IGF1 deficiency in newly diagnosed Graves’ disease patients

Sorina Martin,1,2 Anca Sirbu,1,2,3 Minodora Betivoiu,2 Suzana Florea,4 Carmen Barbu,1,2 Simona Fica1,2

1Endocrinology Department, Carol Davila University of Medicine and Pharmacy; 2Endocrinology Department, Elias Hospital; 3Victor Babes Institute; 4Laboratory Department, Elias Hospital; Bucharest, Romania

ABSTRACT

OBJECTIVE: Thyroid hormones influence the GH/IGF1 axis, but previous studies have reported discrepant results regarding serum IGF1 levels in hyperthyroidism. We have therefore investigated, at diagnosis, the relationship between serum IGF1 levels and the main characteristics of Graves’ disease (GD): severity of hyperthyroidism, goiter size, presence of active Graves’ ophthalmopathy (GO), antithyroid antibodies status and titer. DESIGN AND METHODS: This cross-sectional study included 98 newly diagnosed hyperthyroid patients with GD who presented consecutively at our clinic. The main measured parameters were: TSH, FT4, FT3, TT3, thyroglobulin, anti-thyroid peroxidase antibodies (TPOAb), anti-thyroglobulin antibodies (ATA), thyrotropin receptor antibodies (TRAb), IGF1. Patients were considered IGF deficient if IGF1 z score was ≤-2SD from mean for age. RESULTS: In GD patients, men had higher IGF1 levels (p=0.023) and IGF1 z scores (p=0.013) than women. 18.4% of GD patients were, at diagnosis, IGF1 deficient. Compared to patients without IGF1 deficiency, these patients presented higher thyroglobulin (median=72.55, IQR=116.02 vs median=11.40, IQR=80.74 ng/ml, p=0.002) and FT3 (median=11.30, IQR=7.64 vs median=7.33, IQR=5.72 pg/ml, p=0.027), and lower ATA (median=20, IQR=0 vs median=34.05, IQR=161 iu/ml, p<0.001) levels. Thyroglobulin was independently associated with IGF1 deficiency (AUROC=0.732, 95% CI: 0.620-0.844, p=0.002; cut-off for thyroglobulin=50.40 ng/ml, Se=77.8%, Sp=70%). IGF1 status was not influenced by gender (p=0.084), current smoking (p=0.558), goiter size (p=0.533), active ophthalmopathy (p=0.334), TRAb (p=0.239) or TPOAb status (p=0.367). CONCLUSIONS: Nearly one fifth of newly diagnosed GD patients had IGF1 deficiency. IGF1 deficiency was associated with lower ATA titers, higher thyroglobulin levels and more severe FT3 hyperthyroidism at diagnosis.

Key words: Anti-thyroglobulin antibodies, IGF1, IGF1 deficiency, Graves’ disease, Thyroglobulin

INTRODUCTION

Graves’ disease (GD) is an autoimmune disorder characterized by diffuse goiter and thyrotoxicosis which can be accompanied by ophthalmopathy and, rarely, dermopathy.

While the role of thyrotropin receptor antibodies
(TRAb) and thyrotropin receptor (TSHR) in GD hyperthyroidism is well established and a correlation appears to exist between TRAb levels and the clinical activity of Graves’ ophthalmopathy (GO), whether TRAb or TSHR plays a role in extrathyroidal manifestations of GD has not been resolved. The role of the IGF1/IGF1R pathway in the pathogenesis of GD was recently studied, although the relationship between the thyroid and IGF1 was first described in the 80s. IGF1 and IGF1R may be actively involved early in the pathogenesis of GD. It appears that both TSHR and IGF1R contribute to GD pathogenesis and together may comprise a physical and functional complex in thyroid and orbital tissue of GD patients.

