Thyroid hormones profile in students of Makerere College of Health Sciences in Kampala Uganda

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Abstract: Serum concentrations of thyroxine (T4), triiodothyronine (T3) and Thyroid Stimulating Hormone (TSH) are used to assess thyroid function. It is recommended that each laboratory or hospital should establish its own reference values of T4, T3 and TSH for their clients because these hormones vary with ethnicity, geographical and climatic conditions of a population. There is no documented study which has been done to determine the Thyroid hormones profile in Ugandan general population. This study is one of the first attempts to determine Thyroid hormones profile in healthy Ugandans. The main objective of this study was to determine the thyroid hormones profile of students of the Makerere College of Health Sciences in Kampala, Uganda. A cross sectional descriptive study was done involving 72 students, with the mean age of 24.17 ± 4.48 years. Subjects who volunteered to participate in the study were interviewed; their height and body weight measured, 5ml of blood withdrawn, and sera harvested. FT4 and T3 Radioimmuno Assay (RIA) were done and TSH was assayed using Immunoradiometric Assay (IRMA) technique. The mean serum concentration of FT4 was 17.016 ± 3.847 qmol/L. For T3, mean serum concentration was 1.43 ± 0.825 nmol/L, and mean serum TSH level was 2.412 ±2.284 μIU/ml. Variations of serum concentrations of FT4, T3 and TSH with sex, age, or region of origin were not statistically significant. Serum concentration of TSH increased with increased body mass index (BMI). It was 2.073 ± 1.907 μIU/ml for subjects with BMI of ≤ 24.9 Kg/m², 3.588 ± 1.495 μIU/ml for subjects with BMI of 25 – 29.9 kg/m² and 4.450 ± 0.593μIU/ml for subjects with BMI ≥30kg/m² (P=0.009). However, BMI had no effect on serum concentrations of FT4 and T3. Serum concentrations of T4, T3 and TSH obtained from this study all differ with the values which are currently used as reference ranges in the country. We recommend a similar study involving a population representative of Ugandans to be conducted so as to establish normal reference values of T4, T3 and TSH for Ugandans. We also recommend BMI of patients to be taken into consideration during interpretation of serum TSH concentrations results.

Key words: Thyroxine, triiodothyronine, Thyroid Stimulating Hormone, university students, Uganda

Introduction

The two principal biologically active thyroid hormones, thyroxine (T4) and triiodothyronine (T3) together with Thyroid Stimulating Hormone (TSH) are used to assess thyroid function

Thyroid function tests are used in a variety of clinical settings to assess integrate of the thyroid gland and monitor the treatment of hyper and hypothyroidism.

It has been recognized for some time that levels of thyroid function parameters in healthy subjects show considerable inter-individual variability leading to wide (population-based) laboratory reference ranges, whereas intra-individual variability has a much narrower range (Andersen et al., 2002 ). Since thyroid function vary with ethnicity, geographical and climatic conditions, it is recommended that each laboratory or hospital establishes its own reference values of T4, T3 and TSH serum concentrations for their clients

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(Institute of Isotopes Company Ltd, Budapest. http://www.izotop.hu). Otherwise, choosing unrelated “normal ranges” may lead to a false diagnosis. This is important because clinical management of thyroid conditions depends very much on estimations of serum TSH, T4 and T3 in relation to the reference range. Many times given abnormal values for a given individual may not necessarily be abnormal but misinterpreted according to the designated reference range.

Furthermore, there is increasing evidence that at the population level, small differences in thyroid function are associated with differences in clinically important parameters such as body mass index (BMI) (Knudsen et al. 2005). As different populations can have specific basal metabolic rate due to various climatic conditions and various genetic make-ups, variation in the mean plasma concentrations of T4, T3 and TSH is obvious (Reed 2000). Although the distribution of the mean serum concentrations of T4, T3 and TSH in the various populations shows normal distribution curves, this continuous variation fluctuate because of variations in the living conditions and genetic profile (Meikle et al. 1988).

Unfortunately, there is no known study done to determine the Thyroid hormones profile in healthy Ugandans. The currently used normal references for normal ranges of serum concentrations of T4, T3 and TSH in Uganda, are those set in the Caucasian population, which may not be normal for the Ugandan population. This study therefore was carried out to document T4, T3 and TSH levels for students in Makerere College of Health Sciences in Kampala, Uganda. The findings can serve as reference values for adult Ugandans.

Materials and Methods

Study subjects

A cross sectional descriptive study was done to determine the Thyroid hormones profile of students in Makerere College of Health Sciences of Makerere University, Kampala, Uganda. The study involved male and female students who voluntarily agreed to participate in the study. Non-Ugandan students and those with history of or symptoms and/signs of thyroid diseases were excluded from the study. Students using steroids orally were also excluded. A consecutive sampling procedure was used to include subjects that met the selection criteria. To ensure that the subjects selected were representative of students of Makerere College of Health Sciences, about equal proportions (6.3 – 7.0%) of subjects from each class were recruited for study.

