BRIGHT ARTIFICIAL LIGHT PRODUCES SUBSENSITIVITY TO CLONIDINE

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(Received in final form December 1, 1987)

Summary

The authors used a thermoregulation paradigm to test the hypothesis that chronic treatment with bright artificial light produces subsensitivity to the hypothermic effects of clonidine, an α₂-agonist. One week of treatment produced blunting of the hypothermic response to clonidine (p < 0.00001). These findings are consistent with previous reports that somatic treatments for depression produce subsensitivity of the α₂-receptor.

Seasonal Affective Disorder (SAD) is a syndrome characterized by recurrent depressions which occur annually (1). This syndrome responds to daily treatment with 2 - 6 hours of bright artificial light (1-5). A mechanism accounting for the efficacy of this treatment has not been identified.

Deficient noradrenergic neurotransmission is implicated in the pathophysiology of endogenous depression (6,7). This could be due to an increased density or supersensitivity of α₂-receptors (8-11) which, when activated, decrease the release of norepinephrine (12,13).

Materials and Methods

We used a thermoregulation paradigm to demonstrate that chronic treatment with antidepressants produces supersensitivity to the hypothermic effects of oxotremorine (14,15). We now use this paradigm to verify that bright artificial light produces subsensitivity to the hypothermic effects of clonidine (16-19). We also demonstrate that multiple injections of clonidine do not produce subsensitivity to subsequent injections of itself.

The dependent variable in the experiments reported here was change in core temperature in response to clonidine HCl, an α₂-agonist. These receptors are predominantly but not exclusively of presynaptic origin. Core temperature was measured using a telemetric thermosensor, the model VM Mini-Mitter (Mini-Mitter Co., Sun River, OR). These devices emit radio waves at a
frequency detectable using an AM receiver. Information regarding the validity of this method is available elsewhere (20).

Full-spectrum bright artificial light (11,500 lux) was emitted from a bank of eight 122 cm long Vitalight tubes suspended 50 cm above the animals. The animals were exposed to the light for 24 hours a day for 7 consecutive days.

The light unit used (Duro Test Corp., Bergen, NJ, Model 5599) provides full spectrum light of the variety used to treat SAD (1-5). This unit is used to treat SAD (2). Temperature under the lights was 23 - 23.5°C.

Mini-Mitters were implanted into the peritoneal cavities of 10 adult, male Sprague-Dawley rats (mean weight ± SEM = 391.8 ± 14.2 g). The animals were allowed five days to recover before beginning experimentation. Experiment 1 involved the measurement of the hypothermic response to clonidine HCl, 0.10 mg/kg ip, prior to exposure to bright artificial light (i.e., at baseline), following one week of light exposure and one week after the withdrawal of phototherapy. Core temperature was measured immediately prior to the injection of clonidine and every 10 minutes thereafter for 120 minutes. The animals were subjected to the standard 12 hour light/dark cycle during the baseline and withdrawal phases. Experiment 2 involved the exposure of animals (mean weight ± SEM = 268.6 ± 9.9 g) to standard fluorescent light at an intensity of 300 lux 24 hours a day for 7 days. This was designed to show that changes in circadian rhythms due to light exposure per se do not account for our findings.

Experiment 3 involved the measurement of core temperature in response to clonidine HCl, 0.40 mg/kg ip, at baseline and one and two weeks later in 7 adult, male Sprague-Dawley rats (mean weight ± SEM = 243.6 ± 5.8 g) not exposed to bright, artificial light. These animals were subjected to the normal ambient conditions in our vivarium. This involves exposure to standard fluorescent light at an intensity of 300 lux between 0600 and 1800. The objective of this experiment was to demonstrate that multiple injections of clonidine do not alter the thermic responsiveness to itself. The high dose of clonidine provided a stringent test of the hypothesis that clonidine does not produce subsensitivity to itself when administered weekly.

Data were assessed for significance using student's paired t-test. All measures of variance in the text refer to the standard error of the mean (SEM).

Results

Experiment 1: Mean core temperature at baseline was 37.2 ± 0.31°C. Table I summarizes the mean thermic response over the 12 time points of all 10 animals. Nine (9) of the 10 animals exhibited blunting of the hypothermic response to clonidine at α < 0.04. Further, the mean thermic response of the sample, which was -1.22 ± 0.15°C prior to light exposure and -0.03 ± 0.10°C after chronic light treatment, was significantly blunted (p < 0.00009, t = 6.74, df = 9). One week after the discontinuation of light treatment, the mean hypothermic response of the sample increased to -1.16 ± 0.14°C. This differed significantly from the response during light treatment (p < 0.0003, t = 5.80, df = 9) but did not differ from baseline (p < 0.80, t = 0.24, df = 9).
TABLE I

