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Elevated thyroglobulin level is associated with dysfunction of regulatory T cells in patients with thyroid nodules

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Abstract

Objective: Thyroid nodules are usually accompanied by elevated thyroglobulin (Tg) level and autoimmune thyroid diseases (AITDs). However, the relationship between Tg and AITDs is not fully understood. Dysfunction of regulatory T cells (Tregs) plays an important role in the development of AITDs. We aimed to evaluate the effects of Tg on the function of Tregs in patients with thyroid nodules.

Methods: Tg levels and the functions of Tregs in peripheral blood and thyroid tissues of patients with thyroid nodules from Nanjing First Hospital were evaluated. The effects of Tg on the function of Tregs from healthy donors were also assessed in vitro. The function of Tregs was defined as an inhibitory effect of Tregs on the effector T cell (CD4+CD25−T cell) proliferation rate.

Results: The level of Tg in peripheral blood correlated negatively with the inhibitory function of Tregs (R = 0.398, P = 0.03), and Tregs function declined significantly in the high Tg group (Tg >77 μg/L) compared with the normal Tg group (11.4 ± 3.9% vs 27.5 ± 3.5%, P < 0.05). Compared with peripheral blood, the function of Tregs in thyroid declined significantly (P < 0.01), but the proportion of FOXP3+ Tregs in thyroid increased (P < 0.01). High concentration of Tg (100 μg/mL) inhibited the function of Tregs and downregulated FOXP3, TGF-β and IL-10 mRNA expression in Tregs in vitro.

Conclusions: Elevated Tg level could impair the function of Tregs, which might increase the risk of AITDs in patient with thyroid nodules.

Introduction

Thyroid nodules are very common; up to 65% of the general population have at least one thyroid nodule (1), but most of them have no symptoms, and life expectancy is not affected (2). Recent studies have focused mainly on the relationship between thyroid nodule and thyroid cancer (3, 4). Thyroid nodules are usually accompanied by an increase in thyroglobulin (Tg) (5, 6) and autoimmune thyroid diseases (AITDs), especially Hashimoto's thyroiditis (HT) (7, 8). Interestingly, the levels of Tg in patients with thyroid cancer are always considerably higher than in patients with benign thyroid nodules (9, 10). Meanwhile, the incidence of HT in patients with thyroid cancer was also higher than in patients with benign thyroid nodule (11, 12, 13). Those studies implied...
that there might be some interactions between elevated Tg level and HT.

Tg accounts for approximately 75–80% of the total thyroidal protein and serves as a precursor for thyroid hormones (14). Previous studies have proved that Tg could induce lymphocytic thyroiditis by elevating Tg-Ab titers in BB/Wor rats (15) by activating T cells (16). In addition, a number of studies underlined the important roles of regulatory T cells (Tregs) on the prevention of thyroiditis in animal models. Depletion of Tregs enabled the induction of thyroiditis with mouse Tg in traditionally resistant mice as well as iodide in NOD-H2h4 mice (17, 18). Furthermore, clinical studies have suggested that the numerical and/or functional impairments of Tregs were found in patients with HT (19, 20, 21, 22, 23). We questioned whether elevated Tg level could impair the function of Tregs.

Tregs are important subtypes of T cells that are involved in the modulation of the immune response and play essential roles in the prevention of autoimmune disease (24, 25). Natural Tregs (nTregs) develop in the thymus and represent approximately 5–10% of the total number of peripheral CD4+ T cells (25). They are characterized by a high expression of CD25 and the transcription factor forkhead box P3 (FOXP3). Tregs can suppress the proliferation and activity of autoreactive T cells (26).

The aim of our study was to evaluate the effects of Tg on the function of Tregs and to understand the role of Tg in inducing thyroiditis in patients with thyroid nodules.

Materials and methods

Reagents

Human Tg protein was purchased from Fitzgerald Industries International (Acton, MA, USA) and verified to be >98% pure by SDS-PAGE. Anti-human CD3 (Cat# 555336) and anti-human CD28 (Cat# 555725) antibodies were purchased from eBioscience. Allophycocyanin (APC)-conjugated anti-human CD25 (Cat# 555434), PerCP-Cyanine5.5-conjugated anti-human CD4 (Cat# 560650), PE-conjugated anti-human CD25 (Cat# 555432) and APC-conjugated anti-human FOXP3 (Cat# 560045) antibodies were purchased from BD Biosciences (San Jose, CA, USA) and were all of mouse origin. Tg detection kits were purchased from Siemens Healthcare Diagnostics Inc.

