Testosterone or Dehydroepiandrosterone Sulfate as a Biomarker for Hirsutism in Women with Polycystic Ovary Syndrome

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Hirsutism is a distressing medical problem for women. Most of hirsutism in women is associated with excess androgen, and most cases have PCOS as an underlying cause. Which androgen to be used to evaluate clinical or biochemical hyperandrogenism in women with PCOS is still debated. There are a small number of studies that evaluated androgens in women with PCOS having hirsutism with conflicting results. The objective of this study was to determine which androgen predict hirsutism in women with polycystic ovary syndrome (PCOS). A case-control study was done in Faiha Specialized Diabetes, Endocrine, and Metabolism Center (FDEMC), Basrah, Iraq. A total of 130 women with PCOS (based on Rotterdam criteria) and 70 healthy controls of comparable age (16-40 years) were investigated for androgens (total testosterone, free testosterone, DHEA-S) using Electrochemiluminescence technology assay; excess hair was examined according to the modified Ferriman-Gallwey (mFG) score and a cut-off value of 8 defined hirsutism. In the three groups of women, the first (n=100) included PCOS with hirsutism, the second (n=30) PCOS without hirsutism, and the third (n=70) women without PCOS or hirsutism as healthy control, hirsutism was seen in about 77% of PCOS women mostly of moderate severity; High TT, FT, DHEA-S, and overall androgens were seen in 69%, 76%, 37%, and 99% respectively of our PCOS women with hirsutism. No correlation was found between TT, FT, and DHEA-S and the mFG score. This study provides evidence that presence of hirsutism in women with PCOS was associated with a higher level of biochemical hyperandrogenism than seen in PCOS without hirsutism; however, there was no correlation between the studied androgens and mFG score.

Keywords: Polycystic Ovary Syndrome, biochemical hyperandrogenism, testosterone, DHEA-S, hirsutism.

Hirsutism is a distressing medical problem for women. Overall, it affects about 5-15% of women¹, it is usually associated with cosmetic concern, emotional distress, and depression², and is typically a feature of an underlying condition mostly of hormonal disturbance origin³. Hirsutism can be described as terminal (thick and dark) hair in the face and/or body of women in areas usually having little or no hair¹. In 1981, Hatch, Rosenfield, Kim, and Tredway revised the initial visual score that introduced in 1961 by Ferriman and Gallwey, where nine body areas used, each is graded from...
nil (no substantial terminal hair growth) to four (considerable terminal hair growth) and then added up to a total score of the modified Ferriman-Gallwey (mFG), which is the standard scoring scale for hirsutism currently in use.

Most of hirsutism in women is associated with excess androgen (up to 85%), and most cases (around 70 – 80%) have PCOS as an underlying cause. Other causes of hirsutism constitute about 10 – 15% and include idiopathic hirsutism, thyroid dysfunction, hyperprolactinemia, congenital adrenal hyperplasia, Cushing disease, acromegaly, ovarian and adrenal tumors.

PCOS is an endocrine condition observed during the fertile life of females, with a 6 – 15% prevalence that varies according to the diagnostic criteria used. Hyperandrogenism, chronic anovulation, and polycystic ovarian morphology (PCOM) are characteristic of PCOS.

The European Society for Human Reproduction and Embryology (ESHRE) and the American Society for Reproductive Medicine (ASRM) have proposed the PCOS diagnosis needs two out of the following three features: the Rotterdam criteria:

- Ovulatory dysfunction refers to oligomenorrhea (cycles approximately 35 days apart or nine cycles or less per year) or amenorrhea (absence of menstruation 6 to 12 months after a cycle).
- Clinical or biochemical hyperandrogenemia.
- Polycystic ovaries (as described on ultrasonography): 12 or more follicles in at least one ovary with a diameter of 2-9 mm and/or a total volume of an ovary greater than 10 mL.

Before diagnosing PCOS, disorders can lead to ovulatory dysfunction such as hyperprolactinemia and thyroid dysfunction, and those with androgen excess, including non-classic congenital adrenal hyperplasia (NC-CAH), should be omitted. In women, the ovaries release androstenedione, T, and dehydroepiandrosterone (DHEA) under the influence of pituitary luteinizing hormone (LH); the adrenal glands produce androstenedione, T, DHEA, and dehydroepiandrosterone sulfate (DHEA-S) under the control of pituitary adrenocorticotropic hormone (ACTH). In peripheral tissues (liver, skin, and adipose tissue), the androstenedione and DHEA (ovaries and adrenals), and DHEA-S (adrenals) can be converted into T. Approximately 98 – 99% of circulating T is bound to sex hormone-binding globulin (SHBG) and albumin; the remainder (1–2%) is the free testosterone (FT), which is bioactive androgen that will be converted into DHT by the 5 alpha-reductase enzymes in the target tissue. This DHT has a higher affinity for androgen receptors found in the dermal papilla cells and the hair follicle’s outer root sheath to produce longer thick hair.