<table>
<thead>
<tr>
<th>Animal</th>
<th>Mean Hypothermic Response at Baseline</th>
<th>Mean Hypothermic Response After 1 Week of Light</th>
<th>Mean Hypothermic Response After 1 Week Light Withdrawn</th>
<th>Baseline vs 1 Week Light</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>-0.55 ± 0.09</td>
<td>0.35 ± 0.04</td>
<td>-1.92 ± 0.12</td>
<td>0.000002</td>
</tr>
<tr>
<td>2</td>
<td>-1.81 ± 0.18</td>
<td>0.07 ± 0.04</td>
<td>-1.15 ± 0.11</td>
<td>0.000001</td>
</tr>
<tr>
<td>3</td>
<td>-1.34 ± 0.19</td>
<td>-0.26 ± 0.05</td>
<td>-1.50 ± 0.22</td>
<td>0.003</td>
</tr>
<tr>
<td>4</td>
<td>-1.72 ± 0.19</td>
<td>0.21 ± 0.04</td>
<td>-1.30 ± 0.20</td>
<td>0.000004</td>
</tr>
<tr>
<td>5</td>
<td>-1.10 ± 0.13</td>
<td>0.39 ± 0.06</td>
<td>-1.40 ± 0.12</td>
<td>0.000001</td>
</tr>
<tr>
<td>6</td>
<td>-1.64 ± 0.16</td>
<td>0.27 ± 0.06</td>
<td>-0.62 ± 0.15</td>
<td>0.00002</td>
</tr>
<tr>
<td>7</td>
<td>-1.62 ± 0.19</td>
<td>0.00 ± 0.05</td>
<td>-0.43 ± 0.12</td>
<td>0.000002</td>
</tr>
<tr>
<td>8</td>
<td>-1.15 ± 0.15</td>
<td>0.47 ± 0.04</td>
<td>-1.05 ± 0.16</td>
<td>0.000001</td>
</tr>
<tr>
<td>9</td>
<td>-0.55 ± 0.12</td>
<td>-0.29 ± 0.07</td>
<td>-0.96 ± 0.11</td>
<td>n.s.</td>
</tr>
<tr>
<td>10</td>
<td>-0.69 ± 0.08</td>
<td>-0.41 ± 0.11</td>
<td>-1.16 ± 0.17</td>
<td>0.004</td>
</tr>
</tbody>
</table>

Student's paired t-tests of hypothermic responses of individual animals (n = 10) to clonidine (0.10 mg/kg ip) at baseline (pretreatment), following 7 days of 11,500 lux light treatment, and 7 days following withdrawal of light. Core temperature was measured every 10 minutes for 120 minutes following the injection of clonidine.

Experiment 2: Mean core temperature at baseline was 37.8 ± 0.35°C. The mean thermic response of the sample before and after treatment with 300 lux light was -1.4 ± 0.28°C and -2.2 ± 0.23°C afterwards (p < 0.02, t = 2.94, df = 9).

Experiment 3: Mean core temperature at baseline was 37.6 ± 0.18°C. The baseline thermic response to clonidine was -3.49 ± 0.28°C (p < 0.00005, t = 12.48, df = 6). Rechallenge with clonidine after one and two weeks revealed changes in the thermic response of +0.56 ± 0.46°C (p > 0.30, t = 1.31, df = 5) and +0.28 ± 0.26°C (p > 0.30, t = 1.1, df = 6), respectively.

Discussion

Chronic treatment with bright artificial light (11,500 lux) potently subsensitized the α2-receptor as inferred from measurement of clonidine-induced hypothermia. However, treatment with standard fluorescent light at an intensity of 300 lux had the opposite effect. This suggests that light intensity is a critical variable. Our observations argue against the...
hypothesis that the effects of bright light result from an alteration of circadian rhythms due to constant light exposure. The intensity of light in the rats' cages in our vivarium is 300 lux. The animals demonstrate clear-cut circadian changes in motor activity and core temperature in response to turning the lights on or off. Thus, we propose that the results presented here are due to the effects of high intensity light rather than to light per se. The observation that the control sample exhibited a significant increase in its hypothermic response to clonidine of 0.8 ± 0.28°C (p < 0.02) was unexpected. Certainly, significant blunting of the hypothermic response to clonidine following chronic exposure to 300 lux light would suggest that mere exposure to light or alterations in circadian rhythms affect subsensitivity of the mechanism mediating clonidine induced hypothermia. However, the results are opposite this. Bright light but not standard room light produced subsensitivity. Thus, the data are consistent with the hypothesis that bright artificial light affects subsensitivity of presynaptic α2 receptors.

Our finding is consistent with reports that tricyclic antidepressants (21,22), monoamine oxidase inhibitors (23), electroconvulsive shock (24), and lithium carbonate (25) (treatments for depression) all subsensitize the α2-receptor. Our data do not prove that bright artificial light subsensitizes the presynaptic α2-receptor but are consistent with this possibility. Thus, the capacity to subsensitize α2-receptors is a property common to all treatments for depressive disorders which have been assessed for it.

Patients with SAD are typically treated with full-spectrum light at an intensity of 2500 lux. There are advantages to using higher intensities of bright artificial light in preliminary studies. Our objective was to determine whether bright light, as opposed to standard room lighting, produces subsensitivity to the hypothermic effects of clonidine. Use of a high "dose" decreases the probability of accepting a false null hypothesis ("bright artificial light does not produce subsensitivity to the thermic effects of clonidine"). Now that it has been demonstrated that treatment with full-spectrum light at an intensity of 11,500 lux results in decreased sensitivity to clonidine, and possibly subsensitivity of presynaptic α2-receptors, it would be reasonable to evaluate lower "doses" and the effects of administering light for circumscribed periods of time each day. These studies are now under way.

The light unit we used delivers light at an intensity of 2500 lux at a distance of 122 cm from a patient's face. However, to deliver light at an intensity of 11,500 lux, the unit was suspended 50 cm above the animals. When placed this distance from the face of a patient, there is no discomfort. Thus, it would be feasible to change the current treatment protocols should studies suggest an intensity of light in excess of 2500 lux might be superior.

Acknowledgements

Supported in part by MH00553-02 and NIH 2507RR05383-5.
References