Familial Gynecomastia with Increased Extraglandular Aromatization of Plasma Carbon_{19-}Steroids

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Abstract

We evaluated a family in which gynecomastia occurred in five males in two generations. In each affected subject, gynecomastia and male sexual maturation began at an early age. The ratio of the concentration of plasma estradiol-17\beta to that of plasma testosterone was elevated in each affected subject. In the three siblings with gynecomastia, the transfer constant of conversion of androstenedione to estrone (i.e., the fraction of plasma androstenedione that was converted to estrone as measured in the urine) was 10 times that of normal persons. The transfer constant of conversion of testosterone to estradiol-17\beta in the one subject studied also was 8–10 times that of normal men, whereas the transfer constants of conversion of estrone to estradiol-17\beta and of estradiol-17\beta to estrone were normal. Despite the elevation in extraglandular aromatase activity, there was a normal response of the hypothalamic-pituitary axis to provocative stimuli. This is the second documentation of gynecomastia that is associated with increased extraglandular aromatase activity, and the first time that the defect was found to be familial with a probable X-linked (or autosomal dominant, sex limited) mode of inheritance.

Introduction

Gynecomastia is a common finding among pubertal boys; usually, it is limited to a nodule of subareolar tissue, and ordinarily the breast enlargement begins during the early stages of the pubertal process and thereafter regresses within 6–18 mo. Levels of plasma testosterone (T), estradiol-17\beta (E_2), serum luteinizing hormone (LH), and follicle-stimulating hormone (FSH) in boys with pubertal gynecomastia are not significantly different from those in unaffected boys (1), although a relative increase in the level of plasma E_2 compared with that of T during certain times of the day has been found (2). In a small percentage of apparently normal pubertal boys, persistent and marked gynecomastia develops even though the plasma levels of T, E_2, prolactin, gonadotropins, and testosterone-estradiol binding globulin in such subjects are normal at the time of evaluation (3). Marked and persistent breast enlargement, however, can sometimes be associated with Klinefelter syndrome or an underlying abnormality in male sexual differentiation (4, 5), including partial gonadal dysgenesis or various forms of androgen resistance. In 1977, Hemsell et al. (6) reported the findings in a prepubertal boy who developed gynecomastia as a consequence of extraordinarily high extraglandular aromatization of plasma androstenedione (A) and T. This was the first report of such a metabolic abnormality, and to the best of our knowledge no other affected subjects have been described since. In that study, a family history was not available since the affected boy was an adopted child (MacDonald, P. C., personal communication). Recently, we evaluated a family in which marked gynecomastia developed in three brothers, a maternal cousin, and a maternal uncle. The gynecomastia was unusual in several respects: (a) it began at an early age; (b) from initial studies of plasma steroids, we found that there was an elevation in the ratio of plasma E_2/T; (c) known causes of gynecomastia were excluded in all subjects; and (d) in studies of extraglandular aromatization of plasma C_{19-}steroids, it was found that the fractional conversion of plasma A to estrone (E_1), [P]^A_{E_1}, was 10 times greater than normal in three subjects studied; and, similarly, the fractional conversion of plasma T to E_2, [P]^T_{E_2}, in the only subject studied, was also 8–10 times that found in normal men. Despite the excessive peripheral aromatase activity, there was evidence of normal gonadotropin release by the pituitary and normal Leydig cell function in the affected subjects.

Case histories

The pedigree of a family in which there was marked and persistent gynecomastia in three siblings, a maternal cousin, and a maternal uncle is presented in Fig. 1. In each of the affected subjects, gynecomastia began at the time of other pubertal changes between 10 and 11 yr of age. Breast enlargement progressed steadily, although there was a slow progression in the maturational events of male puberty. In all five subjects, there was a normal 46,XY karyotype. Thyroid and liver function tests as well as serum levels of prolactin were normal in all five subjects (data not shown). There was one additional adult male (subject II-6) in the second generation who did not have gynecomastia. His two sons were 10 and 5 yr of age at the commencement of these studies.

