

# The Anti-Diabetic Potential of Thermally Treated Garlic, Turmeric, and Ginger in Pre-Diabetic Male Wistar Rat Model

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## Abstract

Spices (turmeric (T), ginger (GI), and garlic (GA) (TGG)) have been used for centuries for food preservation, flavors, and medicinal properties. Research suggests that TGG contain potent antioxidants that may prevent and/or delay chronic diseases such as cancer, diabetes, and heart diseases. Heat treatment of spices may potentially increase antioxidative activity by modifying the inherent chemical structure of potent antioxidative compounds in spices. The purpose of this study was to determine the anti-diabetic potential of thermally treated TGG on Wistar male rats. Two-week-old male Wistar rats were randomly assigned to 8 groups (N = 24, n = 3) including control AIN-93G Diet (C) and high fat (HFD) and high sugar (HS) (glucose 10%) diet and treatment HFD/HS diets containing T, GA, GI (1% and 2%) singly for 11 weeks. Weekly feed intake, body weight, and blood glucose levels were recorded. Rats were sacrificed at 13 wks. by CO<sub>2</sub> asphyxiation. Liver, pancreas, adipose (thigh), cecum, femur, urine, and serum samples were collected for quantitative determination of detoxification and antioxidative enzyme analysis, bone mineralization, and cholesterol using standard protocol. Of spice-incorporated diets, rats fed turmeric (1%) exhibited the lowest reduction in blood glucose levels at 90 mg/dL compared to the control 58 mg/dL. Additionally rats fed TGG at both concentrations resulted in an induction of antioxidant (GSH) and antioxidant enzyme (GPx) activity with significantly ( $p \leq 0.05$ ) higher levels compared to the control. Serum total cholesterol levels were lower in spice-incorporated diets compared to control HFD/HS fed rats. Therefore, the use of thermal application on spices presents promise in potentiating the antioxidant effects and thereby their potential health promoting properties.

## Keywords

Type 2 Diabetes, Bioactive Compounds, Wistar Rats, Spices

## 1. Introduction

Diabetes is a chronic disease characterized by hyperglycemia due to insufficient and/or ineffective insulin secretion [1]. Individuals with diabetes are at an increased risk for the development of chronic diseases and health complications such as cardiovascular disease, blindness, amputation, and kidney failure [2]. Prediabetes or Impaired Glucose Tolerance (IGF) is a precursor to diabetes affecting 86 million Americans [3]. Prediabetes is a condition where individuals' blood glucose levels are above normal levels yet not classified as being diabetic. According to Babey *et al.* [4] it is estimated that one in three adults in the US has prediabetes and without intervention 30% of these individuals will develop type 2 diabetes and 70% will develop it within their lifetime. High incidences of diabetes have been attributed to diet, specifically the western diet [5] [6]. The western diet is composed of high fat and high sugar intakes and low intakes of fruits, vegetables, and whole grains. Maintaining a healthy diet is critical in the prevention and/or treatment of diabetes as increased consumption of fruits and vegetables has shown to have a reduced incidence of the development of diabetes [7] [8] [9]. Research has shown that spices, herbs, and their extracts possess antimicrobial, anti-inflammatory, anti-rheumatic, lipid-lowering, hepatoprotective, nephroprotective, antimutagenic and anticancer activities [10] [11] [12]. Research has shown garlic, turmeric, and ginger to contain anti-diabetic properties attributed to the presence of phytochemicals curcuminoids, gingerols, and sulfur containing compounds [13] [14] [15] [16]. Wongsu *et al.* [17] reported that both garlic and turmeric showed potential inhibition against  $\alpha$ -amylase and  $\alpha$ -glucosidases ranging from 0.00% - 58.38% and 6.75% - 100.00%, and may be good candidates to help treat and manage the early stages of hyperglycemia. Additionally these spices have shown to lower serum glucose, insulin, and triglycerides in male wistar rats [13]. Although these spices have been used for centuries for their medicinal properties additional research is required to substantiate these therapeutic and pharmacological claims. Therefore the purpose of this study was to determine the anti-diabetic potential of thermally treated garlic, turmeric, and ginger on pre-diabetic wistar male rats.

