Blend of Roselle Calyx and Selected Fruit Modulates Testicular Redox Status and Sperm Quality of Diabetic Rats

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Abstract

Hyperglycaemia-induced oxidative stress has been reported to be associated with testicular failure leading to sexual dysfunction, impotence and infertility. The effect of blend of roselle calyx and selected fruits on the testicular antioxidant activities and sperm quality of diabetic rats was investigated. Diabetes was induced by a single intraperitoneal injection of alloxan. Treatment lasted for 14 days after induction. The rats were sacrificed by cervical dislocation. Testicular tissues were used for assessment of GSH, catalase, SOD and lipid peroxidation. Sperm cells were analyzed for sperm motility, counts and abnormality. Induction of diabetes led to a significant decrease in GSH level, elevated SOD and catalase activities. These were significantly modified by the blend. The blends were observed to reduce malondialdehyde level. Induction of diabetes led to a significant decrease in the studied sperm quality parameters, treatment with the blend significantly improved these qualities. This study indicates improved testicular antioxidant activities and sperm qualities by single and double doses of the fruit blend suggesting its protective potential against spermatoxic and testicular toxicity in diabetics.

Keywords: Roselle calyx; Fruits; Sperm cells; SOD; CAT; MDA; GSH

Introduction

Diabetes has been described as a disease where the body produces little insulin and/or ceases to produce insulin, or becomes progressively resistant to its action [1]. It is a chronic disorder in metabolism of carbohydrates, proteins, and fat often characterized by hyperglycaemia [2].

The involvement of oxidative stress in the progress of diabetic complications has been well documented as well as its implication in diabetic pathogenesis [3]. Oxidative stress is caused by generation of free radicals such as reactive oxygen species (ROS). In physiological condition, the body synthesizes both free radicals and anti-oxidants [4]. Oxidative stress sets in when an imbalance occurs between these free radicals and antioxidants in favor of the free radicals [5]. Hyperglycaemia – induced oxidative stress have been reported to occur via increased glycolysis; auto-oxidation of glucose and non-enzymatic protein glycation [6].

The testis contains an elaborate array of antioxidant enzymes and free radical scavengers to protect against oxidative stress. This is of great importance as peroxidative damage is currently regarded as the major cause of impaired testicular functions. Hyperglycaemia have been reported to be associated with testicular failure leading to sexual dysfunction, impotence and infertility [6]. The lipids in sperm are the main substrates for peroxidation, which is induced by ROS generated by hyperglycaemia. This will ultimately lead to infertility in most men, characterized by low sperm count and motility [7].

The use of plants in the use in the management of diabetes has been well documented [8]. The medicinal properties of fruits have been reported. Their daily consumption has been associated with reduced risk of cancer, heart disease, premature aging, stress, diabetes, and fatigue (Shibumon). This is primarily due to the integrated action of oxygen radical scavengers such as β- carotene and ascorbic acid, calcium and dietary fiber [9]. Among such fruits are the pawpaw (Carica papaya L.), and grape (Citrus paradisi) fruits. Their anti-diabetic and/or hypoglycemic properties have been reported in several studies [10-12]. Extracts of unripe pawpaw fruits (Carica papaya L.) have been used in treatment of diabetes [10,12], while grape fruits (Citrus paradisi) have been reported to exhibit antidiabetic effect by protecting beta cells in the pancreas [11,12]. However, the post-harvest shelf life of these fruits is very limited due to their perishable nature, thereby leading to wastage [9]. Blending of two or more fruit juices for the preparation of ready-to-serve beverage have been shown to be a convenient alternative for ensuring a longer postharvest shelf life and utilization of these fruits [9,13]. Blending of fruit juice helps in improving flavor, taste, nutritive value, and reduces production cost [9,14]. Blending juices with other medicinal plants such roselle calyx (Hibiscus sabdarifa) and guava leaves (Psidium guajava L) improves the medicinal properties. Guava leaves have been shown to exhibit hypoglycemic effect in normal and diabetic rats [15]. Its ability to inhibit increase plasma glucose level in alloxan-induced diabetic rats has been reported [12]. Locally known as Zoborodo in Northern Nigeria, aqueous extract of roselle calyx has been demonstrated as having beneficial effects on anti-oxidation and lipid-lowering in experimental diabetic studies [16]. A blend of these fruits with the mentioned medicinal plants would therefore form a natural health drink suitable for the treatment and/or management of diabetes and its related complication.

