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CORE

Association between Diabetes Consequences and Quality of Life among Patients with Diabetes Mellitus in the Aseer Province of Saudi Arabia

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Abstract

BACKGROUND: Diabetes mellitus (DM) is a major health-care burden worldwide. AIM: The aim of the study was to explore how the quality of life (QoL) of DM patients could be affected in the Aseer Province of the Kingdom of Saudi Arabia (KSA).

METHODS: A cross-sectional, multicenter study in DM patients of both sexes and all age groups in Aseer Province were done using a validated self-administered questionnaire. The study was conducted between April 1, 2018 and

RESULTS: A total of 418 patients completed our questionnaire, of which 240 (58%) were male and 178 (42%) were female. Furthermore, 50.23% were married and 104 (24.16%) were illiterate. We found that 403 (96.42%) respondents had type-2 DM and 315 (75.35%) had a family history of DM. In addition, 132 (31.57%) respondents were

on monotherapy whereas only 61 (14.59%) were using combination therapy. Hypertension was the most prevalent

CONCLUSION: DM had a significant impact on QoL among patients from Aseer Province in KSA. Our study underscores the importance of generating data on QoL among DM patients

comorbidity (166, 39.71%) and peripheral neuropathy the most prevalent complication of DM (157, 37.56%).

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Introduction

Diabetes mellitus (DM) is a chronic disease that can affect quality of life (QoL) if the desired treatment outcomes are not achieved [1], [2], Therefore, patients with DM should be counseled and educated about the impact of DM on their QoL, and encouraged to adhere to pharmacologic and non-pharmacologic management [3].

The prevalence of DM worldwide has been increasing worldwide and has been projected be ~370 million by 2030 [4]. There are two common types of DM, type 1 and type 2, and the latter represents ~90% of DM worldwide [5]. Over the past four decades, the Kingdom of Saudi Arabia (KSA) has witnessed major socioeconomic changes. As a result, the overall prevalence of DM among adults in KSA is ~25% [6].

QoL represents the ultimate goal of all health interventions [7]. It provides deep insight and information, irrespective of clinical data, on how the patient feels [8]. Evidence suggests that in DM, psychosocial factors play an important part in self-care, acceptance of therapeutic regimens, and treatment success and that metabolic measures (e.g., glycemic control) are poorly correlated with QoL [9]. DM complications have been found to have a negative effect on QoL [10]. Therefore, the main objective of DM treatment is to prevent DM complications and, therefore, improve the QoL of patients [1], [2]. An improvement in QoL not only benefits the patients but also reduces the social, financial, and psychological burden related to DM [1], [2].

The paucity of data related to QoL in the Aseer Province of KSA prompted us to evaluate the QoL of patients with DM and related complications.

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Methods

Ethical approval of the study protocol

The study protocol was approved from the Regional Ethical Committee of the Aseer Province (REC-2018-03-055). The purpose of the study was explained to respondents, and questions were administered only after seeking their consent for participation. The personal information of patients was not collected.

Study design

A cross-sectional study was conducted between April 1, 2018 and November 25, 2018. The study cohort was patients with type-1 and -2 DM in five primary care centers in the cities of Abha and Khamis Mushait in the Aseer Province in Southwest KSA.

Sample size and sampling method

The sample size (n) was calculated based on the Cochran equation. The calculated sample size was 385. A convenient sampling method was used in this study. Patients with DM were invited to participate in this study.

Study instrument and data collection

A self-administered questionnaire was adapted from the literature [11]. This validated self-administered questionnaire comprised 35 questions with four domains: Demographic and socioeconomic factors; clinical characteristics; QoL; and compliance of QoL.

Then, the study was carried out in a systematic and sequential manner. Step 1 involved development and validation of the questionnaire: A focus group of academicians and DM consultants determined the domains affecting QoL. Step 2 comprised a pilot study conducted for validation. The World Health Organization Brief Quality of Life Questionnaire was used for comparison [11]. Step 3 was questionnaire administration; the questionnaire was administered in person during daily visits to health facilities in which the study was carried out.

