EFFECT OF SUPPLEMENTATION OF VITAMIN C ON BLOOD GLUCOSE AND GLYCOSYLATED HEMOGLOBIN LEVELS IN EXPERIMENTALLY-INDUCED DIABETIC RABBITS

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INTRODUCTION

Diabetes mellitus (DM) is a spectrum of common metabolic disorders, arising from a variety of pathogenic mechanisms, all resulting in hyperglycemia. The number of individuals with diabetes is rising rapidly throughout the world. Both genetic and environmental factors contribute to its pathogenesis, which involve insufficient insulin secretion, reduced responsiveness to endogenous or exogenous insulin, increased glucose production, and/or abnormalities in fat and protein metabolism. The resulting hyperglycemia may lead to both acute symptoms and metabolic abnormalities [1]. However, the major sources of the morbidity of diabetes are the chronic complications that arise from prolonged hyperglycemia, including retinopathy, neuropathy, nephropathy, and cardiovascular disease [2].

Diabetes is growing alarmingly in India, home to more than 65.1 million people with the disease in 2013, compared to 50.8 million in 2010, rising to 131.2 million by 2035, according to the International Diabetes Federation [3,4]. According to the World Health Organization criteria, the prevalence of known diabetes was 5.6% and 2.7% among urban and rural areas, respectively [5].

Ascorbic acid is one of the most effective, powerful antioxidants in biological fluids. Vitamin C scavenges physiologically important reactive oxygen species such as superoxide radical anion, hydrogen peroxide, hydroxyl radical and singlet oxygen, and also reactive nitrogen species [6-8]. It regenerates other small molecule antioxidants such as α-tocopherol, reduced glutathione, urate, and β-carotene from their respective radical species and may, therefore, prevent oxidative damage to biological macromolecules, including DNA, lipids, and proteins [9-12]. Vitamin C is structurally similar to glucose and hence possibly may replace it in many chemical reactions, such as non-enzymatic glycosylation of proteins (Maillard reactions).

Up-to-date, available literature suggests conflicting results related to supplementation of Vitamin C and improvement in blood glucose level and glycosylated hemoglobin (HbA1c). However, no study has examined the effects of Vitamin C with metformin in the treatment of Type 2 DM, especially on blood glucose levels in animals. Hence, the present study taken up to see the effect of supplementation of Vitamin C on blood glucose and glycosylated hemoglobin (HbA1c) along with metformin in experimentally-induced diabetic rabbits.

METHODS

The study was conducted at the Department of Pharmacology, Gandhi Medical College, Hyderabad, after obtaining ethical clearance from the Institutional Animal Ethics Committee.

Twenty-four male albino rabbits were randomized into groups. All rabbits were allowed an 1-week acclimatization period to become accustomed to the laboratory conditions. Rabbits were randomly divided into four groups, each comprising six rabbits.

RESULTS:

Twenty-four adult New Zealand white rabbits (1.5-2.5 kg) were divided into four groups, each containing six rabbits. Group 1: Normal control (distilled water); Group 2: Metformin (23.33 mg/kg) is given orally; Group 3: Metformin (23.33 mg/kg) plus Vitamin C 250 mg are given orally; and Group 4: Metformin (23.33 mg/kg) plus Vitamin C 500 mg are given orally. Animals were treated for 30 days. The blood samples were collected on days 0 and 30 from the marginal ear vein of rabbits for the estimation of blood glucose and HbA1c levels.

Statistically analyzed by ANOVA test followed by post hoc Tukey’s test using GraphPad Prism software. Results shown that Vitamin C 500 mg oral supplementation with metformin had very highly significantly reduced HbA1c levels by 40.12% (p = 0.0001***) and blood glucose levels by 49.12% (p = 0.0003***) whereas Vitamin C 250 mg oral supplementation with metformin also significantly reduced HbA1c levels by 25.49% (p = 0.0001***) and blood glucose levels by 42.95% (p = 0.0026**) when compared to metformin alone, which reduced HbA1c levels by 22% (p = 0.0001***) and blood glucose levels by 39.58% (p = 0.0001***).

CONCLUSION:

Oral supplementation of Vitamin C 500 mg/250 mg to the metformin was superior in reducing HbA1c levels and blood glucose levels compared to metformin alone in rabbits. Hence, Vitamin C oral supplementation may be helpful in lowering blood glucose levels and HbA1c levels and improving glycemic control in Type 2 DM.

