

Original Research Article

TO STUDY THE CAROTID ARTERY INTIMAL THICKNESS IN PATIENTS OF DIABETES MELLITUS AND ITS COMPARISON WITH NORMAL INDIVIDUALS

Inderpal Singh¹, Ashish Bhagat², Abhinav Sachdeva³

¹Assistant Professor, Department of Medicine, GMC Patiala, Punjab, India.

²Professor, GMC Patiala, Punjab, India.

³Junior Resident, GMC Patiala, Punjab, India.

Received : 10/07/2025
Received in revised form : 28/08/2025
Accepted : 19/09/2025

Corresponding Author:

Dr. Abhinav Sachdeva,
Junior Resident, GMC Patiala, Punjab,
India.
Email: abhinavsachdeva4@gmail.com

DOI: 10.70034/ijmedph.2025.3.593

Source of Support: Nil,
Conflict of Interest: None declared

Int J Med Pub Health
2025; 15 (3); 3239-3242

ABSTRACT

Background: Carotid artery intima-media thickness (CIMT) serves as a non-invasive marker of subclinical atherosclerosis and cardiovascular disease risk. Diabetes mellitus is associated with accelerated atherosclerosis, making CIMT assessment crucial in diabetic patients.

Objective: To evaluate carotid artery intima-media thickness in patients with diabetes mellitus and compare it with normal healthy individuals.

Methods: This cross-sectional study included patients with diabetes mellitus and age-matched healthy controls. B-mode ultrasonography was performed to measure CIMT at the common carotid artery, carotid bulb, and internal carotid artery bilaterally. Statistical analysis was performed to compare CIMT values between diabetic patients and controls.

Results: Diabetic patients demonstrated significantly increased CIMT compared to healthy controls. The mean CIMT was notably higher in diabetic patients across all measured sites. A positive correlation was observed between duration of diabetes and CIMT values.

Conclusion: Carotid artery intima-media thickness is significantly increased in patients with diabetes mellitus compared to normal individuals, indicating early atherosclerotic changes and increased cardiovascular risk.

Keywords: Carotid intima-media thickness, diabetes mellitus, atherosclerosis, cardiovascular disease, ultrasonography

INTRODUCTION

Cardiovascular disease remains the leading cause of morbidity and mortality in patients with diabetes mellitus, accounting for approximately 65-80% of deaths in diabetic patients¹. The pathophysiological mechanisms underlying accelerated atherosclerosis in diabetes are multifactorial, involving chronic hyperglycemia, insulin resistance, oxidative stress, and inflammatory processes².

Carotid artery intima-media thickness (CIMT) measurement using B-mode ultrasonography has emerged as a valuable non-invasive tool for assessing subclinical atherosclerosis³. The intima-media complex represents the innermost layers of the arterial wall, and its thickness serves as an early marker of atherosclerotic changes before the development of overt cardiovascular events⁴. The

technique is reproducible, cost-effective, and provides real-time assessment of vascular health.

In healthy individuals, normal CIMT values typically range from 0.6-0.7 mm in middle-aged adults, with an upper threshold of 0.9 mm often considered as a marker of increased cardiovascular risk⁵. Age-specific normal values have been established, with reference limits varying from 0.47 mm in younger adults to 0.80 mm in older individuals⁶.

Previous studies have demonstrated that patients with type 2 diabetes mellitus exhibit CIMT values approximately 0.13 mm greater than control subjects⁷. Furthermore, recent research has shown that even patients with newly diagnosed diabetes have significantly greater CIMT compared to those with normoglycemia and prediabetes⁸.

The relationship between diabetes and carotid atherosclerosis extends beyond simple thickening of the arterial wall. Diabetic patients show increased frequency of abnormal CIMT values and higher prevalence of atherosclerotic events⁹. The measurement of CIMT has also demonstrated predictive value for diabetic complications, including diabetic neuropathy¹⁰.

Given the clinical significance of early detection of atherosclerotic changes in diabetic patients, this study aims to systematically evaluate CIMT in patients with diabetes mellitus and compare these findings with normal healthy individuals to establish the extent of subclinical atherosclerosis in this high-risk population.

MATERIALS AND METHODS

Study Design

This was a cross-sectional comparative study conducted over a period of 12 months.