Statistical analysis

Data were analyzed descriptively using SPSS v21 (IBM, Armonk, NY, USA). The Kruskal–Wallis test of significance (p < 0.05) was undertaken to assess the association between adherence and DM-specific QoL. A multiple regression analysis (backward-selection method) was used to identify the independent predictors of DM-specific QoL.

Results

Patient demographics

Out of 439 study participants, 417 (95.2%) completed the questionnaire. Of these 417 participants, 240 (58%) were male and 178 (42%) were female. Most respondents were elderly (159, 38.08%). The lowest number of respondents (21, 5.02%) was <20 years of age. Furthermore, 210 (50.23%) respondents were married. In addition, 55 (13.15%) and 104 (24.16%) respondents were illiterate, whereas 94 (44.48%) had attained education at university level. Furthermore, 150 (35.88%) were unemployed whereas 49 (11.75%) were self-employed (Table 1).

Patient distribution by clinical characteristics and treatment type

Of the 418 people who completed the questionnaire, 403 (96.42%) had type-2 DM and 315 (75.35%) had a family history of DM. Furthermore, 148 (35.40%) had an underlying chronic history of DM from 5 years, whereas 27 (6.45%) were diagnosed recently (<1 year) with DM. In addition, 132 (31.57%) respondents were on monotherapy whereas 61 (14.59%) respondents were using combination therapy involving insulin. All patients had comorbidity, with hypertension being the most prevalent (166, 39.71%), followed by asthma (59, 14.11%) and dyslipidemia (58, 13.88%). Peripheral neuropathy was the most common DM-induced complication (157, 37.56%), followed by diabetic retinopathy (93, 22.25%) (Table 2).

Self-awareness of patients

We found that 226 (54.07%) respondents were well aware of hypoglycemic symptoms. Furthermore, 181 (43.30%) respondents were physically inactive and

Table 1: Demographics of questionnaire respondents

Characteristic	Number	Percentage
Sex		
Male	243	58
Female	174	42
Age (years)		
<20	21	5
20-39	56	13
40-59	136	33
60-79	157	38
>80	47	11
Marital status		
Unmarried	207	50
Married once	56	13
Married multiple times	70	17
Separated or divorced	53	13
Widowed	31	7
Education		
None	104	25
Primary level	101	24
Secondary level	118	28
University	94	23
Work status		
Government employee	78	19
Private employee	42	10
Self-employed	69	17
Unemployed	49	12
Pensioner	179	43

108 (25.84%) were carrying out physical activity for <30 min. In addition, 338 (88.85%) of respondents did not know about their level of glycated hemoglobin, and 368 (88.40%) were unaware of their body mass index. Worryingly, 163 (39.0%) of respondents did not follow their dietician's advice and 332 (79.42) did not use or kept a necklace/bracelet saying that they had DM with them.

Table 2: Patient distribution by clinical characteristics and treatment type

Characteristic	Number	Percentage
Type of DM		
1	399	96
2	18	4
Family history of DM		
Yes	317	76
No	100	24
How long you have been diagnosed with DM?		
<1 year	27	6
1–5 years	147	35
5-10 years	118	28
>10 years	125	30
Type of your therapeutic regimen		
Monotherapy	132	32
Combination therapy including metformin	184	44
Combination therapy without metformin	40	10
Combination therapy including insulin	61	15
Comorbidity		
None	93	22
One comorbidity	187	45
Two comorbidities	93	22
More than two comorbidities	44	11
DM complication		
None	64	15
One complication	80	19
Two complications	188	45
More than two complications	85	20
DM: Diabetes mellitus.		