## 2. Materials and Methods

### *Spice Preparation*

Garlic (*Allium sativum*), ginger (*Zingiber officinale*), and turmeric (*Curcuma longa*) powders were purchased from Monterey Bay Spice Company, (Watsonville, CA). Spices were subjected to direct heat treatments for experimentation. Direct heat-treated turmeric, ginger, and garlic powders were conducted by stovetop toasted as follows garlic 1 minute toasting (70°C - 100°C), ginger 5 minute toasting (70°C - 130°C) and turmeric 5 minute toasting (70°C - 130°C) based upon preliminary chemical analysis. Spices were then incorporated in an HFD/HS AIN-93G diet singly at 1% and 2% concentrations.

### *Animal Care*

A total of 24 two week old Wistar male rats (Harlan Laboratories, Indianapolis, IN) weighing 209 - 253 g were housed in an environmentally controlled animal care facility for the duration of the study. The temperature and relative humidity were kept at  $20^{\circ}\text{C} \pm 1^{\circ}\text{C}$  and 50%. Light and dark cycles were at 12 hours each. The rats were allowed a 2-week acclimation period and fed the American Institute of nutrition diet (AIN-93G) and water [18]. At 6 weeks of age, rodents (Wistar males) were housed in stainless steel cages (2/cage). After acclimation rodents were randomly assigned to groups each consisting of 3 rats followed by the introduction of high fat (40% Lard/soybean) and high sugar diet (10% glucose). Spices and fat incorporated into the basal AIN-93G diet (Table 1) were replaced by corn starch. Weekly weight and blood glucose measurements conducted until the end of the experimental period.

Rats had ad-libitum access to treatment diets throughout the experimental period (11 weeks). At the end of the testing period (11 weeks) a single collection of blood was taken followed by immediate euthanasia by carbon dioxide asphyxiation. Vital organs of each rat were excised, blotted and weighed and the organ/body weight ratios calculated. The animals in the study were handled in accordance with the AAMU guidelines for the protection and care of animals. The Institute of Animal Care and Use Committee (IACUC) committee approved the protocol for the study before beginning experiment.

**Table 1.** Short term animal feed diet.

Ingredient	AIN-93G Composition Diets							
	Control (g)	HFD/HS-Control	HFD/HS-T 1%	HFD/HS-T 2%	HFD/HS-GA 1%	HFD/HS-GA 2%	HFD/HS-GI 1%	HFD/HS-GI 2%
Cornstarch	397.5	257.5	247.5	237.5	247.5	237.5	247.5	237.5
Sucrose	100	100	100	100	100	100	100	100
Casein	200	200	200	200	200	200	200	200
Fiber	50	50	50	50	50	50	50	50
Soybean Oil	70	105	105	105	105	105	105	105
Lard	0	105	105	105	105	105	105	105
Dextrose	132	132	132	132	132	132	132	132
Mineral Mix (G)	35	35	35	35	35	35	35	35
Vitamin Mix	10	10	10	10	10	10	10	10
L-Cysteine	3	3	3	3	3	3	3	3
Choline	2.5	2.5	2.5	2.5	2.5	2.5	2.5	2.5
Garlic	0	0	0	0	10	20	0	0
Ginger	0	0	0	0	0	0	10	20
Turmeric	0	0	10	20	0	0	0	0
TOTAL	1000	1000	1000	1000	1000	1000	1000	1000

### *Organs & Biological Samples Preparation*

During autopsy, blood was taken from rats from each group, and the serum separated and cholesterol and glucose were determined using standard kits (Cayman Chemicals Ann Arbor, MI, USA). Urinary pH was determined by measurement with pH testing strips obtained from EM Science (Gibbstown, N.J.). Test strips were immersed in urine samples for 2 minutes and pH was recorded. Femurs were excised from rats and weights of bones, bone length, and diameter will be recorded. Selected minerals (Mg, Ca, P, Zn, and Fe) were analyzed by inductively coupled plasma (ICP) (AOAC 984.27).

