



## Study of Serum Thyroid Stimulating Hormone Level Among Individuals in An Iodine Deficient Region of Upper Assam- A Hospital Based Study

### KEYWORDS

Iodine, Micronutrient, thyroid stimulating hormone

#### \* Dr. Saurabh Borkotoki

Professor & Head, Department of Biochemistry, Jorhat Medical College, Jorhat-785001, Assam, India  
\* Corresponding Author

#### Dr. Uttara Borkotoki

Associate professor, Department of Microbiology, Jorhat Medical College, Jorhat-785001, Assam, India

#### Dr. Reetwik Kumar Dutta

PGT, Department of Biochemistry, Jorhat Medical College, Jorhat-785001, Assam, India

#### Mr. Rituraj Baruah

Assistant professor, Department of Statistics, Jaganath Barooah College, Jorhat-785001, Assam, India

### ABSTRACT BACKGROUND:

Iodine is an essential micronutrient required for synthesis of thyroid hormones. Jorhat, Golaghat and Sivasagar districts of upper Assam belong to the Sub Himalayan iodine deficient belt with large scale floods every year during the rainy seasons. Inhabitants of this region are at risk of developing altered thyroid status due to iodine deficiency in their food, if not supplemented adequately. Thyroid stimulating hormone (TSH) secreted by the anterior pituitary is regarded as one of the best indicators of thyroid hormone activity. With this background, a cross sectional observational study has been undertaken to evaluate the thyroid status of the population of Jorhat, Golaghat and Sivasagar districts by examining their TSH levels. Subjects selected for the study are apparently euthyroid individuals visiting Jorhat Medical College Hospital from March 2014 to September 2015.

### RESULTS AND OBSERVATION:

A total of 1314 cases of both sexes belonging to different age groups were studied. The mean TSH value obtained is  $2.27\mu\text{IU/L} \pm 1.69\mu\text{IU/L}$ . For females it is  $2.33\mu\text{IU/L} \pm 1.73\mu\text{IU/L}$  and for the males is  $2.19\mu\text{IU/L} \pm 1.64\mu\text{IU/L}$ . 6.925% of the cases studied have values  $>5\mu\text{IU/L}$ . The results are comparable with studies from non-iodine deficient areas.

### Introduction:

Thyroxine (T4) and triiodothyronine (T3) are the two active thyroid hormones. They play critical roles in cell differentiation during development and maintain metabolic homeostasis. Synthesis and secretion of thyroid hormones are under control of thyroid stimulating hormone (TSH) of the anterior pituitary. Under normal conditions, TSH level in blood vary according to thyroid hormone levels. High TSH level indicates lowered thyroid activity and vice versa. Thus blood TSH concentration is a useful physiological marker of thyroid hormone activity [1]. Increased TSH levels lead to hypertrophy of the thyroid gland and in extreme cases manifest as hypothyroid goitre.

Iodine is the essential micronutrient required for synthesis of thyroid hormones. Plants and water derive iodine from the top soil reserves, which is subsequently acquired by the animal kingdom. Flood prone and hilly regions of the world are subjected to leaching of the top soil during rainy season, rendering these regions deficient in iodine. Inhabitants of such regions suffer from iodine deficiency disorders (IDD) due to lack of adequate dietary availability of iodine and resulting in inadequate thyroid hormone synthesis.

Thyroid hormones are essential for intellectual and physical development of an individual. Children born to hypothyroid mothers often result in stunted growth, cretinism and have lower IQ scores [2]. People living in areas affected by severe iodine deficiency may have an intelligence quotient of upto about 13.5 points below that of those from comparable communities in areas where there is no iodine deficiency [3]. Iodine deficiency and the resultant hypothyroidism increases the risk of still births, abortions, increased perinatal

deaths, infant deaths and congenital anomalies [4].

Globally IDD is a major public health problem. Over 2 billion people worldwide is affected with IDD [5]. It is estimated that in India alone, more than 200 million people are at risk of iodine deficiency and 71 million suffer from goiter and other IDD [6].

