

## Thyroid Profile Status of Female Patients with Polycystic Ovary Syndrome (PCOS) in Western Rajasthan

Dhamija Shilpa<sup>1</sup>, Kumar Rohitash<sup>2\*</sup>, Mathur Ranjana<sup>3</sup>

<sup>1</sup>M.Sc (Medicine) Biochemistry,

Department of Biochemistry, Dr. S. N. Medical College, Jodhpur, Rajasthan, India.

<sup>2</sup>M.Sc (Medicine) Biochemistry, Senior Demonstrator,

Department of Biochemistry, R. N. T. Medical College, Udaipur, Rajasthan, India.

<sup>3</sup>MD Biochemistry, Senior Professor & Head,

Department of Biochemistry, Dr. S. N. Medical College, Jodhpur, Rajasthan, India.

### ABSTRACT

**Background:** Polycystic ovarian syndrome (PCOS) is a common endocrine disorder in women with 6-7% prevalence in the world. PCOS produces symptoms in approximately 5% to 10% of women of reproductive age (12–45 years old). The heterogeneous disorder is associated with menstrual irregularities, hyperandrogenemia, obesity, increased insulin levels and insulin resistance. PCOS together with thyroid disorder create independent risks of ovarian malfunction as well as pregnancy associated problem. An abnormal thyroid concentration could trigger alterations in ovulation and then menstruation.

**Objective:** Assessment of serum Thyroid profile Levels in Female patients with Polycystic ovarian syndrome (PCOS) trying to find a correlation between Polycystic ovarian syndrome (PCOS) and the metabolic functions of Thyroid gland, as well as pregnancy associated problem.

**Patients and Methods:** 50 females were included in this study, they were divided into 2 groups, the first group included 25 PCOS female patients of age group (18-30yr) attending the Out Patient Department, Department of Obstetrics and Gynecology, Dr. S.N. Medical College and its associated group of hospitals, Jodhpur. The result will be compared with the second group in which age matched 25 healthy- control female subjects were included. Patients and controls were subjected to fill history taking, duration of the disease, the clinical course and any complications in relation to the disease were noted, measurement of body mass index and assessment of Serum thyroid hormones and Serum TSH level which was measured by using Enzyme Linked Fluorescent Assay Method.

**Results:** PCOS female patients were found with elevated Serum TSH level and Mean serum TSH values were highly significant as compared to healthy control subjects. While Serum T<sub>3</sub> and Serum T<sub>4</sub> levels were almost similar in both groups and there was a non-significant relationship.

**Conclusion:** The subclinical higher level of thyroid stimulating hormone in PCOS female patients may be associated with the development of the disease and may also implicated with other symptoms of the disease. As there is high prevalence of thyroid disorders in PCOS patients, thus all women in their reproductive age with and without PCOS, should have their thyroid function tests evaluated.

**Keywords:** PCOS, Ovarian Malfunction, Obesity, Subclinical Hypothyroidism.

### \*Correspondence to:

**Mr. Rohitash Kumar,**

Senior Demonstrator,

Department of Biochemistry,

R.N.T. Medical College, Udaipur, Rajasthan, India.

### Article History:

**Received:** 29-12-2016, **Revised:** 26-01-2017, **Accepted:** 11-02-2017

### Access this article online

Website: www.ijmrp.com	Quick Response code 
DOI: 10.21276/ijmrp.2017.3.2.025	

### INTRODUCTION

Polycystic Ovarian Syndrome (PCOS) is one of the most common female endocrine disorder.<sup>1</sup> It is a complex, heterogeneous disorder of multifactorial etiology and there is strong evidence that it can be classified as a genetic disease.<sup>2,3</sup> PCOS is thought to be one of the leading causes of female subfertility and the most frequent endocrine problem in women of reproductive age.<sup>4</sup>

This endocrine disorder was first identified in 1935 by Stein and

Leventhal who noticed a condition in women characterized by irregular menstruation, obesity, and hirsutism, in addition to cysts on the women's ovaries and it is also known as Stein – Leventhal syndrome.<sup>5</sup>

Although PCOS is an endocrine disease, it affects many systems of the body resulting in reproductive, metabolic, and psychological consequences.<sup>6</sup>

Women with PCOS are at risk for weight gain.<sup>7-10</sup> Obesity is a common finding of women with PCOS, but it is not part of the diagnostic criteria. Overweight or obesity affects approximately 60-80% of PCOS patients.<sup>11</sup>

PCOS together with thyroid disorder create independent risks of ovarian malfunction as well as pregnancy associated problem. An abnormal thyroid concentration could trigger alterations in ovulation and then menstruation. Initial stages of thyroid malfunction can result in delicate modification in ovulation as well as endometrial receptivity, which consequently may have drastic influence on fertility.<sup>12,13</sup>

