

## Descriptive Assessment of the Clinic-Epidemiological Factors among Thyroid Dysfunction (TD) Patients Diagnosed with Metabolic Syndrome (Mets)

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Conflict of interest: Nil

### Abstract

**Aim:** The aim of the present study was to assess the prevalence and clinical and epidemiological factors of thyroid dysfunction (TD) patients diagnosed with metabolic syndrome (MetS).

**Methods:** The study was conducted in the Department of Medicine, Sadar Hospital, Motihari, Bihar, India for nine months. In this study, we had enrolled 400 patients with MetS. The study was conducted in accordance with the principles of the Declaration of Helsinki, International Council on Harmonization Good Clinical Practice (ICH GCP) guidelines, and Indian regulatory guidelines (Indian Council of Medical Research and Indian GCP guidelines).

**Results:** In this study, we had enrolled 400 patients with MetS. The baseline demographic characteristics of these patients are shown. Of all the enrolled patients, 120 (30%; 95% CI: 23.83-32.32) were diagnosed with TD (mean age (SD): 47.9 (10.96) years; mean BMI: 30 ± 4.94 kg/m<sup>2</sup>), with a higher prevalence among women compared to men (91 (75%) vs. 30 (25%)). Of the 400 MetS patients, overt hypothyroidism was reported in 78 (19.5%) patients and overt hyperthyroidism in 8 (2%) patients.

**Conclusion:** The prevalence of TD in patients with MetS was high, indicating a possible interplay between thyroid status and MetS. The data generated from the present study will aid in establishing a correlation between TD and MetS in Indian patients. This early diagnosis of TD in MetS would help in modifying the disease course by early interventions with appropriate lifestyle modification regimens, as applicable.

**Keywords:** Metabolic Syndrome, Hypothyroidism, Subclinical hypothyroidism, thyroid dysfunction

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### Introduction

Metabolic syndrome constitutes a cluster of risk factors characterized by hypertension, atherogenic dyslipidemia, hyperglycemia, prothrombotic and

proinflammatory conditions. [1] This cluster of metabolic abnormalities is associated with increased risk for atherosclerotic cardiovascular disease and

type 2 diabetes mellitus. [2] The prevalence of metabolic syndrome is increasing all over the world with distinct evidence of high prevalence in India and other South Asian countries. [3]

Metabolic syndrome comprises a group of interrelated metabolic abnormalities that are characterized by central obesity, high triglycerides (TGs), low high-density lipoprotein cholesterol (HDL-C), hypertension and hyperglycaemia. Patients with metabolic syndrome have an increased risk of cardiovascular disease, type 2 diabetes, and all-cause mortality. After adjusting for potential risk factors and each component of metabolic syndrome as a continuous variable, metabolic syndrome was associated with an increased 10-year risk of coronary heart disease. [4] It is estimated that the population attributable rates of metabolic syndrome are approximately 6%-7% for all-cause mortality, 12%-17% for cardiovascular disease, and 30%-52% for diabetes. [5]

Thyroid diseases are among the most prevalent endocrine disorders worldwide. Based on the estimation from various studies, it has been projected that about 42 million people in India suffer from thyroid diseases.[6] MetS is closely associated with thyroid dysfunction (TD) due to the impact of thyroid hormones on lipid metabolism, glucose, blood pressure, and cardiovascular dysfunction.[7] Functional changes in the thyroid gland might have an association with MetS and its related components including obesity, insulin resistance (IR), lipid and glucose metabolism abnormalities, raised blood pressure, and cardiovascular dysfunction. MetS and TD are both characterized by a cluster of common abnormalities such as abdominal obesity, hyperglycemia, hypertension, reduced high-density lipoprotein cholesterol (HDL-C), and elevated triglycerides (TG). Moreover, IR, identified as a basic mechanism for MetS, also plays a role in hypothyroidism. [8]

The occurrence of both the conditions may be compounded to increase the risk for cardiovascular diseases (CVDs).

Both metabolic syndrome and hypothyroidism are independent risk factors for cardiovascular diseases (CVD). Presence of both conditions may be compounded to increase the risk for CVD and a considerable overlap occurs in the pathogenic mechanisms of atherosclerotic cardiovascular disease by metabolic syndrome and hypothyroidism.[9]

The aim of the present study was to assess the prevalence and clinical and epidemiological factors of thyroid dysfunction (TD) in Indian patients diagnosed with metabolic syndrome (MetS).

## Methods

The study was conducted in the Department of Medicine, Sadar Hospital, Motihari, Bihar, India. In this study, we had enrolled 400 patients with MetS. The study was conducted in accordance with the principles of the Declaration of Helsinki, International Council on Harmonization Good Clinical Practice (ICH GCP) guidelines, and Indian regulatory guidelines (Indian Council of Medical Research and Indian GCP guidelines).

