Relative Roles of Follicle-stimulating Hormone and Luteinizing Hormone in the Control of Inhibin Secretion in Normal Men

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Abstract

The glycoprotein hormone inhibin is produced by the Sertoli cells of the testis under the influence of follicle-stimulating hormone (FSH) and is postulated in turn to inhibit FSH secretion. Luteinizing hormone (LH) is not recognized to have an important role in the control of inhibin secretion in any species. To determine the relative roles of FSH and LH in the control of inhibin secretion in man, we examined the effects of selective FSH and LH replacement on serum inhibin levels in normal men whose endogenous gonadotropins were suppressed by testosterone (T). After a 3-mo control period, nine men received 200 mg T enanthate i.m. weekly for 3–9 mo. During T treatment, serum LH and FSH levels were markedly suppressed and serum inhibin levels fell to 40% of control values. While continuing T, 3–5 mo of treatment with purified hFSH (n = 4) or LH (n = 4) increased the respective serum gonadotropin level into the upper normal range and significantly increased inhibin levels back to 64 and 55% of control values, respectively. Supraphysiological LH replacement with high doses of human chorionic gonadotropin (n = 3) returned serum inhibin levels to 63% of control values. In no case did inhibin levels return fully to control levels. In conclusion, serum inhibin levels fell during gonadotropin suppression and were partially and approximately equally restored by either FSH or LH treatment. FSH presumably acts directly on the Sertoli cell to increase inhibin secretion whereas LH may act via increases in intratesticular T levels and/or other factor(s).

Introduction

Inhibin is a glycoprotein produced by the testis and ovary and has been postulated to have an important role in the regulation of pituitary follicle-stimulating hormone (FSH) secretion (1). In the male, in vitro studies have shown that Sertoli cells (2–5) secrete inhibin under the influence of FSH but not luteinizing hormone (LH) (5–9). The effect of LH on testicular inhibin secretion in vivo is unknown. Inhibin (a glycoprotein heterodimer) was recently purified from ovarian follicular fluid (10–12), cloned, and sequenced (13–15). Pure bovine inhibin has been used in the development of sensitive and specific RIA systems applicable to human serum (16, 17), allowing investigation of inhibin levels in women during ovulation induction (18, 19), the normal menstrual cycle (17), and pregnancy (19, 20).

In previous studies, we have demonstrated that both FSH and LH are required to maintain quantitatively normal spermatogenesis in man (21–25). As a result of the suppression of gonadotropins, testosterone (T)1 administration to normal men markedly reduced sperm counts. While continuing T, selective replacement with physiological doses of either highly purified human FSH (22) or LH (23), or supraphysiological doses of human chorionic gonadotropin (hCG) (21, 24), were each associated with only partial restoration of spermatogenesis, indicating a separate role for each gonadotropin. Quantitatively normal spermatogenesis could only be restored with replacement of both gonadotropins (25).

As Sertoli cell function is intimately related to both spermatogenesis and inhibin production, this study aimed to examine the effect on inhibin levels of T-induced gonadotropin suppression in normal men and subsequent selective replacement with FSH or LH. Specifically, we sought to determine whether inhibin secretion decreased during gonadotropin suppression, and, if so, whether inhibin levels could be stimulated by selective replacement of FSH or LH in physiological amounts.

Methods

Subjects

A total of nine normal men (25–40 yr old) volunteered to participate in these studies. The experimental protocols, sperm counts, and gonadotropin levels have been described previously (22–24). The normality of subjects was established by (a) complete medical histories and physical examinations; (b) routine hematological analysis, blood chemistry, coagulation times, and urinalysis; (c) six seminal fluid analyses obtained over a 3-mo period (sperm concentration > 20 million/ml, sperm motility > 50% and > 60% oval forms); and (d) normal basal LH, FSH, and testosterone levels; normal LH and FSH secretory patterns on blood sampling every 20 min over 6 h and normal FSH and LH responses to a 4-h continuous intravenous infusion of 50 μg of gonadotropin-releasing hormone (GnRH).

