Impact of Cyproterone Acetate and Ethinylestradiol on Clomiphene Resistant PCOS Patient with High Serum AMH level

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Abstract:

Background: Polycystic ovary syndrome (PCOS) is the most common endocrinopathy in women of reproductive age. PCOS is characterized by hyperandrogenism and oligomenorrhea, and Polycystic ovary. It is a common cause of female subfertility. Anti-mullerian hormone (AMH) is 2-4 times higher in patients with PCOS compared to normal subfertility patient.

Objective: To assess the impact of metformin and oral contraceptives (OCs) containing cyproterone acetate and ethinyl estradiol on serum anti-Mullerian hormone (AMH) levels, with CC resistant polycystic ovary syndrome (PCOS)

Methods: This prospective randomized controlled trial was conducted at Bangabandhu Sheikh Mujib Medical University (BSMMU), Dhaka during the period of January 2019 to June 2019. Already diagnosed patients of polycystic ovary syndrome with subfertility with high serum AMH level more than 5 ng/ml attending in the OPD of Reproductive Endocrinology and infertility department at BSMMU are the study population. Total 60 subjects fulfilling rotardum criteria and who are treated with 3 cycle CC with 100mg/day for 5 days from day 2 of menstrual cycle but fail to ovulate were recruited for the current study. Those subjects with tubal, and male factors excluded from the study. After enrollment of the patients, randomization will be carried out by blocked randomization method to divide them equally into two groups-Group A number of patients 30 (treated with 2mg cyproterone acetate (CPA) and 35 mcg of ethinylestradiol (EE) and Group B number of patients 30 treated with Metformin). Group A patients will receive treatment with 2mg cyproterone acetate (CPA) and 35 mcg of ethinylestradiol (EE) for 6 months and Group B patients will receive treatment with Metformin for 6 months. After enrollment of the patients, base line hormonal assessment basal level of FSH, LH, thyroid-stimulating hormone (TSH), prolactin (PRL), and testosterone hormones were registered and metabolic assessment included OGTT and fasting insulin level were done, and to identify the changes in clinical and biochemical profile of study population. Clinical evaluation, by history and examination, transvaginal ultrasonography of both group and changes of the above mentioned parameters of both group were recorded after 6 month of completion of treatment.

Results: A Changes in clinical parameters (Acne, BMI, Menstrual cycle) was recorded every three monthly for 6 month. Changes of biochemical parameters (serum LH (D2), serum FSH (D2), serum testosterone, OGTT and fasting insulin), serum AMH were recorded every three monthly for 6 month. Changes of sonographic parameters (ovarian volume, Antral follicular count development, was done monthly for 6 cycles. AMH levels are significantly reduced under treatment with OCP containing 35mg ethinylestradiol plus 2 mg cyproteroneacetate than metformin. AMH was significantly decreased after treatment with 35 mg ethinylestradiol plus 2 mg cyproteroneacetate (p < 0.001 at 6 months), and treatment with metformin did not significantly affect serum AMH levels. AMH was significantly decreased after treatment with metformin (p < 0.005). Serum LH was and OGTT was significantly decreased in both treatment p<0.001.

Conclusion(s): AMH serum levels were significantly decreased under treatment with 35mg ethinylestradiol plus 2mg cyproterone acetate, due to decrease in androgens and suppression of gonadotrophins (LH).

Key Words: Polycystic ovary syndrome, Hyperandrogenism, Anti-Mullerian Hormone, Oral contraceptives, Metformin effect of cyproteroneacetate.

