Antidiabetic effects of *Mangifera indica* Kernel Flour-supplemented diet in streptozotocin-induced type 2 diabetes in rats

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*Food Science & Nutrition 2016; 4(6): 828–839*  
doi: 10.1002/fsn3.348

**Abstract**

Our previous report showed that *Mangifera indica* kernel flour (MIKF) is a rich source of pharmacologically important flavonoids and phenolic acids; and that its methanolic extract inhibits some key enzymes linked to the pathology and complications of type 2 diabetes (T2D) in vitro. Hence, this study evaluated the antidiabetic effects of 10% and 20% MIKF-supplemented diets in T2D in rats. T2D was induced in rats using a high-fat diet (HFD), low-dose streptozotocin (HFD/STZ) model, by feeding the rats with HFD for 2 weeks followed by single dose administration of STZ (40 mg/kg body weight, intraperitoneally). The diabetic rats were later fed the MIKF-supplemented diets, or administered with metformin (25 mg/kg b.w.) for 21 days; the control rats were fed basal diet during this period. Intake of the MIKF-supplemented diets resulted in significant (*P* < 0.05) improvement in the fasting blood glucose, hepatic glycogen, glycosylated hemoglobin, lipid profile, plasma electrolytes, hepatic and pancreatic malonaldehyde, and the liver function markers of the diabetic rats, compared with the diabetic control rats. The ameliorative effect of 20% MIKF-supplemented was comparable (*P* > 0.05) with that of metformin administration in the diabetic rats. It is concluded that *M. indica* kernel flour has antidiabetic effects in T2D rats, and could therefore be a promising nutraceutical therapy for the management of T2D and its associated complications.

**Introduction**

According to World Health Organisation (WHO), about 350 million people have diabetes globally, compared to 153 million in 1980, and 90% of these cases are Type 2 diabetes (T2D) (Danaei et al. 2011). This translates to about 56% increase in global diabetes prevalence from 1980 to date. With this current global prevalence and a projection that 439 million people will have T2D by 2030 (International Diabetes Federation 2009), the disease is rising at an alarming rate and may assume an epidemic proportion in the near future, if research is not intensified more on preventive than curative strategies. The challenge posed by this dreadful prevalence rate of T2D is further compounded by the complexity and complications of the disease. Hence, despite advances in modern medicine and a huge capital investment, T2D remains a major global health problem, as it has no satisfactory therapy in terms of safety and efficacy. As a metabolic disorder characterized by chronic hyperglycemia, resulting from insulin resistance and loss of pancreas ß-cell function (Laakso 2001), T2D is often associated with dyslipidemia (Ahmed and Goldstein 2006).

It is heart-warming to know that the incidence of T2D can be reduced by half among individuals at high risk through lifestyle and pharmacologic interventions (Li et al. 2008). But incidentally, the currently available antidiabetic drugs for the treatment of T2D have certain drawbacks ranging from adverse side effects and development of resistance to lack of responsiveness in many diabetic patients, in addition to their inability to effectively control the hyperlipidemia that is frequently associated with the
disease (Zia et al. 2001). Interestingly, the use of dietary therapy offers hope for a more effective and safer preventative and management strategy for T2D. This has been proven by epidemiological studies that have consistently shown an inverse association between the risk of chronic human diseases such as T2D and the consumption of polyphenolic rich diet (Arts and Hollman 2005).

Our recent study showed that *M. indica* kernel flour is a rich source of pharmacologically important flavonoids (catechin, rutin, quercitrin, quercetin, and kaempferol), and phenolic acids (gallic acid, caffeic, chlorogenic, and ellagic acid) (Irondi et al. 2014). In the same study, we demonstrated that methanol extract of MIKF inhibits some key enzymes (α-amylase, α-glucosidase, and aldose reductase) linked to the pathology and complications of type 2 diabetes (T2D) in vitro. *Mangifera indica* L. (mango) is an important economic tree crop, ranking second as internationally traded tropical fruits, and fifth in total world production among major fruit crops (Food Agriculture Organization FAO 2010). It serves as food and as an important medicinal plant. The fruit which is commonly eaten by man is a rich source of vitamin A, and also contains vitamins B and C; the mesocarp is typically composed of water (84%), sugar (15%), protein (0.5%), fibers and skin (0.5%). The seed kernels contain carbohydrate (80%), fat (10%), and protein (6%) (Purseglove 1991), and are usually discarded as a waste product after the fruit is eaten. However, in India they are consumed by humans during times of food scarcity (Kumar et al. 1991; Purseglove 1991). In the South-Eastern part of Nigeria, they are used for thickening of traditional soups (Eddy and Udoh 2005). Different parts of the *M. indica* plant are used in folk medicine for a wide variety of remedies (Garrido et al. 2004). The extracts have been reported to have several pharmacological activities such as antioxidant (Ribeiro et al. 2008; Maisuthisakul and Gordon 2009), anti-inflammatory (Garrido et al. 2004), antidiabetic (Jain et al. 2014), and immune-modulatory activities (Garcia et al. 2003). This study was therefore designed to evaluate the antidiabetic effects of 10% and 20% *M. indica* kernel flour-supplemented diet in high-fat, low-dose streptozotocin-induced T2D in rats.

### Materials and Methods

#### Sample collection and preparation of kernel flour

Samples of fresh ripe *M. indica* seeds (Figure 1) were purchased in a local farm settlement in Ibadan, Oyo State, Nigeria. The seeds were authenticated at the Department of Botany, University of Ibadan, Nigeria. Subsequently, the seeds were sorted and the kernels were manually removed from the endocarp. The kernel flour was prepared by chopping the fresh kernels were with a stainless kitchen knife and sun-drying for 4 days. Subsequently, the kernels were ground into fine particle size (0.5 mm) using a laboratory grinder to obtain the MIKF. The MIKF was packed in an airtight container and stored at 4°C until analysis.

All the chemicals used for analysis were of analytical grade.

