# Study the Correlation between Mean Platelet Volume and HbA1c in Type 2 Diabetes Mellitus with Special Reference to Microvascular Complication

S.N Bentoor<sup>1</sup>, Gourav Kumar<sup>2</sup>

<sup>1</sup>Professor, Department of Medicine, Shri B M Patil Medical College, BLDE (Deemed To Be University), Vijayapura, Karnataka, India. <sup>2</sup>Junior Resident, Department Of Medicine, Shri B M Patil Medical College, BLDE (Deemed To Be University), Vijayapura, Karnataka, India.

Received: January 2020 Accepted: January 2020

**Copyright:** © the author(s), publisher. Annals of International Medical and Dental Research (AIMDR) is an Official Publication of "Society for Health Care & Research Development". It is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

### ABSTRACT

Background: Diabetes mellitus is the most common endocrine disease characterized by metabolic abnormalities, hyperglycemia and by long term complications. Large platelet are younger, more active and aggregable, have dense granules, secrete more proaggregatory molecules. Platelet activation contributes to trigger thrombus formation and causing microcapillary embolization. Platelet is directly regulated by insulin via functional insulin receptor found on human platelets. Insulin inhibits platelet interaction with collagen and attenuates the platelet aggregation effect of agonist in healthy platelets. MPV, a determinant of platelet function, is a newly emerging risk factor for atherothrombosis. Increased in HbA1c concentration is directly proportional to increased MPV. MPV can emerge as an important, simple, effortless, and costeffective tool for monitoring and for early recognition of patients that could possibly benefit from preventive treatment. Methods: It is a comparative study conducted in 98 patients admitted in Sri B.M Patil Medical College and Hospital, who are already diagnosed to have Type 2 DM with more than five years duration and 98 non-diabetic subjects without known microvascular complication. MPV is measured in cases and control groups. Patients were selected on the basis of inclusion and exclusion criteria and statistically analyzed. The study was conducted between December 2017 to july 2019. Results: In our study, MPV was significantly higher in diabetics with HbA1c levels ≥ 6.5% than in diabetics with HbA1c levels <6.5%. Among the 98 study group participants total 92 patients had one or more microvascular complications. Of the microvascular complication evident in the form of retinopathy, nephropathy and neuropathy, diabetic neuropathy was the most common, followed by diabetic nephropathy. Conclusion: Changes in MPV are seen to be statistically associated with diabetes and its complications. They are easily available, simple, convenient, noninvasive, and easy to interpret method to determine platelet dysfunction and in turn predict the presence of microvascular complications.

Keywords: Diabetes Mellitus, HbA1C, Mean Platelet Volume.

## **INTRODUCTION**

Diabetes mellitus (DM) is the most common endocrine disease characterized by metabolic abnormalities, hyperglycemia, and by long term complications.

Large platelets are younger, more active and aggregable, have denser granules, secrete more proaggregatory molecules.

Platelet is directly regulated by insulin via a functional insulin receptor found on human platelets. Insulin inhibits platelet interaction with collagen and attenuates the platelet aggregation effect of agonist in healthy individuals.

Name & Address of Corresponding Author Dr. Gourav Kumar, Junior Resident, Department Of Medicine, Shri B M Patil Medical College, BLDE (Deemed To Be University), Vijayapura, Karnataka, India. MPV, a determinant of platelet function, is a newly emerging risk factor for atherothrombosis. In many studies it has been found that increased in HbA1c concentration is directly proportional to increased MPV.

## **MATERIALS AND METHODS**

To study correlation between mean platelet volumes (MPV) and HbA1c in type 2 diabetes mellitus. Determine if the MPV in the diabetic patients is higher compared to the non-diabetics. To see if there is a difference in MPV in diabetics with and without microvascular complications

#### **Inclusion Criteria**

- Patients already diagnosed to have Type 2 diabetes mellitus with more than five years duration of diabetes.
- Controls are non-diabetic.

