Circulating Levels of Progesterone as Biochemical Evidence of Ovulation is Method Dependent

IBRAHIM A. HASHIM, PhD, FACB1, and TARIF H. ZAWAWI, FRCPI, FACB2
Department of Pathology and Laboratory Medicine1, King Khalid National Guard Hospital, and Department of Medicine2, Faculty of Medicine and Allied Sciences, King Abdulaziz University, Jeddah, Saudi Arabia

ABSTRACT. Serum progesterone level is used as an indicator of ovulation and several assays are commercially available for its routine measurement. Target values suggestive of ovulation, deficient luteal phase and anovulation have been proposed and are widely sighted in the literature. In this study, we compared the use of two different assays namely: Amerlite (Johnson & Johnson, United Kingdom), and AxSym (Abbott Diagnostics, Germany) for measurement of serum progesterone. In patients being investigated for infertility (n = 93), serum progesterone levels showed good correlation (r = 0.92) between the two methods; however, values were significantly different (P < 0.005). Using published cut-offs levels for progesterone, 28.3%, 60.8%, and 10.9% of patients had ovulatory, anovulatory, and deficient luteal phase, respectively, as measured by the Amerlite assay, compared to 32.6%, 60.8%, and 6.5% when using the AxSym assay. Thus when using the AxSym assay, higher percentage of patients would be classified as having ovulatory cycle. This study indicated that methodological differences on their own right play a significant role in the determination of ovulatory status of patients. The study showed that laboratory-based target values need to be established.

Keywords: Progesterone, Ovulation

Introduction

Circulating progesterone level is used as an indicator of either natural or induced ovula-
tion and several assays are commercially available for its measurement. Disorders of ovulation including anovulation are common and account for 15-20% of patients where progesterone levels are usually low. Target values suggestive of ovulation, deficient luteal phase, and anovulation have also been proposed and are widely sighted in the literature.

Serial ultrasound monitoring and direct observation of ovulation at laparoscopy are used to indicate ovulation with the latter being used only in selected patients whereas ultrasound’s use being time consuming. Although basal body temperatures are used to predict ovulation, it is not reliable.

Serum progesterone measurement is used in fertility studies to assess corpus luteum function, to evaluate placental function during pregnancy, and as an aid in confirming ovulation, it is in the latter that its measurement is widely employed[1].

A number of assays are commercially available and it has been reported[2] that serum progesterone levels greater than 30 nmol/L, between 30 and 18 nmol/L and less than 18 nmol/L were consistent with ovulation, deficient luteal phase, and anovulation, respectively. However, different workers have chosen very different target values to indicate ovulation, varying from 3.2 to 30.0 nmol/L. A large emphasis has been placed on the clinical criteria used in collection time and little was attributed to the now improved progesterone immunoassays as a cause of such wide discrepancies.

This study compared the use of two different progesterone assays in the investigation of ovulation and examined if methodological differences could on their own contribute to discrepancies in patients’ classification.

Materials and Methods

Subjects: Patients (n = 93) being investigated for infertility were entered into the study. Patients age ranged from 18 to 40 years old (median = 26 years old). Blood samples were collected on the 21st day of the menstrual cycle following hospital ethical and research committee approval.

Biochemical Measurements: Serum was separated and stored at -20° C until analysis. Progesterone levels were determined using two different immunoassays: namely; Amerlite immunoassay (Johnson and Johnson, Clinical Diagnostics, Amerham, U.K.) and AxSym immunoassay analyser (Abbott Diagnostics, Germany). Analysis was performed according to the manufacturers’ instructions. Statistical analysis was performed using computer software. Inter- and inter assay imprecision were determined for both assays at three different serum progesterone levels. Linearity studies were performed by serially diluting six samples with 0 nmol/L standard of the Amerlite assay and with the AxSym assay-diluent and assaying for progesterone levels using both assays, respectively.
Results

Serum progesterone levels ranged from 0.2 to 76.6 nmol/L (median 3.9) in the Amerlite assay compared to 0.3 to 103.4 nmol/L (median 4.5) in the AxSym assay. There was good correlation between both assays \((r = 0.92)\) (Fig. 1). However, the data was not normally distributed and Wilcoxon’s non-parametric test showed significant difference between results obtained by the two methods \((P < 0.005)\). Higher values were observed in the AxSym assay (Fig. 2).

Fig. 1 Correlation between serum progesterone results obtained by Amerlite and AxSym immunoassays.
Fig. 2 Patients with ovulation, anovulation and deficient luteal phase as indicated by serum progesterone concentration measured using the two immunoassays.
Using the above progesterone “cut-off” levels\textsuperscript{[11]}. 28.3\%, 60.8\%, and 10.9\% of patients had ovulatory, anovulatory cycles, and efficient luteal phase, respectively, as measured by Amerlite assay as compared to 32.6\%, 60.8\%, and 6.5\% when using the AxSym assay, respectively.

