PARIPEX - INDIAN JOURNAL OF RESEARCH | Volume - 11 | Issue - 05 | May - 2022 | PRINT ISSN No. 2250 - 1991 | DOI : 10.36106/paripex

Journal or A	RIGINAL RESEARCH PAPER	Medicine
ARIPEN IN HC	OSS SECTIONAL, OBSERVATIONAL STUDY TO KNOW E PREVALENCE OF MICROALBUMINURIA AND CROALBUMINURIA, HYPERTENSION AND SLIPIDEMIA AND PREVALENCE OF CAROTID PLAQUES FYPE 2 DIABETES PATIENTS IN A TERTIARY CARE SPITAL KOLKATA, WEST BENGAL.	KEY WORDS: DM- diabetes mellitus, Dyslipidemia, HT- hypertension, obesity.
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Type 2 diabetes is often associated with various coexisting comorbidities like-hypertension and dyslipidemia. They are very much associated with ASCVD. Effects of diabetes are more predicting. If we can control multiple risk factors besides Diabetes, we will be able to tackle the ASCVD more efficiently & effectively. The other risk factors for ASCVD like Hypertension, Dyslipidemia, and obesity are prevalent in our country. There is a combination of 5 risk factors collectively named as syndrome X including Hypertension (BP >130/85 mm Hg), insulin resistant DM, hypertriglyceridemia, decreased HDL cholesterol and increased Waist hip ratio. Syndrome X predispose to atherosclerotic changes in the carotids. High blood pressure is reported in over two-thirds of patients with type 2 diabetes, and its development coincides with the development of hyperglycaemia. Dyslipidemia is frequent among patients with type 2 DM (T2DM) (prevalence > 75%) and is mainly a mixed dyslipidemia [increase in triglycerides (TGs), low high-density lipoprotein cholesterol (HDL-C), and small-dense (atherogenic), low-density lipoprotein cholesterol (LDL-C) particles](2). It was also reported that microalbuminuria or macroalbuminuria increased the mortality rate by 60 - 80% -(3) . Albuminuria can be measured by urinary Albumin-to Creatinine ratio (ACR) in a random spot urine collection-(4). The carotid IMT is significantly higher in diabetic patients than that in non-diabetic patients(5), and the increased IMT can predict future events of silent brain infarction and coronary heart disease in the patients with T2DM(6). Our objective is to find out the prevalence of microalbuminuria and macroalbuminuria, hypertension, dyslipidemia, Diabetic kidney disease (serum Cr>1.5) and presence of plaque in carotid arteries in type 2 diabetes patients in our country. Another objective was to find out the distribution of glycemic parameters (FBS, PPBS, HbAlc) and BMI among study groups.

MATERIALS AND METHODS

Ourangajeb*

ABSTRACT

1. **Study Design:** Cross sectional, Single Centre, hospital based prospective and observational study.

- 2. Place Of Study: Nilratan Sircar Medical College, Kolkata.
- 3. Period Of Study: January 2018 to June 2019.

4. Study Population:

Patients attending OPD and IPD of General Medicine Department, Nilratan Sircar Medical College, Kolkata and fulfill inclusion & exclusion criteria of the study. 5. Sample Size: 50.

6. Sample Selection: Randomized

7. Inclusion Criteria:

- 1. Diagnose case of type 2 diabetes mellitus
- 2.Age above 18 years
- 3.Both sexes

8. Exclusion Criteria:

Type 1	Secondary	Gestational	Connecti	Overt renal
diabetes	diabetes	diabetes	ve tissue	failure
mellitus			disorders	
			or	
			vasculitis	
Valvular	Ischemic	Patients	Urinary	Congestive
heart	heart	with	tract	cardiac
disease or	disease	ischemic	infection	failure
atrial	with	stroke.		
fibrillation	medications	infection		

Method Of Data Collection:

Interview &	Physical	Laboratory	Record
History taking	examination	examination,	analysis.

Parameters To Be Studies:

CLINICAL: 1. Presenting clinical features of type 2 diabetes mellitus as polyuria, polydipsia. 2. Complications of type 2 diabetes mellitus as nephropathy, neuropathy or retinopathy and. 3. Duration of diabetes.