The suggested fundamental physiological abnormality in women with PCOS is the increased intensity and frequency of the LH pulse caused by increased pulsatile gonadotropin-releasing hormone (GnRH) secretion, resulting indirect stimulation of ovarian theca cells to generate androgens. In 2008, the Endocrine Society proposed testing women with abnormally high mFG score for elevated androgen levels. In 2012, the Androgen Excess and Polycystic Ovary Syndrome Society (AE-PCOS) also recommended hormonal analysis for women with a high hirsutism score. There are a small number of studies that evaluated androgens in women with PCOS having hirsutism with conflicting results. The study aims to investigate TT, FT, and DHEA-S as biochemical markers for hirsutism in women with PCOS.

Patients and method

Design and participants

A case-control study was conducted on 200 women of reproductive age (16-40 years) at Faiha Specialized Diabetes, Endocrine, and Metabolism Center (FDEMC) in Basrah, Iraq from September 2019 to September 2020. It was composed of three groups: the first group (n=100) included women with PCOS and hirsutism, the second group (n=30) women with PCOS but no hirsutism, and the third group (n=70) women without PCOS or hirsutism as healthy control. Basrah College of Medicine Ethics Committee had approved the study protocol, and participants gave written informed consent. Pregnant and lactating women were excluded as patients with liver and kidney diseases.

Clinical evaluation

Participants medical history focused on age at menarche, childbearing history, menstrual cycle length, and regularity, presence of galactorrhea, the appearance of excess hair in the face and body, and
the use of medications that may affect hormone parameters during the last three months prior entering the study like oral contraceptive pills and glucocorticoids.

Bodyweight (kg) and height (cm) were measured, and body mass index (BMI) was calculated by the Quetelet’s Index formula. Physical examination included assessing the amount and distribution of terminal hair according to the mFG score by the same trained nurse. Women who had removed the excess hair were informed to come later for a more reliable hair score assessment (12 weeks after laser therapy, four weeks after depilation or waxing, and five days after shaving). PCOS diagnosis was proposed by the Rotterdam criteria. Ovulatory dysfunction was labeled as oligomenorrhea (cycles e” 35 days apart or d”9 cycles per year) or amenorrhea (absence of menses for 6 -12 months after an established cycle). The study defined hyperandrogenism clinically by the presence of hirsutism with the mFG score equal to or more than 8. PCOM was determined by transabdominal pelvic ultrasonography. The diagnosis of PCOS was only made after excluding hyperprolactinemia, thyroid dysfunction, Cushing’s syndrome, and NC-CAH by relevant clinical and laboratory analysis, including prolactin (PRL), thyroid-stimulating hormone (TSH), cortisol, ACTH, and 17-hydroxyprogesterone (17-OHP).

**Laboratory tests**

**Basal hormone measurement**

On the second to the seventh day of menses (for women with regular menstrual cycles or at any day for those with irregular ones), fasting venous blood samples (4 mL in clot-activator gel tube and 3 mL in EDTA tube) were collected at 8:00-10:00 am. After separation by NUVE-NF 800 at 4100 RPM, TT, DHEA-S, PRL, SHBG, TSH, cortisol, ACTH, and albumin were tested by Electrochemiluminescence (ECL) technology assay (Cobas e 411 analyzer-Roche, Germany). 17-OHP was measured by Enzyme-linked immunosorbent assay (ELISA) method (BioTek-USA). Our study normal androgens level were 15-46 ng/dL for TT (d’ 6% CV), 0.13-0.56 ng/dL for calculated FT (<6% CV), and 18-86 nmol/L for SHBG (< 4% CV)(17). FT was calculated from TT, SHBG, and albumin, according to the Vermeulen equation(18). Reference ranges for other hormonal parameters used were those of

![ROC Curve](image)

**Fig. 1.** ROC of androgens in women having PCOS with hirsutism. ROC, receiver operating characteristic curve; TT, total testosterone; FT, free testosterone; DHEA-S, dehydroepiandrosterone sulfate.
the FDEMC laboratory; DHEA-S 145-395 µg/dL (<5%CV) for women age 18-19, 65-380 µg/dL for women ages 20-29, 45-270 µg/dL for women ages 30-40, PRL 4-30 ng/mL (<5%CV), ACTH 10-60 pg/mL (<5%CV), and 17OHP 1-495 ng/dL (<10%CV).

