

2 **REGULATED EFFECTS OF *CAPSICUM FRUTESCENS* SUPPLEMENTED DIET**  
3 **(C.F.S.D) ON FASTING BLOOD GLUCOSE LEVEL, BIOCHEMICAL**  
4 **PARAMETERS AND **BODY WEIGHT** IN ALLOXAN INDUCED DIABETIC**  
5 **WISTAR RATS.**

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14 **ABSTRACT**  
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**Aim of the study:** This study investigates the effects of *Capsicum frutescens* supplemented diet (C.F.S.D) on fasting blood glucose level, biochemical parameters and body weight in alloxan induced diabetic Wistar rats.

**Experimental Design:** 130 – 150g healthy forty male Wistar rats were divided into four groups as following; Group 1 served as a normal control and received normal feed-. Group 2 (Diabetic control) received normal feed-. Group 3 (Diabetic test 1) received normal feed + 1g *Capsicum frutescens*-. Group 4 (Diabetic test 2) received normal feed + 2g *Capsicum frutescens*.

**Place and Duration of study:** This study was carried out in the department of Physiology, Faculty of Basic Medical Sciences, Delta State University, Abraka and the feeding lasted for three weeks. At the end of the experiments, the animals were sacrificed, blood samples were collected and then the serum was further subjected to biochemical analysis using biochemical analyzer (Reflotron Plus).

**Results:** AST, ALT, ALP, GGT, Creatinine, Uric acid, total cholesterol and fasting blood sugar level in serum were increased however the high density lipoprotein cholesterol (HDL-c) of serum was decreased in diabetic control (group 2), compared with non-diabetic control (group 1). The administered *Capsicum frutescens* in the diet at 1g and 2g doses significantly reduced the fasting blood glucose level as well as the serum level of AST, ALT, ALP, GGT, Creatinine, Uric acid, total cholesterol, compared with diabetic control. Serum HDL was also significantly increased when compared with diabetic control  $P < 0.05$ . Decrease in body weight in diabetic control group and increased in body weight of 1g and 2g *Capsicum frutescens* supplemented diet groups were also observed.

**Conclusion:** The observed improvement in the biochemical parameters, blood glucose levels and body weight of alloxan induced diabetic Wistar rats by 1g and 2g *Capsicum*

*frutescens* supplemented diet suggests *Capsicum frutescens* to possess, cardio-protective and anti-diabetic properties.

**Recommendation:** The incorporation of *Capsicum frutescens* as spice in the diet of individuals who are diabetic, hypertensive and obese, is worthy of recommendation.

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21 **Keywords:** Capsicum Frutescens, Fasting Blood Glucose, Liver enzymes, Capsaicin,  
22 Thermogenesis.

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## 25 1. INTRODUCTION

26

27 Diabetes mellitus (DM) has been described as a multifactorial disease that is characterized  
28 by hyperglycemia and lipoprotein disorders [1], increased basal metabolic rate [2], defect in  
29 reactive oxygen species scavenging enzymes, as well as altered metabolism of major food  
30 substances [2]. Diabetes is a major degenerative disease in the world today [3], affecting at  
31 least 15 million people and resulting in complications which include hypertension,  
32 atherosclerosis and microcirculatory disorders. Diabetes mellitus is a syndrome of impaired  
33 carbohydrate, fat and protein metabolism caused by either lack of insulin secretion or  
34 decreased sensitivity of the tissue to insulin

35 At least 80% of Africans depend on plant medicine for their healthcare [4]. Today, medicinal  
36 plants are increasingly being used in most parts of the world as: hypolipidemic [5];  
37 antihypertensive [6]; treatment for skin diseases [7] and hypoglycemic [8], agents.

38 For the past 25 years, epidemiological studies have revealed a diminished occurrence of  
39 chronic diseases in populations consuming diets fortified with fruits and vegetables, [9]. It  
40 has been suggested that antioxidants found in large quantities in fruits and vegetables may  
41 be responsible for this protective effect, [10]. In the past three decades, resulting from  
42 several studies, it has been documented that some common spices can also exert beneficial  
43 health effects, [11; 12]. These beneficial health effects of spices in most instances have  
44 been reported to be as a result of their chemical composition; some of these beneficial  
45 health effects of spices documented are hypolipidemic and antioxidant properties [13]