### *Antioxidant and Detoxification Enzymes*

Liver samples were used to measure detoxification and antioxidant enzymes Glutathione (GSH), Glutathione S-transferase (GST), Glutathione peroxidase (GPx), Superoxide dismutase (SOD), and catalase (CAT) of rat livers according to the manufacturer's instructions (Cayman Chemical Company, Ann Arbor, MI). Briefly, one gram of liver was homogenized in 10ml of TRIS hydrochloric acid mixture. Following homogenization, liver solutions were centrifuged for 20 minutes at 5000 xg at 4°C. The supernatant was decanted into test tubes. Following decantation, the precipitant was washed with an additional amount of TRIS hydrochloric acid and centrifuged for 20 min at 10,000 XG. The supernatant was collected and used for analysis.

### *Statistical Analysis*

Results are presented as means  $\pm$  SEM using SAS system version 9.3. ANOVA was used to determine any significant differences among the treatment groups. Significance was determined at  $P \leq 0.05$ . The means were separated using Tukey's Studentized Range Test.

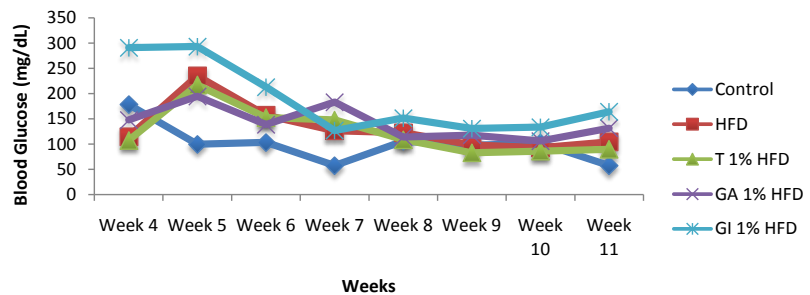
## **3. Results & Discussion**

Blood glucose levels were recorded weekly for all treatment groups for 11 weeks (**Figure 1** and **Figure 2**). When comparing rats fed 1% spice-incorporated diets to the control (**Figure 1**), spice diets resulted in a gradual decline of blood glucose levels throughout the study (11 weeks). Initial blood glucose levels ranged from a high of 291 mg/dL (GI 1%) to a low of 107 mg/dL (T 1%). Final blood glucose levels ranged from a low of 57 mg/dL (control) to a high of 164 mg/dL (GI 1%). Although all spice diets resulted in higher blood glucose levels compared to the control, turmeric 1% (90 mg/dL) diet lowered blood glucose levels to a level closest to control (57.66 mg/dL) fed rats.

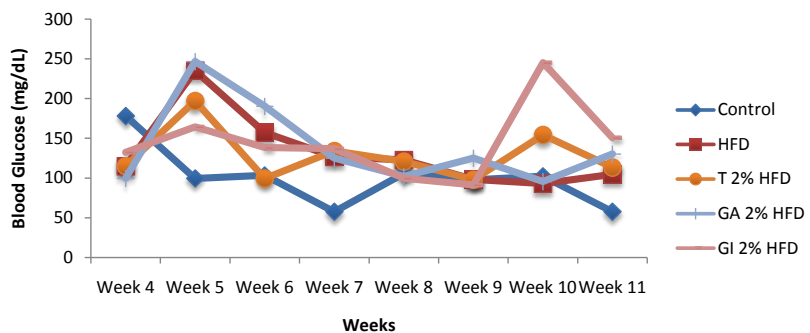
When comparing 2% spice-incorporated diets to the control (**Figure 2**), all spice diets resulted in higher blood glucose compared to the control (no HFD/HS). Initial blood glucose levels ranged from a high of 178 mg/dL (control) to a low of 99 mg/dL (GA 2%). Final blood glucose levels ranged from a low of 57 mg/dL (control) to a high of 150 mg/dL (GI 2%). Among the dietary treatments, rats fed ginger at 1% & 2% showed higher blood glucose levels.

