Tamarindus indica seeds improve carbohydrate and lipid metabolism: An in vivo study

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A R T I C L E  I N F O

Article history:
Received 28 December 2016
Received in revised form 10 March 2017
Accepted 1 June 2017
Available online 6 December 2017

Keywords:
Tamarind seeds
Glucose
Glycogen
Cholesterol
Spontaneous hypertensive rats
Feeding and growth performance

A B S T R A C T

Background: The tamarind seeds have a lot of nutrients that may be used to control cholesterol or glucose levels.

Objective(s): The effects of tamarind seeds (T) on lipid and carbohydrate metabolism in rats were studied. Rats were offered basal diet (BD) with T (2%, 4% or 8%) or without T.

Materials and methods: Feeding and growth performance in rats were measured and samples of liver and blood were analyzed for glycogen content and levels of cholesterol and glucose respectively.

Results: The inclusion of T in the diet influences the feeding and growth performance in rats. The serum cholesterol level was reduced (p < 0.05) in Sprague Dawley (SD) rats fed on basal diet (BD) containing 4% and 8% T (0.24 ± 0.14 g/l and 0.31 ± 0.06 g/l respectively) compared to control (0.79 ± 0.04 g/l). The serum glucose levels in the spontaneous hypertensive rats (SHR) was lower (50.74 ± 2.50 mg/dl; p < 0.05) than control (93.52 ± 10.83 mg/dl) at 4% T. Incorporation of increasing doses of T resulted in linear increase of glycogen storage in livers of SD rats fed on BD and high sucrose diet.

Conclusion: Tamarind seeds can lower blood glucose and serum cholesterol and enhance storage of glycogen in rats.

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1. Introduction

Hypercholesterolemia is an abnormal condition characterized by high cholesterol concentration of low density lipoprotein (LDL) and a lower concentration of functional high density lipoprotein (HDL) which could lead to cardiovascular diseases due to development and increase of atheroma (atherosclerosis) [6]. It could also lead to myocardial infarction (heart attack), stroke and peripheral vascular disease [10]. Generally, the balance in LDL and HDL is genetically determined but can be affected by body build, medications, food choices and other factors [19]. It is suggested that total blood cholesterol (HDL + VLDL) of less than 200 mg/dl is desirable for the body [43]. Serum cholesterol levels of 200–239 mg/dl can be regarded as borderline while values greater than 240 mg/dl are considered high [32].

Hyperglycemia is a metabolic disorder in which the circulation of blood glucose level is excessive in the blood plasma resulting from defects in insulin secretion, insulin action or both [3]. In addition, postprandial glycaemia (related insulinaemia and lipidaemia) has been associated in chronic metabolic diseases such as type 2 diabetes mellitus and cardiovascular disease [9].

Dietary manipulation through the consumption of specific plant materials containing phytochemicals has proven to regulate the level of glucose and lipids in the blood [30]. Studies have shown that Tamarindus indica can be used to reduce visceral fat accumulation and improve hyperlipidemia and hyperglycemia in rats [28,40]. T. indica contains phenolic compounds like catenin, procyanidin B2, epicatechin, tartaric acid, mucilage, pectin, arabinose, xylose, tannins, galactose, glucose, uronic acid and triterpene [7] as well as all essential amino acids except tryptophan [1,25,36]. The tamarind seeds also have similar properties found to be important and used in traditional therapies [25]. Till date, there are intensive bioactivity studies on tamarind pulp. However, the tamarind seed which is basically considered a waste product is an under-utilized resource. Therefore, the aim of the present study is to determine in vivo effects of tamarind seeds inclusion (2%, 4% and 8%) in diets on parameters related to diabetics and hypertension such as blood glucose, glycogen in liver and cholesterol levels in three rat models: normal, hypertensive and exposed to hyperglycemic condition by...
feeding on high sucrose diet (HSD) during 4 weeks of feeding period. Additionally, evaluation of feeding and growth performance of rats was also carried out.

2. Materials and methods

2.1. Materials

2.1.1. Animal feed

Rat chow made from soybean meal was purchased from local feed manufacturer (Gold Coin Malaysia). Tamarind and soybean milk powder (Nutrisoy Inc.) were purchased from local markets.

2.1.2. Experimental animals

Male Sprague-Dawley rats (SD; n = 56) and spontaneously hypertensive rats (SHR; n = 28) out bred of Wistar-Kyoto rats, 6–8 weeks old were randomly selected from University of Malaya animal house. These animals were individually caged (600 × 380 × 200 mm³) at all time and had access to rat chow and fresh water ad libitum.