**Determination of proximate composition of MIKF**

The proximate composition of the MIKF was determined according to Association of Official Analytical Chemists, AOAC (2005) methods. Moisture content was determined by oven-drying at 100–105°C for 18–24 h. Total Nitrogen content (N) was determined by Kjeldahl method, and the protein content was calculated as $N \times 6.25$. Ash content was determined by incinerating 2 g of the MIKF in a preweighed porcelain crucible in a muffle furnace at 600°C for 6 h. Crude fat content of the MIKF was determined using a Soxtec extraction machine. Total carbohydrate was estimated by subtracting the sum of ash, protein, moisture, and fat, from 100 (i.e., $100 - [\% \text{ ash} + \% \text{ protein} + \% \text{ moisture} + \% \text{ fat}]$).

**Formulation of experimental diets**

The formulation of high-fat diet (HFD) containing 40% fat (lard) and 18% protein (casein) and 41% carbohydrate (corn starch) as the percentage of total calorie, was based on Reed et al. (2000). Then the HFD was supplemented with 10% and 20% of the MIKF at the expense of corn starch, to give the *M. indica* kernel flour-supplemented diets (10% MIKF and 20% MIKF diets, respectively). The composition of the experimental diets is shown in Table 1.

| Diet          | Casein (g) | Corn oil (g) | Mineral-vitamin premix (g) | Lard (g) | MIKF (g) | Corn starch (g) |
|---------------|------------|--------------|---------------------------|----------|----------|-----------------|
| Basal         | 19.15      | 10.00        | 4.00                      | –        | –        | 66.85           |
| HFD           | 19.15      | –            | 4.00                      | 40.00    | –        | 36.85           |
| 10% MIKF      | 18.66      | –            | 4.00                      | 38.63    | 10.00    | 28.71           |
| 20% MIKF      | 18.17      | –            | 4.00                      | 37.26    | 20.00    | 20.57           |

MIKF, *Mangifera indica* kernel flour; HFD, high-fat diet 10% MIKF diet supplemented with 10% *M. indica* kernel flour; 20% MIKF, diet supplemented with 20% *M. indica* kernel flour. Casein = 94% protein; lard = 99.9% fat.
Experimental animals and induction of T2D using the HFD/STZ model

Adult male Wistar rats weighing 180–200 g were used for this study. The rats were procured from the experimental animal breeding unit of Department of Veterinary Medicine, University of Ibadan, Nigeria, and were handled in accordance with the Guide for the Care and Use of laboratory animals of our institution. The rats were housed in experimental cages under a 12 h light-dark cycle at an ambient temperature of 25 ± 2°C, to acclimatize for 7 days. During the acclimatization period, the rats were fed rodent chow and water ad libitum. To induce T2D, the rats (excluding the normal control rats) were fed the HFD for 2 weeks (Reed et al. 2000), after which they were injected with single dose streptozotocin (STZ) (40 mg/kg b.w., intraperitoneally, in citrate buffer; pH 4.5) (Parveen et al. 2011); the normal control rats were fed basal diet. The development of hyperglycemia in the rats was confirmed by testing for fasting blood glucose (FBG) 72 h after the STZ injection. The rats that maintained FBG higher than 200 mg/dL were considered diabetic, and were selected for the study.

Animal Groups and experimental design

The rats were divided into five groups of five animals each as follows: group I (normal control), fed basal diet throughout the experiment; group II (diabetic control), fed HFD throughout the experiment; group III (DMT), diabetic rats administered metformin (25 mg/kg b.w., intraperitoneally, in citrate buffer; pH 4.5) (Parveen et al. 2011); the normal control rats were fed basal diet. The development of hyperglycemia in the rats was confirmed by testing for fasting blood glucose (FBG) 72 h after the STZ injection. The rats that maintained FBG higher than 200 mg/dL were considered diabetic, and were selected for the study.

Tissue preparation

The liver and pancreas tissues of the rats were excised immediately after rats were killed, and were perfused with ice-cold saline. They were later homogenized at 4°C with 10 times w/v 0.1 mol/L phosphate-buffer, pH 7.4, in a polytron homogenizer. The homogenate was centrifuged at 800 rpm for 5 min at 4°C to separate the nuclear debris, and an aliquot of the supernatant was used for the determination of malonaldehyde (MDA) level.

Biochemical assays

Fasting blood glucose level was tested using a portable glucometer (Accu Check Active), and the percentage change in FBG level of the rats during the treatment period was calculated. Liver glycogen was determined by the method reported by Ong and Khoo (2000), using anthrone reagent. Glycosylated hemoglobin (HbA1c) level was determined according to the method of Standefer and Eaton (1983). Plasma lipid profile, namely, triglycerides (TRIG), total cholesterol (TC), and high-density lipoprotein (HDL) levels were estimated using commercially available standard kits (Randox Laboratories, Ltd., Crumlin, Co. Antrim, UK), according to the manufacturer’s instruction. Low-density lipoproteins (LDL) and very low-density lipoproteins (VLDL) levels were calculated using the Friedewald formula (Friedewald et al. 1972), as follows:

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LDL = TC - (HDL + VLDL) \\
VLDL = TRIG/5
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Hepatic and pancreatic malonaldehyde level as an index of lipid peroxidation was determined according to the method described by Ohkawa et al. (1979). The level of plasma electrolytes (Mg, Zn, and Cu) quantified according to the method reported by Clegg et al. (1981), using atomic absorption spectrometer. Liver function enzymes [plasma aspartate aminotransferase (AST), alanine aminotransferase (ALT), and alkaline phosphatase (ALP)] activities were determined using Randox laboratory kits. Total protein concentration of the plasma and tissues was determined according to the method of Lowry et al. (1951).

Statistical analysis

Results of replicate experiments were expressed as mean ± standard deviation (SD). Analysis of variance (ANOVA) and least significance difference (LSD) were carried out on the result data at 95% confidence level using SPSS statistical software package, version 17 (SPSS Inc., Chicago).

Results

The proximate composition of the MIKF is presented in Table 2. The result showed that the MIKF was high in carbohydrate (72.73%), but low in protein (4.59%) and ash (1.69%) contents.

