Suppression of Human Growth Hormone Secretion by Melatonin and Cyproheptadine

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ABSTRACT Pituitary growth hormone (GH) release in the rat is stimulated via serotoninergic pathways and can be inhibited by treatment with compounds that act as serotonin antagonists, such as cyproheptadine or the pineal gland hormone, melatonin. To investigate a possible role for serotonin in the control of human GH release, the effects of cyproheptadine and melatonin administration on the GH responses of normal male subjects were examined.

The oral administration of cyproheptadine (8-12 mg daily for 5 days) to normal subjects reduced their GH responses to both insulin-induced hypoglycemia and physical exercise to a highly significant extent. Similarly, the mean GH responses of 10 subjects to insulin-induced hypoglycemia were significantly reduced after the prior oral administration of melatonin (1 g).

The data presented show that serotonin antagonism has a similar effect on GH secretion in man to that observed in the rat and provides further evidence for serotoninergic, and possibly pineal, involvement in the control of human GH secretion.

INTRODUCTION

Evidence for brain catecholamine involvement in growth hormone (hGH) secretion in man was first reported by Blackard and Heidingsfelder (1) who showed that the hGH response to insulin-induced hypoglycemia was influenced by α- and β-adrenergic blockade. Further indication that dopamine or norepinephrine may participate in the release of hGH was provided by Boyd, Lebovitz, and Pfeiffer (2) who showed stimulation of hGH secretion in patients with Parkinson's disease who were administered L-dopa. Many groups have since demonstrated stimulation of hGH release in normal, nonobese subjects after L-dopa administration. However, Müller (3) has suggested that results on L-dopa stimulation of hGH secretion should be interpreted with caution because of the possibility that the effect of L-dopa may be due to final activation of serotoninergic receptors.

There is increasing evidence that serotoninergic pathways may have a more direct role in GH secretory mechanisms than catecholaminergic pathways. Imura, Nakai, and Yoshimi (4) recently demonstrated that hGH secretion is stimulated by relatively small doses of the serotonin (5-hydroxytryptamine) precursor 5-hydroxytryptophan (5-HTP), and, furthermore, on the basis of studies on patients with excess serotonin secretion (carcinoid syndrome), Feldman and Lebovitz (5) have suggested that serotonin may be a stimulator of hGH release. Collu, Fraschini, Visconti, and Martini (6) have demonstrated that intraventricular administration of serotonin causes GH release in the rat. Our studies (7–9) have shown that rat GH secretion is stimulated by intraperitoneal administration of 5-HTP and, furthermore, that it is inhibited by serotoninergic blockade and also by the pineal gland hormone, melatonin, which appears to act by competing with serotonin for serotonin receptor sites (9). The GH release observed in humans after the onset of slow-wave sleep (10) has also been suggested to originate via serotoninergic pathways (3, 11).

The present investigation was aimed at determining whether blockade of brain serotoninergic pathways or receptors in humans suppresses the hGH responses to stimulation and, consequently, at providing further evidence for serotoninergic control of GH secretion in man.

After the studies described herein were completed, Bivens, Lebovitz, and Feldman (12) reported that the hGH response to insulin-induced hypoglycemia was significantly suppressed by pretreatment with the serotonin antagonists cyproheptadine and methysergide. The
conclusions presented by these workers provoked immediate comment and controversy (13, 14). The protocol employed by Bivens et al. (12) was similar to that used in one of the experiments presented in the present study, and thus a comparison of the findings is pertinent.

METHODS

These studies were performed on normal, informed non-obese male volunteers, aged 21–28 yr. All tests on the subjects began at 8:30 a.m. after an overnight fast. Blood samples were taken from an indwelling catheter in an antecubital vein. The patency of the catheter was maintained by a slow isotonic saline infusion. Blood samples were collected before stimulation and then every 15 min throughout the studies, and, except during the 20 min exercise regimen, each subject rested on a bed for the duration of each test. Tests on the same subject were carried out at least 1 wk apart.

