

Testosterone To Estradiol (T/E₂) Ratio, As A Laboratory Predictor In Infertile Women With Polycystic Ovarian Syndrome

Abdelgadir Elmugadam^{*1}, Mohammed M Ahmed², Saber Nugta³, Ahmed Abdalgader², Ghada Elfadil¹

¹Sudan University of Science & Technology,

²Omdurman Islamic University, ³Kordofan University- Khartoum -Sudan

Corresponding author: Abdelgadir Elmugadam. mugadam01@gmail.com

Abstract: Polycystic ovary syndrome is a common endocrine disease in women, characterized by heterogeneous presentation of hyperandrogenism, ovulatory dysfunction, and polycystic ovarian morphology. This study was carried out to assess plasma levels of testosterone, estradiol calcium and cholesterol levels among females with polycystic ovarian syndrome (PCOs).

Forty females aged 18-35 years diagnosed by ultrasonography and Anti-Müllerian hormone (AMH) test in Alsir Abu-Hassan center for fertility in Khartoum State, and forty apparently healthy individuals as control group participated in this study.

Plasma testosterone and estradiol measured by using ELISA, cholesterol measured by BS-380 chemistry analyzer, and calcium by colorimetric method. Results were analyzed using statistical package for social science (SPSS), computer programmed version 21.

The study showed that, the mean plasma levels of testosterone and T/E₂ ratio were significantly increased in PCOs female patients. For testosterone (1.4±1.1 versus 0.87±0.65ng/ml, p-Value =0.011). For T/E₂ ratio (0.019±0.015 versus 0.01±0.008 ng/pg, p-Value =0.022).

Also the finding of this study showed insignificant difference in estradiol levels between PCOs compared with control group. (Mean±SD: 86±39 versus 101±38.6 pg/ml) respectively with P.Value 0.086. The mean plasma levels of cholesterol was significantly increased in PCOs female patients. (Mean±SD: 207.6±33.4 versus 112±33.4 mmol/L) respectively with P.vale 0.000.

The mean level of plasma calcium was significantly decreased (p.value 0.000). It is concluded that: the plasma levels of testosterone, T/E₂ ratio and cholesterol are higher in PCOs female patients, while calcium level decreased.

Keywords: PCOS, testosterone, estradiol, ratio, cholesterol, calcium.

1. INTRODUCTION

Polycystic ovary syndrome (PCOS) is characterized by chronic anovulation and hyperandrogenism which can be present in a different degree of severity. The main physiopathological basis of this syndrome includes Insulin-resistance and hyperinsulinemia⁽¹⁾. Many females with PCOS have an increased risk of insulin resistance which, with the prevalence of obesity, is a powerful risk factor for progression to type 2 diabetes. They also have an increased long-term risk of endometrial hyperplasia /cancer⁽²⁾. PCOS is a heterogeneous disorder (that is, capable of having somewhat different manifestations in different people) and the exact cause not known until now, and it is a very common problem among patients attending infertility clinics. The diagnosis depend on above criteria and rule out other causes of hyperandrogenism. study conducted to assess serum Anti-Müllerian hormone as laboratory predictor in infertile women with PCOS⁽³⁾. 17β-oestradiol is the principle hormone produce by the ovaries also synthesized in the placenta from androgens secreted by the fetal adrenal glands⁽⁴⁾. Dyslipidemia is the most common abnormality in PCOS⁽⁵⁾.

Materials and Methods: This cross sectional study was conducted in Khartoum state, the capital of Sudan, eighty volunteer included to participate in this study, 40 with polycystic ovarian syndrome, 40 healthy subjects without any diseases control group. All subjects were diagnosed with

PCOS, defined as oligomenorrhea (history of no more than eight spontaneous menses per year) and hyperandrogenemia (elevated testosterone level documented within the previous year in an outpatient setting on the basis of local laboratory results. Any menopausal and women receiving contraceptives were excluded. Consent was taken regarding acceptance to participate in the study and re-assurance of confidentiality. Before the specimen was collected, the donors knew that this specimen was collected for research purpose. About 2.5 ml of venous blood were collected from each participant (both case and control). The samples were collected under aseptic conditions and placed in sterile lithium heparin containers and centrifuged for 5 minutes at 3000 RPM to obtain plasma then they obtained sample were kept in plain containers at 2-8°C until the time of analysis. Data was analyzed to obtain means standard deviation and correlation of the sampling using statistical package for social science (SPSS) computer Programmed version 21, t test and person correlation were used for comparison and correlation

Results:

Table (1): Illustrate the mean concentration of testosterone and estradiol and ratio between them serum calcium, albumin in polycystic ovarian syndrome patients and control group.