Patient details and laboratory methods

Blood was collected from healthy donors with normal ranges of free thyroxin (FT4), thyroid stimulating hormone (TSH), thyroid peroxidase antibody (TPO-Ab) and thyroglobulin antibody (TG-Ab) for experiment in vitro. Donors with thyroid nodules (evaluated by ultrasonography), autoimmune diseases, infections or those on any type of drugs were excluded. Blood and normal thyroid tissues surrounding a thyroid nodule or adenoma in patients who underwent thyroid surgery were also collected at Nanjing First Hospital; these were confirmed by pathology. The indications for operation are according to the Chinese Management Guidelines for Patients with Thyroid Nodules and Differentiated Thyroid Cancer (2012) (27). Besides the patients who underwent thyroid surgery, blood from patients with benign thyroid nodule was also collected, while the diagnoses of nodules were supported by fine-needle aspiration biopsy. Patients with thyroid nodules were excluded if they had any systemic disease or used drugs that may either affect Tg level or immune balance (e.g. infection and autoimmune diseases). The Bioethical Committee of the Nanjing First Hospital approved this study, and written informed consent was obtained from each patient and healthy donor.

Thyroid tissue was homogenized and centrifuged at 380g for 10 min. Supernatant (thyroid follicular fluid) was then collected to measure Tg levels. Blood and thyroid Tg levels were measured using chemiluminescent immunometric assays (Elecsys TG II, Roche Diagnostics GmbH) on a Modular Analytics E170 analyzer (Roche Diagnostics GmbH), and the normal reference range was 3.5–77 ng/mL as described in previous study (6). The total protein content of thyroid follicular fluid and plasma samples was measured using a BCA Protein Assay Kit (Sigma Aldrich), following the manufacturer’s instructions. Thyroid volume before surgery was obtained by computing the volumes of both lobes (lobe (mL) = length (cm)×width (cm)×depth (cm)×0.479). Nodules and/or cystic areas were included in the thyroid volume (reference values, 18 mL for females and 25 mL for male patients) (28).

Isolation of peripheral blood and thyroid Tregs

Peripheral blood mononuclear cells (PBMCs) were isolated using Ficoll-Paque density (Sigma Aldrich) gradient centrifugation. For the separation of thyroid mononuclear cells, thyroid specimens were homogenized and passed
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PBMC stimulation with Tg

Isolated PBMCs were counted and cultured in RPMI-1640 medium supplemented with 10% FBS at 37°C under 5% CO₂. Different concentrations of Tg (0, 0.1, 1, 10 and 100 μg/mL) were added; then the cells were incubated for 3 days. Similar to earlier experiments, anti-human CD3 (5 × 10⁻⁸ mol/L) and anti-human CD28 (2.27 × 10⁻⁸ mol/L) antibodies were also added to the wells. After 3 days, Tregs were isolated from the mononuclear cells using a human Treg cell MACS kit, and then RNA was isolated for further analysis.

RNA isolation and real-time PCR

Cells were homogenized in TRIZol reagent (Sigma-Aldrich), and RNA extraction was performed using TaKaRa RNAiso reagent (TaKaRa). For reverse transcription, cDNA was synthesized using a PrimeScript™ RT Master Mix (TaKaRa). The real-time PCR efficiency for each primer pair was calculated using standard curves generated through serial dilution of cDNA from Tregs. Primer sequences are shown in Table 1. PCR reactions were performed using an ABI PRISM 7500 Sequence Detector (Applied Biosystems). The PCR conditions were set as follows: an initial incubation step for 30 s at 95°C followed by 40 cycles of 5 s at 95°C and 34 s at 60°C. The normalized expression values for each transcript were calculated as the quantities of FOXP3, TGF-β and IL-10 mRNA relative to the quantity of β-Actin mRNA using the 2⁻ΔΔCt method. All reactions were performed independently at least three times.

Flow cytometric analysis of Treg cell phenotypes

PBMCs were treated with or without Tg (100 μg/mL) for 3 days. Cells were then washed and stained with PerCP-Cyanine5.5-anti-CD4, PE-anti-CD25 and APC-anti-FOXP3. Cells were analyzed using a FACS Canto™ II (BD Biosciences).

Statistical analysis

Statistical analysis was performed using SPSS 16.0 Software (SPSS, Inc.). Differences between two groups were analyzed using Student’s t-test. The relationships were analyzed using FACS Canto™ II (BD Biosciences). The inhibition rate (%) was calculated using the formula: ((Teffs proliferation alone – Treg-Teffs proliferation with Treg)/Teffs proliferation alone) × 100%.

