

Original Research Article

Antidiabetic and antidiabetic activity of β -carotene on streptozocin-induced diabetic rats

ABSTRACT

Objective: β -carotene is a vital anti-oxidant, has capacity to reduce reactive oxidative stress, metabolic syndrome such as Type2 (T2) Diabetes Mellitus (DM) and prevent inflammation, obesity, Alzheimer and cardiovascular disease in human. In this study, we wanted to evaluate the efficacy of β -carotene on streptozotocin (STZ) induced Type2 (T2) DM rats where metformin used as positive control.

Methods: Diabetes was induced in Wister rats through the intraperitoneal administration of STZ (50 mg/kg b.w.). However anti- hyperlipidemic activities of β -carotene were evaluated by oral dose (10 mg/70 kg b.w.) once daily for 21 days. Metformin (12.1 mg/kg b.w.) was used as a positive control.

Results: Blood samples of rats were drawn by tail vein puncture and cardiac puncture to determine the fasting blood glucose (FBG) and serum level of total cholesterol (TC), triglycerides (TG), low-density lipoprotein (LDL), and high-density lipoprotein (HDL), respectively. The result of individual treatment of β -carotene and metformin is significant ($p < 0.001$) reversal of the diabetes induced increase in BGL, LDL, TC and TG, whereas significantly increased the STZ-induced decrease in HDL, compared to diabetic control.

Conclusion: The monotherapy of β -carotene have a significant antidiabetic and antihyperlipidemic effect and provides a scientific rationale for their use in antidiabetic therapy as a potential antioxidant.

Keywords: *Diabetes Mellitus, Antihyperglycemic, Antihyperlipidemic, Metformin, β -carotene.*

INTRODUCTION

β -carotene is most abundant carotene among all carotenoids in fruits, vegetables and human. Due to the high bioavailability of β -carotene, nearly 17% to 45% injected β -carotene remains intact in plant cells^[1]. It is a vital anti-oxidant and has capacity to reduce reactive oxidative stress and can prevent metabolic syndrome such as T2DM and vascular complication in human^[2-4].

Diabetic mellitus (DM) is most common chronic metabolic disorder. It has high mortality and morbidity rate due to significant effect on microvascular and macrovascular complications^[5]. The disease is rapidly spread and affect at every corner in the world. T2DM is a non- insulin dependent diabetes. More than 90%–95% patient of T2DM does not produce enough insulin or cannot properly use it^[6]. According to the report of World health organization (WHO), around 300 billion or more peoples will be affected by diabetic mellitus in the year of 2025 in all over the world. The DM-induced death reached at 28,065 or 3.61% of total deaths in Bangladesh^[7-8]. While the total focus of diabetes management on hyperglycemia as much as 75% of diabetic patients ultimately dying from vascular disease^[9]. T2DM showed severe lipid metabolism, abnormality persist even after optimal glycemc control has been achieved^[10].

The oxidative stress also induced the pathophysiology of T2DM and vascular complication. It was found that the prooxidant activity significantly increased and antioxidant has been decreased in T2DM. Consequently, the decrease production of reactive oxygen scavenging enzyme named superoxide dismutase (SOD) and catalase have been reported in diabetic animals^[11-16]. The significant increase in reactive oxygen and reduce the scavenging capacity of free radical act as trigger on lipid metabolism that were the leading causes of cardiovascular diseases^[17].

Metformin is most common, old and widely accepted drugs used for management and treatment of T2DM. It is remarkably reduced oxidative stress and dicarbonyl stress in human. Still now it is safe and effective as a mono therapy than combination therapy with other antidiabetic drugs and insulin^[18].

The antioxidant such as N-acetylcysteine, vitamin C, vitamin E and α -lipoic acid are effective in reducing diabetic complications. β -carotene act as a chelator of singlet oxygen and reacts with different species of free radicals (e.g., hydrogen peroxide). The conjugated double chain is responsible for eliminating singlet oxygen^[19-20]. However, our literature review shows that there has been no study on the β -carotene on STZ-induced rats to examine their antidiabetic and antidyslipidemic activities. Hence, this study was undertaken to investigate the effect of β -carotene and metformin on antidiabetic and antidyslipidemic activities in STZ-induced diabetic rats.