**Table 2** and **Figure 3** show the effect of feeding control and treatment diets

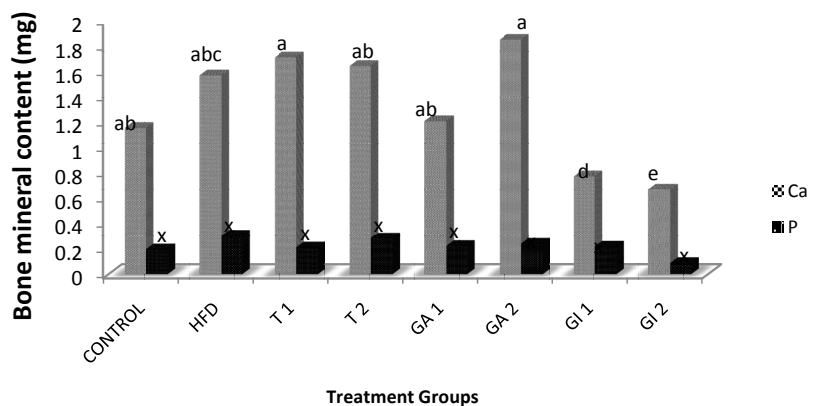
on bone mineralization in Wistar male rats. Bone minerals (Fe, Zn, Mg, Mn, Cu, K, Ca, and P) were analyzed. There were no significant differences found in bone



**Figure 1.** Rat blood glucose data for control, HFD, low (1%) concentration thermally treated groups. Abbreviations: HFD = High fat diet, HS = High sugar, T1 = HFD/HS + turmeric 1%, T 2 = HFD/HS + turmeric 2%, GA 1 = HFD/HS + garlic 1%, GA 2 = HFD/HS + garlic 2%, GI 1 = HFD/HS + ginger 1%, GI 2 = HFD/HS + ginger 2%.



**Figure 2.** Blood glucose data for rats fed control, HFD, high (2%) concentration thermally treated spices. Abbreviations: HFD = High fat diet, HS = High sugar, T1 = HFD/HS + turmeric 1%, T 2 = HFD/HS + turmeric 2%, GA 1 = HFD/HS + garlic 1%, GA 2 = HFD/HS + garlic 2%, GI 1 = HFD/HS + ginger 1%, GI 2 = HFD/HS + ginger 2%.



**Figure 3.** Bone mineralization (calcium and phosphorus) in rats. <sup>abc, x</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet, HS = High sugar, T1 = HFD/HS + turmeric 1%, T 2 = HFD/HS + turmeric 2%, GA 1 = HFD/HS + garlic 1%, GA 2 = HFD/HS + garlic 2%, GI 1 = HFD/HS + ginger 1%, GI 2 = HFD/HS + ginger 2% Ca = calcium, P = phosphorus.

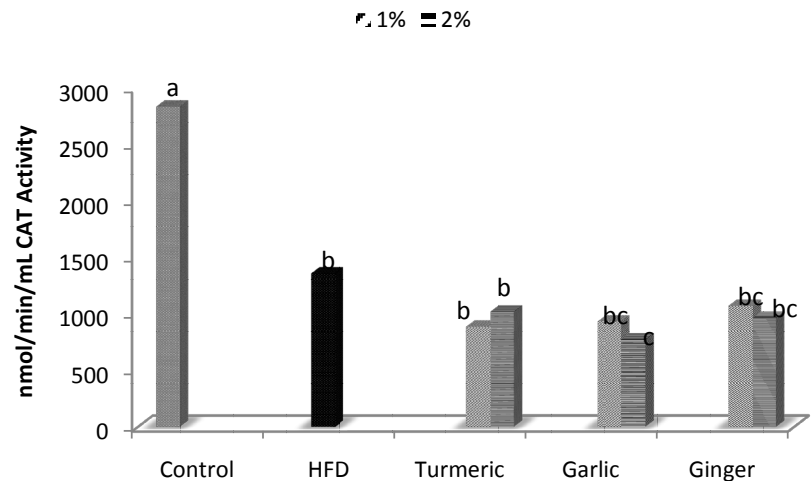
**Table 2.** Bone mineralization in rats.

