



Original Article

Potential Effects of Obesity on Anti-Müllerian Hormone in Polycystic Ovary Syndrome

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ABSTRACT

Background: Polycystic ovary syndrome (PCOS) is a common endocrine abnormality with its known effects on reproduction. It is usually associated with infertility. Early detection is crucial the use of Antimüllerian hormone (AMH) as indicator of ovarian function could be a reliable indicator. The association between PCOS and other metabolism-related disorders (e.g., obesity) is not sufficiently studied.

Aim of the work: we carried out this study aiming to investigate the possible association between AMH, ovarian reserve, AFC with BMI in women with PCOS. Generally, we seek the link between obesity and PCOS.

Methods: The study included 240 women with confirmed diagnosis of PCOS. They were categorized according to body mass index (BMI) into three groups the first for normal BMI (group A; n=100); the second for overweight women (Group B; n=65) and the third for obese women (group C; n=75). All were clinically (by history and examination) and radiologically (pelvic ultrasound) evaluated and then AMH, beside others were measured in plasma and data was compared between groups.

Results: Study groups were comparable regarding their age and incidence of hirsutism, ovarian volume. Hirsutism for example reported in 16%, 15.4% and 25.3% of normal weight, overweight and obese groups, respectively. Otherwise, waist circumference, hip circumference and waist/hip ratio were significantly increased progressively with increasing BMI. Acne also increased in C and B than in normal A group (41.3%, 21.5% vs 16.0%, respectively). By ultrasound, most follicles were peripheral (69.0%, 73.8% and 72.0% in groups A, B and C respectively). Furthermore, AMH and the antral follicular count (AFC) showed progressive increase in B and C groups than normal weight group (5.49 ± 1.20 , 18.61 ± 6.40 ; 8.13 ± 1.88 , 23.73 ± 5.94 versus 2.97 ± 0.66 and 14.96 ± 2.88 , respectively). Finally, AMH was significantly and proportionately correlated with BMI, waist/hip ratio, ovarian volume and AFC. The correlation with BMI, W/H ratio and AFC was powerful ($r > 0.7$). In addition, there was moderate, significant and statistically significant correlation between AFC and each of BMI, W/H ratio and mean ovarian volume. However, the correlation between AFC and age was non-significant.

Conclusion: AMH showed significant increase with obesity, with positive correlation with BMI and ANF. Thus, BMI can play a crucial role in the development of PCOS.

Keywords: Polycystic Ovary; Body Mass Index; Hyperandrogenism; Infertility.



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INTRODUCTION

Polycystic ovary syndrome (PCOS) is a common endocrine disorder affecting 6-10% of women in their fertile period. It is associated with infertility as the incidence of PCOS is reported in 30-40% of infertile women, which increased to 70-80.0% of anovulatory infertile women. In Egypt the prevalence is around 35.0% of overall infertile women ⁽¹⁻⁴⁾.

PCOS is characterized by irregular menstruation, disturbances in glucose metabolism with development of insulin resistance (IR), oxidative stress, hormonal disturbances, androgen excess and polycystic ovaries ^(5,6).

Insulin resistances seem to be a major pathophysiological mechanism in development of polycystic ovaries and may be responsible for reproductive anomalies associated with PCOS ^(7,8).

In addition, abnormal lipids (dyslipidemia) are common findings in PCOS. This lipid profile changes in PCOS include decreased HDL-C (high density lipoprotein cholesterol), elevated triglycerides (TGs), and increased levels of LDL-C (low density lipoprotein cholesterol) ^(9,10).

Anti-Mullerian hormone (AMH) has been introduced as a normal indicator of ovarian function. It is a member of glycoprotein family known as transforming growth factor- β superfamily. The physiological function of anti-Mullerian hormones includes its important role in excessive early follicular growth and the maturation of the dominant follicle (i.e., promotion of follicular arrest). Increased AMH is associated with excessive increase in the antral follicular count (AFC). However, its association with discrimination of PCOS phenotypes and metabolic disturbances ⁽¹¹⁻¹³⁾.

Obesity is a disease defined by excessive storage of triglycerides in fatty cells, which increased in number and size. Body mass index more than 30 kg/m² is the cutoff value for diagnosis of obesity. In addition, body mass index (BMI) was used to categorize obesity into different grades. Overweight and obesity have impacts on the general health as well as reproductive function ^(14,15).