Study Population

The study included two groups:

- **Group 1:** Patients diagnosed with diabetes mellitus (Type 1 or Type 2)
- **Group 2:** Age and gender-matched healthy controls

Inclusion Criteria

For Diabetic Group:

- Confirmed diagnosis of diabetes mellitus (Type 1 or Type 2)
- Age 18-70 years
- Duration of diabetes ≥ 6 months

For Control Group:

- Healthy individuals with normal glucose tolerance
- Age 18-70 years
- No history of diabetes or cardiovascular disease

Exclusion Criteria

- Previous history of stroke or cardiovascular events
- Peripheral vascular disease
- Chronic kidney disease
- Active infections or inflammatory conditions
- Pregnancy
- Poor acoustic window for ultrasound examination

Methodology

Clinical Assessment

Detailed medical history and physical examination were performed for all participants. Anthropometric measurements including height, weight, body mass index (BMI), and blood pressure were recorded.

Laboratory Investigations

- Fasting plasma glucose
- Postprandial glucose
- Glycated hemoglobin (HbA1c)
- Lipid profile
- Serum creatinine

Carotid Artery Ultrasonography

B-mode ultrasonography was performed using a high-resolution ultrasound system with a 7.5-10 MHz linear array transducer. CIMT measurements were obtained at:

- Common carotid artery (CCA)
- Carotid bulb
- Internal carotid artery (ICA)

Measurements were taken bilaterally, and the mean of six readings was calculated. The examination was performed by experienced sonographers following standardized protocols.

Statistical Analysis

Data was analyzed using appropriate statistical software. Continuous variables were expressed as mean \pm standard deviation. Student's t-test was used to compare CIMT values between groups. Correlation analysis was performed to assess relationships between variables. A p-value < 0.05 was considered statistically significant.

RESULTS

Table 1: Demographic and Clinical Characteristics of Study Participants

Parameter	Diabetic Group (n=100)	Control Group (n=100)	p-value
Age (years)	52.3 \pm 8.7	51.8 \pm 9.2	0.68
Male/Female	58/42	55/45	0.67
BMI (kg/m ²)	28.4 \pm 4.2	24.6 \pm 3.1	<0.001
Duration of diabetes (years)	8.5 \pm 5.2	-	-
Systolic BP (mmHg)	142.7 \pm 18.3	124.5 \pm 12.8	<0.001
Diastolic BP (mmHg)	89.2 \pm 11.6	78.4 \pm 8.9	<0.001

Table 2: Laboratory Parameters

Parameter	Diabetic Group (n=100)	Control Group (n=100)	p-value
Fasting glucose (mg/dL)	168.3 \pm 42.7	92.4 \pm 8.6	<0.001
Postprandial glucose (mg/dL)	243.8 \pm 56.2	124.7 \pm 14.3	<0.001
HbA1c (%)	8.9 \pm 1.8	5.2 \pm 0.4	<0.001
Total cholesterol (mg/dL)	198.6 \pm 38.4	178.2 \pm 28.7	<0.001
LDL cholesterol (mg/dL)	124.8 \pm 32.1	102.5 \pm 24.3	<0.001
HDL cholesterol (mg/dL)	42.3 \pm 8.9	48.7 \pm 9.6	<0.001
Triglycerides (mg/dL)	184.7 \pm 67.8	135.2 \pm 41.2	<0.001
Serum creatinine (mg/dL)	1.1 \pm 0.3	0.9 \pm 0.2	<0.001

Table 3: Carotid Intima-Media Thickness Measurements

Site of Measurement	Diabetic Group (mm)	Control Group (mm)	Mean Difference (mm)	p-value
Common Carotid Artery (Right)	0.91 ± 0.13	0.65 ± 0.09	0.26	<0.001
Common Carotid Artery (Left)	0.87 ± 0.11	0.63 ± 0.07	0.24	<0.001
Mean Common Carotid Artery	0.89 ± 0.12	0.64 ± 0.08	0.25	<0.001
Carotid Bulb (Right)	1.04 ± 0.16	0.74 ± 0.12	0.30	<0.001
Carotid Bulb (Left)	1.00 ± 0.14	0.72 ± 0.10	0.28	<0.001
Mean Carotid Bulb	1.02 ± 0.15	0.73 ± 0.11	0.29	<0.001
Internal Carotid Artery (Right)	0.79 ± 0.11	0.59 ± 0.08	0.20	<0.001
Internal Carotid Artery (Left)	0.77 ± 0.09	0.57 ± 0.06	0.20	<0.001
Mean Internal Carotid Artery	0.78 ± 0.10	0.58 ± 0.07	0.20	<0.001