#### **Exclusion Criteria**

Hb <13gm% - males

### Bentoor & Kumar; Correlation between Mean Flatelet Volume and HbA1c in Type 2 Diabetes Mellitus

- Hb <12gm% females
- Non-diabetic subjects with coronary artery disease (ECG changes).
- Patients on antiplatelet drugs such as aspirin and clopidogrel.
- Diabetic subject with less than five year duration of diabetes.

#### Study Design

The study has been carried out in 100 patients who are already diagnosed to have Type 2 DM with more than five years duration and 100 non-diabetic subjects without known microvascular complication.

## **RESULTS & DISCUSSION**

In this study, 100 individuals with T2DM and 100 nondiabetic controls were included. Both groups were age and gender matched. The majority of the individuals were between the age group of 31 60 yrs.

Comparison of Baseline characteristics between two groups								
Variables	Compared	Mean	Std.	t test/Mann	P value	Remark		
	groups		Deviation	Whitney U test				
Age	Study group	62.37	10.996	t=0.78	P=0.408	NS		
	Control group	61.26	11.576					
RBS	Study group	242.03	108.321	U=1169.500	P=0.0001	HS		
	Control group	118.70	27.950					
FBS	Study group	182.70	87.653	U=2486.50	P=0.0001	HS		
	Control group	112.00	20.252					
PPBS	Study group	211.83	83.051	U=1934.500	P=0.0001	HS		
	Control group	133.04	20.383					
HbA1C	Study group	9.069	2.65	U=727.000	<b>P=0.0001</b>	HS		
	Control group	5.723	0.880					
MPV	Study group	12.017	1.227	U=1204.500	<b>P=0.0001</b>	HS		
	Control group	9.737	1.365					
Platelet	Study group	2.541	0.837	U=4938.000	P=0.201	NS		
	Control group	2.433	0.916					
serum	Study group	1.707	1.827	U=4356.500	P=0.009	HS		
creatinine	Control group	0.892	0.272					

Correlation between HbA1C and MPV							
Correlation	Correlation	P value	Remark				
between	coefficient®						
HbA1C(<6.5)	r=0.2645	P=0.3050 NS	Moderate positive correlation.				
and MPV			statistically not significant				
HbA1C(≥6.5)	r=-0.428	P=0.0426	Moderate positive correlation.				
and MPV		Sign	statistically significant.				

The mean MPV of study group is 12.017 and that of control group is 9.737.

The correlation between HbA1C and MPV shows that at HbA1C <6.5, correlation coefficient  $\circledast$  with MPV was r=0.2645 with P=0.3050. Thus, there is

moderate positive correlation which is statistically not significant

At HbA1C  $\geq$  6.5, correlation coefficient® with MPV was r=0.428 with P=0.0426. Thus, there is moderate positive correlation which is statistically significant. It was also observed that the MPV were significantly higher in diabetic individuals with microvascular complications as compared to those without complications

Of the 100 diabetic patients in study group 92 individuals had microvascular complications. They were noted as diabetic retinopathy, diabetic neuropathy and diabetic nephropathy. The diabetic neuropathy was most common as a single microvascular complication. Few patients had any two of the three common microvascular complications.

Whereas, 4 patients had all three microvascular complications. Among all these presentations, diabetic neuropathy was most common as a singlemicrovascular complication at 28%.

Microvascular complications	No. of	Perecentage
	subjects	
Diabetic Retinopathy	7	7%
Diabetic Neuropathy	32	32%
Diabetic Nephropathy	28	28%
Diabetic Retinopathy+ Diabetic	4	4%
Nephropathy		
Diabetic Retinopathy+ Diabetic	7	7%
Neuropathy		
Diabetic Nephropathy+ Diabetic	10	10%
Neuropathy		
Diabetic Retinopathy+ Diabetic	4	4%
Nephropathy+ Diabetic Neuropathy		

# CONCLUSION

Changes in MPV are seen to be statistically associated diabetes and its complications. They are easily available, simple, convenient, noninvasive, and easy to interpret method to determine platelet dysfunction and in turn predict the presence of microvascular complications.