Experimental protocol

All subjects underwent a 3-mo control period during which no hormones were given and only blood samples were collected and semen

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1. Abbreviations used in this paper: bFF, bovine follicular fluid; CV, coefficient of variation; GnRHa, gonadotropin-releasing hormone; hCG, human chorionic gonadotropin; T, testosterone.
analyses performed. After the control period, all subjects received 200 mg testosterone enanthate (Delatestryl, E. R. Squibb and Sons, Princeton, NJ) i.m. weekly until sperm counts were < 2 million/ml, which occurred after 2–4 mo of treatment. The duration of T alone treatment was between 3 and 5 mo (studies 1 and 2) or 9 mo (study 3). While continuing T administration, selective gonadotropin replacement regimens were undertaken as follows.

Study 1. In four subjects, 100 IU hFSH s.c. daily was administered together with T for 3 mo. The hFSH preparation (LER 1577, Lot No. 4) was kindly provided by the National Pituitary Agency, Baltimore, MD. As previously described, this preparation contained < 1% contamination with LH bioactivity. After the FSH plus T period, FSH injections were stopped and T continued until three successive sperm counts were again < 5 million/mℓ; this occurred between 1–5 mo of T therapy. After T treatment, the subjects entered a recovery period until three successive sperm counts were within the subject's own control range.

Study 2. In four subjects, 1,100 IU hLH s.c. daily was coadministered with T for a period of 4–6 mo. The hLH preparation (LER 1549, batch A-3) was also provided by the National Pituitary Agency, Baltimore, MD. This preparation contained < 0.2% contamination with FSH bioactivity as assessed in a rat ovarian augmentation bioassay. All hormones were then stopped and the subjects entered a recovery period until three successive sperm counts returned into the subject's own control range.

Study 3. In three men, 5,000 IU hCG (Profasi, Serono Laboratories, Inc., Braintree, MA) i.m. three times weekly was coadministered with T for a period of 6–7 mo. Both hormones were then ceased and the subjects entered a recovery period until three successive sperm counts were within the subjects' control range.

Measurements. Throughout the study periods, blood samples were obtained each month and semen analyses performed twice monthly. A 6-h urine sample was obtained from each subject at the end of the control, T alone, and T plus gonadotropin periods. Blood samples were obtained immediately before hormone injections and serum was stored at −20°C before assay. For the assay of serum inhibin, the current study made use of residual serum samples from the original three studies (22–24).

Radioimmunoassays. A heterologous RIA for human serum inhibin was used based on a previously described method (17). The assay was modified to use a new inhibin antisera (As 1989), raised in a New Zealand white rabbit to 31 kD bovine follicular fluid (bff) inhibin. The RIA using this antisera showed no interference from castre serum and an enhanced sensitivity when compared with the previous method. This RIA was specific to intact inhibin, showing < 1% cross-reactivity with transforming growth factor β, bovine activin A, and free inhibin subunits obtained after reduction and alklylation of 31 kD bovine inhibin. A partially purified human follicular fluid inhibin preparation was prepared for use as the RIA standard as described (17). This material was calibrated in terms of its in vitro inhibin bioactivity using a bFF inhibin standard preparation itself previously calibrated against an ovine testicular lymph preparation of defined unitage 1 U/mg (26). Male serum samples showed parallel dose-response lines to the RIA standard and a serum pool obtained from women undergoing ovulation induction, which was used as quality control. In the RIA, 200–μl samples were assayed in duplicate. The sensitivity of the assay was 100 U/liter. The interassay coefficient of variation (CV) was 11% and the interassay CV in the upper, mid, and lower portions of the standard curve were 12.0, 3.3, and 4.8%, respectively for five assays. Inhibin immunoactivity has been shown to be stable during repeated freezing and thawing (17). Inhibin levels in the study subject's control sera were ~ 25% lower than those of a recently examined group of normal young men, suggesting the possibility of some loss of immunoactivity during the prolonged storage of the samples used in this study. However, all samples used for this study had been stored for approximately the same length of time, so that any loss of immunoactivity with storage could not account for the differences found between the various hormonal manipulations in these studies.