Hypothyroidism due to iodine deficiency still remains the most common cause of preventable mental deficiency in the world today [7]. Iodination of edible salt is an effective measure to control IDD in iodine deficient regions. In 1983-84, Govt. of India adopted a policy to achieve universal iodization of edible salt by 1992 [8]. In 1998, edible salt iodization was made compulsory and sale of non iodized salt was banned. However, in the year 2000 it was revoked considering compulsion in such matters of individual choice desirable [9].

Jorhat Medical College Hospital is a tertiary care teaching hospital. It caters to the patients from Sivasagar, Jorhat and Golaghat districts of upper Assam. All these districts are heavily flood prone. The Brahmaputra and its tributaries lashes these districts with flood every year during the rainy monsoon months.

Thus with this background, the present study is undertaken to examine the thyroid status of the people of this flood affected region belonging to the sub Himalayan iodine deficient belt by estimating serum TSH levels in individuals visiting Jorhat Medical College Hospital without any apparent thyroid related problems or complaints, critically analyse the findings and compare with observations made in other similar studies.

**Materials and methods:**

**Study design:** cross sectional hospital based

**Cases:**

A total number of 1314 subjects visiting Jorhat Medical College Hospital ,both male and female,were randomly selected,without any apparent thyroid related problems belonging to different age,were estimated for serum TSH levels.

**Exclusion criteria:**

- 1)Pregnancy.
- 2)Patients with diabetes mellitus,hypertension,fever,renal failure,liver cirrhosis,malignant neoplasm and psychological abnormality
- 3)On medications for thyroid disorders

**Study Time:**

Between March 2014 and September 2015.

**Specimen collection for tests:**

Collected 2cc of venous blood in sterile empty vial from each of the study subjects maintaining all routine precautions.

Allowed the samples to clot and serum was separated.

Then serum was shifted to storage tubes and was tested within four hours of collection at room temperature.

Haemolysed samples were discarded.

**Estimation:**

It was carried out in Access Immuno Assay Systems (Beckman Coulter) at the clinical Biochemistry wing of Central Clinical Laboratory, Jorhat Medical College Hospital.

**Assays:**

The Access TSH assay is a two site immunoenzymatic ("Sandwich") assay.

**Calibration:**

Regular calibrations were done every 28 days

**Quality control:**

QC material simulate the characteristics of patient samples are commercially available and supplied by the manufacturers- Beckman Coulter, were used.

Quality control materials were run in every 24 hours time for authenticity of the reports.

These QC materials cover at least two levels of the analyte. The test results were accepted only when quality control results were found to be within acceptable ranges.

**Results :**

Results of the tests were determined automatically by the system's software. The amount of analyte in the sample was determined from the measured light production by means of calibration data.

**Statistical analysis:**

Independent sample t test in SPSS 16 version.

**RESULTS**

1.Overall Mean/SD/Range of TSH values of the study pop-

ulation groups (in  $\mu\text{IU/L}$ )

|       |       |
|-------|-------|
| Mean  | 2.27  |
| SD    | 1.69  |
| Min   | 0.01  |
| Max   | 10.65 |
| Range | 10.64 |

Mean/SD/Range for females (in  $\mu\text{IU/L}$ )

|       |       |
|-------|-------|
| Mean  | 2.33  |
| SD    | 1.73  |
| Max   | 10.65 |
| Min   | 0.03  |
| Range | 10.62 |

Mean/SD/Range for males (in  $\mu\text{IU/L}$ )

|       |      |
|-------|------|
| Mean  | 2.19 |
| SD    | 1.64 |
| Max   | 9.91 |
| Min   | 0.01 |
| Range | 9.9  |

Note: Difference between above two groups are statistically insignificant with P value =0.127 (which is > 0.05)

**2.Table showing number of cases and average TSH values amongst different age groups**

| Age group (years) | Number of cases<br>n=1314 | Mean TSH<br>( $\mu\text{IU/L}$ ) |
|-------------------|---------------------------|----------------------------------|
| <12               | 26                        | 2.65                             |
| 12 to <20         | 122                       | 2.23                             |
| 20 to <40         | 581                       | 2.36                             |
| 40 to <60         | 411                       | 2.22                             |
| $\geq 60$         | 173                       | 2.02                             |

