Reversible Changes in Brain Glucose Metabolism Following Thyroid Function Normalization in Hyperthyroidism


AJNR Am J Neuroradiol 2011, 32 (6) 1034-1042
doi: https://doi.org/10.3174/ajnr.A2449
http://www.ajnr.org/content/32/6/1034
Reversible Changes in Brain Glucose Metabolism Following Thyroid Function Normalization in Hyperthyroidism

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BACKGROUND AND PURPOSE: Patients with hyperthyroidism frequently present with regional cerebral metabolic changes, but the consequences of endocrine-induced brain changes after thyroid function normalization are unclear. We hypothesized that the changes of regional cerebral glucose metabolism are related to thyroid hormone levels in patients with hyperthyroid, and some of these changes can be reversed with antithyroid therapy.

MATERIALS AND METHODS: Relative regional cerebral glucose metabolism was compared between 10 new-onset untreated patients with hyperthyroidism and 20 healthy control participants by using brain FDG-PET scans. Levels of emotional distress were evaluated by using the SAS and SDS. Patients were treated with methimazole. A follow-up PET scan was performed to assess metabolic changes of the brain when thyroid functions normalized.

RESULTS: Compared with controls, patients exhibited lower activity in the limbic system, frontal lobes, and temporal lobes before antithyroid treatment. There were positive correlations between scores of depression and regional metabolism in the cingulate and paracentral lobule. The severity of depression and anxiety covaried negatively with pretreatment activity in the inferior temporal and inferior parietal gyri respectively. Compared with the hyperthyroid status, patients with normalized thyroid functions showed an increased metabolism in the left parahippocampal, fusiform, and right superior frontal gyri. The decrease in both FT3 and FT4 was associated with increased activity in the left parahippocampal and right superior frontal gyri.

CONCLUSIONS: The changes of regional cerebral glucose metabolism are related to thyroid hormone levels in patients with hyperthyroidism, and some cerebral hypometabolism can be improved after antithyroid therapy.

ABBREVIATIONS: BA = Brodmann area; β value = standardized coefficients; FDG = fluorodeoxyglucose; 18F-FDG = fluorine 18 fluorodeoxyglucose; FDR = false discovery rate; FT3 = free triiodothyronine; FT4 = free thyroxine; GLUT-1 = glucose transporter protein type 1; MNI = Montreal Neurological Institute; MRS = MR spectroscopy; PET = positron-emission tomography; SAS = Self-Rating Anxiety Scale; SDS = Self-Rating Depression Scale; SPECT = single-photon emission tomography; SPM = statistical parametric mapping; SPSS = Statistical Package for the Social Sciences; T1 = before antithyroid therapy; T2 = after antithyroid therapy; TRAb = anti-thyroid stimulating hormone receptor antibody; TSH = thyrotropin; VOI = volume of interest

Received September 2, 2010, accepted after revision October 18.

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This work was supported by grants from the National Natural Science Foundation of China (30771024, 30900502, 30800344, and 81070680); the Shanghai Committee of Science and Technology (10JC1401002); the Young Science Fund of Fudan University (09FQ71); the Institutes of Biomedical Sciences, Fudan University (2009056); and the Graduate Innovation Fund of Fudan University (FYF151063).

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DOI 10.3174/ajnr.A2449

Thyroid dysfunctions may be accompanied by numerous neurologic and psychiatric changes. The best well known is cognitive impairment and depression in patients with hypothyroidism as well as an increased risk of cerebrovascular accidents. Also, thyrotoxicosis induces a wide range of cognitive and affective-emotional impairments and is considered 1 common cause of emotional distress. Nervousness, irritability, impulsiveness, poor concentration, easy fatigability, and depressed moods are frequent symptoms in thyrotoxicosis, sometimes suggesting primary neuropsychiatric disease. The links between these clinical phenomena and brain metabolic function are poorly understood.

With advances in brain imaging techniques, a combination of endocrine and psychological testing as well as PET may provide further insight into the underlying mechanisms of the functional neuroendocrine relationships. Functional imaging may contribute to elucidating the potentially different backgrounds of psychiatric complications for patients with hyperthyroidism. To date, there are very few studies on cerebral glucose metabolism dealing with hyperthyroidism-associated changes, which mostly report regional changes of cerebral glucose metabolism before therapy.

