Status of glycemic control and lipid profile in type 2 diabetes mellitus: A descriptive cross-sectional study

Sarbottam Koirala^{1*}, Rano Mal Piryani², Akash Shahi³, Shristi Bhandari⁴, Sanjay Shah⁵, Smita Bashyal⁶

¹Department of Internal Medicine, Seti Provincial Hospital, Dhangadhi, Nepal, ²Department of Pulmonary Medicine and Medical Education, Bilawal Medical College, Pakistan, ³Department of Internal Medicine, Universal College of Medical Sciences, Bhairahawa, Nepal, ⁴Department of Dentistry, Kantipur Dental Home and Institute, Dhangadhi, Nepal, ⁵Department of Anaesthesia, CP Hospital, Dhangadhi, Nepal, ⁶Medical Officer, Crimson Hospital, Tilottama, Nepal

ABSTRACT

Introduction: Dyslipidemia in diabetes mellitus is commonly seen in the form of high total cholesterol (TC), high lowdensity lipoprotein (LDL), high triglyceride (TG) and low high-density lipoprotein (HDL) levels. This study was done to study the association of glycemic control with lipid profile in Type 2 Diabetes Mellitus (T2DM) patients. **Methods**: This was a cross sectional, observational study conducted from December 2018 to June 2020 at Universal College of Medical Sciences and Teaching Hospital, Bhairahawa. It involved 125 patients of type 2 diabetes mellitus (68 males and 57 females), who visited in outpatient department of Internal Medicine. Laboratory tests [(Fasting Blood Sugar (FBS), Post Prandial Blood Sugar (PPBS), Glycosylated Hemoglobin (HbA1c), Renal Function Test (RFT) and Fasting Lipid Profile] were noted. The data was analyzed with SPSS Version 25. Descriptive statistics like frequency, percentage, mean and standard deviation (SD) were calculated. Independent t-test was used to measure the association between glycated hemoglobin with blood glucose level and glycated hemoglobin with lipid profile. **Results:** The study patients were categorized into two groups on the basis of HbA1C level [(good glycemic control <7 (n=33) and poor glycemic control >7 (n=92)]. In good glycemic control the mean TC was 143.30, LDL 71.88, HDL 43.70, VLDL 29.55, TG 134.39 mg/ dl while in poor glycemic control the mean TC was 182.47, LDL 114.85, HDL 32.03. VLDL 36.61, TG 179.33 mg/dl with p-value <0.05. **Conclusions:** There was significant positive association of HbA1c with TC, LDL, VLDL, TG and significant negative association with HDL.

Keywords: Dyslipidemia, HbA1c, lipid profile, Type 2 diabetes mellitus.

*Correspondence:

Dr. Sarbottam Koirala Department of Internal Medicine Seti Provincial Hospital Dhangadi, Nepal Email: koirala.sarbottam@gmail.com ORCID iD: 0000-0001-7951-0190

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INTRODUCTION

Diabetes mellitus (DM) is a group of metabolic disease of abnormal carbohydrate metabolism which is characterized by hyperglycemia due to defect in insulin secretion, action or both.¹. It is mainly classified into type 1, type 2, "other specific types" and gestational diabetes mellitus.² Globally the prevalence among adults over 18 years of age rose from 4.7% in 1980 to 8.5% in 2014.³ It is estimated that in 2011, 366 million people had DM and it would have increased to 522 million by 2030.⁴ The prevalence of DM in Nepal ranges from 6.3-8.5%.^{5.6}

Type 1 Diabetes Mellitus (T1DM) is characterized by autoimmune destruction of pancreatic beta cells which leads to absolute insulin deficiency and accounts for 5-10% of diabetes in adults. They have a longer period of symptoms like polyuria, polydipsia, fatigue than in children.⁷ While in Type 2 Diabetes Mellitus (T2DM), which accounts for >90% of diabetes in adults, it is characterized by hyperglycemia due to progressive loss of insulin secretion from beta cells due to relative insulin deficiency. Most of them are asymptomatic and classical symptoms of diabetes usually appear when there is an

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increase in blood glucose level. It is primarily due to genetics and lifestyle factors like sedentary lifestyles, cigarette smoking and lavish alcohol consumption.^{8,9} Diabetes with dyslipidemia leads to the development of atherosclerotic plaque, accelerating atherogenesis, resulting in an increase in cardiovascular risk and also worsens diabetes and vice versa. There is an atherogenic lipid profile, low high-density lipoprotein (HDL) and high triglyceride (TG) which have a greater risk of cardiovascular disease (CVD) in comparison with non-diabetics.¹⁰

Hence, this study was done to study the association of glycemic control with lipid profile in Type 2 Diabetes Mellitus (T2DM) patients.

