Androgen secretion and characteristics of testicular hCG binding in cryptorchid rats

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Summary. Response of the cryptorchid testis to gonadotrophic stimulation was assessed by comparison of the androgen production capability in vivo and in vitro with that of the normal scrotal testis. Serum androgen concentrations in cryptorchid rats were similar to those in normal rats, and the incremental increase 60 min after 50 i.u. hCG (i.v.) was about 7-fold for both groups. Basal and hCG-stimulated androgen production in vitro was higher for abdominal testes (557 and 3286 ng/pair) than for scrotal tests (157 and 504 ng/pair). Specific binding of hCG by testicular homogenates was slightly higher ($P < 0.05$) for cryptorchid testes when expressed per unit weight, but Scatchard analysis indicated that although hCG binding affinities did not differ ($K_a = 2 \times 10^{10} \text{M}^{-1}$), hCG binding capacity of cryptorchid testes was only 75 ng, compared to 219 ng for scrotal testes. These data indicate that a discrepancy exists between androgen production in vivo and in vitro by cryptorchid testes and that normal serum androgen concentrations are maintained in the presence of decreased numbers of testicular LH/hCG receptors.

Introduction

Induced cryptorchidism in rats results in testicular damage which is accompanied by increased circulating levels of LH and FSH (Swerdloff, Walsh, Jacobs & Odell, 1971; Rager, Zarzychi, Eichner & Gupta, 1975; Gomes & Jain, 1976; de Kretser, Sharpe & Swanston, 1979; Kerr, Rich & de Kretser, 1979). Similar responses are obtained after heat treatment of the testes (Aafjes, Vreeburg & Schenck, 1978). Although testosterone can suppress these elevated levels of gonadotrophins (Aafjes et al., 1978), increases and decreases in circulating concentrations of testosterone after bilateral cryptorchidism have been reported (Amatayakul, Ryan, Uozumi & Albert, 1971; Lloyd, 1972). Clegg (1961) reported that artificial cryptorchidism was followed by a transient increase in the numbers of interstitial and Leydig cells of the rat testis, and the binding capacity for hCG by cryptorchid testes is enhanced on a per unit weight basis (Frowein & Engel, 1975; de Kretser et al., 1979). The objective of the present study, therefore, was to determine the androgen secretory response of cryptorchid rat testes in vivo and in vitro and relate these findings to testicular hCG binding characteristics.

Materials and Methods

Animals. Mature male rats weighing approximately 300 g each were used. The 24 rats in Group 1 were sham operated (controls) and the 24 in Group 2 were made bilaterally cryptorchid as described previously (Schanbacher, Gomes & VanDemark, 1974). The consequences of induced cryptorchidism were assessed 30 days later in 3 ways.
Androgen secretion in vivo. Serum androgen concentrations were measured in 10 control and 10 cryptorchid rats immediately before and 60 min after an intravenous injection of 50 i.u. hCG. Blood was collected and the hormone injected by cardiac puncture under light ether anaesthesia. The animals were subsequently killed to check the position of the testes and whether any adhesions or abnormalities had resulted. Weights of the testes and seminal vesicles were recorded.

Androgen production in vitro. The incubation system described by Dufau, Catt & Tsuruhara (1971) and modified by Bartke, Williams & Dalterio (1977) was used to determine testosterone production in vitro of testicular tissue from 10 control and 10 cryptorchid rats. After death by cervical dislocation, the testes were immediately removed, decapsulated and incubated individually in 5 ml Krebs–Ringer bicarbonate buffer containing 1 mg glucose/ml. One testis from each animal was incubated in the absence of gonadotrophin to estimate basal testosterone production; the other testis was incubated in the presence of 50 mi.u. hCG/ml to estimate stimulated testosterone production. Testes were incubated at 37°C for 4 h and the media were then collected after centrifugation at 1500 g for 5 min.

hCG binding. A procedure similar to that described by Bex & Bartke (1977) for the hamster testis was used to determine gonadotrophin binding to crude homogenates of testes from 4 control and 4 cryptorchid rats. After death by cervical dislocation, testes were excised, decapsulated, weighed to the nearest mg and then homogenized in a ground-glass tissue homogenizer together with 5 ml ice-cold 10 mm-phosphate buffer (PBS) containing 1 mM-EDTA, 140 mM-NaCl and 15.4 mM-NaHCO₃ (pH 7.5). The homogenate was centrifuged at 1500 g, and the resultant tissue pellet was resuspended in PBS at a concentration of 1 g/5 ml. This preparation was used immediately or frozen at −70°C until assayed for hCG binding.