GENEREL & SYSTEMIC EXAMINATION:

LABORATORY INVESTIGATIONS:-

1. The following laboratory parameters were carried out:

Complete haemogr am	Fasting Blood Glucose	PPBS	HBA1 C	Spot urine ACR	Serum electrolytes – Sodium , potassium, chloride etc.
Blood	Urine	Lipid	ECG	Echocard	Brain imaging
creatinine	KE/IVIE	e:.	leads	in selected	cases of CVA.
				cases	

2. Specific Investigation:

Urinary albumin Creatinine ratio(ACR) was calculated by dividing the urinary albumin concentration in micrograms by the urinary creatinine concentration in milligrams. Normal UACR is generally defined <30 mg/g Cr, and increased urinary albumin excretion is defined as >30 mg/g Cr. However, UACR is a continuous measurement, and differences within the normal and abnormal ranges are associated with renal and cardiovascular outcomes.

OBSERVATION AND ANALYSIS:

Age Distribution: 10 cases out of 50 cases in the study are below the age of forty years. (Table-1) comprising 20.0 % of total cases, 25 cases are between 40-60 years comprising 50.0 % of total cases, 15 cases are 60yrs or more comprising 30.0 % of total cases.(Table-2))

Sex Distribution: 22 cases out of fifty cases in the study are Female Comprising 44.0 % of total cases, 28 cases are male comprising 56.0 % of total cases(Table 3)

Fasting Blood Sugar: out of 50 cases, 26 cases are in the range OF 100-126 of FBS, comprising of 52 % of cases, 24 cases are >126, comprising 48 % of FBS (Table 4.)

PPBS distribution: out of 50 cases, 27 cases are in the range of 140-200 of PPBS, comprising of 54 % of cases, 15 cases are >200, comprising 30 % of PPBS (TABLE 5.)

HbAlc: 43 cases out of fifty cases in the study are HbAlc (>6..5) Comprising 86.0 % of total cases, 7 cases are HbAlc

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PARIPEX - INDIAN JOURNAL OF RESEARCH | Volume - 11 | Issue - 05 | May - 2022 | PRINT ISSN No. 2250 - 1991 | DOI : 10.36106/paripex

 (≤ 6.5) comprising 14.0 % of total cases (TABLE 6.)

Total Cholesterol-(TC) - Out of 50 cases, 24 cases are abnormal of TC, comprising of 48 % of cases, 26 cases are normal, comprising 52 % of TC. (Table-7)

TG- triglyceride- Out of 50 cases, 31 cases are abnormal of TG , comprising of 62 % of cases, 19 cases are normal , comprising 38 % of TG (Table-8)

LDL-Cholesterol- Out of 50 cases, 21 cases are abnormal, comprising of 42 % of cases, 29 cases are normal, comprising 58 % of LDL-C (Table-9)

HDL-C- Out of 50 cases, 40 cases are abnormal, comprising of 80 % of cases, 10 cases are normal, comprising 20 % of HDL-C. (Table-10)

Urine ACR- Out of 50 cases, 24 cases are abnormal of Urine ACR, comprising of 48 % of cases, 26 cases are normal, comprising 52 % of Urine ACR (table-11)

Hypertension-30 cases out of fifty cases in the study are no Hypertension Comprising 60.0 % of total cases, 20 cases are Hypertension present comprising 40.0 % of total cases(table-13)

PRESENCE OF CAROTID PLAQUE: 30 cases out of fifty cases in the study are no Presence of plaque Comprising 60.0 % of total cases, 20 cases are Presence of plaque (yes) comprising 30.0 % of total cases (table-14).

CIMT- out of 50 cases, 11 cases are increased of CIMT, comprising of 22 % of cases, 39 cases are normal, comprising 78 % of CIMT (table-15)

SERUM CREATININE: 4 cases out of fifty cases in the study are no Serum creatinine (>1.5) Comprising 8.0 % of total cases, 46 cases are Serum creatinine (NORMAL) comprising 92.0 % of total case -**Table 16**.

BMI distribution-Out of 50 cases, 12 cases are in the range of <22.9 of BMI, comprising of 24 % of cases, 20 cases are 23-27.5, comprising 40 % of BMI and >27.3 BMI is 36 %.Table-17

CIMT * **AGE**: Out of 50 cases 10 cases bellow the age of 40 years, among them none has increased CIMT, 19 cases between 40-60 years among them 6 cases and 10 cases above 60 years, of them 5 have increased CIMT respectably.

