Effect of triiodothyronine (T3) augmentation of acute milnacipran administration on monoamine levels: an in vivo microdialysis study in rats

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Background: Up to 30% of depressed patients are partially or totally resistant to antidepressant therapy. The administration of triiodothyronine (T3) to antidepressant nonresponders can be an effective augmentation strategy, although the mechanism is not fully understood.

Methods: In vivo microdialysis was used to examine the effect of T3 augmentation of the antidepressant, milnacipran. Basal extracellular serotonin, norepinephrine, and dopamine levels were measured before and after acute milnacipran administration in the medial prefrontal cortex and amygdala of rats which had received subchronic (7 days) T3 treatment or control saline.

Results: Subchronic administration of T3 at 0.1 mg/kg significantly increased basal extracellular levels of serotonin in the medial prefrontal cortex, but not in the amygdala. In contrast, subchronic administration of T3 at 0.2 mg/kg did not alter basal extracellular serotonin levels in the medial prefrontal cortex. Basal extracellular levels of norepinephrine and dopamine were not modified by either dose of T3 in either region. Acute administration of milnacipran, a serotonin-norepinephrine reuptake inhibitor, to control animals resulted in a significant increase of extracellular levels of serotonin, norepinephrine, and dopamine. When administered to animals treated subchronically with T3 at 0.1 mg/kg, milnacipran produced an additional increase in extracellular serotonin levels but not in levels of norepinephrine or dopamine in the medial prefrontal cortex of rats.

Conclusion: These results suggest that the mechanism of the augmentation effect of milnacipran by T3 administration occurs via enhancement of serotonergic neurotransmission, but not through noradrenergic or dopaminergic neurotransmission.

Keywords: depression, in vivo microdialysis, milnacipran, monoamines, triiodothyronine

Introduction

Major depression is considered by the World Health Organization to be the fourth most disabling disorder worldwide. Despite successive generations of antidepressant drugs, on average, 30%–45% of depressed patients fail to respond adequately to their initial antidepressant treatment and only 25%–35% achieve full symptom remission. It has been estimated that 10%–30% of depressed patients are partially or totally resistant to antidepressant therapy.

Among the treatment strategies proposed for nonresponders or partial responders to antidepressants, adjunctive administration of triiodothyronine (T3) has been shown to be an effective augmentation therapy. STAR*D research has shown that the augmentation effects of T3 in patients with refractory depression taking selective serotonin reuptake inhibitors, selective serotonin and norepinephrine reuptake inhibitors, or bupropion gave similar remission rates in all cases. However, a recent meta-analysis of four trials where T3 was initiated simultaneously with an selective

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serotonin reuptake inhibitor failed to show any superiority for the augmentation therapy.10

The mechanism of $T_3$ augmentation is still unclear. One week of $T_3$ administration has been shown to result in desensitization of 5-HT$_{1A}$ receptors and an increase in extracellular serotonin levels in the frontal cortex of rats.11,12 Coadministration of imipramine and $T_3$ results in desensitization of 5-HT$_{2c}$ receptors.13 Thus, the effect of $T_3$ augmentation of antidepressant effects has been assumed to involve serotonin neurotransmission.14 However, to our knowledge, there has been no study investigating extracellular levels of serotonin, norepinephrine, and dopamine simultaneously after $T_3$ treatment. A role of the norepinephrine or dopamine neurotransmitter cannot thus be ruled out.

Previous mechanistic studies have investigated the augmentation effects of $T_3$ with tricyclic antidepressants. However, tricyclic antidepressants are not ideal for clarification of the augmentative effects of $T_3$ because these agents block several receptors in addition to their effects on monoamine transporters.15 Therefore, we chose to study the effect of the augmentation effects of $T_3$ on the antidepressant, milnacipran, which inhibits reuptake of serotonin and norepinephrine with similar efficacy but without any receptor interactions.16–18 A recent review19 has suggested hyperactivity of the amygdala and hypoactivity of the prefrontal cortex in depressed patients.

In this study we used in vivo microdialysis in the medial prefrontal cortex and amygdala to measure the basal extracellular levels of serotonin, norepinephrine, and dopamine and levels after acute administration of milnacipran in rats which had received subchronic treatment (for one week) with $T_3$ or with control saline.

