

Glycemic Control of Patients with Type 1 and Type 2 Diabetes Mellitus in Saudi Community

Khalid S Aljabri¹*, Samia A Bokhari¹, Muneera A Alshareef¹, Patan M Khan¹ and Bandari K Aljabri²

¹Department of Endocrinology, King Fahad Armed Forces Hospital, Jeddah, Kingdom of Saudi Arabia ²College of medicine, Um Al Qura University, Makkah, Kingdom of Saudi Arabia

*Corresponding Author: Khalid S Aljabri, Department of Endocrinology, King Fahad Armed Forces Hospital, Jeddah, Kingdom of Saudi Arabia.

Received: May 10, 2018; Published: June 08, 2018

Abstract

Background and Objective: The incidence of type 1 and type 2 diabetes mellitus is growing. Some national studies have measured glycemic control in people with diabetes. The objective of this study was to measure the glycemic control in people with type 1 and type 2 diabetes mellitus.

Methods: A cross section study of patients with type 1 and type 2 diabetes mellitus at the primary health care department and diabetic centre.

Main results: In total, 1612-patients were included in the study. There were 295 (18.3%) diagnosed with T1DM and 1317 (81.7%) with T2DM. There were 636 (39.5%) male and 976 (60.5%) were female with mean age 37.0 ± 7.9 . Mean HbA1c was 8.1 ± 2.3 , with significant differences in mean HbA1c between T1DM and T2DM patients. The correlation of, HbA1c with age and body mass index were (r = 0.05, p = 0.04) and (r = -0.14, p < 0.0001) respectively. HbA1c < 7.0 were achieved in 7.8% and 29.8 of T1DM and T2DM respectively, p = 0.047. There were significant differences in achieved HbA1c < 7.0 between genders for both T1DM and T2DM patients.

Conclusion: We found patients with type 1 and type 2 diabetes have poor glycemic control and that will increase their risk of diabetic complications.

Keywords: Diabetes Mellitus; HbA1c; T1DM and T2DM

Introduction

Diabetes mellitus is a major cause of excess mortality and morbidity. The prevalence and incidence of type 2 diabetes mellitus (T2DM) are increasing worldwide [1]. T2DM patients have a higher risk of developing microvascular and macrovascular disease than the general population. The occurrence of these complications depends largely on the degree of glycemic control as well as on the adequate control of cardiovascular risk factors [2-5]. Type 1 diabetes mellitus (T1DM) caused by destruction of the β -cells of the pancreas through a cellular-mediated autoimmune process [6]. Over the last three decades, the incidence of T1DM has been also on the rise worldwide [7-10]. At the same time, there is a trend towards diabetes being diagnosed at a younger age [11-14]. In Saudi Arabia, T1DM was prevalent in 0.2% of the Saudi males and 0.24% of the Saudi females [15]. Only 15% of patients with T1DM met the glycemic target, > 20% had very poor glycemic control (Hba1c > 8.8%) [16,17]. High prevalence of hyperglycemia (8.9 - 15%) was reported in different regions in Saudi Arabia [18]. The Diabetes Control and Complications Trial (DCCT) showed that good glycemic control has a significant effect on the development of nephropathy, retinopathy, and neuropathy of T1DM [19].

15

Gycosylated hemglobin (HbA1c) is used to evaluate the glycaemic control of diabetic patients [20,21]. HbA1c level < 7.0% as the primary glycaemic control target for diabetics is suggested by the American Diabetes Association (ADA) guidelines [6]. The percentage of patients who reach this objective seems to be notably lower (24%) in the case of less-controlled T2DM patients [22,23]. Increasing HbA1c levels were correlated with macrovascular and microvascular disease whereas decrease in HbA1c level decreases the prevalence of long term complications [24,25]. Therefore, we describe the glycemic control of population with T1DM and T2DM in our institution.

Methods

Older than 12 years old, had T1DM and T2DM were enrolled in the analysis. All patients were from the population of the Primary health and Diabetic Centres at King Fahad Armed Forces Hospital. A complete history and physical examination were taken, to have base-line laboratory assessments including HbA1c. HbA1c was expressed as percentage. High performance liquid chromatography was used. The HbA1c was divided into four groups; < 7.0, 7.0 - 7.9, 8.0 - 8.9 and \geq 9.0.