46 One of such plant that produce spices is *Capsicum frutescens*; a short lived evergreen shrub  
47 that usually grows from 1 to 1.5m in height and 1 to 3cm in basal stem diameter. It is  
48 commonly recognized by its fruit, the large red, orange, or yellow chili peppers that the plant  
49 produces. *Capsicum frutescens* fruits grow as long pods, and when ripe they develop their  
50 characteristic red coloring. This plant originated in south or Central America, then spread

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51 quickly throughout the subtropical regions and still grows wild today. The plant grows in  
52 tropical climates, because it needs a warm, humid climate to survive. It had been reportedly  
53 used in the treatment of various ailments such as diabetes, blood pressure [high/ low],  
54 bronchitis, burning feet, arthritis, etc [14].

55 **Accumulating evidence** has shown multiple pharmacological effects of Capsicum on a  
56 variety of physiological systems such as cardiovascular system, gastro-intestinal tract,  
57 metabolic rate, and pain relief, [15].

58 Previous research **have** shown the Chemo-Protective effect of spices among which are;  
59 *Turmeric, Capsicum frutescens, Cloves* and *Cardamom* **in** correcting iron overload-induced  
60 liver injury, oxidative stress and serum lipid profile in rat model. The incorporation of chili  
61 (*Capsicum frutescens*) in the diet at 2g significantly restored the enzyme activities of the liver  
62 AST, ALT, and ALP to normal level [16].

63 The **chemical substance** in *Capsicum frutescens* that gives **it** the hot and spicy flavor was  
64 identified as capsaicin, [15]. Red chili (*Capsicum frutescens*) is widely used as a spice for  
65 flavoring foods, particularly in South- East Asian and Latin-American countries. **Several**  
66 **studies indicate that capsaicin (red pepper) is an appetite suppressant which can slightly**  
67 **increase metabolism. Spicing up one's foods with capsaicin-containing spices and using red**  
68 **pepper as a condiment can aid in increasing the rate of fat burning or thermogenesis. In an**  
69 **article published in the British Journal of Nutrition, Yoshioka et al (2001)<sup>[17]</sup> concluded that**  
70 **the consumption of red pepper and caffeine can induce a considerable change in energy**  
71 **balance when individuals are given free access to foods.** Pungent capsaicinoids (capsaicin,  
72 dihydrocapsaicin), antioxidant vitamins (ascorbic acid, vitamin E), carotenoids ( $\beta$ -carotene,  
73  $\beta$ - cryptoxanthine) and several organic acids and minerals are the major **active chemical**  
74 **substance found in** *Capsicum frutescens*, [18]. Capsaicin (8-methyl-*N*-vanillyl-6-  
75 nonenamide) is an irritant for mammals, including humans, and produces a sensation of  
76 burning **on** any tissue with which it comes into contact. The burning and painful sensations  
77 associated with capsaicin result from its chemical interaction with sensory neurons.  
78 Capsaicin, as a member of the vanilloid family, binds to a receptor called the vanilloid  
79 receptor subtype 1 (VR1), [19].

80 **However, there are not enough scientific documentation on the effects of *Capsicum***  
81 ***frutescens* supplemented diet on biochemical parameters in a diabetic state. The present**  
82 **study was designed depending on this background.**

83

84 **2. MATERIAL AND METHODS**

85

86 **Chemicals and equipments:**

87 Red Chili (*Capsicum frutescens*), purchased from Abraka market in Ethiopie East local  
88 government area, Delta State, which was authenticated by Dr. (Mrs). N.E. Edema in the  
89 department of Botany, Faculty of Science, Delta State University, Abraka. It was then air-  
90 dried at room temperature ( $22\pm 1^{\circ}\text{C}$ ) for 14 days until a constant weight was attained and  
91 was then blended with the aid of a grinding machine and stored in an airtight container for  
92 use in the experiment. Alloxan monohydrate (Sigma, alpha Aesar, 25g. A15324, CAS:2244-  
93 11-3. Cotton wool, Hand gloves, Dissecting kit, Centrifuge, Pipettes, Growers mash  
94 ,Beakers, Electronic weighing balance, Syringes and needles, Marker pen, Oncall Redii  
95 Glucometer with diagnostic glucose strips, and Reflotron plus kit.

