



## Characteristics of Body Composition in Different Phenotypes of South Indian Women with Polycystic Ovary Syndrome and Relation with Body Image and Psychological Profile

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### ABSTRACT

**Purpose of the study:** To study characteristics of body composition in different phenotypes of South Indian women with polycystic ovary syndrome and relation with body image and psychological profile

**Methods:** Non-comparative cross-sectional open label study to be carried out over the period of 12 months on PCOS patients attending gynecology outpatient department. Personal medical history to be obtained from every woman according to a customized prepared questionnaire. Presence of at least two criteria from clinical, hormonal, and abdominal USG category were considered diagnostic of PCOS. Hirsutism was scored according to modified Ferriman Gallawayscore. Recruited women have their body fat percentage calculated using Health Sense BF 414 Ultra Lite Body Fat Monitor. The bioelectrical impedance analysis (BIA) method was used to estimate body fat percentage.

**Results:** The most common menstrual abnormality identified in study was irregular menstrual cycles followed by a combination of amenorrhea and irregular cycles. The commonest phenotype was ovulatory dysfunction with polycystic morphology in 42.7% cases. The body dysmorphic disorder was the commonest psychological problem in about 25.6% closely followed by major depressive illness.

**Conclusion:** Our study suggested that psychological and neurological factors may play a role in the pathophysiology of PCOS. Therefore, psychological intervention should be an aspect of PCOS management.

**Key Words:** Phenotypes of PCOD, body image, psychological profile



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### INTRODUCTION

The polycystic ovary syndrome (PCOS) is an endocrine metabolic disorder affecting 7% to 10% of women during reproductive age [1]. It is a systemic disorder with reproductive, psychological, cosmetic, and oncologic consequences. It also has a metabolic component involving hyperglycemia and insulin resistance. This results in increased cardiovascular risk and type II diabetes mellitus [2]. Polycystic ovarian syndrome (PCOS) is a highly prevalent disorder [2, 3] affecting multiple aspects of a women's overall health, with long-term effects that transcend well beyond the reproductive age [4, 5]. The symptoms of PCOS include, somewhat variably, hyperandrogenism (HA), ovulatory dysfunction (OD), polycystic ovarian morphology (PCOM), gonadotropic abnormalities, and insulin resistance and compensatory hyperinsulinism. The disorder has high degree of heritability with complex genetic trait.

#### - Epidemiology/ severity of the problem

The syndrome is characterized by chronic anovulation and hyperandrogenism. It is manifested by hirsutism, cystic acne, hair loss, insulin resistance, and weight gain. It is also one of the primary causes of infertility all of which leads to decrease in health-related quality of life [6]. Studies showed PCOS exhibited high rates of psychopathology, with 52.7% of the sample suffering from a psychiatric condition [7].

Asian Indians have higher percentage body fat, abdominal adiposity at lower or similar BMI levels as compared to white Caucasians. In lower or similar BMI levels Asian Indians have higher percentage of abdominal adiposity as compared to white Caucasians. Studies indicate that the cut-off BMI corresponding to the cut-off of percentage body fat is lower for Asian Indians from various parts of India very broad spectrum of clinical manifestations and associated morbidities [8]

Over the last several decades, significant efforts have been made to classify PCOS; however, global consensus regarding a PCOS criterion remains controversial [9-12]. Currently proposed criteria are predominantly based on expert opinion [9-12]

The introduction of Rotterdam criteria led to a substantial increase in the number of patients diagnosed with PCOS, as well as broadened the heterogeneity of PCOS phenotypes as compared with the NIH definition [13]. The distribution and morbidity associated with specific PCOS phenotypes has been the object of extensive research, as reported by studies conducted in Europe [14-17], the Middle East [18], Asia [19], the Americas [20, 21]

As noted above, the presentation of PCOS can be subdivided into four phenotypes: phenotype A: androgen excess and ovulatory dysfunction, phenotype B: androgen excess and polycystic ovarian morphology; phenotype C: ovulatory dysfunction and polycystic ovarian morphology; and phenotype D: ovulatory dysfunction, androgen excess and polycystic ovarian morphology.

