An Unusual Presentation of Altered Thyroid in Male

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Abstract: Altered thyroid function is more common among female and hypothyroidism predominate with clinical features of Weight gain, asthenia due to declined oestrogen in female and declined testosterone in male resulting lowered BMR yielding subcutaneous accumulation of mucin and retention of water and electrolytes. But in the present study male presenting with high TSH, low thyroxin (T4) and normal testosterone presented with body weight less than ideal body weight while person with low TSH, raised thyroxin (T4) and low testosterone presented with weight more than ideal body weight. None was having any other pathology. This study suggest that weight gain or weight loss is secondary to gonadal hormone oestrogen or testosterone which affect BMR and subcutaneous accumulation of mucin or loss of subcutaneous fat. Hence weight gain or weight loss is an effect of altered BMR not the thyroid hormone directly.

Keywords: Hypothyroidism, Hyperthyroidism, TSH, Testosterone, BMR, Subcutaneous deposit, Ideal body weight

1. INTRODUCTION

Endocrinal disorder’s incidence is increasing in geometric progression and alteration in thyroid function remain prime (1,2). Altered thyroid function presents as hypothyroidism and hyperthyroidism, out of which hypothyroidism is very common. Hypothyroidism affects more female than male (3,4) and commonest presenting features as

Weight gain, asthenia, constipation, lowers sexual function, alters spermiogenesis and may yield erectile dysfunction due to altered testosterone in male while hyperthyroidism manifest as weight loss, eye ball prominence, hair fall etc (5,6)

Presentation of hypothyroidism and hyperthyroidism can be summarised as -

<table>
<thead>
<tr>
<th>Hypothyroidism</th>
<th>Hyperthyroidism</th>
</tr>
</thead>
<tbody>
<tr>
<td>•  Weight gain</td>
<td>weight loss</td>
</tr>
<tr>
<td>•  Fatigue</td>
<td>increased appetite</td>
</tr>
<tr>
<td>•  Sensitivity to cold temperatures</td>
<td>nervousness</td>
</tr>
<tr>
<td>•  Depression</td>
<td>restlessness</td>
</tr>
<tr>
<td>•  Dry skin</td>
<td>inability to concentrate</td>
</tr>
<tr>
<td>•  Thinning hair</td>
<td>weakness</td>
</tr>
<tr>
<td>•  Heavy menstrual periods (in female)</td>
<td>irregular heartbeat</td>
</tr>
<tr>
<td>•  Trouble sleeping</td>
<td>difficulty in sleeping</td>
</tr>
<tr>
<td>•  Difficulty concentrating</td>
<td>fine, brittle hair</td>
</tr>
<tr>
<td>•  Pain or swelling of the joints</td>
<td>itching</td>
</tr>
<tr>
<td>•  Constipation</td>
<td>hair loss</td>
</tr>
<tr>
<td>•  High cholesterol levels</td>
<td>nausea and vomiting</td>
</tr>
<tr>
<td>•  Muscle weakness</td>
<td>breast development in men</td>
</tr>
</tbody>
</table>
Presentation of weight loss in hypothyroidism male with raised TSH, low thyroxin (T<sub>4</sub>) and weight gain in raised Thyroxin (T<sub>4</sub>) and very low TSH, created an eagerness to analyse the situation to adjudge the etiological pathogenesis of presenting features in persons with altered thyroid function.

These days increasing incidence of hypothyroidism even in male and presentation adverse to usual manifestations, needed due documentation and analysis for our future reference.

2. MATERIAL & METHODS

Objective of study: To establish reason of variability of myxoedema in hypothyroid and hyperthyroid state. In male with variable testosterone level.

2.1. Material

Cases presenting with varied presentation and shoeing high TSH been selected and analysed at RA. Hospital & Research Centre Warisaliganj and Centre for Endocrine and Metabolism, Aarogyam Punarjeevan, Ram Bhawan, Ara Garden Road, Jagdeo path, Baily Road Patna 14.