Here, we present findings from neuropsychiatric evaluations and cerebral glucose metabolism measurements in individuals with hyperthyroidism before and after antithyroid therapy.
therapy. Of particular interest was the possible correlation of subjective disability with abnormal function in specific brain regions, and the relationship of these changes to thyroid function recovery. We hypothesized that the changes of regional cerebral glucose metabolism are related to thyroid hormone levels in patients with hyperthyroidism, and some of these changes can be reversed with antithyroid therapy.

Materials and Methods

Study Design
Screening procedures included complete medical examinations, routine laboratory evaluations, MR imaging (Magnetom Verio 3T; Siemens, Erlangen, Germany) to exclude participants with potential significant brain diseases, and a neuropsychiatric evaluation. Patients with hyperthyroidism and healthy control subjects were assessed by measurements of behavioral and somatic symptoms and PET scanning twice. For the hyperthyroid group, assessments were performed before and after antithyroid therapy. The initial dose of methimazole was 10 mg, 3 times a day. During treatment, the thyroid hormone levels (FT3 and FT4) were measured every 20 days. The second PET scans were performed when the symptoms had disappeared and the thyroid hormone levels were normal. Then the dose of methimazole was reduced to 10 mg a day. Control subjects underwent the second scan on the same day as the corresponding patients with hyperthyroidism, but they did not receive methimazole treatment. The index of brain glucose metabolism was normalized by using raw counts from the radiotracer FDG as a surrogate marker. Relative activity, as used in this report, refers to this measure. The protocols were approved by the ethics committees of Huashan Hospital, Fudan University in Shanghai, and all studies were performed after informed consent was obtained.

Patients
Ten consecutive patients with Graves disease—caused hyperthyroidism (mean age, 38.7 ± 9.3 years; 5 women; mean TSH-receptor antibodies, 24.68 ± 15.09 U/L; normal range, <1 U/L) were studied at the Endocrine Department of Huashan Hospital, Fudan University. All patients were newly diagnosed. Before entering the study, all subjects were screened and clinically examined by 2 senior investigators. The exclusion criteria were the following: endocrine or autoimmune diseases except for hyperthyroidism, hyperthyroidism crisis, pregnancy, no contraception, intake of any psychotropic medication or β-blocking agents in the 2 months before enrollment, history of bipolar or schizoaffective disorder, and suicide risk (determined on psychiatric interview).

Twenty control subjects underwent the same psychiatric screening procedures as subjects with hyperthyroidism. Inclusion criteria for the age- and sex-matched control group were the following: contraception; no intake of any psychotropic medication, β-blocking agents, thyroid hormones, or antithyroid medication in the past 4 months; and no history of psychiatric, endocrine, neurologic, or other severe medical disease. In both groups, individuals with current dependence on alcohol and illegal drugs or a positive urine test for these substances were excluded.

Data Collection
Anthropometric and other information on age, sex, body-mass index, medication use, diagnosis, and smoking and drinking history were obtained for all patients and control participants. Serum FT3, FT4, TSH, TRAb, and fasting plasma glucose levels were measured on the same investigation day. Fasting plasma glucose was measured by the glucose oxidase method (7600 automatic biochemical analyzer; Hitachi High-Tech Science Systems Corporation, Osaka, Japan). Others were measured by radioimmunoassay with commercially available kits (Vitros Immunodiagnostic Products; Ortho-Clinical Diagnostics, a Johnson & Johnson company, San Jose dos Campos, SP, Brazil).

Psychometric Tests
The emotional distress of each patient was investigated at the beginning and end of the study. Anxiety was assessed by the Zung SAS, and depression was measured by the Zung SDS. Using either the SAS or SDS, we identified 20 items in factor analytic studies for anxiety or depression that corresponded with the definition of the Diagnostic and Statistical Manual of Mental Disorders. Positively and negatively worded items reduce response bias, and reversed items act as a lie scale. Total standard scores ranged from 25 to 100, with higher scores indicative of greater anxiety or depression. Zung stated that total standard scores of ≥45 indicated that participants had “clinically significant” anxiety and ≥50 indicated that respondents could be described as having “clinically significant depression.” The SAS and SDS have been extensively evaluated, showing good sensitivity, specificity, and reliability for diagnosing anxiety and depression.