2.2. Methods

2.2.1. Preparation of ground tamarind seeds

The pulp of fresh tamarind fruits was removed and the seeds were washed out of residual flesh. A total weight of about 8 kg of seeds from the fruits was obtained and the seeds were dried in the oven (50 °C) and ground separately into fine particles of about 50 μm particle size.

2.2.2. Preparation of diets containing ground tamarind seeds

The basal diet (BD) was mixed with ground tamarind seeds at the following concentration: 2%, 4% and 8% w/w. High sucrose diet (HSD) was prepared by adding 30% w/w sucrose to the BD. Equivalent weight of soymilk was added into the diets to balance the difference in nitrogen and energy content as a result of the seeds inclusion. Proximate analysis of rat chow and fresh ground tamarind seeds is shown in Table 1. Furthermore, proximate analysis of the basal diet and high sucrose diet containing different concentration of tamarind seeds are shown in Tables 2 and 3 respectively.

2.2.3. Experimental procedures

Three different and independent experiments were conducted. In the first experiment, SD rats (n = 28) were randomly assigned into seven groups of four each and served with test diets (BD) containing tamarind seeds at various concentrations of 2, 4 and 8%, respectively where 0% served as control in the group. In the second experiment, the same procedures were carried out except that each group was fed on HSD containing 2, 4 and 8% tamarind seeds respectively and 0% tamarind seeds served for control group. In the third experiment, SHR rats were assigned into seven groups accordingly and fed with BD. Initial body weight (IBW), body weight gain (BWG), feed intake (FI) and faecal output from each rat were measured at the beginning and every seven days thereafter for 4 weeks.

The animals were sacrificed by cervical dislocation at the end of the feeding period. Blood samples were collected into heparinized tubes and after centrifugation (2500 g; 10 min 4 °C) the plasma was harvested and kept cold at 4 °C for further analysis. Liver samples of approximately 5 g were collected and stored at −20 °C for liver glycogen estimation.

2.2.4. Feeding performance analysis

1) The amount of weekly diet ingested was calculated as the difference in the total weight of feed offered at the beginning and balance at the end of the week. The weekly data collected were then used to calculate daily feed intake according to Ennouri et al. [15] with the following formula:

\[
\text{Feed intake (g/day)} = \frac{\text{Feed placed} - \text{Feed remaining}}{7 \text{ days}}
\]

2) Fecal dry matter (DM) was determined after drying faeces collected in 24 h at 105 °C to constant weight [15].

3) Macro nutrients digestibility were assessed as the difference between daily DM intake and 24 h DM excretion in faeces according to Ennouri et al. [15]:

\[
\text{Digestibility} = \frac{\text{Dietary DM intake} - \text{Faecal DM excretion}}{\text{Dietary DM intake}} \times 100
\]

4) The feed conversion efficiency (FCE) was determined by the following formula [15]:

\[
\text{FCE} = \frac{\text{Feed consumed (DM) in 28 days (g)}}{\text{Body weight gained in 28 days (g)}}
\]

5) The protein efficiency ratio (PER) is the weight gain of the growing rat divided by total protein intake during the feeding period [15] according to the following formula:

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Table 1

| Materials | Rat chow (±SE) | Tamarind seeds (±SE) |
|-----------|---------------|----------------------|
| Dry matter | 88.00 ± 0.10  | 95.10 ± 0.10         |
| Lipids | 2.20 ± 0.10   | 2.90 ± 0.10          |
| Protein | 19.10 ± 0.10  | 11.80 ± 0.11         |
| Ash | 4.60 ± 0.10   | 4.50 ± 0.00          |
| Total carbohydrates | 62.00 ± 2.03 | 75.80 ± 3.21 |

Data are presented as the mean of four observations ± standard error mean.

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Table 2

| Materials | 0% | 2% | 4% | 8% |
|-----------|----|----|----|----|
| Dry matter | 88.00 ± 0.20 | 88.00 ± 0.10 | 88.8 ± 0.20 | 88.3 ± 0.60 |
| Lipid | 2.30 ± 0.30 | 2.20 ± 0.30 | 2.20 ± 0.30 | 2.20 ± 0.30 |
| Protein | 19.10 ± 0.10 | 19.10 ± 1.10 | 19.10 ± 1.20 | 19.10 ± 1.30 |
| Ash | 4.60 ± 0.10 | 4.60 ± 0.10 | 4.60 ± 0.10 | 4.60 ± 0.01 |
| Total carbohydrates | 62.00 ± 2.00 | 62.10 ± 1.50 | 62.90 ± 2.10 | 62.40 ± 1.80 |

Data are presented as the mean of four observations ± standard error mean.

