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Original Article

Subclinical hypothyroidism ups the risk of vascular complications in type 2 diabetes [☆]Ghada A. Mohamed ^a, Amira M Elsayed ^{b,*}^a Department of Internal Medicine, Assiut Faculty of Medicine, Assiut University, Egypt^b Department of Internal Medicine, Benha Faculty of Medicine, Benha University, Egypt

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ABSTRACT

The incidence of thyroid dysfunction in diabetic patients is higher than that of the general population. Undiagnosed thyroid dysfunction may affect the metabolic control and enhance cardiovascular, and other chronic complication risks in diabetic patients. Few studies have examined the relationship between subclinical hypothyroidism (SCH) and vascular complications of type 2 diabetes. *Objectives:* To find out the relationship between SCH and vascular complications in patients with Type 2 diabetes. *Subjects and Methods:* Our cross sectional study included 110 patients with type 2 DM (45 males and 65 females) who were followed at the Diabetes outpatient Clinics in the state of Kuwait during 6 months period. All patients subjected to complete clinical and laboratory data, including thyroid function tests, total cholesterol (TC), triglyceride (TG), HDL-C, LDL-C, urinary albumin, fundus examination, ECG, and Glycosylated hemoglobin. *Results:* Among 110 patients, 21 (19.1%) Patients had SCH. Patients with SCH were more significantly older, with longer duration of diabetes, higher HbA1c, total cholesterol and LDL-C than euthyroid group. However, gender ($p = 0.076$), BMI ($p = 0.092$), and smoking ($P = 0.715$) were not significantly different between the SCH and euthyroid groups. The SCH group had a higher prevalence of dyslipidemia ($p = 0.017$), diabetic nephropathy ($p = 0.003$) diabetic retinopathy ($p = 0.004$) and IHD ($p = 0.011$) than the euthyroid group while no significant difference in the prevalence of diabetic neuropathy ($p = 0.420$). *Conclusions:* SCH is a common endocrine disorder in patients with Type 2 diabetes. It could be associated with a higher prevalence of vascular complications in type 2 diabetes. We could not prove a relation between SCH and diabetic neuropathy.

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1. Introduction

Subclinical hypothyroidism (SCH) is defined as an asymptomatic state that is characterized by high serum concentration of thyroid-stimulating hormone (TSH) and a normal serum concentration of thyroxine.¹ The incidence of thyroid dysfunction in diabetic patients is higher than that of the general population and up to a third of patients with type-1 diabetes (T1DM) and 4–17% in type- 2 diabetic patients. Undiagnosed thyroid dysfunction may affect metabolic control and enhance cardiovascular, and other chronic complication risks in diabetic patients.² The association between SCH and ischemic heart disease has been noticed in several studies, although the results have been inconsistent^{3,4} and very few studies have focused on the population with diabetes.

Few studies have examined the relationship between subclinical hypothyroidism and microvascular complications in type 2 diabetic patients. To our knowledge, only three studies have investigated the association between SCH and microvascular complications in Type 2 diabetes. Both Yang et al.⁵ and Kim et al.⁶ found that SCH was associated with diabetic retinopathy, although not with nephropathy. By contrast, Chen et al.⁷ reported that patients with Type 2 diabetes and SCH were at increased risk of nephropathy. Moreover, no study to our knowledge has examined the prevalence of neuropathy in association with subclinical hypothyroidism.

Owing to the paucity of data regarding the prevalence of SCH and its potential association with vascular complications in Type 2 diabetes, this study examined the relationship between SCH and vascular complications patients with Type 2 diabetes.

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2. Methods

2.1. Patients

Our cross sectional study included 110 patients with type 2 DM (45 males and 65 females) who were followed at the Diabetes outpatient Clinics in the state of Kuwait during 6 months period were subjected to thyroid function test (FT3, FT4, TSH). Patients with known overt hypothyroidism, known overt or subclinical hyperthyroidism and type 1 diabetes mellitus, pregnancy, Postpartum women, patients with pituitary tumor, thyroid carcinoma or severe heart, liver, kidney and brain diseases were excluded from the study.

2.2. Thyroid function tests and definition

Serum thyrotropin (TSH) levels and free thyroxine levels were measured by chemiluminescent immunoassay. Subclinical hypothyroidism was defined as an elevated TSH level (>4.2 mIU/L) and a normal FT4 level (10–23 pmol/L). Euthyroidism was defined as normal levels of all three parameters: free triiodothyronine, free thyroxine and TSH.

2.3. Clinical examination and laboratory measurements

Relevant data about the patients were collected, including age, gender, duration of diabetes, history of hypertension, BMI, systolic and diastolic blood pressure, total cholesterol (TC), triglyceride (TG), HDL and LDL cholesterol. Glycosylated hemoglobin was measured using an ion-exchange HPLC (Merck-Hitachi9100).

2.4. Assessment of diabetic microvascular and macrovascular complication

The patients were diagnosed as having CAD if they had ischemic changes on a resting electrocardiogram (e.g., abnormal Q waves, ST segment depression or inverted T waves), a history of angina, myocardial infarction, or coronary angiography. Diabetic retinopathy was assessed by ophthalmologists and graded according to the International Clinical Classification of Diabetic Retinopathy.⁸ Diabetic nephropathy was defined as history of diabetes with the presence of albuminuria, impaired glomerular filtration rate, or both.⁹ A single morning urine sample for albuminuria was taken from subjects and albuminuria defined if equal to or more than 30 mcg/mg Creatinine (mcg/mg Cr).¹⁰ Diabetic neuropathy was diagnosed based on abnormalities detected during neuropathic screening test (vibration sense testing, temperature sense, pain perception, monofilament testing and examination of reflexes).