PET and Image Reconstruction
Every subject underwent a PET-CT (with 18F-FDG) scan at the beginning of the study, and all patients with hyperthyroidism underwent a second scan after recovery. Before the PET scan, patients were asked to fast for 8 hours but had free access to water. PET scans were performed with a Biograph 64 PET/CT scanner (Siemens, Erlangen, Germany) in 3D mode. A CT transmission scan was first obtained for attenuation correction. The PET scan was started 45 minutes after an intravenous bolus injection of 185 MBq of 18F-FDG and lasted for 10 minutes. Hanning filters were used during image reconstruction, giving a transaxial and axial cutoff frequency of 0.5. We used normalized count images to measure relative changes in regional glucose metabolism. All studies in patients and healthy controls were performed with the subject’s eyes open in a dimly lit room with minimal auditory stimulation.

Data Analysis
Clinical data analysis was performed with the use of SPSS software, Version 11.5 (SPSS, Chicago, Illinois). For findings before and after therapy, variables were compared by means of a paired Student t test for significant change. All P values presented are 2-tailed, and values <.05 were considered statistically significant.

PET images were preprocessed with SPM (SPM2; Wellcome Department of Cognitive Neurology, London, United Kingdom) implemented in Matlab 6.5.1 (MathWorks, Natick, Massachusetts). Scans from each patient were first realigned between pre- and posttherapy and spatially normalized into MNI brain space with linear and non-linear 3D transformations via the mean image obtained following the realignment. The normalized PET images for patients and healthy volunteers were filtered with a Gaussian kernel of 10 mm (full width at half maximum) in the x, y, and z axes, respectively. Mean signal-intensity differences over the whole brain were removed by proportional scaling.
Table 1: Thyroid measurements and psychological scales in patients with hyperthyroidism before and after antithyroid therapy

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age (yr)</th>
<th>TSH (0.35–5.5 mIU/L)</th>
<th>FT3 (3.5–6.5 pmol/L)</th>
<th>FT4 (10.2–31 pmol/L)</th>
<th>SAS (&lt;45)</th>
<th>SDS (&lt;50)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>57</td>
<td>0.124</td>
<td>3.614</td>
<td>19.58</td>
<td>6.66</td>
<td>42.87</td>
</tr>
<tr>
<td>2</td>
<td>41</td>
<td>0.001</td>
<td>1.71</td>
<td>30.28</td>
<td>4.6</td>
<td>80</td>
</tr>
<tr>
<td>3</td>
<td>42</td>
<td>0.001</td>
<td>0.21</td>
<td>23.03</td>
<td>6.56</td>
<td>67.64</td>
</tr>
<tr>
<td>4</td>
<td>47</td>
<td>0.009</td>
<td>0.36</td>
<td>20.08</td>
<td>6.28</td>
<td>47.61</td>
</tr>
<tr>
<td>5</td>
<td>33</td>
<td>0.001</td>
<td>0.16</td>
<td>21.88</td>
<td>6.3</td>
<td>62.28</td>
</tr>
<tr>
<td>6</td>
<td>35</td>
<td>0.001</td>
<td>4.927</td>
<td>30.8</td>
<td>4.26</td>
<td>95.28</td>
</tr>
<tr>
<td>7</td>
<td>32</td>
<td>0.002</td>
<td>0.35</td>
<td>21.39</td>
<td>6.21</td>
<td>77.97</td>
</tr>
<tr>
<td>8</td>
<td>24</td>
<td>0.001</td>
<td>1.23</td>
<td>25.13</td>
<td>4.1</td>
<td>74.25</td>
</tr>
<tr>
<td>9</td>
<td>33</td>
<td>0.008</td>
<td>0.19</td>
<td>16.11</td>
<td>4.82</td>
<td>35.15</td>
</tr>
<tr>
<td>Mean</td>
<td>38.22</td>
<td>0.016</td>
<td>1.417</td>
<td>23.14</td>
<td>5.53</td>
<td>64.78</td>
</tr>
</tbody>
</table>

* Significant differences were observed between baseline values (T1) and second values (T2) by the Student paired t test.
* P < .05
* P < .001
* P < .005

Effect of Antithyroid Therapy on Regional Glucose Metabolism

To characterize pretherapeutic metabolism in patients with hyperthyroidism with respect to healthy subjects, we performed a group comparison by using a 2-sample t test in SPM2 according to the general linear model at each voxel. The effects of the antithyroid therapy on regional brain metabolism were also estimated by using a paired t test.