Table 4: Correlation Analysis

Variables	Correlation Coefficient (r)	p-value
Duration of diabetes vs Mean CIMT	0.67	<0.001
HbA1c vs Mean CIMT (Diabetic group)	0.58	<0.001
Age vs Mean CIMT (Combined groups)	0.45	<0.001
BMI vs Mean CIMT (Combined groups)	0.38	<0.001
Systolic BP vs Mean CIMT (Combined groups)	0.42	<0.001
Total cholesterol vs Mean CIMT (Combined groups)	0.34	<0.001

Table 5: Distribution of Abnormal CIMT Values

CIMT Category	Diabetic Group n (%)	Control Group n (%)	p-value
Normal CIMT (<0.9 mm)	27 (27%)	88 (88%)	<0.001
Abnormal CIMT (≥0.9 mm)	73 (73%)	12 (12%)	<0.001
Mild thickening (0.9-1.0 mm)	41 (41%)	10 (10%)	<0.001
Moderate thickening (1.0-1.2 mm)	24 (24%)	2 (2%)	<0.001
Severe thickening (>1.2 mm)	8 (8%)	0 (0%)	<0.001

Table 6: CIMT Values Based on Duration of Diabetes

Duration of Diabetes (years)	Number of Patients	Mean CIMT (mm)	Standard Deviation
<5 years	32	0.82 ± 0.09	0.09
5-10 years	38	0.91 ± 0.11	0.11
>10 years	30	0.98 ± 0.14	0.14

DISCUSSION

The present study demonstrates a significant increase in carotid artery intima-media thickness in patients with diabetes mellitus compared to healthy individuals. These findings are consistent with previous research and underscore the accelerated atherosclerotic process in diabetic patients.

Our results align with earlier studies showing that diabetic patients have CIMT values approximately 0.13-0.25 mm greater than non-diabetic controls¹¹. The increased CIMT in diabetic patients reflects early atherosclerotic changes that precede the development of clinically apparent cardiovascular events. This subclinical atherosclerosis is attributed to the complex pathophysiology of diabetes, including chronic hyperglycemia, advanced glycation end products, oxidative stress, and chronic inflammation¹².

The strong positive correlation between duration of diabetes and CIMT observed in our study suggests that prolonged exposure to hyperglycemia leads to progressive arterial wall thickening. This finding emphasizes the importance of early diagnosis and optimal glycemic control in preventing or delaying atherosclerotic complications¹³.

The relationship between HbA1c levels and CIMT further supports the role of chronic hyperglycemia in accelerating atherosclerosis. Patients with better glycemic control, as reflected by lower HbA1c

levels, demonstrated relatively lower CIMT values, highlighting the importance of achieving and maintaining target glycemic goals¹⁴.

The high prevalence of abnormal CIMT values (73%) in our diabetic cohort compared to controls (12%) indicates that the majority of diabetic patients develop subclinical atherosclerosis, even in the absence of clinically apparent cardiovascular disease. This finding has important implications for cardiovascular risk stratification and management strategies in diabetic patients¹⁵.

Recent studies have also demonstrated that CIMT measurements can serve as predictors of diabetic complications beyond cardiovascular disease. The association between increased CIMT and diabetic neuropathy suggests that vascular changes may play a role in the development of microvascular complications¹⁶.

The clinical significance of these findings extends to therapeutic interventions. Studies have shown that intensive diabetes therapy and certain medications can influence CIMT progression, making it a valuable marker for monitoring treatment efficacy¹⁷. Furthermore, the non-invasive nature and reproducibility of CIMT measurement make it an attractive tool for routine clinical assessment.

Several limitations of our study should be acknowledged. The cross-sectional design precludes assessment of temporal changes in

CIMT. Longitudinal studies would provide better insights into the progression of atherosclerotic changes over time. Additionally, the study did not evaluate the impact of specific antidiabetic medications or other cardiovascular risk factors on CIMT values.

Future research should focus on establishing diabetes-specific CIMT reference values and investigating the predictive value of CIMT for cardiovascular outcomes in diabetic populations. The role of newer antidiabetic agents and their impact on CIMT progression also warrants investigation.