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Introduction:
Poly cystic ovary syndrome (PCOS), a multifactorial disorder, is the most common female endocrinopathy of reproductive age group as well as a major cause of anovulatory infertility. It is a heterogeneous condition and affects about 5-10% of female populations [Qiao et al. 2011; Durmont et al. 2015]. Currently, it is thought that PCOS emerges from a complex interaction of genetic and environmental traits [Williams et al. 2016]. It is characterized by defective follicular development, maturation and ovulation and also hormonal deregulation including LH, AMH and/or androgen hypersecretion. The classic symptoms of the disease are due to hyperandrogenism and chronic anovulation. Both of which have substantial psychological, social and economic consequences: Sirmans et al. 2014; Ruan et al. 2017. PCOS is characterized by hyperandrogenism (hirsutism and/or biochemical hyperandrogenemia) and oligo/anovulation, and is also highly associated with obesity and insulin resistance (IR). Anti-Müllerian hormone (AMH) is a member of the transforming growth factor-b (TGF-b) superfamily of glycoproteins that has been found to play an important role in chronic anovulation by inhibiting the initial recruitment of primordial follicles and by promoting follicular arrest. Serum AMH level is 2-4 fold higher in women with PCOS than in healthy women; Villarroel et al. 2011. It is due to increased production of AMH per follicle, not due to increased follicular pool. Cyproterone Acetate plus Ethinylestradiol (CPA/EE) is an oral contraceptive pill (OCP) which contains synthetic estrogen ‘Ethinylestradiol’ and antiandrogen ‘Cyproterone Acetate’. Have traditionally been the first choice for management of PCOS. [Yildiz et al. 2008; Buzney et al. 2014]. CPA/EE are considered to treat androgen dependent conditions i.e. acne, hirsutism and androgenic alopecia, but not indicated solely for contraceptive purpose [Soriano et al. 2016]. Ruan et al. (2017) reviewed that short-term pretreatment using CPA/EE may increase the fertility and improve pregnancy outcome by reducing serum AMH level.

It has been shown that, serum AMH is a reliable diagnostic tool for PCOS [Jacob et al. 2017] than antral follicle count (AFC) [Casadei et al. 2013] as well as useful to establish treatment protocol and to define the best strategy for ovulation induction in infertile women with PCOS [Agathe et al. 2015].

