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CARDIOVASCULAR RISK IN PATIENTS WITH SUBCLINICAL HYPOTHYROIDISM



Medicine

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ABSTRACT

Thyroid hormone regulate every organ system in our body. Metabolic parameters, including lipoprotein metabolism and cardiovascular risk factors like obesity, glucose intolerance or type 2 diabetes, dyslipidemia and hypertension are all influenced by serum TSH level in our body. Several reports have demonstrated a positive correlation between TSH levels and cardiovascular risk factors. C-reactive proteins is one of the markers of inflammation and also considered to be the predictor of cardiovascular disease and may be helpful in the assessment of future development of cardiovascular abnormalities in subclinical hypothyroidism.

Methods: Recently diagnosed 71 subclinical hypothyroidism patients were enrolled in the study and compared with 63 healthy individuals. Patients were divided into two groups on the basis of thyroid stimulating hormone (TSH) concentration. Thyroid profile (TSH, free thyroxine, triiodothyroinine) and C-reactive protein were estimated by enzyme-linked immunosorbent assay. Total cholesterol (TC), triglycerides (TG), and high-density lipoprotein-cholesterol were estimated in the study population by a spectrophotometric method, and low-density lipoproteincholesterol (LDL-C) was measured by the Friedewald formula.

Results and Conclusion: Altered lipid profile, as well as higher concentrations of C-reactive protein, was observed in subclinical hypothyroidism patients as compared to controls. Higher concentration of TC, TG, and LDL was observed in subclinical hypothyroidism patients while the concentration of high-density lipoprotein was lower in patients with subclinical hypothyroidism. The elevated concentration of C-reactive protein was also observed in the patient group. Patients having TSH >10 µIU/ml have a higher risk of developing cardiovascular disease compared to patients having TSH < 10 µIU/ml and increasing concentration of C-reactive protein along with LDL-C may to cardiovascular abnormality.

KEYWORDS

Subclinical hypothyroidism, C-reactive protein, Dyslipidemia, Cardiovascular risk.

INTRODUCTION

Subclinical hypothyroidism (SCH) is a common endocrine disorder presented by mildly elevated thyroid stimulating hormone (TSH) above the reference range along with a normal concentration of free thyroxine (FT4) and triiodothyronine (T3). It is an oscillated state between euthyroidism to hypothyroidism as it may lead to overt hypothyroidism. Thyroid hormones are known to play a significant role in cardiac hemodynamic because the deficient or excessive concentration of thyroid hormones may develop cardiac abnormalities. It is well-established that overt hypothyroidism is associated with cardiovascular symptoms through biochemical analysis. Dyslipidemia is a prominent feature of hypothyroidism established by various evidences, but not for the SCH due to inconsistent findings over the years.

The diversity of the result has raised the question that whether SCH is characterized by cardiovascular symptoms or not. C-reactive protein (CRP), an acute-phase reactant, is an effective and stable tool for the assessment of cardiovascular risk. It is an important marker of various inflammatory disorders like rheumatoid arthritis, myocardial infarction and various others etc. CRP is known to play an important role in the progression of atherosclerosis, a cardiovascular disease caused by the altered mechanism of low-density lipoprotein cholesterol (LDL-C). Therefore, the main objective of this study is an assessment of the role of CRP in SCH patients.

MATERIAL AND METHODS

This cross-sectional study was carried out in Department of Medicine, Darbhanga Medical College and Hospital, Laheriasarai, Bihar. Total 71recently diagnosed SCH patients were enrolled for the study, in which 44 patients were having TSH 6.16- 10 µIU/ml and rest 27 patients were having TSH > 10 µIU/ml termed as Cases-I and Cases-II respectively, compared with 63 healthy individuals as controls. People with a previous history or family history of thyroid diseases, taking thyroid medication, cardiovascular disease, smoking, menopausal women, pregnancy, inflammation, diabetes, hypertension, or alcoholics were excluded from this study. Age group criteria were kept from 21 to 45 years for the study population.

Body mass index (BMI) was calculated by means of height and weight of an individual. The weight of a person is divided by the square of the height. The weight and height of an individual were measured in kilogram (kg) and meter (m), respectively. The person having BMI

18.5 kg/m²to 24.99 kg/m²was considered a normal BMI. Thyroid profile (TSH, FT4 and T3) of SCH patients, as well as controls, was measured by enzyme-linked immunosorbent assay (ELISA). Patients with a concentration of TSH>6.2 $\mu IU/ml$ along with a normal concentration of FT4 and T3 was considered to have SCH. ELISA kits was used for the investigation of thyroid profile estimation.

Total cholesterol (TC), triglycerides (TG), high-density lipoprotein cholesterol (HDL-C), and LDL-C were measured in the study group. TC, TG, and HDL-C were investigated by means of glucose oxidase (CHOD)/peroxidase (POD) method, GPO-PAP method and CHOD-POD/phosphotungstate method, respectively.