**Ovarian ultrasonography**

The same radiologist examined the participants for PCOM using a 4.0 MHz transabdominal transducer. The presence of PCOM was based on visualization of ≥12 follicles in one ovary measuring 2-9 mm in diameter over the peripheral stroma, and/or increased an ovarian volume of 10 mL or more at day two-seven of menstrual cycle or at any day for those with irregular menses. In our study, the ultrasonic examination of the pelvis was transabdominal, not the trans-vaginal one.

**Statistical analyses**

Statistical analysis was done by Statistical package for social science software (SPSS Inc. Chicago, IL, USA) version 25. Using the Kolmogorov–Smirnov method, we tested the distribution of all continuous variables. Since the data were normally distributed, continuous variables were presented as mean ± standard deviations. Different study group’s means were compared using the ANOVA with post hoc. Categorical variables were summarized in number and percentage. A Chi-square test and Fisher’s Exact test were used to compare categorical groups. Correlations between continuous variables were analyzed with Pearson’s correlation. P-value <0.05 was considered statistically significant.

**RESULTS**

The features of the 200 participants included in the study are presented in Table 1. The mean age (±SD) for the first group (women with PCOS and hirsutism) was 24 (±6), for the second group (women with PCOS without hirsutism) was 26 (±8), and for the third group (women without PCOS or hirsutism) was 26 (±7) with no significant difference among the groups (p=0.17), so the three groups of participants were age-matched. BMI for the first group was 32 (±5.9), for the second group was 27.4 (±3.5), and for the third group was 26.9 (±5.7), with a significant statistical difference of the first group from the second and third (p<0.001), and between the first and third groups (p<0.001), but there was no significant difference between the second and third groups (p=0.6). Hirsutism score

<table>
<thead>
<tr>
<th>Variables</th>
<th>PCOS with hirsutism n=100</th>
<th>PCOS with no hirsutism n=30</th>
<th>no PCOS, no hirsutism n=70</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>24±6</td>
<td>26±8</td>
<td>26±7</td>
<td>0.17</td>
</tr>
<tr>
<td>BMI (kg/m²)</td>
<td>32±5.9</td>
<td>27.4±3.5</td>
<td>26.9±5.7</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>mFG score</td>
<td>19±5</td>
<td>5±2</td>
<td>3±2</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Table 1. Age, BMI and hirsutism score of study groups n=200

<table>
<thead>
<tr>
<th>Androgen</th>
<th>PCOS with hirsutism n=100</th>
<th>PCOS with no hirsutism n=30</th>
<th>no PCOS, no hirsutism n=70</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>TT (ng/dL)</td>
<td>50.2±10</td>
<td>39.3±6.3</td>
<td>23.7±7.6</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>FT (ng/dL)</td>
<td>1.04±0.29</td>
<td>0.43±0.07</td>
<td>0.33±0.1</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>DHEA-S (µg/dL)</td>
<td>323.3±99.5</td>
<td>209.8±68</td>
<td>200.6±63.3</td>
<td>&lt;0.001</td>
</tr>
</tbody>
</table>

Table 2. The mean of androgens in study groups n=200

PCOS, polycystic ovary syndrome; SD, standard deviation; BMI, body mass index; mFG, modified Ferriman-Galloway.
for the first group was 19 (± 5), for the second group was 5 (± 2) and for the third group was 3 (± 2), with a significant statistical difference of the first group from the second and third (p <0.001), and between the first and third groups (p <0.001), and there was a significant difference (p < 0.01) between the second and third groups.

The mean TT (±SD), for the first group, was 50.2ng/dL (±10), for the second group was 39.3ng/dL (±6.3) and for the third group was 23.7ng/dL (±7.6). The mean FT (±SD), for the first group, was 1.04ng/dL (±0.29), for the second group was 0.43ng/dL (±0.07) and for third one was 0.33ng/dL (±0.10). The mean DHEA-S (± SD), for the first group, was 323.3µg/dL (±99.5), for the second group 209µg/dL (±68), and the third group was 200.6µg/dL (±63.3), as shown in Table 2.