2. MATERIALS AND METHODS

2.1 Collection of drugs

The metformin and the β -carotene were the gift from Incepta Pharmaceuticals Ltd, Savar, Dhaka, Bangladesh. STZ was purchased from Sisko Research Laboratories Pvt. Ltd., Mumbai, India. Kits for the examination of high-density lipoprotein (HDL), low-density lipoprotein (LDL), total cholesterol (TC), triglycerides (TG) were obtained from Human, Wiesbaden, Germany. All the other chemicals and reagents were of the analytical grade.

2.2 Experimental animals and the lab environment

Three months old, twenty healthy male Wister albino rats (180-220 gm) were procured from Pharmacology Research Laboratory, Department of Pharmacy, Jahangirnagar University, Bangladesh. At the beginning of the experiments, all rats were adjusted in the laboratory environmental condition for one week. The research followed the procedure for laboratory animal care and use (NIH publication No.85-23, revised in 2004). The rats were fed with standard pellets collected from International Centre for Diarrhoeal Disease Research (ICDDR), Bangladesh and fresh drinking water ad libitum during this period. Under normal atmospheric temperature (25 ± 2 °C), humidity and 12 hours light- dark cycles were controlled according to the animal care and welfare guidelines^[21].

2.3 Experimental design

The male Wister albino rats (20) were randomly grouped into 4 groups (A, B, C and D) were 5 rats in each (n=5). The administration of both β -carotene and metformin was given by the oral route.

Group A (Gr. A): Non-diabetic control rats (NC) treated with physiological saline.

Group B (Gr. B): STZ-induced diabetic control rats (DC) received physiological saline.

Group C (Gr. C): Diabetic rats treated with metformin (daily administered 850 mg/70 kg b.w.)^[22].

Group D (Gr. D): Diabetic rats treated with β -carotene (daily administered 10 mg/70 kg b.w.)^[23].

2.4 Induction of T2DM and the measurement of fasting BGL

For the damage of insulin producing pancreatic islets, freshly prepared single dose of STZ (45mg/kg b.w. were dissolved in 0.01 M citrate buffer (pH: 4.5)) were injected intraperitoneally in overnight fasted rats. After the STZ injection, rats were given drinking water supplemented with sucrose (15 g/L) for 48 h to limit early mortality^[23]. Diabetes was confirmed in rats after 3 days of STZ injection. The hypoglycemic

condition was measured by the fasting serum blood glucose level using one-touch glucometer (Glucox TD-4183, Germany). The blood glucose levels >200 mg/dL were considered as diabetic and selected for the study. After three weeks (day 21) treatment with metformin and β -carotene in STZ-induced diabetic rats, the rats were fasted overnight and sacrificed. Blood samples of rats were drawn by tail vein puncture to determine the fasting blood glucose and cardiac puncture for lipid profile

2.5 Biochemical Analysis

At the end of the experiment, the overnight fasted rats were anesthetized with chloroform. 3 mL to 5 mL of fasting blood sample was collected from thoracic artery by a heparinized syringe. After centrifugation at 4000 RPM for 10 minutes, the serum was separated for lipid measurement. Total cholesterol (TC), triglycerides (TG), high density lipoprotein (HDL)-cholesterol and low density lipoprotein (LDL)-cholesterol were determined in the serum using commercial kit reagents (Human, Wiesbaden, Germany), following the company's instruction and the absorbance was read using a UV-Vis spectrophotometer (Shimadzu UV-1280, Kyoto, Japan).

2.6 Statistical Analysis

The average of five samples was reported as the measured value with standard deviation. The difference among the groups was calculated by one-way analysis of variance (ANOVA) with multiple comparison tests, where applicable. The difference between two groups was measured by the student t-test. A p-value at <0.05 was considered to be statistically significant.

3. RESULT

The effects of metformin and β - carotene on the parameters of blood glucose level and lipid profile (TC, TG, LDL, HDL) were performed on streptozotocin induced diabetes rats.

3.1 Hypoglycemic effects of metformin and β - carotene in STZ- induced diabetic rats:

The effect of metformin and β - carotene were separately determined on FBG level in STZ- induced diabetic rats groups at day 0 and day 21 were shown in figure1. At the day 0, the FBG level of STZ - induced diabetic (DC) rats was significantly ($p < 0.05$) higher (20.42 ± 0.816 mmol/L) than those of non-diabetic (NC) rats (5.68 ± 0.159 mmol/L). At the day 21, the fasting BGL of DC group remained higher and significant (18.66 ± 0.51 mmol/L) compared to NC rat groups. After the treatment of β - carotene, FBG level became reduced (13.88 ± 0.816 mmol/L) remarkably in the same manner as that with the standard drug metformin (7.48 ± 0.11 mmol/L)