Women with PCOS are usually treated with an oral contraceptive (CPA/EE), while both lean and obese PCOS with IR might benefit from treatment with metformin. Treatment with CPA/EE s is known to normalise menstrual function and to ameliorate hirsutism and acne, reduces serum AMH level on the opposite, treatment with metformin is beneficial for weight reduction and correction of IR, still the effect on menstrual cycle and hyperandrogenism is rather weak. Few data are available on the impact of the treatment modalities on serum AMH levels in women with PCOS. The aim of the present study was to assess the effect of cyproterone acetate and ethinyl estradiol on serum AMH levels in a well characterized cohort PCOS women in comrison with metformin. Metformin administration in patients with PCOS exerts a differential action on the ovarian AMH levels on the basis of ovulatory response. Changes in AMH levels in antral follicular fluid during metformin treatment could be involved in the local mechanisms mediating the ovulatory restoration. Recently different studies shown that, OCP reduces the AMH level [Kalio et al. 2013]. This could be explained by suppression of LH secretion by OCP. As, luteinizing hormone (LH) can be responsible for the over expression of AMH and AMH specific type 2 receptors (AMHRII) in lutein GCs in anovulatory PCOS. [Pellat et al. 2007; Catteau-Jonard et al. 2008]. So, reduced LH may directly impact on AMH secretion. [Van den Berg et al. 2010]. There is an increased number of pre-antral and antral follicles in the polycystic ovary, many of which individually produce increased amounts of anti-Müllerian hormone (AMH) compared with those in the normal ovary. It is hypothesized that the high AMH concentrations present in women with PCOS play an integral role in causing anovulation due to its inhibitory influence on the actions of follicle-stimulating hormone, which normally promotes follicular development from the small antral to the ovulatory stage. Further indirect evidence to strengthen this hypothesis can be gathered from the practice of ovulation induction. Those with very high concentrations of circulating AMH are less likely to respond with ovulation to treatment with weight loss (Moran et al., 2007), clomiphene citrate or laparoscopic ovarian drilling (Amer et al., 2009). A possible reason that the metformin is less effective and certainly slower than clomiphene in inducing ovulation is the fact that AMH levels only decrease very slowly during this treatment (Fleming et al., 2005; Fabreques et al., 2011). Finally, AMH concentrations decrease during successful low-dose FSH therapy (Catteau-Jonard et al., 2007). Together, this clinical evidence accentuates the fact that the greater the number of antral and pre antral follicles, the higher the AMH concentrations, the more severe the symptoms of PCOS. Panidis et al. (2010) showed that CPA/EE significantly reduces serum AMH level by decreasing androgens and suppression of Serum LH compared to metformin. Branigan and Estes (2003) found that 2 months of CPA/ EE use before repeating CC treatment in CC resistant women produced effective ovulation and pregnancy rates. Salama and Hamza (2019) also found improved ovulation and pregnancy rates by using 42 days pretreatment with CPA/EE before ovulation induction with letrozole in CC resistant patient.
Methods:
This prospective randomized controlled trial study was conducted at Bangabandhu Sheikh Mujib Medical University (BSMMU), Dhaka during the period of January 2019 to June 2019. Already diagnosed patients of polycystic ovary syndrome with subfertility with high serum AMH level more than 5 ng/ml attending in the OPD of Reproductive Endocrinology and Infertility department at BSMMU are the study population. Total 60 subjects fulfilling rotardum criteria and treated with 3 cycle CC with 100 mg/day for 5 days from day 2 of menstrual cycle but fail to ovulate were recruited for the current study. Those subjects with tubal, and male factors excluded from the study. Total 60 PCOS with high serum AMH level patients were included in this study with maintaining inclusion criteria, and among them 6 patients were excluded from the study due to (tubal factors, and male factor). Were considered exclusion criteria for all subjects: an age less than 18 or greater than 35 years; a body mass index (BMI, kg/m2) less than 18 or greater than 25; major medical disorders and/or current or previous use of hormonal and/or metabolic drugs; tubal or male factor infertility or sub-fertility investigated with hysterosalpingography and standard semen analysis. During the study complete history, physical examination, basal level of FSH, LH and testosterone hormones were registered and metabolic assessment included OGTT and fasting insulin level were done, and to identify the changes in clinical, transvaginal ultrasonography and biochemical profile of study population. Clinical evaluation consisted of gynecological examination. Biochemical assessment consisted of complete hormonal, including evaluation of serum follicle stimulating hormone (FSH), luteinizing hormone (LH), thyroid-stimulating hormone (TSH), prolactin (PRL), and metabolic evaluation, including evaluation of fasting glucose and insulin levels. Baseline clinical evaluation consisted of gynecological examination, biochemical parameters and sonographic parameters were recorded. Blood sample were collected in the early morning during follicular phase (Day 2-3 of spontaneous cycle or progesterin induced withdrawal bleeding in case of amenorrhea) for the measurement of serum follicle stimulating hormone (FSH), luteinizing hormone (LH), total testosterone (T), will be measured by using Chemiluminescent Immunoassay. Serum AMH be measured by Enzyme Linked Immunosorbant Assay (ELISA). On the day 2-5 of a menstrual cycle, a baseline transvaginal ultrasound was performed with the urinary bladder empty using transvaginal probe. The ovarian volume and antral follicle count (AFC) were assessed. Side by side absence of any ovarian cyst will be ensured before starting the treatment. The volume of each ovary were calculated by measuring the ovarian diameters (D) in three perpendicular directions. Here, the formula of ellipsoid: D1 x D2 x D3 x 0.5236 will be applied. For AFC, small follicles with a diameter between 2 and 9 mm will be calculated following the recommendations. After enrollment of the patients, randomization were carried out by blocked randomization method to divide them equally into two groups- Group A number of patients 30 (treated with 2 mg cyproterone acetate (CPA) and 35 mcg of ethinylestradiol (EE) and Group B number of patients 30 treated with Metformin. Group A patients will receive treatment with 2 mg cyproterone acetate (CPA) and 35 mcg of ethinylestradiol (EE) for 6 months and Group B patients were receive treatment with Metformin for 6 months. Changes of the above mentioned parameters were recorded of patients after 6 month of treatment. Cyproterone acetate and metformin. The information were collected and recorded in the present questionnaire.
To assess changes of base line parameter and parameter after 6 month of treatment respectively away 2 group. (Ovarian changes was assessed by transvaginal monitoring antral follicular count and size of follicles Hormonal analysis.)