Patients aged 18 to 65 years, with an established diagnosis of MetS based on the National Cholesterol Education Program (NCEP) Adult Treatment Panel (ATP) III criteria (with modified waist), with or without known TD, were invited to participate in the study during their routine clinical visit to the endocrinologists, gastroenterologists, and/or hepatologists. Pregnant patients or patients with a history of jejunoileal bypass, biliopancreatic diversion, extensive small bowel resection, total parenteral nutrition, any forms of chronic liver disease, hepatocellular carcinoma, patients on weight loss therapies or steatogenic drugs, and known HIV-positive cases were excluded from

the study. At the screening visit, the following data were collected in the case report forms: demographics, anthropometric measurements, significant medical (including FPG, serum TG, and HDL-C values from patients' medical records) and surgical history, family history, lifestyle parameters, history of consumption of any obesogenic medicines, vital signs, and details of physical examination.

After obtaining an informed signed consent to participate in the study, the eligible patients from the screening visit were requested to visit the clinic after an overnight fast within 3-10 days of consenting. At this visit (visit 1), abdominal ultrasound examination (USG) was performed in patients who did not consume alcohol or consumed less than 20 g of alcohol per day and had not received corticosteroids, amiodarone, or tamoxifen. Blood samples were collected for assessment of hemogram, coagulogram (activated partial thromboplastin time, thrombin time, and prothrombin time), plasma insulin, plasma glucose, lipid profile (TG, total cholesterol (TC), HDL-C, and low-density lipoprotein cholesterol

(LDL-C)), and thyroid function (free triiodothyronine (FT3), free thyroxine (FT4), and thyroid-stimulating hormone (TSH)). The fasting plasma glucose and plasma insulin were used for the calculation of homeostatic model assessment-established IR (HOMA-IR). The patients were followed up for a mean of one year to check for new diagnoses of TD.

The primary endpoint was the prevalence of TD among patients with MetS. Other endpoints included the percentage of patients with hyperthyroidism, subclinical hyperthyroidism, hypothyroidism, and subclinical hypothyroidism and percentage of patients with TD with respect to individual components of MetS (waist circumference, TG, HDL-C, SBP, DBP, and fasting glucose) and IR (HOMA – IR > 1.64).

#### Statistical Analysis

All statistical analyses were performed using SAS® version 9.2 (SAS Institute Inc., USA). The prevalence of TD in MetS patients was calculated as a number and percentage with 95% CI.

#### Results

**Table 1: Demographics and baseline characteristics of 432 patients with metabolic syndrome**

Parameter	Age ≤ 45 y (N = 180)	Age > 45 y (N = 220)	Total (N = 400)
<b>Age in years</b>			
Mean (SD)	36.4 (6.08)	55.5 (5.52)	47.3 (10.96)
Range	21.0-45.0	46.0-65.0	21.0-65.0
<b>Gender</b>			
Women, N (%)	110 (61.12%)	120 (54.54%)	230 (57.5%)
Men, N (%)	70 (38.88%)	100 (45.46%)	179 (42.5%)
Height in cm, mean (SD)	163.0 (9.02)	160.8 (8.44)	161.7 (8.74)
Weight in kg, mean (SD)	79.4 (13.13)	75.8 (12.29)	77.3 (12.75)
Waist circumference in cm, mean (SD)	98.5 (9.41)	98.7 (9.92)	98.6 (9.70)
Hip circumference in cm, mean (SD)	106.3 (11.07)	104.7 (11.14)	105.4 (11.13)

In this study, we had enrolled 400 patients with MetS. The baseline demographic characteristics of these patients are shown. Of all the enrolled patients, 120 (30%; 95% CI: 23.83-32.32) were diagnosed with

TD (mean age (SD): 47.9 (10.96) years; mean BMI:  $30 \pm 4.94$  kg/m<sup>2</sup>), with a higher prevalence among women compared to men (91 (75%) vs. 30 (25%).

**Table 2: Percentage prevalence of different grades of thyroid dysfunction**

Classification of TD	MetS patients (N = 400) N (%)
Hypothyroidism	70 (17.5)
New overt hypothyroidism	8 (2)
New subclinical hypothyroidism	40 (10)
Hyperthyroidism	8 (2)
New overt hyperthyroidism	0
New subclinical hyperthyroidism	4 (1)
Total number of TD patients	120 (30)

Of the 400 MetS patients, overt hypothyroidism was reported in 78 (19.5%) patients and overt hyperthyroidism in 8 (2%) patients.