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Immunofluorescence staining

Thyroid tissue sections, incubated with monoclonal rabbit anti-FOXP3 (Cell Signaling Technology, dilution 1:500) were subsequently incubated for 30 min with goat anti-rabbit immunoglobulin antibodies conjugated to Alexa-Fluor 488 (DAKO). DAPI (4′,6-diamino-2-phenylindole, KEYGEN, Jiangsu, CN) was used to label nuclear DNA. Appropriate isotype antibodies were used as negative controls. The labeled sections were imaged using a fluorescence microscope.

Functional analysis of Tregs

The main function of Tregs was to inhibit Teffs proliferation. We tested the function of Tregs as described previously (23, 29): Tregs (>97% pure) were cultured with constant amount of Teffs cells at various ratios (Treg:Teffs = 1:2, 1:4, and 1:8 or 1:10) in medium containing 10% FBS at 37°C under 5% CO₂. Anti-human CD3 (5 × 10⁻⁸ mol/L) and anti-human CD28 (2.27 × 10⁻⁸ mol/L) antibodies were added to the cultures. Cells were simultaneously treated with Tg at different concentrations (0, 0.1, 1, 10, and 100 μg/mL) and cultured for 3 days in 24-well plates (5 × 10⁴ cells/well). Thyroid Tregs were also co-cultured with homologous peripheral blood Teffs. To determine cell proliferation, carboxyfluorescein succinimidyl ester (CFSE; BD Biosciences, San Jose, CA, USA) was added to the T cell suspension in an RPMI-1640 medium at a final concentration of 5 × 10⁻⁶ mol/L. Cells were incubated for 15 min at 37°C and then treated with APC-anti-CD25 for another 30 min at 4°C. T cell proliferation was analyzed using FACS Canto™ II (BD Biosciences). The inhibition rate (%) was calculated using the formula: ((Teffs proliferation alone – Treg-Teffs proliferation with Treg)/Teffs proliferation alone) × 100%.

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between Tg levels in the blood and Treg cell functionality were analyzed by multiple linear regression. Differences in the functionality of Tregs from a single individual were analyzed using a paired Student’s t-test. Results were presented as means±s.e.m., or as median (IQR), and a P value<0.05 was considered statistically significant.

Results

Relationship between the functions of Tregs and Tg levels in the peripheral blood

The functions of Tregs and Tg levels in the peripheral blood of 30 patients with benign thyroid nodules were detected at the same time. Linear regression analysis showed that Tg level was negatively correlated with the inhibitory function of Tregs in the peripheral blood (R=−0.398, P=0.03, Fig. 1A). This association remained significant when gender; age; TSH, FT3, FT4 and TG-Ab levels; nodule size and number and thyroid volume were included in a stepwise multiple regression analysis (standardized β=−0.398, P=0.03). We also compared the function of Tregs in the high (Tg >77 μg/L, according to the normal reference range) and normal (Tg ≤77 μg/L) Tg groups and found a significantly declined Tregs function in the high Tg group (11.4 ± 3.9% vs 27.5 ± 3.5%, P<0.05, Fig. 1B). The clinical characteristics of the patients in the two groups are shown in Table 2, and thyrotrophin receptor antibody (TR-Ab), TPO-Ab and Tg-Ab levels were slightly higher in the high Tg group than in the normal Tg group, although the levels of these antibodies in the two groups were within the normal ranges. Moreover, the nodular size and number, and the thyroid volume in the two groups were similar (P>0.05 for all, Table 2).

Inhibitory functions and proportion of thyroid Tregs

The levels of Tg and the ratio of Tg to total protein in the thyroid follicular fluid were significantly higher than those in the peripheral blood (P<0.001, Fig. 2A). We found that FOXP3+ Tregs were concentrated in several thyroid follicles (Fig. 2B). The proportion of FOXP3+ Tregs among CD4+ T cells in thyroid tissue was higher than in peripheral blood (P<0.01, Fig. 2C). We wondered whether elevated Tg level could affect the function of Tregs in the thyroid. Thyroid Tregs were co-cultured with peripheral blood Teffs from the same patient for 3 days in different ratios (1:2, 1:4 and 1:8). The function of peripheral blood Tregs was tested as control. The results showed that the inhibitory function of thyroid Tregs was

Table 1  Primer sequences for real-time PCR.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward (5′→3′)</th>
<th>Reverse (5′→3′)</th>
</tr>
</thead>
<tbody>
<tr>
<td>β-ACTIN</td>
<td>ATCTGCTGGAAGTAGCAGCGA</td>
<td>CCCAGCAAGATGACATGATCAT</td>
</tr>
<tr>
<td>FOXP3</td>
<td>GTGGCCCCAGTGTGGAGAAG</td>
<td>GAGGCCCTTCTGCGATGGA</td>
</tr>
<tr>
<td>TGF-β</td>
<td>AAGCAAGTCGCTTGCTGC</td>
<td>CAGTCGCTACCCGCTTAG</td>
</tr>
<tr>
<td>IL-10</td>
<td>GCTGGAGGACCTTAAGGGTTAC</td>
<td>ATGTCTGGCTTTGTTGC</td>
</tr>
</tbody>
</table>