Thyroid hormones influence the GH/IGF1 axis, having a permissive role in GH/IGF1 action and a specific effect on plasma levels of some of the GH-independent IGF binding proteins (IGFBPs). Previous studies have reported discrepant results regarding serum levels of IGF1 in hyperthyroidism. In 1995, J. Frystyk et al showed that, in rats, hyperthyroidism increases the circulating low-molecular IGFBPs and induces a reduction in free IGF1, which may play a central role in regulation of IGF bioactivity by thyroid hormones. Another study from the same year reported reduced IGFI circulating levels in hyperthyroid patients, possibly due to nutritional factors. Iglesias et al demonstrated in 2001 that hypothyroidism is associated with significant reductions of IGF1 and IGFBP3 and that IGFBP1 is elevated in both hypothyroidism and hyperthyroidism. Co Ng et al showed, in a study performed on adolescent hyperthyroid patients, that hyperthyroidism does not cause alterations in the serum concentrations of either free or total IGF1. In the same study, both serum IGFBP2 and IGFBP3 concentrations were elevated during hyperthyroidism and correlated with serum T4 levels. These abnormalities reversed with normalization of thyroid function. Other studies revealed that hyperthyroidism is associated with increased serum IGF1 levels, decreased GH responsiveness to GHRH at the pituitary level and a lack of suppressive effect of an oral glucose load at the hypothalamic level. IGF1 was positively correlated with free T3 (FT3) and free T4 (FT4) and negatively correlated with TSH. Lakatos et al reported that IGF1 levels were significantly increased in hyperthyroid patients before treatment, correlated with serum FT4 levels and returned to normal after treatment with methimazole. On the other hand, serum IGF and IGFBP levels were not reported to be elevated in GD patients with active GO.

The aim of this study was to investigate the relationship between serum IGF1 levels and the main characteristics of GD (severity of hyperthyroidism, goiter size, presence of active GO, antithyroid antibodies status and titer) in a group of patients with newly diagnosed Graves’ hyperthyroidism.

SUBJECTS AND METHODS

Patients and study protocol

This cross-sectional study included 98 patients with newly diagnosed GD hyperthyroidism, selected out of 239 patients consecutively evaluated for thyrotoxicosis in our clinic between 2008-2012. Exclusion criteria were: other etiologic forms of thyrotoxicosis, serious concomitant illnesses, pregnancy, diabetes mellitus, treatment with drugs known to influence the pituitary-thyroidal axis, pituitary disorders. GD was defined as the presence of biochemical hyperthyroidism (raised serum FT4 and/or FT3 and/or total T3 (TT3) concentration and suppressed TSH) together with two of the following: 1) diffuse goiter on sonographic scan, 2) significant titer of anti-thyroid peroxidase (TPOAb) and/or anti-thyroglobulin antibodies (ATA) and/or TRAb and 3) the presence of GO. At diagnosis, goiter size was categorized by a single observer on the basis of physical examination according to WHO criteria (http://www.who.int/iris/handle/10665/61278. 12.01.2015). Eye disease was classified according to the presence of eye signs in categories 2-6 of the NOSPECS classification and the clinical activity score of GO was calculated.

