CASE REPORT

Diagnostic difficulty in polycystic ovary syndrome due to an LH-β-subunit variant

H Kurioka, K Takahashi, M Irikoma, M Okada, T Ozaki, T Ueda and K Miyazaki
Department of Obstetrics and Gynecology, Shimane Medical University, Izumo 693-8501, Japan

Abstract
We initially failed to confirm a case of polycystic ovary syndrome (PCOS) because underestimation of LH concentrations due to a variant form of this hormone resulted in a misleadingly low LH/FSH ratio. A 26-year-old woman presented to our hospital with infertility. Given the presence of bilateral polycystic ovaries, oligomenorrhea and hirsutism, PCOS was suspected, but a normal LH/FSH ratio as measured by RIA led to diagnostic problems. When we remeasured LH and FSH using a chemical luminescence enzyme immunoassay (CLEIA), the ratio of the LH concentration measured by RIA to that measured by CLEIA was 0.29, and the ratio of LH to FSH measured by CLEIA was 3.3 compared with 0.81 measured by RIA. We then diagnosed PCOS. The point mutations Trp8 to Arg8 and Ile15 to Thr15 in the LH subunit were detected in the corresponding gene. The patient’s LH status represented variant and wild-type LH equally. She was therefore diagnosed as heterozygous for the mutant LH-β. Histologic assessment of ovarian tissue after laparoscopic biopsy was compatible with a polycystic ovary.

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Introduction
Luteinizing hormone (LH) is a member of the glycoprotein hormone family that includes human chorionic gonadotropin, follicle-stimulating hormone (FSH), and thyroid-stimulating hormone. These hormones possess α and β subunits; the β subunit confers the hormonal activity. Both subunits are glycosylated at specific residues, and the degree of glycosylation modulates the biological activity of LH. A certain degree of microheterogeneity occurs in glycosylation, contributing to variability in bioactivity and biochemical properties.

Recently developed ultrasensitive immunoassays (1, 2) using monoclonal antibodies for the measurement of LH have resulted in a new risk of error. These monoclonal antibodies may be highly specific to indigenous LH; some studies have noted restricted reactivity of the monoclonal antibodies in certain populations (3, 4). In previous studies (5, 6) PCR was used to specifically amplify the LH-β gene. Subsequent nucleotide sequencing revealed two point mutations in the gene coding for the LH-β subunit, resulting in two amino acid replacements: Trp6(TCG) was changed to Arg(CGG) and Ile13(ATC) to Thr(ACC). Further examination by restriction fragment length polymorphism revealed the existence of homozygotes and heterozygotes for the mutant LH-β gene among family members, indicating a genetic basis for an abnormal LH molecule. Although the homozygotes varied in degree of clinical ovarian dysfunction, the direct effects of the mutant LH on gonadal dysfunction remained unclear. Several commercially available LH assays using α/β heterodimer-specific monoclonal antibodies fail to detect variant forms of LH, or detect them weakly, potentially leading to errors in LH measurements.

We report here on a patient with polycystic ovary syndrome (PCOS) in whom serum LH concentrations and the LH/FSH ratio appeared to be normal when measured with a conventional immunoassay kit.

Case report
The patient was a 26-year-old nulligravida who had been married for 2 years. She presented to our clinic on 13 July 1995, complaining of infertility. Menarche had occurred at age 13, but oligomenorrhea had been noted. Her history included acute pancreatitis at age 23. Her stature was normal and she was not obese (body mass index: 21.5 kg/m²). Hirsutism was present and her Ferriman score (7) was 10. Pelvic examination was normal, but transvaginal ultrasonography revealed bilaterally polycystic ovaries. From these clinical
features, PCOS was strongly suspected. The results of hormonal determinations by RIA in the early follicular phase (Table 1) indicated a normal LH/FSH ratio, which by Japanese diagnostic criteria (8) was not compatible with PCOS. We re-evaluated the hormonal concentrations using a chemical luminescence enzyme immunoassay (CLEIA; IMMULYZE, Nippon Diagnostic Products Corporation, Tiba, Japan). Comparison of the results obtained by the two methods showed a broad disparity in LH concentrations which by RIA were 6.7 mIU/ml and by CLEIA were 23.0 mIU/ml. The FSH concentrations obtained by the two methods showed almost the same results (by RIA 8.3 mIU/ml and by CLEIA 6.9 mIU/ml). An LH-releasing hormone (LHRH) test (100 μg LHRH injected i.v.) showed a high response of LH measured by CLEIA (Fig. 1), leading to diagnosis of PCOS and variant LH-β. The ratio of the LH concentration found by RIA to that found by CLEIA was 0.29, suggesting that the patient was heterozygous for the mutant LH-β (9).