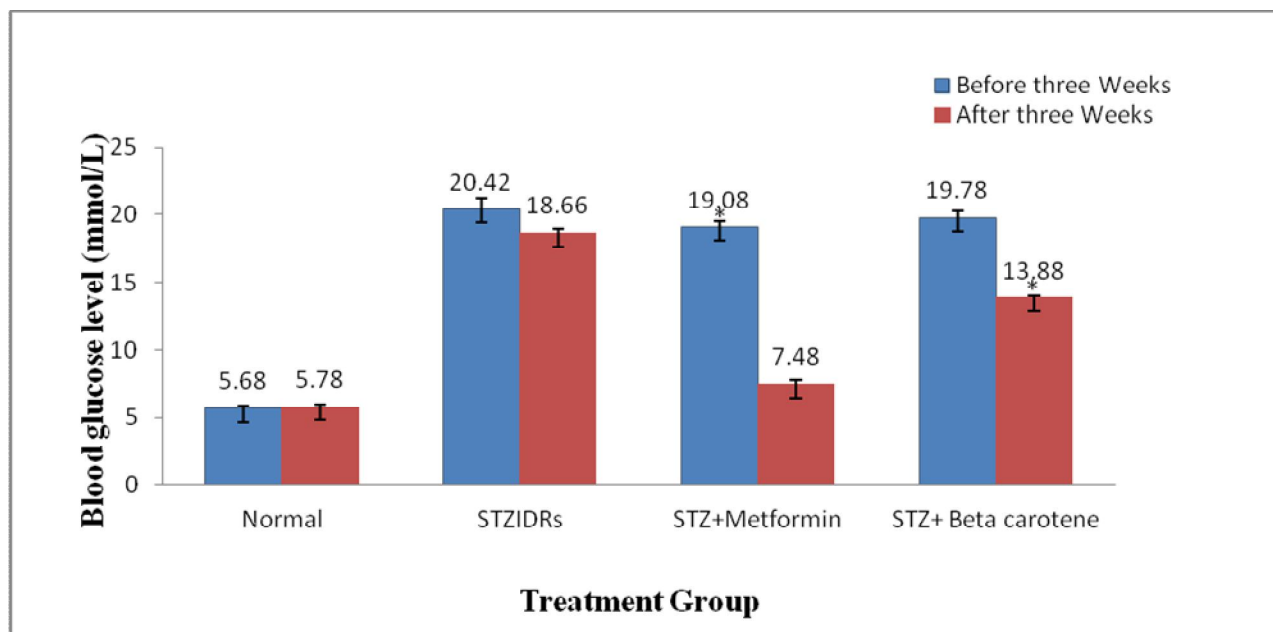


Figure: 1 The hypoglycemic effect of metformin and β - carotene were determined on FBG level in STZ- induced diabetic rats groups and compared with diabetic control (Gr. B) and normal control group (Gr.A) at day 0 and day 21.

*n=5, the values are expressed as mean \pm SEM. $P < 0.05$ compared with normal control group (Gr. A). * $P < 0.05$ compared with diabetic control group (Gr. B).*

3.2 The effect of metformin and β - carotene on Lipoprotein (LDL and HDL) in STZ- induced diabetic rats:

Measurement the effect of metformin and β - carotene of serum lipoprotein (LDL and HDL) on STZ- induced diabetic rats is shown in figure 2. The serum concentration of LDL levels increased and HDL levels were decreased as compared with diabetic controlled rats group (Gr.B). Repeated treatment by metformin and β - carotene were able to decrease ($P < 0.05$) the serum concentration of LDL (87.26 ± 0.021 mg/dL) whereas increase ($P < 0.05$) the level of HDL (30.42 ± 0.033 mg/dL) as compared with STZ- induced diabetic control (24.66 ± 0.03 mg/dL) rats.