Subjects III-3. At the time of initial evaluation, the proband (subject III-3) was 11-1/2 yr old. The early medical history of this boy was unremarkable. Pubic hair and breast development began concomitantly when the boy was 10-yr-old. Height was 156.7 cm (75th percentile) and weight was 63 kg (>95th percentile). Bone age was 14-6/12 yr (>2 SD above the mean...
for age; and, a female fat distribution (Fig. 2) and 18 × 14-cm breasts were present. Pubic hair growth and distribution were that of Tanner Stage III (7) and the stretched penile length was 8.5 cm. The testes were 1.5 × 3 cm bilaterally, with normal contour. Values for plasma T and E₂, and for serum LH and FSH are given in Table I. In Table II, we present the changes in pubertal development and plasma steroids and gonadotropin levels over the course of 1 yr. Plasma T remained somewhat low for the stage of sexual development, whereas the ratio of plasma E₂/T was elevated consistently. An X ray of the skull was normal, as was a CT scan of the adrenal glands.

Subject III-1. The older sibling of the proband was 16-5/12-yr-old at the time of initial evaluation. Height was 175.8 cm (50th percentile), weight was 60.8 kg (50th percentile), and the bone age was normal. Except for gynecomastia, there was a normal male body habitus. The breast tissue was 4 × 4 cm on the right and 6 × 6 on the left. Pubic hair distribution was that of Tanner Stage IV and the penile length was 11 cm. The values for plasma steroids and gonadotropins at the time of the initial visit are given in Table I; and, the results of similar measurements over the next 12 mo of observation are given in Table II. Plasma levels of T and A in this subject were normal for the stage of sexual development.

Subject III-4. During the course of the study, the younger sibling of the proband developed unilateral gynecomastia (at 10-8/12 yr of age). Height was 146 cm (75th percentile) and weight was 38 kg (50th percentile). Physical examination was normal except for the presence of 4.5 × 4.5 cm of breast tissue on the left breast. There was a scant amount of pubic hair; the penile length was 8.5 cm and the testes were 1 × 2.5 cm bilaterally. The results of his endocrine evaluation are given in Table I. In this boy, bone age, determined when he was 11-5/12 yr of age, was 13-6/12 yr, a value > 2 SD above the mean for that age.

Subject III-5. A maternal cousin of the proband was 15 yr of age when he was first evaluated by us. Breast tissue, which was noted first at 11 yr of age, increased steadily in size, and bilateral mastectomy was performed at 12 yr of age. When he was 15-yr-old, height was 179 cm (90th percentile) and weight was 61 kg (50th percentile); at this time, there was a normal male body habitus. Pubic hair distribution was that of Tanner Stage V and the stretched penile length was 11 cm. Testes were 2 × 4.5 cm. The plasma level of T, given in Table I, was normal for that age, but the plasma level of E₂ was elevated. The plasma level of DHA-sulfate (221 μg/dl) was normal.

Figure 1. Pedigree of a kindred with familial gynecomastia.

Figure 2. Subject III-3 at age 11-11/12 yr; breast development present since 10 yr of age. Breasts were 18 × 14 cm and pubic hair distribution was Tanner Stage III.
Subject II-8. The maternal uncle of the proband first noted gynecomastia when he was 10 yr of age and the breast tissue was removed surgically at 12 yr of age. When evaluated by us for the first time, he was a moderately obese 29-yr-old man with fat distribution of a woman. Height was 160 cm (<5th percentile) and weight was 77 kg (>90th percentile). Pubic hair distribution was that of Tanner Stage V. The penile length was 11.5 cm and the testes were 2 × 4 cm bilaterally. As given in Table I, the plasma level of E2 in this subject was elevated, although the plasma T was normal.

Methods

Preparation of radiolabeled steroids for intravenous infusion. [4,14C]A (57.4 mCi/mmol), [6,7,13H]E (60 Ci/mmol), [4,14C]T (51.9 mCi/mmol), [6,7,13H]E (53 Ci/mmol), and [4,14C]E (57.0 mCi/mmol) were obtained from New England Nuclear, Boston, MA. All radiolabeled steroids were purified by paper chromatography and passage through a 0.2-μm Millex-FG filter unit (Millipore Corp., Bedford, MA). Before use, an aliquot of the material was removed for bacteriologic culture and determination of pyrogenicity.

