

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: Assessment of the effects of *Capsicum frutescens* supplemented diet (C.F.S.D) on fasting blood glucose level and biochemical parameters 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 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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24

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 intermediary metabolism of
30 major food substances [2]. Diabetes is a major degenerative disease in the world today [3],
31 affecting at least 15 million people and having 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 rely 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], agent.

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, it has been
42 experimentally documented that several common spices can also exert health beneficial
43 physiological effects, [11; 12]. These physiological effects of spices in most instances have
44 been traced to the bioactive chemicals; some of these physiological effects of spices
45 documented are hypolipidemic and antioxidant properties with beneficial health implications,
46 [13]

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

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

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

59 Previous research had shown the Chemo-Protective effect of spices among which are;
60 *Turmeric, Capsicum frutescens, Cloves* and *Cardamom* on Correcting Iron Overload-
61 Induced Liver Injury, Oxidative Stress and Serum Lipid Profile in Rat Model. The
62 incorporation of chili (*Capsicum frutescens*) in the diet at 2g significantly restored the
63 enzyme activities of the liver AST, ALT, and ALP to normal level. The mean values of lipid
64 profile, the MDA and serum total bilirubin were also reduced, [16].

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

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

86

87 2. MATERIAL AND METHODS

88

89 **Chemicals and equipments:**

90 All chemical used in the research were procured as follows:

91 Red Chili (*Capsicum frutescens*), purchased from Abraka market in Ethiope East local
92 government area, Delta State, which was authenticated by Dr. (Mrs). N.E. Edema in the
93 department of Botany, Faculty of Science, Delta State University, Abraka. It was then air-
94 dried at room temperature ($22\pm 1^{\circ}\text{C}$) for 14 days until a constant weight was attained and
95 was then blended with the aid of a grinding machine and stored in an airtight container for
96 use in the experiment. Alloxan monohydrate (Sigma, alpha Aesar, 25g. A15324, CAS:2244-
97 11-3. Cotton wool, Hand gloves, Dissecting kit, Centrifuge, Pipettes, Growers mash
98 ,Beakers, Electronic weighing balance, Syringes and needles, Marker pen, Oncall Redii
99 Glucometer and Reflorton plus^(R) reflectance photometer (Roch Diagnostic GmbH, D-68298).

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103 **FRESH AND DRIED *CAPISCUM FRUTESCENS* FRUITS**

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106 **PREPARATION OF PEPPER SUPPLEMENTED DIET**

107 1g and 2g *Capsicum frutescens* supplemented diet were prepared weighing 1g and 2g of
108 powdered *Capsicum frutescens* and mixing them with 99g and 98g of animal feed
109 (growers mash) respectively.

110 **COMPOSITION OF THE GROWERS MARSH**

111 Protein-19.0%

112 Fat -2.85%

113 Fibre – 6.00%

114 Calcium – 1.00%

115 Available phosphate – 0.45%

116 Energy – 2875 KGC

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

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119 **HANDLING OF EXPERIMENTAL ANIMALS**

120 Forty (40) **healthy** Male Wistar rats weighing 130-150g were from the International institute
121 of tropical agriculture, (IITA), Ibadan Nigeria. They were acclimatized for 14-days at in the
122 animal house unit in the Department of Pharmacology, Faculty of Basic Medical Science,
123 Delta State University Abraka before the experiment. The rats were kept in well ventilated
124 wooden cages. They were exposed to 12 hours of natural daylight and darkness and fed
125 standard rat feed and water *ad libitum*. **Procedures followed in raising the experimental**
126 **animals were in accordance with the ethical standards of the Institutional Animals Ethics**
127 **Committee (IAEC). And permission for the use of animals and animal protocol was obtained**
128 **from the Research Ethics Committee of Delta State University, Abraka.**

129 **Induction of diabetes**

130 Thirty (30) animals were **fasted** for 24hours (but with free access to water) **and then the**
131 **diabetic model was reproduced by injecting** a single intraperitoneal dose of alloxan
132 monohydrate (150mg/kg) prepared in stock of 1500mg/50ml and a concentration of
133 30mg/ml. **After** three days, rats with fasting blood glucose concentration above 200mg/dl
134 were confirmed diabetic. Diabetic state was maintained for three days for well establishment
135 of diabetes.