The post hoc tests of mean androgen comparison of the study groups revealed the following: for TT, there was a significant difference between the first group and the second group (<.001), and between the first group and the third one (<0.001). Also, there was a significant difference between the second group and the third one (<.001).

Regarding FT, again, there was a significant difference between the first group and second group (<0.001), and between the first group and the third one (<0.001), but there was no significant difference between the second and third groups (p=0.52).

In the case of DHEA-S, there was a significant difference between the first group and second group (<0.001) and between the first group and the third one (<0.001). But, no significant difference between the second and third groups (p=0.6).

The number of participants with high androgen level is shown in Table 3. In the group of women with PCOS and hirsutism, there were 69 patients (69%) with high TT, 76 patients (76%) with high FT, and 37 (37%) patients with high DHEA-S. In women with PCOS and no hirsutism, three patients (10%) had high TT, and ten patients (33.3%) had high FT, while none of them had high DHEA-S level. In the group of women without PCOS or hirsutism, none of the participants had an abnormally high TT or FT or DHEA-S. High overall androgens was seen in 84.61 % of women with PCOS (n=130), in 99 % of the women with hirsutism, and 36.7% of women with PCOS but no hirsutism enrolled in the study.

### Table 3. High Androgens in Participants n=200

<table>
<thead>
<tr>
<th>Androgens level</th>
<th>PCOS with hirsutism n=100</th>
<th>PCOS with no hirsutism n=30</th>
<th>PCOS, no hirsutism n=70</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
<td>n</td>
<td>%</td>
</tr>
<tr>
<td>High TT</td>
<td>69</td>
<td>69</td>
<td>3</td>
<td>10</td>
</tr>
<tr>
<td>High FT</td>
<td>76</td>
<td>76</td>
<td>10</td>
<td>33.3</td>
</tr>
<tr>
<td>High DHEA-S</td>
<td>37</td>
<td>37</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>High overall androgens</td>
<td>99</td>
<td>99</td>
<td>11</td>
<td>36.7</td>
</tr>
</tbody>
</table>

PCOS, polycystic ovary syndrome; TT, total testosterone; FT, free testosterone; DHEA-S, dehydroepiandrosterone sulfate.<sup>a</sup>for high TT levels, there was a significant difference between first and second groups (p <0.001). The significance between other groups was not applicable.<sup>b</sup>for high FT levels, there was a significant difference between first and second groups (p <0.001). The significance between other groups was not applicable.<sup>c</sup>for high DHEA-S levels, statistical deference calculation was not applicable.<sup>d</sup>for any biochemical hyperandrogenism, there was a significant difference between first and second groups (p <0.001). The significance between other groups was not applicable.

### Table 4. Pearson correlation between androgen parameters (TT, FT, and DHEA-S)

<table>
<thead>
<tr>
<th>Variable</th>
<th>r statistics</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>TT</td>
<td>0.099</td>
<td>0.32</td>
</tr>
<tr>
<td>FT</td>
<td>0.072</td>
<td>0.47</td>
</tr>
<tr>
<td>DHEA-S</td>
<td>0.202</td>
<td>0.053</td>
</tr>
</tbody>
</table>

r, Pearson correlation coefficient; TT, total testosterone; FT, free testosterone; DHEA-S, dehydroepiandrosterone sulfate.
For women with PCOS who had hirsutism in the study group, the severity of hirsutism was mild in 29 patients (29%), moderate in 59 patients (59%), and severe in 12 patients (12%).

Figure 1. shows the receiver operating characteristic (ROC) curves for androgen marker; the cut-off values determined from the ROC curve together with diagnostic sensitivity and specificity.

In our patients, TT showed a ROC curve of a good accuracy (area under the curve AUC = 0.944, SE = 0.013, 95%CI (0.918 – 0.970) and $p<0.001$. A TT of 44.3 ng/dL or more had 78% sensitivity and 90% specificity for hirsutism in women with PCOS.

The FT showed a ROC curve with excellent accuracy (area under the curve AUC =1.000, SE=0.0001, 95%CI (1.000-1.000) and $p<0.001$. An FT of 0.54ng/dL or more had 100% sensitivity and 85% specificity for hirsutism in women with PCOS in the study sample.

While, DHEA-S showed a ROC curve of a good accuracy (area under the curve AUS= 0.839, SE=0.026, 95%CI (0.788-0.891) and $p<0.001$. A DHEA-S of229 µg/dL or more had 81% sensitivity and 70% specificity for hirsutism in women with PCOS in the study sample.