In infertile women seeking pregnancy by natural methods, obesity has a significant adverse effect on conception and outcome pregnancy. Increased BMI was associated with reduction in fertility, increased miscarriage rate, when compared to normal-weight females ^(16,17). Thus, reduction of weight before conception may be associated with improvement of fertility and pregnancy

outcome. Poor outcome was also noticed in obese women submitted to assisted reproductive technologies (ARTs) ^(18,19). However, the exact mechanism linking obesity to infertility and poor outcome of pregnancy or ARTs is not yet fully elucidated. Obesity contributes or oligo or anovulation, menstrual disturbances, reduced rate of conception, and poor response to different treatment modalities for infertility. Different indicators of ovarian reserve are used to investigate women regarding their reproductive outcome. These indicators include, but are not limited to, baseline follicle stimulating hormone (FSH, Estradiol₂ (E₂), Inhibin B, Antral follicle count (AFC), ovarian volume and antimullerian hormone (AMH) ⁽²⁰⁾.

Previous studies reported possible association between AMH and obesity. But, with conflicting results ^(21, 22).

We hypothesize that, the AMH linked to body mass index in PCOS can be a good indicator of ovarian reserve and antral follicles count. Thus, we carried out this study aiming to investigate the possible association between AMH, ovarian reserve, AFC with BMI in women with PCOS. Generally, we seek the link between obesity and PCOS.

PATIENTS AND METHODES

This was a prospective comparative study. It was carried out in Damietta Specialized Hospital and Al-Azhar University hospital (New Damietta). It was completed during the duration between the first of March 2019 and the end of August 2020. At first 300 women with confirmed diagnosis of PCOS were screened for eligibility criteria and were asked for participation in the study. Sixty of them were excluded due to different causes (mainly refusal to participate or withdrawn after inclusion according their well). Thus, the final analysis was performed for 240 women.

The diagnosis of PCOS was based on the Rotterdam 2003 criteria. (Clinical hyperandrogenism (Ferriman-Gallwey Score \geq 8) or biochemical hyperandrogenism (elevated total/free testosterone); oligomenorrhea (less than 6-9 menses per year) or oligo-ovulation; polycystic ovaries on ultrasound (\geq 12 antral follicles in one ovary or ovarian volume \geq 10 cm³). At least two of these criteria must be detected to diagnose PCOS. Then patients were categorized according to their BMI into three groups. The first was for women with normal weight (BMI < 25kg/m²) (100 women). The second for overweight women (BMI between 25-3025kg/m²) (65 women) and the third group for obese women (BMI >30 kg/m²) (n=75).

The inclusion criteria were women during their reproductive age (18-35 years), with confirmed diagnosis of PCOS according to previous criteria and they provided consent to participate. On the other side, exclusion criteria were pregnant women and breast feeding.

Methods:

All eligible women were clinically evaluated by complete history taking, clinical and radiological examination. In clinical examination, the general condition was evaluated, and special characters (acne and hirsutism) were recognized and documented. Acne was defined by a history of persistent acne (presence of acne on the most days for 3 years or more), recent acne treatment and presence of more than 10 inflammatory acne lesions). In addition, hirsutism was evaluated by Ferriman-Gallwey (FG) scoring system (score < 8 is indicator of hirsutism). The FG score is used to quantify hirsutism in women. The method was initially published in 1961 by Ferriman and Gallwey. Then modified to reduce body areas for assessment of hair growth from 11 to 9 (23,24). These areas include upper lip, chin, chest, upper and lower back (2 areas), upper and lower abdomen (two areas), upper arms and thighs. The forearm and legs were in the original form but removed in the modified score. In the modified method, hair growth is rated from no growth of terminal hair (stage 0) to extensive hair growth (grade 4) in each of the nine locations. A patient's score may therefore range from a minimum score of 0 to a maximum score of 36. A score of 8 or higher is set as indicator of androgen excess (hirsutism).