#### Distribution of PCOS Phenotypes

Understanding the distribution of PCOS phenotypes is essential in defining the epidemiology of PCOS in a population. Multiple studies from different regions around the world have reported the distribution of phenotypes in clinical cohorts of PCOS patients [22, 23, 24, 25 & 26]. Overall, published data indicate that more than half of PCOS patients identified within the clinical setting demonstrate phenotype A, whereas the other three phenotypes (i.e., B, C and D) have almost equal prevalence. The classic form of PCOS (i.e., phenotypes A and B) constitutes approximately two-thirds of the total of PCOS patients identified within the clinical setting [27]. Unfortunately, few data exist regarding the distribution of phenotypes in women with PCOS identified in medically unbiased (i.e., unselected) populations, which would more accurately reflect the distribution of phenotypes in PCOS in the “natural” state.

The array of symptoms such as obesity, acne, scalp hair thinning, menstrual irregularity, and sub fertility in PCOS contribute to psychological impairment [28]. Studies have evaluated the relation between PCOS and psychiatric disorders; however, most have evaluated psychiatric symptoms based on self-report measures [29]. There remains, therefore, an unclear relationship between PCOS and psychiatric disorders. Cross sectional epidemiological studies have reported that individuals with PCOS are more likely to have anxiety or depressive disorders when compared to those in the general population [30]. Two studies have shown depression, bipolar disorder, anxiety disorders, and binge eating disorder are more frequent among women with PCOS compared with control [31, 32, 33, 34, 35 & 36].

In light of the above mentioned, the purpose of the present study is to perform a study mood (bipolar disorder, dysthymia, or major depressive disorder), obsessive-compulsive spectrum disorders, trauma- and stressor-related disorders, anxiety disorders, psychotic disorders, somatic symptom and related disorders, binge eating disorders and eating disorders among women with different phenotypes of PCOS

As a result of significant body changes like hirsutism, irregular menses, obesity, acne and hair thinning Women with PCOS may report clinically significant symptoms of anxiety or depression [2, 7]. Previous research indicates that alterations in body image may contribute to psychological distress among women with PCOS [37]. The neurophysiologic etiology of anxiety and depression is not fully understood but the dysregulation of the HPA axis has been linked to stress and, although less extensively, its putative association with anxiety and depression disorders has also been studied [38].

#### - Type of study

##### Study design

Non-comparative cross-sectional open label study to be carried out over the period of 12 months on PCOS patients attending gynecology outpatient department. Patients with irregular menses and /or subfertility were enrolled as per inclusion and exclusion criteria after taking written informed consent.

Personal medical history to be obtained from every woman according to a customized prepared questionnaire. Menstrual cycle history documentation including a general review since menarche and a detailed recall of the last 2 to 3 year interval. Ovulatory dysfunction was defined as less than eight cycles per year, and regular menstrual cycle as 21–35 days in length. Clinical examination was performed in each person to confirm diagnosis.

Presence of at least two criteria from clinical, hormonal, and abdominal USG category were considered diagnostic of PCOS. Women with complain of irregular menses or oligomenorrhea (absence of menses for 35-182 days) or amenorrhea (absence of menses for > 182 days), signs or symptoms of hyperandrogenism, abdominal USG showing at least 12 follicles (2-9 mm in diameter) arranged peripherally around a dense core of ovarian stroma or scattered throughout an increased amount of stroma were enrolled in the study. Transvaginal ultrasound (LogIQp3ultrasonic machine, Ge) was used for all patients. Cut-off body mass index (BMI) with body fat as Standard Consensus Statement for Indian population was considered, i.e., Normal BMI: 18.0-22.9 kg/m<sup>2</sup>, Overweight: 23.0-24.9 kg/m<sup>2</sup> Obesity: >25 kg/m<sup>2</sup> BMI ≥ 25 was considered as obese [39].

Hirsutism was scored according to modified Ferriman Gallaway score [40]. Grading of severity based on the score was assessed as <8 normal, 8-14 - moderate,  $\geq 15$  - severe.

Recruited women have their body fat percentage calculated using Health Sense BF 414 Ultra Lite Body Fat Monitor. The bioelectrical impedance analysis (BIA) method was used to estimate body fat percentage. The general principle behind BIA: two or more conductors are attached to a person's body and a small electric current is sent through the body. The resistance between the conductors will provide a measure of body fat between a pair of electrodes, since the resistance to electricity varies between adipose, muscular and skeletal tissue. Fat-free mass (muscle) is a good conductor as it contains a large amount of water (approximately 73%) and electrolytes, while fat is anhydrous and a poor conductor of electric current. Factors that affect the accuracy and precision of this method include instrumentation, subject factors, technician skill, and the prediction equation formulated to estimate the fat-free mass. There is little scope for technician error as such, but factors such as eating, drinking and exercising must be controlled [41] since hydration level is an important source of error in determining the flow of the electric current to estimate body fat. The instructions for use of instruments typically recommended not making measurements soon after drinking or eating or exercising, or when dehydrated. Population-specific equations used for making them more reliable [41].