Statistical Analysis

Univariate analysis of baseline and follow up demography and clinical laboratory endpoints were accomplished using unpaired t-test where appropriate. Chi square (X^2) test were used for categorical data comparison. Pearson correlation was used for correlation. All statistical analyses were performed using SPSS Version 22.0. All P values were based on two-sided tests. The difference between groups was considered significant when P < 0.05.

Results

In total, 1612-patients completed the study. There were 295 (18.3%) diagnosed with T1DM and 1317 (81.7%) with T2DM. There were 636 (39.5%) male and 976 (60.5%) were female with mean age 37.0 ± 7.9 . Baseline characteristics are shown in table 1. Mean HbA1c was 8.1 ± 2.3 with significant differences in mean HbA1c between T1DM and T2DM patients. The correlation of HbA1c with age and body mass index were (r = 0.05, p = 0.04) and (r = -0.14, p < 0.0001) respectively.

Parameters		Total	T1DM	T2DM	P value
n (%)		1612	295 (18.3)	1317 (81.7)	
Age (years)		37.0 ± 7.9	25.9 ± 3.3	39.5 ± 6.3	< 0.0001
Gender	Male	636 (39.5)	96 (15.1)	540 (84.9)	0.007
	Female	976 (60.5)	199 (20.4)	777 (79.6)	
Body mass index (kg/m ²)		31.4 ± 6.0	31.4 ± 6.1	31.3 ± 6.0	0.9
HbA1c		8.1 ± 2.3	7.8 ± 2.3	8.2 ± 2.3	0.02

 Table 1: Characteristics of patients with type 1 and type 2 diabetes mellitus stratified by age, gender, BMI and HbA1c.

 Data are means ± SD or number (%).

HbA1c < 7.0 were achieved in 7.8% and 29.8 of T1DM and T2DM respectively, p = 0.047, figure 1. There were significant differences in achieved HbA1c < 7.0 between genders in both T1DM and T2DM patients. There were significant differences in the groups of HbA1c and the gender groups, figure 2.

Citation: Khalid S Aljabri., *et al.* "Glycemic Control of Patients with Type 1 and Type 2 Diabetes Mellitus in Saudi Community". *EC Endocrinology and Metabolic Research* 3.1 (2018): 14-20.



Figure 1: Characteristics of patients with type 1 and type 2 diabetes mellitus stratified by HbA1c < 7.0 to gender.



Figure 2: Characteristics of patients with type 1 and type 2 diabetes mellitus stratified by HbA1c category to gender.

Citation: Khalid S Aljabri., *et al.* "Glycemic Control of Patients with Type 1 and Type 2 Diabetes Mellitus in Saudi Community". *EC Endocrinology and Metabolic Research* 3.1 (2018): 14-20.

16

Discussion

This study showed the glycemic control in patients with T1DM and T2DM. Generalization to all population could not be due to regionalized characteristics. In addition, it does not evaluate the healthcare services offered in our city. The size of our sample and the crosssection type of the study should be of consideration.

The American Diabetes Association treatment guidelines suggest an HbA1c level < 7.0% as the primary glycaemic control target for diabetics, and a decrease in HbA1c level reduces the prevalence of chronic complications due to the disease [6,22]. In comparison to conventional therapy, the reductions in HbA1c levels achieved in the intensive therapy arm of the DCCT were associated with decreased rates in incidence and progression of microvascular complications. Tight glycaemic control was beneficial in preventing or delaying microvascular complications [19].

In our study, there was nonsignificant an inverse relationship of good glycemic control with older age among patients with T2DM (r = -0.025,p = 0.7), in addition, the average HbAlc of 8.1% was similar to the intensive treatment cohort of the DCCT (8.2%), higher to that described in Belgium with values of 6.6%, a cross-sectional multi-centered study in Europe, Japan and lower than United states, with values of 8.6%, Denmark (mean HbA1c 9.1%) and a cross-sectional study with children and adolescents in France (mean HbA1c 9.0%) and a population based study in Scotland with value of 9.1% [26-30]. To compare our results with those described above conclude that our services are providing satisfactory glycemic control results.

In our study, 92.2% and 70.1% were not able to achieve the American Diabetes Association goal in T1DM and T2 DM respectively and only 11.2% and 56.8% of patients in T1DM and T2DM respectively had a HbA1c concentration less than 8% comparable to other studies [26]. That increases our population risks for complications of diabetes.