Keywords: Alloxan, Blood glucose, Diabetic rabbits, Glycosylated hemoglobin, Metformin, Vitamin C.
Duration of the study
The duration of the study was for 45 days, which included 15 days of the diabetic induction period and the next 30 days of the treatment period.

Experimental animal model and study design
To evaluate the effect of supplementation of Vitamin C on blood glucose and HbA1c levels, the ‘Alloxan-induced diabetic rabbit model' was used.

Procedure
Induction of diabetes
The rabbit was held in a wooden restrainer specially designed for blood collection. Rabbits weighing 1500–2500 g were made diabetic by injecting 100–120 mg/kg body weight of alloxan monohydrate. Two grams of glucose/kg body weight dissolved in 10 cc of distilled water were administered orally to each rabbit before alloxan injection and 4 ml of 25% dextrose is administered i.v after alloxan injection to minimize the anticipated hypoglycemia.

The dose of alloxan monohydrate for each rabbit was selected very carefully not only based on the weight but also the general condition of the animal. The required dose was dissolved in 8 cc of distilled water in a sterile glass beaker and was injected into the marginal ear veins by a butterfly needle mounted on a 5–10 cc syringe contrary to the tuberculin syringe method. Eight days after administration of alloxan, the rabbits having a FBS of more than 200 mg/dl (normal blood glucose levels in alloxan-induced diabetic rabbits for 30 days was observed as follows.

RESULTS
Effect of supplementation of Vitamin C with metformin on blood glucose levels in alloxan-induced diabetic rabbits for 30 days was observed as follows.

DISCUSSION
Vitamin C, a water-soluble vitamin with antioxidant action, plays an important role in scavenging oxidative free radicals from various tissues [15]. Vitamin C is structurally similar to glucose and possibly can inhibit many chemical reactions such as non-enzymatic glycosylation of proteins (Maillard reactions) by competing with glucose [16,17]. High concentration of ascorbic acid can directly inhibit erythrocyte aldose reductase and provide a rationale for the use of oral Vitamin C supplements in diabetes.

Accordingly, based on these properties, the effect of Vitamin C has been investigated by supplementing with metformin in reducing blood glucose and HbA1c levels in the present study. The present preclinical study evaluated the effect of supplementation of Vitamin C on blood glucose and HbA1c levels in ‘Alloxan-induced diabetic rabbit model' for 30 days.

Analysis of blood glucose levels
Table 1 shows before (day 0) and after (day 30) treatment variations in mean blood glucose levels in each group of rabbits. ANOVA analysis followed by post hoc Tukey’s test explains the findings as follows:

- Mean blood glucose levels in the normal control group varied from 113.3 mg/dl to 117.5 mg/dl without much variation during the study with a mean percentage reduction being 3.71% (p = 0.4317).
- Mean blood glucose levels in standard group varied from 263.5 mg/dl to 159.2 mg/dl, which was statistically very highly significant (p < 0.0001***), and the mean percentage reduction was 39.58%.
- Mean blood glucose levels in Test 1 group varied from 284.3 mg/dl to 162.2 mg/dl, which was statistically highly significant (p = 0.0026**), and the mean percentage reduction was 42.95%.
- Mean blood glucose levels in Test 2 group varied from 286.0 mg/dl to 145.5 mg/dl, which was statistically very highly significant (p = 0.0003***), and the mean percentage reduction was 49.12%.

On day 0 (before treatment)
One-way ANOVA analysis showed that there was a highly significant mean difference (p = 0.0013**) between overall groups. Post hoc Tukey’s multiple comparison test showed a highly significant mean difference between the control group and other three groups, whereas no significance between standard, Test 1, and Test 2 groups.

On day 30 (after treatment)
One-way ANOVA analysis shown that there was no significant mean difference (p = 0.2206) between overall groups. Post hoc Tukey’s multiple comparison tests showed no significance between any of the groups.

The above analysis indicated that supplementation of Vitamin C 250 mg and 500 mg with metformin had improved the blood glucose levels than metformin alone, but not to the significant level.