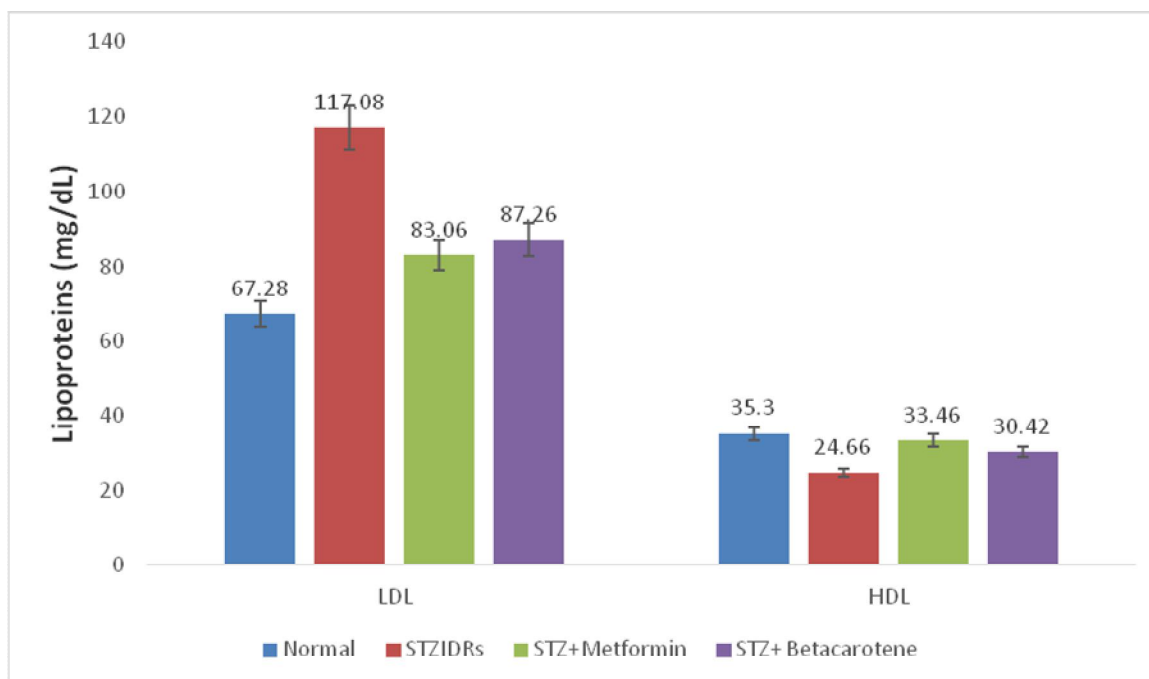


Figure: 2 Effect of metformin and β -carotene on Lipoproteins (LDL and HDL) in normal (Gr.A) and STZ-induced diabetic (Gr. B) rats after three weeks of treatment.

*n=5, the values are expressed as mean \pm SEM. † $P < 0.05$ compared with normal control (Gr. A) group. * $P < 0.05$ compared with diabetic control (Gr.B) group.*

3.3 The effect of metformin and β - carotene on the concentration of serum TC and TG in STZ- induced diabetic rats:

Determination the effect of metformin and β - carotene on the concentration of serum TC and TG level was determined on STZ- induced diabetic rats and is found in figure 3. The serum concentration of TC and TG levels were decreased as compared with diabetic controlled rats group (Gr. B). Repeated treatment by metformin and β - carotene had ability to decrease ($P < 0.05$) the serum concentration of TG (137.16 \pm 0.027 mg/dL) and TC (196.34 \pm 0.025 mg/dL) level as compared with the TG and TC level on STZ- induced diabetic control (Gr. B) rats.