Variable	PCOS N=40 Mean±SD	Control N=40 Mean±SD	P.Value
Estradiol pg/mL	86.0±39.0	101±38.6	0.086
Testosterone ng/ml	1.40±1.10	0.87±0.65	0.011*
T/E ₂ ratio	0.019±0.015	0.01±0.008	0.022*
AMH	10.6±5.92	1.40±0.22	0.000*
Cholesterol mmol/L	207.6±33.4	112±33.4	0.000*
Calcium mg/dL	7.0±0.69	9.2±0.55	0.000*
Albumin g/dL	4.1±0.56	4.5±0.46	0.124

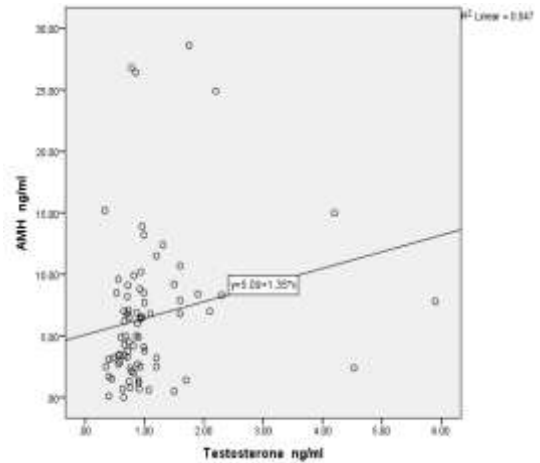


Figure (B)

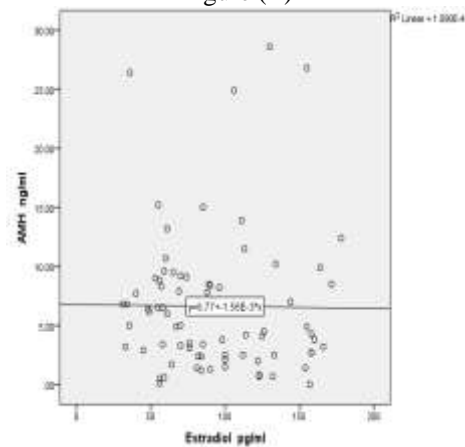


Figure (C)

*Result given in mean ± SD, P-Value ≤ 0.05 Consider significant.

* Independent sample T test was used for comparison

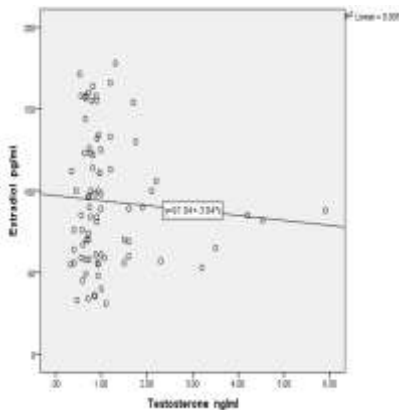


Figure (A)

Figure: (1): (A) Show correlation between testosterone concentration and estradiol ($r = -0.073$, P-Value = 0.522). (B) Show correlation between AMH and Testosterone ($r = 0.216$, P-Value = 0.055). (C) Show correlation between AMH and estradiol ($r = -0.010$, P-Value = 0.927)

Discussion:

Polycystic ovary syndrome (PCOS) disrupt the fertility hormones in females by increasing or decreasing them, this study conducted to get the disrupt of testosterone / estradiol ratio among PCOS female patients. In research, the ages ranged from 17 to 29 years. In the past they thought PCOs presents during the reproductive years only, but now can be diagnosed from fetal life. In this study, the average age of woman was from (17 to 30 years). This finding is consistent with study by ⁽⁶⁾, who had reported that the proportion of women with PCO decreased with age. This can be caused by

a decrease in the number of antral follicles throughout the reproductive years that occurs in normal women, a phenomenon that also applies to patients with PCOS ⁽⁷⁾.