2.5. Statistical analyses

Statistical analyses were performed using SPSS software version 16.0, and results were expressed as mean \pm SD. To compare the means between two groups, Student's *t*-test was used for normally distributed data and the Mann–Whitney test was used for non-normally distributed data. For categorical comparisons, the chi-square test was used. A *P*-value < 0.05 was considered to be statistically significant.

3. Result

Our cross sectional study include 110 patients (45 males and 65 females) with type 2 DM were enrolled in our study with the following characteristics, age, 58.9 ± 10.6 (mean \pm SD) years; duration of DM, 17.7 ± 10.02 years; BMI, 31.8 ± 6.5 kg/m²; HbA1c,

$8.4 \pm 1.8\%$; TC, 4.08 ± 0.93 mmol/l; triglycerides (TG), 1.55 ± 0.76 mmol/l; HDL-C, 1.22 ± 0.38 mmol/l; LDL-C, 2.09 ± 0.75 mmol/l; TSH, 3.02 ± 1.93 mIU/l; T4 (15.28 ± 2.07) pmol/l; urinary albumin creatinine ratio (185.27 ± 379.66) mcg/mg Cr. It was found that (105, 95.5%) of patients had dyslipidemia, (43, 39.1%) had diabetic retinopathy, (47, 42.7%) had diabetic nephropathy, (29, 26.4%) had diabetic neuropathy, (37, 33.6%) had IHD, (82, 74.5%) were hypertensive and (28, 25.5%) were smoker. Among 110 patients, 21 (19.1%) had SCH. Patients with SCH were significantly older, with longer duration of diabetes, higher HbA1c, total cholesterol and LDL-C than euthyroid group. However, gender ($p = 0.076$), BMI ($p = 0.092$), and smoking ($P = 0.715$) were not significantly different between the SCH and euthyroid groups (Table 1). The SCH group had a higher prevalence of dyslipidemia ($p = 0.017$), diabetic nephropathy ($p = 0.003$) diabetic retinopathy ($p = 0.004$) and IHD ($p = 0.011$) than the euthyroid group while no significant difference in the prevalence of diabetic neuropathy ($p = 0.420$) (Table 2 & Fig. 1).

4. Discussion

Vascular endothelial dysfunction is supposed to be an important factor in the pathogenesis of microvascular and macrovascular complications in diabetes. Endothelial dysfunction has also been described in subclinical hypothyroidism as well as in those with upper normal TSH values.^{11,12} The association between microvascular complications and subclinical hypothyroidism and TSH values has not been comprehensively studied. Among the few studies done on the subject, the findings have been opposing. This is a cross-sectional study with a case control analysis which revealed that The prevalence of subclinical hypothyroidism in type 2 diabetic patients was 19.1%, and this is higher than data reported from another study done by Gao et al.¹³ Our present data revealed that the prevalence of subclinical hypothyroidism was 4.5% in males and 14.5% in females with Type 2 diabetes. NHANES III, a population study, reported that the prevalence of subclinical hypothyroidism was 3.4% in males and 5.8% in females.¹⁴ Perros et al. found that the occurrence of subclinical hypothyroidism was 3.3% in men and 4.6% in women with Type 2 diabetes.¹⁵ The incidence of subclinical hypothyroidism in our study was greater than those of both these studies and revealed a dominance of females. The patients with SCH had a significant longer duration of diabetes in comparison to the euthyroid group ($p = 0.007$). In contrast, Kim et al. didn't found any significant difference in the duration of diabetes between euthyroid and SCH patients.⁶ Although obesity affects hypothalamic-pituitary-thyroid axis directly or indirectly leading to variations in thyroid function

Table 1

Comparison between the participants with Euthyroid and SCH in type 2 diabetic patients regarding the different clinical parameters.

	Euthyroid (n = 89)	SCH (n = 21)	P value
Age (years)	57.78 \pm 11.045	63.43 \pm 6.95	0.027
Sex (M/F)	40/49	5/16	0.076
Duration of DM(years)	16.64 \pm 10.245	22.10 \pm 7.77	0.007
BMI (kg/m ²)	31.27 \pm 6.19	34.2143 \pm 7.36	0.092
HbA1c (%)	8.17 \pm 1.72	9.15 \pm 1.78	0.025
Total cholesterol (mmol/l)	3.85 \pm 0.79	5.05 \pm 0.83	<0.001
Triglycerides (TG) (mmol/l)	1.54 \pm 0.81	1.57 \pm 0.50	0.322
LDL cholesterol (mmol/l)	1.89 \pm 0.61	2.93 \pm 0.73	<0.001
HDL cholesterol (mmol/l)	1.17 \pm 0.39	1.41 \pm 0.27	0.001
TSH (mIU/l)	2.24 \pm 0.74	6.37 \pm 1.88	<0.001
T4 (pmol/l)	15.49 \pm 2.11	14.38 \pm 1.66	0.027
UAE	126.43 \pm 303.67	434.63 \pm 546.92	0.011
Dyslipidemia (%)	97.8	85.7	0.017
HTN (%)	76.4	66.7	0.357
Smoking (%)	24.7	28.6	0.715

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