We found that 372 (89%) respondents were satisfied with their medications and that 90 (22%) had an unpleasant experience with their existing drug therapy. Furthermore, 193 (46.45%) respondents felt worrisome about the possible side effects of their drug therapy and 95 (22.8%) used to skip their medications at times, with 194 (46.6%) of them experiencing some side effects from their drugs. In addition, 52.6% of patients "felt isolated" for being served a different diet and 64.3%

Table 3: Linear regression multivariable analysis for the relationship between demographic characteristics and overall QoL score

Variable	Standardized coefficient (beta)	95% CI	p-value
Sex			
Male	Reference	Reference	Reference
Female	-0.251	-8.8244.033	0.000
Age (years)			
<20	Reference	Reference	Reference
20-39	-0.048	-7.756-4.232	0.564
40-59	-0.290	-13.3112.326	0.000
60-79	-0.509	-18.7197.833	0.005
>80	-0.307	-18.4186.121	0.000
Marital status			
Unmarried	Reference	Reference	Reference
Married once	0.145	1.704-9.031	0.004
Married multiple times	-0.010	-3.726-3.035	0.841
Separated/divorced	-0.153	-9.5402.051	0.002
Widowed	-0.050	-7.1032.267	0.311
Education			
None	Reference	Reference	Reference
Primary level	0.098	-0.349 to 6.143	0.080
Secondary Level	0.264	4.278 to 10.529	0.000
University	0.416	9.292 to 15.90	0.000
Work status			
Government employee	0.252	3.91-12.44	0.000
Private employee	0.055	-2.63-7.284	0.358
Self-employed	-0.106	-7.99-0.754	0.105
Unemployed	Reference	Reference	Reference
Pensioner	-0.103	-6.40-1.120	0.168

QoL: Quality of life. CI: Confidence interval.

"hated being watched for that they ate." Furthermore, 77.9% "felt bad for not being able to eat what they like." More than 50% of respondents felt depressed because of the underlying disease and 91.1% worried about their children getting affected with the same disease (Tables 3 and 4).

Patient-related QoL

More than 70.0% of respondents stated that DM affected their general health and routine activities, such as working, shopping, and travelling. However, 23.4% of respondents stated that DM did not affect their capacity to work. Furthermore, 12% of respondents were undecided if DM affected their work capacity, whereas 66.26% felt strongly that absence from work was because of DM. More than 50% of respondents attributed their disassociation from family/social occasions to DM. In addition, 60.0% of Table 4: Linear regression multivariable analysis of the relationship between clinical characteristics and overall QoL score

Variable	Standardized coefficient (beta)	95% CI	p-value
Type of DM			
1	Reference	Reference	Reference
2	0.094	-0.145-12.142	0.056
Family history of DM			
Yes	Reference	Reference	Reference
No	0.234	4.146-9.704	0.000
How long you have been diagnose	d with DM?		
<1 year	0.040	-3.134-7.253	0.436
1–5 years	0.233	3.197-9.152	0.000
5–10 years	0.083	-0.8245.458	0.148
>10 years	Reference	Reference	Reference
Therapeutic regimen			
Monotherapy	Reference	Reference	Reference
Combination therapy including metformin	-0.31	-8.6063.154	0.000
Combination therapy without metformin	0.028	-3.127-5.50	0.584
Combination therapy including insulin	-0.102	-0.060-7.341	0.054
Comorbidity			
None	0.231	2.56-11.46	0.002
One comorbidity	0.25	-3.44-4.70	0.762
Two comorbidities	-0.27	-5.25-3.63	0.720
Three or more than three comorbidities	Reference	Reference	Reference
DM complications			
None	0.455	12.207-19.702	0.000
One complication	0.089	-0.648-6.469	0.109
Two compilations	0.117	-0.036-5.925	0.047
Three complications	Reference	Reference	Reference

QoL: Quality of life, CI: Confidence interval.

respondents stated that their disease had an impact on routine activities such as long talks, conversations, prayers, sleeping, and travelling on long road trips, and 29.2% stated that DM did not affect their relationship with friends/wife/children/coworkers. However, 50% of respondents were undecided on whether DM affected their sexual health, and 50% of respondents felt that DM affected their sexual health (Table 5).