The average food intake of the rats is presented in Table 3. The results showed that there was no significant (P > 0.05) difference in the daily food intake of the various treatment groups.
Table 2. Proximate composition of the *M. indica* kernel flour (MIKF).

| Nutrients         | MIKF                  |
|-------------------|-----------------------|
| Moisture (%)      | 7.31 ± 0.04           |
| Protein (%)       | 4.59 ± 0.47           |
| Fat (%)           | 13.68 ± 0.01          |
| Ash (%)           | 1.69 ± 0.03           |
| Total carbohydrate (%) | 72.73 ± 0.45         |

Values represent mean ± standard deviation of replicate determinations (n = 3).

Table 3. Average food intake of rats fed *M. indica* kernel flour-supplemented diets.

| Treatment groups | Average food intake (g/rat/day) |
|------------------|--------------------------------|
| Normal control   | 5.52 ± 0.46a                  |
| Diabetic control | 6.01 ± 0.47a                  |
| DMT              | 5.68 ± 0.42a                  |
| D10%MIKF         | 5.88 ± 0.8a                   |
| D20%MIKF         | 5.94 ± 0.51a                  |

Values represent mean ± standard deviation of replicate determinations (n = 5). Values with the same superscript letters along the column are not significantly (P > 0.05) different. DMT, diabetic rats treated with metformin; D10% MIKF, diabetic rats fed 10% *M. indica* kernel flour-supplemented high fat diet (HFD); D20% MIKF, diabetic rats fed 20% *M. indica* kernel flour-supplemented HFD.

Table 4. Changes in average body weights (g) of type 2 diabetic rats fed soup thickener flour-supplemented diets.

| Rat group        | Initial weight | Final weight | Weight gain/loss (%) |
|------------------|----------------|--------------|----------------------|
| Normal control   | 200.88 ± 14.09 | 209.48 ± 14.12 | 4.28a                |
| Diabetic control | 200.62 ± 16.56 | 194.1 ± 17.02 | −3.25b               |
| DMT              | 197.12 ± 14.11 | 200.04 ± 14.25 | 1.48c                |
| D10%MIKF         | 204.04 ± 12.29 | 207.74 ± 12.35 | 1.81b                |
| D20%MIKF         | 202.88 ± 14.13 | 207.12 ± 13.86 | 2.09b                |

Values represent mean ± standard deviation of replicate determinations (n = 5). Values with different superscript letters along the column differ significantly (P < 0.05) different. DMT, diabetic rats treated with metformin; D10%MIKF, diabetic rats fed 10% *M. indica* kernel flour-supplemented high fat diet (HFD); D20%MIKF, diabetic rats fed 20% *M. indica* kernel flour-supplemented HFD.

Table 5. Changes in fasting blood glucose (FBG) level (mg/dL) of rats during the dietary intervention.

| Treatment group | Day 0   | Day 7   | Day 14  | Day 21  | % change |
|-----------------|---------|---------|---------|---------|----------|
| Normal control  | 90.6 ± 5.22a | 91.8 ± 4.97a | 91.4 ± 5.46a | 91.6 ± 4.67a | +1.10 (Increase) |
| Diabetic control| 378 ± 7.84a  | 383.6 ± 8.29a | 392.2 ± 7.40ab | 394.6 ± 7.73b | +4.39 (Increase) |
| DMT             | 383 ± 20.29a | 203.2 ± 16.42b | 177.2 ± 9.09c  | 146.8 ± 8.17d | −61.67 (Decrease) |
| D10%MIKF        | 353.2 ± 18.45a | 234.6 ± 9.42b  | 181.2 ± 6.22c  | 141.2 ± 7.89d | −60.02 (Decrease) |
| D20%MIKF        | 356.6 ± 15.52a | 223 ± 9.38b   | 163.8 ± 5.93c  | 125.00 ± 6.82d | −64.95 (Decrease) |

Values with different superscript letters in the same row differ significantly (P < 0.05). % change = (FBG at Day 21 – FBG at Day 0) × 100/FBG at Day 0. DMT, diabetic rats treated with metformin; D10%MIKF, diabetic rats fed 10% *M. indica* kernel flour-supplemented high fat diet (HFD); D20%MIKF, diabetic rats fed 20% *M. indica* kernel flour-supplemented HFD.
control group, the diabetic rats had significant ($P < 0.05$) increase in their plasma total cholesterol, triglycerides, LDL, and VLDL; with a concomitant significant ($P < 0.05$) decrease in their plasma HDL concentration. However, in comparison with the diabetic control group, the diabetic groups fed MIKF-supplemented diets had significantly ($P < 0.05$) lower levels of plasma total cholesterol, triglycerides, LDL, and VLDL; with an attendant significant ($P < 0.05$) increase in HDL. The 20% MIKF-supplemented diet was more effective in improving the lipid profile of the diabetic rats than the 10% MIKF-supplemented diet and metformin.

The levels of peroxidation of the liver and the pancreas of the rats, reported as MDA concentration are presented in Figure 4A and B. The diabetic groups had significantly ($P < 0.05$) higher MDA levels than the normal control group. The MDA levels of the diabetic groups fed 20% MIKF-supplemented diets had significantly ($P < 0.05$) lower levels of plasma total cholesterol, triglycerides, LDL, and VLDL; with an attendant significant ($P < 0.05$) increase in HDL. The 20% MIKF-supplemented diet was more effective in improving the lipid profile of the diabetic rats than the 10% MIKF-supplemented diet and metformin.

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The plasma electrolytes [magnesium (Mg), zinc (Zn), and copper (Cu)] concentrations of the rats are presented in Table 6. Mg and Zn decreased significantly ($P < 0.05$) in the diabetic groups compared with the normal control group. However, the plasma Mg and Zn levels of the diabetic groups fed the MIKF-supplemented diets were significantly ($P < 0.05$) higher than those of the diabetic control group; indicating that the diets were able to improve the plasma Mg and Zn levels of the rats. Conversely, the diabetic groups had significantly ($P < 0.05$) higher plasma Cu level than the normal control group, with the diabetic control group having the highest level.