Materials and methods

Animals

Male Sprague-Dawley rats, obtained from the Shizuoka Laboratory Animal Centre (Shizuoka, Japan), were housed in groups of four and maintained on a 12-hour light-dark cycle (light phase 6.30 am to 6.30 pm) in a temperature-controlled environment ($22 ^\circ C \pm 1 ^\circ C$) with free access to food and water. Experiments began after a 10-day period of acclimatization. The Hokkaido University School of Medicine Animal Care and Use Committee approved all procedures, which complied with the Guide for the Care and Use of Laboratory Animals, Hokkaido University School of Medicine.

Drugs

3,3',5-Triiodo-L-thyronine sodium salt ($T_3$); O-(4-hydroxy-3-iodophenyl)-3,5-diiodo-L-tyrosine sodium salt; Sigma, St Louis, MO) was dissolved in 1 M NaOH with saline, which was used to dilute the solution to the required concentrations (0.1 and 0.2 mg/mL). The final concentration of NaOH was 5 mM for each dose of $T_3$; 5 mM NaOH was used as the vehicle for $T_3$. Milnacipran (Asahi Kasei Corporation, Shizuoka, Japan) was dissolved in distilled water, which was used as the vehicle for milnacipran, to achieve a final concentration of 10 mg/mL.

Drug administration

The animals were divided into two groups. Control group rats were given the vehicle subcutaneously once daily in the morning for 7 days and the $T_3$ group rats were given $T_3$ 0.1 mg/kg or 0.2 mg/kg subcutaneously once daily in the morning for 7 days. On day 7, 2 hours after the last subcutaneous injection, the animals were implanted stereotactically in the medial prefrontal cortex or the amygdala with guide cannulae and microdialysis probes (see below). The rats were administered milnacipran 10 mg/kg intraperitoneally or saline 180 minutes after the first dialysate samples were collected. The 0.1 mg/kg and 0.2 mg/kg doses of $T_3$ were chosen because administration of these doses for one week has been shown to increase extracellular serotonin levels in the frontal cortex of rats.11

Microdialysis procedures

The experiments were performed according to a procedure described previously.20 Under anesthesia with pentobarbital (30 mg/kg intraperitoneally), AG-4 and AG-8 guide cannulae (Eicom Corporation, Kyoto, Japan) were stereotactically implanted into rats at the surface of the medial prefrontal cortex or amygdala at the following coordinates relative to bregma from the stereotaxic atlas of Paxinos and Watson:21 A + 3.2, ML + 0.8, DV +1.0 mm; A – 2.8, ML + 5.0, + DV 7.4 mm, respectively. Dialysis probes with a 0.22 mm outer diameter (A–I–4–03, A–I–4–8–02; Eicom Corporation) were then inserted into the guide cannulae so that 3.0 mm and 2.0 mm of the probes were exposed, respectively, in the medial prefrontal cortex and amygdala tissues. Rats were housed individually after the operation. Microdialysis was performed the next day in the freely moving rats. Perfusion was started 20 hours after surgery using artificial cerebrospinal fluid (145 mM NaCl, 3.0 mM KCl, 1.3 mM CaCl$_2$, 1.0 mM MgCl$_2$) at a flow rate of 2 µL per minute. Following an initial perfusion for 2 hours, dialysate samples were collected in vials containing 50 µL of 50 mM acetic acid every 20 minutes for 480 minutes. The recovery rates of serotonin, norepinephrine, and dopamine in vitro at 30°C were as follows: medial prefrontal cortex, 14.5%, 12.9%,
Effect of subchronic treatment with T<sub>3</sub> on basal extracellular concentrations of serotonin, norepinephrine, and dopamine in the medial prefrontal cortex and amygdala

In the medial prefrontal cortex, basal extracellular serotonin concentrations in the T<sub>3</sub> 0.1 mg/kg group were significantly higher than those of the control group (P < 0.05, Table 1). No significant differences in basal extracellular concentrations of norepinephrine or dopamine were found between the control and T<sub>3</sub> 0.1 mg/kg groups. No significant differences were found between the control and T<sub>3</sub> 0.2 mg/kg groups in basal extracellular levels of any of the monoamines. In the amygdala, no significant difference was found in basal extracellular serotonin, norepinephrine or dopamine concentrations between the control and the T<sub>3</sub> 0.1 mg/kg groups.