Table 1: Statistical analysis (ANOVA) showing a comparison of blood glucose levels (mg/dl) between different groups on day 0 (before treatment) and day 30 (after treatment) followed by between groups Student’s t-test

| Samples collected at days       | Groups                  | mean±SD               | F-value | p-value |
|--------------------------------|-------------------------|-----------------------|---------|---------|
| Day 0 (before treatment)       | Normal control          | 113.3±1.15            | 7.69    | 0.0013**|
| Day 0 (before treatment)       | Standard                | 263.5±5.24            |         |         |
|                                | Test 1                  | 284.3±13.1            |         |         |
|                                | Test 2                  | 286.0±45.15           |         |         |
| Day 30 (after treatment)       | Normal control          | 117.5±0.89            | 1.601   | 0.2206  |
| Day 30 (after treatment)       | Standard                | 159.2±23.30           |         |         |
|                                | Test 1                  | 162.2±5.91            |         |         |
|                                | Test 2                  | 145.5±45.66           |         |         |

Post hoc Tukey’s multiple comparison tests

| Groups compared                | Day 0       | Day 30      |
|--------------------------------|-------------|-------------|
| Control group versus the standard | 5.020**     | 2.586       |
| Control group versus Test 1     | 5.717**     | 2.772       |
| Control group versus Test 2     | 5.772**     | 1.738       |
| Standard versus Test 1          | 0.6965      | 0.1862      |
| Standard versus Test 2          | 0.7522      | 0.848       |
| Test 1 versus Test 2            | 0.05572     | 1.034       |

**p = 0.0013
Analysis of HbA1c levels

Table 2 shows before (day 0) and after (day 30) treatment variations mean HbA1c levels in each group of rabbits and ANOVA analysis followed by post hoc Tukey’s test.

• Mean HbA1c levels in the normal control group varied from 2.18% to 2.20% without much variation during the study, and the mean percentage reduction was 4.54% (P = 0.0770).

• Mean HbA1c levels in standard group varied from 4.97% to 3.93%, which were statistically very highly significant (P < 0.0001***), and the mean percentage reduction was 22%.

• Mean HbA1c levels in Test 1 group varied from 5.07% to 3.82%, which were statistically very highly significant (P < 0.0001***), and the mean percentage reduction was 25.49%.

• Mean HbA1c levels in Test 2 group varied from 5.08% to 3.05%, which were statistically very highly significant (P < 0.0001***), and the mean percentage reduction was 40.12%.

On day 0 (before treatment)

One-way ANOVA analysis showed that there was a very highly significant mean difference (P = 0.0001*** between overall groups. Post hoc Tukey’s multiple comparison test showed very highly significant mean difference between the control group and other three groups, whereas no significance between standard, Test 1, and Test 2 groups.

On day 30 (after treatment)

One-way ANOVA analysis showed that there was a very highly significant mean difference (P = 0.0003*** between overall groups. Post hoc Tukey’s multiple comparison test showed very highly significant mean difference between the control group and standard group, highly significant mean difference between standard and Test 2 groups. There is no much difference between standard, Test 1, and similarly test group 1 and 2.

The above analysis indicated that supplementation of Vitamin C 500 mg with metformin has significantly improved the HbA1c levels than metformin alone, whereas Vitamin C 250 mg with metformin had better efficacy than that of metformin alone.

Bar diagram 1 shows mean blood glucose levels in different groups of rabbits on day 0 and day 30, as detailed in Table 1. It indicates that the blood glucose lowering effect in Test 1 and Test 2 groups is better than that of the standard group.

Bar diagram 2 shows percentage reductions of mean blood glucose levels in 30 days. Percentage reduction in Test 2 (49.12%) is greater than that of standard, whereas percentage reduction in Test 1 (42.95%), which is similar to that of standard (39.58%).

The results of this study are in agreement with previously published data showing betterment in glycemic control with Vitamin C supplementation [18,19]. Another study shown that there is an inverse relationship between Vitamin C and HbA1c levels indicating that supplementation with Vitamin C improves glycemic control [20] which can be indirectly correlated with the dose of Vitamin C in the present study.

Kukner et al. study showed that Vitamin C prevented glomerular damage seen in diabetes when combined with Vitamin E attributing to their antioxidant action. Supplements with Vitamin C increased urinary oxalate levels, but there is no evidence that they produce kidney stones and even up to 10 g; also, they do not exert any toxic or side effects [21].