This paper reports the effect of blends of these fruits with roselle calyx (Hibiscus sabdarifa) and guava leaves on the testicular antioxidant activities and sperm quality as well as blood glucose level of diabetic rats.

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Materials and Methods

Plant materials

Improved varieties of unripe pawpaw fruits (Carica papaya), grape fruits (Citrus paradisi) as well as guava leaves (Psidium guajava) were identified and obtained from National Horticultural Research Institute (NIHORT), Ibadan, Nigeria. Roselle calyx (Hibiscus sabdarifa) was obtained from Mushin market in Lagos, Nigeria. The varieties of pawpaw, guava, and grape fruit identified and purchased were Cg variety of pawpaw, sweet grape and local guava respectively.

Preparation of extracts

Extracts of pawpaw fruit, guava leaf and grape fruit were produced using hot extraction methods as described by Okafor et al. [17]. Briefly, unripe pawpaw fruits were washed, peeled, seeds were removed and the fruits were cut into smaller sizes. This was boiled for 30 minutes at 60°C. The boiled fruits were then minced in a warring blender with the water in which they were boiled to get slurry, which was later filtered using a fine sieve to get the extract. Grape fruits were washed, peeled, cut into smaller pieces; seeds were removed and minced in a warring blender into slurry. The slurry was then filtered using a fine sieve. The guava leaves and the roselle calyx (Hibiscus sabdarifa) were sorted, washed and boiled in water separately for 20 minutes, and the extracts were decanted.

Blend formulation

Varying proportion of the extracts were mixed together to get a blend as depicted on table 1. Roselle calyx (Hibiscus sabdarifa) was used as a carrier and part of the formulation. The blend was pasteurized at 80°C.

Animals

Twenty male albino rats of wister strain weighing about 150-200 g were used for the study. They were fed on standard rat pellet diet and allowed to adapt for one week. They were provided water ad libitum and maintained under standard laboratory conditions of natural photo period of 12-hr light - dark cycle. The animals used in the present study were maintained in accordance with the approval of the Animal Ethical Committee, Federal Institute of Industrial Research, Lagos, Nigeria

Induction of diabetes

Diabetes was induced by a single intraperitoneal injection of 150 mg/kg of alloxan monohydrate in normal saline water in a volume of about 3 mL. After 72 hours, the diabetic rats (glucose level > 150 mg/dl) were treated with a double dose compared to the first week of treatment. However, there was a 32.90% increase in the blood glucose level of rats treated with 2.5 ml of the blend led to 13.25% reduction of GSH activity in the testes of the experimental rats (p<0.05) reduction of GSH activity in the testes of the experimental rats in the second week.

Experimental design

The rats were divided into four groups, each consisting of five animals.

Group 1 – Pelletized mouse chows

Group 2 – Diabetic (Untreated)

Group 3 – Diabetic + 2.5 ml of blend/bw

Group 4 – Diabetic + 5 ml of blend/bw

The rats were monitored daily for food and water intake, and body weight. The fruit blends were orally administered. Blood glucose levels of the rats were monitored on weekly basis with a glucometer. Treatment lasted for 14 days after induction. At the end of the treatment trials, the rats were fasted overnight and sacrificed by cervical dislocation.

Preparation of tissue homogenates

The testes were removed, rinsed in ice-cold 1.15% KCI solution to wash off excess blood, blotted dry with filter paper and weighed. The organs were homogenized in four parts of homogenizing buffer and centrifuged at 10,000 g for 15 min in a ultracentrifuge at a temperature of -2°C to get the mitochondrial fraction. The supernatant (post-mitochondrial fraction) was decanted and stored at -4°C for subsequent analysis. Each time the supernatant was outside the freezer, it was kept in ice bags.