### Sample size

Non random sampling, Sequential inclusion of the women who met the study criteria, Inclusion [42, 43]

- 1) Clinical or biochemical hyperandrogenism Increased serum androgens and/or progressive hirsutism
- 2) Oligo-/anovulation Oligo-/amenorrhea for at least 2 years, or primary amenorrhea by age 16 years.
- 3) Polycystic ovarian Morphology Ovarian volume  $>10$  cm<sup>3</sup>.

### Exclusion

Patients having any major systemic illness, congenital adrenal hyperplasia, hyperprolactinemia, acromegaly, functional hypothalamic amenorrhea, and patients receiving drugs for any other systemic illness. In patients complaining of amenorrhea, pregnancy was ruled out whenever necessary.

### Withdrawal criteria

The patients were withdrawn from study due to Loss of follow up, Withdrawal of consent

## METHODOLOGY

Ultrasound (LogIQp3ultrasonic machine, Ge) was used for all patients in defining the ovarian volume and morphology. The psychiatric illness were identified using standardized questionnaires

### Data management & statistical analysis

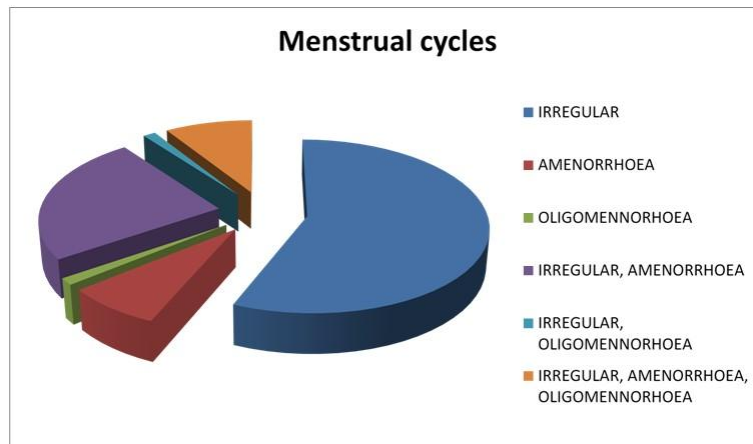
Data was collected in outpatient departments of obstetrics and gynecology

### Statistical analysis

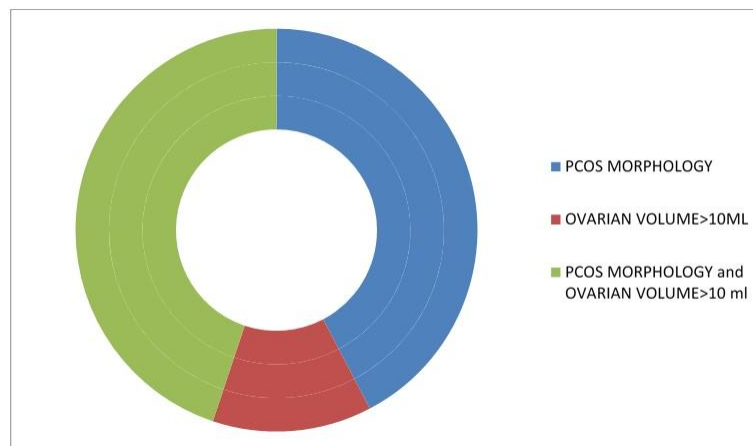
Data analysis to be performed using the SPSS. The Student's t-test to be used for inter-group comparisons of continuous variables. Fisher's least significant difference (LSD) post hoc test to be used to determine significant differences between groups. Categorical variables to be compared using chi-square tests. Results are expressed as mean  $\pm$  SEM and statistical significance for all analyses was defined as a two-tailed P-value of  $<0.05$ . Data analyzed using Fisher's exact test. P value  $< 0.05$  to be considered significant.