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Table 3

| Materials | 0% | 2% | 4% | 8% |
|-----------|----|----|----|----|
| Dry matter | 87.80 ± 0.20 | 87.70 ± 0.10 | 87.90 ± 0.20 | 87.80 ± 0.60 |
| Lipid | 2.20 ± 0.30 | 2.10 ± 0.30 | 2.20 ± 0.30 | 2.20 ± 0.30 |
| Protein | 19.10 ± 0.10 | 19.10 ± 1.10 | 19.10 ± 1.00 | 19.10 ± 1.40 |
| Ash | 4.60 ± 0.30 | 4.60 ± 0.10 | 4.60 ± 0.20 | 4.60 ± 0.00 |
| Total carbohydrates | 61.90 ± 2.00 | 61.90 ± 1.50 | 62.80 ± 2.10 | 62.10 ± 1.80 |

Data are presented as the mean of four observations ± standard error mean.
2.2.5. Enzymatic determination of total cholesterol

The total cholesterol was determined using commercial kits from Chemo Lab (Malaysia). The cholesterol level was evaluated by mixing thoroughly 10 μl of serum or standard solution with 1.0 ml of kit reagent. The mixture was allowed to stand for 5 min at 37 °C prior to absorbance reading at 500 nm. The cholesterol concentration was calculated using the following formula:

\[
\text{Total cholesterol} = \left( \frac{\text{Abs}_{\text{sample}}}{\text{Abs}_{\text{standard}}} \right) \times \text{standard concentration}
\]

2.2.6. Estimation of blood glucose concentration

Serum glucose level was determined according to modified Trinder method [15]. Serum glucose content was estimated by mixing 0.1 ml of serum with 1 ml of water, 1 ml of 5.0% zinc sulfate and 1 ml of 0.25 N sodium hydroxide. The mixture was then centrifuged (2500 g, 10 min) and the supernatant (1 ml) was transferred into a test tube containing 1 ml of alkaline copper reagent followed by boiling in water bath for 10 min. The mixture was cooled by placing the tubes under running water for 3 min. Arsenomolybdate reagent (1 ml) was added to the resultant solution and the volume was made up to 10 ml with water. The optical density was read at 500 nm against a blank set at zero. The glucose concentration in the samples was then calculated from a glucose calibration curve which was also run at the same time with the glucose analysis.

2.2.7. Estimation of liver glycogen content

Liver glycogen content was determined according to Vats et al. [42]. Liver samples (200 mg) were rinsed with ice-cold saline and then solubilized by incubating with 2 ml of 30% potassium hydroxide at 55 °C for 30 min. The solubilized liver tissue (0.2 ml) was placed on ice bath and then neutralized with 0.2 ml of 1 M HCl, 0.8 ml of water and 2 ml anthrone reagent (0.2 g anthrone/100 ml of 95% H2SO4). The mixture was then incubated at 100 °C for 10 min. Absorbance was measured at 620 nm and the liver glycogen content (mg glycogen/g tissue) was calculated using glucose standard curve.

2.2.8. Statistical analysis

All results presented are means of three independent measurements. Data were presented as mean ± standard error mean using one-way analysis of variance (ANOVA) by SPSS software version 16. The statistical significance was tested at \( p < 0.05 \) using post hoc Tukey’s analysis at 95% least significant difference (LSD).

3. Results

3.1. Feeding and growth performance in rats

3.1.1. Feed intake and body weight gain of SD rats fed on BD containing tamarind seeds

The control group consumed feed of 63 ± 4 g/kg BW/day during the first week of feeding and this value reduced gradually to 23 ± 10 g/kg BW/day by the end of week four (Fig. 1). The inclusion of 2% tamarind seeds into BD resulted in reduced feed intake (FI) to 38 ± 15 g/kg BW/day \( (p < 0.05) \) during week 1 of feeding whereas BD containing 4% or 8% tamarind seeds was not different from control. Intake of diets containing tamarind seeds during the next 3 weeks was not different from control except at 4% and 8% of tamarind seeds in week 2, and 2% in week 3 \( (p < 0.05; \text{Fig. 1}) \).