DNA analysis using PCR with direct sequencing by a method previously described (9) was employed. Genomic DNA was isolated from peripheral blood lymphocytes using Sepa Gene nucleic acid isolation reagents (Sanko Junyaku, Tokyo, Japan). The gene encoding the LH-β subunit was sequenced, and two base changes in the N-terminal region were identified. The first mutation (codon 8, TCG to CGG) changes tryptophan to arginine, and the second (codon 15, ATC to ACC) changes isoleucine to threonine. The patient's LH status represented variant and wild-type LH equally. No other infertility factors were recognized. She was treated with clomiphene citrate (Clomid; Shionogi Seiyaku, Osaka, Japan) for two courses of 100 mg/day for 5 days and two courses of 150 mg/day for 5 days without successful ovulation. Two cycles of controlled ovarian hyperstimulation using pure FSH (FERTINORM P; Serono Japan, Tokyo, Japan) were attempted, but she did not conceive and developed mild ovarian hyperstimulation syndrome which was treated in the outpatient clinic. Laparoscopy on 12 August 1997 showed polycystic ovaries measuring 4×3.5×3.0 cm on both sides (Fig. 2). The histological appearance of a biopsy specimen was

### Table 1 Hormonal examinations in the early follicular phase.

<table>
<thead>
<tr>
<th>Date</th>
<th>Substance (method)</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>16 August '95</td>
<td>LH (SPAC-S LH)</td>
<td>4.0 mIU/ml</td>
</tr>
<tr>
<td></td>
<td>FSH (SPAC-S FSH)</td>
<td>7.6 mIU/ml</td>
</tr>
<tr>
<td></td>
<td>PRL (SPAC-S PRL)</td>
<td>5.3 ng/ml</td>
</tr>
<tr>
<td></td>
<td>E₂ (coat-a-count E₂)</td>
<td>28 pg/ml</td>
</tr>
<tr>
<td></td>
<td>T (total testosterone kit)</td>
<td>33.6 ng/dl</td>
</tr>
<tr>
<td>25 February '96</td>
<td>E₁ (estrone test set)</td>
<td>99 pg/ml</td>
</tr>
<tr>
<td></td>
<td>E₂ (coat-a-count E₂)</td>
<td>45 pg/ml</td>
</tr>
<tr>
<td></td>
<td>T (total testosterone kit)</td>
<td>20.1 ng/dl</td>
</tr>
<tr>
<td></td>
<td>ASD (coat-a-count ASD)</td>
<td>2.4 ng/ml</td>
</tr>
<tr>
<td></td>
<td>DHEA-S (coat-a-count DHEA-S)</td>
<td>1830 ng/ml</td>
</tr>
</tbody>
</table>

E₂, estradiol; E₁, estrone; T, testosterone; ASD, androstenedione; DHEA-S, dehydroepiandrosterone-sulfate.

![Figure 1](image1.png)  
**Figure 1** An LHRH test (100 μg LHRH injected i.v.) showed an increased response of LH as measured by CLEIA.

![Figure 2](image2.png)  
**Figure 2** Laparoscopic appearance of enlarged polycystic ovaries. (A) right; (B) left.
compatible with PCOS. She underwent laparoscopic ovarian electrocautery.

Discussion

Controversy surrounds the diagnosis of PCOS, with different authors using differing criteria to define the syndrome. In the UK, polycystic ovaries detected on an ultrasonogram are generally accepted as the unifying diagnostic criterion. Additional biological abnormalities (elevated serum concentrations of LH, androgens, and/or insulin) may occur together with the characteristic sonographic appearance (10). However, the diagnosis of PCOS was originally based on clinical symptoms and biochemical parameters. Some investigators use a single biochemical marker such as elevated serum LH concentration (11) or high androgen levels (10) to define PCOS. Recent studies claim that there is a tendency to overestimate PCOS in the USA (12).