METHODS

This study was conducted in the Department of Internal Medicine. A cross-sectional, observational study was carried out on patients with known case of T2DM visiting outpatient department. The ethical clearance was obtained from Institutional Review Committee (IRC) of Universal College of Medical Sciences and Teaching Hospital (Ref. No. UCMS/IRC/213/18). Patients already diagnosed as T2DM and under treatment, aged more than 25 years and patients who gives written consent were enrolled in the study. Type 2 diabetic patients who were not undergoing treatment for the same, aged less than 25 years, those patients with familial dyslipidemia and the patients who did not give written consent were excluded from the study.

Sample Size was calculated using the following formula

$$n = \frac{Z^2 pq}{d^2}$$

where, n= Sample size, Z= 1.96 at 95% confidence interval, p(expected prevalence)=8.2 $\%^{11}$, q=1-p=91.9 %, d=5% (Maximum tolerable error). The calculated sample size was 116. We have taken 125 as total sample size.

Patient was counselled regarding the procedure. Under aseptic condition, about 6 ml of fasting blood samples (overnight fast between 8 to 12 hours) were drawn from the median cubital vein on the anterior forearm into clot activator/ separating gel tubes. Fasting blood sugar, postprandial blood sugar, blood urea, serum sodium and serum potassium, blood sugar and fasting lipid profile was done by Humastar 600 machine by using spectrophotometry method. Glycated Hemoglobin was determined using GPP-100 HbA1C kit used on GPP-100 Specific Protein Analyzer. Blood glucose was determined by GOD-PAP (glucose oxidase and 4-aminoantipyrine). Cholesterol was determined after enzymatic hydrolysis and oxidation. Triglyceride was estimated using GPO-PAP (glycerine phosphate oxidase) method. LDL was estimated by using Friedwald formula LDL = Total cholesterol - (HDL-C + TG/5).

The data collected were entered and analyzed in Statistical Package for Social Sciences (SPSS) Version 25.0. Descriptive statistics like frequency, percentage, mean and standard deviation (SD) were calculated. Independent t-test was used to measure the association between glycated hemoglobin with blood glucose level and glycated hemoglobin with lipid profile. The level of statistical significance was set at p-value <0.05 at 95% CI.

RESULTS

A total of 125 patients presented in the OPD with the diagnosis of T2DM were included in the study and analyzed systematically. Among 125 patients, majority of the patients, 53 (42.40%) were from age group, 46 to 55 years, followed by 56 to 65 years of age group 27 (21.60%). The least common age group was, patients above 76 years 4(3.20%). The mean age of study patients was 53.31 and standard deviation was 10.83 years with a range of minimum 25 to 86 years. (Table 1)

Table 1: Distribution of study patients according to agegroup (N=125)

Age group (in years)	Frequency	Percentage (%)
26-35	7	5.60%
36-45	22	17.60%
46-55	53	42.40%
56-65	27	21.60%
66-75	12	9.60%
>76	4	3.20%

Among 125 patients enrolled in study visiting outpatient department, 43(34.40%) had tingling sensation of limb, 42(33.60%) had polyuria while 21(16.80%) had diabetic foot (wound/ulcer). Weight loss was the least common symptom associated with 9(7.20%) patients. (Figure 1)

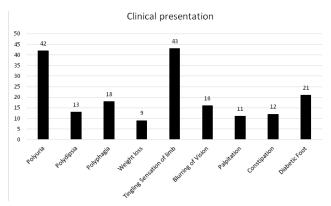


Figure 1: Clinical presentation at the time of study (N=125)

Among the patients studied, diabetic neuropathy was the most common complication of T2DM seen in 33(26.4%), followed by diabetic retinopathy 16(12.8%) while glaucoma was the least common seen in 4(3.2%). (Table 2)

Table 2: Complications at the time of presentation inpatients studied (N=125)

Complications	Frequency	Percentage (%)
Diabetic Neuropathy	33	26.4%
Diabetic Nephropathy	14	11.2%
Diabetic Retinopathy	16	12.8%
Autonomic Neuropathy	9	7.2%
Cataract	12	9.6%
Glaucoma	4	3.2%

The mean FBS with good glycemic control (i.e. HbA1C \leq 7) was 131.87 and with poor glycemic control (i.e. HbA1C>7) mean was 218.97 and mean difference was 87.09 and its p-value is <0.001 which is statistically significant.

