Original Research Article

A study on serum lipid and malondialdehyde levels among diabetic patients

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ABSTRACT

Background: Elevated levels of lipid peroxide in diabetes mellitus may be due to the alteration of function of erythrocytes membrane. This inhibits the activity of superoxide dismutase enzyme leading to accumulation of superoxide radicals which cause the maximum lipid peroxidation and tissue damage in diabetes. The objectives was to study population and matched control group.

Methods: This study was done among 50 NIDDM, 50 IDDM and 50 controls at Thanjavur Medical College, Tamil Nadu, India for a period of one year at the Department of Diabetology after getting the informed consent and IEC clearance. This study included all ambulatory NIDDM and IDDM patients without any complications. The following investigations like serum malondialdehyde, blood sugar, HBA1C, serum lipid profile, blood urea, serum creatinine, urine albumin and sugar were done by standardized procedures and reagents after getting the detailed history and examination.

Results: Among NIDDM group 78% were between 6.4 to 8 categories whereas in IDDM group only 28% were in this 6.4 to 8 category (HBA1C). Comparison of serum MDA values among three groups were done by ANOVA with two groups separately and it was highly significant. Multiple comparison of mean difference of MDA and lipid values among all the three groups showed statistically significant results with p value at 0.05.

Conclusions: Lipid profile is increased in poor glycemic controlled patients (both IDDM and NIDDM patients) and it is reflected in high serum malondialdehyde levels.

Keywords: Glycemic status, HBA1C, IDDM, NIDDM, Serum lipid levels, Serum malondialdehyde

INTRODUCTION

Diabetes mellitus is a group of metabolic disease characterized by hyperglycaemia resulting from defects in Insulin secretion, Insulin action or both.¹ It is a complex disease where carbohydrate, protein and fat metabolism is impaired. Diabetes is an “ice berg” disease. The number of cases of diabetes worldwide is estimated to be around 150 million. It is estimated that 20 percent of the current global diabetic population resides in South-East Asia Region. In India, the prevalence of disease in adults was found to be 2-4 percent in rural and 4-11.6 percent in urban dwellers. High frequencies of impaired
glucose tolerance, shown by studies ranging from 3.6-9.1 percent indicate the potential for further rise in prevalence of DM in the coming decades.3

**Classification**

- Type 1 Diabetes
  - Immune mediated,
  - Idiopathic.
- Type 2 Diabetes
  - Other specific types,
  - Gestational Diabetes Mellitus (GDM),
  - Impaired Glucose Tolerance (IGT).

IDDM onset is typically abrupt and is usually seen in individual less than 30 years.5 Immune mediated and β cells of pancreas are destroyed, usually associated with ketosis, Exogenous insulin is required to reverse the catabolic state. NIDDM is more common than IDDM, gradual in onset and occurs mainly in the middle aged and elderly.3

Diabetes is better known for its complications affecting the vascular system, kidney, retina, lens, peripheral nerves and skin which are extremely costly in terms of longevity and quality of life.6

Lipid peroxidation is elevated in Diabetes.7 Diabetes is usually accompanied by increased production of free radicals or reactive oxygen species which produces oxidative stress.7 The occurrence of free radical induced lipid peroxidation causes considerable change in the cell membrane.8 Peroxidation of lipid membrane has been related to the pathogenesis of many degenerative diseases such as atherosclerosis.9 Atherosclerosis is the most common complication of diabetes.10

Free radicals damage lipids by initiating a process called lipid peroxidation.11

The decomposition of lipid peroxides forms many cytotoxic compounds like malondialdehyde (MDA).

So, oxidative stress can be measured by monitoring the changes in malondialdehyde.6,7 Degree of lipid peroxidation was measured in terms of MDA.

The objectives were to assess the level of lipid peroxide among diabetic patients and to find out the correlation between lipid peroxide with glycaemic control (HBA1C).

**METHODS**

Participants of the study group were selected from the outpatient’s population of Department of Diabetology, Thanjavur Medical College, Thanjavur, Tamil Nadu, India. 100 patients were selected for this study. Out of which 50 patients belong to NIDDM and 50 to IDDM group. 50 persons served as healthy control. Informed consent was obtained before starting the study and IEC clearance was also, obtained.

**Inclusion criteria**

All ambulatory NIDDM and IDDM patients without any complications.

**Exclusion criteria**

Smokers, alcoholics, renal failure, bronchial asthma, history suggestive of complications of DM (angioopathy, cardiopathy, retinopathy, nephropathy).