Japanese diagnostic criteria for PCOS (8) use a combination of ultrasound, clinical symptoms and an LH/FSH ratio >1.0. LH concentrations are an important consideration in diagnosing PCOS. An LH variant with two point mutations in the β-subunit gene was recently reported in Japan (5, 13).

In the present case, LH immunoassays using various antibodies gave different results. Ultrasensitive immunoassays for LH (1, 2), based on the sandwich technique and the use of monoclonal antibodies, have revealed some interesting aspects. The risk that monoclonal antibodies may be too specific has been pointed out (4), especially in cases where the protein or glycoprotein, like LH, is clearly microheterogeneous. With two anti-LH monoclonal antibodies directed to epitopes, only intact dimers were found which had restricted reactivity with LH at various rates (14) in a randomly selected population. The carrier frequency of the variant LH-β allele tended to be more common in populations from Western Europe as compared with those from Asia. Underestimation of LH concentrations in a serum sample can impair detection of the ovulatory LH peak or artefactually decrease the LH/FSH ratio, leading to failure to diagnose PCOS. Patients homozygous or heterozygous for the LH-β gene mutation were discovered by PCR and restriction fragment length polymorphism, an expensive approach not suitable for routine examination.

Nilsson et al. (14) performed two immunofluorometric assays using monoclonal antibodies: assay 1 recognized only wild-type LH, while assay 2 recognized variant and wild-type LH equally. In assay 1, the capture monoclonal antibody recognized a conformational epitope present in the intact α/β-LH dimer but not in the subunits, and the detection monoclonal antibody recognized the α subunit. Assay 2 used two LH-β-subunit-specific monoclonal antibodies. The ratio of results from assay 1 to those from assay 2 indicated the LH status: wild-type >0.9, heterozygous 0.2–0.9, and homozygous <0.15. Similarly, we established (9) a screening method for this genetic variant of LH, measuring hormonal concentrations by IMMULYSE LH (9), a random access instrument using recently developed CLEA technology. IMMULYSE LH is a solid-phase test which uses an alkaline phosphatase-conjugated polyclonal antibody as a tracer and polystyrene beads coated with monoclonal antibodies specific for LH. LH values measured by IMMULYSE LH ordinarily show a good correlation with SPAC-S LH, which uses RIA. SPAC-S LH, purchased from Daiichi Radioisotope Laboratory (Tokyo, Japan), uses an immunoradiometric assay method (1) and two different monoclonal antibodies, which react with the β subunit and intact dimer respectively. The relationship is linear: \[ y = 0.982x + 1.396, \ r = 0.970, \] where \( y \) is the LH value found by IMMULYSE LH, and \( x \) is the LH value found by SPAC-S LH. Furui et al. (5) have shown that measurement by SPAC-S LH is affected by point mutations of the LH-β gene. Therefore, we believe that the difference between tests might be due to poor recognition of LH in such patients by intact LH dimer monoclonal antibodies. Adding serum from SPAC-S LH, including LH coated with anti-LH β monoclonal antibodies, to the IMMULYSE LH preparation resulted in decreased LH values.

From these results, we concluded that our patient represented an immunologically anomalous LH variant. We have reported (9) that a ratio of SPAC-S LH to IMMULYSE LH of less than 0.5 or equal to 0.5 is abnormal, permitting recognition of PCOS occurring together with a point mutation of the LH-β gene. When a patient with bilateral polycystic ovaries by ultrasound, oligomenorrhea and hirsutism shows a normal LH/FSH ratio, the presence of a genetic LH variant should be suspected. Rajkhowa et al. (15) have reported that occurrence of the variant was not generally increased in women with PCOS and the presence of the variant did not alter the clinical or hormonal expression of the disorder in women with PCOS, but was over-represented among obese women with PCOS. Variant LH without PCOS has been associated with elevated total serum testosterone (16). The biological attributes of variant and wild-type LH appear to differ by in vitro bioassay and by circulatory half-time measurements (16). The in vitro bioactivity of the LH variant was significantly increased. The higher bioactivity but faster elimination of the variant LH could change the kinetics of its action in vivo. Connections between the mutation LH-β and infertility are not well known. The LH variant may contribute to some diseases affecting the pituitary–gonadal axis, although atypical clinical features were not seen in the present case of PCOS coexisting with the LH variant.
References


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