Production rate of estrone (PR-E2) and the transfer constant of conversion of [3H]E to [3H]E. These studies were conducted in subjects III-1, III-3, and III-4 at ages 17-2/12, 12-2/12, and 11-1/12 yr, respectively. Subject III-3 was given 12.4 μCi [13C]A and 5.68 μCi [3H]E, dissolved in 112 ml of ethanol (6%) in normal saline, one-sixth of it being infused as a bolus intravenously, and the remainder being delivered as a constant intravenous infusion over 10 h by a Harvard pump. Subject III-1 was given 5.02 μCi [13C]A and 2.71 μCi [3H]E, and subject III-4 was given 6.39 μCi [13C]A and 3.45 μCi [3H]E. In the latter two studies, the radiolabeled steroids were dissolved in 100 ml of ethanol (6%) in normal saline and were infused intravenously over 20 min. In each study, urine was collected for 72 h after commencement of the infusion of radiolabeled steroids. The [3H]E and PR-E2 were computed from the 14H/14C ratio and the specific activity of E2 isolated from urine after hydrolysis of glucuronosides with β-glucuronidase as previously described (6, 8).

Production rate of estradiol (PR-E2) and the transfer constant of conversion of [3H]E to [3H]E. This study was conducted only in subject III-3 at the time when he was 12-2/12 yr old. The radiolabeled tracers, 9.45 μCi [13C]T and 2.21 μCi [3H]E, were dissolved in 55 ml of ethanol (6%) in normal saline. 10 ml of this solution was delivered intravenously as a bolus and 45 ml was delivered intravenously at a constant rate of infusion for 10 h by a Harvard pump. Urine was collected for 72 h. The [3H]E and PR-E2 were computed from the 14H/14C ratio and the specific activity of E2 isolated from urine after hydrolysis of glucuronosides with β-glucuronidase (6, 8).

Transfer constant of conversion of plasma E1 to [3H]E and the transfer constant of conversion of plasma E2 to E1 [3H]E. Subject III-1 was studied when he was 12-3/12 yr old. At this time, 1.57 μCi [3H]E and 8.18 μCi [3H]E were dissolved in 57 ml of ethanol (6%) in normal saline. 10 ml of this solution was injected intravenously as a bolus and 47 ml was infused at a constant rate for 10 h by use of a Harvard pump. Urine was collected for 72 h. The [3H]E and the [3H]E were computed from the 14H/14C ratio of E1 and E2 isolated

### Table I. Hormonal Evaluation of Subjects with Familial Gynecomastia

<table>
<thead>
<tr>
<th>Subjects</th>
<th>Age (yr)</th>
<th>Tanner stage</th>
<th>T (ng/dl)</th>
<th>E2 (ng/dl)</th>
<th>LH (ng/ml)</th>
<th>FSH (ng/ml)</th>
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</thead>
<tbody>
<tr>
<td>Brother III-1</td>
<td>16-5/12</td>
<td>IV</td>
<td>58</td>
<td>3.6</td>
<td>58</td>
<td>314</td>
</tr>
<tr>
<td>Proband III-3</td>
<td>11-11/12</td>
<td>III</td>
<td>32</td>
<td>6.2</td>
<td>35</td>
<td>131</td>
</tr>
<tr>
<td>Brother III-4</td>
<td>10-8/12</td>
<td>II</td>
<td>13</td>
<td>2.8</td>
<td>19</td>
<td>107</td>
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<tr>
<td>Maternal uncle II-8</td>
<td>29</td>
<td>V</td>
<td>506</td>
<td>9.8</td>
<td>38</td>
<td>109</td>
</tr>
<tr>
<td>Maternal cousin III-5</td>
<td>15</td>
<td>V</td>
<td>342</td>
<td>6.2</td>
<td>30</td>
<td>71</td>
</tr>
<tr>
<td>Normal adult men (mean±SD or range)</td>
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<td></td>
<td>575±150</td>
<td>1-6</td>
<td>50±14</td>
<td>187±89</td>
</tr>
</tbody>
</table>