Data collection

The participants were given self-administered questionnaires to provide information on demographic characteristics. Height in the nearest centimetre was measured using a height ruler; and weight in the nearest gram was measured using analog person weighing scale. Body Mass Index (BMI) was computed as weight in kilograms divided by the height in meters squared.

Blood samples were obtained, centrifuged and sera harvested. The sera were stored at -20°C and analyzed within three days. Serum free T4 (FT4) and T3 were determined by radioimmunoassay (RIA), while serum TSH was determined by immunoradiometric assay (IRMA). The RIA and IRMA kits were bought from Institute of Isotopes Company Ltd of Budapest, Hungary.
**Ethical considerations**

The study was approved by Makerere College of Health Sciences Research and Ethics Committee on behalf of Ugandan National Council for Sciences and Technology. All participants gave written informed consent.

**Data analysis**

Data was analysed with SPSS version 11.0. The relationships between serum concentrations of FT4, T3 and TSH; and age, region of origin and BMI were determined using one way ANOVA. Association of hormone levels with sex was determined by Student’s t-test. Statistical significance level was fixed at p < 0.05.

**Results**

A total of 72 Ugandan students aged 19 – 37 years were recruited in the study. Their mean age was 24.17 ± 4.48 years. Forty two (58.3%) of the subjects were males and 30 (41.7%) were females. Majority (48.6%) of the subjects were from central region; 10 (13.9%) were from northern, 12 (16.7%) from eastern and 15 (20.8%) were from western region. Only 1 student (1.4%) was underweight (BMI <18.5kg/m²). Majority, 58 (80.6%) who had normal BMI of 18.5 ≤24.9kg/m² and 10 (13.9%) were overweight (BMI 25 – 29.9kg/m²). Only 3 (4.2%) were obese (BMI ≥30 Kg/m²).

**Table 1: Mean serum concentrations of FT4, T3 and TSH by Age**

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>FT4 (pmol/L)</th>
<th>T3 (nmol/L)</th>
<th>TSH (µIU/ml)</th>
</tr>
</thead>
<tbody>
<tr>
<td>≤20</td>
<td>17.87±4.063</td>
<td>1.31±0.879</td>
<td>2.71±2.404</td>
</tr>
<tr>
<td>21 – 25</td>
<td>17.83±4.243</td>
<td>1.45±0.812</td>
<td>2.45±2.259</td>
</tr>
<tr>
<td>26 – 30</td>
<td>15.24±2.916</td>
<td>1.62±0.788</td>
<td>2.24±2.056</td>
</tr>
<tr>
<td>31+</td>
<td>15.59±1.882</td>
<td>1.99±0.536</td>
<td>1.84±1.328</td>
</tr>
<tr>
<td>P- value</td>
<td>0.077</td>
<td>0.091</td>
<td>0.817</td>
</tr>
</tbody>
</table>

The mean serum concentration of FT4 was 17.016 ± 3.847 pmol/L. For T3, mean serum concentration was 1.43 ± 0.825 nmol/L, and mean serum TSH level was 2.412 ±2.284 µIU/ml. There was no statistically significant difference in mean serum concentrations of FT4, T3 and TSH between males and females. The variations of levels of these hormones with age were not statistically significant (Table 1). There was no relationship between serum concentrations of these hormones with region of origin (Table 2).

**Table 2: Mean serum concentrations of FT4, T3 and TSH by Region of Origin**

<table>
<thead>
<tr>
<th>Mean concentration of serum</th>
<th>Central</th>
<th>Northern</th>
<th>Eastern</th>
<th>Western</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>FT4 (pmol/L)</td>
<td>16.97±4.14</td>
<td>16.52±3.76</td>
<td>15.77±2.18</td>
<td>18.35±4.01</td>
<td>0.389</td>
</tr>
<tr>
<td>T3 (nmol/L)</td>
<td>1.32±0.77</td>
<td>1.26±0.86</td>
<td>2.01±0.94</td>
<td>1.43±0.77</td>
<td>0.061</td>
</tr>
<tr>
<td>TSH (µIU/ml)</td>
<td>1.86±1.17</td>
<td>3.57±3.13</td>
<td>3.39±3.09</td>
<td>2.13±1.73</td>
<td>0.107</td>
</tr>
</tbody>
</table>

Serum concentration of TSH increased with increased BMI (Table 3). It was 2.073 ± 1.907 µIU/ml for subjects with BMI of ≤ 24.9 Kg/m², 3.588 ± 1.495 µIU/ml for subjects with BMI of 25–29.9kg/m² and 4.450 ± 0.593µIU/ml for subjects with BMI ≥30kg/m² (P=0.009). FT4
decreased with increased BMI. It was $17.435 \pm 3.954$ μmol/L for subjects with $<25$ Kg/m², $16.024 \pm 2.931$ μmol/L for subjects with $25–29.9$ Kg/m² and $12.365 \pm 1.020$ μmol/L for subjects with $\geq 30$ Kg/m². However this relationship was not statistically significant (Table 3). Serum T3 levels, increased with increased BMI, though the relationship was not statistically significant (Table 3). It was $1.360 \pm 0.812$ nmol/L for subjects with $<25$kg/m², $1.531 \pm 0.930$ nmol/L for subjects with $25 – 29.9$ Kg/m² and $2.335 \pm 0.589$ nmol/L for subjects with $\geq 30$ kg/m² ($P=0.253$).