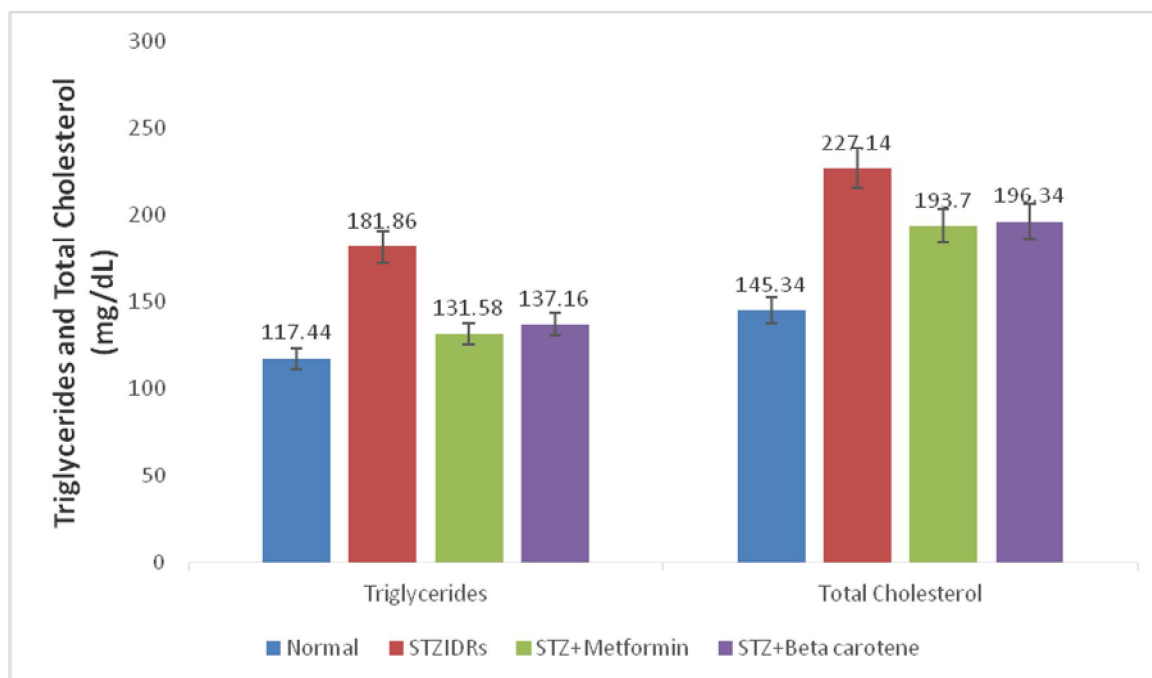


Figure: 3 Effect of metformin and β -carotene on triglyceride (TG) and total cholesterol (TC) in normal and STZ-induced diabetic rats after three weeks of treatment.

*n=5, the values are expressed as mean \pm SEM. † $P < 0.05$ compared with normal control group (Gr. A) * $P < 0.05$ compared with diabetic control group (Gr. B).*

4. DISCUSSION:

Diabetes is an abundant metabolic disorder found all over the world. Insulin resistance and hyperglycemia could associate with vascular complication. Non- insulin dependent T2DM leads to the change the plasma lipoprotein concentration^[24]. We investigated the antidiabetic and antihyperlipidemic activity of β -carotene on STZ- induced diabetic rats.

From our short-term study confirmed that, the effect of the oral intake of β -carotene in a single dose (10 mg/70 kg b.w) in STZ- induced diabetic rats was significantly reduced the fasting BGL level. The values of fasting BGL after 21 days became turn down (13.88 ± 0.816 mmol/L) remarkably in the same manner as that with the standard drug metformin (7.48 ± 0.11 mmol/L). Still now, metformin is an old and magical first line of oral drug used worldwide for the treatment of T2DM. It can improve the endothelial dysfunction, oxidative stress, insulin resistance, lipid profiles, fat redistribution and increased peripheral glucose uptake and utilization^[25], reduced intestinal absorption of glucose^[26] and translocation of glucose transporter GLUT4 from the circulation^[27]. Beside this, long term treatment of metformin has some side effects also^[28]. To avoid the complication of metformin researcher, hunt natural or herbal product or those substances which has ability to have antihyperglycemic and antihyperlipidemic activity. In this sense, the therapies based on an antioxidant β -carotene had the ability to improve the efficacy of glycemic control promoted by metformin to avoid the transition of insulin therapy is of great interest^[29]. Also, it was found that the mono therapy of antioxidant; like as N-acetylcysteine, vitamin C, and α -lipoic acid had capability to reduced diabetic complications^[30]. In this consequence, the novel therapies based on single dose of β -carotene have an ability to control hyperglycemia might be used as antihyperglycemic therapy.

After day 21, we found that monotherapy of β -carotene reduce the serum fasting BGL with significant decrease in the serum lipid profiles, such as LDL, TC and TG level. However, monotherapy of β -carotene slightly increased the level of serum HDL against the diabetic control (Gr. B). The similar results also found that β -carotene can reduce low-density lipoprotein cholesterol (LDL-c), and very-low-density lipoprotein cholesterol (VLDL-c); and an increase in high-density lipoprotein cholesterol (HDL-c).

Moreover, it can improve insulin resistance and insulin binding receptor protein on cell wall in human ^[31-34], also noted that the bioavailability of β -carotene is low through the gut, though rodents (rats and mice) can readily convert β -carotene to vitamin A ^[35]. Therefore, administration of β -carotene at higher doses required needs more attention in our future research.