### Table II. Sequential Hormonal Evaluation of Subjects with Familial Gynecomastia

<table>
<thead>
<tr>
<th>Age (yr)</th>
<th>Subject III-1</th>
<th>Subject III-3</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sexual development</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pubic hair (Tanner Stage)</td>
<td>IV</td>
<td>IV</td>
</tr>
<tr>
<td>Penile length (cm)</td>
<td>11</td>
<td>12</td>
</tr>
<tr>
<td>Testicular size (cm)</td>
<td>2 × 4.5</td>
<td>2 × 4.5</td>
</tr>
<tr>
<td>Breast diameter: right/left (cm)</td>
<td>4.0/6.5</td>
<td>5.0/9.5</td>
</tr>
<tr>
<td>Hormonal profile</td>
<td></td>
<td></td>
</tr>
<tr>
<td>T (ng/dl)</td>
<td>58</td>
<td>104</td>
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<tr>
<td>A (ng/dl)</td>
<td>74</td>
<td>66</td>
</tr>
<tr>
<td>E2 (ng/dl)</td>
<td>3.6</td>
<td>3.1</td>
</tr>
<tr>
<td>DHA sulfate (μg/dl)</td>
<td>131</td>
<td>133</td>
</tr>
<tr>
<td>LH (ng/dl)</td>
<td>58</td>
<td>40</td>
</tr>
<tr>
<td>FSH (ng/dl)</td>
<td>341</td>
<td>64</td>
</tr>
</tbody>
</table>

Familial Gynecomastia and Increased Aromatase Activity 1785
from urine after hydrolysis of glucuronosides with β-glucuronidase (6, 8).

Circadian variation in the levels of plasma T and A and of serum LH and FSH. Subjects III-1 and III-3 were admitted to the Pediatric Clinical Research Unit (The Johns Hopkins Hospital and School of Medicine) when they were 17-3/12 and 12-10/12 yr old, respectively. Beginning at 12:00 noon on the day of admission, blood was collected every 3 h for the determination of the levels of plasma T and A, and serum LH and FSH.

Thyrotropin-releasing hormone (TRH) stimulation test and luteinizing hormone-releasing hormone (LHRH) stimulation test. Pituitary function tests were conducted in subjects III-1 and III-3. TRH (500 μg) was given as a bolus intravenously and thereafter blood samples were collected at 0, 15, 30, 45, and 60 min for the determination of thyroid-stimulating hormone (TSH) and prolactin. LHRH (100 μg) was given intravenously as a constant infusion for 3 h. Blood samples were obtained at -60, -30, 0, 30, 60, 90, 120, 150, and 180 min for the determination of LH and FSH.

Human chorionic gonadotropin (hCG) stimulation test. Leydig cell function was evaluated in subjects III-1 and III-3. All blood samples were collected at 0900 a.m. on the appropriate day when the subjects were 17-3/12 and 12-10/12 yr old, respectively. Base-line blood samples were obtained for the determination of plasma levels of T, A, and E2. hCG was given at a dose of 5,000 IU/d intramuscularly for 5 d. On the sixth day after commencing hCG treatment, blood was withdrawn for the determination of the plasma levels of T, A, E2, dehydroepiandrosterone (DHA), DHA-sulfate, 17-hydroxyprogesterone (17-OHP), and progesterone (P).

Routine radioimmunoassays (RIA)s. Assays for serum LH and FSH (9), plasma T and A (10), 17-OHP, P, and DHA (11) were conducted as described previously. Plasma DHA-sulfate was assayed by use of the Radioscience Laboratories (Carson, CA) RIA kit. E2 was determined by Smith-Kline Clinical Laboratories (King of Prussia, PA) as previously described (12). Serum prolactin was determined by use of the Abbott Laboratories (Chicago, IL) prolactin RIA kit.

Results

The PR-E2 and [P]BU[17/4C] (Table III). The [P]BU[17/4C] was similar in the three siblings; the values were 0.16, 0.16, and 0.17. These values are ~ 10 times those found in normal adult men (13) and 15 times those found in two normal prepubertal boys (6). The PR-E2 in subjects III-1, III-3, and III-4 were 65 μg, 216 μg, and 23 μg/d, respectively. For comparison, the mean PR-E2 of four normal adult men was 58 μg/d with a range of 48–65 μg/d (13), and that of a normal 8-yr-old boy was 24 μg/d (6).

The PR-E2 and [P]BU[17/4C] (Table III). The [P]BU[17/4C], determined in subject III-3, was 0.03. This value is ~ 8-10-fold greater than the fractional conversion of T to E2 previously determined by MacDonald et al. (13) in four normal adult men (range: 0.0028–0.004; mean value = 0.0037). The production rate of E2 in this subject was 150 μg/d, a value that is severalfold greater than the value of 44 μg/d (range: 35–47 μg/d) for normal adult men (13). The [P]BU[17/4C] ratios of urinary E1 and E2 were the same after the infusion of [P]BU[17/4C] and [P]BU[17/4C] (Table III), which suggested that E2 was the principal product of the aromatization of T.