The body mass index was calculated by the equation $BMI = \text{weight in kg} / \text{height in m}^2$). Other measures include waist/hip ratio and presence of central adiposity. The pelvis was examined for the presence of adnexal masses. Then pelvic ultrasound was performed to investigate ovarian criteria of PCOS (12 or more follicles measuring 2-9 mm and/or an increased ovarian volume of $>10\text{cm}^3$). It was performed in the early follicular phase (day 1-3) in lithotomy position as described elsewhere. Number and site of antral follicles was counted in the longest section of the ovary, and for better results, it was counted in the whole ovary by taking a 2D sweep across whole ovary. This method is very feasible and reliable when number of follicles is much more than in PCOS.

Finally, laboratory estimation of serum levels of AMH and

testosterone were performed.

Statistical analysis of data: The collected data were organized in an excel sheet, then tabulated and statistically analyzed using a software computer package (the statistical package of social science (SPSS), version 18 (IBM ©SPSS Inc, USA), running on IBM compatible personal computer (PC). Qualitative data were summarized by the relative frequency and percentage and compared by the appropriate test (Chi square (X^2) or Mann Whitney test). Quantitative data represented by the arithmetic mean and standard deviation (SD), minimum and maximum (difference is the range); and for comparison between groups, the one-way analysis of variance (ANOVA) test was used. P value < 0.05 was considered significant.

RESULTS

In the current work, study groups were comparable regarding patient age and incidence of hirsutism. However, waist circumference, hip circumference and waist/hip ratio were significantly increased progressively with increasing BMI. Acne also increased in groups C and B than in group-A (41.3%, 21.5% vs 16.0%, respectively) (Table 1).

The study groups showed non-significant differences between groups regarding ovarian volume (right, left and mean values). In addition, most follicles were peripheral (69.0%, 73.8% and 72.0% in groups A, B and C respectively). The difference between groups B and C were significantly indifferent. Furthermore, AMH and the antral follicular count (AFC) showed progressive increase in overweight and obese women than normal weight group (5.49 ± 1.20 , 18.61 ± 6.40 ; 8.13 ± 1.88 , 23.73 ± 5.94 versus 2.97 ± 0.66 and 14.96 ± 2.88 , respectively) (Table 2).

AMH was significantly and positively (proportionately) correlated with body mass index (BMI), waist/hip ratio, ovarian volume and antral follicular count. The correlation with BMI, W/H ratio and AFC was powerful ($r > 0.7$), while it was moderate with other variables. However, no significant correlation was reported between AMH and patient age (Table 3).

In addition, there was moderate, significant and statistically significant correlation between AFC and each of BMI, W/H ratio and mean ovarian volume. However, the correlation between AFC and age was non-significant (Table 4).

Table (1): Demographic and clinical data of study groups

| Variable | Group A (<25) (n= 100) | Group B (25-29) (n=65) | Group C (>30) (n=75) | Test | p |
|---------------------|---------------------------|---------------------------|-------------------------|---------|---------|
| Age (years) | 27.25±2.43 | 27.35±2.30 | 27.57±2.23 | 0.412 | 0.661 |
| Waist Circumference | 62.68±1.39 | 72.47±3.50 | 84.54±2.24 | 1798.20 | <0.001* |
| Hip circumference | 84.73±3.44 | 86.04±3.53 | 91.77±3.11 | 99.49 | <0.001* |
| W/H ratio | 0.74±0.03 | 0.85±0.04 | 0.92±0.04 | 573.11 | <0.001* |
| Acne (n,%) | 17(17.0%) | 14(21.5%) | 31(41.3%) | 14.10 | <0.001* |
| Hirsutism (n,%) | 16(16.0%) | 10(15.4%) | 19(25.3%) | 3.113 | 0.211 |

Table (2): Comparison between study groups regarding ovarian volume, site of follicle, AMH and AFC

| Variable | Group A (<25) (n= 100) | Group B (25-29) (n=65) | Group C (>30) (n=75) | F | p |
|------------------|---------------------------|---------------------------|-------------------------|-------|---------|
| Right OV | 12.41±1.09 | 12.42±0.93 | 12.63±0.96 | 1.20 | 0.303 |
| Left OV | 12.63±1.02 | 12.56±0.94 | 12.70±0.98 | 0.361 | 0.698 |
| Mean OV | 12.52±1.01 | 12.49±0.90 | 12.67±0.96 | 0.716 | 0.490 |
| Site of follicle | Peripheral | 69(69.0%) | 48(73.8%) | 0.482 | 0.768 |
| | Mixed | 31 (31.0%) | 17(26.2%) | | |
| AMH | Mean±SD | 2.97±0.66 | 5.49±1.20 | 339.6 | <0.001* |
| | Min.–Max. | 1.8-4.7 | 3.6-9.3 | | |
| AFC | Mean±SD | 14.96±2.88 | 18.61±6.40 | 64.47 | <0.001* |
| | Min.–Max. | 10.0 – 22.0 | 13-34 | | |