All treated groups had a marginal increase in BWG of between 15 and 47 g during the 4 weeks of feeding (Fig. 2). The increase in BWG was linear and similar (25–47 g) for groups consumed 0%, 2% and 4% tamarind seeds whereas group fed on 8% showed the least increase in BWG (15 g). The average BWG (g) to IBW (kg) during the entire feeding period showed that control group had the lowest BWG (57 ± 15 g/kg BW) compared to those fed on tamarind seeds (105 ± 27 \( p < 0.05 \), 69 ± 22 \( p > 0.05 \) and 71 ± 38 \( p > 0.05 \) g/kg BW) for 2%, 4% and 8% respectively (data not shown).

3.1.2. Feed intake and body weight gain of SD rats fed on HSD containing tamarind seeds

Average daily FI of treated groups fed on HSD containing 0%, 4% and 8% of tamarind seeds ranged 50–60 g/kg BW at the first week (Fig. 3). These values reduced \( (p > 0.05) \) to about 33–48 g/kg BW during the subsequent 3 weeks of feeding. Treated group fed on HSD containing 2% of tamarind seeds had lower \( (p < 0.05) \) initial FI (30 g/kg BW) that remained for the first 3 weeks of feeding before increasing to 44 g/kg BW at the 4th week (Fig. 3).

All treated groups were increased in BWG compared to IBW during the feeding (Fig. 4). The difference in IBW and final body
weight of the rats was in the range of 20%–40%. The inclusion of tamarind seeds (2% and 4%) into the diet reduced \( p < 0.05 \) average BWG. The average BWG of the control group \( (282 \pm 36 \text{ g/kg BW}; \ p < 0.05) \) was higher than treated groups \( (155 \pm 22, 118 \pm 36 \text{ and } 156 \pm 26 \text{ g/kg BW}) \) for 2%, 4%, and 8% tamarind seeds respectively (data not shown).

3.1.3. Feed intake and body weight gain of SHR fed on BD containing tamarind seeds

FI of the control group increased significantly from 42 g/kg BW to 73 g/kg BW \( (p < 0.05) \) during the first three weeks of feeding (Fig. 5). However, this value reduced \( (p < 0.05) \) to about 55 g/kg BW by week 4. The inclusion of tamarind seeds (2%, 4% and 8%) to the BD did not affect the FI compared to control (Fig. 5).

Treated group fed on BD containing 4% of tamarind seeds showed the highest BWG (Fig. 6). All rats showed an increase in BWG within 15–25% by the end of the feeding. However, increasing the inclusion of tamarind seeds to 8% into the diet reduced (\( p < 0.05 \)) average BWG \( (95 \pm 18 \text{ g/kg BW}) \) compared to control \( (134 \pm 21 \text{ g/kg BW}) \) (data not shown).

3.1.4. Effect of tamarind seeds on the digestibility of diets in rats

BD was more digestible than when sucrose was included (Fig. 7). Furthermore, BD was more digestible when consumed by SD rats than by SHR. The inclusion of tamarind seeds did not change the digestibility of BD in SD rats and SHR. However, the digestibility of HSD tended to improve \( (p < 0.05) \) with increasing inclusion of tamarind seeds, the optimum effect was achieved at 8% \( (80.46 \pm 1.23\%) \) compared to control \( (76.86 \pm 0.42\%) \).

3.1.5. Effect of tamarind seeds on feed conversion efficiency (FCE) of diets in rats

SD rats group fed on BD showed the least FCE \( (27 \pm 5) \) followed by SHR fed on BD \( (14 \pm 4) \) and SD fed on HSD \( (5.3 \pm 0.2) \) (Fig. 8). The inclusion of 2% tamarind seeds increased \( (p < 0.05) \) FCE up to 50% in SD rats fed on BD. The addition of the tamarind seeds did not
improve ($p > 0.05$) FCE in SD rats and SHR fed on HSD and BD respectively.

### 3.1.6. Effect of tamarind seeds on protein efficiency ratio (PER) of diets in rats

Inclusion of 2% tamarind seeds resulted in the highest PER ($0.54 \pm 0.20$; $p < 0.05$) in SD rats fed on BD compared to control ($0.19 \pm 0.02$; Fig. 9). SD rats fed on HSD had the highest PER ($0.90 \pm 0.03$) but the inclusion of tamarind seeds (4% and 8%) reduced ($p < 0.05$) PER to the lowest values (0.5). The inclusion of tamarind seeds in BD had no significant effect on the PER of SHR as compared to control (Fig. 9).