	Fe (mg)	Zn (mg)	Mg (mg)	Mn (mg)	Cu (mg)	K (mg)
CONTROL	0.00015 <sup>a</sup> ± 0	0.00048 <sup>ab</sup> ± 0	0.0085 <sup>ab</sup> ± 0	0.000013 <sup>a</sup> ± 0	0.000016 <sup>b</sup> ± 0	0.0062 <sup>ab</sup> ± 0
HFD/HS	0.00033 <sup>a</sup> ± 0	0.00067 <sup>a</sup> ± 0	0.012 <sup>a</sup> ± 0	0.0000046 <sup>a</sup> ± 0	0.000014 <sup>b</sup> ± 0	0.0090 <sup>a</sup> ± 0
HFD/HS + T 1%	0.00019 <sup>a</sup> ± 0	0.00043 <sup>ab</sup> ± 0	0.0078 <sup>ab</sup> ± 0	ND	0.0000092 <sup>b</sup> ± 0	0.0047 <sup>ab</sup> ± 0
HFD/HS + T 2%	0.00029 <sup>a</sup> ± 0	0.00062 <sup>ab</sup> ± 0	0.011 <sup>ab</sup> ± 0	0.0000107 <sup>a</sup> ± 0	0.000040 <sup>a</sup> ± 0	0.0090 <sup>a</sup> ± 0
HFD/HS + GA 1%	0.00024 <sup>a</sup> ± 0	0.00046 <sup>ab</sup> ± 0	0.0083 <sup>ab</sup> ± 0	0.0000032 <sup>a</sup> ± 0	0.000014 <sup>b</sup> ± 0	0.0055 <sup>ab</sup> ± 0
HFD/HS + GA 2%	0.00026 <sup>a</sup> ± 0	0.00049 <sup>ab</sup> ± 0	0.0089 <sup>ab</sup> ± 0	0.0000034 <sup>a</sup> ± 0	0.000021 <sup>b</sup> ± 0	0.0051 <sup>ab</sup> ± 0
HFD/HS + GI 1%	0.00025 <sup>a</sup> ± 0	0.00046 <sup>ab</sup> ± 0	0.0084 <sup>ab</sup> ± 0	ND	0.000019 <sup>b</sup> ± 0	0.0050 <sup>ab</sup> ± 0
HFD/HS + GI 2%	0.000058 <sup>a</sup> ± 0	0.00018 <sup>ab</sup> ± 0	0.0036 <sup>ab</sup> ± 0	ND	0.0000067 <sup>b</sup> ± 0	0.0021 <sup>c</sup> ± 0

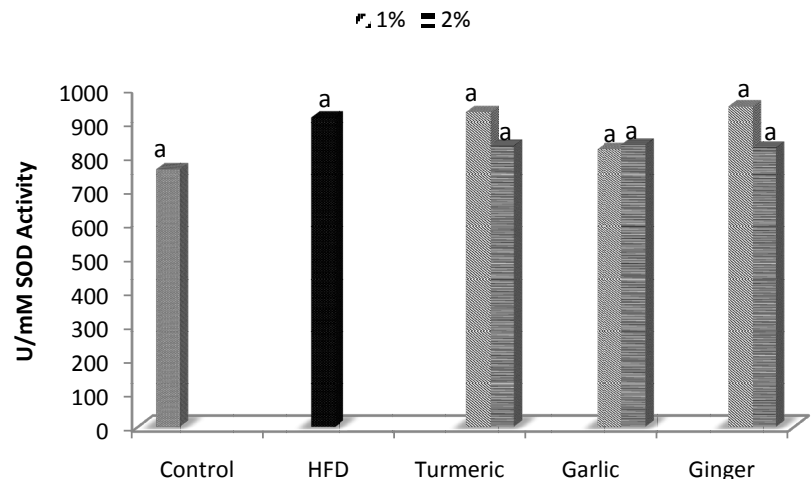
<sup>abc</sup>Values with superscripts in columns are significantly different ( $p \leq 0.05$ ). Results expressed as Means  $\pm$  SEM,  $n = 3$ . Abbreviations: T = Turmeric, GA = Garlic, GI = Ginger, HFD = High fat diet, HS = High sugar, Mn = manganese, Cu = copper, K = potassium.