Table (3): Correlation between AMH and other variables

| | AMH | |
|----------------------|-------------------------|------------|
| | Pearson Correlation (r) | p |
| Age | 0.026 | 0.694 (NS) |
| BMI | 0.850 | <0.001* |
| Waist hip ratio | 0.736 | <0.001* |
| Right ovarian volume | 0.414 | <0.001* |
| Left ovarian volume | 0.393 | <0.001* |
| Mean ovarian volume | 0.412 | <0.001* |
| AFC | 0.818 | <0.001* |

Table (4): Correlation between AFC and other studied parameters

| | AFC | |
|---------------------|-------------------------|---------|
| | Pearson Correlation (r) | P |
| Age | 0.014 | 0.828 |
| BMI | 0.579 | <0.001* |
| Waist hip ratio | 0.504 | <0.001* |
| Mean ovarian volume | 0.404 | <0.001* |

DISCUSSION

The potential association between serum AMH and body mass index is poorly studied in patients with PCOS. Thus, the current work was designed to investigate such relationships. Included women were categorized into

normal body mass index (group A), overweight (group B) and Obese (group C).

Previous studies had described increased prevalence of obesity in infertile women with or without PCOS ⁽²⁵⁾. These reflected the importance of investigating the relationship

between obesity and PCOS. This explains the design and conduct of this study

The mean age in the study groups was 27.25 ± 2.43 , 27.35 ± 2.30 and 27.57 ± 2.23 years in groups A, B and C respectively, with no significant difference between groups. These results are comparable to **Cengiz *et al.*** ⁽²⁶⁾ who reported non-significant differences between obese and non-obese women with PCOS as regard to their age.

Acne was significantly increased with obesity (progressive increase with BMI). However, the hirsutism showed non-significant difference between groups. In addition, the overall prevalence of hirsutism associated with PCO in this study was (18.8%). This value is low when compared to previous studies (39.0% and 29.0%) ^(27,28). This lower incidence of hirsutism in the current work than previous studies may be attributed to the different selection and diagnostic criteria for hirsutism. However, others reported non-significant difference between obese and non-obese women with PCOS in the study of **Dos Reis *et al.*** ⁽²⁹⁾. These results are in line with the current work. In addition, **Legro *et al.*** ⁽³⁰⁾ reported higher prevalence and severity of hirsutism with increased body weight. However, the difference between groups was not significant from the statistical point of view.

Results of the current work showed non-significant differences between study groups regarding ovarian volume (right, left or mean values). However, **Balen *et al.*** ⁽³¹⁾ reported that increased ovarian volume is a characteristic feature of PCOS, which was not reflected in the current work. Again, this may be different selection and diagnostic criteria.

Most of study females (about three fourths), had peripheral distribution of follicles in the ovary, with no significant difference between groups. These confirm the results of **Guraya *et al.*** ⁽²⁷⁾, who reported peripheral distribution of follicles among 82.8 of women with PCOS

AMH was significantly increased in groups B and C than group A (i.e., there was progressive increase of AMH

with increased BMI). These results are comparable to previous studies stated that, plasma AMH was significantly different between normal, overweight and obese adolescent females with PCOS. Interestingly, the same situation was reported for women without PCOS, reflecting the association between increased BMI and higher serum levels of AMH ⁽³²⁻³⁴⁾.

However, contradictory results are reported by **Cengiz *et al.*** ⁽²⁶⁾ who did not find significant difference between normal and obese women with PCOS as regard to AMH levels. This can be explained by the different age groups, as those authors included adolescent girls.

On the extreme side, **Legro *et al.*** ⁽³⁰⁾ reported significant reduction of AMH with increased body mass index, especially in severe obesity. This can be attributed to different selection criteria as they compared morbid obesity to non-obese PCOS. Their results are reported by other researchers ⁽³⁵⁾.