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99 **FRESH AND DRIED *CAPISCUM FRUTESCENS* FRUITS**

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102 **PREPARATION OF PEPPER SUPPLEMENTED DIET**

103 1g and 2g *Capsicum frutescens* supplemented diet were prepared by weighing 1g and 2g  
104 of blended *Capsicum frutescens* and mixing them with 99g and 98g of animal feed  
105 (growers mash) respectively.

106 **COMPOSITION OF THE GROWERS MARSH**

107 Protein-19.0%

108 Fat -2.85%

109 Fibre – 6.00%

110 Calcium – 1.00%

111 Available phosphate – 0.45%

112 Energy – 2875 KGC

113 (Animal Care Services Konsult (NIG) LTD).

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## 115 **HANDLING OF EXPERIMENTAL ANIMALS**

116 Forty (40) **healthy** Male Wistar rats weighing 130-150g were **acquired** from the International  
117 institute of tropical agriculture, (IITA), Ibadan Nigeria. They were acclimatized for 14 days  
118 **before commencement of** the experiment. The rats were kept in well ventilated wooden  
119 cages, **in a room with optimal humidity and temperature, and fed growers marsh, with** water  
120 *ad libitum*. **Procedures followed in raising the experimental animals were in accordance with**  
121 **the ethical standards of the Institutional Animals Ethics Committee (IAEC). And permission**  
122 **for the use of animals and animal protocol was obtained from the Research Ethics**  
123 **Committee of Delta State University, Abraka.**

### 124 **Induction of diabetes**

125 Thirty (30) animals were **fasted** for 24hours (but with free access to water) **and then the**  
126 **diabetes was by injecting** a single intraperitoneal dose of alloxan monohydrate (150mg/kg)  
127 prepared in stock of 1500mg/50ml and a concentration of 30mg/ml. **After** three days, rats  
128 with fasting blood glucose concentration above 200mg/dl were **considered** diabetic and  
129 **selected for the experiment**

## 130 **EXPERIMENTAL PROCEDURE**

131 **Diabetic rats** were randomly **divided** into 3 different groups and **rats the were not induced**  
132 **were grouped as normal control (Group 1) as following;**

133 **Group 1:** Non diabetic rats received normal diet (non-diabetic control)

134 **Group 2:** Diabetic rats received normal diet (diabetic control)

135 **Group 3:** Diabetic rats received 1g *Capsicum frutescens* supplemented diet (test 1 group)

136 **Group 4:** diabetic rats received 2g *Capsicum frutescens* supplemented diet (test 2 group).

137

138 **Each animal was fed a 5g meal formulated by mixing 1g and 2g *Capsicum frutescens* with**  
139 **99g and 98g animal feed** and treatment was done twice daily for twenty one days. Rats'  
140 initial body weight prior to commencement of treatment was recorded. Inclusion criteria in  
141 this study were non diabetic animals (which served as positive control), and animals with  
142 evidence of diabetes. Exclusion criteria include those animals that died during **induction of**  
143 **diabetes and the treatment period**. Thus higher numbers of animals were allocated to groups  
144 1, 2 and 3.

## 145 **BLOOD COLLECTION AND BIOCHEMICAL ASSAY**

146 After twenty one days of treatment, **the rats** were anaesthetized using chloroform and then  
147 sacrificed. Blood samples was collected by cardiac puncture were delivered into lithium  
148 heparin bottles. The tubes were then centrifuged at 4000rpm for ten minutes to obtain clear

149 serum which were later subjected to biochemical evaluation for ALT, AST, ALP, GGT, URIC  
 150 ACID, CREATININE, HDL, and TOTAL CHOLESTEROL using Reflotron plus kit.  
 151 Fasting blood glucose level was determined with the aid of glucose analyzer machine  
 152 (Oncall- Redii glucometer) by collecting blood samples from tail veins of overnight fasted  
 153 animals. Values were expressed in mg/dl.

### 154 STATISTICAL ANALYSIS

155 The result of this study were expressed as mean  $\pm$  SEM, and were analyzed by one way  
 156 analyses of variance (ANOVA) using statistical package for social science (SPSS, 16).  
 157 Difference between the means were tested with post Hoc- Tukey's test for multiple  
 158 comparison and significance was considered when  $p < 0.05$ . Student's dependent t-test was  
 159 used to analyze the significant difference between body weight before treatment and after  
 160 treatment.

