to separate the coat from the cotyledon. The dehulled  
119 seeds were milled to fine soybean flour using an attrition mill. The fine soybean flour was then  
120 defatted using cold extraction with n-hexane. The defatted flour was then air-dried and the  
121 clumps broken into fine flour, then sieved through a mesh screen.

### 122 Processing of ginger powder

123 Fresh ginger roots were sorted and washed to remove soil and other foreign materials  
124 then sliced to thin layers and dried in an oven at 60°C for 24 hours before milling to powder.

### 125 Formulation of unripe plantain, soybeans and ginger flour blends:

126 Five samples were prepared from the combinations of unripe plantain, defatted soybean  
127 and ginger as blends:  
128  
129

- 130 A=100% unripe plantain  
 131 B=80% unripe plantain, 14% soybean, 6% ginger  
 132 C=70% unripe plantain, 26% soybean, 4% ginger  
 133 D= 60% unripe plantain, 38% soybean, 2% ginger  
 134 E= 50% unripe plantain, 50% soybean

135 **Induction of Diabetes in Wister rats**

136  
 137 A total of 35 adult male albino rats with mean body weight of 219.07g were obtained  
 138 from the disease free stock of the animal house, attached to Ambrose Alli University. The rats  
 139 were separated into seven groups with five rats per group including NC and DC as follows in  
 140 table 1.

141 **Table 1: Rat Groups and Treatments**

Groups	Number of rats	Treatments
A	5	STZ-induced diabetic rats fed with 100% unripe plantain,
B	5	STZ-induced diabetic rats fed with 80% unripe plantain, 14% soybean, 6% ginger),
C	5	STZ-induced diabetic rats fed with 70% unripe plantain, 26% soybean, 4% ginger) and
D	5	STZ-induced diabetic rats fed with 60% unripe plantain, 38% soybean, 2% ginger)
E	5	STZ-induced diabetic rats fed with 50% unripe plantain and 50% soybean),
NC	5	Not induced and fed with rat pellet
DC	5	Induced and fed with rat pellet).

142  
 143 Prior to experimentation, the rats were acclimatized to laboratory condition and fed with  
 144 rat pellet and water ad libitum for a week. Diabetes was induced in rats by a single intraperitoneal  
 145 (I.P) injection of freshly prepared solution of streptozotocin (0.1g dissolved in 5ml of freshly  
 146 prepared sodium citrate buffer 0.1M, pH 4.5) at a dose of 40 mg/kg body weight after fasting for  
 147 12 hours. Good hygiene was maintained by constantly cleaning and removal of faeces and spilled  
 148 feeds from cages daily. Fasting blood glucose (FBG) was determined using Accucheck Active  
 149 glucometer, Roche Germany, with blood obtained from the tail vein of the rats. This test was  
 150 repeated on day 5, 10, 15, 20 and 25. Diabetes was confirmed in STZ treated rats with blood  
 151 glucose concentrations  $\geq 200$  mg/dl.

152  
 153  
 154 **Collection and analysis of blood**

155 The rats were anaesthetized with chloroform vapour, twelve hours(12 h) after last day of feed  
156 administration, and blood samples were collected by cardiac puncture into a set of plain and  
157 fluoride oxalate sample bottles.

158

### 159 **Hematological parameters**

160 The packed cell volume (PCV) was measured by the micro hematocrit centrifuge.  
161 Hemoglobin (Hb) concentration was determined by the cyanomethemoglobin technique [25].  
162 The white blood cell components were also determined.

163

### 164 **Lipid Profile Studies**

165 Blood sample was centrifuged to collect plasma which was used to estimate total  
166 cholesterol, high density lipoprotein cholesterol (HDL-C), low density lipoprotein cholesterol  
167 (LDL-C), and triglycerides (TG) using commercial kits obtained from Randox Laboratories, UK.

168

169

## **RESULTS AND DISCUSSION**

170

### **Mineral Composition of Formulated Food**

171 Table 2 depicts the composition of the studied minerals. Food blend E had the highest  
172 potassium content (1099.42ppm), this was followed by D (944.79ppm) while the lowest  
173 potassium content was observed in A (704.80ppm). The highest potassium observed in food blend  
174 E could be attributed to its high inclusion of soybean (50%) which is known to be a rich source  
175 of potassium. Potassium is an important mineral in the body that regulates fluid balance, muscle  
176 contraction and nerve signals. High potassium may reduce blood pressure and water retention,  
177 protect against stroke and prevent osteoporosis and kidney stones.