Radiolabelled hCG was prepared by iodination of purified hCG (CR 119: 11 600 i.u./mg) with 0.5 mCi ¹²⁵I (New England Nuclear, buffered preparation). Chloramine T (5 µg) was used to iodinate 2.5 µg hCG in a reaction vessel for 30 sec. The radiolabelled hCG was separated from free iodide on a 1 × 22 cm Bio-Gel P-100 column eluted with PBS. Specific activity of ¹²⁵I-labelled, hCG determined by self-displacement in the rat testis receptor assay, was estimated at 47 µCi/µg hCG (CR 119). Sensitivity of the binding-inhibition curve was <0.5 ng hCG.

The binding of hCG was determined for 100 µl aliquots (20 mg) of testicular homogenate. Each tube received 55 000 c.p.m. (528 pg) of ¹²⁵I-labelled hCG and either 0 or 100 i.u. hCG in a total of 0.5 ml PBS. All samples were run in duplicate at 20°C. Following a 20 h incubation 2 ml ice-cold PBS were added, and the samples were centrifuged at 1500 g for 15 min. The supernatants were removed and the sediments counted in a spectrometer (No. 5385, Packard). Specific binding of ¹²⁵I-labelled hCG was calculated as total binding (c.p.m.) minus non-specific binding (c.p.m.). Self displacement curves for a representative normal and cryptorchid rat were used to generate the Scatchard plots (Scatchard, 1949) and calculate the binding affinity and number of binding sites for each preparation.

Concentrations of androgen (testosterone + 5α-dihydrotestosterone) in serum and incubation media were determined by radioimmunoassay (Schanbacher, 1976). The sensitivity was 0.2 ng/ml and the intra-assay coefficient of variation was <12% for each duplicate.

Differences between means were tested statistically by Student’s t test.

Results

Testicular weight was reduced markedly after 30 days of induced cryptorchidism (Table 1), but seminal vesicle weight was maintained and serum androgen concentration was not significantly different from that in controls. The ability of the cryptorchid testis to secrete testosterone in response to hCG appeared normal because serum androgen increased similarly.
Testosterone secretion and hCG binding in cryptorchid rats

Table 1. Effect of surgically induced, bilateral cryptorchidism on weights of the testes and seminal vesicles and serum androgen response to hCG by rats in vivo

<table>
<thead>
<tr>
<th>Rat</th>
<th>Wt of paired testes (g)</th>
<th>Wt of seminal vesicles (g)</th>
<th>Serum androgen (ng/ml) Before hCG</th>
<th>Serum androgen (ng/ml) After hCG*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham-operated</td>
<td>3.81 ± 0.06</td>
<td>1.16 ± 0.10</td>
<td>6.8 ± 0.7</td>
<td>42.0 ± 4.4</td>
</tr>
<tr>
<td>Cryptorchid</td>
<td>1.25 ± 0.08†</td>
<td>0.93 ± 0.07</td>
<td>6.0 ± 0.6</td>
<td>40.1 ± 4.7</td>
</tr>
</tbody>
</table>

Values represent mean ± s.e.m. for 10 normal and 10 cryptorchid rats.

* At 60 min after 50 i.u. hCG given i.v.
† Significantly different from corresponding value for control rats: P < 0.01.

As shown in Table 2, basal androgen production in vitro was considerably higher for abdominal testes than for scrotal testes. Stimulation by hCG enhanced androgen production for both groups, but the response by cryptorchid testes was significantly greater.

Table 2. Effect of surgically induced, bilateral cryptorchidism on basal and hCG-stimulated (50 mi.u./ml) androgen production by the rat testis in vitro over 4 h

<table>
<thead>
<tr>
<th>Testes</th>
<th>Basal testosterone production</th>
<th>Stimulated testosterone production</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Per g testis (ng)</td>
<td>Per paired testes (ng)</td>
</tr>
<tr>
<td>Scrotal</td>
<td>49 ± 12</td>
<td>157 ± 39</td>
</tr>
<tr>
<td>Abdominal</td>
<td>732 ± 70**</td>
<td>557 ± 186*</td>
</tr>
</tbody>
</table>

Values represent mean ± s.e.m. for 10 scrotal and 10 abdominal testes, each from individual rats.

Significantly different from corresponding value for scrotal testes: * P < 0.05; **P < 0.01.