The mean PPBS with good glycemic control was 179.15 and with poor glycemic control was 302.21 and mean difference was 123.06 and its p-value is <0.001 which is statistically significant. (Table 3)

Table 3: Association of HbA1C with blood glucose in studypatients (N=125)

Glycated Hemoglobin (%)	≤7 (n=33)	>7 (n = 92)	Mean difference	p value*
FBS (mg/dL)	131.87±29.58	218.97±71.32	87.09	<0.001*
PPBS (mg/dL)	179.15±32.67	302.21±82.81	123.06	<0.001*

Independent t-test; *p-value<0.05 signifies statistical significance

Similarly, in patients with good glycemic control the mean of TC was 143.30 and with poor control was 182.47 and its p value is <0.001 which is statistically significant. The mean of LDL in good glycemic control was 71.88 and with poor control was 114.85 and its p value is <0.001 which is statistically significant. The mean of HDL in good control was 43.70 and with poor control was 32.03 and its p-value is <0.001 which is statistically significant. The mean of VLDL in good control was 29.55 and with poor control was 36.61. The mean of TG in good control was 134.39 and with poor control was 179.33 and its p-value is <0.001 which is statistically significant. (Table 4)

Table 4: Patients lipid profile in relation with good andpoor glycemic control

Parameters	HbA1C≤7% (n=33)	HbA1C>7% (n = 92)	Mean dif- ference	p value*
Total Cholesterol (mg/ dL)	143.30±23.36	182.47±35.39	39.16	<0.001*
LDL (mg/dL)	71.88±19.02	114.85±38.82	42.96	< 0.001*

HDL (mg/dL))	43.70±8.78	32.03±11.66	11.66	<0.001*
VLDL (mg/dL)	29.55±12.21	36.61±10.72	7.06	0.05
TG (mg/dL)	134.39±41.15	179.33±46.96	44.93	< 0.001*

Independent t-test; *p-value<0.05 signifies statistical significance

DISCUSSION

In this study the mean age of patients was 53.31 years with standard deviation of 10.83 and range of 25 to 86 years. The majority of patients were of age group 46 to 55 years. Similar result was found in a study conducted in Manipal College of Medical Sciences, Nepal by Pokharel et al., where mean age was 52.7 with standard deviation 10.50.¹² There was slight male preponderance in this study with male 54.4% male and 45.6% female. In study done by Memon et al., 67.8% were male and 32.14% were female with male preponderance which correlates with our study.¹³ Tingling sensation of limb 34.4% followed by polyuria 33.6%, diabetic foot 16.8%, polyphagia 14.4%, blurring of vision 12.8%, polydipsia 10.4%, constipation 9.6%, palpitation 8.8% and weight loss 7.2% which correlates with study done by Singh et al.¹⁴

The present study on T2DM patients showed that there was positive association of TC, LDL, VLDL and TG and negative association of HDL in relation with HbA1C. Prabhavathi et al. conducted a similar study where there was positive association of TC, LDL, TG and negative association with HDL which correlates with our study.¹⁵ Similarly, Jain et al.¹⁶ Sarkar et al.¹⁷ Khan et al.¹⁸ and Hussain et al.¹⁹ also found similar result in their study.

The limitation of this study was that it was a singlecentered study, so the findings could not be generalized. Also, the type of antidiabetic drugs and their dosages were not specified. Hence, their individual effect could not be assessed.

CONCLUSIONS

Glycemic control with glycated hemoglobin has helped in predicting the dyslipidemia in patients. So, it can be used as a screening method for T2DM patients to prevent further cardiovascular events and timely management.

CONFLICTS OF INTEREST: None declared

SOURCE OF FUNDING: None

AUTHORS CONTRIBUTION

SK did concept designing, definition of intellectual content, literature search, RMP, AS did review and editing SB did data acquisition and analysis, manuscript preparation; SS and SB did concept designing, literature search, data analysis and manuscript review.