Several reasons may contribute to failure of not achieving the American Diabetes Association goal. In clinical practice compared to trial settings, barriers to tight glycaemic control exist. More intensive insulin regimen even in young people with type 1 diabetes was the major factor in producing good glycemic control in the DCCT [31].

Optimizing glycaemic control measures were recently introduced. These include an intensive education programme; Dose Adjustment for Normal Eating (DAFNE), DAFNE may produce short-term but not long-term improvements in HbA1c levels [32]. Along with considerable nurse educator resources, a highly motivated patient is needed. Many patients move away from the clinic area or do not comply with regimens long-term.

Poor glycemic control was shown in diabetic patients who frequently missed appointments than those who missed none. Patients had a difference that is clinically relevant when missing more than 30% of scheduled visits, a HbA1c value 0.7 point higher relative to those with perfect attendance, a. missing appointments could have a direct effect on clinical outcomes by reducing continuity of care, measuring clinical variables or adjust medications, delaying the appropriate timing of interventions and screenings, and lacking a trusting providerpatient relationship. Frequent appointments cannot be offered as in clinical trials, due to resource limitations [33,34].

Many patients in the DCCT were treated with continuous subcutaneous insulin infusions (CSII), in contrast to no patients in our study. Achieving better glycaemic control was shown in some studies with the use of CSII over multiple daily injections in others not [35,36]. CSII is more expensive, requires an experienced diabetes team who can provide regular and frequent input into the ongoing care of the patient and a highly motivated patient without psychological problems. Studies with T1DM patients demonstrated that up to 62% of patients maintained a good glycemic control. Fluctuations of glycemia and, consequently, affect HbAlc could be attributed to different factors associated to general population can cause and upset the maintenance of good glycemic control [27-29]. The high prevalence of poor glycemic control shown in this study, reflects needs for more efforts for improvement.

Citation: Khalid S Aljabri., *et al.* "Glycemic Control of Patients with Type 1 and Type 2 Diabetes Mellitus in Saudi Community". *EC Endocrinology and Metabolic Research* 3.1 (2018): 14-20.

17

The burden of T2DM is on the increase worldwide, affecting more than 8% of the global adult population [37]. The risk of diabetic complications in patients with T2DM is strongly associated with the level of glycemia. The risk of microvascular and perhaps macrovascular complications of diabetes could be reduced by achieving tighter glycemic control [38]. There was nonsignificant an inverse relationship of glycemic control with age among patients with T2DM (r = -0.002, p = 0.9). That it could be due to developmental effect where cohorts mature, and their glycemic control improves as they age and may also be related to rapid changes in lifestyles witnessed in Saudi over the past five decades. More than half (63.4%) of participants with T2DM had poor glycemic control which is similar to that reported in Saudi Arabia where half of the studied populations had poor glycemic control [39]. In USA, data from National Health and Nutrition Examination Surveys reported that 42% - 50% of people with diabetes met the HbA1c target of 7% [40,41]. In the United Kingdom, a series of retrospective analysis of data found that 76% - 79% of patients had HbA1c > 7.5% [42].

Conclusion

These data indicate that many patients with T1DM and T2DM have poor glycemic control where they will be at high risk of diabetic complications. A structured primary and secondary intervention programs will be required to effectively manage this disease. More national studies are needed to assess glycemic control among diabetic patients in Saudi Arabia.

Acknowledgments

We are grateful to the staffs from the diabetic centre at King Fahad Armed Forces Hospital for their valuable contributions in data collection.

Conflict of Interest

The authors have no conflict of interest to disclose.