CIMT * **DURATION OF DM:** From the Pie diagram 18, it is observed that there is maximum increased CIMT for 8 cases comprising of 16 % for > 10 years of duration of DM.

DISCUSSION:

A study by **Gupta et al**(7) found tobacco use, obesity, high blood pressure, high low-density lipoprotein (LDL) cholesterol, low high-density lipoprotein (HDL) cholesterol, and diabetes to correlate with vascular events. Study, showed 92.5% prevalence of high CIMT in T2DM patients with ischemic stroke, 80% prevalence in T2DM patients without any stroke and 20% prevalence in non-diabetics. These prevalence rates vary as per the accompanying risk factors. As per the ultrasonography measurements, it was observed that among patients with increased CIMT (>0.8 mm), male: female ratio was 3.75:1, suggesting that males are more vulnerable to have increased CIMT. As the age and BMI progresses, the risk of having increased CIMT goes up. In all age groups, males scored over females in having increased CIMT. Smokers had statistically significant higher prevalence of having increased CIMT. Hypertensive patients had significantly higher chances of getting increased CIMT than nonhypertensives. But though the subjects with duration of T2DM >10 years had a higher prevalence of increased CIMT, it was statistically not significant in comparison to subjects with duration of T2DM <10 years. As the age and BMI progresses, the risk of having increased CIMT goes up. In all age groups, males scored over females in having increased CIMT. Smokers had statistically significant higher prevalence of having increased CIMT. Any increase in BMI above normal

weight levels is associated with an increased risk of being diagnosed as having complications of diabetes mellitus. For men, the increased risk of these complications occurred at higher BMI levels than in women. Ocular complications occurred at higher BMI levels than other complication types in both men and women(8).

Hypertensive patients had significantly higher chances of getting increased CIMT than nonhypertensives. But though the subjects with duration of T2DM >10 years had a higher prevalence of increased CIMT, it was statistically not significant in comparison to subjects with duration of T2DM <10 years.

The three glycemic parameters (FPG, PPPG, and HbA1C) and the lipid parameters like total cholesterol, LDL cholesterol, and triglyceride were all significantly higher in patients with increased CIMT. Though HDL cholesterol had a negative association with the occurrence of increased CIMT, it was not significant. Diercks et al. similarly showed that urine albumin excretion was strongly related to subclinical atherosclerosis in the presence of increased CIMT in patients with T2DM (9). The Framingham study and Multiple Risk Factor Intervention Trial (MRFIT) showed a 2-3 fold elevation in the risk of clinically evident atherosclerotic disease in patients with T2DM(10). The risk factors predispose to increased IMT and correlate well with lack of coronary artery disease, whereas an increasing CIMT above normal level (> 0.8 mm) is associated with increasingly severe coronary artery disease, an increased risk of myocardial infarction and also stroke.

CONCLUSION-

Most common age group of diabetes was 40- 60yrs (50%). Male predominance of diabetes was seen 56% v/s 44%. Regarding glycemic parameters FBS>126, PPBS>200, HbA1c>6.5% were associated with 48%, 30%, 86% respectively. Hypertension was coexisting with diabetes in 40% cases. In our study diabetes patients are mostly having abnormalities of hypertriglyceridemia, then low HDL, then total cholesterol, then LDL cholesterol. Which tally with Indian lean diabetics-abdominal obesity. 27.3% cases having high BMI >27.3. Not very significant patients were having significant kidney disease (8%). Atherosclerotic plaque (30%) was not very common in our study. CIMT increased only in 22% v/s 78%. Risk of carotid atherosclerosis increased with age and duration of diabetes >10 years.

Limitation of the study- STUDY DESIGN: A longitudinal study with baseline and follow up in type 2 diabetic patients would have been more suited to determine the relationship between the occurrence of atherosclerosis and Syndrome X. SELECTION BIAS: The study being a hospital-based study, there is always a chance of selection bias SAMPLE SIZE: A larger number of study populations would have made this study more accurate and leaded further weightage to the results. 50 patients were too small to come into a conclusion. REFERRAL BIAS: Referral bias may have influenced the study result in this study.

Conflict Of Interest-There is no conflict of interest.

Acknowledgement: I acknowledge my co-author for his sincere effort.