Bar Diagram 1: Day-wise variation in blood glucose levels in various study groups. Y-axis represents blood glucose levels

Table 2: Statistical analysis (ANOVA) showing a comparison of HbA1c levels (%) between different groups on day 0 (before treatment) and day 30 (after treatment) followed by between groups Student t-test

| Samples collected at days | Groups                  | mean±SD       | F-value | p-value      |
|---------------------------|-------------------------|---------------|---------|--------------|
| Day 0 (before treatment)  | Normal control           | 2.13±0.53     | 32.22   | <0.0001***   |
|                          | Standard                | 4.97±0.46     |         |              |
|                          | Test 1                  | 5.07±0.78     |         |              |
|                          | Test 2                  | 5.08±0.65     |         |              |
| Day 30 (after treatment) | Normal control           | 2.28±0.69     | 9.830   | 0.0003***    |
|                          | Standard                | 3.93±0.29     |         |              |
|                          | Test 1                  | 3.82±0.75     |         |              |
|                          | Test 2                  | 3.05±0.56     |         |              |

Post hoc Tukey’s multiple comparison tests

| Groups compared          | Day 0     | Day 30    |
|--------------------------|-----------|-----------|
| Control group versus standard | 11.06***  | 6.753***  |
| Control group versus Test 1 | 11.45***  | 6.275**   |
| Control group versus Test 2 | 11.52***  | 3.138     |
| Standard versus Test 1    | 0.3973    | 0.4775    |
| Standard versus Test 2    | 0.4635    | 5.915**   |
| Test 1 versus Test 2      | 0.06621   | 3.138     |

***p=0.0001
Abdel-Wahab et al. study showed that the Vitamin C supplementation in obese hyperglycemic (ob/ob) mice significantly reduced glycated hemoglobin, plasma glucose, and insulin concentrations from 26% to 48% compared to untreated control ob/ob mice. Vitamin C supplementation also significantly reduced the total insulin content and extent of insulin glycation in the pancreas of ob/ob mice by 42-45% and also significantly reduced circulating glycated insulin by 80%. It demonstrates Vitamin C supplementation that can decrease insulin glycation and amelioration of the obesity-diabetes syndrome in ob/ob mice [22]. Cunningham et al. study showed that Vitamin C supplements of both 100 mg and 600 mg had significantly reduced sorbitol accumulation in erythrocytes in IDDM diabetic patients for 58 days. In vitro studies showed that ascorbic acid inhibited aldose reductase activity. Hence suggested, Vitamin C is superior over aldose reductase inhibitors [23].

The exact mechanism by which Vitamin C produces glycemic control is not known. Vitamin C scavenges important reactive oxygen species derived from glucose auto-oxidation, protein glycosylation, and polyol pathway that are involved in the generation of oxidative stress, implicated in the origin of both Types 1 and 2 DM. Vitamin C probably may also inhibit excess glucose uptake by cells of various tissues by competing for insulin. Within cells, Vitamin C possibly may compete with glucose for proteins and thus preventing protein glycation. Thus, Vitamin C supplementation for diabetic subjects may provide a simple means of preventing and ameliorating the complications of diabetes and helpful in maintaining tight glycemic control.

However, studies with a larger sample size and longer follow-up period together with the measurement of plasma Vitamin C levels and other related antioxidant levels are necessary. There is a need for further studies on complicated and uncomplicated Type 2 DM to elucidate the exact role of Vitamin C supplementation in Type 2 DM. Further studies should be done to understand its exact mechanism of action.

CONCLUSION

Finally, based on the above study, conclusions drawn are as follows:
- Supplementation of Vitamin C 500 mg to the metformin was superior in reducing HbA1c levels and blood glucose levels compared to metformin alone in rabbits.
- Supplementation of Vitamin C 250 mg to the metformin was beneficial in reducing HbA1c levels and blood glucose levels compared to metformin alone in rabbits.

Our study would suggest that treatment with Vitamin C with metformin was well tolerated. The cheaper cost and improvement in FBS and HbA1c in Type 2 DM after oral supplementation make it a particularly attractive therapeutic adjuvant in the treatment of Type 2 DM.

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AUTHORS’ CONTRIBUTIONS

Dr. P. Vijayakrishna raised the idea, designed the study, and participated in statistical analysis, Dr. A. Naveen drafted the manuscript; Dr. K. Indira participated in the study conception and manuscript revision; Dr. S. Shirisha involved in data collection and statistical analysis.

CONFLICTS OF INTEREST

All authors have none to declare.

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