Evaluation of thyroid function must be undertaken in any woman presenting with menstrual irregularities.<sup>14</sup>

The most obvious connection between these two disorders (Thyroid and PCOS) is the increased BMI and insulin resistance. Increase in BMI is an integral part of PCOS and is seen in large majority (54-68%) of these cases.<sup>15</sup>

PCOS is believed to be a heterogeneous dysfunction of multifactorial etiology. It can also be linked to primary hypothyroidism in 6.3% of PCOS patients.<sup>16,17</sup>

Thyroid gland dysfunction manifested as hypothyroidism is a common disorder has been reported in PCOS patients. The clinical features of hypothyroidism also include weight gain, menstrual irregularities and infertility.<sup>11</sup> Most of the times hypothyroidism is subclinical and diagnosed first time during evaluation of PCOS.<sup>7,18</sup>

The main pathophysiological components of PCOS are gonadotropic dysfunction and insulin resistance. It has been found that both of these components are related to BMI.<sup>19-21</sup>

## MATERIALS and METHODS

### Study Design and Population

The present study represents a case-control study. The study was carried out in 25 cases, between the age group (18-30 year) and 25 age and gender matched controls who attended the Out

Patient Department, Department of Obstetrics and Gynaecology, Dr. S.N. Medical College and its associated group of hospitals, Jodhpur, Rajasthan, India.

The institutional ethical committee approved the study protocol. Informed consent was obtained from all the participants. A thorough clinical and symptomatic examination of all the patients was done with the help of treating clinicians and was recorded to rule out and confirm the presence of disease by evidence of symptoms. The detailed information regarding family background, personal history, dietary habits and physical activity of each subject was recorded. Family history for obesity and hypothyroidism was taken.

### Methods

Informed consent was taken from patients and controls. A pre-structured Performa was used to collect the data. Baseline data including age, gender, BMI (kg/m<sup>2</sup>), detailed medical history, clinical examinations. Sample was collected from women patients between the age group 18 – 30years, during 2<sup>nd</sup> or 3<sup>rd</sup> day of menstrual cycle, venous blood sample was drawn from antecubital vein of each subject by using aseptic techniques. Blood samples were collected in vial was allowed to clot. The serum was separated from the clotted specimen by centrifugation at 3000 rpm for 10 minutes and serum was estimated for T<sub>3</sub>, T<sub>4</sub> and TSH by using Enzyme Linked Fluorescent Assay Method.

### Statistical analysis

The data assembled for different anthropometric and biochemical parameters were subjected to suitable statistical analysis to compute central tendencies (mean) and accompanying measures of variability statistics (standard deviation) etc for all the groups. The magnitude of inter group differences for each of the parameters was quantified by using students't' test values (Student's 't' test). On the basis of t-values, 'p' values (probability) were determined to make out significance of variance between the mean values of individual parameters among the groups of subjects studied.

Table 1: Mean BMI (kg/m<sup>2</sup>) of the subjects studied

S. No.	Group studied	BMI (Mean ± S.D.) [Range]
1	Healthy Control (25)	22.18 ± 1.97 [20.06-26.84]
2	PCOS patients (25)	25.19 ± 1.14 [22.72-27.11]

Table 2: Statistical analysis of BMI among the groups studied

S. No.	Group Compared	Healthy Control v/s PCOS patients
1	t – value	6.61
2	p – value	<0.0001 (HS)

HS = Highly Significant

Table 3: Mean Serum Thyroid Profile of the subjects studied

S.No.	Thyroid profile Tests	Healthy Control (25)	PCOS patients (25)	
1.	Serum T <sub>3</sub>	Mean ± S.D.	1.53 ± 0.29 nmol/L	1.64 ± 0.33 nmol/L
		Range	1.04-2.02	1.10-2.26
2.	Serum T <sub>4</sub>	Mean ± S.D.	93.05 ± 9.75 nmol/L	97.82 ± 17.67 nmol/L
		Range	78.59-113.87	60.98-123.12
3.	Serum TSH	Mean ± S.D.	2.15 ± 0.84 µIU/mL	5.06 ± 1.21 µIU/mL
		Range	1.19-3.78	3.51-7.87

**Table 4: Statistical analysis of Thyroid Profile Tests among the groups studied**

S. No.	Thyroid profile Tests	Group Compared	Healthy Controls v/s PCOS patients
1.	Serum T <sub>3</sub>	t – value	1.25
		p – value	0.2167 (NS)
2.	Serum T <sub>4</sub>	t – value	0.24
		p – value	1.18 (NS)
3.	Serum TSH	t – value	9.878
		p – value	<0.0001 (HS)