161

### 162 3. RESULTS AND DISCUSSION

163

164 **Table 1:**

165 **Effects of *Capsicum frutescens* supplemented diet on biochemical parameters of**  
 166 **alloxan induced diabetic Wistar.**

	<b>Group 1: Non- Diabetic control</b>	<b>Group 2: Diabetic control</b>	<b>Group 3: Diabetic +1g C.F.S.D</b>	<b>Group 4: Diabetic + 2g C.F.S.D.</b>
<b>Creatinine (IU/L)</b>	0.42 $\pm$ 0.03	0.94 $\pm$ 0.17 <sup>a</sup>	0.4 $\pm$ 0.3 <sup>b</sup>	0.54 $\pm$ 0.07 <sup>b</sup>
<b>Uric acid (IU/L)</b>	5.49 $\pm$ 0.2	7.87 $\pm$ 0.85 <sup>a</sup>	5.03 $\pm$ 0.2 <sup>b</sup>	6.3 $\pm$ 0.7
<b>GGT (IU/L)</b>	223.4 $\pm$ 7.5	275.0 $\pm$ 10.7 <sup>a</sup>	221.8 $\pm$ 6.4 <sup>b</sup>	224.8 $\pm$ 6.0 <sup>b</sup>
<b>AST (IU/L)</b>	278.4 $\pm$ 19.6	325.2 $\pm$ 26.1	247.2 $\pm$ 10.8 <sup>b</sup>	251.8 $\pm$ 12.3
<b>ALP (IU/L)</b>	251 $\pm$ 6.81*	316.4 $\pm$ 37.7*	327.6 $\pm$ 27.6*	243.8 $\pm$ 4.53*
<b>ALT (IU/L)</b>	61.7 $\pm$ 1.03*	128.2 $\pm$ 32.97*	98.98 $\pm$ 8.74*	87.86 $\pm$ 8.54*
<b>HDL (mg/dl)</b>	47.98 $\pm$ 1.8 <sup>ns</sup>	43.1 $\pm$ 2.8	46.8 $\pm$ 1.6 <sup>ns</sup>	46.0 $\pm$ 1.4 <sup>ns</sup>
<b>T.Cholesterol (mg/dl)</b>	85.6 $\pm$ 5.6	79.2 $\pm$ 4.4	101.6 $\pm$ 3.3 <sup>b</sup>	61.5 $\pm$ 3.4 <sup>abc</sup>
<b>InitialBlood glucoselevel (mg/dl)</b>	88.8 $\pm$ 6.22	380.2 $\pm$ 16.6	363.8 $\pm$ 24.3 <sup>d</sup>	382.2 $\pm$ 14.7 <sup>d</sup>
<b>FinalBlood glucoselevel (mg/dl)</b>	94.8 $\pm$ 6.18 (6.8%)	370.0 $\pm$ 19.81 <sup>a</sup> (-2.63%)	182.8 $\pm$ 16.82 <sup>abd</sup> (-49.8%)	146.6 $\pm$ 14.8 <sup>bd</sup> (-61.6%)

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167 Values are expressed as mean  $\pm$  Standard error of mean (S.E.M), n=10 \*P<0.05;  
 168 Significant as determined by one way analysis of variance. Significant difference (<sup>abc</sup>P  
 169 < 0.05): (a) compared to group 1, (b): to group 2, (c): to group 3.<sup>d</sup>P<0.05: Significant  
 170 when initial and final fasting blood glucose level were compared in groups 3 and 4.  
 171 Values in parenthesis depict the percentage change in FBGL when initial and final  
 172 values were compared. Significant difference (<sup>ns</sup>P< 0.05) HDL, comparing groups 1, 3,  
 173 4 with group 2 .

174  
 175 **AST- (Aspartate Transaminase)**

176 **ALT- (Alanine amino Transaminase)**

177 **,ALP- (Alkaline Phosphatase)**

178 **GGT- (Gamma Glutamyl Transpeptidase)**

179 They are all liver enzymes(biomarkers) of liver damage.

180

181

182 **Table 2:**

183 **Effects of *Capsicum frutescens* (C.F.) supplemented diet on body weight of alloxan**  
 184 **induced diabetic rats.**

	Body weight before treatment Week 0 (g)	Body weight after treatment Week 3 (g)
Group 1 (Normal control)	131 $\pm$ 9.8	195 $\pm$ 17.2 (48.9%)
Group 2 (Diabetic control)	140 $\pm$ 9.6	120 $\pm$ 7.9 (-16.7%)
Group 3 (Diabetic, 1g C.F.S.D)	125 $\pm$ 6.7	134 $\pm$ 19.2 (7.2%)
Group 4 (Diabetic, 2g C.F.S.D)	140 $\pm$ 7.2	152 $\pm$ 16.9 (8.5%)

185 Values are expressed as mean  $\pm$  Standard error of mean (SEM), n = five animals per  
 186 group. C.F: *Capsicum frutescence*.