Table 1 Effect of subchronic treatment with T<sub>3</sub> (for 7 days) on baseline concentrations of serotonin, norepinephrine, and dopamine in the medial prefrontal cortex and the amygdala

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>T&lt;sub&gt;3&lt;/sub&gt; 0.1 mg/kg</th>
<th>T&lt;sub&gt;3&lt;/sub&gt; 0.2 mg/kg</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n = 7</td>
<td>n = 6</td>
<td>n = 6</td>
</tr>
<tr>
<td>Medial prefrontal cortex T&lt;sub&gt;3&lt;/sub&gt; 0.1 mg/kg</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serotonin</td>
<td>0.793 ± 0.084</td>
<td>1.349 ± 0.234*</td>
<td></td>
</tr>
<tr>
<td>Norepinephrine</td>
<td>1.171 ± 0.107</td>
<td>1.060 ± 0.150</td>
<td></td>
</tr>
<tr>
<td>Dopamine</td>
<td>0.246 ± 0.027</td>
<td>0.220 ± 0.024</td>
<td></td>
</tr>
<tr>
<td>Medial prefrontal cortex T&lt;sub&gt;3&lt;/sub&gt; 0.2 mg/kg</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serotonin</td>
<td>1.08 ± 0.22</td>
<td>1.12 ± 0.37</td>
<td></td>
</tr>
<tr>
<td>Norepinephrine</td>
<td>0.880 ± 0.164</td>
<td>0.827 ± 0.171</td>
<td></td>
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<tr>
<td>Dopamine</td>
<td>0.242 ± 0.039</td>
<td>0.237 ± 0.037</td>
<td></td>
</tr>
<tr>
<td>Amygdala T&lt;sub&gt;3&lt;/sub&gt; 0.1 mg/kg</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Serotonin</td>
<td>0.259 ± 0.024</td>
<td>0.342 ± 0.035</td>
<td></td>
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<tr>
<td>Norepinephrine</td>
<td>0.561 ± 0.072</td>
<td>0.760 ± 0.139</td>
<td></td>
</tr>
<tr>
<td>Dopamine</td>
<td>0.504 ± 0.039</td>
<td>0.606 ± 0.121</td>
<td></td>
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</tbody>
</table>

*P < 0.05 compared with control group.

Notes: Values represent mean ± standard error of the mean (pg per 20-minute fraction); n = number of animals in each group; data are calculated from Figures 1–3.
Effect of acute milnacipran administration on extracellular concentrations of serotonin, norepinephrine, and dopamine in the medial prefrontal cortex and amygdala after control and subchronic T₃ treatments

Acute intraperitoneal administration of milnacipran 10 mg/kg to saline-treated (control) animals resulted in an increase in extracellular concentrations of serotonin, norepinephrine, and dopamine in the medial prefrontal cortex (Figures 1 and 2) and in the amygdala (Figure 3).

Following intraperitoneal administration of milnacipran 10 mg/kg to rats treated with subchronic T₃, two-way ANOVA with repeated measures (0–300 minutes) of extracellular serotonin concentrations in the medial prefrontal cortex (Figure 1A) showed significant interaction between T₃ treatment and time after milnacipran administration [F(15,165) = 1.992, P = 0.0185] and a significant main effect of time [F(15,165) = 13.111, P < 0.0001]. However, the main effect of T₃ treatment after milnacipran administration was not significant [F(1,11) = 4.758, P = 0.0518].

The T₃ 0.1 mg/kg group showed significantly higher concentrations of extracellular serotonin than those of the control group at two time points, ie, at 40 and 100 minutes (P < 0.05, Figure 1).

Notes: Values represent the mean ± standard error of the mean (pg per 20-minute fraction). Black arrows indicate milnacipran administration. [n = 6 (control group), n = 6 (T₃ 0.2 mg/kg group)].
For the extracellular dopamine concentrations in the medial prefrontal cortex (Figure 1C), two-way ANOVA with repeated measures (0–300 minutes) showed a significant main effect of time [F(15,165) = 8.449, P < 0.0001]. There was neither a significant main effect of subchronic T₃ 0.1 mg/kg treatment and time after milnacipran administration [F(15,165) = 1.107, P = 0.3537].

For the extracellular dopamine concentrations in the amygdala (Figure 3A), two-way ANOVA with repeated measures (0–300 minutes) showed a significant main effect of time [F(15,165) = 15.234, P < 0.0001]. There was neither a significant main effect of T₃ 0.1 mg/kg treatment after milnacipran administration [F(1,11) = 0.25, P = 0.8769] nor an interaction between subchronic T₃ 0.1 mg/kg treatment and time after milnacipran administration [F(15,165) = 1.107, P = 0.3537].