## RESULTS

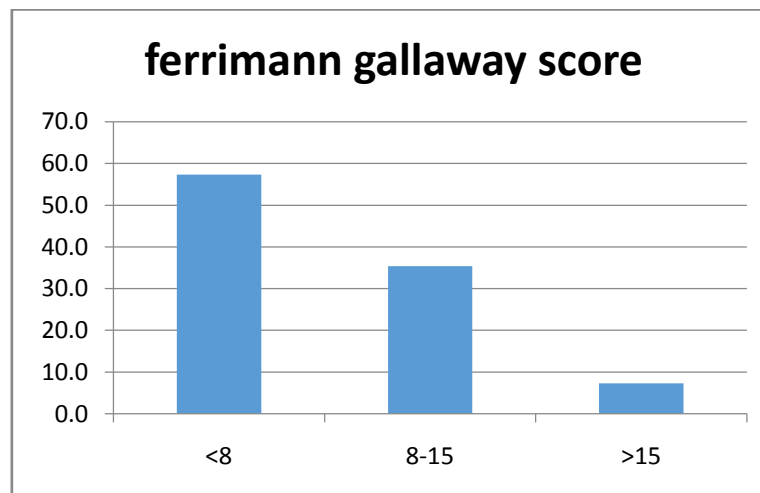
The most common menstrual abnormality identified in study was irregular menstrual cycles followed by a combination of amenorrhea and irregular cycles. On ultrasound pcos morphology was found 42.3% and ovarian volume was more than 10ml in 12.8%. Simultaneous occurrence of both the features occurred in 44.9% Hirsutism of more than 8 ferrimangallaway score was seen in 42.7% cases. In this study we demonstrated the commonest phenotype was ovulatory dysfunction with polycystic morphology in 42.7% cases. Classical phenotype D: ovulatory dysfunction, androgen excess and polycystic ovarian morphology was demonstrated in 28.8% cases we also could demonstrate the body dysmorphic disorder was the commonest psychological problem in about 25.6% closely followed by major depressive illness.