Serum TSH, FT4, FT3, TT3, thyroglobulin, TPOAb and ATA were assayed at a single laboratory using a two-site, solid-phase, enzyme-labeled chemiluminescent immunometric assay (Immulite 2000, Siemens Healthcare Diagnostics Products Ltd.). Reference ranges: TSH=0.4-4.0 miu/L, FT4=0.7-1.85 ng/dL, FT3=1.8-4.2 pg/mL, TT3=84-172 ng/dL, thyroglobulin <55 ng/ml, TPOAb=10-35 iu/mL and ATA <40 iu/mL. Serum TRAb levels were measured using a second generation enzyme immunoassay, ELISA.
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kit (DRG Instruments GmbH, Germany). Reference range: <1iu/L negative, >1.5iu/L positive. Antithyroid antibody status was defined as follows: ATA positive= patients with ATA serum levels >40iu/ml; TPOAb positive= patients with TPOAb serum levels >35 iu/ml; TRAb positive= patients with TRAb serum levels >1.5iu/L. Serum tumor necrosis factor alpha (TNF alfa) was measured using an ELISA kit (DRG Instruments GmbH, Germany); reference range: 4.6-12.4 pg/ml. Serum high-sensitivity C-reactive protein (HS-CRP) was assayed using a two-site, solid-phase, enzyme-labeled chemiluminescent immunometric assay (Immulite 2000, Siemens Healthcare Diagnostics Products Ltd.); reference range: <11 mg/L. Serum IGF-1 was assayed using a two-site, solid-phase, enzyme-labeled chemiluminescent immunometric assay (Immulite 2000, Siemens Healthcare Diagnostics Products Ltd.). The analytical sensitivity was 20 μg/l, the intra-assay CV was 2.3-3.9% and the inter-assay CV was 3.7-8.1%. Since IGF1 levels are age-dependent, we calculated the standard deviation score of IGF1 levels according to age (z score). Age-adjusted IGF1 values were referenced from a previously published study by Elmlinger et al\(^\text{18}\) that used the same assay as in our study. Patients were considered IGF1 deficient if IGF1 z score was ≤−2 SD from the mean for age.\(^\text{19}\)

The study protocol was approved by the ethics committee of Elias Hospital. The study was conducted according to the standards of good clinical practice and the Declaration of Helsinki and all study patients signed an informed consent.

**Data presentation and statistical analyses**

Descriptive data are presented as means ±SD, medians with interquartile range (IQR) or percents. IGF1 and IGF1 z score were normally distributed. Because thyroglobulin and ATA were not normally distributed, their levels were transformed to their logarithms before regression analysis and graphical representation. Between-groups comparisons were carried out using parametric (independent sample t-test, one-way ANOVA) or nonparametric (Mann-Whitney U-test, Kruskal-Wallis one-way ANOVA) tests, as appropriate. \(\chi^2\) test and Fisher’s exact test were used to compare proportions in large and small groups, respectively. Relations between continuous variables were analyzed using Pearson’s correla-

**RESULTS**

A total number of 98 GD patients were evaluated: median age 42 (IQR=23) years, 78 (79.6%) women, 33 (33.7%) current smokers, 65 (66.3%) with medium/large goiters (grade 2 WHO goiter grading system) and 24 (24.5%) with active GO. Mean IGF1 at diagnosis was 169.90±82.46 ng/ml and IGF1 z score -0.33±1.63. After the initial laboratory evaluation, all patients received treatment with methimazole at a mean dose of 30.15±7.54 mg/day. The demographic, clinical and laboratory characteristics of the 98 patients are summarized in Table 1.

At diagnosis, men had significantly higher mean serum IGF1 levels (207.13±92.45 vs 160.35±77.49 ng/ml, \(p=0.023\)) and IGF1 z scores (-0.46±1.68 vs -0.54±1.56 ng/ml, \(p=0.013\)) compared to women. Current smokers and nonsmokers had similar serum IGF1 levels (\(p=0.901\)) and IGF1 z scores (\(p=0.663\)). There was no significant association between goiter size at diagnosis and serum IGF1 levels (\(p=0.895\)) or IGF1 z scores (\(p=0.766\)). There was no significant difference regarding serum IGF1 levels (\(p=0.136\)) or IGF1 z scores (\(p=0.070\)) between GD hyperthyroid patients with active GO compared to patients without GO.

No correlations were found between serum inflammatory markers HS-CRP, TNF alfa and IGF1 levels (\(p=0.131\), respectively \(p=0.551\)) or IGF1 z scores (\(p=0.432\), respectively \(p=0.629\)).
Eighty-nine (89) (90.81%) patients were TRAb positive, 66 (67.30%) were TPOAb positive and 38 (38.80%) were ATA positive. TRAb and TPOAb positive patients had similar serum IGF1 levels (p=0.440, respectively p=0.640) and IGF1 z scores (p=0.489, respectively p=0.785) when compared to TRAb and TPOAb negative patients, respectively. ATA positive patients had significantly higher serum IGF1 levels (191.82±72.49 vs 156.01±85.89 ng/ml, p=0.036) and IGF1 z scores (0.12±1.33 vs -0.63±1.74 ng/ml, p=0.024) compared to ATA negative patients.