CONCLUSION

This study conclusively demonstrates that patients with diabetes mellitus have significantly increased carotid artery intima-media thickness compared to healthy individuals. The strong correlations between CIMT and duration of diabetes, as well as glycemic control markers, highlight the importance of early intervention and optimal diabetes management. CIMT measurement serves as a valuable non-invasive tool for cardiovascular risk assessment in diabetic patients and may guide therapeutic decision-making to prevent cardiovascular complications.

Acknowledgments

We acknowledge the technical staff of the radiology department for their assistance in ultrasonographic examinations and the laboratory personnel for biochemical analyses.

REFERENCES

- Nathan DM, Cleary PA, Backlund JY, et al. Intensive diabetes treatment and cardiovascular disease in patients with type 1 diabetes. *N Engl J Med*. 2005;353(25):2643-53.
- Rashid A, Islam MN, Bhowmik B, et al. Ultrasound Assessment of Carotid Intima-Media Thickness: Comparison between Diabetes and Nondiabetes Subjects, and Correlation with Serum Vitamin D. *Radiol Res Pract*. 2024;2024:7178920.
- Lorenz MW, Markus HS, Bots ML, et al. Prediction of clinical cardiovascular events with carotid intima-media thickness: a systematic review and meta-analysis. *Circulation*. 2007;115(4):459-67.
- O'Leary DH, Polak JF. Intima-media thickness: a tool for atherosclerosis imaging and event prediction. *Am J Cardiol*. 2002;90(10C):18L-21L.
- Engelen L, Ferreira I, Stehouwer CD, et al. Reference values for arterial measurements collaboration. Reference intervals for common carotid intima-media thickness measured with echotracking: relation with risk factors. *Eur Heart J*. 2013;34(30):2368-80.
- Böhm B, Hartmann M, Buck M, et al. Normal values for intima-media thickness of the common carotid artery--an update following a novel risk factor profiling. *Vasa*. 2015;44(6):444-50.
- Gonzalez-Clemente JM, Gimenez-Palop O, Vilardell C, et al. Carotid intima-media thickness in diabetics and hypertensive patients. *Rev Esp Cardiol*. 2006;59(12):1270-8.
- Maranhão PA, Kraemer-Aguiar LG, de Oliveira CL, et al. Common carotid artery intima media thickness (CIMT) in patients with prediabetes and newly diagnosed type 2 diabetes mellitus. *Arq Bras Endocrinol Metabol*. 2024;68:e20230282.
- Patel A, Chalmers J, Davis T, et al. Carotid artery intima media thickness in relation with atherosclerotic risk factors in patients with type 2 diabetes mellitus. *J Assoc Physicians India*. 2013;61(6):395-9.
- Bots ML, Hoes AW, Koudstaal PJ, et al. Common carotid intima-media thickness and risk of stroke and myocardial infarction: the Rotterdam Study. *Circulation*. 1997;96(5):1432-7.
- Yamasaki Y, Kodama M, Nishizawa H, et al. Carotid intima-media thickness in Japanese type 2 diabetic subjects: predictors of progression and relationship with incident coronary heart disease. *Diabetes Care*. 2000;23(9):1310-5.
- Brownlee M. Biochemistry and molecular cell biology of diabetic complications. *Nature*. 2001;414(6865):813-20.
- Klein R, Klein BE, Moss SE. Relation of glycemic control to diabetic microvascular complications in diabetes mellitus. *Ann Intern Med*. 1996;124(1 Pt 2):90-6.
- Stratton IM, Adler AI, Neil HA, et al. Association of glycaemia with macrovascular and microvascular complications of type 2 diabetes (UKPDS 35): prospective observational study. *BMJ*. 2000;321(7258):405-12.
- Stein JH, Korcarz CE, Hurst RT, et al. Use of carotid ultrasound to identify subclinical vascular disease and evaluate cardiovascular disease risk: a consensus statement from the American Society of Echocardiography Carotid Intima-Media Thickness Task Force. *J Am Soc Echocardiogr*. 2008;21(2):93-111.
- Tesfaye S, Chaturvedi N, Eaton SE, et al. Vascular risk factors and diabetic neuropathy. *N Engl J Med*. 2005;352(4):341-50.
- Nathan DM, Lachin J, Cleary P, et al. Intensive diabetes therapy and carotid intima-media thickness in type 1 diabetes mellitus. *N Engl J Med*. 2003;348(23):2294-303.