The alanine aminotransferase, AST and ALP activities (Table 7) were consistently significantly ($P < 0.05$) higher in the diabetic groups than in the normal control group. When compared with the diabetic control group, there was significant ($P < 0.05$) decrease in the activities of these three enzymes in the diabetic groups fed MIKF-supplemented diets; suggesting that intake of the diets attenuated the level of the hepatocellular damage that might have been caused by STZ administration in the diabetic rats. Relative to the normal control group, there was significant ($P < 0.05$) reduction in the protein level of the diabetic groups. However, intake of the MIKF-supplemented diets resulted in a significant ($P < 0.05$)
Figure 3. Effect of MIKF-supplemented diets on the lipid profile of the rats. (A) total cholesterol; (B) triglycerides; (C) high-density lipoprotein-cholesterol (HDL); (D) low-density lipoprotein-cholesterol (LDL-C); (E) very low-density lipoprotein-cholesterol (VLDL). Bars with different letters differ significantly ($P < 0.05$). DMT, diabetic rats treated with metformin; D10%MIKF, diabetic rats fed 10% *M. indica* kernel flour-supplemented high fat diet (HFD); D20%MIKF, diabetic rats fed 20% *M. indica* kernel flour-supplemented HFD.
increase in the plasma total protein levels of the diabetic groups when compared with the diabetic control group.

**Discussion**

In this study, we evaluated the antidiabetic activity of *M. indica* kernel flour (MIKF)-supplemented diets in T2D rats, having reported in a recent study that MIKF is a rich source of pharmacologically important flavonoids and phenolic acids, and that its methanol extract inhibits some key enzymes (α-amylase, α-glucosidase, and aldose reductase) linked to the pathology and complications of T2D in vitro (Irondi et al. 2014). In view of the adverse side effects and high cost of synthetic antidiabetic drugs, research emphasis has continued to increase on alternative ways of preventing and managing T2D that could be more effective, safer and affordable. Dietary intervention has proven to be one of such ways (Oboh et al. 2010), as the consumption of polyphenol-rich diets has been reported to protect against development of degenerative diseases including diabetes (Arts and Hollman 2005).

The use of HFD/STZ model to induce T2D in rats has been reported by previous studies (Reed et al. 2000; Parveen et al. 2011). In this model, the HFD leads to hyperinsulinemia, insulin resistance and/or glucose intolerance, and the subsequent STZ administration, a β-cell toxin, results in a severe reduction in functional β-cell mass (Szkudelski 2001; Lenzen 2008). The STZ, a known diabetogen (Marianna et al. 2006), then brings about the destruction of β-cells of the islets of Langerhans, thereby leading to a large-scale reduction in insulin release. Consequently, the deficiency of insulin causes high glucose level in the blood (hyperglycemia) (Grover et al. 2002)
Table 7. Effect of *M. indica* flour-supplemented diet on liver function markers of the rats.

| Treatment group          | Alanine aminotransferase (U/L) | Aspartate aminotransferase (U/L) | Alkaline phosphatase (U/L) | Protein (g/dL) |
|--------------------------|---------------------------------|---------------------------------|---------------------------|---------------|
| Normal control           | 23.42 ± 1.01<sup>a</sup>        | 55.86 ± 1.47<sup>a</sup>        | 76.91 ± 2.06<sup>d</sup>  | 7.87 ± 0.13<sup>d</sup> |
| Diabetic control         | 51.25 ± 1.65<sup>a</sup>        | 100.21 ± 2.12<sup>a</sup>       | 146.88 ± 3.34<sup>a</sup> | 3.21 ± 0.11<sup>d</sup> |
| DMT                      | 30.51 ± 1.16<sup>c</sup>        | 63.12 ± 1.26<sup>c</sup>        | 84.14 ± 1.83<sup>c</sup>  | 5.03 ± 0.21<sup>b</sup> |
| D10%MIKF                 | 33.60 ± 1.91<sup>b</sup>        | 66.28 ± 1.91<sup>b</sup>        | 87.23 ± 2.32<sup>b</sup>  | 5.15 ± 0.16<sup>c</sup> |
| D20%MIKF                 | 27.98 ± 1.46<sup>d</sup>        | 60.60 ± 1.57<sup>d</sup>        | 81.68 ± 2.20<sup>c</sup>  | 5.35 ± 0.22<sup>b</sup> |

Values represent mean ± standard deviation of replicate determinations (*n* = 5). Values with different superscript letters along the column differ significantly (*P* < 0.05). DMT, diabetic rats treated with metformin; D10%MIKF, diabetic rats fed 10% *M. indica* kernel flour-supplemented diet; D20%MIKF, diabetic rats fed 20% *M. indica* kernel flour-supplemented HFD.

and other metabolic aberrations associated with T2D such as increased cholesterol (Kaur et al. 2011). Usually, hypoinsulinemia is characterized by a reduction in glucose utilization by the tissues, and excessive hepatic glycogenolysis and gluconeogenesis, which all constitute the fundamental mechanism underlying hyperglycemia in DM (Jayasri et al. 2008). Thus, diabetic rats induced using this model show hyperglycemia, hyperlipidemia, and insulin resistance that are similar to the natural history and metabolic characteristics of T2D in human (Srinivasan et al. 2005).

The levels of the proximate parameters of the MIKF (Table 2) are in agreement with the range (moisture: 10.48%; ash: 2.20%; fat: 8.30%, protein: 4.49%; carbohydrate: 72.52%) earlier reported by Eddy and Udoh (2005).

The decrease in the body weight of the diabetic control rats may be due to dehydration and increased muscle wasting (Ene et al. 2007); and ongoing gluconeogenesis, in which fats and proteins are catabolized (Prabhu et al. 2008).