mineralization for Fe, Zn, Mg, Mn, and P. Rats fed HFD/HS + Turmeric 2% had significantly ( $p \leq 0.05$ ) higher levels of copper in bones compared to other rats fed treatment and control fed diets. While rats fed HFD/HS + Ginger 2% had significantly ( $p \leq 0.05$ ) lower levels of potassium compared to other treatment and control fed rats. Bone Ca levels (**Figure 3**) were significantly ( $p \leq 0.05$ ) lower in rats fed HFD/HS + ginger 1% & 2%, compared to other groups. Ginger is more fibrous as a spice compared to both garlic and turmeric. Studies [19] [20] have shown that fiber consumption may interfere with calcium absorption and bioavailability, due to fiber chelating calcium making it unavailable for use and then excreted.

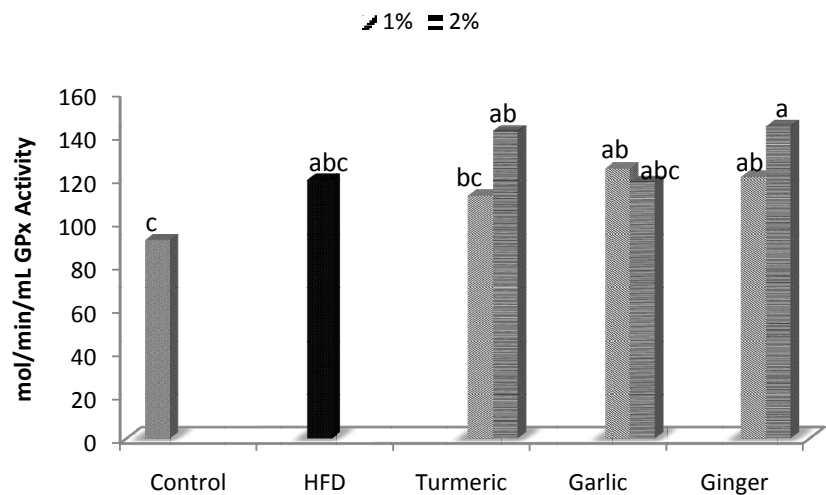
Antioxidant and detoxification enzyme activities in Wistar rats (hepatic tissues) are reported in **Figures 4-8**. Catalase activity (CAT) was significantly ( $p \leq 0.05$ ) higher in control (2839.14 nmol/min/mL) fed rats compared to the rats fed treatment diets (**Figure 4**). Rats fed HFD/HS diets had significantly ( $p \leq 0.05$ ) higher CAT activity compared to rats fed HFD/HS + Garlic 2% (766.99 nmol/min/mL). However, there were no significant differences among the other treatment groups. There were no significant differences seen in SOD activity (**Figure 5**). However, all treatment diets resulted in higher SOD activity compared to the control. SOD activity ranged from a low of 761.32 U/mL (control diet) to a high of 947.68 U/mL (HFD/HS + Ginger 1% diet). GPx activity (**Figure 6**) was significantly ( $p \leq 0.05$ ) induced in rats fed spice-incorporated diets compared to the control. Highest GPx activity was seen in rats fed HFD/HS + Ginger 2% diet (144.32 mol/min/mL). SOD serves as a first line of defense, and catalyzes the conversion of the superoxide radical into  $H_2O_2$ , which further breaks down to  $H_2O$  and  $O_2$  by CAT and GPx. Our findings showed that rats fed spice-incorporated diets showed a low induction of SOD yet significantly ( $p \leq 0.05$ ) higher induction of GPx for detoxification. There was also no dose dependent response observed in the induction of enzymatic activity between 1% and 2% diets.