Study 1. Insulin-induced hypoglycemia control study. All 13 volunteers were subjected to this study. After collection of three basal blood samples, insulin (Novo Industri A/S, Denmark, neutral porcine crystalline; 0.1 U/kg) was given intravenously at time “0” minutes. Blood samples were collected during the 2 h following insulin administration.

Study 2. Insulin-induced hypoglycemia after cyproheptadine administration. Seven of the subjects took part in this study. 5 days before the commencement of a second insulin hypoglycemia test, each subject began a regimen of oral cyproheptadine (Periactin, Merck, Sharp & Dohme, West Point, Pa.). On the 1st and 2nd days 4 mg was taken twice daily and on the 3rd, 4th, and 5th days 4 mg was taken three times daily. On the day of testing (6th day) a final 4 mg was taken 1–2 h before commencement of the test so that the cumulative dose of cyproheptadine for the subjects completing the course was 56 mg. Due to severe side effects of the cyproheptadine experienced by one subject (extreme drowsiness) he was withdrawn from the study before completing the course. An insulin hypoglycemia test was then carried out on the remaining six subjects in an identical manner to that used in the control study.

Study 3. Exercise after cyproheptadine administration. The six subjects from study 2 (above) renewed oral cyproheptadine 2 days after the insulin hypoglycemia test and for 5 days followed an identical regimen to that above (i.e., 8 mg for 2 days, then 12 mg for 3 days). A final tablet (4 mg) was taken 1–2 h before commencement of testing. After collecting basal blood samples the volunteers were subjected to a standardized physical exercise regimen consisting of 20 min pedaling on a bicycle ergometer (Monarch) at a load of 600 kilopondmeters (kpm)/min. The time of starting exercise was taken as 0 min, and blood samples were collected during the following 2 h.

Study 4. Physical exercise control study. After 1 wk without medication the six subjects from study 3 presented for a second exercise study to determine their control responses. Due to difficulties with venous catheterization, one subject was unable to be tested. The remaining subjects were given an identical exercise regimen to that of study 3.

Study 5. Insulin-induced hypoglycemia after oral melatonin. This study group comprised 10 subjects. At 60 and 30 min before administration of insulin each subject was administered a 500-mg capsule of synthetic melatonin (total 1.0 g). Insulin (0.1 U/kg) was administered at time 0 min using the same protocol as described in the control study.

Plasma glucose was estimated by a Technicon AutoAnalyzer (Technicon Instruments Corp., Tarrytown, N. Y.) using the ferricyanide method for reducing sugars. Serum hGH was measured by the radioimmunoassay technique of Molinatti et al. (15), and the results are expressed in microunits of the WHO International Reference Preparation for hGH radioimmunoassay. Serum hGH levels in control and test samples for each individual were estimated in the same assay. Statistical analysis was carried out using Student’s t test. Results are expressed as the mean ± the SEM.

RESULTS

The serum hGH response to insulin-induced hypoglycemia and the effect of cyproheptadine treatment on this response is illustrated in Fig. 1. Following cyproheptadine the mean serum hGH levels are significantly reduced at all times after 30 min. The mean peak level of hGH was earlier in the cyproheptadine study and was reduced from 90±15.0 to 46.4±10.0 µU/ml (P<0.025). Cyproheptadine was found to also affect the hGH responses to physical exercise as illustrated in Fig. 2. While the subjects commented that they experienced greater difficulty in completing the exercise regimen with cyproheptadine treatment, their GH responses were markedly lower. One of the five subjects failed to show a rise in serum GH levels with exercise either on or off cyproheptadine, and his result was not included in the study. The other four subjects showed a normal response to the control exercise regimen, but two of these showed a peak serum response of less than 5 µU/ml with cyproheptadine. The small number of subjects completing this study prevented a high level of significance being obtained at individual sample times, but when the total amount of GH secreted by each subject was esti-
Figure 2 The effect of cyproheptadine on the serum hGH response to physical exercise in four normal subjects. The broken line shows their response after cyproheptadine and the continuous line their control response. Means±SEM are shown. (*) $P<0.05$.