Bibliography

- 1. Mata M., et al. "The cost of type 2 diabetes in Spain: The CODE-2 study". Gaceta Sanitaria 16.6 (2002): 511-20.
- American Diabetes Association. "Implications of the United Kingdom Prospective Diabetes Study". *Diabetes Care* 25.1 (2002): S28-S32.
- Holman RR., et al. "10-year follow-up of intensive glucose control in type 2 diabetes". The New England Journal of Medicine 359 (2008): 1577-1589.
- 4. Gaede P., *et al.* "Effect of a multifactorial intervention on mortality in type 2 diabetes". *The New England Journal of Medicine* 358 (2008): 580-591.
- UK Prospective Diabetes Study (UKPDS) Group. "Intensive blood-glucose control with sulphonylureas or insulin compared with conventional treatment and risk of complications in patients with type 2 diabetes (UKPDS 33)". *Lancet* 352.9131 (1998): 837-853.
- 6. American Diabetes Association. "Classification and Diagnosis of Diabetes: Standards of Medical Care in Diabetes—2018". *Diabetes Care* 41.1 (2018): S13-S27.
- 7. C Gong., *et al.* "Trends in childhood type 1 diabetes mellitus incidence in beijing from 1995 to 2010: a retrospective multicenter study based on hospitalization data". *Diabetes Technology and Therapeutics* 17.3 (2015): 159-165.
- J Tuomilehto., et al. "Increase in incidence of insulin-dependent diabetes mellitus among children in Finland". International Journal of Epidemiology 24.5 (1995): 984-992.

Citation: Khalid S Aljabri., *et al.* "Glycemic Control of Patients with Type 1 and Type 2 Diabetes Mellitus in Saudi Community". *EC Endocrinology and Metabolic Research* 3.1 (2018): 14-20.

- 9. TheWriting Group for the SEARCHforDiabetes inYouth Study Group. "Incidence of diabetes in youth in theUnited States". *The Journal of the American Medical Association* 297.24 (2007): 2716-2224.
- 10. King H and Rewers M. "WHO ad Hoc Diabetes Reporting Group. Global estimates for the prevalence of diabetes mellitas and impaired glucose tolerante in adults". *Diabetes Care* 16.1 (1993): 157-177.
- 11. Soltesz G., *et al.* "Worldwide childhood type 1 diabetes incidence what can we learn from epidemiology?" *Paediatric Diabetes* 8.6 (2007): 6-14.
- 12. Maahs DM., et al. "Chapter 1: epidemiology of type 1 diabetes". Endocrinology Metabolism Clinics of North America 39.3 (2010): 481-497.
- 13. Patterson CC., *et al.* "Incidence trends for childhood type 1 diabetes in Europe during 1989-2003 and predicted new cases 2005-20: A multicenter prospective registration study". *Lancet* 373.9680 (2009): 2027-2033.
- 14. Tuomilehto J., *et al.* "Record-high incidence of type 1 (insulin-dependent) diabetes mellitus in Finnish children". *Diabetologia* 42.6 (1999): 655-660.
- 15. El-Hazmi MA., *et al.* "Diabetes mellitus and impaired glucose tolerance in Saudi Arabia". *Annals of Saudi Medicine* 16.4 (1996): 381-385.
- 16. Livingstone SJ., *et al.* "Risk of cardiovascular disease and total mortality in adults with type 1 diabetes: Scottish registry linkage study". *PLoS Medicine* 9.10 (2012): e1001321-e1001321.
- 17. Swedish National Diabetes Register. Annual Report (2012).
- 18. El-Hazmi MA and Warsy AS. "Comparative study of hyperglycemia in different regions of Saudi arabia". *The Annals of Saudi Medicine* 21 (1989): 43-48.
- The Diabetes Control and Complications Trial. "The effect of intensive treatment of diabetes on the development and progression of long-term complications in insulin-dependent diabetes mellitus". The Diabetes Control and Complications Trial Research Group". The New England Journal of Medicine 329.14 (1993): 977-986.
- 20. DeFronzo RA., *et al.* "Determination of glucose tolerance in impaired glucose at baseline in the Actos Now for Prevention of Diabetes (ACT NOW) study". *Diabetologia* 53.3 (2010): 435-445.
- 21. Goldstein DE. et al. "Tests of glycemia in diabetes". Diabetes Care 27 (2004):1761-1773.
- 22. Khaw KT., *et al.* "Association of hemoglobin A1c with cardiovascular disease and mortality in adults: the European prospective investigation into cancer in Norfolk". *Annals of Internal Medicine* 141.6 (2004): 413-420.
- 23. Mata-Cases M., *et al.* "Treatment of hyperglycaemia in type 2 diabetic patients in a primary care population database in a Mediterranean area (Catalonia, Spain)". *Journal of Diabetes and Metabolism* 5 (2014): 338.
- 24. Fowler MJ. "Microvascular and macrovascular complications of diabetes". Clinical Diabetes 26.2 (2008): 77-82.
- 25. XuD., et al. "Large scale simulation of red blood cell aggregation in shear flows". Journal of Biomechanics 46.11 (2013): 1810-1817.
- 26. Mortensen HB and Hougaard P. "Comparison of metabolic control in a crosssectional study of 2,873 children and adolescents with IDDM from 18 countries". *Diabetes Care* 20.5 (1997): 714-720.