High level of glucose concentration in blood stimulates ROS production that leads to increased oxidative stress. The antioxidants play vital role against ROS through the antioxidant defense system of the body. Also, diabetes and the increased oxidative stress related protein are the major risk factors for the development of cardiovascular disease ^[36]. The β -carotene is reported to reduce oxidative stress and increase antioxidant activity, mainly glutathione-related defense systems in lead-exposed worker ^[37]. Plasma triglyceride and total cholesterol was increased by diabetes, but it was reduced by β -carotene. It has been reported that the higher plasma concentration of β -carotene could improved insulin resistance and help to reduction of body adiposity ^[38-39]. Also, that the administration of β -carotene suppresses the elevation of lipid peroxidation (LPO) and reduces the symptoms of DM in the STZ-induced diabetic rats ^[40].

In our experiment, we have selected β -carotene as a mono therapy, as β -carotene was reported to be defensive against diabetes via improving insulin sensitivity ^[41] and also shielding against DR ^[42] and cardiovascular diseases ^[43]. Justifying it with the findings ^[44-45], our results showed that β -carotene markedly reduced serum triglycerides, total cholesterol and other lipid indices of diabetic risk in diabetic rats and thus suggest antihyperglycemic and antidyslipidemic potentials of β -carotene in reducing the incidence of diabetic vascular complications through the normalization of lipid metabolism in diabetic condition. More work is required to truly understand how β -carotene works at the molecular level in individuals with T2DM.

5. CONCLUSION

β -carotene is an anti-oxidant and it had an effect on oxidative stress in diabetic rats. From our experiment it may be concluded that β -carotene had anti-glycemic activity on STZ- induced diabetic rats as well as anti- hyperlipidic activity by reducing the glycooxidative stress in blood. Therefore, the β -carotene it is very promising antidiabetic and antihyperlipidemic drug like as traditional antidiabetic drugs metformin. Further studies are needed to know how β -carotene works at molecular level.

7. REFERENCES

- [1] Shete V, Quadro L. Mammalian metabolism of β -carotene: gaps in knowledge. *Nutrients*. 2013; 27;5(12):4849-68. doi: 10.3390/nu5124849.
- [2] Mounien L, Tourniaire F, Landrier JF. Anti-Obesity Effect of Carotenoids: Direct Impact on Adipose Tissue and Adipose Tissue-Driven Indirect Effects. *Nutrients*. 2019; 11;11 (7):1562. doi: 10.3390/nu11071562.
- [3] Rodriguez-Concepcion M, Avalos J, Bonet ML, Boronat A, Gomez-Gomez L, Hornero-Mendez D, Limon MC, Meléndez-Martínez AJ, Olmedilla-Alonso B, Palou A, Ribot J, Rodrigo MJ, Zacarias L, Zhu C. A global perspective on carotenoids: Metabolism, biotechnology, and benefits for nutrition and health. *Prog Lipid Res*. 2018; 70:62-93. doi: 10.1016/j.plipres.2018.04.004.
- [4] Conrad O, Perera & Gan Mei Yen, Functional Properties of Carotenoids in Human Health, *International Journal of Food Properties*, 2007; 10:2, 201- 230, DOI: [10.1080/10942910601045271](https://doi.org/10.1080/10942910601045271)
- [5] Patel DK, Kumar R, Prasad SK, Sairam K, Hemalatha S. Antidiabetic and in vitro antioxidant potential of *Hybanthus enneaspermus* (Linn) F. Muell in streptozotocin-induced diabetic rats. *Asian Pac J Trop Biomed*. 2011;1(4):316-22. doi: 10.1016/S2221-1691(11)60051-8.