FOXP-3, Forkhead box P3; IL-10, interleukin-10; TGF-β, transforming growth factor-β.
weaker than that of the peripheral blood Tregs \((P<0.01, \text{Fig. 3A and B}).\)

### Effects of Tg on the function of Tregs

To determine the impact of Tg on the function of Tregs, Tregs and Teffs from peripheral blood of healthy donors were co-cultured for 3 days in various ratios (1:2, 1:4 and 1:8) with different concentrations of Tg (Fig. 4A). We found that a high concentration of Tg (100\(\mu\)g/mL) significantly suppressed the inhibitory function of Tregs compared with control \((P<0.05).\) Conversely, low concentration of Tg (1\(\mu\)g/mL) promoted the inhibitory function of Tregs compared with control \((P<0.01\) with Table 2)

<table>
<thead>
<tr>
<th>Clinical characteristic</th>
<th>Normal Tg group</th>
<th>High Tg group</th>
<th>(P) value</th>
<th>Normal range</th>
</tr>
</thead>
<tbody>
<tr>
<td>(n)</td>
<td>21</td>
<td>9</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Gender (male, %)</td>
<td>23.8</td>
<td>22.2</td>
<td>1.000</td>
<td>–</td>
</tr>
<tr>
<td>Age (year)</td>
<td>49.0 ± 2.4</td>
<td>52.7 ± 4.3</td>
<td>0.430</td>
<td>–</td>
</tr>
<tr>
<td>TSH (mIU/L)</td>
<td>2.1 ± 0.2</td>
<td>2.0 ± 0.3</td>
<td>0.770</td>
<td>0.35–4.94</td>
</tr>
<tr>
<td>FT3 (pmol/L)</td>
<td>4.4 ± 0.1</td>
<td>4.0 ± 0.3</td>
<td>0.161</td>
<td>2.63–5.70</td>
</tr>
<tr>
<td>FT4 (pmol/L)</td>
<td>15.7 ± 0.4</td>
<td>15.5 ± 1</td>
<td>0.792</td>
<td>9.0–19.0</td>
</tr>
<tr>
<td>TR-Ab (IU/L)</td>
<td>&lt;0.3</td>
<td>0.3 (0.4, 0.5)</td>
<td>–</td>
<td>&lt;1.75</td>
</tr>
<tr>
<td>TPO-Ab (IU/L)</td>
<td>5.0 (5.0, 7.7)</td>
<td>7.1 (5.0, 12.7)</td>
<td>0.255</td>
<td>&lt;34.0</td>
</tr>
<tr>
<td>Tg-Ab (IU/mL)</td>
<td>11.2 (10.0, 20.4)</td>
<td>21.3 (11.3, 24.9)</td>
<td>0.095</td>
<td>&lt;115.0</td>
</tr>
<tr>
<td>Nodular size (mm)</td>
<td>17.9 ± 4.1</td>
<td>20.9 ± 6.8</td>
<td>0.695</td>
<td>–</td>
</tr>
<tr>
<td>Nodular number</td>
<td>1.8 ± 0.1</td>
<td>1.8 ± 0.1</td>
<td>0.928</td>
<td>–</td>
</tr>
<tr>
<td>Thyroid volume (mL)</td>
<td>11.00 ± 1.2</td>
<td>38.59 ± 23.45</td>
<td>0.274</td>
<td>Females &lt;18 mL</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Male &lt;25 mL</td>
</tr>
</tbody>
</table>

FT3, free triiodothyronine; FT4, free thyroxin; \(n\), number; Tg, thyroglobulin; Tg-Ab, thyroglobulin antibody; TPO-Ab, thyroid peroxidase antibody; TR-Ab, thyrotrophin receptor antibody; TSH, thyroid-stimulating hormone.

