**ORIGINAL RESEARCH PAPER** 

# **INTERNATIONAL JOURNAL OF SCIENTIFIC RESEARCH**

# A STUDY OF LIPID PROFILE AND ITS RELATIONSHIP WITH BLOOD GLUCOSE LEVEL IN METABOLIC SYNDROME

General Medicine		
Dr. Tarang Shah	3 <sup>rd</sup> Year Resid LG Hospital,	lent, Department of General Medicine, AMC MET Medical college & Sheth Ahmedabad.
Dr. Abhi Doshi*		lent, Department of General Medicine, AMC MET Medical college & Sheth Ahmedabad. *Corresponding Author
Dr. Nilay Suthar	Professor, De Hospital, Ahr	epartment of General Medicine, AMC MET Medical college & Sheth LG nedabad.
Dr. Dipak Solanki		ofessor, Department of General Medicine, AMC MET Medical college & spital, Ahmedabad.
Dr. Ajay Rathod		ofessor, Department of General Medicine, AMC MET Medical college & spital, Ahmedabad.

# ABSTRACT

Background: Metabolic syndrome (MetS) and its associated factors such as dyslipidemia and hyperglycemia are associated with increased risk of cardiovascular disease (CVD).

Aims and Objective: To assess lipid profile and its relation with blood glucose levels in patients with MetS.

Materials and Methods: This cross-sectional study included 320 patients with MetS. Anthropometry, lipid profile, blood glucose, and presence of MetS (NECPAPT3) were determined.

**RESULT**: High LDL-C was the most common lipid abnormality observed in these patients. High LDL-C (>100 mg/dl) that is including suboptimal and high level were observed in 75.93% patients. Whereas high total triglyceride, high cholesterol, and low HDL level were found in 59.68%, 47.18%, 41.87% respectively. On intergroup comparison of NFG, IFG, and T2DM among patients with MetS, no statistically significant difference in lipid levels was observed.

**Conclusion:** we conclude that dyslipidemia is a common feature of MetS, and a large number of patients had more than one individual lipid abnormality. Most common dyslipidemia was high LDL-C and least was low HDL. Pattern of the dyslipidemia was similar in all three groups based on blood glucose levels.

# **KEYWORDS**

# INTRODUCTION

Metabolic syndrome is defined as per the NATIONAL CHOLESTEROL EDUCATION PROGRAMME AND ADULT TREATMENT PANEL 3 [ NCEP ATP3 ], Central obesity : waist circumference >102cm (M), >88cm (F), Hypertriglyceridemia : triglyceride level >=150 mg/dl or specific medication, Low HDL cholesterol : <40 mg/dl and <50 mg/dl for men and women respectively, or specific medication, Hypertension: blood pressure >=130 mmhg systolic or >=85 mmhg diastolic or specific medication, Fasting plasma glucose level >=100 mg/dl or specific medication or previously diagnosed type 2 diabetes.

Obesity, particularly abdominal obesity, is associated with resistance to the effects of insulin on peripheral glucose and fatty acid utilization, often leading to type 2 diabetes mellitus. Insulin resistance, the associated hyperinsulinemia and hyperglycemia, and adipocyte cytokines (adipokines) may also lead to vascular endothelial dysfunction, an abnormal lipid profile, hypertension, and vascular inflammation, all of which promote the development of atherosclerotic cardiovascular disease (CVD).

The co-occurrence of metabolic risk factors for both type 2 diabetes and CVD (abdominal obesity, hyperglycemia, dyslipidemia, and hypertension) suggested the existence of a "metabolic syndrome". Other names applied to this constellation of findings have included syndrome X, the insulin resistance syndrome, the deadly quartet, or the obesity dyslipidemia syndrome [12]. Genetic predisposition, lack of exercise, and body fat distribution all affect the likelihood that a given obese subject will become overtly diabetic or develop CVD.

Many studies available in literature show the association between lipid profile and MetS-associated variables. But there is paucity of data regarding relationship of blood glucose levels and lipid profile in MetS. In this study, lipid pattern and its relation to blood glucose levels in patients with MetS was investigated.

## MATERIALAND METHOID

This cross-sectional study was conducted in the Postgraduate

Department of Medicine, AMC MET Medical College, Ahmedabad, India. Informed written consent was obtained after explaining the nature of the study to the patients.