### Discussion

Metabolic syndrome is a constellation of abnormalities, including increase in weight (obese), having hypertension, abnormal lipid profile with elevated triglycerides and low values of high-density lipoproteins, increased values of fasting blood sugars. Patients of metabolic syndrome had a higher risk of developing diabetes and cardiovascular diseases in future. Thyroid dysfunction is common among patients of metabolic syndrome. In various studies conducted in India, Nepal, Middle East and African countries the prevalence of thyroid dysfunction in metabolic syndrome patients is in the range of 21- 51%. [10-13] Metabolic syndrome can be associated with endocrine and non-endocrine disorders and has widespread consequences. Alterations in thyroid functions, though well known, are not recognized clinically and there is inconsistency in thyroid functions in metabolic syndrome. [14]

Oxidative stress, chronic inflammation, and angiogenesis are believed to enhance the pathogenesis of MetS.[15] The important components of MetS, such as hyperglycemia and inflammation, upsurge the production of reactive oxygen species (ROS) resulting in increased oxidative stress with overactivation of nicotinamide adenine dinucleotide phosphate (NADPH) oxidase. [16,17] The main ROS is the

superoxide anion, produced by NADPH oxidase [19]. Hypermetabolic state in hyperthyroidism may accelerate free radical production in mitochondria and induce changes in the antioxidant defense system. In hypothyroidism, associated oxidative stress is the consequence of reduced capacity of the antioxidative defense.

The lesser number of subclinical hypothyroidism cases (10%) reported in our study could be attributed to the high enrolment of patients with known cases of hypothyroidism (17.5%), who were already on levothyroxine therapy. The overall rate of hypothyroidism at baseline (17.5%) plus new overt hypothyroidism (2%) and new subclinical hypothyroidism (10%) was calculated as 25.70%. In agreement with our findings, a similar prevalence of hypothyroidism was reported in MetS population in other Indian studies as well, viz., Kota et al. (26%). [18]

In our study, women with MetS had a higher incidence of TD in comparison to men. This corroborates reports from other studies where women outnumbered men in terms of prevalence of TD in MetS [27–29]. Both men and women in the higher age group (>45 years of age) had a higher incidence of TD. Our results were in agreement with other reports where there was a tendency of increasing TD with aging across both genders. [19,20] Thus, age and gender could represent significant risk factors for TD in MetS patients, prompting a detailed clinical and laboratory evaluation in these groups.

In this study, the MetS components observed in patients diagnosed with TD were high waist circumference, reduced HDL-C, raised HOMA-IR, systolic blood pressure, diastolic blood pressure, fasting glucose, and TG. A higher proportion of females had waist circumference above the cutoff (>80 cm) as compared to men. Though the other studies have also reported an association between TD and components of MetS, but it is still debatable. A Nigerian study reported MetS to be significantly associated with higher FT4. [21] Kota et al. found the significant association between subclinical hypothyroidism and MetS with the relationship between TSH levels and TC, TG, LDL, and HDL-C levels among Indian patients. [18]

It should be noted that while exploring the relationship between TD and components of MetS, most studies have focused on the subclinical hypothyroidism. Further, it should be noted that the pattern of TD in MetS and its relationship with components may vary upon geographic locale, age, gender, diet, and genetics, and environmental factors. [20,22,23,24]

### Conclusion

Thyroid dysfunction is an important entity as a complication in metabolic syndrome patients. From various studies, it is a known fact that the incidence of thyroid hormone abnormality is more in females as compared to males. The prevalence of TD in patients with MetS was high, indicating a possible interplay between thyroid status and MetS. The data generated from the present study will aid in establishing a correlation between TD and MetS in Indian patients. This early diagnosis of TD in MetS would help in modifying the disease course by early interventions with appropriate lifestyle modification regimens, as applicable. However, future large sample-sized prospective studies are warranted which could evaluate the impact of TD

management in terms of reduction in MetS and its related components.

### References

1. Shantha GP, Kumar AA, Jeyachandran V, Rajamanickam D, Rajkumar K, Salim S, Subramanian KK, Natesan S. Association between primary hypothyroidism and metabolic syndrome and the role of C reactive protein: a cross-sectional study from South India. *Thyroid research*. 2009 Dec;2(1):1-7.
2. Akinboro AO, Akinyemi SO, Olaitan PB, Raji AA, Popoola AA, Awoyemi OR, Ayodele OE. Quality of life of Nigerians living with human immunodeficiency virus. *The Pan African Medical Journal*. 2014;18.
3. Gyawali P, Takanche JS, Shrestha RK, Bhattarai P, Khanal K, Risal P, Koju R. Pattern of thyroid dysfunction in patients with metabolic syndrome and its relationship with components of metabolic syndrome. *Diabetes & metabolism journal*. 2015 Feb 1;39(1):66-73.
4. Misra A, Khurana L. Obesity and the metabolic syndrome in developing countries. *The Journal of Clinical Endocrinology & Metabolism*. 2008 Nov 1;93(11\_supplement\_1):s9-30.
5. Lu J, Wang L, Li M, Xu Y, Jiang Y, Wang W, Li J, Mi S, Zhang M, Li Y, Wang T. Metabolic syndrome among adults in China: the 2010 China noncommunicable disease surveillance. *The Journal of Clinical Endocrinology & Metabolism*. 2017 Feb 1;102(2):507-15.
6. Unnikrishnan AG, Menon UV. Thyroid disorders in India: An epidemiological perspective. *Indian journal of endocrinology and metabolism*. 2011 Jul;15(Suppl2):S78.
7. Liu YY, Brent GA. Thyroid hormone crosstalk with nuclear receptor signaling in metabolic regulation. *Trends in Endocrinology & Metabolism*. 2010 Mar 1;21(3):166-73.