REFERENCES

- American Diabetes Association (ADA). Introduction. Diabetes Care. 2017;40(January):S1-S2. DOI: 10.2337/ dc17-S001
- Baynest HW. Classification, Pathophysiology, Diagnosis and Management of Diabetes Mellitus. J Diabetes Metab. 2015;06(05):1-9. DOI: 10.4172/2155-6156.1000541
- Sarwar N, Gao P, Kondapally Seshasai SR, Gobin R, Kaptoge S, Angelantonio DE, et al. Diabetes mellitus, fasting blood glucose concentration, and risk of vascular disease: a collaborative meta-analysis of 102 prospective studies. Lancet. 2010;375(9733):2215-22. DOI: 10.1016/S0140-6736(10)60484-9 PMID: 20609967.
- Chamnan P, Simmons RK, Forouhi NG, Luben RN, Khaw KT, Wareham NJ, et al. Incidence of type 2 diabetes using proposed HbA1C diagnostic criteria in the European prospective investigation of Cancer-Norfolk cohort: Implications for preventive strategies. Diabetes Care. 2011;34(4):950-6. DOI: 10.2337/dc09-2326 PMID: 20622160.
- Shakya-Vaidya S, Aryal UR, Upadhyay M, Krettek A. Do non-communicable diseases such as hypertension and diabetes associate with primary open-angle glaucoma? Insights from a case-control study in Nepal. Global Health Action. 2013;6:22636. DOI: 10.3402/gha. v6i0.22636 PMID: 24192283.
- Sharma SK, Ghimire A, Radhakrishnan J, Thapa L, Shrestha NR, Paudel N, et al. Prevalence of hypertension, obesity, diabetes, and metabolic syndrome in Nepal. International Journal of Hypertension. 2011;2011:821971. DOI: 10.4061/2011/821971 PMID: 21629873.
- Karjalainen J, Salmela P, Ilonen J, Surcel HM, Knip M. A comparison of childhood and adult type I diabetes mellitus. New England Journal of Medicine. 1989;320(14):881-6. DOI: 10.1056/ NEJM198904063201401 PMID: 2648146.
- 8. Ripsin CM, Kang H, Urban RJ. Management of blood glucose in type 2 diabetes mellitus. American Family Physician. 2009;79(1):29-36. PMID: 19145963.
- 9. Hu FB, Manson JE, Stampfer MJ, Coldotz G, Liu SL,

Solomon CG, et al. Diet, lifestyle and the risk of type 2 diabetes mellitus in women. New England Journal of Medicine. 2001;345(11):790-7. DOI: 10.1056/ NEJMoa010492 PMID: 11556298.

- Windler E. What is the consequence of an abnormal lipid profile in patients with type 2 diabetes or the metabolic syndrome? Atheroscleros Supplements. 2005;6(3 spec.iss.):11-4. DOI: 10.1016/j. atherosclerosissup.2005.06.003 PMID: 16046281.
- 11. Gyawali B, Sharma R, Neupane D, Mishra SR, van Teijlingen E, Kallestrup P. Prevalence of type 2 diabetes in Nepal: A systematic review and meta-analysis from 2000 to 2014. Glob Health Action. 2015;8. DOI: 10.3402/ gha.v8.29088 PMID: 26613684.
- Pokharel DR, Khadka D, Sigdel M, Yadav NK, Acharya S, Kafle RC, et al. Prevalence and pattern of dyslipidemia in Nepalese individuals with type 2 diabetes. BMC Research Notes. 2017;10(1):1-11. DOI: 10.1186/ s13104-017-2465-4
- Farooq Memon DF, Rathi DN, Ali Rahoo DQ, Maheshwari DB, Maheshwari DM, Memon DF. HbA1C and lipid profile; correlation with the complexity of coronary artery disease. The Professional Medical Journal. 2017;24(05):650-655. DOI: 10.29309/ TPMJ/2017.24.05.1294
- 14. Singh NK, Shah NK, Bhandari A, Pandey S, Sharma SJ. Presentations and complications of diabetes patients presenting to diabetic clinic of Eastern Nepal. Journal of College of Medical Sciences-Nepal. 2014;9(3):25-30. DOI: 10.4103/2230-8210.152790 PMID: 25932400.
- Prabhavathi K, Kirthana Kunikullaya U, Goturu J. Glycosylated haemoglobin(HbA1c) A marker of circulating lipids in type 2 diabetic patients. J Clin Diag Res. 2014;8(2):20-23. DOI: 10.7860/jcdr/2014/7742.3996 PMID: 24701471.
- Jain HR, Shetty V, Singh GS, Shetty S. A Study of Lipid Profile in Diabetes Mellitus. International Journal of Scientific Study. 2016;4(4):56-61. DOI: 10.17354/ ijss/2016/615
- Sarkar S, Meshram A. Hba1C and Lipid Profile Levels in the Known Type 2 Diabetic Group in the Rural Region of Vidarbha, Maharashtra, India. Journal of Evidence Based Medicine and Healthcare. 2017;4(32):1915-20. DOI: 10.18410/jebmh/2017/374

18. Khan HA, Sobki SH, Khan SA. Association between

page 103

glycaemic control and serum lipids profile in type 2 diabetic patients: HbA1c predicts dyslipidaemia. Clinical and Experimental Medicine. 2007;7(1):24-29. DOI: 10.1007/s10238-007-0121-3 PMID: 17380302.

19. Hussain A, Ali I, Ijaz M, Rahim A. Correlation between

hemoglobin A1c and serum lipid profile in Afghani patients with type 2 diabetes: Hemoglobin A1C prognosticates dyslipidemia. Therapeutic Advances in Endocrinology and Metabolism. 2017;8(4):51-7. DOI: 10.1177/2042018817692296 PMID: 28507727.