To explain the ovulatory derangements in PCOS, **Jonard and DeWailly** ⁽³⁶⁾ reported that, these changes are due to different etiologies (changes): 1) increased follicular growth in early phases, which lead to a larger than normal reserve of these follicles; 2) inability to select one follicle of this pool leading to what is known as follicular arrest.

AMH is known to as an inhibitor of the initial follicle recruitment ⁽³⁷⁾ and causes follicular arrest ⁽³⁸⁾. In addition, AMH levels are increased in PCOS. Thus, it is postulated that its involvement in PCOS-associated anovulation depends only on the follicular arrest ⁽³⁶⁾. However, Jonard and DeWailly reported that the higher AMH production in PCOS is due to increased number of small follicles. However, this was not supported by other studies.

Searching correlation between AMH and other variables revealed significantly positive correlation with BMI, W/H ratio and ovarian volume. However, the correlation with age was statistically non-significant.

These results are contradictory to **Skalba *et al.*** ⁽³⁹⁾ who

cannot find any significant correlation between BMI and serum AMH in obese women. But they can find such correlation in women with normal weight PCOS. However, in the current work, we studied the correlation among all study females and do not calculate correlation separately for each group. However, the same authors reported a weak impact of body mass index on the AMH values in women in their reproductive age. It is important to confirm that, their patients were younger than women of the current work.

In the present study, antral follicular count (AFC), was significantly increased with higher BMI. This cannot be confirmed in the previous study **Legro *et al.*** ⁽³⁰⁾. Nonetheless they reported lower AFC in obese than non-obese women. This may be due to difference in selection criteria or methods used to estimate AFC.

In the present study, correlation between AFC was significantly positive with BMI, W/H ratio, mean ovarian volume and AMH. But it was not significant with patient age. These results are in line with previous studies reporting a direct and significant correlation between AFC and AMH. They explained higher values of AMH in PCOS by higher number of early antral follicles ⁽⁴⁰⁾. However, others stated that the increase in AMH is largely due to the increase of AMH by each follicle, not due to increased number of early antral follicles ⁽⁴¹⁾. Anyway, the changes in antral follicles either in count or size and early maturation are responsible for increased AMH. Thus, AMH can be used as an indicator for ovarian aging due to its correlation with the number and early antral follicles ⁽⁴⁰⁾.

Elmashad ⁽⁴²⁾ also reported a significant positive correlation between AMH levels with ovarian volume and the AFC in PCOS, which was not surprising as ovarian volume reflects the number of small antral follicles present in PCOS, which are the only source of AMH. Furthermore, a significant correlation was reported between serum values of AMH in PCOS and hyperandrogenism in the current and previous studies ^(40,43).

In short, the current work showed increased levels of AMH in PCOS with obesity. These values positively correlated with BMI and AFC. Thus, we can say that body weight can play a role in pathogenesis and development of PCOS. However, the cause effect relationship needs further investigation in future studies.

Conflict of interest: None

Financial disclosure: None

REFERENCES

1. Sanad AS. Prevalence of polycystic ovary syndrome among fertile and infertile women in Minia Governorate, Egypt. *Int J Gynaecol Obstet.* 2014 Apr;125(1):81-2. doi: 10.1016/j.ijgo.2013.09.025.
2. Pirotta S, Joham A, Grieger JA, Tay CT, Bahri-Khomami M, Lujan M, Lim SS, Moran LJ. Obesity and the Risk of Infertility, Gestational Diabetes, and Type 2 Diabetes in Polycystic Ovary Syndrome. *Semin Reprod Med.* 2020 Nov;38(6):342-351. doi: 10.1055/s-0041-1726866.
3. Rafique M, Nuzhat A, Al-Jaroudi D. Risk of Infertility Index in Women with Polycystic Ovarian Syndrome. *J Coll Physicians Surg Pak.* 2020 Nov;30(11):1188-1192. doi: 10.29271/jcpsp.2020.11.1188.
4. Hamoda ME, Megahed AM, Oun AM. Effects of Different Methods of Laparoscopic Ovarian Drilling and its Outcome in Patients with Polycystic Ovary Syndrome. *Int J Med Arts* 2021; xx: xx-xx [Article in Press]. DOI: 10.21608/IJMA.2021.78712.1328.
5. Zhou L, Ni Z, Yu J, Cheng W, Cai Z, Yu C. Correlation Between Fecal Metabolomics and Gut Microbiota in Obesity and Polycystic Ovary Syndrome. *Front Endocrinol (Lausanne).* 2020 Sep 8; 11:628. doi: 10.3389/fendo.2020.00628.
6. Louwers YV, Laven JSE. Characteristics of polycystic ovary syndrome throughout life. *Ther Adv Reprod Health.* 2020 Mar 18;14:2633494120911038. doi: 10.1177/2633494120911038.
7. Fonseka S, Subhani B, Wijeyaratne CN, Gawarammana IB, Kalupahana NS, et al. Association between visceral adiposity index, hirsutism and cardiometabolic risk factors in women with polycystic ovarian syndrome: A cross-sectional study. *Ceylon Med J.* 2019 Sep 30;64(3):111-117. doi: 10.4038/cmj.v64i3.8958.
8. Pereira-Eshraghi CF, Chiuzan C, Zhang Y, Tao RH, McCann M, Neugut YD, et al. Obesity and Insulin Resistance, Not Polycystic Ovary Syndrome, Are Independent Predictors of Bone Mineral Density in Adolescents and Young Women. *Horm Res Paediatr.* 2019;92(6):365-371. doi: 10.1159/000507079.