Table 5: Mean QoL scores in various domains

Mean ± SD
38.63 ± 18.43
35.25 ± 14.447
49.57 ± 25.69
41.17 ± 12.662

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Discussion

QoL evaluation is a fundamental component when providing healthcare in patients with DM. In recent years, there has been growing appreciation of the patient's perspective on health, disease, and medical treatments. DM prevalence in KSA is underestimated. Only a few national/regional DM registries are available to support DM research and provide reliable data on various aspects of the disease [11]. We investigated QoL among patients with DM, with a focus on the clinical, mental and psychosocial factors that affect QoL.

Most respondents were elderly, married, and illiterate. A sex-specific significant difference in QoL was not found among study participants. These results are in accordance with data from Brown *et al.*, who reported that neither sex, education level nor age affected QoL significantly [12]. However, in western countries, lower QoL in women suffering from DM was reported by Rubin and Peyrot in 2004 [3]. Similar results have been reported by Redekop *et al.* and Aghamollaei *et al.* in 2002 and 2003, respectively [13], [14], [15].

We found that 96.42% of respondents had type-2 DM, with 75.35% having a family history of DM. Most respondents had an underlying chronic history of DM. Family history was found to be a stressful factor and reduced their QoL because they felt that their children were equally susceptible to the disease. Hypertension and asthma were common comorbidities, whereas peripheral neuropathy and retinopathy were common complications of DM. Moreover, respondents with neuropathic pain experienced a reduction in sleep. walking, and ability to undertake domestic duties. DM impairs all dimensions of health [11], [15]. Brown et al. reported that the type and duration of DM had an impact on QoL. DM complications, such as retinopathy and peripheral sensory neuropathy, have been associated with poor QoL in several studies, and painful diabetic neuropathy has a significant impact on QoL [16], [17].

Satisfaction with use of medications has a key role in terms of treatment compliance. Several studies have reported that long-term maintenance of drug therapy and glycemic control is dependent on adherence to medication and lifestyle changes. Most respondents (89%) were satisfied with the medications they were taking [18]. However, factors such as unpleasant taste and the side effects of medications had an impact on medication adherence among study respondents. More than 50% of respondents felt depressed because of their underlying disease and were worried about their children getting affected by DM. The results of our study are in accordance with those of Farias et al. [19]. They suggested that the QoL of patients with DM may influence treatment adherence, improve clinical outcomes satisfactorily, and reduce the prevalence of morbidity and mortality and disease progression. The relationship between QoL and treatment adherence remains controversial. Some scholars have shown the highest QoL level in patients with DM to be associated with better treatment adherence, whereas other researchers have not identified this association [19], [20].

Self-awareness and management of the complications and symptoms of DM have been positively correlated with QoL [21]. Self-management activities demand require effort, which many people find difficult to incorporate into their daily life [22], [23]. More than 50% of respondents were found to be self-aware about symptom management but these respondents were also found to be physically inactive. These results are in accordance with other studies that have reported that the duration of DM is associated with the level of DM knowledge [24]. Yun et al. and Kamel et al. observed a linear relationship between overall knowledge of DM and DM management. However, in another study, it was reported that people with DM lacked knowledge and, consequently, had low levels of self-care practices [25], [26]. Another study reported that exercise was associated with improved QoL among people with DM [27].

DM affects the QoL of the patient and his/her family due to its chronic nature, and its severity has been found to be associated with lower QoL [21]. Social isolation and dietary restriction had a negative impact on QoL among study respondents: >50% of respondents felt isolated for being served a different diet to that of other people. Moreover, monitoring what DM patients eat was found to have a negative impact on their QoL in the present study. Seventy-percent of respondents faced difficulties in routine activities, such as working, shopping, travelling, and praying.

Conclusion

DM had a significant impact on QoL among patients from the Aseer Province of KSA. Moreover, DM patients should be involved in awareness and counseling sessions to gain more knowledge regarding DM and its consequences. Our study underscores the importance of generating more data on QoL among DM patients.

Authors' Contributions

SMA and MJA conceived the study. AO, NAK, and YAH were involved in domain selection and questionnaire validation. MA, SSA, EV, and KA helped in data collection and statistical analyses. All authors participated in writing or reviewing the manuscript.

All authors declare that there are no conflicts of interest.

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