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# Polycystic Ovarian Syndrome in a population of obese adolescents

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## Background

• Polycystic Ovarian Syndrome (PCOS) is the most frequent endocrine disorder in women of reproductive age with a prevalence of  $8.5-12\%^{1}$ .

• Establishing clinically relevant diagnostic criteria for adolescents is challenging because of overlap with physiological puberty i.e acne, irregular menses.

## **Patients & Methods**

- Prospective observational study of 21 patients.
- Inclusion criteria:
  - $\checkmark$  from menarche to 20 years of age

• PCOS is closely linked to obesity and strongly associated with hypertension & metabolic problems (dyslipidemia & type 2 diabetes).

## Aims

• To determine the prevalence of PCOS in a population of obese adolescents using the Androgen Exess Society (AES) criteria<sup>3</sup> compared to those using NIH, Rotterdam, and Sultan diagnostic criteria for PCOS.

• To compare the clinical & biological features of PCOS vs. non-PCOS obese patients using AES criteria.

- ✓ <u>or</u> primary amenorrhea
- ✓ BMI > +2 SDS (WHO criteria)
- tumors, antidiabetic medication, oral contraceptives.

## Definitions

- PCOS defined by AES criteria<sup>3</sup>:
  - ✓ clinical and/or biochemical
    - hyperandrogenism
  - ✓ AND oligo-amenorrhea and/or

ultrasonographic criteria<sup>4</sup>:

 $\geq$  12 follicles (2-9 mm diameter)

and/or ovarian volume  $\geq$  10 mL.



#### PCOS on ultrasound (Balen et al, 2003)

## Results

Frequency of PCOS according to different diagnostic criteria

Patient characteristics (AES criteria)

Hormonal profile (AES criteria)

Serum AMH and inhibin B levels (AES criteria)



## **Frequency observed for** each AES criterion



	Non- PCOS N=12	PCOS N=9	P value
Age (years)	14.4 ±1.7	15.1 ±0.9	0.34
Weight (kg)	90.5 ±11.8	88.1 ±6.2	0.58
Height (cm)	161 ±6.8	163 ±5.6	0.49
BMI Z-Score (SDS)	3.1 ±0.6	2.7 ±0.2	0.12
Medical History			
Age of menarche (yrs)	11.4 ±1.4	12.2 ±1.0	0.17
FH of PCOS/hirsutism/ oligomenorrhea	17%	0	0.20
Macrosomia	20%	0	0.18
Small for gestational age	0	25%	0.09
Clinical characteristics			
Hypertension	17%	22%	0.75
Acanthosis nigricans	58%	67%	0.70
<ul> <li>Clinical hyperandrogenism</li> <li>Acne</li> <li>Hirsutism</li> </ul>	58 42 25	78 55 22	0.35 0.53 0.88
Irregular menses <ul> <li>Amenorrhea</li> <li>Oligomenorrhea</li> </ul>	17% 0 17%	78% 22% 56%	<b>0.05</b> 0.08 0.06
PCO morphology by pelvic ultrasound	17%	78%	0.05

	Non- PCOS N=12	PCOS N=9	P value
LH (mUI/L)	5.3 ±3.3	10.6 ±6.6	0.029
FSH (mUI/L)	4.2 ±2.3	5.2 ±0.93	0.27
LH/FSH ratio	1.8 ±1.3	2.0 ±1.14	0.63
AMH (pmol/l)	16.6 ±16.2	35.2 ±14.7	0.050
Inhibin B (pg/ml)	57.5 ±37	108.3 ±9.8	0.023
Total T LCMS (nM)	0.9 ±0.4	1.3 ±0.4	0.047
SHBG (nmol/l)	24 ±11	20 ±8.6	0.37
DHEAS (µmol/l)	4.9 ±3.3	4.2 ±1.6	0.53
$\Delta$ 4 Andro (nM)	4.3 ±1.8	9.9 ±5.2	0.002
17-OH P (nM)	1.7 ±1.3	2.0 ±1.4	0.57
Leptin (µg/l)	48 ±20	45 ±12	0.73
Cortisol (nM)	267 ±76	325 ±111	0.17



Fig. 3: AMH and Inhibine B levels comparing PCOS and non-PCOS patients. Dark lines are shown as median, light lines are mean.

## Metabolic profile (AES criteria)

- Trend: **1** insulin resistance
- Significantly higher total cholesterol in PCOS group.

	Non-PCOS N=12	PCOS N=9	P value
HOMA-IR	5.46 ±2.4	7.19 ±4.4	0.27
GlycemiaT120 min (mM)	6.53 ±1.2	6.84 ±1.2	0.55
Cholesterol T (mM)	3.62 ±0.5	4.13 ±0.6	0.05

#### Triglycerides (mM) $0.97 \pm 0.5$ $1.36 \pm 0.5$ 0.13 $2.46 \pm 0.6$ 0.06 LDL (mM) $2.02\pm\!\!0.4$

## **Discussion & Conclusions**

- $\checkmark$  Rates of PCOS varies widely according to the criteria employed (AES vs. Sultan)
- V PCOS (AES) was frequent in this small cohort of obese adolescents suggesting that PCOS is likely under-estimated.
- $\checkmark$  Our data suggest an increased metabolic risk in adolescents with PCOS irrespective of obesity.
- $\checkmark$  AMH appears to be a useful diagnostic tool for PCOS.
- ✓ Difficulty assessing PCO morphology via pelvic ultrasound raises the question of using pelvic MRI in obese adolescents.
- ✓ Questions remain whether or not widespread, systematic screening of PCOS in obese adolescents is warranted, is clinically useful, and at what time points.

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