STATISTICALANALYSIS

All the variables (age, BMI, TSH, FT4, T3, lipid profile, and CRP) were expressed in mean \pm standard deviation (SD). One-way ANOVA was used for the differentiation of all the parameters between the various groups (Control group, Cases-I and Cases-II). A p<0.05 was considered statistically significant. IBM SPSS version 20 (Statistical package for social sciences) was used for the statistical analysis.

This study shows a statistical difference among the various groups regarding the different variables. The age of all the participants was not significantly different statistically. Thyroid profile (TSH and T3) was significantly different among the groups except the FT4. BMI was significantly different among the groups and was highest in Cases-II and lowest in control groups (Table 1).

Lipid profile was significantly different among the groups. TC, TG, and LDL-C were higher in SCH groups compared to control group and highest in Cases-II. HDL-C was significantly different among the groups. The lowest concentration of HDL-C was observed in the second group of SCH patients. The concentration was also lower in Cases-I comparing to control group. The level of CRP was significantly different among the groups as its level was comparatively increasing in each group, from Cases-I to the Cases-II (Table 2).

Table 1: Baseline characteristics among the various groups ‡

Sl. No.	Parameters	Control (0.39-6.16 mIU/ml)	Case I (6.16-10 mIU/ml)	Case II (>10.0 mIU/ml)	p-value				
1	Age (years)	35.46±3.77	35.75±3.71	36.25±2.96	0.628				
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All the variables were expressed as mean±SD. ‡By analysis of variance, SD: Standard deviation, BMI: Body mass index, TSH: Thyroid stimulating hormone, FT4: Free thyroxine, T3: Triiodothyroinine

Table 2: Lipid profile and CRP among the various groups ‡

2	BMI (kg/m ²)	22.68±1.82	26.4±2.13	29.64±3.83	< 0.01
3	TSH (mIU/ml)	2.75±0.78	8.20±0.82	14.86±2.95	< 0.01
4	FT4 (ng/dl)	1.15±0.20	1.15±0.24	1.25±0.30	0.196
5	T3 (ng/dl)	1.08±0.28	0.95 ± 0.18	0.93±0.15	< 0.01

All the variables were expressed as mean±SD, ‡By analysis of variance, CPR: C-reactive protein, TC: Total cholesterol, TG: Triglycerides, HDL: High-density lipoprotein, LDL: Low-density lipoprote

This study shows that SCH patients are characterized by increased

BMI, lipid profiles, and CRP level. Rotterdam Study specified that

DISCUSSION

subclinical hypothyroidism is a strong predictor of myocardial infarction and risk of atherosclerosis. Elevated TSH is found to be associated with increased BMI in SCH patients. Srideviet al. supported our study by signifying altered concentration of lipid profile as characterized by significantly increased level of TC, a higher concentration of TG and elevated LDL-C. Similar to this study Erdemet al. observed that the concentration of HDL-C was significantly lower in SCH patients compared to controls.Study ofKvetnyet al. also support this study that SCH might be a risk factor for the development of cardiovascular diseases due to increased level of TG and CRP. CRP, a plasma protein, predominantly synthesized in liver, emerged as an additional cardiovascular risk factor was found to be significantly higher in the SCH group compared to control group. CRP has been reported to bind to oxidized LDL and actively participates in atherogenesis by direct influence on complement system, vascular cell activation, lipid accumulation and thrombosis. The level of CRP was significantly higher in SCH patients promising the approach for further development of cardiac risk, supported by Gaoet al. study. Gupta et al. also observed the increased concentration of inflammatory markers in SCH patients. While contrary to it other studies defined that SCH has no relation with dyslipidemia. Luboshitzkyet al. described that CRP does not play any role to contribute to the increased risk of cardiovascular disease in SCH patients. Toruneret al. described that SCH is characterized by dyslipidemia, but this association does not depend on the range of

TSH. This study proposed that SCH patients having TSH>10 µIU/ml

have a higher risk of future development of cardiac abnormalities

matched with Hernandez-Mijares and Marwaha study. In addition to this patients having SCH with TSH<10 µIU/ml also showed some indication of altered lipid fractions, comparing to control group

CONCLUSION

supported by Regmi study.

The outcome of this study specifies that SCH patients are characterized by increased BMI, dyslipidemia, and a higher concentration of CRP. This significant increase in CRP level might be the cause of development of cardiac symptoms in the coming future. Cardiovascular abnormalities can be developed in the later stage of disease. Although there are remarkable changes in different variables in SCH patients having TSH > 10 µIU/ml, Patients with SCH in which TSH range was <10 µIU/ml also has some alteration in different variables compared to normal healthy adults. Increased BMI, altered lipid fractions and abnormal CRP might be helpful in the early detection to diagnose cardiovascular risk in SCH patients. The sample size of this study was small, so the further studies with larger number of patients will be helpful to establish the fact.

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