In rats treated with subchronic T₃ 0.2 mg/kg (Figure 2), two-way ANOVA with repeated measures (0–300 minutes) showed a significant main effect of time for the extracellular serotonin, norepinephrine, and dopamine concentrations [F(15,150) = 15.068, P < 0.0001; F(15,150) = 12.058, P < 0.0001; F(15,150) = 4.241, P < 0.0001, respectively]. However, there was neither a main effect of subchronic T₃ 0.2 mg/kg treatment after milnacipran administration [F(1,10) = 0.036, P = 0.853; F(1,10) = 0.206, P = 0.660; F(1,10) = 0.325, P = 0.581, respectively] nor an interaction between subchronic T₃ 0.2 mg/kg treatment and time after milnacipran administration [F(15,150) = 0.463, P = 0.955; F(15,150) = 0.664, P = 0.816, respectively].

In the amygdala (Figure 3), for extracellular serotonin and norepinephrine concentrations, two-way ANOVA with repeated measures (0–300 minutes) showed a significant main effect of time [F(15,75) = 1.843, P = 0.044; F(15,75) = 5.197, P < 0.0001, respectively]. There was neither a main effect of subchronic T₃ 0.1 mg/kg treatment after milnacipran administration [F(1,5) = 0.0008, P = 0.993; F(1,5) = 0.848, P = 0.399, respectively] nor an interaction between subchronic T₃ 0.1 mg/kg treatment and time after milnacipran administration [F(15,75) = 0.547, P = 0.905; F(15,75) = 0.342, P = 0.988, respectively].

For the extracellular dopamine concentrations in the amygdala (Figure 3C), there was neither a main effect of subchronic T₃ 0.1 mg/kg treatment after milnacipran administration [F(1,5) = 0.165, P = 0.701] nor an interaction between subchronic T₃ 0.1 mg/kg treatment and time after milnacipran administration [F(15,75) = 1.325, P = 0.209]. The main effect of time was not significant [F(15,75) = 1.217, P = 0.279].

The area under the curve (0–180 minutes) for extracellular serotonin in the subchronic T₃ 0.1 mg/kg group was significantly higher than that of the control group (527.4 ± 101.1 and 282.9 ± 46.5 pg * minutes, respectively, P < 0.05). The area under the curve (0–180 minutes) for extracellular norepinephrine and dopamine were not significantly different between the T₃ and control groups.

For the extracellular norepinephrine concentrations in the medial prefrontal cortex (Figure 1B), two-way ANOVA with repeated measures (0–300 minutes) showed a significant main effect of time [F(15,165) = 15.234, P < 0.0001]. The area under the curve (0–180 minutes) for extracellular serotonin in the subchronic T₃ 0.1 mg/kg treatment group showed a significant main effect of time [F(15,165) = 15.234, P < 0.0001]. However, there was neither a main effect of subchronic T₃ 0.1 mg/kg treatment after milnacipran administration [F(1,11) = 0.25, P = 0.8769] nor an interaction between subchronic T₃ 0.1 mg/kg treatment and time after milnacipran administration [F(15,165) = 1.107, P = 0.3537].

**Figure 3** Effect of acute intraperitoneal administration of milnacipran 10 mg/kg on extracellular concentrations of serotonin (A), norepinephrine (B), and dopamine (C) in the amygdala after subchronic treatment with T₃ 0.1 mg/kg.

**Notes:** Values represent the mean ± standard error of the mean (pg per 20-minute fraction). Black arrows indicate milnacipran administration. [n = 3 (control group), n = 4 (T₃ 0.1 mg/kg group)].
Discussion
Subchronic (7 days) administration of T₃ 0.1 mg/kg significantly increased basal extracellular levels of serotonin in the medial prefrontal cortex (Table 1). This finding is consistent with previous studies¹¹,¹² which have shown that T₃ administered daily for 7 days over the range 0.02–0.2 mg/kg increases basal extracellular levels of serotonin in the rat cortex, in a dose-dependent manner. Subchronic administration of T₃ at 0.1 mg/kg had no effect on basal extracellular levels of norepinephrine or dopamine (Table 1). Although several studies have investigated the role of serotonin in the mechanism of T₃ augmentation,¹¹–¹³ to our knowledge, the present study is the first to examine extracellular serotonin, norepinephrine, and dopamine levels simultaneously after T₃ treatment. It is currently believed that the effect of T₃ on serotonin neurotransmission is integral to the mechanism of T₃ augmentation of antidepressant effects.¹⁴ The results of the present study strongly support this notion, and the absence of effect on the levels of norepinephrine or dopamine makes it unlikely that these neurotransmitters are involved.