178 Food blend A had the highest sodium content (75.65ppm), this was followed by B  
179 (67.19ppm) while the lowest content (47.80ppm) was observed in E. sodium is essential for life. It  
180 helps to control the body's fluid balance. It send nerve impulses and affects muscle function.

181 Food blend E had the highest calcium content (804.02ppm), this was followed by D  
182 (626.91ppm) and C (435.71ppm) while the lowest calcium content (236.16ppm) was observed in  
183 food blends A. calcium plays an important role in muscle contraction, transmitting messages  
184 through the nerves and the release of hormones. Calcium is also important mineral in the  
185 formation of teeth and bones

186 Food blend E had the highest iron content (141.49ppm), this was followed by D  
187 (121.42ppm) and C (114.64ppm) while the lowest content (28.60ppm) was observed in food  
188 blends A. Iron is an important component of haemoglobin, the substance in red blood cell,  
189 responsible for carrying oxygen and transports it throughout the body.

190 The mineral content (potassium, calcium and iron except sodium) of the blends, increased  
191 with increasing soybean inclusion level (Table 2), depicting that soybean is rich in these minerals.

192

193

194

**Table 2:- Mineral composition of formulated food blends**

Blends	Minerals (ppm)				195
	Potassium	Sodium	Calcium	Iron	
A	704.80 <sup>a</sup>	75.65 <sup>a</sup>	236.16 <sup>c</sup>	28.60 <sup>e</sup>	
B	931.82 <sup>b</sup>	67.19 <sup>b</sup>	430.77 <sup>d</sup>	92.89 <sup>d</sup>	
C	942.17 <sup>c</sup>	66.00 <sup>b</sup>	435.71 <sup>c</sup>	114.64 <sup>c</sup>	
D	944.79 <sup>b</sup>	62.08 <sup>c</sup>	626.91 <sup>b</sup>	121.42 <sup>b</sup>	
E	1099.42 <sup>a</sup>	47.80 <sup>d</sup>	804.02 <sup>a</sup>	141.49 <sup>a</sup>	
SEM	0.05	0.54	0.06	0.05	

196 Means with the same letters down the column are not significantly different (P&gt;0.05)

197 A=100% unripe plantain

198 B=80% unripe plantain, 14% soybean, 6% ginger

199 C=70% unripe plantain, 26% soybean, 4% ginger

200 D= 60% unripe plantain, 38% soybean, 2% ginger

201 E= 50% unripe plantain, 50% soybean

202 SEM= Standard error of mean

203

204 **Phytochemical Properties**

205 Table 3 shows the phytochemical compositions of the blends. The lowest tannin content  
 206 (tannin 0.27mg/100g) was observed in food blend A and was followed by blend B (0.55  
 207 mg/100g). Blends C, D and E had the same tannin content (0.61mg/100g).

208 The highest alkaloid content (6.43%) was observed in blend A and was followed by B  
 209 (6.23%), C (5.99%), D (5.75%), and E (4.84%) in that decreasing order.

210 Blends B and C had the same flavonoid content(0.42 mg/100g) which was higher than  
 211 the other blends. The lowest flavonoid content (0.11mg/100g) was observed in blend A; this was  
 212 followed by E(0.31mg/100g) and D (0.35mg/100g).

213 Blend A (0.16 mg/100g) had the lowest saponin content and was followed by B  
 214 (2.39mg/100g), C (3.99mg/100g), D (4.22mg/100g), and E (6.33mg/100g) in that decreasing  
 215 order.

216 Saponins are known to possess both beneficial(cholesterol lowering) and deleterious  
 217 (cytotoxic permeabilization of the intestine and paralysis of the sensory system) properties [26].  
 218 Flavonoids, alkaloids and tannins are polyphenolic compounds with antioxidant properties. In  
 219 addition, phenolic compounds existing in plants are also responsible for their contribution to  
 220 colour, sensory and antioxidant properties of food [27].