Serum FSH and LH were determined by RIA as described previously (21) using reagents distributed by the National Pituitary Agency and using the reference preparation LER 907. In the FSH RIA, the first antibody was rabbit anti–human FSH batch No. 5 and the tracer was HS-1 radiolabeled with 125I using chloramine T. The sensitivity of the assay was 25 μg/liter. The intraassay CV was 7.3% and the interassay CV was 9.7%. Urinary FSH excretion was measured by RIA after acetone precipitation of urine (27) as described previously (21) using the Second International Reference Preparation of human menopausal gonadotropin as the reference standard. In the LH RIA, the tracer was purified hCG (supplied courtesy of Dr. C. Alvin Paulsen, Department of Medicine, University of Washington) that was radiolabeled with 125I by chloramine T. The sensitivity of the assay was 6 μg/liter and the intra- and interassay CVs were 5.5 and 8.4%, respectively.

Bioassay. Serum LH levels were measured by in vitro bioassay using a modification (21) of the procedures described by Van Damme et al. (28) and Dufau et al. (29). T production was measured from dispersed Leydig cells isolated from immature Swiss Webster mice (age 5–7 wk). The reference standard was LER 907. All samples were run in duplicate and the detection limit of the assay was 100 μg/liter. The mean intra- and interassay CVs for pooled human sera were 14 and 24%, respectively.

Seminal fluid analysis. Sperm concentrations were determined using a Coulter counter (Coulter Electronics Inc., Hialeah, FL), whereas concentrations < 15 million/mℓ were confirmed by direct determination using a hemocytometer as described.

Statistical analysis. Monthly inhibin levels during each phase of treatment were averaged for each patient. To ensure that each hormonal manipulation had sufficient time to both be effective and to have been adequately washed out before the next study period, only results obtained during the following time periods were included in these analyses: (a) control period, all values until the day of commencement of T; (b) T alone and T plus gonadotropin periods, all values obtained after 2 wk of T or gonadotropin treatment until the day of commencement of the next study period; (c) subsequent periods on T alone, all values obtained at least 2 wk after cessation of gonadotropin therapy until the day of the next hormonal administration or within 2 wk of the last T injection in the early recovery phase; and (d) recovery period, all values obtained at least 4 wk after the cessation of T.

Mean monthly serum LH and FSH levels and mean sperm concentrations for each period of study were calculated for each patient as described. Sperm concentrations were not normally distributed and values were log-transformed before statistical analysis.

Mean hormonal values for each patient during the various treatment periods were obtained. Analysis of variance with repeated measures was used to detect a significant change across the treatments. The existence of a significant difference between treatments was determined by paired t test with correction for multiple comparisons using the Bonferroni method. Values are expressed as mean±SEM.

Results

Study 1. The changes in serum inhibin levels in the four subjects receiving 100 IU hFSH daily are shown in Fig. 1. Inhibin levels decreased to 35% of control values after T administration (432±34 vs. 152±14 U/liter, P < 0.02). During hFSH treatment, inhibin levels increased (P < 0.05) to between 60 and 76% control values with a mean level of 278±16 U/liter. Nonetheless, inhibin levels were still significantly lower than control (P < 0.05). After cessation of hFSH, inhibin levels again fell to 150±10 U/liter on T alone, then rose to 418±44 U/liter after cessation of T (NS vs. control).

Serum-immunoreactive FSH levels were suppressed from control levels (98±21 μg/liter) to undetectable (< 25 μg/liter) levels during T treatment. Urinary FSH excretion was sup-
pressed to levels comparable to those of prepubertal or hypo-
gonadotropic subjects (21). After hFSH treatment, FSH levels
then increased to just above the upper limit of the normal
range (273±44 μg/liter). Urinary FSH excretion was similarly
restored into the upper normal range. Serum LH bioactivity
was markedly suppressed to the limit of assay detection
throughout both T and T plus FSH treatment.