Specific binding of hCG by testicular homogenates of cryptorchid rat testes (47.4 ± 2.9% c.p.m./20 mg tissue) was greater (P < 0.05) than that determined for testicular homogenates of control rats (38.5 ± 2.0%). Scatchard plots of actual binding data are presented in Text-fig. 1. Increasing mass of hCG was added to duplicate tubes containing 20 mg testicular tissue and the linear plots of B/F (125I-labelled hCG) versus B (pM) demonstrated that association constants were similar for scrotal testes (K_s = 1.6 × 10^-10 M^-1) and cryptorchid testes (K_s = 2.1 × 10^-10 M^-1). The number of binding sites/20 mg scrotal and abdominal testicular tissue was 25 and 32.5 fmol, respectively. Based on average testicular weights, maximal hCG binding was approximately 62.5 ng/g scrotal testicular tissue (219 ng/paired testes) and approximately 81 ng/g abdominal testicular tissue (75 ng/paired testes).

Discussion

Gross changes occur in the germinal epithelium after induced cryptorchidism, but changes in the morphological and biochemical functions within the interstitial compartment are ill-defined (Bergh & Damber, 1978). Increased numbers (Clegg, 1961) and size of Leydig cells (Damber, Bergh & Janson, 1978; Kerr et al., 1979) have been reported for abdominal testes. Combined with a change in the pituitary–testicular endocrine axis (Swerdloff et al., 1971; Rager et al., 1975; Gomes & Jain, 1976; de Kretser et al., 1979) of the cryptorchid rat, Leydig cell damage

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and malfunction seem apparent. Hauger, Chen, Kelch & Payne (1977) have concluded that the normal testis depends on anterior pituitary hormones for maintenance of testicular LH receptors and testosterone secretion and so the imbalance in gonadotrophin secretion imposed by cryptorchidism may have affected both.

A transient decrease in serum testosterone has been noted after induced cryptorchidism (Gomes & Jain, 1976), but near-normal androgen concentrations were attained in the present study by 30 days after surgery. The responses of normal and cryptorchid rats to intravenous hCG were also equivalent in this study, although Kerr et al. (1979) reported a diminished testosterone response to hCG administration in cryptorchid rats.

Basal and hCG-stimulated androgen production in vitro confirm the findings of de Kretser et al. (1979) and emphasize the need to investigate further the differences between in-vitro and in-vivo testosterone production by scrotal and abdominal testes.

**Text-fig. 1.** Scatchard plots for specific binding of 125I-labelled hCG to testicular homogenates of a normal male rat (●) and a cryptorchid rat (○). The affinity constant ($K_a$), calculated from the slopes, was approximately $2 \times 10^{10}$ M$^{-1}$ for both scrotal and abdominal testes. The number of binding sites, taken from the x intercepts, was 1.25 pmol/g scrotal testis and 1.62 pmol/g abdominal testis. Molecular weight of hCG was taken as 50 000.

Scatchard analysis of testicular hCG binding indicates that receptor affinity is not affected by cryptorchidism but that capacity to bind hCG is decreased. Total binding capacity was nearly 3 times greater in scrotal testes than in abdominal testes. Frowein & Engel (1975) obtained a similar value of 95 ng/g normal scrotal testis tissue, and found enhanced binding capacity in cryptorchid testes relative to normal when expressed on a per unit protein basis, although Hagenäs, Ritzén, Svensson, Hansson & Purvis (1978) and de Kretser et al. (1979) reported a substantial decrease in binding capacity when expressed on a per testis basis. Importantly, the association constant ($K_a = 10^{10}$ M$^{-1}$) and number of binding sites ($Q = 10^{-12}$ mol/testis) reported previously for the intact rat (Catt & Dufau, 1973a; Frowein & Engel, 1975)
and that reported here are similar. The paradoxical relationship between hCG binding capacity and in-vitro androgen production by cryptorchid testes is misleading because ultrastructural characteristics of the Leydig cells from abdominal testes suggest that these cells are extremely steroidogenically active (Kerr et al., 1979). The phenomenon of "spare receptors" described by Catt & Dufau (1973b) seems to be an inherent characteristic of the normal rat testis and that elevated levels of gonadotrophins in cryptorchid rats decrease testicular LH/hCG receptors by autoregulation. A moderate reduction in LH/hCG receptors in the normal rat testis does not seem to affect androgen production capability (Purvis, Torjesen, Haug & Hansson, 1977; Pirke, Vogt & Geiss, 1978; Purvis, Clausen, Brandtzæg & Hansson, 1978; Sharpe & McNeilly, 1978). Although this conclusion can explain the normal androgen response to hCG in vivo, it fails to explain the enhanced response of the cryptorchid testis in vitro. The dose-response study conducted by de Kretser et al. (1979) suggests that the discrepancy between testosterone production in vivo and in vitro might depend on the degree of gonadotrophic stimulation being given; but this and other hypotheses need critical evaluation.

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References


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