Serum IGF1 was positively correlated with ATA levels (r= 0.306, p=0.002) and negatively correlated with log serum thyroglobulin (r= -0.394, p<0.001) (Figure 1A). IGF1 z scores were positively correlated with ATA levels (r= 0.306, p=0.002) and negatively correlated with log serum thyroglobulin (r= -0.426, p<0.001) (Figure 1B) and FT3 (r= -0.314, p=0.008) (Figure 2).

Simple linear regression analysis showed that in GD patients, higher FT3 (p=0.044) and thyroglobulin (p<0.001), and respectively lower ATA (p=0.027) levels are associated, at diagnosis, with lower IGF1 z scores (Table 2).

IGF1 deficiency (IGF1 z score ≤-2 DS) was present in 18 (18.4%) patients. Compared to patients without IGF1 deficiency (mean IGF1 67.74±28.96 ng/ml, IGF1 z score -2.64±0.51) presented, at diagnosis, higher thyroglobulin (medi-

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### Table 1. Demographic, clinical and laboratory characteristics of the 98 newly diagnosed GD patients, according to the presence of IGF1 deficiency

<table>
<thead>
<tr>
<th>Parameter</th>
<th>All (N=98)</th>
<th>No IGF1 deficiency (N=80)</th>
<th>IGF1 deficiency (N=18)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age at diagnosis median (IQR), (years)</td>
<td>42 (23)</td>
<td>42 (21)</td>
<td>46 (35)</td>
<td>0.680</td>
</tr>
<tr>
<td>Current smoker N (%)</td>
<td>33 (33.7)</td>
<td>28 (35)</td>
<td>5 (27.8)</td>
<td>0.558</td>
</tr>
<tr>
<td>BMI mean±SD, (kg/m²)</td>
<td>25.25±6.48</td>
<td>24.49±5.53</td>
<td>28.55±9.07</td>
<td>0.023</td>
</tr>
<tr>
<td>Gender N (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Females</td>
<td>78 (79.6)</td>
<td>61 (76.2)</td>
<td>17 (94.4)</td>
<td>0.084</td>
</tr>
<tr>
<td>Ophthalmopathy N (%)</td>
<td></td>
<td></td>
<td></td>
<td>0.533</td>
</tr>
<tr>
<td>Grade 0</td>
<td>3 (3.1)</td>
<td>3 (3.8)</td>
<td>0 (0)</td>
<td></td>
</tr>
<tr>
<td>Grade 1</td>
<td>30 (30.6)</td>
<td>23 (28.7)</td>
<td>7 (38.9)</td>
<td></td>
</tr>
<tr>
<td>Grade 2</td>
<td>65 (66.3)</td>
<td>54 (67.5)</td>
<td>11 (61.1)</td>
<td></td>
</tr>
<tr>
<td>TSH median (IQR), (miu/L)</td>
<td>0.006 (0.010)</td>
<td>0.006 (0.011)</td>
<td>0.005 (0.005)</td>
<td>0.510</td>
</tr>
<tr>
<td>FT4 median (IQR), (ng/dl)</td>
<td>3.57 (3.35)</td>
<td>3.35 (3.20)</td>
<td>4.66 (3.18)</td>
<td>0.235</td>
</tr>
<tr>
<td>TT3 median (IQR), (ng/ml)</td>
<td>331.16 (293.60)</td>
<td>318.50 (283.00)</td>
<td>417.00 (302.00)</td>
<td>0.168</td>
</tr>
<tr>
<td>FT3 median (IQR), (pg/ml)</td>
<td>8.26 (6.30)</td>
<td>7.33 (5.72)</td>
<td>11.30 (7.64)</td>
<td>0.027</td>
</tr>
<tr>
<td>Thyroglobulin median (IQR), (ng/ml)</td>
<td>25.45 (88.55)</td>
<td>11.40 (80.74)</td>
<td>72.55 (116.02)</td>
<td>0.002</td>
</tr>
<tr>
<td>TPOAb median (IQR), (iu/ml)</td>
<td>77.15 (637.62)</td>
<td>234.70 (940.25)</td>
<td>57.45 (578.60)</td>
<td>0.270</td>
</tr>
<tr>
<td>ATA median (IQR), (iu/ml)</td>
<td>20.00 (51.00)</td>
<td>34.05 (161.00)</td>
<td>20.00 (0)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>TRab median (IQR), (iu/L)</td>
<td>6.12 (14.45)</td>
<td>8.50 (15.40)</td>
<td>8.51 (32.86)</td>
<td>0.690</td>
</tr>
<tr>
<td>HS-CRP median (IQR), (mg/L)</td>
<td>2.00 (5.10)</td>
<td>1.91 (5.07)</td>
<td>2.65 (4.51)</td>
<td>0.388</td>
</tr>
<tr>
<td>TNFα median (IQR), (pg/ml)</td>
<td>7.50 (5.12)</td>
<td>7.55 (5.07)</td>
<td>7.50 (5.05)</td>
<td>0.752</td>
</tr>
</tbody>
</table>