Determination of oxidative stress parameters

Lipid peroxidation was determined by measuring malondialdehyde (MDA) formed by thiobarbituric acid reaction (TBAR) [18]. Catalase (CAT) activity was estimated by measuring the rate of decomposition of H2O2 [19]. The level of superoxide dismutase (SOD) activity was determined by the method of Kakkar et al. [20], while the method of Ellman [21] was adopted in estimating the activity of reduced glutathione (GSH).

Evaluation of sperm quality

The tail of the epididymis was cut into small pieces in 2 mL of normal saline and sperm collected by squeezing it gently on clean slide [22]. Sperm motility and count were evaluated using conventional methods [23]. Sperm abnormality was carried out by microscopical examination of the seminal smears stained with Eosin and Nigrosin stain. Presence of epithelial cells, triple phosphate crystals and oil droplets were also investigated.

Statistical analysis

Statistical significance was established using one-way analysis of variance (ANOVA), and data were reported as mean ± standard deviation. Significant difference was established at P<0.05. Statistical analyses were carried out using SPSS for Windows, version 15.0 (SPSS Inc., Chicago, IL).

Results

Blood glucose level

Induction of diabetes led to an increase in the blood sugar level as depicted in figure 1. Treatment with 2.5 ml of the blend led to 13.25% reduction of the blood glucose level which was statistically significant (p<0.05) at the first week of treatment, while at a double dose (5.0 ml) a 56.62% reduction was observed. A further 33.95% reduction was observed in the single dose group (group 3) in the second week. However, there was a 32.90% increase in the blood glucose level of rats treated with a double dose compared to the first week of treatment.

Antioxidant activities

There were significant differences (p<0.05) in all the studied antioxidant parameters. Induction of diabetes led to a significant (p<0.05) reduction of GSH activity in the testes of the experimental rats...
abnormality. The concentration of pus cells was observed to be constant in all experimental groups. The concentration of epithelial cells were observed to increase in the untreated diabetic group, this was reduced in both treated groups. Triple phosphate crystal was observed in the sperm cells of the untreated diabetic group but absent in the treated. Oil droplets were observed in the sperm cells of the groups 1 and 4 respectively.

**Discussion**

The antioxidant capacity of juices (fruit blends) has been linked to in vivo protection from oxidative stress in numerous studies [24]. Various studies have also demonstrated the ability of these blends to scavenge free radicals and protect against lipid peroxidation *in vitro* [25]. These studies emphasized the protective effect of fruits against oxidative damage. These antioxidant activities can be explained in part by their dietary components especially the phytochemicals and vitamin C [26]. The testes are susceptible to oxidative stress despite the low oxygen tensions that typify the testicular cells. This is attributed to abundance of highly unsaturated fatty acids and the presence of potential ROS – generating systems [27]. The testes have developed

Sperm quality

The effects of the blend at both single and double doses are shown in table 2. The sperm cells were observed to be grayish and opaque. They were also observed to be watery. Sperm count, and motility were significantly decreased in the untreated diabetic group in comparison to the control. These were observed to increase in the treated groups. However, treatment with single dose (2.5 ml) had no effect on the motility. Sperm abnormality was observed to increase in the untreated group. Treatment with single and double dose had no effect on the

![Figure 1: Blood glucose level of experimental groups. Values = mean ± SD; n = 5. Note: Values = mean ± SD; n = 5. a= statistical significant (p < 0.05) as compared with group 1; b = statistical significant (p < 0.05) as compared with group 2; c = statistical significant (p < 0.05) as compared with group 3; d = statistical significant (p < 0.05) as compared with group 4.](image1)

![Figure 2: Effect of blends on GSH activities of testes in diabetic rats. Values = mean ± SD; n = 5. Note: Values = mean ± SD; n = 5. a= statistical significant (p < 0.05) as compared with group 1; b = statistical significant (p < 0.05) as compared with group 2; c = statistical significant (p < 0.05) as compared with group 3; d = statistical significant (p < 0.05) as compared with group 4.](image2)