221 The low phytochemical values (Table 3) recorded in this study are significantly lower than  
 222 (P<0.05) the results of Elezuet *al.*[28] who recorded significant values (saponin 1.827, flavonoid  
 223 0.981 and tannin 1.577) in unripe plantain flour. However, he further reported that the levels of  
 224 saponin in the flour are quite too low to cause any deleterious effects.

225 **Table 3:- Phytochemical Properties of formulated food blends**

Blends	Phytochemicals			
	Tannin (mg/100g)	Alkaloids (%)	Flavonoids (mg/100g)	Saponin (mg/100g)

A	0.27 <sup>c</sup>	6.43 <sup>a</sup>	0.11 <sup>d</sup>	0.16 <sup>c</sup>
B	0.55 <sup>b</sup>	6.23 <sup>b</sup>	0.42 <sup>a</sup>	2.39 <sup>d</sup>
C	0.61 <sup>a</sup>	5.99 <sup>c</sup>	0.42 <sup>a</sup>	3.99 <sup>c</sup>
D	0.61 <sup>a</sup>	5.75 <sup>d</sup>	0.35 <sup>b</sup>	4.22 <sup>b</sup>
E	0.61 <sup>a</sup>	4.84 <sup>e</sup>	0.31 <sup>c</sup>	6.33 <sup>a</sup>
SEM	0.008	0.014	0.005	0.008

226 Means with the same letters down the column are not significantly different (P>0.05)

227 A=100% unripe plantain

228 B=80% unripe plantain, 14% soybean, 6% ginger

229 C=70% unripe plantain, 26% soybean, 4% ginger

230 D= 60% unripe plantain, 38% soybean, 2% ginger

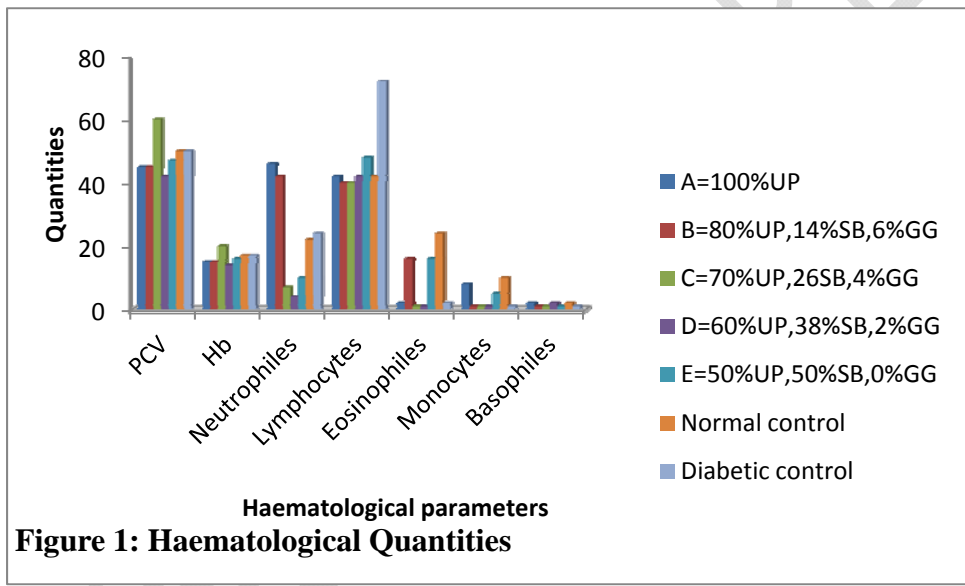
231 E= 50% unripe plantain, 50% soybean

232 SEM= Standard error of mean

233

234

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245

246 Food and nutrients play vital role in the normal functioning of the body. In this study,  
 247 plant materials such as unripe plantain, soybean and ginger were used to formulate food blends  
 248 with the aim of studying its effect on the haematological parameters, lipid profile and blood  
 249 glucose level of diabetic rats.