Circadian variation of plasma hormones. Because the levels of plasma T were lower than expected for the respective Tanner stage of pubertal development in subjects III-1, III-3, and III-4 at the time of initial evaluation (Table I), and because the levels of plasma T and A remained low during the course of a year of observation (Table II), the circadian variation in plasma T and A, and serum LH and FSH, was investigated in subjects III-1 and III-3. A diurnal variation in T and A was observed (Fig. 3).

LHRH stimulation and TRH stimulation tests. Pituitary function tests were conducted in subjects III-1 and III-3 when they were 17-3/12 and 12-10/12 yr old, respectively. After LHRH administration, serum LH rose from a base-line level of 29 ng/ml to a maximum value of 142 ng/ml at 180 min in subject III-1 and from 28 ng/ml to 170 ng/ml in subject III-3. These results are in the normal range (14). After TRH administration, the peak serum level of TSH (subject III-1 = 19.0 μIU/ml; subject III-3 = 16.0 μIU/ml) and the peak serum level of prolactin (subject III-1 = 40 ng/ml; subject III-3 = 41 ng/ml) were normal (15, 16).

hCG stimulation test. The results of the hCG stimulation tests conducted in subjects III-1 and III-3 are given in Table IV. The plasma E2 levels were disproportionately elevated compared with those of plasma T in both the base-line and the post hCG-treatment samples (17). In addition, there was no abnormal accumulation of plasma DHA, P, 17-OHP, or A after hCG stimulation (data not shown).

Discussion

In 1977, Hemsell et al. (6) described the findings in a prepubertal boy who developed gynecomastia. Breast development in this

Table III. Transfer Constants of Conversion of Plasma Cyp-Steroids to Estrogens and Urinary Production Rate of Estrogens in Sibs with Familial Gynecomastia

<table>
<thead>
<tr>
<th>Subject</th>
<th>III-1</th>
<th>III-3</th>
<th>III-4</th>
<th>Normal adult men</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (yr)</td>
<td>16</td>
<td>12</td>
<td>11</td>
<td>26-35</td>
</tr>
<tr>
<td>[P]BU[17/4C]</td>
<td>0.16</td>
<td>0.16</td>
<td>0.17</td>
<td>0.016</td>
</tr>
<tr>
<td>PR-E2 (μg/24 h)</td>
<td>65</td>
<td>216</td>
<td>23</td>
<td>55</td>
</tr>
<tr>
<td>[P]BU[17/4C]</td>
<td>—</td>
<td>0.03</td>
<td>—</td>
<td>0.0037</td>
</tr>
<tr>
<td>PR-E2 (μg/24 h)</td>
<td>—</td>
<td>150</td>
<td>—</td>
<td>44</td>
</tr>
<tr>
<td>[P]BU[17/4C]</td>
<td>0.47</td>
<td>—</td>
<td>0.53</td>
<td></td>
</tr>
<tr>
<td>[P]BU[17/4C]</td>
<td>0.97</td>
<td>—</td>
<td>0.93</td>
<td></td>
</tr>
</tbody>
</table>

Figure 3. Circadian variation of plasma T, A, serum FSH, and LH in two siblings with familial gynecomastia.
subject began at 8-7/12 yr of age and progressed slowly. The rate of linear growth after the appearance of gynecomastia was accelerated and the bone age was advanced appreciably. Pubic hair was noted first at 10-5/12 yr and levels of plasma T ranged between 270 and 410 ng/dl from 12-6/12 to 13 yr. A persistent elevation in plasma levels of E₁ and E₂ also was demonstrated. After excluding an adrenal or testicular source of excess estrogen secretion, the elevated estrogen levels were shown to be due to an increase in the peripheral conversion of plasma C₁₀-steroids to estrogens. The fractional conversions of plasma A to E₁ and of plasma T to E₂ were 50 times those found in normal young men. In addition, 94% of the extraglandular aromatization occurred in extrahepatic sites. Furthermore, the E₁ produced in peripheral tissue from the aromatization of A underwent extensive sulfurylation in the tissue site(s) of aromatization before its release into blood (6).