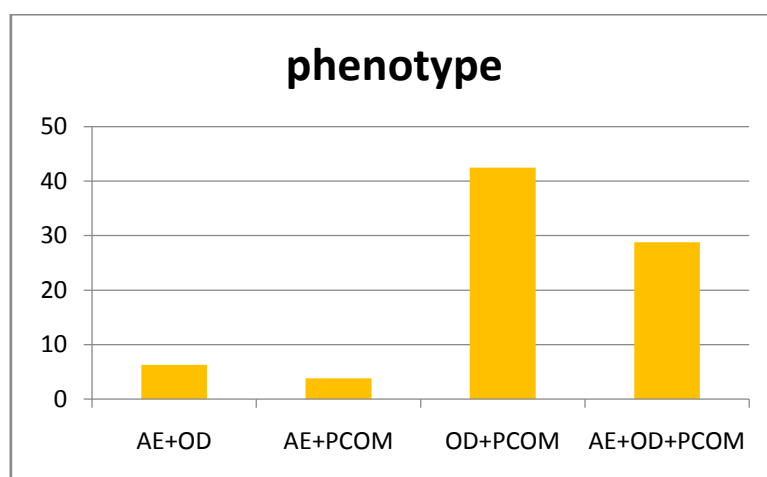
**Figure: 1**



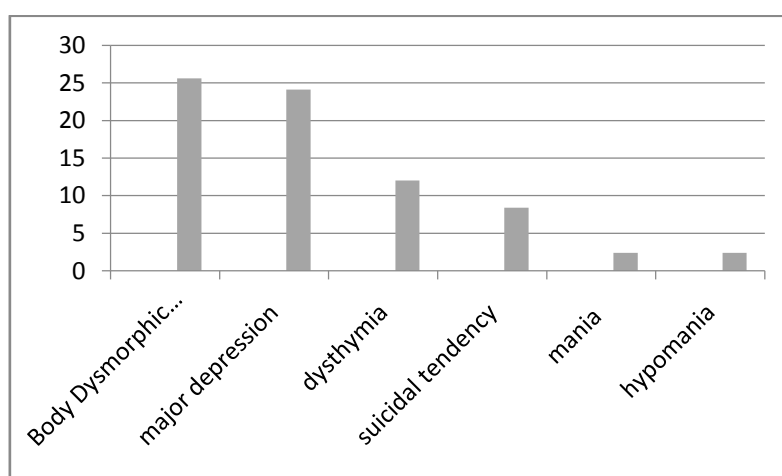
**Figure: 2**



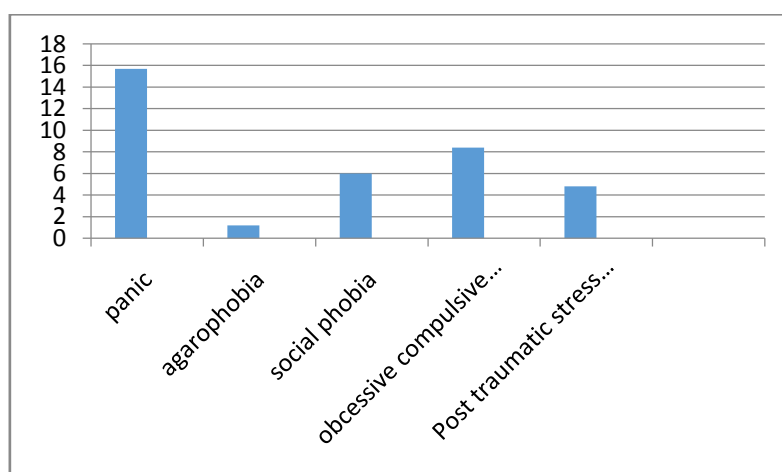
**Figure: 3**



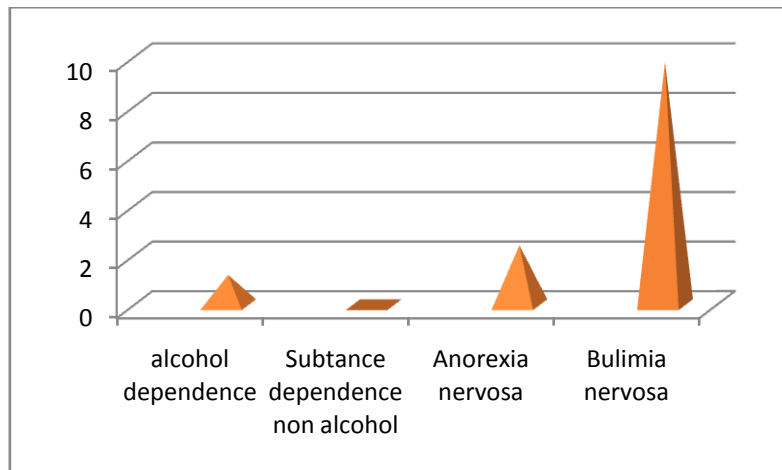
**Figure: 4**



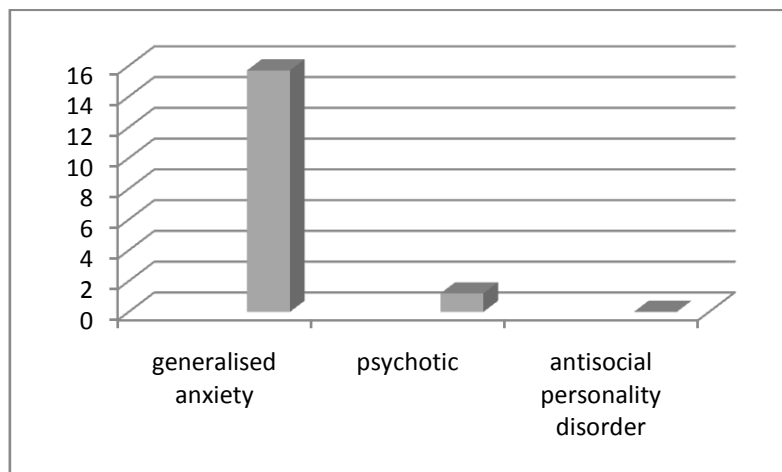
**Figure: 5**



**Figure: 6**



**Figure: 7**



**Figure: 8**

## DISCUSSION

Polycystic ovary syndrome (PCOS) being one of the most common endocrine disorders, empirical evidence shows relationship between psychological factors and physiological changes in PCOS women [44]. In this study, we demonstrated that levels of anxious and depressive symptoms were significantly higher in patients with PCOS the incidence of phobia also is higher. Mc Cook et al. [46] For instance, depression, increased incidence of psychological disorders, increased susceptibility to external pressures, and decreased quality of life could be observed commonly in PCOS patients [47]. The prevalence of depression in PCOS women has been reported to be as high as 24.1 [48].

Elsenbruch et al. [49] reported that PCOS patients have significantly higher chance of developing obsessive-compulsive disorder, interpersonal stress and depressive symptoms than healthy population.