To evaluate significant differences, we used an approach with the peak threshold set at P = .001 (uncorrected) for whole brain regions. For hypothesis generation, we reported only clusters that survived an FDR correction at P < .05. Significant regions were localized by Talairach Daemon software (Research Imaging Imaging Center, University of Texas, Austin, Texas) after using a MNI-to-Talairach conversion tool. SPM maps for increased or decreased metabolism were overlaid on a standard MR imaging brain template.

Correlation of Emotional Distress with Glucose Metabolism in Patients with Hyperthyroidism before Antithyroid Therapy

Using SPM2, we correlated baseline SAS and SDS scores with PET images to evaluate whether specific brain regions would correlate with the measures of emotional distress in patients with hyperthyroidism. As an exploratory analysis, hypothesis-driven searches were conducted and considered significant at P < .001, uncorrected for multiple comparisons. SPM maps showing significant correlation between glucose metabolism and emotional distress were overlaid on a standard MR imaging brain template.

Correlation between Changes in Glucose Metabolism and Clinical Outcome in Patients with Hyperthyroidism

To quantify regionally specific metabolic changes, we constructed a 4-mm-radius VOI within the image space, centered at the peak voxel of clusters that were significant in SPM analysis. The VOIs were predefined on the
**Group Differences in Relative Brain Activity before Treatment**

Compared with euthyroid controls, patients with hyperthyroidism showed no significant increases of regional cerebral glucose metabolism. In contrast, patients with hyperthyroidism showed a significantly decreased metabolism in the limbic system, temporal lobe, and frontal lobe, mainly in the parahippocampal gyrus (bilateral BA 35 and left BA 28), right uncus (BA 36), left cingulate gyrus (BA 31), bilateral fusiform gyrus (BA 37), right precentral gyrus (BA 4/6), right superior and middle frontal gyrus (BA 6) (FDR-corrected P < .05; Table 2 and Fig 1).

**Correlation of Emotional Distress with Relative Brain Activity in Patients with Hyperthyroidism before Treatment**

The correlation analysis yielded a significantly negative correlation between the anxiety score in the SAS and the metabolic activity in the left inferior parietal gyrus (BA 40), while the depression score was positively correlated to the metabolic activity of the left cingulate gyrus (BA 31) and paracentral lobule (BA 5). A negative correlation was found between the depression score and the left temporal lobe, mainly the inferior temporal gyrus (BA 20) (uncorrected P < .001; Table 3 and Fig 2).

**Treatment-Related Changes in Relative Cerebral Activity**

The second set of PET scans was performed in all patients when the symptoms had disappeared and the thyroid hormone levels (FT3 and FT4) were normal. Compared with the baseline findings in the hyperthyroid state, the treatment induced a significant cluster of increased metabolic activity in the left parahippocampal gyrus (BA 28/35), right superior frontal gyrus (BA 6), and left fusiform gyrus (BA 37). However, compared with euthyroid controls, the glucose metabolism was still significantly decreased in the left parahippocampal gyrus (BA 35) and fusiform gyrus (BA 37) (FDR-corrected P < .05; Table 4 and Figs 3 and 4). No significant increase was identified for other regions.

**The Associations between the Relative Brain Activity and Thyroid Hormone Levels and Emotional Symptoms of Patients**

The decrease in both FT3 and FT4 was associated with increased relative glucose metabolic rates in the left parahippocampal gyrus (BA 28) and the right superior frontal gyrus (BA 6) (P < .05; Table 5). Other rehabilitation regions were affected by a single thyroid hormone. An increase in the left parahippocampal gyrus activity (BA 35) was positively correlated with a decrease in FT3 (β = 0.820, P = .009), and an increase in the left fusiform gyrus activity (BA 37) was positively correlated with a decrease in FT4 (β = 0.814, P = .012). Furthermore, the increase in activity from the first scan to the second in the right superior frontal gyrus (BA 6) (β = 0.239, P = .013) and the left fusiform gyrus (BA 37) (β = 0.539, P = .007) was positively correlated with a decrease of anxiety symptoms. There was also a negative correlation between the decrease in depressive symptoms and the increase in relative activity in the right superior frontal gyrus (BA 6) (β = −0.505, P = .003).