### REFERENCES

- Kodiatte TA, Manikyam UK, Rao SB, Jagadish TM, Reddy M, Lingaiah HKM, et al. Mean platelet volume in Type 2 diabetes mellitus. J Lab Physicians. 2012;4(1):5-9.
- Zuberi BF, Akhtar N, Afsar S. Comparison of mean platelet volume in patients with diabetes mellitus, impaired fasting glucose and non-diabetic subjects. Singapore Med J. 2008;49(2):114-6.
- 3. Guariguata L, Whiting DR, Hambleton I, Beagley J, Linnenkamp U, Shaw JE. Global estimates of diabetes prevalence for 2013 and projections for 2035. Diabetes research and clinical practice. 2014;103(2):137-49.
- 4. Hudspeth B. The burden of cardiovascular disease in patients with diabetes. The American journal of managed care. 2018;24(13 Suppl):S268-s72.
- Classification and Diagnosis of Diabetes: Standards of Medical Care in Diabetes-2018. Diabetes care. 2018;41(Suppl 1):S13-s27.

Bentoor & Kumar; Correlation between Mean Platelet Volume and HbA1c in Type 2 Diabetes Mellitus

- 6. Prasad RB, Groop L. Genetics of type 2 diabetes-pitfalls and possibilities. Genes (Basel). 2015;6(1):87-123.
- Hemminki K, Li X, Sundquist K, Sundquist J. Familial risks for type 2 diabetes in Sweden. Diabetes care. 2010;33(2):293-7.
- Skyler JS, Bakris GL, Bonifacio E, Darsow T, Eckel RH, Groop L, et al. Differentiation of Diabetes by Pathophysiology, Natural History, and Prognosis. Diabetes. 2017;66(2):241-55.
- Aly TA, Ide A, Jahromi MM, Barker JM, Fernando MS, Babu SR, et al. Extreme genetic risk for type 1A diabetes. Proceedings of the National Academy of Sciences of the United States of America. 2006;103(38):14074-9.
- Hu X, Deutsch AJ, Lenz TL, Onengut-Gumuscu S, Han B, Chen WM, et al. Additive and interaction effects at three amino acid positions in HLA-DQ and HLA-DR molecules drive type 1 diabetes risk. Nature genetics. 2015;47(8):898-905.
- Cooper JD, Howson JM, Smyth D, Walker NM, Stevens H, Yang JH, et al. Confirmation of novel type 1 diabetes risk loci in families. Diabetologia. 2012;55(4):996-1000.
- Long SA, Cerosaletti K, Wan JY, Ho JC, Tatum M, Wei S, et al. An autoimmune-associated variant in PTPN2 reveals an impairment of IL-2R signaling in CD4(+) T cells. Genes and immunity. 2011;12(2):116-25.
- Colli ML, Moore F, Gurzov EN, Ortis F, Eizirik DL. MDA5 and PTPN2, two candidate genes for type 1 diabetes, modify pancreatic beta-cell responses to the viral by-product doublestranded RNA. Human molecular genetics. 2010;19(1):135-46.
- 14. Törn C, Hadley D, Lee HS, Hagopian W, Lernmark Å, Simell O, et al. Role of Type 1 Diabetes-Associated SNPs on Risk of Autoantibody Positivity in the TEDDY Study. Diabetes. 2015;64(5):1818-29.
- 15. Insel RA, Dunne JL, Atkinson MA, Chiang JL, Dabelea D, Gottlieb PA, et al. Staging presymptomatic type 1 diabetes: a scientific statement of JDRF, the Endocrine Society, and the American Diabetes Association. Diabetes care. 2015;38(10):1964-74.

How to cite this article: Bentoor SN, Kumar G. Study the Correlation between Mean Platelet Volume and HbA1c in Type 2 Diabetes Mellitus with Special Reference to Microvascular Complication. Ann. Int. Med. Den. Res. 2020; 6(2):ME06-ME08.

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