The present study found no effects on basal extracellular serotonin levels in the amygdala (Table 1) after subchronic administration of T₃ 0.1 mg/kg, in contrast with the increase found in the medial prefrontal cortex (Table 1). A previous study¹¹ has reported that subchronic administration of T₃ at 0.1 mg/kg increased basal extracellular serotonin levels in the frontal cortex of rats, but not in the hippocampus. The effects of T₃ on increases in extracellular serotonin thus appear to be sensitive, not only to dose but to the brain region studied. A more detailed study of brain regions sensitive to the effects of T₃ administration might be helpful in clarifying the mechanism of action of T₃ augmentation.

Results of the present study showed that, in contrast with the effect of subchronic administration of T₃ 0.1 mg/kg, similar administration of T₃ 0.2 mg/kg did not alter basal extracellular serotonin levels (Table 1). This differs from previous findings.¹² However, higher doses of T₃ 0.5 mg/kg have been shown to have no effect on extracellular serotonin levels in the frontal cortex of rats,¹² suggesting a bell-shaped relationship between the dose of T₃ and the increase in serotonin levels. The nature of the bell-shaped curve has not been studied in detail, but it is possible that experimental differences between studies (strain of rats, for example) could result in a leftward shift of the bell-shaped curve, offering a possible explanation of the different results obtained.

As we have reported previously,¹⁶,²² acute administration of the antidepressant, milnacipran, to rats receiving saline resulted in an increase in basal extracellular concentrations of serotonin, norepinephrine, and dopamine in the medial prefrontal cortex (Figures 1 and 2) and amygdala (Figure 3). In rats treated subchronically with T₃ 0.1 mg/kg, acute administration of milnacipran resulted in an additional increase in extracellular serotonin levels, but not in norepinephrine or dopamine levels. This suggests that the augmentation effect of T₃ administration in a selective serotonin and norepinephrine reuptake inhibitor, such as milnacipran, occurs through enhancement of serotonergic neurotransmission, but not through noradrenergic and dopaminergic neurotransmission.

The present study suffers from several weaknesses. A one-week administration of T₃ was used since this is the duration typically used in similar animal studies.¹¹,¹³,¹⁴ However, a recent study reported that coadministration of T₃ and the selective serotonin reuptake inhibitor, fluoxetine, for 3 weeks caused greater increases in neurogenesis in the hippocampus than fluoxetine administration alone.¹⁴ The effects of one week of administration of T₃ may thus not be optimal for extrapolating to the effects occurring with long-term treatment as used clinically.

In addition, in the present study, the effect of milnacipran was studied acutely. Subchronic administration of milnacipran to rats has been shown,²⁵ however, to enhance noradrenergic transmission beyond that achieved with acute administration while no such effect is seen on serotonergic or dopaminergic transmission. Thus, it may be interesting to study longer durations of administration of both T₃ and milnacipran in future studies.

Another weakness of the present study is that it is exclusively neurochemical. Simultaneous behavioral studies could possibly give a “functional window” for studying the phenomenon. Unfortunately, the constraints of in vivo microdialysis are not compatible with behavioral paradigms.

In conclusion, subchronic administration of T₃ 0.1 mg/kg significantly increased basal extracellular levels of serotonin in the medial prefrontal cortex, but not in the amygdala. Subchronic administration of T₃ at 0.2 mg/kg showed no effect on extracellular levels of serotonin in the medial prefrontal cortex, suggesting a bell-shaped dose-effect relationship. Acute administration of milnacipran, which in control animals significantly increased levels of serotonin, norepinephrine, and dopamine, caused an additional increase in extracellular serotonin levels, but not in norepinephrine or dopamine levels, in the medial prefrontal cortex in rats treated subchronically with T₃ 0.1 mg/kg. This suggests that the augmentation effect of T₃ administration on selective serotonin and norepinephrine reuptake inhibitors, such as
milnacipran, occurs via enhancement of serotonergic neurotransmission, but not through noradrenergic and dopaminergic neurotransmission.

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Disclosure
The authors report no conflicts of interest in this work.

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