9. Liu Q, Xie YJ, Qu LH, Zhang MX, Mo ZC. Dyslipidemia involvement in the development of polycystic ovary syndrome. *Taiwan J Obstet Gynecol*. 2019 Jul;58(4):447-453. doi: 10.1016/j.tjog.2019.05.003.
10. Macut D, Bjekić-Macut J, Savić-Radojević A. Dyslipidemia and oxidative stress in PCOS. *Front Horm Res*. 2013;40:51-63. doi: 10.1159/000341683.
11. Zhao Y, Zhao Y, Wang C, Liang Z, Liu X. Diagnostic value of anti-müllerian hormone as a biomarker for polycystic ovary syndrome: a meta-analysis update. *Endocr Pract*. 2019 Oct;25(10):1056-1066. doi: 10.4158/EP-2019-0098.
12. Lie Fong S, Laven JSE, Duhamel A, Dewailly D. Polycystic ovarian morphology and the diagnosis of polycystic ovary syndrome: redefining threshold levels for follicle count and serum anti-Müllerian hormone using cluster analysis. *Hum Reprod*. 2017 Aug 1;32(8):1723-1731. doi: 10.1093/humrep/dex226.
13. Hwang YI, Sung NY, Koo HS, Cha SH, Park CW, Kim JY, et al. Can high serum anti-Müllerian hormone levels predict the phenotypes of polycystic ovary syndrome (PCOS) and metabolic disturbances in PCOS patients? *Clin Exp Reprod Med*. 2013 Sep;40(3):135-40. doi: 10.5653/term.2013.40.3.135.
14. Oldfield AL, Kazemi M, Lujan ME. Impact of Obesity on Anti-Müllerian Hormone (AMH) Levels in Women of Reproductive Age. *J Clin Med*. 2021 Jul 20;10(14):3192. doi: 10.3390/jcm10143192.
15. Sahmay S, Usta T, Erel CT, Imamoğlu M, Küçük M, Atakul N, Seyisoğlu H. Is there any correlation between amh and obesity in premenopausal women? *Arch Gynecol Obstet*. 2012 Sep;286(3):661-5. doi: 10.1007/s00404-012-2363-x.
16. Ramezanzadeh F, Kazemi A, Yavari P, Nasr-Esfahani MH, Nejat S, Rahimi-Foroshani A, Saboor-Yaraghi A. Impact of body mass index versus physical activity and calorie intake on assisted reproduction outcomes. *Eur J Obstet Gynecol Reprod Biol*. 2012 Jul;163(1):52-6. doi: 10.1016/j.ejogrb.2012.03.035.
17. Soritsa D, Mäestu E, Nuut M, Mäestu J, Migueles JH, Läänelaid S, et al. Maternal physical activity and sedentary behaviour before and during in vitro fertilization treatment: a longitudinal study exploring the associations with controlled ovarian stimulation and pregnancy outcomes. *J Assist Reprod Genet*. 2020 Aug;37(8):1869-1881. doi: 10.1007/s10815-020-01864-w.
18. Candeloro M, Di Nisio M, Ponzano A, Tiboni GM, Potere N, Tana M, Rutjes AWS, Porreca E. Effects of Obesity and Thrombophilia on the Risk of Abortion in Women Undergoing In Vitro Fertilization. *Front Endocrinol (Lausanne)*. 2020 Dec 23; 11:594867. doi: 10.3389/fendo.2020.594867.
