

Sefika Burcak POLAT¹, Nagihan UGURLU², Fatma YULEK², Huseyin SIMAVLI³, Reyhan ERSOY¹, Ozcan EREL⁴, Bekir CAKIR¹,

¹Yildirim Beyazıt University, Ataturk Education and Research Hospital, Department of Endocrinology and Metabolism, Ankara, TURKEY

²Yildirim Beyazıt University, Faculty of Medicine, Department of Ophthalmology, Ankara, TURKEY

³Izzet Baysal Hospital, Department of Ophthalmology, Bolu, TURKEY

⁴Yildirim Beyazıt University, Faculty of Medicine, Department of Clinical Biochemistry, Ankara, TURKEY

BACKGROUND

➤ Diabetic retinopathy (DR) is the leading cause of blindness in the world.

➤ Retinopathy and nephropathy can still progress in diabetics despite optimal metabolic control.

➤ This suggests that factors other than hyperglycemia, such as abnormal hemostatic parameters, may play a role in the disease pathogenesis. Based on this hypothesis, several markers of hypercoagulation including fibrinogen, plasma activator inhibitor (PAI), and alpha-2-anti plasmin have been identified.

➤ The aim of the present study was to determine whether different degrees of DR (proliferative or non-proliferative) were associated with abnormally modulated hemostatic parameters in patients with type 1 diabetes mellitus (DM).

MATERIALS AND METHODS

➤ 52 type 1 diabetic patients and 40 healthy controls were enrolled in the study. Patients were then subdivided into three categories. Group I was defined as those without retinopathy, group II with non-proliferative retinopathy (NPRP), and group III with proliferative retinopathy (PRP).

➤ We have compared these subgroups with each other and the control group (group IV) according to the serum fibrinogen, plasminogen, α 2-anti-plasmin, and PAI.

RESULTS

➤ We detected that PAI-1 levels were higher in the diabetic groups than control, but this was not statistically significant whereas serum fibrinogen ($p=0.224$) and plasminogen ($p=0.224$) were similar between the diabetic and control groups.

➤ Alpha-2-anti-plasmin in groups I, II, and III was higher compared to the control group ($p<0.01$, $p<0.05$, and $p<0.001$, respectively) and the positive correlation identified between serum α 2-anti-plasmin and HbA1c levels ($r=0.268$, $p=0.031$).

Table: Biochemical and hemostatic measurements of diabetic patients and healthy controls

Variables	Group I (n:21)	Group II (n:18)	Group III (n:13)	Group IV (n:40)	p-değeri
HbA1c	9,2 (6,6-11,0) ^a	9,2 (6,0-11,0) ^b	8,5 (5,6-12,0) ^c	5,3 (4,4-6,3) ^{abc}	<0,001
PAI	182,7 (9,4-271,0)	158,0 (7,9-255,2)	162,1 (7,9-246,7)	155,6 (107,2-236,9)	0,209
α 2Antiplasmin	245,0 (31,0-650,0) ^a	202,0 (55,0-904,0) ^b	418,0 (42,0-1184,0) ^c	115,5 (23,0-591,0) ^{abc}	0,004
Plasminogen	90,0 (69,0-152,0)	102,0 (55,0-141,0)	107,0 (57,0-133,0)	87,0 (68,0-121,0)	0,244
Fibrinogen	259,8 \pm 72,6	277,3 \pm 62,8	307,1 \pm 81,3	272,1 \pm 42,3	0,224

a: Difference between Group I and Group IV is significant ($p<0,01$), b: Difference between Group II and Group IV is significant ($p<0,05$), c: Difference between Group II and Group IV is significant ($p<0,001$).

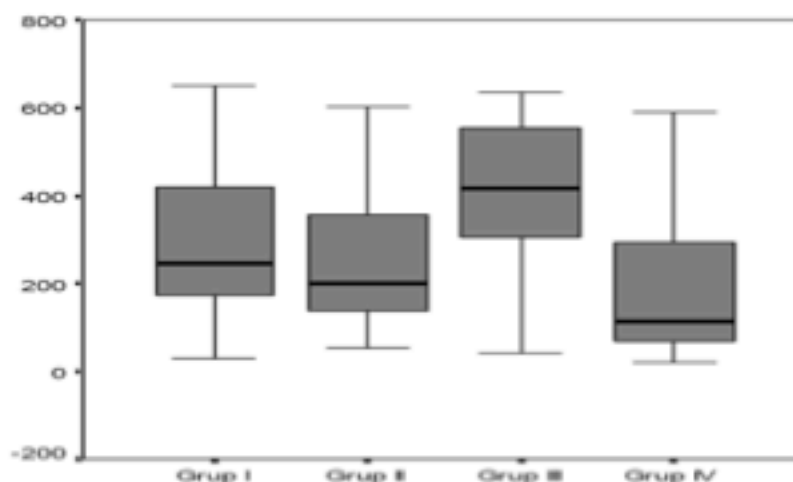


Figure: Expression of α 2-antiplasmin in the diabetic and control groups

CONCLUSION:

➤ To our knowledge there are only a small number of studies measuring α 2- antiplasmin levels in type 1 diabetes.

➤ A positive correlation between α 2 anti-plasmin with HbA1c suggests that fibrinolytic markers may improve with disease regulation, and better glycemic control.

➤ High α 2-anti-plasmin level might be a novel risk factor for development of DR. Confirmation of these data would allow a better understanding of the pathogenesis of DR.