250 The analysis of variance showed significant difference (P<0.05) in the packed cell  
 251 volume (PCV) and haemoglobin (Hb) level of the diabetic rats (Figure 1). The highest PCV and  
 252 Hb level (60%, 20g/dl respectively) were observed in rat fed blend C that contains 70% unripe  
 253 plantain, 26% soybean and 4% ginger. This was significantly higher (P<0.05) than the  
 254 normal control (not induced) (50%, 17g/dl) fed conventional feeds. The increase in PCV and

255 Hbin diabetic ratsshowed that the formulated blends were able to raise the PCV and Hb above  
 256 50% and 17g/dl.

257 The degree of anemia in diabetic patients can be associated with a number of factors such  
 258 as glomerular filtration rate andglycated h (HbA1c) level. Thomas *et al.*[29] reported that anemia  
 259 is due to diminished erythropoietin production by failing kidneys and increased non enzymatic  
 260 glycosylation of red blood cell (RBC) membrane protein. In this study, increase in PCV and Hb  
 261 level of some of the diabetic rats does not depict occurrence of anemia rather shows its potency  
 262 in the management of the ailment (diabetes). This could be attributed to the phytochemicals and  
 263 mineral present in the blends. The antioxidant properties of these phytochemicals especially  
 264 flavonoids have been reported in several studies. Onat *et al.*[30] reported the anti-sickling  
 265 properties. This according to Palacios *et al.* [31] it prevents oxidation of RBC and Hb that often  
 266 lead to haemolysis. According to Egunyomiet *al.* [32] it may also stimulate formation or  
 267 secretionof erythroprotein in the stem cells of the animals as evidenced by the increased level of  
 268 PCV and Hb. There is no significant difference ( $P < 0.05$ ) in the lymphocytes of the formulated  
 269 blends (A and D) from the normal control. The diabetic control rat had lymphocytes (72%)  
 270 significantly higher ( $P < 0.05$ ) than every otherrat. The high lymphocytes level could be attributed  
 271 to unknown infection.The values of Neutrophiles, Ecsinophiles, Basophiles and  
 272 Monocytesobtained in rats fed with blends C, D and E were significantly lower ( $P < 0.05$ ) than the  
 273 normal control rats.

#### 274 Changes in Blood Glucose and Body Weight of Streptozotocininduced Wister Rats

275 **Table 4: Blood glucose of Streptozotocin rats**

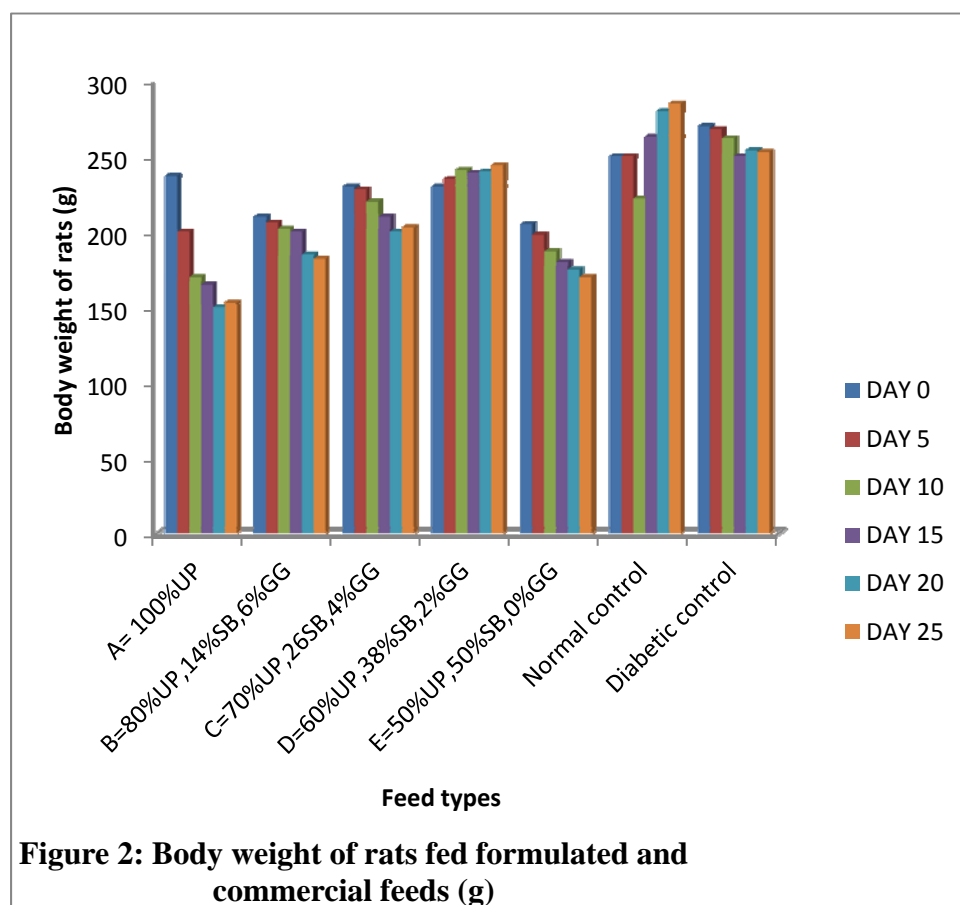
Food blends	Days						SEM
	0	5	10	15	20	25	
A	104.40 <sup>mno</sup>	272.00 <sup>h</sup>	334.00 <sup>e</sup>	504.00 <sup>b</sup>	409.00 <sup>c</sup>	413.20 <sup>c</sup>	
B		93.00 <sup>nop</sup>	245.00 <sup>i</sup>	301.00 <sup>f</sup>	562.00 <sup>a</sup>	559.00 <sup>a</sup>	506.00 <sup>b</sup>
C	107.00 <sup>lmn</sup>	286.00 <sup>g</sup>	73.00 <sup>q</sup>	99.00 <sup>lmno</sup>	101.00 <sup>mno</sup>	85.00 <sup>p</sup>	
D		103.00 <sup>lmno</sup>	307.00 <sup>f</sup>	99.00 <sup>mno</sup>	114.00 <sup>lm</sup>	103.00 <sup>lmno</sup>	90.00 <sup>op</sup> 4.15
E	109.00 <sup>lmn</sup>	247.00 <sup>i</sup>	370.00 <sup>e</sup>	392.00 <sup>d</sup>	375.00 <sup>e</sup>	402.00 <sup>cd</sup>	
NC	108.00 <sup>lmn</sup>	110.00 <sup>lmn</sup>	133.00 <sup>k</sup>	109.00 <sup>lmn</sup>	106.00 <sup>mno</sup>	103.00 <sup>lmno</sup>	
DC	120.00 <sup>l</sup>	229.0 <sup>j</sup>	184.00 <sup>k</sup>	214.00 <sup>k</sup>	225.00 <sup>jk</sup>	283.00 <sup>gh</sup>	