19. Mintziori G, Nigdelis MP, Mathew H, Mousiolis A, Goulis DG, Mantzoros CS. The effect of excess body fat on female and male reproduction. *Metabolism*. 2020 Jun;107:154193. doi: 10.1016/j.metabol.2020.154193.
20. Anderson KS, Segars JH. Predicting fertility with antimüllerian hormone: is a cutoff value adequate? *Fertil Steril*. 2012 Dec;98(6):1421-2. doi: 10.1016/j.fertnstert.2012.08.054.
21. Georgopoulos NA, Saltamavros AD, Decavalas G, Piouka A, Katsikis I, Panidis D. Serum AMH, FSH, and LH levels in PCOS. *Fertil Steril*. 2010 Feb;93(3):e13; author reply e14. doi: 10.1016/j.fertnstert.2009.10.006.
22. Park AS, Lawson MA, Chuan SS, Oberfield SE, Hoeger KM, Chang RJ. Serum anti-müllerian hormone concentrations are elevated in oligomenorrheic girls without evidence of hyperandrogenism. *J Clin Endocrinol Metab*. 2010 Apr;95(4):1786-92. doi: 10.1210/jc.2009-2106.
23. Ferriman D, Gallwey JD. Clinical assessment of body hair growth in women. *J Clin Endocrinol Metab*. 1961 Nov; 21:1440-7. doi: 10.1210/jcem-21-11-1440.
24. Lunde O, Grottum P. Body hair growth in women: normal or hirsute. *Am J Phys Anthropol*. 1984 Jul;64(3):307-13. doi: 10.1002/ajpa.1330640313.
25. Barber TM, Golding SJ, Alvey C, Wass JA, Karpe F, Franks S, McCarthy MI. Global adiposity rather than abnormal regional fat distribution characterizes women with polycystic ovary syndrome. *J Clin Endocrinol Metab*. 2008 Mar;93(3):999-1004. doi: 10.1210/jc.2007-2117.
26. Cengiz H, Ekin M, Dagdeviren H, Yildiz Ş, Kaya C, Kanawati A. Comparison of serum anti-Müllerian hormone levels in normal weight and overweight-obese adolescent patients with polycystic ovary syndrome. *Eur J Obstet Gynecol Reprod Biol*. 2014 Sep; 180:46-50. doi: 10.1016/j.ejogrb.2014.06.018.
27. Guraya SS. Prevalence and ultrasound features of polycystic ovaries in young unmarried Saudi females *Journal of Microscopy and Ultrastructure* 2013; 1:30–34
28. Diamanti-Kandarakis E, Kouli CR, Bergiele AT, Filandra FA, Tsianateli TC, Spina GG, Zapanti ED, Bartzis MI. A survey of the polycystic ovary syndrome in the Greek island of Lesbos: hormonal and metabolic profile. *J Clin Endocrinol Metab*. 1999 Nov;84(11):4006-11. doi: 10.1210/jcem.84.11.6148.
29. dos Reis RM, Foss MC, de Moura MD, Ferriani RA, Silva de Sá MF. Insulin secretion in obese and non-obese women with polycystic ovary syndrome and its relationship with hyperandrogenism. *Gynecol Endocrinol*. 1995 Mar;9 (1): 45-50. doi: 10.3109/09513599509160190.