The study was prospective and randomised. Randomisation was non-blind and was based on patients’ chronological presence at the outpatient endocrine infirmary, namely the first one in Group A the second in Group B etc. sixty (60) women with PCOS were randomly divided into two groups: Group A (n=30, age 18-35 years, BMI 19-25 kg/m2), which comprised women were treated for 6 months with an OC containing 35 mcg ethinylestradiol plus 2 mg cyproterone acetate. Group B (n=30, age, BMI same range), which comprised women who were treated for 6 months with metformin 500 mg 8 hourly.
After treatment clinical evaluation, blood sampling, transvaginal ultrasonography, were done. Blood samples were collected during on the 2-3 day of cycle (reflecting 3 and 6 months of treatment), while patients were on OCP or metformin. Biochemical assessment consisted of complete hormonal, including evaluation of serum follicle stimulating hormone (FSH), luteinizing hormone (LH), thyroid-stimulating hormone (TSH), prolactin (PRL), total testosterone (T), and metabolic evaluation, including evaluation of fasting glucose and insulin levels. Serum AMH levels were assessed.

Finally, the ovarian dimensions, volume and morphology and the number of antral follicles (follicular diameter ranged from 2 to 9 mm) were evaluated bilaterally by
transvaginal ultrasonography. The antral follicle number per ovary, defined as the average for the total number of antral follicles counted from both ovaries, was also calculated monthly for 6 month.

Changes in clinical parameters (Acne, BMI, Menstrual cycle) was recorded after 6 month. Changes of biochemical parameters (serum LH (D2), serum FSH (D2), serum testosterone, OGTT and fasting insulin), serum AMH were recorded every three monthly for 6 month. Changes of sonographic parameters (ovarian volume, Antral follicular count development, was done after 6 month.

**Results:**

This prospective study was carried out in the Department of Reproductive endocrinology and infertility Department of Bangabandhu Sheikh Mujib Medical University, Dhaka, between January 2019 to June 2019. Total 60 PCOS with high serum AMH level patients were included in this study with maintaining inclusion and exclusion criteria, due to (tubal factors, and male factor).

During the study complete history, physical examination, basal level of FSH, LH and testosterone hormones were registered and metabolic assessment included OGTT and fasting insulin level were done, and to identify the changes in clinical and biochemical profile and TVS findings following ciproterone acetate in PCOS patients with high AMH level. Statistical analyses were carried out by using the Statistical Package for Social Sciences version 23.0 for Windows (SPSS Inc, Chicago, Illinois, USA) in personal computer. The p value <0.05 were considered as statistically significant

Table I shows characteristics of the study population. It was observed that before treatment mean age of study population was found 26.8±3.6 years in Group-A and 25.5±2.8 in Group-B. Mean BMI was found 26.3±4.9 kg/m2 in Group-A and 25.69 ± 2.34 in Group-B. Hirsutism 10.23 ± 3.56 in Group-A and 9.59 ± 1.47 in Group-B. Ovarian volume 12.34 ± 1.67 in Group-A and 11.89 ± 2.46 in Group-B.

Number of follicle 14.27 ± 2.13 in Group-A, 13.45 ± 1.78 in Group-B.