Our study suggested that psychological and neurological factors may play a role in the pathophysiology of PCOS [49]. These findings indicate that PCOS patients usually have obvious psychological problems. Our study demonstrated that anxiety and depression are also significantly associated with PCOS. Therefore, psychological intervention should be an aspect of PCOS management [50]

**Conflict of Interest:** The authors declare that we have no conflict of interest.

## REFERENCES

1. 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
2. Azziz R, Carmina E, Dewailly D, Diamanti-Kandarakis E, Escobar-Morreale HF et al. (2006) Positions statement: criteria for defining polycystic ovary syndrome as a predominantly hyperandrogenic syndrome: an Androgen Excess Society guideline. *J ClinEndocrinolMetab* 91: 4237-45.
3. Yildiz BO, Bozdag G, Yapici Z, Esinler I, Yarali H. Prevalence, phenotype and cardiometabolic risk of polycystic ovary syndrome under different diagnostic criteria. *Hum Reprod* 2012;27:3067-73.



4. Puurunen J, Piltonen T, Morin-Papunen L, Perheentupa A, Jarvela I, Ruokonen A, et al. Unfavorable hormonal, metabolic, and inflammatory alterations persist after menopause in women with PCOS. *J ClinEndocrinolMetab* 2011;96:1827–34.
5. Shaw LJ, BaireyMerz CN, Azziz R, Stanczyk FZ, Sopko G, BraunsteinGD, et al. Postmenopausal women with a history of irregular menses and elevated androgen measurements at high risk for worsening cardiovascular event-free survival: results from the National Institutes of Health–National Heart, Lung, and Blood Institute sponsored Women’s Ischemia Syndrome Evaluation. *J ClinEndocrinolMetab* 2008;93:1276–84.
6. Elsenbruch S, Hahn S, Kowalsky D, Offner AH, SchedlowskiM, Mann K, et al. Quality of life, psychosocial well-being, and sexual satisfaction in women with polycystic ovary syndrome. *J Clin Endocrinol Metab* 2003; 88:5801–7.
7. Arshad Hussain, Rajesh Kumar Chandel, Mohd Ashraf Ganie1, Mansoor Ahmad Dar, Yasir Hassan Rather,Zaid Ahmad Wani, Javid Ahmad Shiekh et al Prevalence of Psychiatric Disorders in Patients witha Diagnosis of Polycystic Ovary Syndrome in Kashmir Indian Journal of Psychological Medicine 2015 :37 :66-70
8. Jedel E, Waern M, Gustafson D, Landen M, Eriksson E, Holm G, et al. Anxiety and depression symptoms in women with polycystic ovary syndrome compared with controls matched for body mass index. *Hum Reprod* 2010; 25:450–6.
9. Dunaif A. Polycystic ovary syndrome in 2011: genes, aging and sleep apnea in polycystic ovary syndrome. *Nat Rev Endocrinol.* 2012;8(2):72–74.
10. Rotterdam ESHRE/ASRM-Sponsored PCOS Consensus Workshop Group.Revised 2003 consensus on diagnostic criteria and long-term health risks related to polycystic ovary syndrome. *FertilSteril* 2004;81:19–25.
11. Rotterdam ESHRE/ASRM-Sponsored PCOS Consensus Workshop Group. Revised 2003 consensus on diagnostic criteria and long-term health risks related to polycystic ovary syndrome (PCOS). *Hum Reprod* 2004;19:41–7.
12. Ali AT. Polycystic ovary syndrome and metabolic syndrome. *CeskaGynekol.* 2015;80(4):279–289.
13. Broekmans FJ, Knauff EA, Valkenburg O, Laven JS, Eijkemans MJ, Fauser BC. PCOS according to the Rotterdam consensus criteria: change in prevalence among WHO-II anovulation and association with metabolic factors. *BJOG* 2006;113:1210–7.