30. Legro RS, Brzyski RG, Diamond MP, Coutifaris C, Schlaff WD, Alvero R, et al; National Institute of Child Health and Human Development Reproductive Medicine Network. The Pregnancy in Polycystic Ovary Syndrome II study: baseline characteristics and effects of obesity from a multicenter randomized clinical trial. *Fertil Steril*. 2014 Jan;101(1):258-269.e8. doi: 10.1016/j.fertnstert.2013.08.056.
31. Balen AH, Laven JS, Tan SL, Dewailly D. Ultrasound assessment of the polycystic ovary: international consensus definitions. *Hum Reprod Update*. 2003 Nov-Dec;9(6):505-14. doi: 10.1093/humupd/dmg044.
32. Gracia CR, Freeman EW, Sammel MD, Lin H, Nelson DB. The relationship between obesity and race on inhibin B during the menopause transition. *Menopause*. 2005 Sep-Oct;12(5):559-66. doi: 10.1097/01.gme.0000172268.24949.94.
33. Freeman EW, Gracia CR, Sammel MD, Lin H, Lim LC, Strauss JF 3rd. Association of anti-mullerian hormone levels with obesity in late reproductive-age women. *Fertil Steril*. 2007 Jan;87(1):101-6. doi: 10.1016/j.fertnstert.2006.05.074.
34. Piouka A, Farmakiotis D, Katsikis I, Macut D, Gerou S, Panidis D. Anti-Mullerian hormone levels reflect severity of PCOS but are negatively influenced by obesity: relationship with increased luteinizing hormone levels. *Am J Physiol Endocrinol Metab*. 2009 Feb;296(2):E238-43. doi: 10.1152/ajpendo.90684.2008.
35. Panidis D, Katsikis I, Karkanaki A, Piouka A, Armeni AK, Georgopoulos NA. Serum anti-Müllerian hormone (AMH) levels are differentially modulated by both serum gonadotropins and not only by serum follicle stimulating hormone (FSH) levels. *Med Hypotheses*. 2011 Oct;77(4):649-53. doi: 10.1016/j.mehy.2011.07.005.
36. Jonard S, Dewailly D. The follicular excess in polycystic ovaries, due to intra-ovarian hyperandrogenism, may be the main culprit for the follicular arrest. *Hum Reprod Update*. 2004 Mar-Apr;10(2):107-17. doi: 10.1093/humupd/dmh010.
37. Durlinger AL, Visser JA, Themmen AP. Regulation of ovarian function: the role of anti-Müllerian hormone. *Reproduction*. 2002 Nov;124(5):601-9. doi: 10.1530/rep.0.1240601.
38. Durlinger AL, Gruijters MJ, Kramer P, Karels B, Kumar TR, Matzuk MM, Rose UM, de Jong FH, Uilenbroek JT, Grootegoed JA, Themmen AP. Anti-Müllerian hormone attenuates the effects of FSH on follicle development in the mouse ovary. *Endocrinology*. 2001 Nov;142(11):4891-9. doi: 10.1210/endo.142.11.8486.
39. Skalba P, Cygal A, Madej P, Dąbkowska-Huć A, Sikora J, Martirosian G, Romanik M, Olszanecka-Glinianowicz M. Is the plasma anti-Müllerian hormone (AMH) level associated with body weight and metabolic, and hormonal disturbances in women with and without polycystic ovary syndrome? *Eur J Obstet Gynecol Reprod Biol*. 2011 Oct;158(2):254-9. doi: 10.1016/j.ejogrb.2011.06.006.
40. Laven JS, Mulders AG, Visser JA, Themmen AP, De Jong FH, Fauser BC. Anti-Müllerian hormone serum concentrations in normoovulatory and anovulatory women of reproductive age. *J Clin Endocrinol Metab*. 2004 Jan;89(1):318-23. doi: 10.1210/jc.2003-030932.
41. Catteau-Jonard S, Pigny P, Reyss AC, Decanter C, Poncelet E, Dewailly D. Changes in serum anti-mullerian hormone level during low-dose recombinant follicular-stimulating hormone therapy for anovulation in polycystic ovary syndrome. *J Clin Endocrinol Metab*. 2007 Nov;92(11):4138-43. doi: 10.1210/jc.2007-0868.
42. Elmashad AI. Impact of laparoscopic ovarian drilling on anti-Müllerian hormone levels and ovarian stromal blood flow using three-dimensional power Doppler in women with anovulatory polycystic ovary syndrome. *Fertil Steril*. 2011 Jun;95(7):2342-6, 2346.e1. doi: 10.1016/j.fertnstert.2011.03.093.
43. Pigny P, Merlen E, Robert Y, Cortet-Rudelli C, Decanter C, Jonard S, Dewailly D. Elevated serum level of anti-mullerian hormone in patients with polycystic ovary syndrome: relationship to the ovarian follicle excess and to the follicular arrest. *J Clin Endocrinol Metab*. 2003 Dec;88(12):5957-62. doi: 10.1210/jc.2003-030727.