This study included 320 patients with MetS, whereas those with history of CVD, thyroid disorders, or currently on lipid-lowering agents were excluded. Detailed history was noted and clinical examination was carried out. Body mass index (BMI), waist circumference (WC), and systolic and diastolic blood pressure were measured using standard methods. Laboratory assessment included venous blood samples in a fasted state for the determination of components of the lipid profile [total cholesterol (TC), HDL-C, and TG] and blood glucose levels. The serum glucose was measured using the glucose oxidase/peroxidase method and the lipid profile by the enzymatic colorimetric method. LDL-C was calculated from the formula of Friedewald., Patients were categorized into three groups depending on their fasting blood glucose levels. Group I comprised patients with normal fasting glucose (NFG; <100 mg/dL), Group II had patients with IFG status (100-125 mg/dL), and Group III had patients with T2DM ( $\geq 126 \text{ mg/dL}$ ).

#### Statistical Analysis:

Intergroup comparisons were done using Pearson's  $\chi$ 2-test, and mean values were compared using analysis of variance. Statistically significant differences were reported at p < 0.05.

## RESULT

The baseline characteristics of 320 patients with MetS show that their mean age (years) was 58.59  $\pm$  10.74, BMI (kg/m2) 24.5  $\pm$  5.46, and WC (cm) 91  $\pm$  10.48. Biochemical analysis showed that mean fasting blood sugar was 124  $\pm$  52.34 mg/dL whereas HDL, TG, TC, and LDL-C were 41  $\pm$  9.42, 159  $\pm$  34.19, 198  $\pm$  28.89, and 117  $\pm$  24.85 mg/dL, respectively (Table 1).

# Table 1 : Baseline Characteristic Of Patients (n=320)

Tuble 1 · Dusenne Characteristic Off attents (il 520)			
PARAMETERS	MEAN ±SD		
AGE ( years )	$58.59 \pm 10.17$		
BMI (kg/m2)	$24.5\pm5.46$		
International Journal of Scientific Research	- 69		

## Volume-8 | Issue-11 | November - 2019

WAIST CIRCUMFERENCE (cm)	$91 \pm 10.48$
SYSTOLIC BP (mmhg)	$135 \pm 19.77$
DIASTOLIC BP (mmhg)	$85 \pm 9.73$
FASTING BLOOD SUGAR (mg/dl)	$124 \pm 52.34$
SERUM HDL – C (mg/dl)	$41 \pm 9.42$
SERUM TRIGLYCERIDE (mg/dl)	$159 \pm 34.19$
TOTAL SERUM CHOLESTEROL (mg/dl)	$198\pm28.89$
SERUM LDL – C (mg/dl)	$117 \pm 24.85$

# TABLE 2: Prevalence of optimal, suboptimal, and high lipid levels in metabolic syndrome (*n* = 320)

LIPID PROFILE	OPTIMAL	SUBOPTIMAL	HIGH
TG *	129 (40.31%)	159 (49.69%)	38 (11.86%)
LDL-C**	81 (25.31%)	230 (71.86%)	9 (0.029%)
TC***	169 (52.81%)	130 (40.62%)	21 (0.006%)

\* TG: Optimal - <150 mg/dL; Suboptimal - 150–199 mg/dL; High:  $\geq 200 \text{ mg/dL}$ .

\*\* LDL-C: Optimal - <100 mg/dL; Suboptimal - 100−159 mg/dL; High:≥160 mg/dL.

\*\*\* TC: Optimal - <200 mg/dL; Suboptimal - 200–239 mg/dL; High: ≥240 mg/Dl

## Table 3 : Prevelence Of Hdl C In Mets (n=320)

HDL – C	,	Low, <40 mg/dL	Total
SUBJECT	186 (58.13%)	134 (41.87%)	320

 Table 4 :distribution Of Dyslipidemia In Relation To Blood Glucose

 Level

LIPID		IFG (n=106)	T2DM	P VALUE
PARAMETER	(n=112)		(n=102)	
$TG \ge 200$	11 (9.82%)	15 (14.15%)	12 (11.76%)	0.7166
mg/dL				
LDL-C≥	6 (5.35%)	3 (2.83%)	4 (3.92%)	0.1159
160 mg/dL				
$TC \ge 240$	12 (10.71%)	4 (3.77%)	5 (4.90%)	0.9521
mg/dL				
HDL-C < 40	49 (43.75%)	60 (56.60%)	52 (50.98%)	2.8127
mg/dL				

TG < 150 mg/dL, LDL < 100 mg/dL, TC < 200 mg/dL), and HDL-C > 40mg/dL were observed in 40.31%, 25.31%, 52.81%, and 58.13% patients, respectively. TG  $\ge$  200 mg/dL, TC  $\ge$  240 mg/dL, and LDL-C  $\ge$  160 mg/dL were observed in 11.86%, 0.006%, and 0.029% patients, respectively, suggesting that many patients had more than one lipid abnormality (Tables 2 and 3).