Table 1. Americ	an Diabetic Ass	ociation C	lassification
of albuminuria			

Spot Collection	Timed	24-hr	Category
	Collection	Collection	
Less than 30	Less than 20	Less than	Normal
mcg/mg	mcg/min	30 mg	
creatinine			
30-300 mcg/mg	20-200	30-300 mg	Microalbumin
creatinine	mcg/min		uria

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More than 300	More th	an 200	More th	an Clinical				
mcg/mg	mcg/mi	n	300 mg	ng albuminuria				
creatinine								
Table 2. Age Dis	stributio	n						
AGE (IN YEARS)) FREQ	FREQUENCY		PERCENT				
<40 YEARS	10		20	0.0				
40-60 YEARS	25		50	0				
NO VEADO	16		20	0				
>60 YEARS 15 30.0				0.0				
Table 3. Sex dist	ribution							
SEX	FREC	DIENC	Y P	ERCENT				
	00	201110						
FEIMALE	22		44	44.0				
MALE	28	28		56.0				
Table 4FBS dis	stributio	n						
FBS	FREC	DUENC	Y P	ERCENTAGE				
<100	0	ç • • .		0				
100	0		0.	0.0				
100-126	26		52	52.0				
>126	24		48	48.0				
TABLE 5. PPBS	distribut	tion						
DDDC	TDT	JUENC	v D	FDCENTACE				
FFD3	I KEQ	SOFIAC:	· P.					
<140	8		16	5.0				
140-200	27		54	1.0				
>200	15		30	0.0				
			1.2					
TABLE 6. HbAlc	: distrib	ution						
HbAlc	FREÇ	QUENC	Y P	ERCENT				
> 6.5	43		86	6.0				
≤ 6.5	7		14	4.0				
TABLE 7. TC dis	stributio	n						
TC	FREQ	QUENC	Y P	ERCENTAGE				
ABNORMAL	24		48	3.0				
(>200 mg/dl)								
DESIRARIE	26		50	2.0				
(<200 mg/dl)	20		02	1.0				
(<200 mg/ai)								
TABLE 8. TG								
TG	FREC	DUENC	Y P	ERCENTAGE				
ABNORMAL.	31	•	60	2.0				
(> 150 mg/dl)	01		02	1.0				
(> 150 mg/m)								
DESIRABLE	19	19		3.0				
(<150 mg/dl)								
Table 9-I.DIDie	stributio	n_						
			7 D					
LDL-C	FREQ	QUENC	Y P.	ERCENTAGE				
ABNORMAL	21		42	2.0				
(>100 mg/dl)								
DESIRABL	29		58	58.0				
(< 100 mg/dl)				00.0				
(< 100 mg/ ai)								
TABLE 10. HDL-	C DISTI	ABUTIC	TABLE 10. HDL-C DISTRIBUTION					
UDI C								
HDR-C	FREQ	QUENC	Y P	ERCENTAGE				
ABNORMAL	40	QUENC	Y P	ERCENTAGE				
ABNORMAL (<50 mg/dl)	40	QUENC	Y P	ERCENTAGE				
ABNORMAL (<50 mg/dl)	40	QUENC	Ý P 80	ERCENTAGE				
ABNORMAL (<50 mg/dl) DESIRABLE	40 10	QUENC	Y P : 80 20	ERCENTAGE 0.0				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl)	40 10	QUENC	Y P 80 20	ERCENTAGE D.0 D.0				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN	FREQ 40 10 IE ACR I	QUENC	Y P 80 20 BUTION	ERCENTAGE 0.0 0.0				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN	FREQ 40 10 IE ACR I FREQ	QUENC'	Y P 80 20 BUTION Y P	ERCENTAGE				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN Urine ACR	FREQ 40 10 IE ACR I FREQ 24	QUENC' DISTRII QUENC'	Y P: 80 20 BUTION Y Y P:	ERCENTAGE 0.0 ERCENTAGE				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN Urine ACR ABNORMAL	FREQ 40 10 IE ACR I FREQ 24	QUENC' DISTRII QUENC'	Y P: 80 20 BUTION Y Y P: 48 48	ERCENTAGE 0.0 0.0 ERCENTAGE 3.0				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN Urine ACR ABNORMAL (> 30 mcg/mg	FREQ 40 10 IE ACR I FREQ 24	QUENCY DISTRII QUENCY	Y P: 80 20 BUTION P: Y P: 48 48	ERCENTAGE 0.0 0.0 ERCENTAGE 3.0				
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ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN Urine ACR ABNORMAL (> 30 mcg/mg Creatinine) NORMAL	FREQ 40 10 IE ACR I FREQ 24 26	QUENC DISTRII QUENC	Y P: 80 20 BUTION 48 48 52	ERCENTAGE 0.0 0.0 ERCENTAGE 3.0 2.0				
ABNORMAL (<50 mg/dl) DESIRABLE (> 50 mg/dl) TABLE 11. URIN Urine ACR ABNORMAL (> 30 mcg/mg Creatinine) NORMAL (<30 mcg/mg	FREQ 40 10 IE ACR I FREQ 24 26	QUENC' DISTRII QUENC'	Y P 80 20 BUTION Y Y P 48 52	ERCENTAGE 0.0 0.0 ERCENTAGE 3.0 2.0				
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TABLE 15. CIMT DISTRIBUTION					
CIMT		FREQUENCY	PI	ERCENTAGE	
INCREASED (> 0.8 mm)		11 22.0		.0	
NORMAL (<= 0.8 mm)		39 78.0		.0	
Table 16: SERUM C	CRE/	TININE			
SERUM CREATININE FREQUENCY PERCENT			PERCENT		
>1.5 (ABNORMAL)		4		8.0	
≤ 1.5 (NORMAL)		46		92.0	
Table 17. BMI DISTRIBUTION					
BMI	FRE	QUENCY	PI	ERCENTAGE	
≤ 22.9	12		24	.0	
23 - 27.5	20		40.0		
> 27.5	18		36	.0	