Mitigated by integrating the area under the individual response curves obtained with and without cyproheptadine, there was found to be a highly significant difference between the control and cyproheptadine treatment means for both the exercise and hypoglycemia studies. This is shown in Fig. 3. In the exercise study cyproheptadine treatment resulted in a reduction of approximately 50% in total hGH secreted over the time period covered ($P<0.0125$).

The effect of melatonin administration on the GH response to insulin-induced hypoglycemia in the 10 subjects tested is illustrated in Fig. 4. The GH response of only one subject failed to be suppressed by the melatonin. The mean GH response of all subjects was significantly reduced by 60 min and was reduced significantly at later sampling times. The mean peak hGH level for the 10 subjects was reduced from 94.7±12.8 to 55.7±11.9 μU/ml ($P<0.025$) with melatonin administration. The ability of melatonin to cause a small rapid paradoxical rise in serum GH levels 30 min after administration is seen in Fig. 4 at time 0 where the serum GH level for the melatonin study is significantly ($P<0.05$) higher than that of the control study. Untoward side-effects were not noted after the melatonin administration.

Whereas both the cyproheptadine and melatonin regimens caused a significant alteration in the GH responses to insulin-induced hypoglycemia, neither treatment altered the degree of hypoglycemia achieved. This is shown in Table I.

DISCUSSION

The results of this investigation demonstrate that the oral administration of either the serotonin antagonist, cyproheptadine, or the pineal gland hormone, melatonin, result in a highly significant suppression of the normal hGH release after stimulation. The hGH responses to both insulin-induced hypoglycemia and physical exercise were significantly reduced by cyproheptadine treatment in all parameters measured. With the dose of cyproheptadine employed, the mean hGH responses were reduced by approximately 50%. The inability of the cyproheptadine regimen to suppress completely the GH responses could indicate either that the GH response is only partly due to activation of serotonin receptors or, alternatively, that blockade of brain serotonin recep-
The Degree of Hypoglycemia Induced by Insulin* after Cyproheptadine and Melatonin Administration in Normal Subjects

<table>
<thead>
<tr>
<th>Study</th>
<th>No. of subjects</th>
<th>Fall in plasma glucose, mg/100 ml</th>
<th>mean ± SEM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Insulin alone</td>
<td>6</td>
<td>54.6 ± 3.8</td>
<td></td>
</tr>
<tr>
<td>Cyproheptadine and insulin</td>
<td>6</td>
<td>54.5 ± 3.1</td>
<td></td>
</tr>
<tr>
<td>Insulin alone</td>
<td>10</td>
<td>52.1 ± 3.1</td>
<td></td>
</tr>
<tr>
<td>Melatonin and insulin</td>
<td>10</td>
<td>54.2 ± 2.5</td>
<td></td>
</tr>
</tbody>
</table>

The mean basal fasting plasma glucose for all studies was 88.25 ± 1.10 mg/100 ml and all subjects were within 2 ± SD of this value at commencement of each test.

* 0.1 U/kg.

tors was incomplete at the dose level of cyproheptadine selected. The results presented here showing suppression of hGH secretion by cyproheptadine are in excellent agreement with those presented by Bivens et al. (12) for insulin-induced hypoglycemia. Bivens et al. (12) proposed that the effect of cyproheptadine (and methysergide) was exerted at the level of the hypothalamus. This proposal provoked the suggestion by Essman, Sherman, and Kolodny (13) that these serotonin antagonists were unlikely to be able to exert a central effect, suggesting further that their actions were confined to peripheral receptor-site blockade of serotonin. However, we believe that there is ample evidence demonstrating that serotonin can cross the blood-brain barrier to exert central serotonin antagonism at doses well below the mean lethal dose (LD₅₀) for laboratory animals. Van Riezen (16) has studied the pharmacological effects of cyproheptadine in mice, and in those parameters which are considered to involve serotonin receptors in the central nervous system, namely nialamide-induced hyperactivity and 5-HTP-induced head twitch and tremors, cyproheptadine exerted marked antagonism at doses as low as 1 mg/kg (representing 2.5% of the LD₅₀). In a study of spinal neuronal activity using acute spinal cats, Banna and Anderson (17) have shown that small doses of cyproheptadine and methysergide completely inhibit the 5-HTP-induced increase in monosynaptic spike amplitude and spontaneous motoneuronal discharge. These neuronal responses to the serotonin antagonists were dose related but showed no correlation with their cardiovascular effects. The ability of cyproheptadine to inhibit hyperthermia produced by administration of bacterial pyrogen has been suggested by Feldberg (18) to be caused by its antagonism of serotonin at a hypothalamic level. Furthermore, we have shown (7) that, in the rat, serum GH levels already elevated by pentobarbital administration are rapidly suppressed to base-line levels by cyproheptadine treatment.