- [6] Li WL, Zheng HC, Bukuru J, DeKimpe N. Natural medicines used in traditional Chinese medical system for therapy of diabetic mellitus. *Journal of Ethnopharmacology*, 2004; 92, 1-21.
- [7] Sy GY, Cissé A, Nongonierma RB, Sarr M, Mbodj NA, Faye B. Hypoglycaemic and antidiabetic activity of acetonic extract of *Vernonia colorata* leaves in normoglycaemic and alloxan-induced diabetic rats. *J Ethnopharmacol*. 2005; 98(1-2):171-5.
- [8] Web World Health Rankings. Bangladesh: Diabetes Mellitus. Available from: <https://www.worldlifeexpectancy.com/bangladesh-diabetes-mellitus>. [Last accessed on 10 Apr 2021]
- [9] Basha B, Samuel SM, Triggler CR, and Ding H. Endothelial Dysfunction in Diabetes Mellitus: Possible Involvement of Endoplasmic Reticulum Stress? *Exp Diabetes Res* 2012; 2012:481840.
- [10] O'Brien T, Nguyen TT, Zimmerman BR. Hyperlipidemia and diabetes mellitus. *Mayo Clin Proc*. 1998; 73(10):969-76. doi: 10.4065/73.10.969. PMID: 9787748
- [11] Beladi Mousavi SS, Nasri H, Rafieian-Kopaei M, Tamadon MR. Metformin improves diabetic kidney disease. *J Nephroarmacol*. 2012; 1(1):1-2.
- [12] Ayodhya S, Kusum S, Saxena A. Hypoglycaemic activity of different extracts of various herbal plants. *Int J Res Ayurveda Pharm* 2010; 1:212.
- [13] Abdollahi M, Farshchi A, Nikfar S, Seyedifar M. Effect of chromium on glucose and lipid profiles in patients with type 2 diabetes; a meta-analysis review of randomized trials. *J Pharm Pharm Sci*. 2013; 16(1):99-114. doi: 10.18433/j3g022.
- [14] Rafieian-Kopaei M, Nasri H. Ginger and diabetic nephropathy. *J Renal Inj Prev*. 2013; 1;2(1):9-10. doi: 10.12861/jrip.2013.05.
- [15] Nasri H, Rafieian-Kopaei M. Tubular Kidney Protection by Antioxidants. *Iran J Public Health*. 2013; 42(10):1194-6.
- [16] Ardalan MR, Sanadgol H, Nasri H, Baradaran A, Tamadon MR, Rafieian-Kopaei R. Vitamin D therapy in diabetic kidney disease; current knowledge on a public health problem. *J Parathyroid Dis* 2014; 2:15-7.
- [17] Basha B, Samuel SM, Triggler CR, Ding H. Endothelial dysfunction in diabetes mellitus: possible involvement of endoplasmic reticulum stress? *Exp Diabetes Res*. 2012; 2012:481840. doi: 10.1155/2012/481840. Epub 2012 Feb 28.
- [18] Rojas LB, Gomes MB. Metformin: an old but still the best treatment for type 2 diabetes. *Diabetol Metab Syndr*. 2013; 15;5(1):6. doi: 10.1186/1758-5996-5-6.
- [19] Namitha KK, Negi PS. Chemistry and biotechnology of carotenoids. *Crit Rev Food Sci Nutr*. 2010; 50(8):728-60. doi: 10.1080/10408398.2010.499811.
- [20] Fiedor J, Burda K. Potential role of carotenoids as antioxidants in human health and disease. *Nutrients*. 2014; 27;6(2):466-88. doi: 10.3390/nu6020466.
- [21] Roy P, Islam M, Islam MS, Rashid M. Beneficial Effects of Combination Therapy of Sitagliptin and β -carotene Drugs on Streptozotocin-induced Diabetic Rats. *Bangladesh pharm J* 2020; 23: 87-95
- [22] Guide for the Care and Use of Laboratory Animals. 8th Edition. Washington DC: The National Academies Press, 2011.