REFERENCES:

- Ferrannini E, Cushman WC. Diabetes and hypertension: the bad companions. 1. Lancet. 2012 Aug 11;380(9841):601-10. doi: 10.1016/S0140-6736(12)60987-8. PMID:22883509.
- Athyros VG, Doumas M, Imprialos KP, Stavropoulos K, Georgianou E, Katsimardou A, Karagiannis A. Diabetes and lipid metabolism. Hormones 2 (Athens). 2018 Mar;17(1):61-67. doi: 10.1007/s42000-018-0014-8. Epub 2018 Apr 16.PMID:29858856
- Jackson CE, Solomon SD, Gerstein HC, Zetterstrand S, Olofsson B, Michelson EL, Granger CB, Swedberg K, Pfeffer MA, Yusuf S, McMurray JJ; CHARM Investigators and Committees 2009 Albuminuria in chronic heart failure: 3 prevalence and prognostic importance. Lancet 374:543–5501. Tuttle KR, Bakris GL, Bilous RW, et al. Diabetic kidney disease: a report from an
- 4. ADA Consensus Conference. Diabetes Care 2014;37:2864-2883.
- Ito H, Komatsu Y, Mifune M, Antoku S, Ishida H, Takeuchi Y, et al. The estimated 5. GFR, but not the stage of diabetic nephropathy graded by the urinary albumin excretion, is associated with the carotid intima-media thickness in patients with type 2 diabetes mellitus: a cross-sectional study. Cardiovasc Diabetol.2010;9:18[DOI][PubMed].
- Temelkova-Kurktschiev TS, Koehler C, Leonhardt W, Schaper F, Henkel E, Siegert G, et al. Increased intimal-medial thickness in newly detected type 2 6. diabetes:risk factors.Diabetes Care. 1999;22(2):333-8[PubMed].
- 28. Gupta R, Joshi P, Mohan V, Reddy KS, Yusuf S. Epidemiology and causation of 7. coronary heart disease and stroke in India. Heart. 2008;94:16–26. [PubMed]. Gray N, Picone G, Sloan F, Yashkin A. Relation between BMI and diabetes
- 8. mellitus and its complications among US older adults. South Med J. 2015 Jan;108(1):29-36. doi: 10.14423/SMJ.000000000000214. PMID: 25580754; PMCID: PMC4457375.
- Diercks GF, Stroes ES, Van Boven AJ, Van Roon AM, Hillege HL, De Jong PE, et al. 9. Difference in the relation between urinary albumin excretion and carotid intima-media thickness in nondiabetic and type 2 diabetic subjects. Diabetes Care. 2002; 25(5): 936-7.
- 10. Kannel WB, McGee DL. Diabetes and glucose tolerance as risk factors for cardiovascular disease: The Framingham study. Diabetes Care. 1979;2:120-6. [PubMed].

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