**Procedure**

Detailed history and complete clinical examination were done in all the cases. For all the patients, fasting and post prandial blood samples and fasting urine samples were collected. For blood sugar estimation, blood collected in fluorinated tube. For other investigations in plain tube samples were collected. The following investigations like serum malondialdehyde, Blood sugar (fasting and post prandial), HbA1C, serum lipid profile, blood urea, serum creatinine, urine albumin and sugar. All the procedures for measurements of above biochemical values were done using standardized reagents and instruments.

Appropriate statistical methods like descriptive statistics, chi-square tests and ANOVA were applied using SPSS software version 20.

**RESULTS**

Totally 150 study participants were studied of which 50 from NIDDM, 50 from IDDM and 50 as controls. Majority were from 30 to 50 years age group with equal number of males and females and 90% of the study participants in all the three groups had BMI between 18 to 25 categories. Glycemic status with HBA1C values classified between less than 6.4, 6.5 to 8 and more than 8. Among NIDDM group 78% were between 6.4 to 8 categories whereas in IDDM group only 28% were in these 6.4 to 8 categories.

**Serum malondialdehyde result analysis**

Comparison of serum MDA values among three groups were done by ANOVA with two groups separately shown in Tables 1 and 2 and it was highly significant. The difference of MDA values is more pronounced among control and diabetic groups of patients (p <0.0001). Similarly, the MDA mean values among IDDM and NIDDM also showed the significant difference but not much in control group. Multiple comparison of mean difference of MDA values among all the three groups control, NIDDM and IDDM simultaneously showed significant results with p value at 0.05 (Table 3).
The statistical test between three groups in Table 3 showed the mean difference was 1.233 with the 95% CI range between 0.99 to 1.48 with highly significance. This clinically correlates with the poor glycemic control among diabetes patient’s high values of MDA than the control group (p <0.001).

The scatter plot between serum lipid levels and HbA1C levels shows that both the variables are mutually increasing as the HbA1c levels are more than 7 which suggests that the lipid levels are affected by the glycemic status among diabetic individuals.

Figure 2 depicts the TC levels with the glycemic status (HBA1c), predicts the relationship between both the variables. When the patients are going in for poor glycemic control the lipid peroxidation metabolite that is MDA and total cholesterol values increase which is the predictor for poor glycemic status among diabetic patients.

**DISCUSSION**

The mean value of plasma MDA is high in diabetic patients when compared to control group. Increased lipid peroxidation in diabetes mellitus is due to excess formation of free radicals. Hyperglycaemia in diabetics causes increased glycation of protein which itself act as a source of free radicals. Metabolic derangements in...
diabetes lead to an increase in concentration of oxidizable substrates and compromised detoxification pathways. The study shows that cases on insulin as therapeutic regime (IDDM) had lower mean MDA level (4.46 μmol/L) as compared to those on oral hypoglycaemics (NIDDM) (4.8 μmol/L) indicating lesser level of oxidative stress in diabetics on insulin. Considering MDA levels among cases on the basis of their glycaemic status, significant correlation is seen between well controlled and poorly controlled diabetics (both in IDDM and NIDDM). MDA is higher in individuals with poor glycaemic control compared to good glycaemic control. For every 1% reduction in HbA1C, one can expect 35% reduction in microvascular complications, which can be attributed to decrease in oxidative stress on treatment.13 The metabolic parameters such as total cholesterol, triglycerides, LDL and VLDL values were more in diabetic groups than the control groups. Mean value of serum HDL is decreased in diabetic group compared to control. Most common lipid disorder observed in DM is the presence of high plasma triglyceride and low HDL cholesterol.14 Insulin is the principal antilipolytic regulator, acting on hormone sensitive lipase. Without its action as in DM, lipolysis in adipose tissue is increased. As a result, there is increased availability of NEFAS for re-esterification in the liver to produce more triglycerides. Lipoprotein lipase activity is less in insulin deficiency resulting in diminished triglyceride clearance, impaired lipolysis of VLDL and reduced formation of HDL particles.15 Insulin increases the number of LDL receptor. In insulin deficiency, the level of LDL receptors is low, which causes the increase in LDL cholesterol. LDL oxidation plays an important role in atherogenesis.16-25 The uptake of LDL by macrophage (to form foam cell) is increased by oxidation of LDL, Derivatization of ApoB by glycosylation and reaction with malondialdehyde.