**3. Table showing Average TSH level among different age groups (Male vs Female)**

| Age groups<br>(years) | Mean TSH level<br>( $\mu\text{IU/L}$ ) |        | P<br>Value |
|-----------------------|--|--------|------------|
|                       | MALE                                   | FEMALE |            |
| <12                   | 2.85                                   | 1.822  | 0.227      |
| 12 - <20              | 2.23                                   | 2.24   | 0.964      |
| 20 - <40              | 2.31                                   | 2.39   | 0.561      |
| 40 - <60              | 2.08                                   | 2.36   | 0.124      |
| $\geq 60$             | 1.97                                   | 2.08   | 0.698      |

Note: Statistically comparison between Males & Females are insignificant with all the values >0.05

#### 4. Table showing number of cases and their relative prevalence with TSH levels >5 $\mu$ IU/L and <0.3 $\mu$ IU/L in males and females.

| TSH level ( $\mu$ IU/L) | Male (n=578) | Female (N=736) |
|-------------------------|--------------|----------------|
| <0.3                    | 17 (2.94%)   | 24 (3.26%)     |
| >5                      | 40 (6.92%)   | 51 (6.93%)     |

#### DISCUSSION:

The present study was carried out on 1314 apparently euthyroid individuals (both male and female) of different age groups from a known iodine deficient region (Jorhat, Golaghat and Sivasagar districts of Upper Assam) showed mean TSH value to be  $2.27\mu\text{IU/L} \pm 1.69\mu\text{IU/L}$ . This finding is complete to similar other studies elsewhere [10,11,12]

In our study mean  $\pm$ SD TSH values in females ( $2.33\mu\text{IU/L} \pm 1.73\mu\text{IU/L}$ ) was little higher than the males ( $2.25\mu\text{IU/L} \pm 1.63\mu\text{IU/L}$ ). Similar finding is also reported by Aghini-Lombardi F, et al. [13]

When TSH levels in different age groups were examined in relation to sex, no statistically significant result was obtained on comparison across the different age groups. However, an increasing trend of mean TSH values was observed in females as age advances till it is 60 years. The values are also higher than their male counterparts without any statistical significance.

In our study we have considered  $5\mu\text{IU/L}$  as the upper limit of normal TSH level. Upper limit of TSH normal value is a matter of intense debate, still this value is comparable to various studies conducted worldwide [11,12,13,14,15,16,17,18,19,20]. In the present study, 91 (6.925%) cases had TSH levels  $>5\mu\text{IU/L}$ . On further analysis, 51 (6.93%) were females and 40 (6.92%) were male subjects. Thus, TSH values above  $5\mu\text{IU/L}$  does not show any sex preference in the present study.

TSH values above  $5\mu\text{IU/L}$  is an important indicator of subacute hypothyroidism. In fact, subacute hypothyroidism is considered to be the most prevalent thyroid disorder affecting 3-15% of the adult population and is a matter of serious morbidity [12,21]. This prevalence of higher TSH value in our study was conducted in a iodine deficient region. But the results of the study shows a very similar pattern of TSH values obtained in non-iodine deficient region. Even though our study subjects consisted apparently euthyroid subjects, the similarity of prevalence of higher TSH level cases with studies of iodine sufficient region gives a reflection of a good thyroid status of the population of this region. This is an important observation. Probably, awareness against iodine deficiency disorders is the cause of this encouraging observation which is definitely an outcome of sustained and result oriented efforts of the Govt. and NGOs against IDD.

However, it is too early to comment. There is definitely a scope for an elaborate, community based study on thyroid status among the population of this region.