276 Means with same superscript down the column and along the row are not significantly different  
 277 ( $P > 0.05$ )

- 278 A=100% unripe plantain
- 279 B=80% unripe plantain, 14% soybean, 6% ginger
- 280 C=70% unripe plantain, 26% soybean, 4% ginger
- 281 D= 60% unripe plantain, 38% soybean, 2% ginger
- 282 E= 50% unripe plantain, 50% soybean
- 283 NC= Normal control, fed conventional feed (not induced)
- 284 DC= Diabetic control, fed conventional feed (induced)

285





**Figure 2: Body weight of rats fed formulated and commercial feeds (g)**

298

299 Blood glucose and body weight were monitored for total duration of 25 days. At 5 days  
 300 interval blood glucose level and body weight were determined. The initial measurements were  
 301 taken before induction at day 0 for glucose level and body weight. The various rat groups had  
 302 blood glucose level between 93-120mg/dl and body weight between 205-270g (day 0). They were  
 303 induced and fed formulated food blends and water ad libitum.

304 On the 5<sup>th</sup> day, all the induced groups had significant increase ( $P < 0.05$ ) in glucose  
 305 level  $> 200$ mg/dl (Table 4) with corresponding decrease in body weight (Figure 2). Thus the rats  
 306 were considered diabetic at  $\geq 200$ mg/dl. The rat fed blend D had the highest blood glucose level  
 307 307mg/dl. NC rats had the lowest blood glucose level (110.0 mg/dl) and showed no significant  
 308 ( $P > 0.05$ ) change throughout the period of experiment.