Although PCOS classically presents during the reproductive years with menstrual irregularities, hyperandrogenism and metabolic complications, the origin of the disorder probably occurs very early starting from fetal life. In utero exposure to elevated testosterone levels coupled with gestational hyperglycemia may contribute to early differentiation of PCOS or may lead to amplification of the phenotype in genetically predisposed individuals. The spectrum of presentation of PCOS phenotype changes across the life span of a given individual. Improved understanding of the disease

spectrum has allowed us to identify endocrine and metabolic changes in the very young subject with high risk of developing PCOS⁽⁸⁾

The present study revealed significant increase in mean testosterone and T/E₂ ratio levels among case when compared to control group with p-value 0.011 and 0.022 respectively, while estradiol showed insignificant difference with p-value 0.086. This finding agreed with study done by (9) (Shang-Gwo et al., 2008), who reported that polycystic ovary syndrome (PCOS) had higher levels of T/E₂ ratio as compared to control group. Also similar to study done on by (10), who approved that (PCOS) had higher levels of testosterone as compared to control and estradiol not significant difference in PCOs compared to control. Because the PCOs female produce excess LH and AMH hormone which can inhibit the ovarian production of estradiol, also inhibit the aromatase enzyme that can lead to prevent conversion of testosterone to estradiol in peripheral tissues.

In the present study of PCOs, there are no correlations were found between plasma levels of testosterone and estradiol. This finding was agreed with studies done by (10).

The result showed that, there was no correlations were found between AMH and plasma levels of testosterone and estradiol. This finding was agreed with estradiol and disagreement with testosterone in studies done by (11). Also the result showed that, there was no correlations were found between age and plasma levels of testosterone and estradiol and T/E₂ ratio. In our study, we found that there was highly elevation in the concentration of AMH in patients samples when compared with control group and the difference was significant (P. value 0.000), also the highly reduction in concentration of serum calcium was found and the difference was significant also (P. value 0.000). Several studies showed that the level of AMH in plasma was significantly increased in patients with polycystic ovarian syndrome but there was significant decrease level of calcium among them. These findings were similar with that obtained by (12). The increased level of AMH was explained by the fact that AMH concentrations directly reflecting the increased number of early antral follicles. Moreover, the magnitude of AMH elevations in PCOS is associated with the extent of disease (13).

Dyslipidemia is one of the important risk factor associated with PCOS. Abnormal lipid metabolism is one of the main metabolic characteristics of PCOS patients. The result of this study show that PCOS patients had higher total cholesterol concentration when compared with control group which are similar to the results observed in PCOS patients in another study (14) also in agreement with some studies suggested that PCOS patients are hyperlipidemic with higher total cholesterol compare to control group (7,15).

The reduced level of calcium was explained by the fact that Females with polycystic ovarian syndrome have vitamin D

deficiency, 83% of all PCOS patients showed vitamin D deficiency while 35% were severely deficient (16). Another study showed the opposite observations in which they suggest that abnormalities in calcium homeostasis may be responsible, in part, for the arrested follicular development in women with PCOS and may contribute to the pathogenesis of PCOS (17).

Also it showed that there were no significantly difference between serum albumin and estradiol levels among female patients with polycystic ovarian syndrome compared to healthy individuals (P.value0.283) and (P.value0.086) respectively. The possible explanation of the decreased level of estradiol is the evidence that decreased aromatase activity may be a possible mechanism underlying the arrested follicular growth in PCOS. This was suggested by the study which showed that follicles in women with PCOS contain low levels of estradiol, aromatase mRNA and aromatase activity. PCOS follicular fluid contains one or more endogenous inhibitors of aromatase activity. 5 α -androstane-3, 17-dione, a 5 α - reduced androgen, is a competitive inhibitor of aromatase activity; it is markedly elevated in PCOS follicular fluid (18). In addition, 5 α -reductase activity is substantially higher in PCOS follicles than in control follicles, leading to increased production of 5 α - androstane-3, 17-dione in women with PCOS. Collectively, the decreased estradiol production and increased androgen production in PCOS may be a result of elevated 5 α -reductase activity and decreased aromatase activity (18).

Conclusion

from the results and finding of this study, it is concluded that the plasma levels of testosterone and T/E₂ ratio are higher in PCOs female patients. Decreased level of calcium and estradiol as well as albumin also decreased which may increase the risk of cardiovascular disease, osteoporosis and delay the course of treatment and recover of ovulatory problems. The plasma levels of total cholesterol is increase in PCOs female patient

Acknowledgement

The authors are grateful acknowledge the staff members of Al-sir Abu-elhasa Center for Fertility for their contribution in diagnosis and sample collection which were essential steps in completion of this study.

Conflicts of Interest:

The authors declare no conflicts of interest regarding the publication of this paper.