**Figure 2**

TG and Tregs in thyroid tissue. Thyroid tissues were taken from seven patients who had undergone surgery for benign thyroid nodules. The concentrations of Tg (A) and the ratio of Tg/total protein in the thyroid follicular fluid and peripheral blood were measured by chemiluminescent immunometric assay. (B) Immunofluorescence microscopy of normal thyroid sections. DAPI was used to label nuclear DNA and FOXP3 was used to label Tregs. Original magnification ×200. (C) Thyroid mononuclear cells were stained with anti-CD4-PerCP-Cyanine5.5, anti-CD25-PE, and anti-FOXP3-APC, as well as PBMCs. The proportion of CD4*CD25*FOXP3* cells in the CD4* T cell population was determined by flow cytometry. The experiment was performed seven times independently. **\(P<0.01\).**
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Figure 3
Suppressive functionality of thyroid Tregs. (A) Thyroid tissues were taken from six patients who had undergone surgery for benign thyroid nodules. Thyroid Tregs were co-cultured with peripheral blood effector T cells (Teffs, CD4^+ CD25^- T cells) at different ratios (1:2, 1:4, and 1:8) for 3 days. Teffs were labeled with CFSE and analyzed by flow cytometry. (B) The inhibitory function of Tregs were analyzed using the inhibition rate (%) of Teffs proliferation and was calculated using the equation: ([proliferation of Teffs – proliferation of Teffs plus Treg]/proliferation rate of Teffs alone) × 100%. The experiment was performed independently six times.

Treg/Teff = 1/2 and P<0.05 with Treg/Teff = 1/4 and 1/8, Fig. 4B). In order to further investigate the underlying mechanisms of Tg on the function of Tregs, we isolated Tregs from peripheral blood of healthy donors and treated them with different concentrations of Tg (0, 0.1, 1, 10 and 100 μg/mL) for 3 days. We found that a low concentration of Tg (1 μg/mL) significantly stimulated the mRNA expression of FOXP3, TGF-β and IL-10; P for all was <0.01 (Fig. 5). However, higher concentrations of Tg (100 μg/mL) inhibited the mRNA expression of FOXP3, TGF-β and IL-10 (P<0.01) (Fig. 5). To examine the direct effect of Tg on Teffs proliferation, we also added 100 μg/mL Tg to Teffs without Treg. The proliferation rate of Teff in 100 μg/mL Tg group and blank group were 56.2±6.8% vs 65.7±9.1%, respectively, P=0.055.

Discussion

In the present study, we found a correlated dysfunction of Tregs with Tg level in patients with benign thyroid nodules and a significantly decreased function of Tregs in the high Tg group (Tg >77 μg/L) compared with the normal Tg group. Serum Tg levels could be elevated in most proliferative thyroid diseases (5, 30), and the numerical and/or functional impairments of Tregs have been shown to be involved in the pathogenesis ofAITDs in humans (18, 23, 31, 32). The mechanism of this high incidence ofAITDs might be due to the high level of Tg, which impairs the function of Tregs in thyroid nodules and cancer.

We also confirmed that high levels of Tg could affect the function of Tregs in vitro, with decreased FOXP3, IL-10 and TGF-β expressions. FOXP3 was not only a molecular marker and cell lineage specification factor for Tregs, but also critical for the regulatory activity of Tregs (33). Higher levels of FOXP3 expression suppresses the activity of non-regulatory T-cells in rodents (34). In humans, mutations in the FOXP3 gene are linked to Tregs deficiency, causing a severe systemic autoimmune disease called immune dysregulation, polyendocrinopathy, enteropathy and X-linked syndrome (35). IL-10 and TGF-β are soluble factors that are expressed by Tregs, and they have direct suppressive effects on Teffs (36, 37). High levels of Tg affecting the function of Tregs might be due to the inhibitory effects of Tg on the expression of FOXP3, IL-10 and TGF-β in Tregs. Inversely, low levels of Tg could improve the function of Tregs and stimulate the mRNA expression of FOXP3, TGF-β and IL-10. Previous studies suggested that 100% CBA/J mice developed thyroiditis with positive Tg-Ab when high concentration of Tg (100μg) was injected i.v. once (17), but only 17% of the mice developed thyroiditis after low concentration of Tg (40 μg daily) injection for 10 days (38). None of the mice developed thyroiditis when they were injected with the low concentration of Tg (20 μg daily) and TSH (0.251U daily) (38). The results of these studies might be due to the effects of the different concentrations of Tg on Tregs.

Although high concentrations of Tg suppressed Treg cell function, low concentrations had the opposite effect in vitro, suggesting that Tg may act differently to
regulate Treg cell function depending on whether it is in the thyroid tissue or peripheral blood. The bidirectional effects of Tg on Treg cells were similar to the effects of Tg on the growth of the thyroid follicular epithelial cells (39, 40). The induction of thyroid cell growth was noted only at lower concentrations of Tg, whereas the inhibition of cell growth and follicular function were induced dose dependently at higher concentrations of Tg.

Suzuki et al. suggested that different recognition systems, rather than the two different domains of the Tg protein,
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Endocrine Connections

8:4

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