NS = Non Significant; HS = Highly Significant

## RESULTS

The mean Body Mass Index (BMI) of the healthy control subjects was  $22.18 \pm 1.97$ ; which varies from 20.06 to 26.84 kg/m<sup>2</sup>. It was  $25.19 \pm 1.14$  in PCOS patients which varies from 22.72 to 27.11 kg/m<sup>2</sup>. (Table 1)

A statistically high-significant difference ( $p < 0.0001$ ) was observed in BMI of PCOS patients ( $t = 6.61$ ) when results were compared with the healthy control subjects. (Table 2)

Serum triiodothyronine (T<sub>3</sub>) level varies from 1.04 to 2.02 nmol/L in healthy control subjects and 1.10 to 2.02 nmol/L in PCOS patients. Mean triiodothyronine was  $1.53 \pm 0.29$  in healthy controls and  $1.64 \pm 0.33$  in PCOS patients. (Table 3) Mean serum triiodothyronine (T<sub>3</sub>) values were non-significant ( $t=1.25$ ,  $p=0.2167$ ) in PCOS patients, as compared to healthy control subjects. (Table 4)

Serum tetraiodothyronine (T<sub>4</sub>) level varies from 78.59 to 113.87 nmol/L in healthy control subjects and 60.98 to 123.12 nmol/L in PCOS patients. Mean serum tetraiodothyronine (T<sub>4</sub>) level was  $93.05 \pm 9.75$  nmol/L in healthy controls and  $97.82 \pm 17.67$  in PCOS patients. (Table 3) Mean serum tetraiodothyronine (T<sub>4</sub>) were non-significant ( $t=0.24$ ,  $p=1.18$ ) in PCOS patients, as compared to healthy control subjects. (Table 4)

The serum TSH levels varied from 1.19-3.78  $\mu$ IU/mL in healthy control subjects and 1.30-7.87  $\mu$ IU/mL in PCOS patients. Mean serum TSH level was  $2.15 \pm 0.84$   $\mu$ IU/mL in healthy control subjects and  $4.00 \pm 1.86$  in PCOS patients. (Table 3) Mean serum TSH values were highly significant ( $t=4.53$ ,  $p<0.001$ ) in PCOS patients, as compared to healthy control subjects. (Table 4)

## DISCUSSION

Polycystic ovarian syndrome (PCOS) is an intense problem which manifests later as infertility, obesity, insulin resistance, dyslipidemia, endothelial dysfunction and overt diabetes mellitus. PCOS is often associated with abnormalities of other endocrine glands. Women with PCOS may be four times more likely to suffer from hypothyroidism.<sup>22</sup>

In our study, we found that the BMI values in the cases were significantly higher than in the control group. Thathapudi S et al (2014) observed a statistically high-significant difference ( $p<0.0001$ ) in BMI of PCOS patients as compared to healthy control subjects.<sup>23</sup> Ibraheem NJ et al (2015) also observed a statistically highly significant difference ( $p=0.02$ ) in BMI of PCOS patients ( $28.9 \pm 4.1$  kg/m<sup>2</sup>) as compared to healthy control subjects ( $26.0 \pm 2.5$  kg/m<sup>2</sup>).<sup>24</sup>

Hypothyroidism can aggravate PCOS symptoms. It can lead to low levels of sex hormone binding globulin (SHBG) which in turn can lead to higher concentrations of free testosterone throughout the body. High level of testosterone is one of the factors which

contribute to PCOS symptoms like infertility, polycystic ovaries, hirsutism, male pattern hair loss and acne. If hypothyroidism is diagnosed and treated early, some of PCOD symptoms may diminish.<sup>25</sup>

Our study showed that the mean TSH level were significantly higher in the cases than those in the control group which was consistent with Janssen OE *et al* (2004).<sup>26</sup> This study did not show a statistically significant difference between cases and controls regarding T<sub>3</sub> and T<sub>4</sub> levels which was consistent with study by Shirsath A et al (2015).<sup>27</sup>

Mean serum TSH values were highly significant ( $t=4.53$ ,  $p<0.001$ ) in PCOS patients in our study. Similar result was obtained by Janssen OE et al (2004)<sup>26</sup> in his study, which demonstrated a threefold higher prevalence of hypothyroidism in patients with PCOS.