A number of similarities exist between the subjects of the present study and the subject described by Hemsell et al. The five affected family members in our study developed gynecomastia at an early age (10–11 yr), starting concomitantly with the sexual maturational processes of male puberty. This sequence is in contrast to that of the physiologic pubertal gynecomastia that usually begins 6–12 mo after the appearance of pubic hair. In addition, despite the relatively early appearance of physical changes, presumed to be due to adenarche, pubertal development progressed slowly in the subjects with gynecomastia in this study. 2 yr after the appearance of pubic hair in subject III-3, the levels of plasma T still were quite low. The low levels of plasma T in subject III-1 when he was 17 yr of age are also striking considering that pubertal development had commenced at 11 yr of age. In comparison to males with physiologic gynecomastia and Tanner Stage III-IV pubertal development, where a T/E₂ ratio of 140:1 (ng/dl: ng/dl) has been reported (1), there was a disproportionate elevation of plasma E₂ levels compared with those of plasma T in our subjects, the T/E₂ ratios being 16:1 in subject III-1 and 5:1 in subject III-3. Known causes of gynecomastia were we excluded on the basis of a normal 46,XY karyotype, normal or low levels of serum gonadotropins and prolactin, normal serum levels of thyroxine, and normal liver function tests. A scan (CT) of the adrenal glands in subject III-3 was normal.

The fractional conversion of [³H]Bu₁ÆEI in subjects III-1, III-3, and III-4 was 10 times that found in normal young adult men, and 15 times that of prepubertal boys. PR-E₁ in the proband (III-3) was markedly elevated (216 µg/d), whereas in his older brother (III-1) it was 65 µg/d, which was well within the range for normal adult men (13). Unfortunately, the production rates of A were not determined, and therefore we cannot determine the rate of production of E₁ from the conversion of A to E₁. In fact, the variation in the plasma levels of A from day to day (Table II) and within a 24-h period (Fig. 2) would have made it impossible to estimate accurately the production rate of A. One possible explanation for the difference in PR-E₁ in subjects III-1 and III-3 might be that the production rate of A in subject III-3 was greater because of obesity (18). Although the PR-E₁ in subject III-1 was in the normal adult range, it was higher than expected for a pubertal boy. The presence of mild gynecomastia was consistent with this moderate increase in PR-E₁.

The PR-E₁ of subject III-4 at 11-1/12 yr was normal for a prepubertal boy, although the fractional conversion of A to E₁ in this subject was elevated. At that time, there were a few strands of pubic hair but the genitalia of this boy were otherwise entirely prepubertal, as were the plasma levels of T, E₂, LH, and FSH. We assume that the production rate of A was variable during the early stage of his adenarche, and that the gynecomastia resulted from intermittent exposure to estrogen arising from the extraglandular conversion of adrenal A to E₁.

The [³H]/[¹⁴C] ratios of urinary E₁ and E₂ were the same after the infusion of [³H]E₂ and [¹⁴C]A, which indicated that estrone (or possibly estrone-sulfate) was the product of aromatization of plasma A.

The fractional conversion of T to E₂ in subject III-3 was eight times that found in normal adult men. The urinary production rate of E₂, 150 µg/24 h, was above the normal range. It should be stressed that [³H]Bu₁⁰¹ and [³H]Bu₁ÆEI represent the sum of all routes of conversion of T to E₂, and of A to E₁, respectively. E₂ can arise from T by direct conversion or indirectly by the pathway T → A → E₁ → E₂. Similarly, E₁ can arise from A by direct conversion or indirectly by the pathway A → T → E₂ → E₁.

After determining that the estrogen imbalance observed in the subject of this study was due to increased extraglandular aromatization, we speculated as to whether E₁ in increased amounts might act to diminish the release of LH, and therefore diminish testosterone production. We also sought to ascertain whether the increased E₂ levels might act to directly inhibit the production of T by Leydig cells.