**Figure 4.** Catalase activity in wistar male rats. <sup>abc</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet.

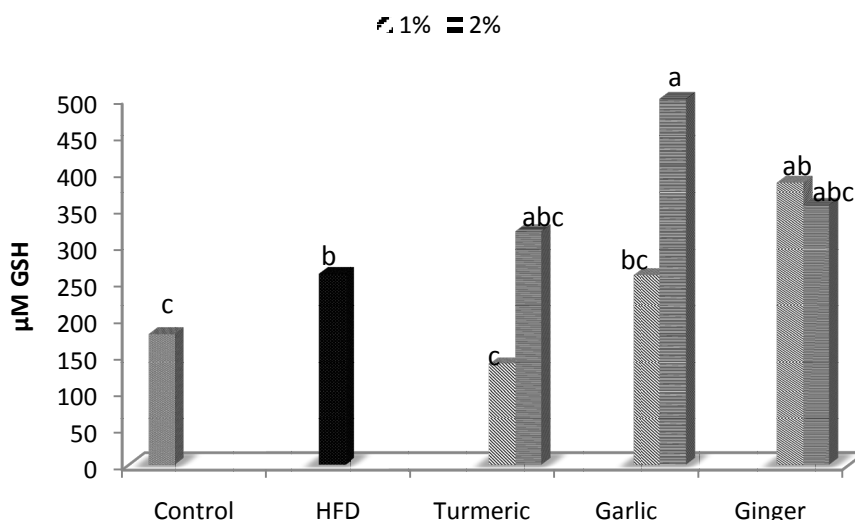


**Figure 5.** Superoxide dismutase activity in wistar male rats. <sup>abc</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet.

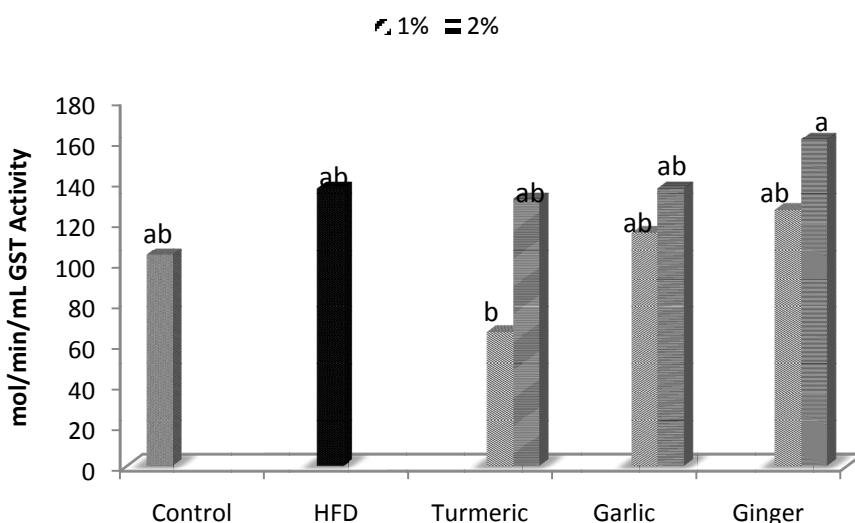


**Figure 6.** Glutathione peroxidase activity in wistar male rats. <sup>abc</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet.





**Figure 7.** Glutathione levels in wistar male rats. <sup>abc</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet.



**Figure 8.** Glutathione S-transferase activity in wistar male rats. <sup>abc</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet.

The route of detoxification among endogenous enzymes expressed may vary depending on a number of factors including vehicle introduced, dietary components, and genetic predispositions. Wohaieb & Godin [21] showed that enzymes (SOD, GPx, and CAT) in insulin treated and non-treated diabetic Wistar rats, significantly ( $p \leq 0.05$ ) higher induction of SOD and CAT compared to GPx. While Suryanarayana *et al.* [22] reported similar results compared to our study where they fed curcumin to STZ-induced Wistar rats, reporting significantly ( $p \leq 0.05$ ) higher GPx activity in liver tissues compared to CAT activity.