In fewer other studies same results obtained, like Baul SN et al (2013) observed a statistically highly-significant relation ( $p<0.05$ ) in serum TSH levels between normal individuals ( $2.67 \pm 3.11$  mIU/ml) and PCOS patients ( $4.55 \pm 2.66$  mIU/ml).<sup>28</sup> Shirsath A et al (2015) also reported a highly significant relation ( $p<0.05$ ) in Thyroid stimulating hormone (TSH) levels ( $\mu$ IU/mL) between normal individuals ( $1.26 \pm 0.45$ ) and PCOS patients ( $7.13 \pm 1.60$ ).<sup>27</sup>

The study also evaluated that there is no relationship between thyroid hormones and PCOS disease which is similar with few old literatures. Shirsath A et al (2015) reported a non-significant relation ( $p>0.05$ ) in triiodothyronine (T<sub>3</sub>) levels (ng/ml) between normal individuals ( $1.26 \pm 0.21$ ) and PCOS patients ( $1.30 \pm 0.16$ ).<sup>27</sup> Abdeisalam A and Ibrahim W (2015) also observed a non-significant relation ( $p>0.05$ ) in triiodothyronine (T<sub>3</sub>) levels between normal healthy subjects and PCOS patients. Our findings of serum triiodothyronine (T<sub>3</sub>) levels were in accordance with the above studies.<sup>29</sup>

Shirsath A et al (2015) reported a non-significant relation ( $p>0.05$ ) in tetraiodothyronine (T<sub>4</sub>) levels ( $\mu$ g/dl) between normal individuals ( $7.52 \pm 1.82$ ) and PCOS patients ( $7.87 \pm 1.38$ ).<sup>27</sup>

## CONCLUSION AND RECOMMENDATIONS

The positive correlation between serum TSH levels and the presence of higher BMR in our study threw bright light that female patients with PCOS must be investigated for the presence of Subclinical hypothyroidism which is proved to be metabolic risk factors. As there is high prevalence of thyroid disorders in PCOS patients, all women in their reproductive age with and without PCOS, should have their thyroid function tests evaluated. Secondly correcting subclinical hypothyroidism will lead to improvement of overall hormonal and metabolic health of these females.