Data are presented as percents, mean±SD or median (IQR) according to the type of the variable and the normality of distribution. P value for comparison between GD patients with and without IGF1 deficiency. IGF1 deficiency: IGF1 z score ≤-2 DS.

GD: Graves’ disease; N: number of cases; IGF1: Insulin-like growth factor 1; IQR: interquartile range; SD: standard deviation; BMI: body mass index; TSH: thyroid-stimulating hormone; FT4: free thyroxine; FT3: free triiodothyronine; TT3: total triiodothyronine; TPOAb: anti-thyroperoxidase antibodies; ATA: anti-thyroglobulin antibodies; TRAb: thyroid-stimulating hormone receptor antibodies; HS-CRP: high-sensitivity C-reactive protein; TNF alpha: tumor necrosis factor alpha.
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Figure 1. Correlation between IGF1 (A) and IGF1 z score (B) and log-transformed serum thyroglobulin levels.

Figure 2. Correlation between IGF1 z score and FT3 at diagnosis.

Table 2. Univariate linear regression analysis of FT3, thyroglobulin and anti-thyroglobulin antibodies on IGF1 z score.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Coefficient</th>
<th>95% CI</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>FT3 (pg/ml)</td>
<td>-0.023</td>
<td>-0.045 - 0.001</td>
<td>0.044</td>
</tr>
<tr>
<td>Log thyroglobulin (ng/ml)</td>
<td>-0.224</td>
<td>-0.357 - -0.091</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Log ATA (iu/ml)</td>
<td>0.241</td>
<td>0.028 - 0.454</td>
<td>0.027</td>
</tr>
</tbody>
</table>

IGF1: Insulin-like growth factor 1; FT3: free triiodothyronine; Log thyroglobulin: logarithmic transformed thyroglobulin; Log ATA: logarithmic transformed anti-thyroglobulin antibodies.

Figure 3. Mean log transformed thyroglobulin (A) and antithyroglobulin antibody (B) values, at diagnosis, according to IGF1 status.

Univariate binary logistic regression analysis identified log thyroglobulin (OR: 2.97, 95% CI: 1.403-6.299, p=0.004) and log ATA (OR: 0.114, 95% CI: 0.016-0.802, p=0.029), but not FT3 (p=0.14) as independent predictors of IGF1 deficiency (Table 3). Thyroglobulin was significantly associated with the presence of

an=72.55, IQR=116.02 vs median=11.40, IQR=80.74 ng/ml, p=0.002 (Figure 3A), FT3 (median=11.30, IQR=7.64 vs median=7.33, IQR=5.72 pg/ml, p=0.027) (Figure 4) and lower ATA levels (median=20, IQR=0 vs median=34.05, IQR=161 iu/ml, p<0.001) (Figure 3B) (Table 1).
IGF1 deficiency (AUROC=0.732, 95% CI: 0.620-0.844, p=0.002; cut-off for thyroglobulin=50.40 ng/ml, Se=77.8%, Sp=70%) (Figure 5).