14. Baldani DP, Skrgatic L, Simunic V, Zlopasa G, Canic T, Trgovcic I. Characteristics of different phenotypes of polycystic ovary syndrome based on the Rotterdam criteria in the Croatian population. *CollAntropol* 2013;37: 477–82.
15. Belosi C, Selvaggi L, Apa R, Guido M, Romualdi D, Fulghesu AM, et al. Is thePCOS diagnosis solved by ESHRE/ASRM 2003 consensus or could it include ultrasound examination of the ovarian stroma? *Hum Reprod* 2006;21:3108–15.
16. Cupisti S, Haeberle L, Schell C, Richter H, Schulze C, Hildebrandt T, et al.The different phenotypes of polycystic ovary syndrome: no advantages for identifying women with aggravated insulin resistance or impaired lipids. *ExpClinEndocrinol Diabetes* 2011;119:502–8.
17. Dewailly D, Catteau-Jonard S, Reyss AC, Leroy M, Pigny P. Oligoanovulation with polycystic ovaries but not overt hyperandrogenism. *J ClinEndocrinolMetab* 2006;91:3922–7.
18. Mehrabian F, Khani B, Kelishadi R, Kermani N. The prevalence of metabolic syndrome and insulin resistance according to the phenotypic subgroups of polycystic ovary syndrome in a representative sample of Iranian females. *J Res Med Sci* 2011;16:763–9.
19. Hsu MI, Liou TH, Chou SY, Chang CY, Hsu CS. Diagnostic criteria for polycystic ovary syndrome in Taiwanese Chinese women: comparison between Rotterdam 2003 and NIH 1990. *FertilSteril* 2007;88:727–9.
20. Shroff R, Syrop CH, Davis W, Van Voorhis BJ, Dokras A. Risk of metabolic complications in the new PCOS phenotypes based on the Rotterdam criteria. *FertilSteril* 2007;88:1389–95.
21. Melo AS, Vieira CS, Romano LG, Ferriani RA, Navarro PA. The frequency of metabolic syndrome is higher among PCOS Brazilian women with menstrual irregularity plus hyperandrogenism. *ReprodSci* 2011;18:1230–6.
22. Kim JJ, Hwang KR, Choi YM, Moon SY, Chae SJ, Park CW, et al. Complete and metabolic profiles of a large consecutive cohort of untreated Korean women with polycystic ovary syndrome. *FertilSteril* 2014; 101:1424–30.
23. Diamanti-Kandarakis E, Panidis D. Unravelling the phenotypic map of polycystic ovary syndrome (PCOS): a prospective study of 634 women with PCOS. *ClinEndocrinol (Oxf)* 2007;67:735–42.
24. Guastella E, Longo RA, Carmina E. Clinical and endocrine characteristics of the main polycystic ovary syndrome phenotypes. *FertilSteril* 2010;94: 2197–201.
25. Yilmaz M, Isaoglu U, Delibas IB, Kadanali S. Anthropometric, clinical and laboratory comparison of four phenotypes of polycystic ovary syndrome based on Rotterdam criteria. *J ObstetGynaecol Res* 2011;37:1020–6.
26. Guo M, Chen ZJ, Macklon NS, Shi YH, Westerveld HE, Eijkemans MJ, et al.Cardiovascular and metabolic characteristics of infertile Chinese women with PCOS diagnosed according to the Rotterdam consensus criteria. *Reprod Biomed Online* 2010;21:572–80.
27. Wild RA, Carmina E, Diamanti-Kandarakis E, Dokras A, Escobar-Morreale HF, Futterweit W, et al. Assessment of cardiovascular risk and prevention of cardiovascular disease in women with the polycystic ovary syndrome: a consensus statement by the Androgen Excess and PolycysticOvary Syndrome (AE-PCOS) Society. *J ClinEndocrinolMetab* 2010;95:2038–49.
28. Naqvi SH, Moore A, Bevilacqua K, et al. Predictors of depression in women with polycystic ovary syndrome. *Arch WomensMent Health.* 2015;18(1):95–101.