Citation: Khalid S Aljabri., *et al.* "Glycemic Control of Patients with Type 1 and Type 2 Diabetes Mellitus in Saudi Community". *EC Endocrinology and Metabolic Research* 3.1 (2018): 14-20.

- 27. Dorchy H., *et al.* "Glycated hemoglobin and related factors in diabetic children and adolescents under 18 years of age: a Belgian experience". *Diabetes Care* 20.1 (1997): 2-6.
- Mortensen HB., *et al.* "A nation-wide cross-sectional study of urinary albumin excretion rate, arterial blood pressure and blood glucose control in Danish children with type 1 diabetes mellitus. Danish Study Group of Diabetes in Childhood". *Diabetic Medicine* 7.10 (1990): 887-897.
- 29. Rosilio M., *et al.* "Factors associated with glycemic control. A cross-sectional nationwide study in 2,579 French children with type 1 diabetes. The French Pediatric Diabetes Group". *Diabetes Care* 21.7 (1998): 1146-1153.
- 30. Scottish Study Group for the Care of the Young Diabetic. "Factors influencing glycemic control in young people with type 1 diabetes in Scotland: a population-based study (DIABAUD2)". *Diabetes Care* 24.2 (2001): 239-244.
- 31. "Diabetes Control and Complications Trial: Effect of intensive diabetes treatment on the development and progression of long-term complications in adolescents with insulin dependent diabetes mellitus". *Journal of Paediatric* 125.2 (1994): 177-188.
- 32. "Training in flexible, intensive insulin management to enabledietary freedom in people with type 1 diabetes: dose adjustment for normal eating (DAFNE) randomised controlled trial". *British Medical Journal* 325 (2002): 746.
- Parchman ML., et al. "Continuity of care, selfmanagement behaviors, and glucose control in patients with type 2 diabetes". Medical Care 40.2 (2002): 137-144.
- 34. Andrew J., et al. "Missed Appointments and Poor Glycemic Control an Opportunity to Identify High-Risk Diabetic Patients". Medical Care 42.2 (2004): 2110-2115.
- 35. Saurbrey N., *et al.* "Comparison of continuous subcutaneous insulin infusion with multiple insulin injections using the NovoPen". *Diabetic Medicine* 5.2 (1988): 150-153.
- 36. Hanaire-Broutin H., *et al.* "Comparison of continuous subcutaneous insulin infusionand multiple daily injection regimens using insulin lispro in type 1 diabetic patients on intensified treatment: a randomized study. The Study Group for the Development of Pump Therapy in Diabetes". *Diabetes Care* 23.9 (2000): 1232-1235.
- 37. International Diabetes Federation. IDF Diabetes Atlas, 5th edition. (2011).
- Stratton IM., et al. "Association of glycaemia with macrovascular and microvascular complications of type 2 diabetes (UKPDS 35): prospective observational study". BMJ 321.7258 (2000): 405-412.
- 39. Abdel-Gayoum A and Musa AS. "The effect of glycemic control on serum lipids and calcium phosphate profiles in patients with type 2 diabetes mellitus". *The Egyptian Journal of Biochemistry and Molecular Biology* 27 (2009): 79-92.
- Resnick HE., et al. "Achievements of American Diabetes Association Clinical practice Recommendations among U.S. Adults with Diabetes, 1999-2002 The National Health and Nutrition Examination survey". Diabetes Care 29.3 (2006): 531-537.
- 41. Saaddine JB., et al. "Improvements in diabetes processes of care and intermediate outcomes: United states, 1988-2002". Annals of Internal Medicine 144.7 (2006): 465-474.
- 42. Fox KM., *et al.* "Prevalence of Inadequate Glycemic Control Among patients with type 2 diabetes in the United Kingdom general practice research data base: A series of retrospective analysis of data from 1998 through 2002". *Clinical Therapeutics* 28.3 (2006): 388-395.

Volume 3 Issue 1 June 2018 ©All rights reserved by Khalid S Aljabri., *et al*.

20