Case 1: Male Hindu aged 25 yrs, non-vegetarian, average built resident of, Manglam Bihar Apartment, Ara Graden, Patna 14 presented with –

Marked debility, Exertional Heaviness in the chest, Sleeplessness, leg cramps, uneasiness

Pain in chest, loss of appetite, general debility

Afebrile, Systemic examination shows no abnormality, No muscle stiffness, No myxoedema

Blood pressure 100/60 mm Hg, Blood Sugar: Random 123 mg %; HbA<sub>1c</sub>: 5.7

Serum Cholesterol Normal Thyroid Function: TSH 18.11 IU/L, T<sub>4</sub>: 7.6ug/dl

Vitamin D- 21.68ng/dl Vitamin B<sub>12</sub> low, Serum Calcium: 8.83mg/dl

Serum Testosterone: Normal 481ug/dl

ECG: Low voltage

Urine: RBC ++, Albumin ++

Case 2:

A 45 yrs male pure vegetarian resident of Old Bank Road, Warisaliganj (Nawada), average built complains of progressive decline weight, loss of appetite, general debility, uneasiness, constipation, occasional heaviness in the chest on strenuous exertion

BP: 100/50 mm Hg, Blood Sugar: (F) 78mg%; (PP): 123 mg % Serum Cholesterol - N,

TSH 28.12IU/L, T<sub>4</sub>: 7.2ug/dl

HbA1C: 5.2 Serum Testosterone 540

ECG Low voltage tracing

Case No: 3

A Mohamden male aged 43 years resident of Takya par Warisaliganj (Nawada), Weight 89 Kg, non-vegetarian, pain in chest, pain in abdomen, vertigo, exertional dyspnoea

Serum TSH 0.1 IU/dl; T<sub>4</sub>: 14ug/dl Serum testosterone 168

Blood pressure 140/80 mm Hg; Blood sugar(F) : 80mg%, HbA1C : 5.0

Serum Calcium: Vitamin D3: 18.4ng/dl

ECG: Normal sinus rhythm
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Case 04:

A 52 yrs old Hindu male presenting with weight gain, exertional dyspnoea, occasional pain in chest non vegetarian presented with -

Body weight: 86 Kg, Blood pressure: 150/100 mm Hg; HbA1C: 4.9

Thyroid profile: TSH: 3.17uIU/dl; T4: 12.2 ug/dl; Serum Calcium 10.12mg%

Serum testosterone: 294.95 ng/dl; Vitamin D: 17.52 ng/dl

ECG: Normal tracing

2.2. Method

All the three persons presenting with altered manifestation been clinically examined and thoroughly investigated for other possible adjuvant presentation responsible for the altered presentations.

Patients were adjudged for altered thyroid state as per -

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Thyroxin (T4)</th>
<th>Thyroid function</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>Normal</td>
<td>Normal thyroid function</td>
</tr>
<tr>
<td>Raised</td>
<td>Normal</td>
<td>Subclinical hypothyroidism</td>
</tr>
<tr>
<td>Raised</td>
<td>Low</td>
<td>Overt hypothyroidism</td>
</tr>
<tr>
<td>Low/ Normal</td>
<td>Low</td>
<td>Central hypothyroidism</td>
</tr>
<tr>
<td>Very Low</td>
<td>High</td>
<td>Hyperthyroidism</td>
</tr>
</tbody>
</table>

Based on the value of thyroxin (T4) and Thyroid stimulating Hormone (TSH) level hypothyroidism been categorized as –

Types of hypothyroid state | Characteristics
---|---
Central : If the TSH level is normal or low and serum free T4 levels are low, this is suggestive of central hypothyroidism (not enough TSH or TRH secretion by the pituitary gland or hypothalamus). There may be other features of hypopituitarism, such as menstrual cycle abnormalities and adrenal insufficiency. There might also be symptoms of a pituitary mass such as headaches and vision changes. Central hypothyroidism should be investigated further to determine the underlying cause
Overt : TSH levels are high and T4 and T3 levels are low. Overt hypothyroidism may also be diagnosed in those who have a TSH on multiple occasions of greater than 5µU/L, appropriate symptoms, and only a borderline low T4. It may also be diagnosed in those with a TSH of greater than 10µU/L
Sub clinical elevated serum TSH level, but with a normal serum free thyroxine level. In adults it is diagnosed when TSH levels are greater than 5 µU/L and less than 10µIU/L. The presentation of subclinical hypothyroidism is variable and classic signs and symptoms of hypothyroidism may not be observed or people with subclinical hypothyroidism, a proportion will develop overt hypothyroidism each year. In those with detectable antibodies against thyroid peroxidase (TPO), this occurs in 4.3%, while in those with no detectable antibodies, this occurs in 2.6%. Those with subclinical hypothyroidism and detectable anti-TPO antibodies who do not require treatment should have repeat thyroid function tests more frequently (e.g. yearly) compared with those who do not have antibodies
3. OBSERVATION

Both cases (Case 1 & 2) shows all the findings suggestive of hypothyroidism but not weight gain or myxoedema rather weight loss while 3rd and 4th cases having hyperthyroidic state present with myxoedema and weight gain.