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190 **DISCUSSION**



191 The action of capsaicin is mediated by TRPV1 (vanilloid receptor), which belongs to an ion  
192 channel group. TRPV1 when activated permits cations to pass through the cell membrane and  
193 into the cell resulting in depolarization of the neuron stimulating it to signal the brain. By  
194 binding to the TRPV1 receptor, the capsaicin molecule produces the same sensation that  
195 excessive heat or abrasive damage would cause, explaining why the spiciness of capsaicin  
196 is described as a burning sensation. The inflammation resulting from exposure to Capsaicin  
197 is believed to be the result of the body's reaction to nerve excitement rather than just  
198 chemical burn or any direct tissue damage when chili peppers are the source of exposure.  
199 Capsaicin is the chemical in chili peppers that contributes to their spiciness; capsaicin  
200 stimulates a receptor found in sensory neurons, creating the heat sensation and subsequent  
201 reactions like redness and sweating.

202 In the study by Yoshioka et al (2001)<sup>[17]</sup>, 8.6g and 7.2g red pepper, added to lunch and  
203 dinner respectively and caffeine consumption significantly reduced the cumulative ad libitum  
204 energy intake and increased energy expenditure. Almost 1000 additional calories per day  
205 were burned by combining caffeine consumption with substances containing red pepper.

206 The New York Daily News published an article, "15 fat-burning foods" about the capsaicin  
207 and caffeine combination that simply states "men who consume coffee and red pepper-  
208 packed snacks and meal burned almost 1000 more calories a day than the control group".

209 Yasser (2008)<sup>[22]</sup> found that capsaicin can create "heat" in a more direct manner by altering  
210 the activity of a muscle protein called SERCA. Normally, muscle contraction is initiated  
211 following the release of a wave of calcium ions from a compartment called the sarcoplasmic  
212 reticulum. SERCA then actively pumps the calcium back into the sarcoplasmic reticulum  
213 (using ATP energy), causing muscle relaxation and renewing the cycle. Capsaicin, however  
214 can attach to SERCA and "uncouple" this pumping activity, that is, the protein still burns ATP  
215 energy but does not use it to pump calcium. Instead, all the ATP energy is given off as heat.  
216 This uncoupling known as thermogenesis, is one important method of staying warm and is  
217 most often seen in hibernating animals. Yasser noted also that capsaicin is the first natural  
218 compound known to augment the thermogenesis process. The findings further explained  
219 how capsaicin intake can increase metabolism and body temperature. The study also noted  
220 that though relatively high amounts of capsaicin (probably more than someone could eat),  
221 was required to effectively achieve the desired result, but the structure of capsaicin could be  
222 used as a model to design more potent compounds that might have clinical use such as  
223 treating hypothermia.

224 Avraham et al (2008)<sup>[23]</sup>, in their study titled, “ Cannabinoids and capsaicin improve liver  
225 function following thioacetamide-induced acute injury in mice”, reported an improvement  
226 both in liver pathology and function.

227 In the present study, there was an observed decrease in body weight of the Wistar rats in  
228 the treated groups compared to the normal control group (table 2). However, an increase in  
229 body weight was observed when the treated groups were compared to the diabetic control  
230 group. Though, capsicum frutescens has been reported to aid the rate of fat burning [17], in  
231 a diabetic state it can actually reduce the rate of loss of the body’s protein (muscles). This is  
232 possibly achieved through the activities of the antioxidant vitamins such as ascorbic acids  
233 and vitamin E present in capsicum frutescens [18], which helps to counteract the effect of  
234 the reactive oxygen species.