Table 3. Univariate binary logistic regression analysis of the factors associated with IGF1 deficiency.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Odds Ratio</th>
<th>95% CI</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Log ATA (iu/ml)</td>
<td>0.114</td>
<td>0.016-0.802</td>
<td>0.029</td>
</tr>
<tr>
<td>Log Thyroglobulin (ng/ml)</td>
<td>2.97</td>
<td>1.403-6.299</td>
<td>0.004</td>
</tr>
</tbody>
</table>

IGF1: Insulin-like growth factor 1; Log ATA: logarithmic transformed anti-thyroglobulin antibodies; Log thyroglobulin: logarithmic transformed thyroglobulin.

DISCUSSIONS

In the present study, we investigated serum IGF1 levels in a group of hyperthyroid patients with newly diagnosed GD, without prior antithyroid drug treatment. In order to remove the effect of age on serum IGF1 levels we also analyzed IGF1 z scores. Our data showed that in newly diagnosed hyperthyroid GD patients, IGF1 z score is negatively correlated with the severity of FT3 hyperthyroidism and serum thyroglobulin levels and is positively correlated with ATA levels. Thyroglobulin level at diagnosis was significantly associated with the presence of IGF1 deficiency.

The relationship between the thyroid and the GH/IGF1 axis is bidirectional. In patients with autoimmune thyroid diseases, thyroid hormones modulate the synthesis and/or the secretion of IGF1 and IGFBP3, and this function is apparently not mediated by GH. On the other hand, IGF1 and IGF1R play an important role in autoimmunity, the relationship between immune system and growth factors having recently been characterized.

Previous studies reported discrepant results regarding circulating IGF1 levels: lower, unmodified or even higher values in hyperthyroid compared to euthyroid patients. These discrepant results may be due to different laboratory tests used, changes in the number of IGFR1, an interaction between the anabolic effect of IGF-related peptides and the catabolic thyrotoxic condition or changes associated with nutritional signals and/or energy balance.

To the best of our knowledge, there are only six studies that evaluated the plasma IGF-related peptides profile in newly diagnosed hyperthyroid GD patients. The previous studies were conducted on a small number of patients, mostly of different ages and genders. The relationship between IGF-related
peptides and antithyroid antibodies status and titer has been analyzed in only two studies. 20,26

In our population of 98 GD patients, serum IGF1 levels and IGF1 z scores were higher in men compared to women. Previous studies, not performed in hyperthyroid populations, have reported similar, 27 higher 28 or even lower 29 IGF1 levels in men.

Current smoking and goiter size did not influence serum IGF1 levels and IGF1 z scores in our population. In an earlier study conducted on 130 healthy adults, smoking, a known risk factor for GD and GO, was independently and positively associated with serum IGF1 levels. 30 However, a recent study found no correlation between tobacco use and serum IGF1 levels, concluding that smoking does not appear to be a major confounder of the reported clinical associations between IGF1 and specific disease entities. 31

Our 24 hyperthyroid patients with GD and active GO did not present different serum IGF1 levels or IGF1 z scores when compared to hyperthyroid GD patients without GO. Only one study has investigated serum IGF1, IGF2, IGFBP1, IGFBP2 and IGFBP3 in euthyroid GD patients with active GO and reported that IGF1 related peptides are not higher compared to those in healthy adults. 32 Thus, previously reported retrobulbar elevated IGF1 levels seem to be independent of serum IGF1 concentration, probably being an expression of an auto/paracrine activity.