Thus, on the basis of this evidence we support the proposal by Bivens et al. (12) that cyproheptadine may exert its effects on hGH secretion by hypothalamic serotonin antagonism. However, we do not discard the possibility that cyproheptadine might act directly on the pituitary gland. Such a mechanism would imply that serotonin mediates the release of hGH from the pituitary, and in this regard it is noteworthy that large concentrations of serotonin have been detected in the bovine pituitary gland and median eminence (19). Moreover, the pituitary gland has been shown to concentrate peripherally administered melatonin (20). The suppression of hGH after serotonin antagonism could thus be a consequence of an action at either or both the hypothalamus or the pituitary gland.

The effect of acute administration of melatonin on the hGH response to insulin-induced hypoglycemia was similar to that observed for cyproheptadine. Melatonin was selected as a possible inhibitor of brain serotonin receptors in this investigation and in studies on rat GH secretion (7–9) because it is an O-methylated derivative of serotonin. We have previously hypothesized (21) and recently shown (22, 23) that O-methylation of dopamine results in derivatives that inhibit the actions of dopamine by competing for its hypothalamic receptor sites. We subsequently proposed (24) that, by analogy with dopamine, O-methylated derivatives of serotonin such as O-methylserotonin (5-MT) and melatonin would antagonize the actions of serotonin at a receptor-site level. The structural relationships between dopamine and serotonin and their O-methyl derivatives are illustrated in Fig. 5.

![Di-O-methyl dopamine (DOMPA)](http://www.jci.org)

**Suppression of hGH Secretion by Melatonin and Cyproheptadine**

![Figure 5](http://www.jci.org) The structures of dopamine, serotonin, and their O-methylated derivatives.
While oral administration of melatonin rapidly results in a small but significant elevation of serum hGH levels (25), the present data show this treatment to exert a significant inhibitory effect on the hGH response following insulin-induced hypoglycemia. We propose that the mean serum GH response after melatonin, shown in Fig. 4 represents a combination of two effects. First there is a stimulatory effect on serum hGH possibly reflecting initial activation of brain serotonin receptors by the melatonin (the early part of this effect can be seen in Fig. 4 where the GH response curve for the insulin-induced hypoglycemia study following melatonin is significantly higher than the control curve at time 0). Second, there is a subsequent blockade by melatonin of serotoninergic pathways normally activated by hypoglycemia. The net effect is significant suppression of the hGH response. The situation suggested is in concert with our proposal (9) that melatonin can act as a general blocker or competitive inhibitor of serotonin at its receptor sites. The findings explain the results of Starr (26) who reported decreased serum hGH levels in patients with sarcoma who were given long-term infusions of melatonin. The results for melatonin could also provide an explanation for the observation of Krieger and Glick (27) showing that blind human subjects do not exhibit slow-wave sleep-induced secretion of hGH. This phenomenon may be due to increased pineal synthesis and secretion of melatonin following blindness. Blinding or constant darkness results in a large increase in the activity of the melatonin-synthesizing enzymes in the rat pineal (28, 29), and recently it was shown (30) that the concentration of a melatonin-like substance increases in the plasma of humans kept in darkness.

The data of the present study do not allow us to say whether the observed effect of melatonin on GH secretion is pharmacological or whether it reflects a physiological mechanism, but it does allow for the possibility of pineal involvement in the control of GH secretion from the pituitary gland.

The results of this investigation show that GH secretion in man is suppressed by serotoninergic blockade and thus provide further evidence for the involvement of serotonin in the neuroendocrine regulation of hGH release.

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REFERENCES