136 **EXPERIMENTAL PROCEDURE**

137 **Diabetes** mellitus **rats** were randomly **allotted** into 3 different groups and **non diabetic rats as**
138 **normal control (Group 1) as following;**

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

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

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

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

143

144 **Each animal was fed a 5g meal formulated by mixing 1g and 2g *Capsicum frutescens* with**
145 **99g and 98g animal feed** and treatment was done twice daily for twenty one days. Rats'
146 initial body weight prior to commencement of treatment was recorded. Inclusion criteria in
147 this study were; non diabetic that were not induced with diabetes (which served as positive
148 control), and animals with evidence of diabetes. Exclusion criteria include those animals that
149 died during the maintenance of diabetes. Thus higher numbers of animals were allocated to
150 groups 1, 2 and 3.

151 **BLOOD COLLECTION AND BIOCHEMICAL ASSAY**

152 After twenty one days of treatment, all overnight fasted rats were anaesthetized using
153 chloroform and then sacrificed. Blood samples collected by cardiac puncture were delivered

154 into lithium heparin bottles. The tubes were then centrifuged at 4000rpm for ten minutes to
 155 obtain clear serum which were later subjected to biochemical evaluation for ALT, AST, ALP,
 156 GGT, URIC ACID, CREATININE, HDL, and TOTAL CHOLESTEROL using Reflotron plus
 157 kit.

158 Fasting blood glucose level was determined with the aid of glucose analyzer machine
 159 (Oncall- Redii glucometer) by collecting blood samples from tail veins of overnight fasted
 160 animals. Values were expressed in mg/dl.

161 STATISTICAL ANALYSIS

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

168 3. RESULTS AND DISCUSSION

171 **Table 1:**
 172 **Effects of *Capsicum frutescens* supplemented diet on biochemical parameters of**
 173 **alloxan induced diabetic Wistar.**

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

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(mg/dl)

174 Values are expressed as mean \pm Standard error of mean (S.E.M), n=10 *P<0.05;
175 Significant as determined by one way analysis of variance. Significant difference (^{abc}P
176 < 0.05): (a) compared to group 1, (b): to group 2, (c): to group 3.^dP<0.05: Significant
177 when initial and final fasting blood glucose level were compared in groups 3 and 4.
178 Values in parenthesis depict the percentage change in FBGL when initial and final
179 values were compared. Significant difference (^{ns}P< 0.05) HDL, comparing groups 1, 3,
180 4 with group 2 .

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182 **AST- (Aspartate Transaminase)**

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

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

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

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

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188

189 **Table 2:**

190 **Effects of *Capsicum frutescens* (C.F.) supplemented diet on body weight of alloxan**
191 **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%)

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

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197 **DISCUSSION**

198 The action of capsaicin is mediated by TPRV1 (vanilloid receptor), which belongs to an ion
199 channel group. VR1 when activated permits cations to pass through the cell membrane and
200 into the cell resulting in depolarization of the neuron stimulating it to signal the brain. By
201 binding to the VR1 receptor, the capsaicin molecule produces the same sensation that
202 excessive heat or abrasive damage would cause, explaining why the spiciness of capsaicin
203 is described as a burning sensation. The inflammation resulting from exposure to Capsaicin
204 is believed to be the result of the body's reaction to nerve excitement rather than just
205 chemical burn or any direct tissue damage when chili peppers are the source of exposure.

206 Capsaicin is the chemical in chili peppers that contributes to their spiciness; capsaicin
207 stimulates a receptor found in sensory neurons, creating the heat sensation and subsequent
208 reactions like redness and sweating.