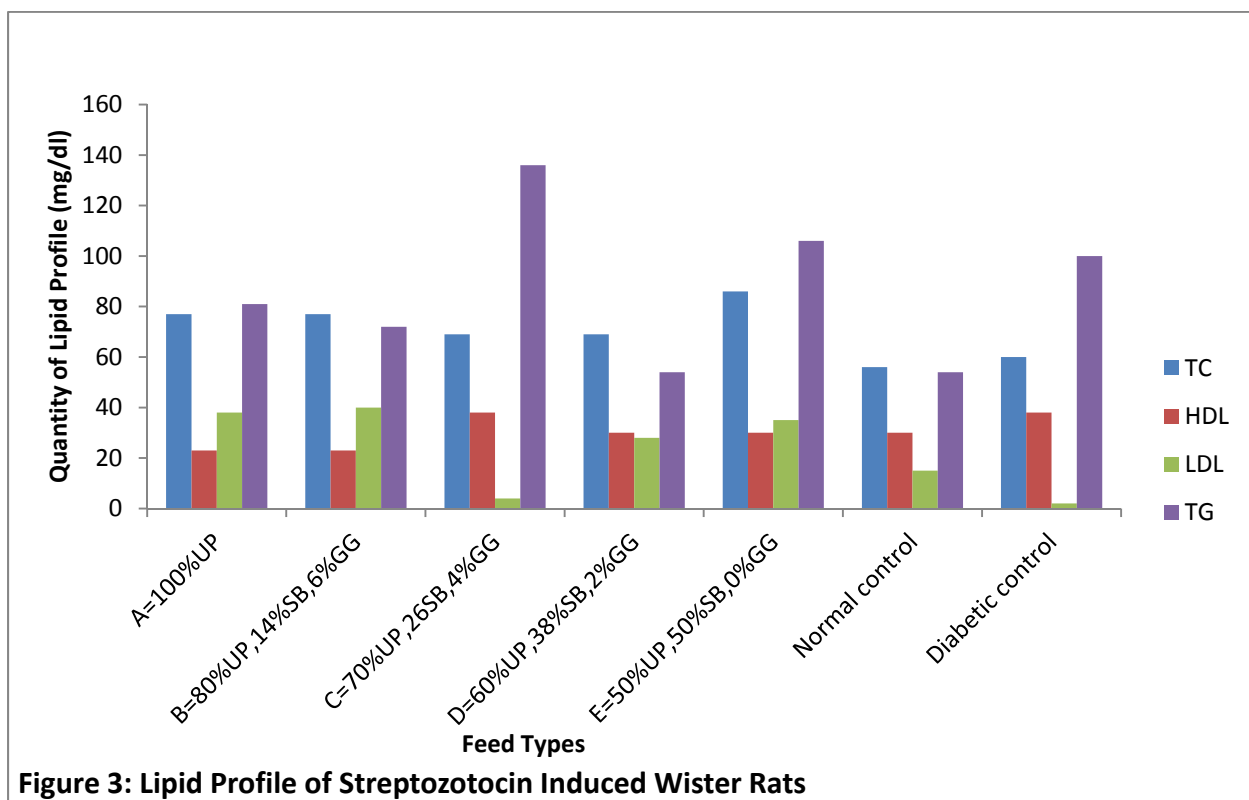
309 There was steady significant increase ( $P < 0.05$ ) in the blood glucose of group A, B, E and  
 310 DC throughout the period of this experiment. The results showed that at 0% and 6% inclusion of  
 311 ginger in blends A and B respectively, the rats were hyperglycemic. This demonstrated that the  
 312 inclusion of ginger at 6% causes rise in BGL. Significant reduction ( $P < 0.05$ ) was observed in the  
 313 blood glucose level of rats fed with blends C and D from 286 to 73mg/dl and 307 to 99mg/dl  
 314 from day 5 to 10 respectively. This same trend was observed in blood glucose level for 15, 20 and  
 315 25 days with rats fed blends C (99, 101 and 85mg/dl) and D (114, 103 and 90mg/dl) respectively,

