Diabetogenic effect of gluten in Wistar albino rats: a preliminary preclinical screening

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Abstract

Background and aims. Gluten-related disease affects less than 1% population and is not considered of relevance at the public health level. However, the consumption of a gluten-free diet has been most commonly adopted as a special diet worldwide in the recent past. In the present study, we investigated the association of gluten intake and diabetes in Wistar albino rats.

Methods. Thirty adult Wistar rats were randomly divided into five groups: control, diabetic, and test treated with pure gluten (100, 200 and 400 mg/kg body weight). Diabetes was induced in rats by intraperitoneal injection of Streptozotocin (65 mg/kg) after a dose of nicotinamide (110 mg/kg). Body weight, fasting blood glucose levels, postprandial blood glucose levels and histopathology of the pancreas were compared.

Results. Fasting blood glucose levels and postprandial blood glucose were significantly higher in diabetes animals but there were no significant changes in gluten treated groups. Other parameters were not significantly changed among different groups.

Conclusions. Gluten at doses 100 mg/kg, 200 mg/kg and 400 mg/kg is not a diabetogenic diet and hence it needs not be excluded from diet for the prevention and management of Type 2 diabetes mellitus.

Keywords: animals, blood glucose, diabetes mellitus, glutens, niacinamide, streptozocin

Introduction

Diabetes mellitus (DM) is a metabolic disorder of several etiologies characterized by prolonged hyperglycemia with disturbances of carbohydrates, fat and protein metabolism resulting from defects in insulin secretion, insulin action, or both [1]. It is a growing public health concern worldwide. The prevalence of diabetes is increasing fast becoming an epidemic in some countries of the world [2]. Type-2 diabetes mellitus (T2DM) is characterized by insulin insensitivity due to insulin resistance, diminishing insulin production and subsequent pancreatic β-cell failure. Lifestyle factors and genetics play a role in the development of T2DM [3]. Various aspects of diet causing insulin resistance have been hypothesized, over the long term, to influence the risk of T2DM. It has been observed in metabolic studies that carbohydrates with a high glycemic index (a qualitative indicator of carbohydrate’s ability to raise blood glucose levels) seem to increase insulin demand and accentuate hyperinsulinemia [4]. Type-1 diabetes mellitus (T1DM) is an autoimmune disorder caused by the destruction of the insulin-producing β-cells of the pancreas followed by absolute insulin deficiency and hyperglycemia [5].

The taxonomic classification of cereals sub classifies wheat, rye, and barley under the gluten tribe. Prolamin which is further separated into gliadins
Materials and methods

Chemicals

Standard drug Streptozotocin and nicotinamide were purchased from Sigma Chemical Company. St. Louis, USA. Streptozotocin was dissolved in citrate buffer (pH 4.5) and Nicotinamide in normal saline. Streptozotocin was dissolved in citrate buffer (pH 4.5), Nicotinamide in normal saline.

Pure wheat gluten was procured from the Vintop products Pvt. Ltd, Bangalore, India. Nutrition constituents of the procured gluten powder are as follows; total calories (0), total fat (0 g), cholesterol (0 mg), sodium (0 mg), total carbohydrate (0 g), protein (85 g), vitamin A (0%), vitamin C (0%), calcium (0%) and iron (0%). Since gluten was not clearly soluble in distilled water, 1% gum acacia was used to dissolve gluten.

Animals

Adult Wistar rats of either sex were obtained from the Central Animal House, A. J. Institute of Medical Sciences, Mangaluru, India bearing registration number 1075/PO/Re/S/07/CPCSEA. Prior to experiment ethical approval was obtained from the Institutional Animal Ethical Committee under the reference number IAEC/02/2016/CPCSEA.

Rats were housed in clean polypropylene cages, six rats in each cage, in a controlled environment (24 ± 2°C) with 12-hour light and dark cycle. They were fed with commercial pelleted chow and water ad libitum. All efforts were made to minimize the animal suffering and the number of animals used. Following experimentation, all the animals were rehabilitated at the Central Animal House facility. Committee for the Purpose of Control and Supervision of Experiments on Animals (CPCSEA) guidelines were followed throughout the experiment.