**Discussion**

This is the first brain functional imaging study about the cerebral metabolic changes in Graves disease before and after antithyroid treatment. In this study, adult patients with previously untreated hyperthyroidism showed less relative activity in multiple brain regions than healthy control subjects. The differences reached statistical significance in the limbic system (parahippocampus, cingulate gyrus, and uncus), temporal and frontal cortices, and all brain centers integral to the neurocircuitry of clinical behavior, mood, and memory regulation. After antithyroid therapy, the patient group displayed a normalization of brain metabolism in some regions, and statistical differences between the 2 groups were no longer detected. Treatment with methimazole specifically increased regional activity in the left parahippocampus and the right superior frontal gyrus. There also was an association between changes in relative brain activity and disease severity, measured by FT3, FT4, and psychometric scores. The findings suggest that thyroid hormones participate in the modulation of cerebral glucose metabolism in the mature brain. The cerebral changes induced by thyrotoxicosis could recover when thyroid hormone levels were normal.

First reported in the 1950s, the effects of hyperthyroidism on cerebral activity described as cerebral blood flow with normal oxygen consumption increased. These early findings...
were interpreted as the result of decreased vascular resistance, instead of primary changes in brain function. Only decades later did investigators discover significant abnormalities in brain metabolism in the hyperthyroid state. MR spectroscopy was used to demonstrate a relatively decreased choline/creatinine ratio of the right prefrontal cortex in patients with hyperthyroidism, a finding thought to reflect an altered brain cholinergic-adrenergic balance in hyperthyroidism. Recent functional brain imaging studies by using SPECT and FDG-PET demonstrated that thyroid hormones affect regional brain metabolism in patients with hyperthyroidism. These studies showed a significantly reduced blood flow of the bilateral temporal-parietal cortex or a decreased metabolism of the uncus and the inferior temporal gyrus. Increased breathing and heart rates might inhibit metabolism of the brain stem. Also, in our study, hyperthyroidism-associated regional hypometabolism was noted.

Compared with previous imaging studies of hyperthyroidism, our study detected altered metabolism in regions beyond the uncus, temporal cortex, and prefrontal cortex. In addition to the posterior cingulate and the inferior parietal lobe, new cerebral regions correlated with anxiety and depression scales were identified. These may be due to differences in patient samples, severity of disease, and imaging methodology. In the previous studies, the number of patients was relatively small or the state of hyperthyroidism was mild. Furthermore, different neuroimaging techniques, (eg, proton MR spectroscopy and SPECT with lipophilic radiotracers) were used in the studies of patients with hyperthyroidism. Most interesting, the

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**Table 3: The relationship of cerebral activity with SAS and SDS in patients with hyperthyroidism**

<table>
<thead>
<tr>
<th>Relationship</th>
<th>Voxel Levels</th>
<th>Uncorrected Z Score</th>
<th>Peak Voxel*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>SAS</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Parietal lobe</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Inferior parietal gyrus (BA 40)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>&lt;.001</td>
<td>4.83</td>
<td>-48 -41 44</td>
</tr>
<tr>
<td>Left</td>
<td>&lt;.001</td>
<td>3.63</td>
<td>-57 -33 37</td>
</tr>
<tr>
<td>Limbic lobe</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cingulate gyrus (BA 31)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>&lt;.001</td>
<td>4.25</td>
<td>-12 -29 44</td>
</tr>
<tr>
<td>Frontal lobe</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Paracentral lobule (BA 5)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>&lt;.001</td>
<td>3.44</td>
<td>-6 -40 55c</td>
</tr>
<tr>
<td>Temporal lobe</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Inferior temporal gyrus (BA 20)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Left</td>
<td>&lt;.001</td>
<td>3.86</td>
<td>-55 -45 -15</td>
</tr>
</tbody>
</table>

*a Uncorrected P value < .001.
*b Talairach coordinates.
*c Positive covariations; the others are negative covariations.