The ameliorative effect of MIKF-supplemented diets on the hyperglycemic conditions of the diabetic rats observed in this study could be attributed to the flavonoids and phenolic acids present in the flour as earlier reported (Irondi et al. 2014). These phenolic compounds are known to exert their antidiabetic activity through several mechanisms including the inhibition of carbohydrate-hydrolyzing enzymes (Iwai et al. 2006). The increase in the FBG levels of the diabetic rats was accompanied by a reduction in their hepatic glycogen concentration, relative to the normal control group. Under normal metabolic and physiological conditions, insulin enhances intracellular glycogen deposition by stimulating activities of glycogen synthase and inhibiting glycogen phosphorylase (Shivanna et al. 2013). This regulatory function of insulin is compromised under the hypoinsulinemic condition that is caused by the diabetogenic activity of STZ; this may account for the depleted hepatic glycogen level in the diabetic rats observed in this study. This finding is in conformity with that of Ahmed et al. (2010), who reported a reduction in hepatic glycogen level in STZ-induced diabetic rats. Interestingly, this aberration was mitigated in the diabetic rats fed MIKF-supplemented diets, suggesting that there may have been increased insulin release; increased synthesis of glycogen synthase and prevention of its inactivation (Selvan et al. 2008), due to the MIKF. Other researchers have also reported the antidiabetic activity of different parts of *M. indica*. For instance, Murugananandan et al. (2005) reported that the chronic administration of mangiferin at 10 and 20 mg/kg doses, once daily for 28 days, lowered the fasting plasma glucose level of STZ-induced diabetic rats. Similarly, various extracts of the stem-bark were reported to possess antihyperglycemic effect in diabetic rats (Ojewole 2005; Bhowmik et al. 2009). On the other hand, the decrease in the FBG level of the diabetic group treated with metformin may be attributed to the antihyperglycemic activities of metformin including increase in insulin sensitivity, enhancement of peripheral glucose uptake, suppression of glucose production by the liver, increase in fatty acid oxidation, and decrease in absorption of glucose from the gastrointestinal tract (Kirpichnikov et al. 2002; Collier et al. 2006).

Glycosylated hemoglobin (HbA1c) is a routine index of T2D, being a biomarker for identifying stress hyperglycemia and diabetic hyperglycemia (Kundu et al. 2013). The glycosylation of hemoglobin occurs progressively and irreversibly during the 90–120 days life span of the erythrocytes (Saudek et al. 2005). It is therefore a more comprehensive measure of total glycemic exposure than FBG due to the representation of blood glucose level in both the postprandial and fasting state (Rohlfing et al. 2000). The increase in HbA1c observed in the diabetic rats relative to the normal control rats may be attributed to the higher rate of glycation of hemoglobin under the hyperglycemic condition occasioned by HFD/STZ administration. This is in line with the reports of cooccurrence of diabetic hyperglycemia with elevated HbA1c by other researchers (Bernadette et al. 2008). However, intake of the MIKF-supplemented diets led to a reduction in the HbA1c levels of the diabetic rats, when compared with
the diabetic control group; this further gave credence to the antihyperglycemic effect of the MIKF.

Dyslipidemia is a common biochemical feature of T2D due to mostly insulin resistance and insulin deficiency (Chahil and Ginsberg 2006), and it is one of the major risk factors for cardiovascular disease (CVD) in DM (Chehade et al. 2013). Abnormalities in lipid metabolism resulting from increased free fatty acid release from insulin-resistant fat cells leads to alterations in plasma lipid profile in diabetes (Chehade et al. 2013). Under normal metabolic conditions, insulin modulates lipid metabolism by activating lipoprotein lipase, which hydrolyzes triglycerides to release fatty acids and glycerol. Then the fatty acids are oxidized for fuel or reesterified for storage in body tissues. Contrarily, the lipoprotein lipase is inactivated under a condition of insulin deficiency and/or resistance, and this leads to hypertriglyceridemia. The alterations in the plasma lipid profile (elevated triglycerides, total cholesterol, LDL and VLDL; and reduced HDL) of the diabetic rats in this study are in agreement with the alterations in lipid profiles of HFD-STZ-induced diabetic rats reported by other researchers (Zhang et al. 2010). Increased LDL level enhances the deposition of cholesterol in the arteries and aorta, and consequently the development of coronary heart disease in diabetic patients, since LDL helps to transport cholesterol from the liver to body tissues (Pedersen 2001). HDL, on the other hand, helps to transport endogenous cholesterol and cholesteryl esters from other body tissues to the liver and steroidalogenic tissues where they are metabolized and excreted. Thus, HDL is regarded as a beneficial lipoprotein that prevents cholesterol deposition in the system, thereby preventing atherosclerosis (Xu et al. 2005). The observed decrease in the plasma triglyceride, total cholesterol, LDL, and VLDL; and a concomitant increase in the HDL levels of diabetic rats fed MIKF-supplemented diets may be due to improved insulin secretion by the islet cells of the pancreas.

Lipid peroxidation is an oxidative deterioration of polyunsaturated lipids, involving reactive oxygen species and transition metal ions, which yields diverse cytotoxic products, most of which are aldehydes, such as MDA (Shalaby and Shanab 2013). MDA and other products resulting from the peroxidation polyunsaturated fatty acids, can disrupt the function of the membrane, increase tissue permeability, and inactivate membrane-bound receptors and enzymes (Rahman 2003). Excessive lipid peroxidation will in turn cause elevated generation of free radicals, which are harmful to the body cells (Perez-Gutierrez and Mota Flores 2010). The increased MDA levels of the liver and pancreas of the diabetic rats, when compared with that of the normal control group, indicate that there was oxidative damage (peroxidation) to the lipids of these two key tissues involved in glucose metabolism; and this could be attributed to the hyperlipidemic state of the diabetic rats (Annadurai et al. 2014). However, this was ameliorated in the diabetic rats fed MIKF-supplemented diets. This observation suggests that the MIKF-supplemented diets might have the ability to protect the pancreas and liver from oxidative damage.

The reduced plasma concentrations of Mg and Zn, and increased Cu concentration observed in the diabetic groups, relative to the normal control group, is consistent with the findings of Viktorinová et al. (2009), who reported lower levels of plasma Mg and Zn, but higher Cu level in diabetic patients than in normal patients. Lower Zn (Masood et al. 2009) and lower Mg (Diwan et al. 2006) have also been reported in diabetic state in comparison with control subjects. It is well-known that Zn and Mg enhance insulin function and promote glucose entry into the cells. Masood et al. (2009) stated that the hypersecretion of insulin associated with obese T2D patients could cause hyperzincuria; hence, patients with T2D are more likely to have suboptimal Zn status. Hypomagnesemia in diabetes is attributed to osmotic renal losses from glycosuria, decreased intestinal absorption, and redistribution of Mg from the plasma into blood cells caused by insulin effect (Wälti et al. 2003; Guerrero-Romero and Rodriguez-Morán 2006). STZ-induced diabetes has been reported to increase Cu content associated with the metallothionein of rat liver and kidney tissues (Orhan et al. 2011).