During T therapy, sperm counts declined markedly from a
mean of 99 million/ml to <2 million/ml in all subjects. Dur-
ing hFSH treatment, sperm counts increased significantly
to 33±7 million/ml. In no case did sperm counts consistently
reach the individual’s control values. Upon withdrawal of
hFSH, counts again fell to <1 million on T alone. In two
subjects, sperm counts were obtained after the recovery period
and both were within the individual’s own control range.

Study 2. The changes in serum inhibin in the four subjects
receiving hLH treatment are shown in Fig. 2. During T treat-
ment, inhibin levels decreased to 40% of control values
(432±60 vs. 172±40 U/liter, P < 0.02), hLH treatment led to a
significant (P < 0.05) increase in inhibin levels to 55% of con-
trol levels (240±60 U/liter), which was still significantly (P
< 0.02) lower than control values. In the recovery phase, inhibin
levels returned to control levels (418±64 U/liter).

Serum LH bioactivity fell markedly during T treatment
(390±20 vs. 120±10 μg/liter, P < 0.001). LH injections re-
stored LH bioactivity to the control range when samples were
collected immediately before an LH injection (400±40 μg/
liter) and a peak level, obtained 8 h after injection, at the upper
limit of the normal male range (750±150 μg/liter). Serum FSH
was suppressed from control values (98±12 μg/liter) to unde-
tectable levels (<25 μg/liter) where it remained until the recov-
ery period. Similarly, urinary FSH excretion was sup-
pressed into the prepubertal range throughout T±hLH treat-
ment.

Sperm concentrations were markedly suppressed by T
(98±17 million/ml to <4 million/ml) and partially restored by
hLH treatment (19±4 million/ml after 3 mo of treatment).
In no case did the sperm count consistently reach the individ-
ual’s control range. After recovery, sperm counts returned to
control levels in both subjects in whom this was assessed.

Study 3. The changes in serum inhibin in the three sub-
jects receiving hCG treatment are shown in Fig. 3. During T
therapy, inhibin levels decreased to 38% of control values
(492±170 vs. 186±86 U/liter). hCG treatment led to a signifi-
cant (P < 0.05) increase in inhibin levels to 63% of control
values (308±90 U/liter), which was still lower than control
levels. In the recovery phase, inhibin levels returned to control
levels (498±90 U/liter).

Serum LH immunoactivity fell during T treatment from
28±4 to 7±1 μg/liter (at the limit of assay sensitivity). hCG
administration led to a marked elevation in serum LH-like
immunoactivity (>200 μg/liter) due to cross-reactivity of
hCG in the LH RIA. Serum FSH was suppressed from control
levels (125±34 μg/liter) to undetectable (<25 μg/liter) levels
during both T alone and hCG + T administration. Urinary
FSH excretion was suppressed to prepubertal levels during
both of these periods.

Sperm counts were markedly suppressed by T (82±9 mil-
lion/ml to <1 million/ml) and substantially restored by hCG
treatment (22±4 million/ml after 3 mo of hCG). In some sam-
ples, sperm counts were within the subject’s control range;
however, the mean sperm count remained below control
values.