316 having normal blood glucose <200mg/dl. This shows the potency of the blends C (70% unripe  
317 plantain, 26% Soybean and 4% ginger) and D (60% unripe plantain, 38% soybean and 2%  
318 ginger) in the management of the ailment (diabetes). This could be attributed to the combination  
319 levels of the plant materials particularly the inclusion of ginger at 4% and 2% in blend C and D  
320 respectively. Ginger provides an amount of potassium that could help stroke and diabetes and  
321 adult requires 2000mg of potassium each day. Potassium is important for diabetic patients and  
322 those at the risk of it. The findings of recent study published by researchers from university of  
323 Sydney in 2012 revealed that ginger extract helps to increase cell absorption of glucose even  
324 independent of insulin ([www.naturalnews.com](http://www.naturalnews.com)). The predominant pungent compound in ginger is  
325 responsible for its benefit to humans [33]. According to Andalluet *al.* [34] ginger has a  
326 therapeutic benefit of lowering fasting serum blood glucose level in Type 2 diabetes. According  
327 to Singh *et al.* [35] many of the putative activities of ginger (antioxidant, anti-inflammatory,  
328 hepatoprotective, antiobesity) are often associated with the etiology and pathophysiology of  
329 Type 2 diabetes, which suggest the possibility that ginger may not have a direct effect on  
330 diabetes but acts indirectly by suppressing factors that lead to impaired glucose control. Thus,  
331 was supported by a study showing that ginger root powder (200mg/kg body weight) in type 2  
332 diabetic rat model reversed symptoms of metabolic syndrome, blood glucose, blood lipid and  
333 decreased oxidative stress [36]. Although blend B had ginger inclusion at 6%, the glucose level  
334 was >200mg/dl throughout the period of this experiment. This shows that ginger inclusion at 6%  
335 could result in hyperglycemic condition. However at day 10, a rise was observed in the glucose  
336 level of rats fed blend A (272-334mg/dl), B (245-301mg/dl) and E (247-370mg/dl) with  
337 corresponding decrease in body weight (Figure 2). Thus this indicates that the formulation for A,  
338 B and E could not control the diabetic condition. The DC rats fed with conventional rat feed  
339 increased in blood glucose level and body weight steadily throughout the period of experiment,  
340 while the body weight decreased from 228 to 220g and increased from 235 to 241g in rats fed  
341 blends C and D respectively.

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### Effect of Food Blends (Diet) on Serum Lipid Profile



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**Figure 3: Lipid Profile of Streptozotocin Induced Wister Rats**

359 Serum lipid concentration of streptozotocin induced rats fed with food blends and  
360 conventional feed in this study is shown in figure 3. From the results, serum high density  
361 lipoprotein cholesterol (HDLc) concentration in rats fed blends C (38mg/dl) and NC  
362 (38mg/dl) were same but higher and significantly different ( $P < 0.05$ ) from HDLc of rats fed with  
363 other blends.

364 Total cholesterol (TC) increased in all the blends. However, the lowest concentration in  
365 TC was observed in blends C and D (Figure 3). Thus this depicts that blends C (69mg/dl) and D  
366 (69mg/dl) are better having lower cholesterol concentration.

367 For low density lipoprotein cholesterol (LDLc) significant increase ( $P < 0.05$ ) was  
368 observed in all the blends. However, blend C (4.0mg/dl) was next to NC (2.0mg/dl) while the  
369 highest was observed in blend B (40.0mg/dl).

370 The increased in LDLc, TC, and decreased in HDLc agrees with the findings of  
371 Adaramoye *et al.* [37] for diabetic rats. Besides, the formulated diets are plant materials  
372 containing phytochemicals. [38] reported that action of plant extract in reducing plasma

373 cholesterol concentration could be due to the ability of one or more of the phytochemicals in the  
374 plant to activate the functioning enzymes of the rats responsible for cholesterol absorption.

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## CONCLUSION

379 In this research work, it was observed that the blends of unripe plantain, soya beans and  
380 ginger in adequate proportion (C=70% unripe plantain, 26% soybean, 4% ginger or D= 60%  
381 unripe plantain, 38% soybean, 2% ginger) could help to reduce blood glucose, improve  
382 haematological parameters and lipid profile. The mineral content (potassium, calcium and iron  
383 except sodium) of the blends increased with increasing soybean inclusion level, depicting that  
384 soybean is rich in these minerals. Significant reduction ( $P < 0.05$ ) was observed in the blood  
385 glucose level of rats fed blends C and D from 286 to 85mg/dl and 307 to 90mg/dl respectively.  
386 The lowest concentration of TC was observed in blends C and D. This depicts that blends C  
387 (69mg/dl) and D (69mg/dl) are better and preferred to the other blends. In addition, blend C also  
388 had the least value (4.0mg/dl) of low density lipoprotein cholesterol (LDLc). Hence, blend C  
389 is most preferred to prevent and control diabetes as well as improve the health status of diabetic  
390 patients.

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### Ethical Approval:

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394 As per international standard or university standard ethical approval has been collected  
395 and preserved by the authors.

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