### Table I

**Clinical presentations of the study participants.**

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Group-A</th>
<th>Group-B</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (in years)</td>
<td>26.8 ± 3.6</td>
<td>25.5 ± 2.8</td>
</tr>
<tr>
<td>BMI (Kg/m2)</td>
<td>26.3 ± 4.9</td>
<td>25.69 ± 2.34</td>
</tr>
<tr>
<td>Waist circumference (cm)</td>
<td>88.34 ± 2.15</td>
<td>87.40 ± 3.68</td>
</tr>
<tr>
<td>Hirsutism</td>
<td>10.23 ± 3.56</td>
<td>9.59 ± 1.47</td>
</tr>
<tr>
<td>Ovarian volume</td>
<td>12.34 ± 1.67</td>
<td>11.89 ± 2.46</td>
</tr>
<tr>
<td>Number of follicle</td>
<td>14.27 ± 2.13</td>
<td>13.45 ± 1.78</td>
</tr>
</tbody>
</table>

### Table II

**Baseline the hormonal parameter of both group (before treatment).**

<table>
<thead>
<tr>
<th>Variables</th>
<th>Group-A</th>
<th>Group-B</th>
</tr>
</thead>
<tbody>
<tr>
<td>OGTT mmol/L</td>
<td>8.7 ± 1.0</td>
<td>7.13 ± 2.4</td>
</tr>
<tr>
<td>FSH (mIU/ml)</td>
<td>5.40 ± 1.56</td>
<td>6.04 ± 1.75</td>
</tr>
<tr>
<td>LH (mIU/ml)</td>
<td>14.8 ± 6.64</td>
<td>13.76 ± 6.86</td>
</tr>
<tr>
<td>Serum Testosterone (ng/ml)</td>
<td>81.79 ± 33.22</td>
<td>83.14 ± 22.14</td>
</tr>
<tr>
<td>AMH (ng/dl)</td>
<td>9.10 ± 2.95</td>
<td>8.90 ± 3.49</td>
</tr>
<tr>
<td>HOMA IR</td>
<td>1.95 ± 1.4</td>
<td>2.11 ± 1.18</td>
</tr>
</tbody>
</table>
Mean base line OGTT mmol/L was found in 8.7 ± 1.0 Group-A, 7.13 ± 2.4 Group-B. Mean FSH (mIU/ml) Group-A 5.40 ± 1.56 and Group-B 6.04 ± 1.75mIU/ml. Mean LHmIU/ml level was found in 14.8 ± 6.64 Group-A, and 13.76 ± 6.86 Group-B. Mean testosterone(ng/ml) was found in 81.79 ± 33.22 Group-A, and 83.14 ± 22.14 Group-B. Mean AMH (ng/dl) 9.10 ± 2.95 Group-A, and 8.90 ± 3.49 Group-B. HOMA  IR  3.95 ± 1.4 Group-A and 3.11 ± 1.18 Group-B.

**Table III**

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group -A</th>
<th>P-Value</th>
<th>Group -B</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum LH</td>
<td>4.02 ± 1.05</td>
<td>&lt; 0.001</td>
<td>7.47 ± 3.06</td>
<td>= 0.001</td>
</tr>
<tr>
<td>Serum FSH</td>
<td>3.88 ± 2.35</td>
<td>=0.002</td>
<td>4.14 ± 6.32</td>
<td>=0.012</td>
</tr>
<tr>
<td>Serum Testosterone</td>
<td>50.78 ± 14.86</td>
<td>=0.003</td>
<td>55.59 ± 27.23</td>
<td>=0.004</td>
</tr>
<tr>
<td>OGTT</td>
<td>5.07 ± 2.37</td>
<td>&lt;0.001</td>
<td>6.14 ± 4.25</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>HOMA IR</td>
<td>2.90 ± 1.76</td>
<td>=0.152</td>
<td>2.60 ± 1.53</td>
<td>= 0.342</td>
</tr>
<tr>
<td>Serum AMH</td>
<td>3.88 ± 0.84</td>
<td>&lt;0.001</td>
<td>6.42 ± 3.57</td>
<td>&lt;0.005</td>
</tr>
<tr>
<td>BMI</td>
<td>21.05 ± 1.99</td>
<td>0.967</td>
<td>21.67 ± 2.30</td>
<td>0.990</td>
</tr>
</tbody>
</table>