Inter-assay imprecision ranged from 25.0\%, 5.0\%, and 8.5\% at 1.1, 15.8, and 65.0 nmol/L, respectively, in the Amerlite assay as compared to 9.2, 6.1, and 5.9 at 3.2, 17.5, and 69.9 nmol/L, respectively, in the AxSym assay.

Linearity studies were performed and showed good recovery and parallel responses (data not shown).

**Discussion**

This study compared the use of two different immunoassays for serum progesterone measurement. Assays were based on competitive binding between serum progesterone and assay progesterone. In the Amerlite assay, horseradish peroxidase-conjugated progesterone competes with serum progesterone for anti-progesterone antibody, the latter is captured by a second antibody coated onto a 96 well-plate, whereas, in the AxSym assay, immobilized progesterone competes with serum progesterone for anti-progesterone antibody labeled with alkaline phosphates. The final signal in both assays is inversely proportional to the concentration of serum progesterone.

The Amerlite assay had been calibrated against a pure progesterone preparation and values were checked to relative to Gas-chromatography-Mass Spectrometer (GCMS) calibrated samples. The standard used is different from that in the Abbott assay; however, it was said to have good correlation with GCMS results (r = 0.994, Abbott Diagnostics, personal communication). The Amerlite immunoassay showed poor precision at low progesterone levels (< 15 nmol/L), whereas, both immunoassays showed good precision at higher progesterone levels.

Although results obtained by both assays showed good correlation, results were significantly different. It is possible that the nature of both assays and the presence of competitors / inhibitors in serum may contribute to the discrepancy. Reported cross-reactivity studies by both manufacturers were comparable.

Patients classified as having anovulatory cycles were the same by both assays; however, more patients had higher progesterone results by AxSym immunoassay and thus were classified as having ovulatory cycles.

The study highlighted that, in addition to the published discrepancy in serum progesterone “cut-off” limits, methodological differences play a significant role and that further studies are required to investigate such discrepancy. Laboratory based progesterone target values should, therefore, be established.
References


دراسة لمستوى هرمون البروجسترون في مصل الدم
واستخدام القيم كمؤشر حدوث التبويض

إبراهيم هاشم وطريف هشام زواوي
قسم علم الأمراض وال킹بريات الطبية، كلية الطب والعلوم الطبية،
جامعة الملك عبد العزيز
جدة - المملكة العربية السعودية

المستخلص. هناك عدة وسائل تحليلية متوفرة تجاريا وتستخدم روتينيا لقياس مستوى هذا الهرمون. أما النسب والمقدار المستهدفة التي تدل على حدوث التبويض، أو نقص الطور النموتي (الطور الأصفر)، أو عدم التبويض فإنه يأتي ذكرها يتوسع في الأدبيات المنشورة. وفي دراستنا هذه قمنا بإجراء مقارنة للنتائج المستخلصة من استخدام نوعين من الوسائل التحليلية وهما:

1- أمولاتي (جونسون آند جونسون) - المملكة المتحدة
2- أكسيم (أبوت ديانستك) - ألمانيا

ولقد تم قياس مستوى البروجسترون لثلاثة وتسعين مريضة بشكين من تأخر الحمل أو العقم ووجدنا أن هناك ترابط جيدا بين الطريقتين (R=0.92) على الرغم من الاختلاف الهام بين القيم (P<0.005). واستنادا على قيم البروجسترون المشورة وجدنا أنه عند استخدام وسيلة أمولاتي (Amerlite) للمتالك كانت النتائج كالتالي:

1- (3,6٪) من المرضى دلت النتائج على حدوث التبويض لديهن.
2- (8,6٪) من المرضى دلت النتائج على عدم حدوث التبويض لديهن.
3- (9, 10%) من المريضات دلت النتائج على قصور في الطور اللوبتي (الطور الأصفر - Luteal phase - لديهن.
أما في حالة استخدام وسيلة أكسيم (AxSym) فقد كانت النتائج كالآتي:
1. (6, 2%) من المريضات في حالة التلوث
2. (8, 20%) من المريضات على عدم التلوث
3. (5, 6%) من المريضات لديهن قصور في الطور اللوبتي.

ومن ذلك نخلص أنه باستخدام وسيلة أكسيم نحصل على نسبة أعلى لحدوث التلوث لدى المريضات. وعلى هذا فإن هذه الدراسة مؤشر على أن اختلاف الطرق المستخدمة في الفحوصات يلعب دوراً كبيراً في تحديد فترة التلوث لدى المريضات. ولذا فإن كل مختص أن يحدد النسب والقيم المستهدفة الخاصة به.