References:

1. Lizneva D., Suturina L., Walker W., Brakta S., Gavrilova-Jordan L., Azziz R. Criteria, prevalence, and phenotypes of polycystic ovary syndrome. *Fertilityand Sterility*. 2016;106(1):6–15

2. **Simon Walker**, Geoffrey Beckett, Pater Rae, Peter Ashby. (Ed.) (2013).WHITBY'S CLINICAL BIOCHEMISTRY Lecture Notes. 9Th editions. UK. John Wiley & Sons. 141-147.
3. **Tayrab, E.**, Ali ,M., Modawe, G.A., Naway, L(2014). Serum Anti-Müllerian hormone as laboratory predictor in infertile women with and without polycystic ovary syndrome. 2(3):61–6.
4. **William J Marshall**, Stephen K, Bangert Marta Lapsley. (2012). *Clinical chemistry*, seventh edition. China. Elsevier.258-272.
5. **Lergo ,R.**, Kunselman, A., Dunaif, A(2001).Prevalence and predictors of dyslipidaemia in women with polycystic ovary syndrome, *Am JMed.*, 111: 607-613.
6. **Johnstone ,E.** Rosen, M.Neril ,R.Trevithick, D.Sternfeld, B.Murphy, R(2010)The polycystic ovary post-rotterdam: a common, age-dependent finding in ovulatory women without metabolic significance. *endocrinol metab.*;95(11):4965–72.
7. **Madhu, L.**Vijay Bhaskar, S. Sharma, A. Sumapreethi.(2012). Marker of oxidative stress and serum lipid in PCOS ,journal of evaluation of medical and dental science .vol 1.P.769.
8. **Belinda George** and M Ganapathi Bantwal. (2016). Polycystic Ovary Syndrome (PCOS) - From in Utero to Menopause. *Diabetes & Obesity International Journal*. 1(2).107.
9. **Shang-Gwo Horng**, Tzu-Hao Wang, Hsin-Shih Wang. (2008).Estradiol-to-Testosterone Ratio Is Associated with Response to Metformin Treatment in Women with Clomiphene Citrate- Resistant Polycystic Ovary Syndrome (PCOS). *Chang Gung Med J* .31. 477-83.
10. **Elisabeth Lerchbaum**1,2*, Verena Schwetzl , Thomas Rabe2 , Albrecht Giuliani3 , Barbara ObermayerPietsch(2014). Hyperandrogenemia in Polycystic Ovary Syndrome: Exploration of the Role of Free Testosterone and Androstenedione in Metabolic Phenotype. *PLOS ONE*; 9 (10). 1 – 12.
11. **Adel F., Begawy.** , Akmal, N., El-Mazny. , Nermeen A., Abou-Salem. , Nagwa, E and. El-Taweel. (2010). Anti-Mu" llerian hormone in polycystic ovary syndrome and Norma-ovulatory women: Correlation with clinical, hormonal and Ultrasonographic parameters. *Middle East Fertility Society Journal*. (15).253-258.
12. **Al-Hakeem H K**, Kadhemi M A. (2009). Is lipid profile in women with polycystic ovary syndrome related to serum calcium or magnesium? *Journal of Karbala University*; 7(2):48-53
13. **Laven J S**, Mulders A G, Visser J A, Themmen A P et al (2004). Anti-Mullerian hormone serum concentrations in normoovulatory and an ovulatory women of reproductive age. *J Clin Endocrinol Metab*; 89:318–323.
14. **Valkenburg ,O.**, Steegers-Theunissen, R.P., Smedts,H.P., Dallinga-Thie ,G.M., Fauser ,B.C., Westervel,E.H (2008).A more atherogenic serum lipoprotein profile is present in women with polycystic ovary syndrome: A case-control study. *J clin endocrinol metab*;93:470-6.
15. **Cristian-Ioan ,I** , Nicolae, C, Dan. M(2012).lipid parameter in polycystic ovary syndrome .*Applied medical informatics* .vol.31 .P.27-32.
16. **Sirmans S M**, Pate K A (2013). Epidemiology, diagnosis, and management of polycystic ovary syndrome. *Clin Epidemiol*; 6: 1-13.
17. **Firouzabadi R d**, Aflatoonian A, Modarresi S et al (2012). Therapeutic effects of calcium & vitamin D supplementation in women with PCOS. *Complement Ther Clin Pract*; 18(2): 85-88.
18. **Horng S D**, Wang T H, Wang H S (2008). Estradiol-to-testosterone ratio is associated with response to metformin treatment in women with clomiphene citrate-resistant Polycystic Ovary Syndrome (PCOS) *Chang Gung Med J*; 31(5).24-28.