**Figure 7** and **Figure 8** show GSH level and GST (liver) activities in Wistar male rats fed control and treatment diets. The levels of antioxidant glutathione ranged from a low of 137.86  $\mu\text{M}$  (HFD/HS + Turmeric 1%) to a high ( $p \leq 0.05$ ) of GSH at 500  $\mu\text{M}$  (HFD/HS + Garlic 2%). Both turmeric and garlic incorpo-

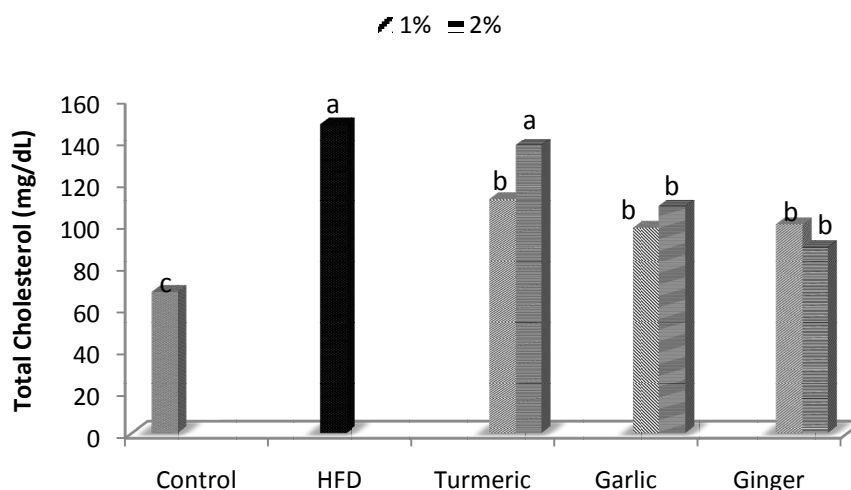


rated diets resulted in a 2-fold increase (dose response) with increasing from 1% to 2% level. Rats fed the control diet were seen to have lower levels of GSH compared to other treatment groups and was significantly ( $p \leq 0.05$ ) lower compared to rats fed diets with 2% garlic and 1% ginger. However, there was no significant induction of GST activity among control and treatment groups. Among spice-incorporated diets, 2% diets seemed to result in higher GST activity compared to their 1% counterparts, with a 2-fold increase in GST activity between 1% & 2% turmeric fed rats. The primary function of GST is for the detoxification of xenobiotic compounds that are conjugated to glutathione; these compounds are then excreted from the organism [23]. Decreased levels of GSH have been reported among individuals with diabetes [24] [25] [26].

Serum cholesterol levels in Wistar rats are shown in **Figure 9**. Cholesterol levels ranged from a low of 67 mg/dL (control) to a high of 146 mg/dL (HFD/HS). There was a significant ( $p \leq 0.05$ ) increase in cholesterol levels in all treatment groups compared to that of the control (normal fat & sugar). Among rats fed spice-incorporated diets, rats fed ginger exhibited lowest cholesterol levels. Rats fed spice incorporated diets resulted in significantly ( $p \leq 0.05$ ) lower cholesterol levels compared to HFD/HS fed rats with the exception of rats fed turmeric 2%.

#### 4. Conclusion

*In vivo* pre-diabetic model was utilized for the assessment of the effects of incorporating, garlic, turmeric, and ginger at select concentrations using high fat and sugar diets on Wistar male rats. This study explored the effects of thermally treated TGG on male wistar rats using a pre-diabetic (HFD/HS) model for 11 weeks. Unhealthy dietary and lifestyle patterns for extended periods of time may lead to the development of both pre-diabetes and Type 2 Diabetes. Our study showed that diets that incorporated TGG had lower weight gain compared to rats fed HFD/HS alone. Additionally, rats fed spice-incorporated diets showed a



**Figure 9.** Total cholesterol in wistar male rats. <sup>abc</sup>Bars with superscripts are significantly different ( $p \leq 0.05$ ). Abbreviations: HFD = High fat diet.

gradual reduction in serum glucose and cholesterol levels over time. Research [27] [28] [29] suggest that these anti-diabetic properties exhibited by spices may be attributed to the ability of spices to delay gastric emptying, reduce glucose absorption, and reducing inflammation by inactivating the pro-inflammatory factor NF $\kappa$ - $\beta$  via its I $\kappa$ - $\beta$  inhibitor. Induction of selected enzymes' (CAT, SOD, GPx, GST) serve as internal health promoting functions that combat ROS and may prevent cellular damage. Phytochemicals within spices may aid in the activation of these detoxifying enzymes. Rats fed TGG diets showed lower activities of GPx and GST along with increased levels of GSH.

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