Table IV

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Group -A Mean - SD</th>
<th>P-Value</th>
<th>Group -A Mean - SD</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total number of follicles</td>
<td>8.24 ± 1.46</td>
<td>&lt;.001</td>
<td>10.88 ± 1.06</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>Ovarian Volume</td>
<td>6.12 ± 1.89</td>
<td>&lt;.001</td>
<td>9.87 ± 2.02</td>
<td>&lt; 0.453</td>
</tr>
</tbody>
</table>

Total number of follicle and Ovarian Volume was significantly decreased under CA/EE treatment compared to metformin (p< 0.001).

Mean base line OGTT mmol/L was found in 8.7 ± 1.0 Group-A , 7.13 ± 2.4 Group-B. Mean FSH (mIU/ml) Group-A 5.40 ± 1.56 and Group-B 6.04 ± 1.75mIU/ml. Mean LHmIU/ml level was found in 14.8 ± 6.64 Group-A, and 13.76 ± 6.86 Group-B. Mean testosterone(ng/ml) was found in 81.79 ± 33.22 Group-A, and 83.14 ± 22.14 Group-B. Mean AMH (ng/dl) 9.10 ± 2.95 Group-A, and 8.90 ± 3.49 Group-B. HOMA  IR  3.95 ± 1.4 Group-A and 3.11 ± 1.18 Group-B.

**Results:**

AMH was significantly decreased after treatment with 35 µg ethinylestradiol plus 2 mg cyproterone acetate (p <0.001 at 6 months), and treatment with metformin did not significantly affect serum AMH levels. AMH was significantly decreased under CA/EE treatment compared to metformin (p = 0.005). Serum LH and OGTT was significantly decreased with both group of treatment p< 0.001.