29. Annagur BB, Kerimoglu OS, Tazegul A, Gunduz S, Gencoglu BB. Psychiatric comorbidity in women with polycystic ovary syndrome. *J Obstet Gynaecol Res.* 2015;41(8):1229–1233.
30. Dokras A. Mood and anxiety disorders in women with PCOS. *Steroids.* 2012;77(4):338–341.
31. Davari-Tanha F, Hosseini Rashidi B, Ghajarzadeh M, Noorbala AA. Bipolar disorder in women with polycystic ovarian syndrome (PCO). *Acta Med Iran.* 2014;52(1):46–48.
32. Kerchner A, Lester W, Stuart SP, Dokras A. Risk of depression and other mental health disorders in women with polycystic ovary syndrome: a longitudinal study. *Fertil Steril.* 2009;91(1):207–212.
33. Zueff LN, da Silva Lara LA, Vieira CS, Martins Wde P, Ferriani RA. Body composition characteristics predict sexual functioning in obese women with or without PCOS. *J Sex Marital Ther.* 2015;41(3):227–237.
34. Asik M, Altinbas K, Eroglu M, et al. Evaluation of affective temperament and anxiety-depression levels of patients with polycystic ovary syndrome. *J Affect Dis.* 2015;185:214–218.
35. Moran LJ, Deeks AA, Gibson-Helm ME, Teede HJ. Psychological parameters in the reproductive phenotypes of polycystic ovary syndrome. *Hum Reprod.* 2012;27(7):2082–2088.
36. Hollinrake E, Abreu A, Maifeld M, Van Voorhis BJ, Dokras A. Increased risk of depressive disorders in women with polycystic ovary syndrome. *Fertil Steril.* 2007;87(6):1369–1376.
37. Himelein MJ, Thatcher SS. Depression and body image among women with polycystic ovary syndrome. *J Health Psychol.* 2006;11(4): 613–625.
38. Mueller SC, Ng P, Sinaii N, et al. Psychiatric characterization of children with genetic causes of hyperandrogenism. *Eur J Endocrinol.* 2010;163(5):801–810.
39. Misra A, Chowbey P, Makkar BM, Vikram NK, Wasir JS, Chadha D, et al. Consensus statement for diagnosis of obesity, abdominal obesity and the metabolic syndrome for Asian Indians and recommendations for physical activity, medical and surgical management. *J Assoc Physicians India.* 2009;57:163–70. [PubMed: 19582986]
40. De Ugarte CM, Woods KS, Bartolucci AA, Azziz R. Degree of facial and body terminal hair growth in unselected black and white women: towards a populational definition of hirsutism. *J Clin Endocrinol Metab* 2006;91:1345–50
41. Brown, Stanley P.; Miller, Wayne C. and Eason, Jane M. (2006) Exercise physiology: Basis of Human Movement in Health and Disease, 2nd Ed., p. 324, Lippincott Williams & Wilkins
42. Lizneva. Criteria, prevalence, and phenotypes of PCOS. *Fertil Steril* 2016.
43. Fauser BC, Tarlatzis BC, Rebar RW, Legro RS, Balen AH, Lobo R, et al. Consensus on women's health aspects of polycystic ovary syndrome (PCOS): the Amsterdam ESHRE/ASRM-Sponsored 3rd PCOS Consensus Workshop Group. *Fertil Steril* 2012;97:28–38.e25.
44. Ramasubbu R. Insulin resistance: a metabolic link between depressive disorder and atherosclerotic vascular diseases. *Medical Hypothesis.* 2002;59:537–551.
45. Elsenbruch S, Hahn S, Kowalsky D, et al. Quality of life, psychosocial well-being, and sexual satisfaction in women with polycystic ovary syndrome. *J Clin Endocrinol Metab.* 2003;88:5801–5807.
46. Mc Cook JG, Reame NE, Thatcher SS (2005) Health-related quality of life issues in women with polycystic ovary syndrome. *J Obstet Gynecol Neonatal Nurs* 34:12–20
47. Coffey S, Mason H (2003) The effect of polycystic ovary syndrome on health related quality of life. *Gynecol Endocrinol* 17:379–386
48. Rasgon NL, Rao RC, Hwang S, et al. Depression in women with polycystic ovary syndrome: clinical and biochemical correlates. *J Affect Disord.* 2003;74:299–304.
49. Dokras A, Clifton S, Futterweit W, Wild R (2011) Increased risk for abnormal depression scores in women with polycystic ovary syndrome: a systematic review and meta-analysis. *Obstet Gynecol* 117:145–152
50. Wang Y, Cao Y, Chong L et al (2005) Polycystic ovary syndrome and psychological analysis of the investigation. *Anhui Med Univ (in Chinese)* 40:277–279

Proforma

**Characteristics of body composition in different phenotypes of South Indian women with polycystic ovary syndrome and relation with body image and psychological profile**

NAME		AGE		HOSPITAL NUMBER	
HEIGHT		WEIGHT		BMI	
OVERWEIGHT	CLASS 1 25-29.9	CLASS 2 30-34.9	CLASS 3 35-39.9	CLASS 4/MORBID >40	
WAIST HIP RATIO					
BODY FAT %					
MENSTRUAL CYCLES	IRREGULAR	AMENORRHOEA	OLIGOMENORRHOEA		
USS		PCOS MORPHOLOGY		OVARIAN VOLUME >10ML	



FERRIMAN GALLAWAY	<8	8-15	>15
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PHENOTYPE	AE+OD	AE+PCOM	OD+PCOM	AE+OD+PCOM
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Body dysmorphic disorder
Major depressive episode
Dysthymia
Suicidality
Maniac
Hypomaniac
Panic
Agoraphobia
Social phobia
Obsessive compulsive disorder
Posttraumatic stress disorder
Alcohol dependence/abuse
Substance dependence (non-alcohol)
Psychotic disorders
Anorexia nervosa
Bulimia nervosa
Generalized anxiety disorder
Anti social personality disorder