Elevated blood levels of ALT, AST, and ALP indicate liver damage by toxicants or disease condition (Singh et al. 2001), including T2D (Idris et al. 2011). The increase in the activities of ALT, AST, and ALP in the diabetic rats relative to the normal control rats is consistent with the findings of Nwanjo (2007), who also reported an elevation in the activities of liver function enzymes in diabetic rats; and this may be due to necrosis of the liver, and the consequent leakage of these enzymes from the liver into circulation (Mansour et al. 2002). The decrease in the activities of ALT, AST, and ALP in the diabetic rats fed MIKF-supplemented diets, in comparison with the diabetic control group indicates a possible hepatoprotective effect of the MIKF (Iwalokun et al. 2006).

The reduced plasma protein levels of the diabetic rats, in comparison with that of the normal control rats could be attributed to decreased protein synthesis coupled with increased muscle proteolysis (Gray and Cooper 2011). Insulin regulates protein metabolism by stimulating protein synthesis and retarding protein degradation (Murray et al. 2000). Thus, diabetic hypoinsulinemia results in decreased protein synthesis in all tissues due to decreased production of alkaline phosphatase in absolute or relative deficiency of insulin (Murugan and Pari 2007). The decreased plasma protein was, however, restored toward normal in diabetic rats fed MIKF-supplemented diets,
probably as a result of the potential of the MIKF to improve the secretion, sensitivity and action of insulin in the diabetic rats.

The overall antidiabetic activity of the MIKF may be attributed to its polyphenols, including the flavonoids and phenolic acids, as earlier reported (Irondi et al. 2014). In addition to modulating carbohydrate metabolism by inhibiting α-glucosidase and α-amylase, the polyphenols are known to exhibit antidiabetic effect through other mechanisms such as reduction in intestinal absorption of dietary carbohydrate; improvement of β-cell function and insulin action; stimulation of insulin secretion; and antioxidant and anti-inflammatory properties (Cabrera et al. 2006; Iwai et al. 2006).

Conclusion

The M. indica kernel flour-supplemented diets improved the levels of fasting blood glucose, hepatic glycogen, glycosylated hemoglobin, lipid profile, plasma electrolytes, malonaldehyde, and the liver function biomarkers of the diabetic rats. M. indica kernel flour could therefore be a promising nutraceutical therapy for the management of T2D and its associated complications.

Conflict of Interest

None declared.

References

Ahmed, O. M., A. Abdel-Moneim, I. Abulyazid, and A. M. Mahmoud. 2010. Antihyperglycemic, anti-hyperlipidemic and antioxidant effects and the probable mechanisms of action of Ruta graveolens and rutin in nicotinamide/streptozotocin diabetic albino rats. Diabeto. Croat. 39:15–32.

Annadurai, T., A. Vasanthakumar, P. Geraldine, and P. A. Thomas. 2014. Variations in erythrocyte antioxidant levels and lipid peroxidation status and in serum lipid profile parameters in relation to blood haemoglobin A1c values in individuals with type 2 diabetes mellitus. Diabetes Res. Clin. Pract. 105:58–69.

Arts, I. C. W., and P. C. H. Hollman. 2005. Polyphenols and disease risk in epidemiologic studies. Am. J. Clin. Nutr. 81:317–325.

Association of Official Analytical Chemists, AOAC. 2005. Official methods of analysis of association of analytical chemists international. Pp.152–154 252 in Horwitz, W. ed. 18th ed. AOAC International, Washington D.C.

Bernadette, B. A., C. Sam, J. Chong, C. U. Culing, W. Clinton, R. Tatjana, S. V. Mitchell, et al. 2008. Diabetes, fasting glucose levels and risk ischaemic and vascular events. Diab. Care 31:1132–1137.

Bhowmik, A., L. A. Khan, M. Akhter, and B. Rokeya. 2009. Studies on the antidiabetic effects of Mangifera indica stem-barks and leaves on nondiabetic, type 1 and type 2 diabetic model rats. Bangladesh J Pharmacol. 4:110–114.

Cabrera, C., R. Artacho, and R. Giménez. 2006. Beneficial effects of green tea - a review. J. Am. Coll. Nutr. 25:79–99.

Chahil, T. J., and H. N. Ginsberg. 2006. Diabetic dyslipidaemia. Endocrinol. Metab. Clin. North Am. 35:491–510.

Chehade, J. M., M. Gladysz, and A. D. Mooradian. 2013. Dyslipidaemia in type 2 diabetes: prevalence, pathophysiology, and management. Drugs 73:327–339.

Clegg, M. S., C. L. Keen, B. Lonnerdal, and L. S. Hurley. 1981. Influence of ashing techniques on the analysis of trace elements in animal tissue. Biol. Trace Elem. Res. 3:107–115.

Collier, C. A., C. R. Bruce, A. C. Smith, G. Lopaschuk, and D. J. Dyck. 2006. Metformin counters the insulin-induced suppression of fatty acid oxidation and stimulation of triacylglycerol storage in rodent skeletal muscle. Am. J. Physiol. Endocrinol. Metab. 291:E182–E189.

Danaei, G., M. Finucane, Y. Lu, G. M. Singh, M. J. Cowan, C. J. Paciorek, et al. 2011. National, regional, and global trends in fasting plasma glucose and diabetes prevalence since 1980: systematic analysis of health examination surveys and epidemiological studies with 370 country-years and 2.7 million participants. Lancet 378:31–40.

Diwan, A. G., A. B. Pradhan, D. Lingojiwar, K. K. Krishna, P. Singh, and S. I. Almelkar. 2006. Serum zinc, chromium and magnesium levels in type-2 diabetes. Int. J. Diab. Dev. Ctries. 26:122–123.