**Discussion:**

AMH is produced by the granulosa cells of early developing follicles and reflects the continuous, non-cycling growth of small follicles in the ovary. AMH has been found to be increased in the serum of women with PCOS. AMH levels are not influenced by hormone al fluctuations and remain constant throughout the menstrual cycle, making it a promising diagnostic marker for patients with PCOS [19]. The aim of the present study was to assess the impact of CEA/EE on serum AMH levels in women with PCOS. The data of the present study clearly demonstrate a significant decrease of serum AMH levels under treatment with an OC containing 35 mg ethinylestradiol plus EE while no statistically significant change was noted under treatment with metformin 500 mg hourly. Our data are clearly contra-dictory to the data previously presented by Somunkiran et al. who reported no change in serum AMH level after 6 months of treatment with 35mg ethinylestradiol plus EE. This discrepancy might be attributed to the different selection of patients with PCOS between the two studies. Somunkiran et al. recruited their PCOS patients according to the criteria of the Rotterdam PCOS workshop group, namely the presence of two of the following three criteria: oligomenorrhea or amenorrhea, clinical or biochemical signs of hyperandrogenism, and ultrasonographic polycystic ovarian morphology, while patients with PCOS in our study were recruited to fulfill (and) the stricter criteria proposed in 1990 by the National Institute of Child Health and Human Development Conference on PCOS. Never theless, in the present study, serum AMH levels after 6 months of treatment with 35mg ethinylestradiol
plus 2 mg cyproterone acetate were significantly related to serum testosterone. In a previous study, we have shown that AMH levels were higher in anovulatory and hyperandrogenemic women with NIH-defined ‘classical’ PCOS, compared to anovulatory women with PCOS morphology on ultrasound but normal androgen. Serum AMH levels at baseline 6 months treatment with 35mg ethinylestradiol and 2 mg cyproterone acetate, with metformin 500 mg 8 hourly. Mean base line AMH (ng/dl) 9.10 ±2.95 Group-A, and 8.90 ± 3.49 Group-B . Serum AMH group A 3.88 ± 0.84 p value < 0.001 after 6 month. Serum AMH Group-B 6.42 ± 3.57and p value <0.005 after 6 month. AMH was significantly decreased after treatment with 35 µg ethinylestradiol plus 2 mg cyproterone acetate (p <0.001 at 6 months), and treatment with metformin did not significantly affect serum AMH levels p value < 0.005 after 6 month. Levels. In this study, the strong positive association between LH and AMH levels, the significantly higher LH concentrations in women with ‘severe’ PCOS along with the highest levels of serum AMH, Therefore, the findings of the present study add additional evidence that AMH levels reflect the severity of PCOS, traditionally defined by its two cardinal elements, i.e. oligo-anovulation and hyperandrogenemia.16 Somunkiran et al. [14 Ovulatory disorders in women with PCOS are caused by an increased early follicular growth, resulting in a large reserve of follicles and/or a defective follicular selection, leading to follicular arrest. Since intra-ovarian androgens are also responsible for defective follicular selection and follicular arrest, it has been proposed that the intra-ovarian hyperandrogenism by increasing the AMH intra-ovarian level could exert an inhibiting effect on the selection process [8]. The excess in AMH production by poly cystic ovaries might be the result of the increased number of follicles ≤ 9 mm in diameter caused by the intra-ovarian excess of androgens4. We have previously shown that increased serum LH levels was the most significant independent link between PCOS-associated disorders of ovulation and the observed increase in serum AMH as it contributed as much as 18% in the variance of circulating AMH.7 It is postulated that premature LH action on the granulosa cells of anovulatory women with PCOS con-trIBUTE to the follicular arrest, being the link between PCOS-associated disorders of ovulation and the observed increase in ovarian production of AMH.11 In the present study BMI was not modified in a both groups after 6 months of treatment, therefore the major effect of metformin was not achieved and thus additional time should be given in or derto observe the net effect of metformin administration on body weight reduction, IR amelioration and eventually its effect on serum AMH levels. Therefore, the findings of this study suggest that although obesity and, to a lesser extent IR, play a role in modulating serum AMH levels, the decrease in. Finally, the results of the present study should be reinterpreted with caution taking into account the limitations of this study. A systematic review Badawy et al. 42 and Elmashad43 concluded that ovarian volume has little clinical application in prediction of poor pregnancy response. However, another recent review commented on the value of ovarian volume with regard to its easy execution, and therefore could be included in preparatory protocols providing data for continuity of research.

**Conclusion:**

AMH serum levels reflect the severity of PCOS and are significantly increased in its ‘classical’ phenotypic forms, based on consensus 2015. Moreover, AMH levels are significantly decreased under treatment with OCS containing 35 ethinylestradiol plus 2 mg Cyproteroneacetate, through decrease in serum androgens and the suppression of pituitary gonadotropins (LH) there by release of negative feedback effect of LH and androgen on hypothalamus is released. So compensatory release of FSH, resulting correction of anovulation which is assessed by folliculometry, So further research is needed to evaluate the response of ovulation induction pretreatment serum AMH level with Ciproterone Acetate and estradiol with Patient PCOS having high some AMH levels who one clouephane resident.

**References**


26. Carlsen SM, Vanky E, Fleming R. Anti-Mu llerian hormoneconcentrations in androgen-suppressed women with poly cystic ovary syndrome. Hum Reprod 2009;24:1732–1738.92D. Panidis et al. Gynecol Endocrinol Downloaded from informahealthcare.com by Hacettepe Univ. on 01/27/12 For personal use only. Endocrin Downloaded from informahealthcare.com by Hacettepe Univ. on 01/27/12 For personal use only.