### 3.2. Serum cholesterol concentration of rats fed on BD and HSD containing tamarind seeds

SD rats fed on BD had blood cholesterol of $0.79 \pm 0.04$ g/l (Fig. 10). The blood cholesterol level was lowered ($p < 0.05$) with the inclusion of tamarind seeds at all doses in BD. The inclusion of different doses of tamarind seeds did not affect significantly the cholesterol level of the SD rats fed on HSD. SHR (control group) had the highest blood cholesterol ($1.19 \pm 0.05$ g/l). The addition of tamarind seeds resulted in lowering of blood cholesterol but a significant effect was only seen at 2% of tamarind seeds inclusion ($0.91 \pm 0.14$ g/l; Fig. 10).

### 3.3. Serum glucose concentration of rats fed on BD and HSD containing tamarind seeds

The lowest blood glucose was shown in SD rats fed on BD ($41.72 \pm 12.46$ mg/dl) followed by SHR fed on BD ($93.52 \pm 10.83$ mg/dl) and SD rats fed on HSD ($133.08 \pm 31.24$ mg/dl) (Fig. 11). The addition of tamarind seeds did not affect serum glucose levels except for SHR, which showed a significant reduction ($p < 0.05$) of serum glucose to $50 \pm 3$ mg/dl at 4% tamarind seeds inclusion. However, the inclusion of 4% tamarind seeds in HSD resulted in increased ($p < 0.05$) serum glucose level in SD rats ($250 \pm 44$ mg/dl) compared to control.

### 3.4. Liver glycogen content of rats fed on BD and HSD containing tamarind seeds

All control groups fed on diet with no tamarind seeds showed similar liver glycogen content ($0.95 \pm 1.27$ mg/g; Fig. 12). Incorporation of increasing doses of tamarind seeds resulted in a linear increase of glycogen storage in livers of SD rats fed on BD with
maximum level recorded at 8% tamarind seeds (3.43 ± 0.55 mg/g, \( p < 0.05 \)). There was also evidence of increased liver glycogen storage with increasing tamarind seeds inclusion for SD rats fed on HSD ranged from 2–2.5 mg/g compared to control (Fig. 12). The inclusion of tamarind seeds brought about an increased \(( p < 0.05)\) liver glycogen content at 2% and 8% (1.45 ± 0.14 and 1.80 ± 0.53 mg/g respectively) than control in SHR (Fig. 12).

4. Discussion

4.1. Effects of tamarind seeds on feed intake and body weight gain

Feed intake can be affected by diet palatability, flavor and odor \([22]\) whereas body weight gain is determined by the energy and nitrogen content of food consumed \([41]\). Previous attempts on tamarind seeds inclusion in the diet showed lower feed intake in cow \([8]\) and pig \([27]\). This has been suggested to be due to the presence of tannin \([16]\) in seed which in general could cause feed avoidance as the percentage inclusion of seed in the compound feed increased. In the present study, the inclusion of 2–8% tamarind seeds in the diets was shown not to affect the feed intake of the SD rats during the period which is in agreement with Kumar and Bhattacharya \([24]\). Nevertheless, there were some variations \((10–40\%)\) in the quantity of feed consumed \((g/kg BW)\) per day between control and treatments during the trial and this occurred possibly due to differences in adaptations of the rats to test diets \([38]\). In general these variations did not result in significant change in feed intake over the period in normal SD rats; however in SHR, the negative effect of the tamarind seeds secondary compounds (tannins) resulted in lowering voluntary feed intake after two weeks of feeding possibly by decreasing palatability of the ration because of astrigent effects of tannin on the oral cavity \([45]\).

The inclusion of tamarind seeds had no effect on body weight gain of SD rats fed on BD although there were some variations in body weight gain between the treated and control which may be attributed to the differences in animal's efficiency in converting absorbed nutrients into body mass \([29]\). Previous study showed that rats fed on high carbohydrate meal \(e.g.\) sucrose) gained more weight than rats fed on normal rat chow \([17]\) which was also seen in rats fed on HSD in the present study. However, the apparent beneficial effect of sucrose on body weight gain was lowered when tamarind seeds were included in the diet. Tannins in these feed may have acted as anti-nutritive factors since it was demonstrated that this plant's secondary metabolites lower absorption and post digestive assimilation of nutrients into body mass in animals \([5,39]\). In SHR fed on BD, the inclusion of tamarind seeds lowered the body weight gain of the rats when compared with control and this may be explained by reduced feed intake by SHR (Fig. 3).