8. Singh BM, Goswami B, Mallika V. Association between insulin resistance and hypothyroidism in females attending a tertiary care hospital. *Indian Journal of Clinical Biochemistry*. 2010 Apr;25(2):141-5.
9. Kota SK, Meher LK, Krishna SV, Modi KD. Hypothyroidism in metabolic syndrome. *Indian journal of endocrinology and metabolism*. 2012 Dec 1;16(Suppl 2):S332-3.
10. Shrestha S, Das BKL, Baral N, et al. Association of metabolic syndrome and its components with thyroid dysfunction in females. *Int J Diab Dev Countries* 2007;27(1):24-6.
11. Uzunlulu M, Yorulmaz E, Oguz A. Prevalence of subclinical hypothyroidism in patients with metabolic syndrome. *Endocr J* 2007; 54(1):71-6.
12. Deshmukh V, Farishta F, Bhole M. Thyroid dysfunction in patients with metabolic syndrome: a cross-sectional, epidemiological, Pan-India study. *International journal of endocrinology*. 2018 Dec 25;2018.
13. Gyawali P, Takanche JS, Shrestha RK, et al. Pattern of thyroid dysfunction in patients with metabolic syndrome and its relationship with components of metabolic syndrome. *Diabetes Metab J* 2015;39(1):66-73.
14. Chugh K, Goyal S, Shankar V, Chugh SN. Thyroid function tests in metabolic syndrome. *Indian journal of endocrinology and metabolism*. 2012 Nov;16(6):958.
15. Mahjoub S, Masrou-Roudsari J. Role of oxidative stress in pathogenesis of metabolic syndrome. *Caspian journal of internal medicine*. 2012;3(1):386.
16. Hopps E, Noto D, Caimi G, Aversa MR. A novel component of the metabolic syndrome: the oxidative stress. *Nutrition, Metabolism and Cardiovascular Diseases*. 2010 Jan 1; 20(1):72-7.
17. Vanessa Fiorentino T, Prioletta A, Zuo P, Folli F. Hyperglycemia-induced oxidative stress and its role in diabetes mellitus related cardiovascular diseases. *Current pharmaceutical design*. 2013 Oct 1;19(32):5695-703.
18. Meher LK, Raveendranathan SK, Kota SK, Sarangi J, Jali SN. Prevalence of hypothyroidism in patients with metabolic syndrome. *Thyroid Research and Practice*. 2013 May 1;10(2):60.
19. Meng Z, Liu M, Zhang Q, Liu L, Song K, Tan J, Jia Q, Zhang G, Wang R, He Y, Ren X. Gender and age impacts on the association between thyroid function and metabolic syndrome in Chinese. *Medicine*. 2015 Dec;94(50).
20. Tehrani FR, Tohidi M, Dovom MR, Azizi F. A population-based study on the association of thyroid status with components of the metabolic syndrome. *J Diabetes Metab*. 2011;2 (8):156-68.
21. Udenze I, Nnaji I, Oshodi T. Thyroid function in adult Nigerians with metabolic syndrome. *Pan African Medical Journal*. 2014;18(1).
22. Shantha GP, Kumar AA, Jeyachandran V, Rajamanickam D, Rajkumar K, Salim S, Subramanian KK, Natesan S. Association between primary hypothyroidism and metabolic syndrome and the role of C reactive protein: a cross-sectional study from South India. *Thyroid research*. 2009 Dec;2(1):1-7.
23. Tseng FY, Lin WY, Lin CC, Lee LT, Li TC, Sung PK, Huang KC. Subclinical hypothyroidism is associated with increased risk for all-cause and cardiovascular mortality in adults. *Journal of the American College of Cardiology*. 2012 Aug 21;60(8):730-7.
24. Margute T. G., Ferreira P. C., Almeida I. M. M., Denardin C., Silva T. Q. M. da, Margute T. G., Maione M. S., Rossato A. R., & Santos, I. F. dos. Use of tricyclic antidepressants in trigeminal neuralgia. *Journal of Medical Research and Health Sciences*. 2022; 5(5): 2008–2012.