Although we could not assess directly the effects of elevated estrogen production on the release of LH by the pituitary, we gathered data that is suggestive that the function of the hypothalamic-pituitary axis was normal in these subjects. It is known that in normal pubertal boys, there is a marked diurnal variation in levels of plasma T, with peak values occurring at night, ~4–8 h after the onset of sleep (19, 20). The magnitude of the diurnal variation in plasma T decreases gradually as puberty progresses until the level of plasma T is similar throughout the day. Large and Anderson (2) found the same diurnal variation for plasma T levels in boys with gynecomastia and in boys in whom there was a physiologic delay in the onset of puberty. As presented in Fig. 2, the diurnal variation in plasma T was normal in subjects III-1 and III-3, which suggested that the normal rhythmicity in LH release was preserved. The marked diurnal variation in plasma T in our subjects also may provide an explanation for the apparent discrepancy between the degree of sexual development observed and the low levels of plasma T obtained at the time of regular clinic visits. Additional evidence in support of normal pituitary
function in the subjects of this study was obtained by the
finding of a normal response in serum LH to LHRH treatment
(14) and a normal response in serum prolactin after the
infusion of TRH (16).

With regard to Leydig cell function in these subjects, by
hCG stimulation we found that the Leydig cells were capable
of producing testosterone in normal amounts in spite of
elevated levels of E2. This is in contrast to some findings in
men (21) and in experimental animals (22) that are suggestive
that E2 acts directly to inhibit the secretion of T by Leydig
cells. These studies, however, do not rule out a possible defect
in T secretion in our patients, since base-line levels of plasma
T were low for Tanner Stage in these subjects. Unfortunately,
we do not have information about sperm production in the
subjects of this study and we cannot predict the effect of the
increased extraglandular aromatase activity on fertility.

We also studied the response of plasma levels of E2 to
hCG administration. In these studies, hCG was administered
intramuscularly in a daily dose of 3,000 U/M2 of body surface
area, with a maximum of 5,000 U, for 5 d. By use of this
regimen in a group of normal adult men, the mean level of
plasma T rose from 612 ng/dl to 1,404 ng/dl (unpublished
data). By use of a similar regimen in a group of normal men,
Weinstein et al. (16) found that plasma T rose from 770 to
1750 ng/dl, and that the mean level of plasma E2 rose from
3.0 to 10.4 ng/dl. In our subjects (III-1 and III-3), the increase
in plasma T after hCG was normal. The rise in plasma E2
levels after hCG stimulation, however, was three times greater
than that observed in normal men, which suggested again that
in these subjects there was increased extraglandular aromatase
activity.

We have presented evidence that in several men of a single
family, there was increased extraglandular aromatase activity
compared with that found in normal men. Several possibilities
exist to explain this abnormality. The defect could be due to
a gene mutation resulting in an enzyme complex with increased
activity, or to an increase in the intracellular concentration of
enzyme due to increased production or decreased catabolism
of the enzyme.

Hemsell et al. (6) proposed that the increased extraglandular
aromatization was due to a persistence of fetal levels of both
aromatase and sulfokinase enzyme activities. In view of the
difference in the level of peripheral aromatase activity observed
in our subjects (10 times normal), compared with the activity
in the subject described by Hemsell et al. (50 times normal),
it is possible that one is dealing with different underlying
defects in the two families. It is of interest that George et al.
(23) demonstrated that the increased estrogen formation in the
Sebright Bantam chicken is due to an abnormal elevation in
the level of peripheral aromatase activity.

Aromatase activity is widely distributed in several extra-
glandular tissues including skin, fat, muscle, kidney, bone, and
certain areas of the central nervous system. We could not
determine whether the abnormality in our patients resides in
one tissue alone or whether the defect is present in all tissues.
Hemsell et al. (6) reported that <6% of the aromatization
observed in their subject occurred in intrahepatic sites. This is
consistent with the recent observation that aromatase activity
in liver accounts for only a minimal fraction of the extraglan-
dular aromatization in normal adult men (24).

The affected members of the family of this study include
three siblings, a maternal cousin, and a maternal uncle. We
found increased fractional conversion of androstenedione and
testosterone to estrogens in the three siblings, but we have not
studied the affected maternal uncle and maternal cousin. Both
the history and the hormonal profile of the uncle and cousin
are suggestive that in these men there also is excessive extra-
glandular aromatase activity. The pattern of inheritance of this
disorder in the family studied is suggestive of an X-linked
recessive or a sex-limited autosomal dominant trait. Further
studies will be necessary to elucidate the underlying molecular
and genetic defects in affected persons.

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