![Figure 3: Effect of blends on SOD activities of testes in diabetic rats. Values = mean ± SD; n = 5. Note: Values = mean ± SD; n = 5. a= statistical significant (p < 0.05) as compared with group 1; b = statistical significant (p < 0.05) as compared with group 2; c = statistical significant (p < 0.05) as compared with group 3; d = statistical significant (p < 0.05) as compared with group 4.](image3)

![Figure 4: Effect of blends on CAT activities of testes in diabetic rats. Values = mean ± SD; n = 5. Note: Values = mean ± SD; n = 5. a= statistical significant (p < 0.05) as compared with group 1; b = statistical significant (p < 0.05) as compared with group 2; c = statistical significant (p < 0.05) as compared with group 3; d = statistical significant (p < 0.05) as compared with group 4.](image4)
Increased SOD and CAT activities due to oxidative stress have been reported [34]. Their observed increase in the testes of untreated diabetic rats might be attributed to hyperglycemia - induced oxidative stress. The importance of SOD in controlling O₂⁻ leakage from testicular mitochondria has been reported [35]. Gu and Hecht [35] further reported that its mRNA is higher in the testes than the liver. CAT catalyzes the decomposition of hydrogen peroxide (H₂O₂) into less reactive gaseous oxygen and water molecules [36]. The present study demonstrated significant decreases in SOD and CAT activities in testes of rats treated with blends at both single and double dose.

Induction of lipid peroxidation (LPO) in diabetics has been observed in numerous tissues both in vitro and in vivo [37]. The testes are potent target owing to the high concentration of polyunsaturated fatty acids. In the present study, induction of diabetes enhanced the level of testicular lipid peroxidation in accordance to previous studies. The increased MDA level in the untreated diabetic rats depicts peroxidation of the lipids of the testicular tissues. Reduction of the MDA level by both doses of the blend reflects their protective potentials against hyperglycemia – induced peroxidation of testicular membrane lipids.

Increasing evidence suggests that diabetes has an adverse effect on male reproduction function and oxidative stress may be involved [7]. This is evidenced by the sperm quality of the untreated diabetic rats. Their low sperm motility and count as well as percentage abnormality corresponds to previous studies on the sperm quality of diabetic rats [7,27,38]. Single dose of the blend was observed not to improve the sperm motility. However, at double dose the cell motility was restored. Insufficient sperm motility is a common cause of infertility, thus the improved motility at a double dose of the blend signifies improvement of the sperm quality and improved fertility. The observed improvement of the sperm count by both single and double doses of the blend portrays its potency in the management of the diabetic-induced spermatotoxic and impotency in males. This could be attributed to the ability of the blend to improve the testicular antioxidant activities as evidenced by the enhanced antioxidant markers. The reduced concentration of epithelial cells and absence of triple phosphate crystals in the semen in the testes of the treated group further portrays the potency ability of the blend. Epithelial cells from the genitourinary tract, as well as leukocytes and immature germ cells are often present in ejaculates other than the spermatooza which may be clinically relevant [39].

**Conclusion**

This paper indicates improved testicular antioxidant activities and sperm qualities by single and double doses of the fruit blend suggesting its protective potential against spermatotoxic and testicular toxicity in diabetic male rats.

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Table 2: Effect of blend on sperm quality of diabetic rats.

| Parameters               | Group 1 | Group 2 | Group 3 | Group 4 |
|--------------------------|---------|---------|---------|---------|
| Motility                 | 50%     | nil     | nil     | nil     |
| Sperm Count              | 800 × 10⁶ ml | 40 × 10⁶ ml | 75 × 10⁶ ml | 72 × 10⁶ ml |
| Abnormal Count           | 1%      | 10%     | 10%     | 10%     |
| Epithelial Cells         | *       | **      | *       | *       |
| Triple Phosphate Crystal | nil     | nil     | nil     | nil     |
| Oil Droplet              | **      | nil     | nil     | ***     |

Figure 5: Effect of blends on LPO in testes in diabetic rats. Values = mean ± SD; n = 5. Note: Values = mean ± SD; n = 5; a = statistical significant (p < 0.05) as compared with group 1; b = statistical significant (p < 0.05) as compared with group 2; c = statistical significant (p < 0.05) as compared with group 3; d = statistical significant (p < 0.05) as compared with group 4.
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