<table>
<thead>
<tr>
<th>Table3: Mean serum concentrations of FT4, T3 and TSH by BMI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean serum concentration of</td>
</tr>
<tr>
<td>----------------------------</td>
</tr>
<tr>
<td>FT4 (μmol/L)</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>T3 (nmol/L)</td>
</tr>
<tr>
<td>TSH (μIU/ml)</td>
</tr>
</tbody>
</table>

* Statistically significant

Discussion

Although the mean serum concentration of FT4 was comparable to $17.5 \pm 4.8$ pmol/L found by Samollow et al. (2004) and higher than $14.83 \pm 2.2$ μmol/L found by Institute of Isotopes Company Ltd among healthy blood donors in Hungary, the T3 level was slightly lower than those found by the same studies. The mean TSH level was comparable to that reported by Hollowell et al. (2002) for the Mexican-American” disease-free” population and that obtained by Samollow et al. (2004). Other studies found TSH to vary from 0.4–4.2μIU/ml (Nelson et al., 1993; Fisher, 1996).

Differences between the results from this study and the reference values which are currently being used in Uganda laboratories are accounted for by race, age of subjects and climatic condition. In this study, all the study subjects were black Africans, while the healthy blood donors used by Institute of Isotopes were Caucasians. Subjects in this study aged 19 – 37 years, with a mean age of 24.17 ± 4.48 year, while the subjects used by Institute of Isotopes Company Ltd aged from 19–69 years, with a mean age of 33.4 ±11.7 years. This study was conducted in Uganda which is in the tropics, and the climate differs from the temperate climate of Hungary where the reference values for serum concentration of FT4, T3 and TSH used in Uganda were established. Differences between this study results and those reported by Fisher (1996) and Nelson et al. (2004), can be explained by the differences in race, climate and probably life style and diets.

This study like many others before shows no difference in serum concentration of FT4, T3 and TSH between males and females (Nelson et al., 1993; Fisher, 1996; Samollow et al., 2004; Fathzadeh et al., 2005). Although other studies had found that serum concentrations of FF4, T3 and TSH decrease with age (Yoshida et al., 1989; Nelson et al., 1993; Fisher, 1996; Myers et al., 2006), this study found no statistically significant change of these hormones with ages. Lack of significant variation in serum concentrations of these hormones with age in this study could be due to the narrow age range for the subjects (19–37 years). Even other researchers reporting on adults (20 – 50 years) found no significant change with age in serum T4, TSH, but modest decrease in T3 (Nelson et al., 1993; Delange & Fisher, 1995; Fisher 1996).
There was no relationship between students’ origin and mean serum concentrations of FT4, T3 and TSH. This is due to the fact that all regions of Uganda are within the same (tropical) climatic condition. Another reason is the government policy whereby all table salts in the Ugandan markets are iodinated to supports easy formation of T3 and T4 in the population. This makes it possible to have iodinated salt in more than two-third of all households in Uganda.

Overweight (BMI 25-29.9 kg/m²) and obese students (>30kg/m²), had average TSH above that of the students of normal BMI. This shows an increased TSH concentration with increased BMI. This finding is similar to those reported by other scholars (Knudsen et al. 2005; Bhowmick et al. 2007; Ortega et al. 2007). The lack of significant association between BMI and T4 and T3 levels as been reported by Manji et al. (2006). Yet in children T4 has been found to be significantly higher in obese children compared to those of normal weight (Bhowmick et al. 2007). A negative association between BMI, and T3 and T4 has been reported by Myers et al.(2006) while others showed that free T3 and free T4 concentrations being lower in obese women (Sari et al. 2003), which is in contrary to our findings.

The mechanism of increased TSH in overweight than normal students could be due to a high level of leptin in overweight students than other group because TSH concentration is associated with percentage body fat and leptin concentration. Leptin which is produced mainly in the adipose tissue, in low doses cause increase in serum TSH concentration (Ortiga-Carvalho et al. 2002). It stimulates biosynthesis of Thyrotropin Releasing Hormone (TRH) and influences the activity of the hypothalamic–pituitary–thyroid axis (Nillni et al. 2000; Ortiga-Carvalho et al. 2002). Although plasma leptin levels were not determined in these subjects, the results indirectly suggest the fact that a possible association between leptin and the hypothalamic–pituitary–thyroid axis do exist.

In conclusion, the hormonal levels for Ugandans obtained from this study differ from the reference values which are currently used in the country. We therefore recommend a similar study involving bigger sample which is more representative of Ugandans to be conducted so as to establish normal reference values of T4, T3 and TSH for Ugandans. Since TSH was observed to increase with BMI, it is important that body mass index is taken into consideration during interpretation of serum TSH concentrations results.

Acknowledgements

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Declaration of interest

There is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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References