#### REFERENCES:

- Fauci AS, Braunwald E, Kasper DL et al. (2008) Harrison's Principles of Internal medicine. 17<sup>th</sup> ed. USA: The McGraw-Hill Companies, Inc. page no.-2225

- Grewal E, Khadgawat R, Gupta N et al. (2013) Assessment of iodine nutrition in pregnant north Indian subjects in three trimesters Indian J Endocrinol Metab. 2013; 17 : 289-93
- WHO, ICCIDD, UNICEF. Assessment of Iodine Deficiency Disorders and monitoring their elimination : A guide to programme managers 2<sup>nd</sup> edition. (2001) Geneva : World Health Organisation.; 2001. total no of pages 124. Report no. WHO/NHD/01.1
- Singh MB, Fotedar R, Lakshminarayana J. (2009) Micronutrient deficiency status among women of desert areas of Western Rajasthan .India. Public Health Nutr. 12: 624-9
- WHO. Iodine status : Worldwide WHO Global Database on Iodine Deficiency Disorders (2004). Geneva : World Health Organisation ; 2004. total no of pages 48.
- Tiwari BK. Revised Policy Guidelines on National Iodine Deficiency Disorder revised edition October 2006 (2006). New Delhi : IDD and Nutrition cell Directorate General of Health Services Ministry of health and Family Welfare Government of India; October 2006. total no of pages 38.
- Kapil U. Goiter in India and its prevalence. (1998) Journal of Medical Science and family planning. 3: 46-50
- Bhat P N M, Arnold F, Gupta K et al. National Family Health Survey ( NFHS-3) 2005-06 India volume I. (2007) Deonar Mumbai : International Institute for population sciences (II PS) and macro international. September 2007. total no of pages 540.
- Government of India. Withdrawal of restriction on sale of common salt for direct human consumption. [internet] available at <http://pib.nic.in/archive/releases98/lyr2000/rmay2000/r11052000.html>
- Dika H , Kasolo J, Bimenya G. (2010) Thyroid hormones profile in Students of Makerere College of Health Sciences in Kampala Uganda. Tanzania Journal of Health Research. January 12(1):1-7
- Lee KY , Kim JE , On HJ et al . (2011) Serum TSH level in healthy Koreans and the Association of TSH with serum lipid concentration and metabolic syndrome. Korean J Intern Med. 26:432-439
- Deshmukh V , Behl A, Joshi H et al. (2013) Prevalence , clinical and biochemical profile of Subclinical hypothyroidism in normal population in Mumbai. Indian Journal of Endocrinology and Metabolism. May-Jun 2013 ; 17(3): 454-459.
- Aghini - Lombardi F, Antonangeli L, Martino E et al. (1999) The spectrum of Thyroid disorders in an Iodine - deficient community : The Pescopagano survey. The Journal of Clinical Endocrinology and metabolism . 84(2):561-566
- Tunbridge WMG, Evered DC, Hall R et al. (1977) The Spectrum of Thyroid disease in the community : the Whickham Survey . Clin Endocrinol. 7: 481-93
- Vanderpump MPJ, Tunbridge WMG, French JM, et al. (1995) The incidence of thyroid disorders in the community: a twenty year follow up of the Whickham Survey. Clin Endocrinol. 43:55-68
- Canaris GJ, Manowitz NR , Mayor G et al. (2000) The Colorado thyroid disease prevalence Study. Arch Intern Med. 160: 526-34
- Hollowell JG , Staehling NW, Dana Flanders W et al. (2002) Serum TSH, T<sub>4</sub> and thyroid antibodies in the United States Population ( 1994-1998) : National Health and Nutrition Examination survey ( NHANES III). J Clin Endocrinol Metab. 87: 489-99
- Spencer CA , Lo Prosti JS, Patel A, et al. (1990) Application of a new Chemiluminometric thyrotropin assay to Subnormal measurement . J Clin Endocrinol Metab. 70: 453-60
- Hamilton TE, Oavis S, Onstad L et al. (2008) Thyrotropin levels in a population with no clinical autoantibody or ultrasonographic evidence of Thyroid disease: implications for the diagnostic of subclinical hypothyroidism. Journal of Clinical Endocrinology and Metabolism. 96: 1224-1230
- Wartofsky L, Dickey RA. (2005) The evidence for a narrower thyrotropin reference range is compelling. Journal of clinical Endocrinology and Metabolism.. 90: 5483-5488
- Ayala AR , Danese MD , LandenSon PW. (2000) When to treat mild hypothyroidism. Endocrinol Metab Clin J Am. 29:399-415