In both Case 1 and 2, serum testosterone remained in normal range with low voltage ECG while in 3rd and 4th cases serum testosterone was lower than normal, with normal ECG and blood sugar status was normal in all the 4 cases.

<table>
<thead>
<tr>
<th>Particulars</th>
<th>Case I</th>
<th>Case II</th>
<th>Case III</th>
<th>Case IV</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (in years)</td>
<td>29</td>
<td>45</td>
<td>43</td>
<td>52</td>
</tr>
<tr>
<td>Body weight (in Kg)</td>
<td>42</td>
<td>53</td>
<td>89</td>
<td>86</td>
</tr>
<tr>
<td>Blood pressure(mmHg)</td>
<td>100/60</td>
<td>100/50</td>
<td>140/80</td>
<td>150/100</td>
</tr>
<tr>
<td>TSH (µIU/dL)</td>
<td>18.11</td>
<td>28.12</td>
<td>0.1</td>
<td>3.17</td>
</tr>
<tr>
<td>Thyroxin (T4)µg/dl</td>
<td>7.6</td>
<td>7.2</td>
<td>14</td>
<td>12.2</td>
</tr>
<tr>
<td>Blood Sugar (mg%)</td>
<td>123 (R)</td>
<td>123(PP)</td>
<td>80(F)</td>
<td>132(PP)</td>
</tr>
<tr>
<td>HbA1C</td>
<td>5.6</td>
<td>5.2</td>
<td>5.0</td>
<td>4.9</td>
</tr>
<tr>
<td>Serum cholesterol</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
<td>Normal</td>
</tr>
<tr>
<td>Serum Calcium</td>
<td>Lowered</td>
<td>Lowered</td>
<td>Normal</td>
<td>10.12mg%</td>
</tr>
<tr>
<td>Serum Testosterone</td>
<td>298</td>
<td>540</td>
<td>168</td>
<td>198.78</td>
</tr>
<tr>
<td>Vitamin D3(ng/dl)</td>
<td>21.68</td>
<td>13.67</td>
<td>12.65</td>
<td>17.54</td>
</tr>
<tr>
<td>ECG</td>
<td>Low voltage</td>
<td>Low voltage</td>
<td>Normal</td>
<td>Normal</td>
</tr>
</tbody>
</table>

4. RESULT

In male hypothyroidism presenting without myxoedema and hyperthyroidism with myxoedema

5. DISCUSSION

Thyroid disease affects women more than men i.e.- 5-8 women per men .Male hormones are considered protective against hypothyroidism in male low testosterone level predispose for hypothyroidism.  

Fatigue, cold intolerance, weight gain are typical presentation of hypothyroidism but may present with other associated features -low libido, depression, high cholesterol, loss of executive function, reduced frequency of morning erection, difficulty in growing facial hair, erectile dysfunction and also loss of muscle mass. In addition thyroid hormone can normalize testosterone.

Normally in hypothyroidism myxoedema is a result of accumulation of mucous polysaccharide, hyaluric acid and chondroitin sulphuric acid in subcutaneous tissue which exerts osmotic pressure and water retention, solely due to reduced catabolic process. 

TSH secretion is under the control of Hypothalamus by its hormone Thyrotropin releasing Hormone (TRH) which acts via thyroid receptor activating Adenyl cyclase to increase Cyc AMP.

Decrease in thyroxin level and increased TSH level affects testosterone which affect basal metabolic rate and alters glucose metabolism leading to accumulation of mucin and retention of Sodium and water in the skin resulting in myxoedema, thus myxoedema in thyroid depends on testosterone value, as in 3rd and 4th cases presenting with myxoedema and overweight having low TSH and increased thyroxin level with low testosterone while in other 2 cases high testosterone with low thyroxin and high TSH level fails to lower BMR thus fails to promote subcutaneous deposit of mucin, and retention of water and electrolyte.
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6. CONCLUSION

Altered thyroid function alters testosterone secretion in male, which alters BMR thus in case of hypothyroidism raised TSH and low thyroxin (T4), if testosterone remain normal BMR remain unchanged, thus fails to cause subcutaneous deposit of mucin and retention of water and electrolyte, but in case of lowered TSH (<0.1) and high thyroxin (T4) if testosterone level remain lower than normal, then BMR gets lowered and cause subcutaneous deposit of mucin and retention of water and electrolyte, causing myxoedema. Thus conclusively body weight gain or loss in thyroid function alteration is due to altered BMR which depends on gonadal hormone level than on TSH.

REFERENCES


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