The correlation between biochemical markers and hirsutism is illustrated in table 4. A negligible correlation was noted between the hirsutism score and measured androgens (TT, FT, and DHEA-S).

**DISCUSSION**

Hirsutism is usually a sign of different medical conditions, and PCOS accounts for about 80% of them(1), where it is considered a useful marker of biochemical hyperandrogenism(5). In this study, we investigated biochemical markers in women with PCOS to predict hirsutism. The mean age of the enrolled women with PCOS and hirsutism was 24±6 years, which coincided with the finding of a previous study from the same center on women in Basrah in 2017(20). There was a significant difference in BMI between the group of women with PCOS and hirsutism and other groups (women with PCOS without hirsutism and women in the control group). A similar finding of a significantly higher BMI in 409 women with PCOS than 7057 women in the non-PCOS group was seen in a population-based observational study in 2013(21). The high prevalence of obesity in our data among women with PCOS may represent an overall obesity trend in our locality population(22).

The mean mFG score for enrolled women in different study groups was comparable to that seen by an Italian study in 2006(23); the difference in means of hirsutism score between the groups of women with no hirsutism in the study although it was statistically significant, but clinically was not significant as the scores were below the cut-off that defined hirsutism used during enrollment.

We observed higher mean values for TT, FT, and DHEA-S in PCOS groups than non-PCOS, which was also seen in Turkish and Italian studies in 2011 and 2016(24, 25).

Women with hirsutism had significantly higher androgens mean values than women without hirsutism, a finding similar to a study from the Aristotle University of Thessaloniki, Greece, in 2013(26).

The high androgen levels seen in women with PCOS were comparable to a study by Chang and colleagues in 2005, where TT, FT, and DHEA-S were high in about 50%, 88%, and 34% of women with PCOS, respectively(11), and to Ibanez and associates study in 2017, where about 70% of women with PCOS have elevated FT, and 20 - 30% have high DHEA-S(27).

Women with hirsutism in our data had higher TT, FT, and DHEA-S levels, which were higher than those seen by some authors(26, 29); the different biochemical assay methods used for androgen estimation and different cut-off values used in the study can explain the diversity in androgens levels.

In a systematic review published in 2006, biochemical hyperandrogenism was found in approximately 60-80% of women with PCOS, which was comparable to our result(30).

The documented strong association between hirsutism and biochemical hyperandrogenism(5) was also seen in our study, where overall high androgens noted in almost all women included with hirsutism.

Hirsutism was identified in about three-quarters of women with PCOS in our study, and this was consistent with similar studies from Saudi Arabia and Turkey(12, 31). Hirsutism was mild in one-third of patients, moderate in about two-thirds, and...
severe in less than one-fifth of patients, which was also seen by Chhabra and co-workers in 2012 who founded mild hirsutism in 32.5%, moderate scores in 52.5%, and a severe degree in 15%\(^3\).

Although the DHEA-S role in PCOS is not yet well clarified, several studies have documented high DHEA-S in about 20-30 percent of patients with PCOS, which was consistent with our findings. However, this does not necessarily reflect an excess adrenal function; it may result from DHEA conversion or represent an inherited genetic abnormality, so interpretation needs caution\(^3\).

An Italian cross-sectional study in 2016 showed that the percentages of women with PCOS and hirsutism who had elevated DHEA-S were higher than women with PCOS and no hirsutism with even increased serum mean values as established in our study\(^24\). The current study cut-off values for TT, FT, and DHEA-S were different from those proposed by the University of Duisburg-Essen in 2007 and a Chinese cross-sectional in 2012\(^34, 35\). This difference in cut-off values can be explained by either different assay methods for androgens measure mentor various PCOS phenotypes.

No significant correlation was reported in this study between the androgens (TT, FT, and DHEA-S) and hirsutism score, as was seen in several other studies\(^13, 23, 24\), and this can be explained by the fact that the local androgens level, the sensitivity of the pilosebaceous units, and the activity of androgen receptor are the principal determining factors for hirsutism severity rather than the level of circulating androgens\(^36\).

The study limitations were recruitment of women was from a tertiary referral center and thus may not sufficiently represent the general population, and TT was measured by ECL rather than the liquid chromatography with tandem mass spectrometry method.

**CONCLUSION**

since it demonstrated the highest sensitivity and specificity in the study, we recommend FT as a valuable marker for hirsutism in women with PCOS, even though we couldn’t find a correlation between any of the studied androgens with hirsutism severity.

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**Conflict of Interest**

None.

**Funding Source**

None.

**REFERENCES**