Our study showed no association between TRAb or TPOAb status and titer and serum IGF1 levels or IGF1 z scores. Consistent with our data, Zimmermann-Belsing et al, in a study of 24 hyperthyroid women with GD, did not identify TPOAb and TRAb as significant predictors for total IGF1. 26

In our newly diagnosed GD hyperthyroid patients, higher FT3 and thyroglobulin, and respectively lower ATA serum levels were associated with lower IGF1 z scores (Figures 1, 2). Previous data regarding the relationship between IGF system and thyroid hormones in GD patients are discrepant. Thyroid hormones were either positively correlated with IGFBP3 and/or IGF1 14,11,20 or negatively correlated with IGFBP3 and ALS. 26

Eighteen (18) (18.4%) of our GD patients had, at diagnosis, IGF1 deficiency, defined as IGF1 z score ≤−2 DS. Interestingly, these patients had higher thyroglobulin and FT3, and respectively lower ATA levels (Figure 3, 4). Univariate binary logistic regression analysis identified thyroglobulin and ATA, but not FT3, as independent predictors for IGF1 deficiency. We identified a cut-off level for serum thyroglobulin of 50.40 ng/ml as a strong predictor for IGF1 deficiency, with Se=77.8% and Sp=70%. Patients with thyroid disorders may present with, even several years after reaching euthyroidism, other complications such as: memory impairment, 33 depression, 35 fatigue or body composition changes, 35 signs and symptoms that are also described in patients with GH deficiency. It is known that in thyrotoxicosis protein catabolism predominates, leading to a slight decrease in serum total protein concentration. 36 Furthermore, it was demonstrated that serum IGF1 levels are low in severe catabolic states such as acute postoperative stress, burns, cystic fibrosis or HIV 37 and that the recovery from the catabolic states is associated with increased levels of IGF1. 38 Thus, thyrotoxicosis, a catabolic state, could be associated with a decrease of IGF1 levels and clinical manifestations similar to partial GH deficiency, potentially corroborating our results. These findings are however in contradiction with the data from other studies reporting high IGF1 levels in hyperthyroidism, that decreased after ATDs treatment. 14,26 Zimmermann-Belsing et al suggest that the complex IGF system seems intact in thyrotoxic patients, but changes in body composition and the regulation of leptin and insulin secretion during treatment of autoimmune thyroid disease influence IGF related peptides leaving the patient in a state somewhat similar to partial GH deficiency. 26

The proinflammatory cytokines TNF alfa and IL1 beta often act as negative regulatory signals that temper the action of hormones and growth factors and may induce a state of IGF1 resistance. 39 However, we found no correlation between inflammatory markers HS-PCR, TNF alfa and IGF1 or IGF1 z scores in our GD patients.

Our study has some strengths as well as several limitations. Although we studied a relatively large population of GD patients, we cannot exclude that we could have missed associations of smaller magnitude due to low statistical power. Another limitation is that we did not compare the GD patients IGF1/IGF1 z-score.
values with a control group of healthy individuals, without pathologies known to influence the GH/IGF1 axis. Also, we could not extensively study the IGF protein system (IGF1, free IGF1, IGF2, IGFBP3, ALS) and we did not follow the dynamics of IGF1 after ATD treatment. Nevertheless, we believe that our results contribute some important novel information and deserve further validation in large-scale prospective studies.

In conclusion, the present study has shown for the first time the presence of IGF1 deficiency in nearly one fifth of newly diagnosed GD patients. The IGF1 deficient GD patients’ phenotype was characterized by lower ATA titers, higher thyroglobulin levels and more severe FT3 hyperthyroidism at diagnosis.

ACKNOWLEDGEMENT

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CONFLICTS OF INTEREST

The authors have no multiplicity of interest to disclose.

Note

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REFERENCES

5. Tramontano D, Cushing GW, Moses AC, Ingbar SH, 1986 Insulin-like growth factor-I stimulates the growth of rat thyroid cells in culture and synergizes the stimulation of DNA synthesis induced by TSH and Graves’-IgG. Endocrinology 119: 940-942.
16. Werner SC, 1977 Modification of the classification of the eye changes of Graves’ disease: recommendations of the ad hoc committee of the American Thyroid Asso-