Induction of diabetes in experimental animals

Diabetes was induced by administering streptozotocin (65 mg/kg body weight; i.p), following 15 min nicotinamide administration (110 mg/kg body weight; i.p.) [12,13]. Random blood glucose was measured after 72 h of administration of streptozotocin-nicotinamide and animals blood glucose levels higher than 200 mg/dL, referred to as diabetic animals, were included in group II as diabetic control and group III to V was given the gluten treatment [14,15].

Experimental design

Animals were randomly divided into five groups (each group containing 6 rats) as follows:

Group I (Normal Control), animals with a normal diet, received 1% of gum acacia orally(equivalent volume of gluten) for 90 days.

Group II (Diabetic Control), streptozotocin-nicotinamide induced diabetic animals with a normal diet, received 1% of gum acacia orally (equivalent volume of gluten) for 90 days.

Group III (Test G100), animals which received gluten dissolved in 1% gum acacia at doses of 100 mg/kg body weight for 90 days.

Group IV (Test G200), animals which received gluten dissolved in 1% gum acacia at doses of 200 mg/kg body weight for 90 days.

Group V (Test G400), animals which received gluten dissolved in 1% gum acacia at doses of 400 mg/kg body weight for 90 days.

Animal groups are illustrated in Figure 1.
All the drugs were administered once daily orally. Body weights of rats were checked on the first day, then every week for the first one month and every month for another 3 months. The animals were also monitored for any behavioral changes during the study period.

**Blood collection and biochemical assessment**

Blood was collected from the tail vein of animals. Biochemical parameters like fasting blood glucose levels and postprandial blood glucose levels were checked on the first day, then every week for the first one month and every month for another 3 months.

Blood glucose levels were determined by a glucometer (Elegance CT-X10, convergent technologies, Germany) and biochemical assay kits (Pars Azmoon, Iran), with a sensitivity of 0.1 mg/dl.

After the completion of the experiment, i.e. after 90 days, rats were sacrificed after giving euthanasia, dissection was performed and head of the pancreas was separated and was sent to a pathology laboratory for histopathological examination.

**Statistical analysis**

Mean values of all the parameters were calculated. The data are presented as mean ± SD. The comparison between the diabetic group and the test groups was performed using one-way ANOVA. A p-value of less than 0.05 was considered significant. All data were analyzed statistically using GraphPad Prism 6 software (San Diego, CA, USA).

**Results**

Random blood glucose level was significantly increased in group II after two diabetic inductions when compared to Group I (Table I). However, there was no significant change in the mean fasting blood sugar levels in any of the test groups (group III, IV and V) compared to the normal control group on 8th to 90th day of the experiment. Similarly, mean postprandial blood sugar levels were significantly increased in diabetes animals but did not change in gluten treated test animals of group III, IV and V (Table II). On the 8th day, random blood glucose level was increased but no significant while postprandial blood sugar levels found significant higher diabetes animals.

**Table I.** Fasting Blood Sugar (mg/dL) in control, diabetic and gluten treated rats.

| Group          | 1st Day | 8th Day | 15th Day | 21st Day | 28th Day | 60th Day | 90th Day |
|----------------|---------|---------|----------|----------|----------|----------|----------|
| 1. Normal Control | 80±3.26 | 78.33±2.28 | 80±3.14 | 79.67±2.99 | 79.57±3.53 | 79.37±2.60 | 84±1.71 |
| 2. Diabetic Control | 83±6.51 | 142.7±47.02 | 160.67±38.48* | 204.33±73.31* | 124.5±24.47* | 329±66.61** | 337.17±69.48** |
| 3. Gluten G100   | 88.67±2.47 | 96.17±5.07 | 84.5±4.96 | 70.83±3.02 | 78.57±3.35 | 66.57±4.49 | 65.33±3.47 |
| 4. Gluten G200   | 91±5.62 | 99.5±2.58 | 82.83±3.95 | 70.33±2.19 | 78.33±4.84 | 82.5±3.73 | 68.5±1.59 |
| 5. Gluten G400   | 97.5±9.99 | 103.67±3.46 | 81.83±2.27 | 80±2.10 | 79.5±2.85 | 79.17±2.46 | 73.5±2.55 |

Observations are Mean ± SEM, ANOVA followed Dunnett’s multiple comparison test, p value < 0.05 *, p value < 0.01 **
There was no significant change in body weight between Group I and Group II on from the first week to three-month comparison. Gluten treated test group (group III, IV and V) showed a slight increase in body weight but it was not statistically significant when compared to Group I and II (Table III).

