

INTRODUCTION

- Nephropathy, retinopathy, neuropathy are long-term microvascular complications of diabetes.
- Glycated hemoglobin (HbA1c), used as a glycemic control indicator, have proven to be indicative in the development of microvascular complications.

AIM

In this study, it was aimed to evaluate the contribution of HbA1c variability to the development of complications.

METHOD

Twenty one cases with type 1 diabetes mellitus (T1DM) who developed microvascular complications and 39 cases without complications, that were similar in terms of age, gender, age of diagnosis, insulin treatment regimen, insulin doses (U / Kg), and mean HbA1c (%) levels were included in the study. Student t test and Mann Whitney U methods were used for statistical analysis and ROC analysis were used to determine limit values.

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GLYCATED HEMOGLOBIN VARIABILITY AND MICROVASCULAR COMPLICATIONS IN PATIENTS WITH TYPE 1 DIABETES MELLITUS

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RESULTS

Mean age of 21 cases (15 girls, 6 boys) who developed microvascular complications was 18.11 ± 4.39 years, and the mean duration of diabetes was 5.87 ± 3.93 years.

Mean age and duration of diabetes of 39 control patients (24 girls, 15 boys) were 13.25 ± 5.22 and 4.63 ± 3.33 years respectively.

Nephropathy was detected in 17 cases, neuropathy in 8 cases, and retinopathy in 1 case. Nephropathy occured at a mean age of 11.52 ± 4.12 years; neuropathy at 14.13 ± 5.68 years and retinopathy at 18.41 years.

Mean age of Type1 DM diagnosis in the complication group was 5.87 ± 3.93 years and 4.63 ± 3.33 years in the control group (p = 0.20).

Insulin infusion pump therapy (CSII) was initiated 3.12 ± 2.91 years after the diagnosis of diabetes and mean duration of CSII use was 7.45 ± 3.78 years. In the complication group 40% of the patients and in the control group 32.5 % were using CSII.

The mean HbA1c during follow up was similar in both of the groups (8.60% ± 0.63 vs 8.84% ± 1.32%). Adjusted HbA1c-Standard deviation (SD) and HbA1c-Variation coefficient (CV) values were 1.30 ± 0.65 and 14.36 ± 6.23 in the group with complications, and 0.91 ± 0.37 and 10.59 ± 4.01 in the control group (p <0.05).

In the ROC-analysis for microvascular complications, the limit value HbA1c-CV value was 11.99%. (sensitivity: 61.9%, specificity: 71.9; Youden index J: 0.3370). This value for HbA1c-SD was 0.9699. (sensitivity: 71.43%, specificity: 66.67; Youden index J: 0.3810)

CONCLUSIONS

This study has shown that longterm fluctuations in HbA1care associated with the development of microvascular complications in type 1 diabetes.

However, new studies with more patients are needed in this area.

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	Patients with complication n=21	Patients without complication n=39	þ	OR (95% CI)	þ
Male/female (n)	15/6	24/15	0.44	0,64(0.204-2.012	0,44
Age (yr)	18.11±4.39	13.25±5.22	0.001*	1.224(1.075-1.393)	0.002*
Age at diagnosis (yr)	5.87 ± 3.93	4.63 ± 3.33	0.20	1.102(0.950-0.279)	0.20
Duration of DM (yr)	12.2 ± 4.08	8,62 ± 4.19	0.002*	1.224(1.064-1.409)	0.005*
CSII/MDT(%)	40/60	32.5/67.5	0.56	1.385(0.455-4.213)	0.56
Number of HbA1c measurements	26.19±10.98	17.87±10.5	0.006*		
Mean HbA1c (%)	8.60 ± 0.63	8.84 ± 1.32	0.349	1.320(0.742-2.349)	0.345
HbA1c- SD	1.30 ± 0.65	0.91 ± 0.37	0.006*	4.602(1.37-5.45)	0.014*
HbA1c- CV	14.36 ± 6.23	10.59 ± 4.01	0.006*	1.165(1.033-0.313)	0.013*

REFERENCES





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