4.2. Effects of tamarind seeds on digestibility, FCE and PER

Digestibility is described as the difference in feed intake and fecal excretion in relation to feed intake \([29]\). Thus, when consumption of feed is high and fecal excretion is also high, the value of digestibility is low i.e. feed is not properly digested and/or absorption is impaired. The values of digestibility from the present study ranged from 77 to 80% which are comparable to previous report using normal rats \([15]\) and hypercholesterolemic rats \([29]\). The addition of tamarind seeds in HSD may however provide additional beneficial effect of the phytochemical compounds of tamarind seeds.

FCE is defined as a measure of an animal’s efficiency in converting feed mass into increased body mass, which is expressed as the mass of the food eaten divided by the body mass gain over a specified period of time \([29]\). Low values of FCE implies high efficiencies and vice versa. The lowest FCE of control groups were recorded in BD-fed SD rats followed by BD-fed SHR and HSD-fed SD rats. This shows that the same diet may have different effect on rats of different physiological status and thus the same rat may perform differently on different diet \([21,38]\). The increased \(( p < 0.05)\) FCE up to 50% in SD rats fed on BD containing 2% of tamarind seeds may associated with high growth performance (Fig. 2).

PER relates the body weight gain over the protein consumed with the implication that a high PER value indicates an efficient feed as a protein source \([15]\). There was an evidence of improved PER in BD-fed SD but it occurred only at 2% of tamarind seeds (Fig. 9). In general, tamarind seeds inclusion in the diet did not have a profound effect on PER improvement. In fact, tamarind seeds inclusion may even reduce PER when the diet also contains sucrose.
4.3. Effects of tamarind seeds on serum cholesterol, glucose and liver glycogen content in rats

According to American Heart Association [4] high total cholesterol content (>2.5 g/l) indicates hypercholesterolemia while values less than 2.0 g/l is considered normal. The lowering of blood cholesterol as a result of tamarind seeds inclusion in the BD fed SD rats and SHR may be due to the presence of the phytochemicals. Tamarind seeds contain phenolic compounds such as phytosterols [11,44] in a concentration of 590 mg/kg dry weight [26]. These phytosterols, especially the beta-sitosterols are recognized to decrease plasma lipoprotein and cholesterol levels [20] via reducing the cholesterol solubility and absorption across the intestinal barrier [18,44]. In fact, phytosterols’ hydrophobicity is more readily to mix with bile salt and acid micelles [34] than can animal cholesterol [13]. This causes the excretion of a greater part of unabsorbed cholesterol particularly the low density lipoprotein with the faeces [35].

Blood sugar concentration in the body has a range of 64.8–104.4 mg/dl in a normal person [2]. Blood glucose is the primary source of energy in the body obtained from carbohydrates as a result of certain conditions like illness, stress, surgery or intake of a particular substance [11]. Phenolic compounds in plants can have blood glucose lowering capacity [12,15,31,46]. However, tamarind seeds appeared not to be a strong suppressor of blood glucose elevation in SD rats fed on HSD and BD as compared to control. This difference could be due to the physiological conditions of the animals [37]. The difference in response of these rats reflects the importance of several facets of blood glucose homeostasis.

Glycogen is stored in the liver or muscles and it functions as secondary energy storage in animals which is reconverted back to glucose by glycogenolysis in low energy state [33]. The increase of glucose intake or use additives or phenolic compounds could enhance glycogen storage [15]. In the present study, the inclusion of tamarind seeds had positive effects on glycogen storage which occurred in a dose dependent manner. The improved liver glycogen storage could be associated with the presence of phenolic compounds in tamarind seeds.

5. Conclusion

The inclusion of tamarind seeds into the diet influenced the feeding and growth performance to some extent. The inclusion of tamarind seeds (4% and 8%) in basal diet lowered the cholesterol levels of normal rats. In addition, 4% of tamarind seeds suppressed high blood glucose of SHR. Increased doses of tamarind seeds enhanced glycogen storage in the liver in all the three models of rats. Thus, tamarind seeds could be useful in treating people with hyperglycemia and/or hypercholesteremia. The present investigation may discover the promising values of tamarind seeds as a source of energy, protein as well as bioactive phytochemical for health improvement. Further studies are needed to identify the principal phenolic compounds in tamarind seeds which are responsible for lowering cholesterol and blood glucose levels in rats.

Sources of funding

IPPP University of Malaya.

Conflict of interest

None.

Acknowledgement

We are grateful to the Vice Chancellor University of Malaya, Prof Datuk Dr. Gauth Jasmon and to the IPPP University of Malaya for their dynamic support and also to the Dean faculty of science, University of Malaya Professor Datin Dr. Saadah, Abdul Rahman for his kind gestures.

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