Eddy, N. O., and C. L. Udoh. 2005. Proximate evaluation of nutritional value of some soup thickeners. ChemClass J. 2:12–14.

Ene, A. C., E. A. Nwankwo, and L. M. Samdi. 2007. Alloxan-induced diabetes in arts and the effects of black caraway (Caramcarvi L.) oil on their body weight. ChemClass J. 2:12–14.

Estimation of the concentration of low- density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. Clin. Chem. 18:499–502.

Food Agriculture Organization FAO. 2010. Statistical database. Rome, Italy. http://faostat.fao.org (accessed 15 December 2011).

Friedewald, W. T., R. I. Levy, and D. S. Fredrickson. 1972. Estimation of the concentration of low-density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. Clin. Chem. 18:499–502.

García, D., J. Leiro, R. Delgado, et al. 2003. Mango L. extract (Vimang) and mangiferin modulate mouse humoral immune response. Phytother. Res. 17:1182–1187.

Garrido, G., D. Gonzalez, Y. Lemus, D. Garcia, D. Garcia, J. Leiro, R. Delgado, et al. 2003. In vivo and in vitro anti-inflammatory activity of Mango L. extract (Vimang®). Pharmocol. Res. 50:143–149.

Gray, S. P., and M. E. Cooper. 2011. Diabetic nephropathy in 2010: alleviating the burden of diabetic nephropathy. Nat. Rev. Nephrol. 7:71–73.

Garrido, G., D. Gonzalez, Y. Lemus, D. Garcia, D. Garcia, J. Leiro, R. Delgado, et al. 2003. In vivo and in vitro anti-inflammatory activity of Mango L. extract (Vimang®). Pharmocol. Res. 50:143–149.

Gray, S. P., and M. E. Cooper. 2011. Diabetic nephropathy in 2010: alleviating the burden of diabetic nephropathy. Nat. Rev. Nephrol. 7:71–73.
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Grover, J. K., S. Yadav, and V. Vats. 2002. Medicinal plants of India with antidiabetic potential. J. Ethnopharmacol. 81:81–100.

Guerrero-Romero, F., and M. Rodriguez-Morán. 2006. Hypomagnesemia, oxidative stress, inflammation and metabolic syndrome. Diab/Metab. Res. Rev. 22:471–476.

Idris, A. S., K. F. H. Mekky, B. E. E. Abdalla, and K. A. Ali. 2011. Liver function tests in type 2 Sudanese diabetic patients. Int. J. Nutr. Metab. 3:17–21.

International Diabetes Federation. 2009. The diabetes atlas, 4th ed. Brussels, International Diabetes Federation.

Iridoni, A. E., G. Oboh, A. A. Akindahunsi, A. A. Boligon, and M. L. Athayde. 2014. Phenolic composition and inhibitory activity of Mango and Horse-eye bean seeds extracts against key enzymes linked to the pathology and complications of type 2 diabetes. Asian Pac. J. Trop. Biomed. 4:903–910.

Iwai, K., M. Y. Kim, A. Onodera, and H. Matsue. 2006. Alpha-glucosidase inhibitory and antihyperglycemic effects of polyphenols in the fruit of Vibrurnum dilatatum Thunb. J. Agric. Food Chem. 54:4588–4592.

Iwalokun, B. I., J. A. Alibi-Sofunde, T. Odunala, O. A. Magbagbeola, and A. I. Akinwande. 2006. Hepatoprotective and antioxidant activities of Vernonia amygdalina on acetaminophen-induced hepatic damage in mice. J. Med. Foods 9:526–530.

Jain, C., A. Singh, P. Kumar, and K. Gautam. 2014. Anti-diabetic potential of flavonoids and other crude extracts of stem bark of Mango Linn: a comparative study. J. Sci. Innovative Res. 3:21–27.

Jayasri, M. A., S. Gunasekaran, A. Radha, and T. L. Mathew. 2008. Antidiabetic effect of Costus pictus leaves in normal and streptozotocin-induced diabetic rats. Int. J. Diab. Metab. 16:117–122.

Kaur, G., P. Kamboj, and A. N. Kalia. 2011. Antidiabetic and anti-hypercholesterolemic effects of aerial parts of Sida cordifolia Linn. on Streptozotocin-induced diabetic rats. Indian J. Nat. Prod. Resour. 2: 428–434.

Kirpichnikov, D., S. I. McFarlane, and J. R. Sowers. 2002. Metformin: an update. Ann. Intern. Med. 137:25–33.

Kumar, S., D. K. Das, and U. S. Prasad. 1991. Free amino acid composition in fruit of two mango cultivars during maturation and ripening. Indian J. Plant Physiol. 34:1–6.

Kundu, D., A. Roy, T. Mandal, U. Bandyopadhyay, E. Ghosh, and D. Ray. 2013. Relation of microalbuminuria to glycosylated haemoglobin and duration of type 2 diabetes. Nig. J. Clin. Pract. 16:216–220.

Laakso, M. 2001. Insulin resistance and its impact on the approach to therapy of T2D. Int. J. Clin. Prac. Suppl. 121: 8–12.

Lenzen, S. 2008. The mechanisms of alloxan- and streptozotocin-induced diabetes. Diabetol 51:216–226.

Li, G., P. Zhang, J. Wang, E. W. Gregg, W. Yang, Q. Gong, et al. 2008. The long-term effect of lifestyle interventions to prevent diabetes in the China Da Qing diabetes prevention study: a 20-year follow-up study. Lancet 371:1783–1789.

Lowry, O. H., N. J. Rosenbrough, A. L. Farr, and R. J. Randall. 1951. Protein measurement with folin phenol reagent. J. Biol. Chem. 193:265–270.

Mausuthisakul, P., and M. H. Gordon. 2009. Antioxidant and tyrosinase inhibitory activity of mango seed kernel by product. Food Chem. 117:332–341.

Mansour, H. A., A. Newairy, S. A. Youssef, and M. I. Sheweita. 2002. Biochemical study on the effects of some Egyptian herbs in alloxan-induced diabetic rats. Toxicology 170:221–228.