- [23] Zannah S, Islam M, Ali Y, Asaduzzaman M, Sarwar MS, Al Bari AA, Rashid M. Antidiabetics in Combination with Hydroxychloroquine Improve Antioxidant and Hepatoprotective Activities in Alloxan-Induced Diabetic Rats. *Bangladesh pharm J* 2015;18: 72-77.
- [24] Ira J. Goldberg, Diabetic Dyslipidemia: Causes and Consequences, *The Journal of Clinical Endocrinology & Metabolism*, 2001; 86(1): 965–971, <https://doi.org/10.1210/jcem.86.3.7304>
- [25] Viollet B, Guigas B, Garcia NS, Leclerc J, Foretz M, Andreelli F: Cellular and molecular mechanisms of metformin: an overview *Clin Sci (Lond)* 2012;122: 253-70.
- [26] Minamii T, Nogami M, Ogawa W. Mechanisms of metformin action: In and out of the gut. *J Diabetes Investig* 2018; 9: 701-703
- [27] Yang J, Holman GD. Long-term metformin treatment stimulates cardiomyocyte glucose transport through an AMP-activated protein kinase-dependent reduction in GLUT4 endocytosis. *Endocrinology* 2006; 147: 2728-36.
- [28] Ghosal S and Ghosal S, The Side Effects of Metformin - A Review, *J Diabetes Metab Disord* 2019; 6: 030 DOI: 10.24966/DMD-201X/100030
- [29] Khanam, K., Rajia, S., Yeasmin, M., Morshed, M. and Haque, R. "Synergistic Effect of β -Carotene and Metformin on Antihyperglycemic and Antidyslipidemic Activities in Streptozotocin-Induced Diabetic Rats", *Journal of Pharmaceutical Research International*, 2021; 33(51A): 79-87. doi: 10.9734/jpri/2021/v33i51A33471.
- [30] Ruhe RC, McDonald RB. Use of Antioxidant Nutrients in the Prevention and Treatment of Type 2 Diabetes. *J Am Coll Nutr* 2001;20:363S-9.
- [31] Amengual J, Gouranton E, Helden YGJ, Hessel S, Ribot J, Kramer E, Kiec-Wilk B, Razny U, Lietz A, Wyss A, et al. Beta-carotene reduces body adiposity of mice via BCMO1. *PLoS ONE*, 2011;6: e20644.
- [32] Beydoun MA, Chen X, Jha K, Beydoun HA, Zonderman AB, Canas JA. Carotenoids, vitamin A, and their association with the metabolic syndrome: A systematic review and meta-analysis. *Nutr. Rev.* 2019; 77: 32–45.
- [33] Canas JA, Lochrie A, McGowan AG, Hossain J, Schettino C, Balagopal PB. Effects of mixed carotenoids on adipokines and abdominal adiposity in children: A pilot study. *J. Clin. Endocrinol. Metab.* 2017; 102: 1983–1990.
- [34] Asemi Z, Alizadeh SA, Ahmad K, Goli M, Esmailzadeh A. Effects of beta-carotene fortified symbiotic food on metabolic control of patients with type 2 diabetes mellitus: A double-blind randomized cross-over controlled clinical trial. *Clin. Nutr.* 2016; 35: 819–825.
- [35] Szuszkiewicz-Garcia MM, Davidson JA. Cardiovascular disease in diabetes mellitus: Risk factors and medical therapy. *Endocrinol Metab Clin of N Am* 2014;43: 25-40.
- [36] Braunwald E. Biomarkers in heart failure. *N Engl J Med.* 2008; 358(20):2148-59. doi: 10.1056/NEJMra0800239.
- [37] Kasperczyk S, Dobrakowski M, Kasperczyk J, Ostalowska A, Zalejska-Fiolka J, Birkner E. Beta-carotene reduces oxidative stress, improves glutathione metabolism and modifies antioxidant defense systems in lead-exposed workers. *Toxicology and Applied Pharmacology* 2014;280:36- 41.

[38] Canas JA, Lochrie A, McGowan AG, Hossain J, Schettino C, Balagopal PB. Effects of mixed carotenoids on adipokines and abdominal adiposity in children: A pilot study. *J. Clin. Endocrinol. Metab.* 2017; 102: 1983–1990. doi: 10.1210/jc.2017-00185.

[39] Canas JC, Damaso L, Altamare A, Killen K, Hossain J, Balagopal PB. Insulin resistance and adiposity in relation to serum β -carotene levels. *J. Pediatr.* 2012;161:58–64. doi: 10.1016/j.jpeds.2012.01.030.

[40] Furusho T, Kataoka E, Yasuhara T, Wada M, Innami S. Administration of betacarotene suppresses lipid peroxidation in tissues and improves the glucose tolerance ability of streptozotocin-induced diabetic rats. *Int J Vitam Nutr Res* 2002;72:71-6.

[42] Roohbakhsh A, Karimi G, Iranshahi M. Carotenoids in the treatment of diabetes mellitus and its complications: a mechanistic review. *Biomed Pharmacother* 2017;91:31-42

[43] Fatahlipour M, Hadis Fathalipour H, Nowrouzi-Sohrabi OSP, Hossein M, Hassanipour S. Therapeutic Role of Carotenoids in Diabetic Retinopathy: A Systematic Review Diabetes, Metabolic Syndrome and Obesity: Targets and Therapy 2020;13:2347-58.

[44] Gammone MA, Pluchinotta FR, Bergante S, Tettamanti G, D'Orazio N. Prevention of cardiovascular diseases with carotenoids. *Front Biosci* 2017;9:165-71.

[45] Seo JS, Lee KS, Jang HJ, Quan Z, Yang KM, Burri BJ. The effect of dietary supplementation of β -carotene on lipid metabolism in streptozotocin-induced diabetic rats. *Nutrition research*, 2004; 24:1011-21

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