Discussion

Our results show that serum inhibin levels in normal men
decrease after the suppression of gonadotropin levels by tes-
tosterone administration and are partially restored by either
FSH or LH replacement. During T therapy, levels of FSH
immunoactivity and LH bioactivity were suppressed into the
prepubertal range. Serum inhibin levels were suppressed to
levels similar to those observed in early puberty and in subjects
with Kallmann’s syndrome before pulsatile GnRH treatment
(unpublished observations). Similarly, a substantial fall in tes-
ticular inhibin content has been reported after hypophysec-
tomy in rats (30). GnRH agonist administration to women
also markedly reduced serum estradiol and inhibin levels as a
consequence of functional gonadotropin withdrawal (31). These
studies underline the trophic effect of gonadotropins on
gonadal inhibin production.
Selective human FSH replacement stimulated inhibin secretion in these gonadotropin-suppressed normal men. The highly purified hFSH preparation used had minimal LH contamination as confirmed by the suppressed levels of serum LH bioactivity throughout treatment. Inhibin levels were not returned to control values despite serum FSH levels two- to threefold higher than the subjects' control values, suggesting that FSH alone is incapable of restoring inhibin secretion to normal. This trophic action of LH on human inhibin secretion is in keeping with extensive animal studies. FSH replacement stimulated testicular inhibin content and production in hypophysectomized rats (30). Sertoli cells in culture secrete inhibin bioactivity (5–7) and inhibin-like immunoactivity (8, 9) in response to FSH. Similarly, ovarian inhibin production in vivo is stimulated by FSH-containing agents in rats (32) and humans (18).

In the second study, hLH replacement therapy also partially restored inhibin secretion in these gonadotropin-suppressed normal men. The replacement regimen achieved physiological levels of serum LH bioactivity. The hCG replacement regimen used in the third study has been shown previously to achieve a sixfold elevation in serum LH bioactivity (21). Serum inhibin levels increased somewhat more during hCG treatment than during the hLH regimen but were still lower than control levels. These data indicate that LH-like bioactivity alone, even at supraphysiological levels, cannot restore inhibin secretion to normal levels.

The stimulatory effect of LH upon inhibin secretion was an unanticipated finding in view of previous animal data. hCG replacement had no effect on testicular inhibin production or content in hypophysectomized rats (30). LH had no effect on the secretion of inhibin bio- (4) or immunoactivity (8, 9) by Sertoli cells in culture. The mechanism of the LH effect in this study is unclear. During T suppression in man, despite elevations in serum T, intratubular T levels fall dramatically (33). This result from suppression of LH levels with the concomitant loss of local T production by the Leydig cells of the testicular interstitium. LH replacement thus could potentially mediate the rise in inhibin secretion by restoring local T levels. Androgens have been reported to increase Sertoli cell inhibin secretion in vitro (4), although there are several studies to the contrary (6–9). High-dose T treatment in hypophysectomized male rats partially restored inhibin production rates (34) as assessed by the acute accumulation of inhibin in the testis after unilateral efferent duct ligation (35). Finally, other factor(s) produced by the Leydig cell, or other interstitial cells, may observe the LH effect on inhibin production. For example, androgen-dependent hormonal factors arising from the peritubular cells have been described to affect Sertoli cell production of androgen binding protein (36).

The failure of LH, hCG, or FSH individually to restore inhibin levels to normal is of interest. It is possible that restoration to normal inhibin secretion requires a synergistic effect of FSH and LH; it is also possible that administration of gonadotropins in a more physiological, pulsatile pattern would have been more effective in stimulating inhibin. The response of serum inhibin shows strikingly similar qualitative changes to those described for the sperm counts in these men. Sperm counts were markedly suppressed during T treatment and were partially restored by replacement of either gonadotropin. Inhibin secretion by the Sertoli cell may be coupled to the spermatogenic process and therefore represent an appropriate feed-back factor upon FSH secretion as generally proposed. Alternatively, inhibin itself may be involved in the process of spermatogenesis as a local growth factor and therefore would be anticipated to rise in parallel with the quantitative outcome of spermatogenesis.

In conclusion, we have shown that in normal men whose gonadotropins were suppressed by testosterone, serum inhibin levels were partially restored by either LH or FSH treatment, demonstrating a role for both gonadotropins in the regulation of inhibin secretion. FSH presumably acts directly upon the Sertoli cell to enhance inhibin secretion, whereas LH may act via an increase in local T production or by other factor(s).

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