235 Significant reduction in FBGL in 1g (group 3) and 2g (group 4) C.F.S.D treated groups may  
236 be attributed to the presence of hypoglycemic agents in *Capsicum frutescens*. Studies had  
237 shown that *Capsicum frutescens* is used to treat diabetes mellitus by traditional healers in  
238 Jamaica, [24]. Pharmacokinetic and effect of Capsaicin in *Capsicum Frutescens* on  
239 decreasing plasma glucose level in a crossover study of 12 healthy volunteers by performing  
240 the OGTT while receiving placebo or 5 grams of capsicum had been documented [25].

241 Impaired carbohydrate utilization in the **diabetes** also leads to accelerated lipolysis, which  
242 results in elevated plasma triglycerides levels (hyperlipidemia), [26]. The observed  
243 abnormalities of triglyceride and HDL metabolism are in accordance with reports on early  
244 manifestation of insulin resistance, the precursor to diabetes [27; 28]. **In this study, treatment**  
245 **with 2g capsicum frutescens resulted in a reduction in serum level of total cholesterol when**  
246 **compared to the control groups. Manjunatha and Srinivasan (2008), in the study also**  
247 **reported the hypolipidemic and antioxidant potency of capsicum frutescens.**

248 Individuals with type 2 diabetes had also been reported to have a higher incidence of liver  
249 function test abnormalities **compared to** non diabetic individuals. Mild chronic elevations of  
250 transaminases often reflect underlying insulin resistance. Diabetes mellitus can arise as a  
251 result of insulin insufficiency, which is associated with altered activity of various liver  
252 enzymes, [20]. Grossi, *et al.*, (1998)<sup>21</sup> had also reported that values of serum ALP can be  
253 raised in diabetic patients. The liver releases alanine aminotransferase (ALT) and an  
254 elevation in **serum levels** are an indicator of liver damage, [28]. The levels of aspartate  
255 aminotransferase (AST), alanine amino transaminase (ALT) and alkaline phosphatase (ALP)  
256 had been reported to be increased in alloxan-induced diabetic rats, [29]. Increased in serum  
257 liver enzymes parameters in diabetic control group observed in the present study

258 corroborates these findings. Reduction in liver enzyme levels in group 3 (1g, C.F.S.D.) and 4  
259 (2g C.F.S.D.) clearly indicates the therapeutic **potency** of *Capsicum frutescens* against  
260 increased in serum liver enzyme parameters **seen in** alloxan induced diabetes. In previous  
261 research, *Capsicum frutescens* had been documented to protect against iron overload liver  
262 injury by reducing plasma liver parameters levels to normal, [16].

263 There was a significant increase in serum creatinine level of group 2. An increase in plasma  
264 creatinine levels may be a sign of impaired renal function which is associated with diabetes.  
265 The elevation in the plasma creatinine concentration indirectly suggests kidney damage  
266 specifically the renal filtration mechanism, [30]. Significant reduction observed in the serum  
267 creatinine levels of the diabetic rats treated with 1g and 2g C.F.S.D in this study suggests  
268 protective effect by *Capsicum frutescens* against kidney disorders associated with diabetes  
269 mellitus.

#### 270 **4. CONCLUSION**

271  
272 In this study, increase in serum liver enzymes (AST, ALT, ALP, GGT), increased in serum  
273 uric acid, creatinine, total cholesterol, fasting blood glucose level and reduced high density  
274 lipoprotein (HDL) cholesterol associated with alloxan induced diabetes mellitus were  
275 reversed after treatment with 1g and 2g *Capsicum frutescens* supplemented diet. Such  
276 remarkable changes observed in this study could be traced to the chemical substances  
277 [capsaicin, dihydrocapsaicin, antioxidant vitamins (ascorbic acid, vitamin E), carotenoids ( $\beta$ -  
278 carotene,  $\beta$ - cryptoxanthine) and several organic acids and minerals present in *Capsicum*  
279 *frutescens*.**The thermogenic and protein sparing properties of capsicum frutescens has been**  
280 **reported by several authors and results from this study also lends credence to this fact. It's**  
281 **therefore recommended that *Capsicum frutescens* be added as spices to the food of obese**  
282 **individual as well as diabetic patients for its hypoglycemic properties, inducing of increase**  
283 **energy utilization as well as being cardio-protective by its effect on plasma lipids.**

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#### 289 **AUTHORS' CONTRIBUTIONS**

290 Author 1 designed the study and wrote the first draft of the manuscript. Author 2 managed  
291 the literature searches; author 3 performed the statistical analysis and managed the  
292 analyses of the study. All authors read and approved the final manuscript.

293

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