Histopathological report of test groups revealed that there was no increase in the number and size of the islets, no necrosis of the β-cells, and no inflammatory lesions in the exocrine parenchyma and there was no degranulation of the β-cells. These reports suggested that there were no diabetic changes in the β-cells of the head of the pancreas in test groups with gluten at doses 100 mg/kg, 200 mg/kg and 400 mg/kg body weight (Figure 2).

### Discussion

Diabetes mellitus is one of the chronic disorders which accounts for about 10% of the diseases in the world’s population [16]. Among different types of diabetes, T2DM is the commonest one. To start with, it presents as an increase in the glucose levels of blood and in the long term it can result in severe impairment and failure of various organs [17,18]. By controlling T2DM effectively, we can prevent long term complications associated with the disease.

The various risk factors like genetic, environmental, and metabolic are interrelated and contribute to the progression of T2DM [19]. A major goal in the management includes lifestyle management which comprises of diet control, physical activity and oral hypoglycemic
Gluten is referred as a complex protein which is found in certain grains like wheat, rye, and barley. Gluten altered the intestinal microbiota that maintains the intestinal permeability [16]. These bacteria produce short-chain fatty acids by the breakdown of dietary fibers which includes butyrate, acetate, and propionate. Butyrate and acetate induce intestinal permeability through lipoxygenase activation via histone acetylation [21]. Butyrate can also boost the number and functions of regulatory T cells which are known to suppress inflammatory responses [22]. Therefore, increased permeability leads to the production of a number of cytokines that can, in turn, promote inflammation and damage in the small intestine. Many studies have demonstrated the correlation between the intake of gluten and the subsequent development of T2DM [4,7,10]. Gluten is established to cause the destruction of β-cells in T1DM which is autoimmune in nature [9]. However, the scientific evidence associating gluten with T2DM is still lacking.

In the present study, pure gluten was used to determine its effect on blood glucose levels in adult WISTAR Albino strain rats over a period of time and to determine the possible damage to the β-cells of pancreatic tissue. Streptozotocin-Nicotinamide induced T2DM is a model that shares characteristic features with human T2DM. When nicotinamide was administered along with the streptozocin, it gives partial protection to insulin-secreting cells against streptozotocin, hence mimicking a T2DM like state [23].

The present study was conducted to evaluate the possible diabetogenic activity of gluten at doses 100 mg/kg, 200 mg/kg and 400 mg/kg in rats. The results suggest that administration of gluten at doses 100 mg/kg, 200 mg/kg and 400 mg/kg body weight for 90 days did not show any significant increase in the blood glucose levels and body weight in rats. Similarly, recently Zong et al., also showed an inverse association between gluten intake and T2DM risk among large cohort healthy US population [24].

Whereas streptozocin, a standard drug which was given along with the nicotinamide, significantly increased the blood glucose levels in rats. The results suggest that administration of gluten at doses 100 mg/kg, 200 mg/kg and 400 mg/kg body weight for 90 days did not show any significant increase in the blood glucose levels and body weight in rats.

Histopathological report of test groups revealed that there was no diabetic changes in the β-cells of the head of the pancreas in test groups with gluten at doses 100 mg/kg, 200 mg/kg and 400 mg/kg body weight. Gluten intakes have no significant adverse effects on the incidence of T2DM or weight gain [25].

Still, there is a debate over the belief that simply avoiding or eliminating highly processed foods (including highly processed gluten-containing foods) may or may not have health benefits [26]. Limiting gluten intake may limit the beneficial nutrients like cereal fibers, micronutrients and minerals in the diet [27]. Therefore, careful investigation on how factors associated with this diet affect a variety of symptoms, including gastrointestinal function, cognition, and overall well-being is required.

**Conclusions**

Our preliminary results have shown that gluten has no effect on increasing the blood sugar level and body weight at doses 100 mg/kg, 200 mg/kg and 400 mg/kg in the experimental rats which were used in this study. The histopathological study which was done after the study also did not show any changes in the β-cells of the islets of the pancreas. The study infers that gluten may not a diabetogenic diet, but further studies required to see long-term effect of gluten consumption.

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