209 In the study by Yoshioka et al (2001)^[17], 8.6g and 7.2g red pepper, added to lunch and
210 dinner respectively and caffeine consumption significantly reduced the cumulative ad libitum
211 energy intake and increased energy expenditure. Almost 1000 additional calories per day
212 were burned by combining caffeine consumption with substances containing red pepper.

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

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

229 used as a model to design more potent compounds that might have clinical use such as
230 treating hypothermia.

231 Avraham et al (2008)^[23], in their study titled, “ Cannabinoids and capsaicin improve liver
232 function following thioacetamide-induced acute injury in mice”, reported an improvement
233 both in liver pathology and function.

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

242 Significant reduction in FBGL in 1g (group 3) and 2g (group 4) C.F.S.D treated groups may
243 be attributed to the presence of hypoglycemic agents in *Capsicum frutescens*. Studies had
244 shown that *Capsicum frutescens* is used to treat diabetes mellitus by traditional healers in
245 Jamaica, [24]. Pharmacokinetic and the effect of Capsaicin in *Capsicum Frutescens* on
246 decreasing Plasma Glucose level in a crossover study of 12 healthy volunteers by
247 performing the OGTT while receiving placebo or 5 grams of capsicum had been documented
248 [25].

249 Impaired carbohydrate utilization in the diabetic also leads to accelerated lipolysis, which
250 results in elevated plasma triglycerides levels (hyperlipidemia), [26]. The observed
251 abnormalities of triglyceride and HDL metabolism are in accordance with reports on early
252 manifestation of insulin resistance, the precursor to diabetes [27; 28]. From the result of this
253 study, 2g C.F.S.D treated group elicited reduction in serum level of total cholesterol than 1g
254 treated group. The physiological effects of most spices had been documented to exhibit
255 hypolipidemic and antioxidant properties with beneficial health implication, [13].

256 Individuals with type 2 diabetes had also been reported to have a higher incidence of liver
257 function test abnormalities than non diabetic individuals. Mild chronic elevations of
258 transaminases often reflect underlying insulin resistance. Diabetes mellitus can arise as a
259 result of insulin insufficiency, which is associated with altered activity of various liver
260 enzymes, [20]. Grossi, *et al.*, (1998)²¹ had also reported that values of serum ALP can be
261 raised in diabetic patients. The liver releases alanine aminotransferase (ALT) and an

262 elevation in plasma concentrations are an indicator of liver damage, [28]. The levels of
263 aspartate aminotransferase (AST), alanine amino transaminase (ALT) and alkaline
264 phosphatase (ALP) had been reported to be increased in alloxan-induced diabetic rats, [29].
265 Increased in serum liver enzymes parameters in diabetic control group observed in the
266 present study corroborates these findings. Reduction in liver enzyme levels in group 3 (1g,
267 C.F.S.D.) and 4 (2g C.F.S.D.) clearly indicates the therapeutic role of *Capsicum frutescens*
268 against increased in serum liver enzyme parameters correlated with alloxan induced
269 diabetes. In previous research, *Capsicum frutescens* had been documented to protect
270 against iron overload liver injury by reducing plasma liver parameters levels to normal, [16].
271 There was a significant increase in serum creatinine level of group 2. An increase in plasma
272 creatinine levels may be a sign of impaired renal function which is associated with diabetes.
273 The elevation in the plasma creatinine concentration indirectly suggests kidney damage
274 specifically the renal filtration mechanism, [30]. Significant reduction observed in the serum
275 creatinine levels of the diabetic rats treated with 1g and 2g C.F.S.D in this study suggests
276 protective effect by *Capsicum frutescens* against kidney disorders associated with diabetes
277 mellitus.

278 **4. CONCLUSION**

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

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297 **AUTHORS' CONTRIBUTIONS**

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

301

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