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were interpreted as the result of decreased vascular resistance, instead of primary changes in brain function. Only decades later did investigators discover significant abnormalities in brain metabolism in the hyperthyroid state. MR spectroscopy was used to demonstrate a relatively decreased choline/creatinine ratio of the right prefrontal cortex in patients with hyperthyroidism, a finding thought to reflect an altered brain cholinergic-adrenergic balance in hyperthyroidism. Recent functional brain imaging studies by using SPECT and FDG-PET demonstrated that thyroid hormones affect regional brain metabolism in patients with hyperthyroidism. These studies showed a significantly reduced blood flow of the bilateral temporal-parietal cortex or a decreased metabolism of the uncus and the inferior temporal gyrus. Increased breathing and heart rates might inhibit metabolism of the brain stem. Also, in our study, hyperthyroidism-associated regional hypometabolism was noted.

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**Fig 1.** Group differences in regional cerebral activity before treatment (FDR-corrected at \( P < .05 \)). The figures are depicted in neurologic orientation. The gray-scale image is a T1 structural MR image that is representative of MNI space. Blue areas represent where the patients with hyperthyroidism exhibited lower activity relative to control subjects.
methods used to analyze data from the PET studies varied. Many PET studies were based on a region-of-interest approach that examined select regions originally reported by earlier studies. The region-of-interest method requires that regions be manually delineated, a subjective and technically laborious process for which operators must be rigorously trained.

Also, methods for manually defining anatomic landmarks differed among studies; this difference may have lead to inconsistencies in reported volume changes. Additionally, an a priori hypothesis is required because researchers must choose specific regions to investigate on the basis of prior evidence. Therefore, critically implicated regions may have been overlooked in most previous studies using region-of-interest approaches. However, a whole brain based analysis by using a voxel-level analysis method was performed in our study to localize more areas not previously examined in earlier region-of-interest studies. A key feature of this rapid and automated method of analysis is that it examines differences in gray matter throughout the brain, without the need to prespecify regions of interest for investigation. This unbiased approach is of value both to confirm the validity of the hypothesis developed in region-of-interest studies and to reveal gray matter differences in areas not previously considered.

Our findings are also different from neuroimaging studies of mood disorders in euthyroid humans. Neuroimaging studies of major emotional dysfunction in euthyroid humans have identified neurophysiologic abnormalities in multiple areas of the frontal and temporal cortices, the insula, the amygdala, and related parts of the striatum and thalamus. Some of these abnormalities persist following symptom remission, and postmortem studies demonstrated that these regions had cortex volume reductions and histopathologic changes. Besides these regions, we found significant hypoxia in the cingulate gyrus, involved in behavior, learning, and memory. Some of the hypoactive regions, such as the left parahip-

Fig 2. Covariation of relative brain activity before treatment in the hyperthyroid group with psychometric scores (uncorrected $P < .001$). Blue areas exhibit negative correlation and red areas exhibit positive correlation between the psychometric score and normalized glucose metabolism in the patients with hyperthyroidism.

Table 4: Treatment effects: comparison of changes in relative cerebral activity between the first and second PET assay in patients with hyperthyroidism (effect of treatment with methimazole) and differences between patients with hyperthyroidism after treatment and control subjects

<table>
<thead>
<tr>
<th>Patients with Hyperthyroidism</th>
<th>Voxel Levels</th>
<th>Corrected $P$ Value</th>
<th>Z Score</th>
<th>Peak Voxel*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Increase</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Parahippocampal gyrus (BA 35)$^b$</td>
<td>.006</td>
<td>4.41</td>
<td>-22</td>
<td>-5</td>
</tr>
<tr>
<td>Left</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Parahippocampal gyrus (BA 28)$^b$</td>
<td>.022</td>
<td>4.17</td>
<td>-18</td>
<td>-11</td>
</tr>
<tr>
<td>Left</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Superior frontal gyrus (BA 6)$^b$</td>
<td>.009</td>
<td>4.33</td>
<td>14</td>
<td>7</td>
</tr>
<tr>
<td>Right</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fusiform gyrus (BA 37)$^b$</td>
<td>.005</td>
<td>4.46</td>
<td>-40</td>
<td>-63</td>
</tr>
<tr>
<td>Left</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
| Cured compared with healthy controls
| Parahippocampal gyrus (BA 35)$^b$ | .015        | 4.29                | -22     | -5          | -25        |
| Left                        |             |                     |         |             |
| Parahippocampal gyrus (BA 28) | .068        | 3.97                | -18     | -11         | -21        |
| Left                        |             |                     |         |             |
| Superior frontal gyrus (BA 6) | .395        | 3.79                | 14      | 7           | 60         |
| Right                       |             |                     |         |             |
| Fusiform gyrus (BA 37)$^b$                           | .040        | 4.09                | -40     | -63         | -14        |
| Left                        |             |                     |         |             |

* Talairach coordinates.
$b$ Survived at FDR-corrected $P < .05$. 