Marianna, M., D. Chaonan, H. QingPing, L. Yanling, and L. Ping-A. 2006. Streptozotocin-induced diabetes causes astrocyte death after ischemia and reperfusion injury. Diabetes 55:349–355.

Masood, N., G. H. Baloch, R. A. Ghori, I. A. Memon, M. A. Memon, and M. S. Memon 2009. Serum zinc and magnesium in type-2 diabetic patients. J. Coll. Physicians Surg. Pak. 19:483–486.

Murray, R. K., D. K. Granner, and P. A. Mayes. 2000. Harper’s biochemistry, 25th ed. Appleton and Lange, Stamford, CT.

Murugan, P., and L. Pari. 2007. Influence of tetrahydrocurcumin on hepatic and renal functional markers and protein levels in experimental type 2 diabetic rats. Basic Clin. Pharmacol. Toxicol. 101:241–245.

Muruganandan, S., K. Scrini, S. Gupta, P. K. Gupta, and J. Lal. 2005. Effect of mangiferin on hyperglycemia and atherogenicity in streptozotocin diabetic rats. J. Ethnopharmacol. 97:497–501.

Nwanjo, H. U. 2007. Studies on the effect of aqueous extract of Phyllanthus niruri on plasma glucose level and some hepatospecific markers in diabetic Wistar rats. Int. J. Lab. Med. 2:1–18.

Oboh, G., A. J. Akinremi, A. O. Ademiluyi, and S. A. Adefegha. 2010. Inhibitory effects of aqueous extract of two varieties of ginger on some key enzymes linked to type-2 diabetes in vitro. J. Food Nutr. Res. 49:14–20.

Okawa, H., N. Ohishi, and K. Yagi. 1979. Assay for lipid peroxides in animal tissues by thiobarbituric acid reaction. Analy. Biochem. 95:351–358.

Ojewole, J. A. O. 2005. Antiinflammatory, analgesic and hypoglycemic effects of Mangifera indica Linn. (Anacardiaceae) stem-bark aqueous extract. Methods Find. Exp. Clin. Pharmacol. 27:547–554.

Ong, K. C., and H. E. Khoo. 2000. Effects of myricetin on glyceremia and glycogen metabolism in diabetic rats. Life Sci. 67:1695–1705.

Orhan, N., A. Berkkan, D. D. Orhan, M. Aslan, and F. Ergun. 2011. Effects of Juniperus oxycedrus ssp. oxycedrus on tissue lipid peroxidation, trace elements (Cu, Zn, Fe)
and blood glucose levels in experimental diabetes. J. Ethnopharmacol. 133:759–764.

Parveen, K., R. Khan, and W. A. Siddiqui. 2011. Antidiabetic effects afforded by Terminalia arjuna in high fat-fed and streptozotocin-induced type 2 diabetic rats. Int. J. Diab. Metab. 19:23–33.

Pedersen, T. R. 2001. Low density lipoprotein cholesterol lowering is and will be the key to the future of lipid management. Am. J. Cardiol. 87:8B–12B.

Perez-Gutierrez, R. M., and J. M. Mota Flores. 2010. Attenuation of hyperglycaemia and hyperlipidaemia in streptozotocin-induced diabetic rats by chloroform extract of fruits of Ferocactus latispinus and Ferocactus histrix. Bol. Latinoam. Caribe Plant. Med. Aromat. 9:475–484.

Prabhu, K. S., R. Lobo, and A. Shirwaikar. 2008. Antidiabetic properties of the alcoholic extract of Sphaeranthus indicus in streptozotocin-nicotinamide diabetic rats. J. Pharm. Pharmacol. 60:909–916.

Puressglove, J. W. 1991. P. 29Tropical crops: dicotyledons. Harlow, Longman Scientific and Technical Publishers, Harlow.

Rahman, I. 2003. Oxidative stress, chromatin remodeling and gene transcription in inflammation and chronic lung diseases. J. Biochem. Mol. Biol. 36:95–109.

Reed, M. L., K. Meszaros, L. J. Entes, M. D. Claypool, J. G. Pinkett, T. M. Gadbois, et al. 2000. A new rat model of type 2 diabetes: the fat-fed, streptozotocin-treated rat. Metabolism 49:1390–1394.

Ribeiro, S. M. R., L. C. A. Barbosa, J. Queiroz, H. M. Knodler, and A. Schieber. 2008. Phenolic compounds and antioxidant capacity of Brazilian mango (Mango L.) varieties. Food Chem. 110:620–626.

Rohlfing, C. L., R. R. Little, H. Wiedmeyer, J. D. England, R. Madsen, M. I. Harris, et al. 2000. Use of GHb (HbA1c) in screening for undiagnosed diabetes in the U.S. population. Diab. Care 23:187–191.

Saudek, C. D., R. R. Kalyani, and R. L. Derr. 2005. Assessment of glycaemia in diabetes mellitus: haemoglobin A1c. J. Assoc. Physicians India 53:299–305.

Selvan, V. T., L. Manikandan, K. G. P. Senthil, R. Suresh, and B. Kakoti. 2008. Antidiabetic and antioxidant effect of methanol extract of Artanema sesamoides in streptozotocin-induced diabetic rats. Int. J. Appl. Res. Nat. Prod. 1:25–33.

Shalaby, E. A., and S. M. M. Shanab. 2013. Antioxidant compounds, assays of determination and mode of action. Afr. J. Pharm. Pharmacol. 7:528–539.

Shivan, N., M. Naika, F. Khanum, and V. K. Kaul. 2013. Antioxidant, anti-diabetic and renal protective properties of Stevia rebaudiana. J. Diab. Complications 27:103–113.

Singh, N. S., P. Vats, S. Suri, R. Shyam, M. M. L. Kumria, S. Ranganathan, et al. 2001. Effect of an antidiabetic extract of Catharanthus roseus on enzymic activities in streptozotocin induced diabetic rats. J. Ethnopharmacol. 76:269–277.

Appendix

Figure A1. Mangifera indica L. (mango) seeds.