**CONCLUSION**

Oxidative stress is observed more in diabetes mellitus patients. Lipid peroxidation is the marker for oxidative stress, and it is statistically significant in this study. Compared to NIDDM, IDDM patients with good glycemic control have low level of serum malondialdehyde.

Lipid profile is increased in poor glycemic controlled patients (both IDDM and NIDDM patients) and it is reflected in high serum malondialdehyde levels. It is found to be statistically significant in this study group. Strict glycemic control is needed for reducing oxidative stress in diabetes mellitus patients and to prevent its complications.

Further scope of this study is extending its size and follow up for assessing Diabetes mellitus complications in various sub groups. Also, the effect of starting either moderate or high-intensity statins based on the patient’s risk profile should be studied intensively.

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**Conflict of interest:** None declared

**Ethical approval:** The study was approved by the Institutional Ethics Committee

**REFERENCES**

1. Bishop ML, Fody EP, Schoeff LE. Clinical Chemistry Technique: Principle correlations. 6th ed. Woltes Kluwee (India) New Delhi; 2009: 13;314.
2. Altomare E, Vendemiale G, Chicco D, Proacci V, Cirelli F. Increased lipid peroxidation in type 2 poorly controlled diabetic patients. Diabete & metabolism. 1992;18(4):264-71.
3. K Park. Park’s text book of preventive and social Medicine. 20th ed. 2009;6:342-343.
4. Burtis CA, Ashwood ER, Bruns DE. Tietz text book of clinical chemistry and molecular diagnostics. 2012; 25: 854.
5. Anderson SC, Cockayne S. Clinical chemistry concept and applications. 2003: 10; 159.
6. Rask-Madsen C, King GL. Vascular complications of diabetes: mechanisms of injury and protective factors. Cell Metab. 2013;17(1):20-33.
7. Lester Packer, Peter Rosen, Chler HJT, King GL, Angelo A. Antioxidants in diabetes management. UNESCO MCBN 15A. 2000;3:42-5.
8. Agarwal S, Banejee S, Chattejee SN. Effects of oxygen and ferrous sulphate induced lipid peroxidation in liposomal membrane. Ind J Biochem Biophysics. 1985;21:331-4.
9. Chatterjee SN, Agarwal S, Amikumar, Membrane lipid peroxidation and its pathological consequence. Ind J Biochem Biophysics. 1988;25:31.
10. Steiner G. Artherosclerose, the major complication of diabetes. In: urica M, Hollenberg CH, Steriner G, eds. Comparison of type 1 and type 11 Diabetes: Similarities and Dissimilates in etiology, Pathogenesis and Complications. New York: Plenum press: 1985; 277-297.
11. Marshall WT, Bangert SK. Clinical Biochemistry, Metabolic and Clinical aspects, 2nd ed. Churchill Livingstone; 2008: 946-947.
12. Suryawanshi NP, Bhutey AK, Nagdeote AN, Jadhav AA, Manoorkar GS. Study of lipid peroxide and lipid profile in Diabetes Mellitus. Ind J Clin Biochem. 2006;21:126-30.
13. Sayday SH, Ebarhardt, MS, Loria cm. Age and burden of death attributable to diabetes in the united states. Am Epideiol. 2002;156:714-9.
14. Harmel AP, Mathur R. Davinson’s diabetes mellitus diagnosis and treatment, 5th ed. 2004; 8: 246.
15. Cathcard MK, Folcik VA. Lipoygenases and atherosclerosis. Protection versus pathogenesis. Free Radic Biol Med. 2000;28:1726-34.
16. Kuhn H, Chan L, The role of 15-Lipooxygenase in atherogenesis; pro and antiatherogenic actions, Corropin Lipidol. 1997;8:111-17.
17. Yia-Herttuala S, Palinski W, Rosenfeld ME. Evidence for the presence of oxidatively modified
low-density lipoprotein in atherosclerotic lesions of rabbit and man. J Clin Invest. 1989;84:1086-95.
18. Parthasarathy S, Rankin SM. Role of oxidized LDL in atherogenesis. Prog Lipid Res. 1992;31:127-43.
19. Berliner TA, Navab M, Fogelman AM. Basic mechanisms oxidation, inflammation and genetics. Correlation. 1995;91:2488-96.
20. Berliner TA, Heinecke JW. The role of oxidized LP in atherogenesis. Free Rad Biol Med. 1996;20:707-27.

21. Steinberg D. Oxidative Modification of LDL and atherogenesis. Circulation. 1997;95:1062-71.

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