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Pocampal gyrus (BA 28) and the right superior frontal gyrus (BA 6), became normal and correlated positively with the decrease of FT3 and FT4 (Tables 4 and 5). This finding may implicate primary effects of thyroid hormones that result in the emotional, motivational, cognitive, and behavioral manifestations of affective disorders. Also, the observed correlations to the cerebral metabolism may be directly induced by the elevated thyroid hormone concentrations and were not a secondary epiphenomenon due to hyperthyroid-associated mood changes. The effect on the cerebrum was mediated by...
actions of thyroid hormones on thyroid hormone nuclear receptors. Therefore, with increased disease severity, there is lower metabolism in high-concentration regions of T3 nuclear receptors. Thyroid hormones (age-dependent) also modulate the expression of the cerebral GLUT-1, suggested by the reduced 55-kDa GLUT-1 isoform in both hyperthyroid and hypothyroid young rats. The GLUT-1 isoform located at the blood-brain barrier is essential for glucose uptake, and a hyperthyroidism-induced decrease in its expression might explain the cortical hypometabolism noted in patients with thyrotoxicosis.

There were 2 unexpected findings in our study. One was that the emotional distress–associated regions were not completely consistent with hyperactive regions before treatment. This finding might be due to the fact that global cerebral metabolism is affected by multiple factors, so a single correlation analysis could not reflect the changes of the whole brain. The other finding was that treatment with methimazole improved activity in the limbic system and the frontal and temporal lobes and the association between improved symptoms and activity changes when the euthyroid state is reached. Thyroid hormones are active in modulating metabolic function in the adult brain, and abnormal cerebral metabolism might be one of the possible mechanisms in mood disorder. The behavioral symptoms and subjective disturbances that many patients with hyperthyroidism have may be an integral part of the symptoms of the illness, just as perspiration and heat intolerance are. Our findings indicate that the mood changes caused by Graves disease respond favorably to treatment with regard to clinical symptoms and PET findings, and functional imaging may contribute to elucidating the potentially different background of psychiatric complications.

Acknowledgments
We sincerely thank the patients who participated.

References

Table 5: Associations between the relative brain activity and thyroid hormone levels as well as emotional symptoms of patients

<table>
<thead>
<tr>
<th>Region</th>
<th>FT3 Value</th>
<th>FT3 P Value</th>
<th>FT4 Value</th>
<th>FT4 P Value</th>
<th>SAS Value</th>
<th>SAS P Value</th>
<th>SDS Value</th>
<th>SDS P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Parahippocampal gyrus (BA 35) Left</td>
<td>0.820</td>
<td>.009*</td>
<td>-0.069</td>
<td>.787</td>
<td>0.347</td>
<td>.065</td>
<td>0.820</td>
<td>.698</td>
</tr>
<tr>
<td>Parahippocampal gyrus (BA 28) Left</td>
<td>0.994</td>
<td>.014b</td>
<td>-0.519</td>
<td>.019b</td>
<td>-0.222</td>
<td>.310</td>
<td>0.712</td>
<td>.055</td>
</tr>
<tr>
<td>Superior frontal gyrus (BA 6) Right</td>
<td>-0.359</td>
<td>.007b</td>
<td>1.573</td>
<td>&lt;.001b</td>
<td>0.239</td>
<td>.013b</td>
<td>-0.505</td>
<td>.003b</td>
</tr>
<tr>
<td>Fusiform gyrus (BA 37) Left</td>
<td>-0.111</td>
<td>.456</td>
<td>0.014</td>
<td>.012b</td>
<td>0.539</td>
<td>.007b</td>
<td>-0.236</td>
<td>.189</td>
</tr>
</tbody>
</table>

a Absolute values of β value represent the effect of independent variables on the dependent variable in a multiple regression analysis.
b Survived at P < .05.