Analysis of distribution of dyslipidemia showed that hypertrigly ceridemia (TG  $\geq$  200 mg/dL) was present in 9.82%, 14.15%, and 11.76% patients with NFG, IFG, and T2DM, respectively.Low HDL-C was observed in NFG (43.75%), IFG (56.60%) and T2DM (50.98%) of patients. High LDL-C was observed in 5.35%, 2.83% and 3.92% patients with NFG, IFG and T2DM, respectively. High TC was observed in 10.71%, 3.77% and 4.90% of patients with NFG,IFG and T2DM, respectively. On intergroup comparison, differences were not found to be statistically significant (Table 4).

#### DISCUSSION

In this study, the relationship between glucose levels and lipid pattern in patients with MetS was examined. High LDL-C was the most common lipid abnormality observed in these patients.

High LDL-C (>100 mg/dl) that is including suboptimal and high level were observed in 75.93% patients. Whereas high total triglyceride, high cholesterol, and low HDL level were found in 59.68%, 47.18%, 41.87% respectively.

High levels of TG and low levels of HDL-C in patients with MetS result from decreased clearance of these lipoproteins from the circulation. Lipoprotein lipase (LPL) is a major enzyme responsible for clearing TG-containing lipoproteins from the circulation, and insulin resistance is associated with impaired LPL activity. Hepatic lipase, which is responsible for clearing HDL particles from the circulation, shows increased activity in the presence of insulin resistance and causes HDL-C levels to decline.[15] A low level of HDL-C is an important risk factor for CVD. The cardio-protective

effects of HDL-C have been attributed to its role in reverse cholesterol transport, its effects on endothelial cells, and its antioxidant activity.

Elevated levels of LDL-C are a major risk factor for CVD and its reduction is prime target of pharmacotherapy. The positive relationship between first or subsequent attacks of coronary heart disease is observed over a broad range of LDL-C levels. The higher the level of LDL-C, the greater the risk is.

In a study carried out on Indian population with T2DM, hypertrigly ceridemia and high serum LDL-C levels ( $\geq 100 \text{ mg/dL}$ ) were recorded as major components of dyslipidemia, and most of these patients had mixed dyslipidemia. These findings are in concurrence with the results of this study, whereas others have recorded normal levels of LDL-C.

On intergroup comparison of NFG, IFG, and T2DM among patients with MetS, no statistically significant difference in lipid levels was observed. Insulin-resistant individuals not having diabetes mellitus are likely to have lipid profiles that are nearly identical to those seen in the large majority of patients with T2DM as observed in this study

## CONCLUSION

From results of this study, we conclude that dyslipidemia is a common feature of MetS, and a large number of patients had more than one individual lipid abnormality. Most common dyslipidemia was high LDL-C and least was low HDL. Pattern of the dyslipidemia was similar in all three groups based on blood glucose levels.