## REFERENCES

1. Fauser BCJM, Tarlatzis BC, Rebar RW, Legro RS, Balen AH, Lobo R et al (2012); Consensus on women's health aspects of polycystic ovary syndrome (PCOS): the Amsterdam 91 ESHRE/ASRM-Sponsored 3rd PCOS Consensus Workshop Group. *Fertility and Sterility*; 97:28-38.
2. Legro RS, Strauss JF (2002); Molecular progress in infertility: polycystic ovarysyndrome. *Fertil Steril*; 78(3): 569 – 576.
3. Diamanti-Kandarakis E et al (2006); Indices of Low-Grade Inflammation in Polycystic Ovary Syndrome. *Annals of the New York Academy of Sciences*; 1092(1):175-186.
4. Boomsma CM, Fauser BC, Macklon NS (2008); Pregnancy complications in women with polycystic ovary syndrome. *Semin Reprod Med*; 26:72-84
5. Stein IF, Leventhal ML (1935); Amenorrhea associated with bilateral polycystic ovaries. *Am J Obstet Gynecol*; 29:181-191.
6. Garad R, Teede HJ & Moran L (2011); an evidence-based guideline for Polycystic ovary syndrome. *Australian Nursing Journal*: 19(4),30.
7. Teede H, Deeks A & Moran L (2010); Polycystic ovary syndrome: a complex condition with psychological, reproductive and metabolic manifestations that impacts on health across the lifespan. *BMC Med*; 8,41.
8. March WA, Moore VM, Willson KJ, Phillips DI, Norman RJ, Davies MJ. (2010); The prevalence of polycystic ovary syndrome in a community sample assessed under contrasting diagnostic criteria. *Hum Reprod (Oxford, UK)*; 25(2):544–51.
9. Boyle JA, Cunningham J, O'Dea K, Dunbar T & Norman RJ (2012); Prevalence of polycystic ovary syndrome in a sample of Indigenous women in Darwin, Australia. *Med J Aust*; 196(1),62-16.
10. Legro RS, Arslanian SA, Ehrmann DA, Hoeger KM, Murad MH, Pasquali R et al (2013); Diagnosis and treatment of polycystic ovary syndrome: an endocrine society clinical practice guideline. *J Clin Endocrinol Metab*, 98(12), 4565-4592.
11. Azziz R et al (2004). The prevalence and features of the polycystic ovary syndrome in an unselected population. *The Journal of Clinical Endocrinology and Metabolism*; 89(6):2745-49.
12. Krassas GE (2000); Thyroid disease and female reproduction. *Fertility and sterility*; 74(6):1063-70.
13. Sinha U, Sinharay K, Saha S, Longkumer TA, Baul SN, Pal SK (2013); Thyroid disorders in polycystic ovarian syndrome subjects: A tertiary hospital based cross-sectional study from Eastern India. *Indian Journal of Endocrinology and Metabolism*; 17(2):304-9.
14. Gaitonde DY, Rowley KD, Sweeney LB (2012); Hypothyroidism: an update. *Am Fam Physician*; 86(3):244–51.
15. Lim SS, Davies MJ, Norman RJ, Moran LJ (2012); Overweight, obesity and central obesity in women with polycystic ovary syndrome: A systematic review and meta-analysis. *Hum Reprod Update*; 18:618-37.
16. Vilar L, Freitas MC, Naves LA, Casulari LA, Azevedo M, Montenegro R, Jr et al (2008); Diagnosis and management of hyperprolactinemia: results of a Brazilian multicenter study with 1234 patients. *Journal of endocrinological investigation*; 31(5):436-44.
17. McDermott MT (2009); In the clinic. Hypothyroidism. *Annals of internal medicine*; 151(11):l1c61.
18. Nelson VL, Legro RS, Strauss JF III, McAllister JM (1999); Augmented androgen production is stable steroidogenic phenotype of propagated theca cells from polycystic ovaries. *Mol Endocrinol Metab* ;86:5925-33.
19. Dale PO, Tanbo T, Vaaler S and Abyolm H (1992); Body weight, hyperinsulinemia and gonadotropin levels in the polycystic ovarian syndrome, evidence of two distinct population. *Fertil. Steril*; 3: 487–491.
20. Fulghesu AM, Cucinelli F, Pavone V, Murgia F, Guido M, Caruso A et al (1999). Changes in luteinizing hormone and insulin secretion in polycystic ovarian syndrome. *Hum Reprod*; 14 (3): 611-617.
21. Moran C, Garcia-Hernandez E, Barahona E, Gonzalez S & Bermudez JA (2003); Relationship between insulin resistance and gonadotropin dissociation in obese and nonobese women with polycystic ovary syndrome. *Fertility and Sterility*; 80(6): 1466-1472.
22. Bussen S, Steck T, Dietl J (2000); Increased prevalence of thyroid antibodies in euthyroid women with a history of recurrent in-vitro fertilization failure. *Hum Reprod*; 15(3): 545-8.
23. Thathapudi S et al (2014); Anthropometric and Biochemical Characteristics of Polycystic Ovarian Syndrome in South Indian Women Using AES-2006 Criteria. *Int J Endocrinol Metab*. 12(1): e12470
24. Ibraheem NJ (2015); Prevalence Elevated Day3 FSH/LH Ratio, Prolactin Variation and Central obesity with Menstrual Irregularities Among Primary Infertile Women in Babylon Province. *Journal of Babylon University/Pure and Applied Sciences*; (23):641.
25. Franks S, Gharani N, McCarthy M (2001); Candidate genes in polycystic ovarysyndrome. *Hum Reprod Update*; 7(4): 405-10.
26. Janssen OE, Mehlmauer N, Hahn S, Offner AH, Gartner R (2004); High prevalence of autoimmune thyroiditis in patients with polycystic ovary syndrome. *European journal of endocrinology*; 150(3):363-9.
27. Shirsath A, Aundhaka N, Kamble P (2015); Does the thyroid hormonal levels alter in polycystic ovarian disease? A comparative cross sectional study. *Indian Journal of Basic and Applied Medical Research*; 4: 265-271.
28. Baul NS et al (2013); Thyroid disorders in polycystic ovarian syndrome subjects: A tertiary hospital based cross-sectional study from Eastern India. *Indian Journal of Endocrinology and Metabolism*, 17;304-309.
29. Ibrahim W, abdelsalam KEA (2015); Levels of FSH LH, SHBG, Total Testosterone, and LH/FSH ratio in Sudanese patients with polycytic ovary syndrome in relation by body mass index. *International Journal of Current Research* 2015; 7(1):119-22.

**Source of Support:** Nil. **Conflict of Interest:** None Declared.

**Copyright:** © the author(s) and publisher. IJMPP is an official publication of Ibn Sina Academy of Medieval Medicine & Sciences, registered in 2001 under Indian Trusts Act, 1882.

This is an open access article distributed under the terms of the Creative Commons Attribution Non-commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

**Cite this article as:** Dhamija Shilpa, Kumar Rohitash, Mathur Ranjana. Thyroid Profile Status of Female Patients with Polycystic Ovary Syndrome (PCOS) in Western Rajasthan. *Int J Med Res Prof*. 2017; 3(2):122-25. DOI:10.21276/ijmrrp.2017.3.2.025