#### REFERENCES

- Alberti KG, Zimmet P, Shaw J. Metabolic syndrome—a new world-wide definition. A Consensus Statement from the International Diabetes Federation. Diabet Med. 2006;23(5):469–80.
- Haffner SM, Stern MP, Hazuda HP, Mitchell BD, Patterson JK. Cardiovascular risk factors in confirmed pre-diabetic individuals: does the clock for coronary heart disease start ticking before the onset of clinical diabetes? JAMA. 1990;263:2893–8.
- Chahil TJ, Ginsberg HN. Diabetic dyslipidemia. Endocrinol Metab Clin North Am. 2006;35:491–510.
   Jeppesen J, Hein HO, Suadicani P, Gyntelberg F. Relation of high TG-low HDL
- Jeppesen J, Hein HO, Suadicani P, Gyntelberg F. Relation of high IG-low HDL cholesterol and LDL cholesterol to the incidence of ischemic heart disease. An 8-year follow-up in the Copenhagen Male Study. Arterioscler Thromb Vasc Biol. 1997;17:1114–20.
- Assmann G, Schulte H. Relation of high-density lipoprotein cholesterol and triglycerides to incidence of atherosclerotic coronary artery disease (the PROCAM experience). Prospective Cardiovascular Munster study. Am J Cardiol. 1992;70:733–7.
- Kawamoto R, Tabara Y, Kohara K, Miki T, Kusunoki T, Takayama S, et al. Relationships between lipid profiles and metabolic syndrome, insulin resistance and serum high molecular adiponectin in Japanese community-dwelling adults. Lipids Health Dis. 2011;10:79.
- Goodarzi MT, Mohammadian M, Borzouei S, Hassanzadeh T. Association between plasma cholesteryl ester transfer protein activity and lipid profiles in metabolic syndrome in an Iranian population. Int Res J Biol Sci. 2014;3(4):87–90.
- Friedewald WT, Levy RI, Fredrickson DS. Estimation of the concentration of low density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. Clinical Chem. 1972:18:499–502.
- Alberti KG, Eckel RH, Grundy SM, Zimmet PZ, Cleeman JI, Donato KA, et al. Harmonizing the metabolic syndrome: a Joint Interim Statement of the International Diabetes Federation Task Force on Epidemiology and Prevention; National Heart, Lung, and Blood Institute; American Heart Association; World Heart Federation; International Atherosclerosis Society; and International Association for the Study of Obesity. Circulation. 2009;120(16):1640–5.
   National Cholesterol Education Program (NCEP) Expert Panel on Detection,
- National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III). Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III) Final Report. Circulation. 2002;106:3143.
- Alberti KG, Zimmet PZ. Definition, diagnosis and classification of diabetes mellitus and its complications. Part diagnosis and classification of diabetes mellitus provisional report of a WHO consultation. Diabet Med. 1998;15(7):539–53.
   Bal SS, Khurana D, Sharma A, Lal V, Bhansali A, Prabhakar S. Association of metabolic
- Bal SS, Khurana D, Sharma A, Lal V, Bhansali A, Prabhakar S. Association of metabolic syndrome with carotid atherosclerosis in the young North Indian population. Diabetes Metab Syndr. 2011;5(3):153–7.
- Despres JP, Lemieux I. Abdominal obesity and metabolic syndrome. Nature. 2006;444:881–7.
- Maheux P, Azhar S, Kern PA, Chen YD, Reuven GM. Relationship between insulinmediated glucose disposal and regulation of plasma and adipose tissue lipoprotein lipase. Diabetologia. 1997;40:850–8.
- Adiels M, Olofsson SO, Taskinen MR, Boren J. Overproduction of very low-density lipoproteins is the hallmark of the dsylipidemia in the metabolic syndrome. Arterioscler Thromb Vase Biol. 2008;28:1225–36.
- Assmann G, Gotto AM Jr. HDL Cholesterol and protective factors in atherosclerosis. Circulation. 2004;109:III-8–III-14.
   Pandya H, Lakhani JD, Dadhania J, Trivedi A. The prevalence and pattern of
- Pandya H, Lakhani JD, Dadhania J, Trivedi A. The prevalence and pattern of dyslipidemia among type 2 diabetic patients at rural based hospital in Gujarat, India. Indian J Clin Pract. 2012;22(12):36–44.
- U.K. Prospective Diabetes Study 27: Plasma lipids and lipoproteins at diagnosis of NIDDM by age and sex. Diabetes Care. 1997;20(11):1683–7.
- Cowie CC, Howard BV, Harris MI. Serum lipoproteins in African Americans and whites with non-insulin dependent diabetes in the US population. Circulation. 1994;90(3):1185–93.
- Ginsberg HN. Insulin resistance and cardiovascular disease. J Clin Invest. 2000;106:453–8.
- 21. Alberti KG, Eckel RH, Grundy SM, et al. Harmonizing the metabolic syndrome: a joint interim statement of the International Diabetes Federation Task Force on Epidemiology and Prevention; National Heart, Lung, and Blood Institute; American Heart Association; World Heart Federation; International Atherosclerosis Society; and International Association for the Study of Obesity. Circulation 2009; 120:1640.

70