Serum concentrations of afamin are elevated in patients with polycystic ovary syndrome

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Abstract
Oxidative stress seems to be present in patients with polycystic ovary syndrome (PCOS). The aim of this study was to evaluate the correlation between characteristics of PCOS and serum concentrations of afamin, a novel binding protein for the antioxidant vitamin E. A total of 85 patients with PCOS and 76 control subjects were investigated in a pilot cross-sectional study design between 2009 and 2013 in the University Hospital of Essen, Germany. Patients with PCOS were diagnosed according to the Rotterdam ESHRE/ASRM-sponsored PCOS Consensus Workshop Group. Afamin and diagnostic parameters of PCOS were determined at early follicular phase. Afamin concentrations were significantly higher in patients with PCOS than in controls (odds ratio (OR) for a 10 mg/ml increase in afamin Z 1.3, 95% CI Z 1.08–1.58). This difference vanished in a model adjusting for age, BMI, free testosterone index (FTI), and sex hormone-binding globulin (SHBG) (OR Z 1.05, 95% CI Z 0.80–1.38). In patients with PCOS, afamin correlated significantly with homeostatic model assessment-insulin resistance (HOMA-IR), fasting glucose, BMI, FTI, and SHBG (P < 0.001), but in a multivariate linear model, only HOMA-IR remained significantly associated with afamin (P = 0.001). No correlation was observed between afamin and androgens, LH, FSH, LH/FSH ratio, antral follicle count, ovarian volume, or anti-Müllerian hormone. In conclusion, elevated afamin values may indicate a state of oxidative stress and inflammation, strongly associated with IR and offering an indicator of impaired glucose tolerance in patients with PCOS irrespective of obesity.

Key Words
- vitamin E-binding protein
- afamin
- polycystic ovary syndrome
- oxidative stress
- insulin resistance

Introduction
The polycystic ovary syndrome (PCOS) is one of the most frequent endocrine disorders, found in up to 8% of women of reproductive age (1). According to the Rotterdam ESHRE/ASRM-sponsored PCOS Consensus Workshop Group, the diagnosis is confirmed, when two of the following three diagnostic criteria are present: i) hyperandrogenism, ii) polycystic ovaries with ≥12 sonographically measured small follicles with a diameter between 2 and 9 mm and/or an ovarian volume >10 ml, and iii) oligo- or anovulation (2).

New insights into the pathogenesis of PCOS identify oxidative stress as one possible cause, therefore offering
Afamin in patients with PCOS

Subjects and methods

Study population and study design

A total of 161 women, aged 18–46 years, were consecutively enrolled in the study. Of them, 85 patients suffered from PCOS, and the remaining 76 PCOS-free patients served as controls in a cross-sectional study design. All patients were treated at the University Hospital of Essen, Department of Obstetrics and Gynecology, between 2009 and 2013. The study was conducted according to the Declaration of Helsinki for Medical Research Involving Human Patients and approved by the local research ethics committee (No. 11-4643, No. 11-4688). Informed written consent was obtained from all patients.

PCOS was diagnosed according to the Rotterdam ESHRE/ASRM-sponsored PCOS Consensus Workshop Group 2003, i.e. if two of the following criteria were present: menstrual cycle disorders (oligomenorrhea, defined as cycles lasting longer than 3 days, or amenorrhea, defined as cycles lasting longer than 3 months), clinical or biochemical signs of hyperandrogenism (hirsutism) with a Ferriman–Gallwey score of more than seven (17) or obvious acne or alopecia (18) or an elevated total testosterone (normal range 0.5–2.6 nmol/l) and/or DHEAS (normal range 6–123 µg/dl) and/or androstenedione (normal range 0.3–3.3 ng/ml), and sonographically diagnosed polycystic ovaries (at least one ovary with at least 12 follicles with a diameter of 2–9 mm each or a volume > 10 ml).

Patients with endocrine disorders other than PCOS were excluded. None of the participants had taken hormonal contraceptives for at least 3 months before entering the study. Patients with tubal sterility, male causes of sterility, and patients with recurrent miscarriages but without any endocrine disorders or causes served as controls. Endocrine variables as well as afamin were determined between the second and fifth day of the menstrual cycle or after induction of artificial bleeding in cases of amenorrhea. Free testosterone index (FTI) was calculated using the formula: (total testosterone/SHBG) × 100. BMI was calculated as weight/(height)^2. In cases of elevated 17-hydroxyprogesterone, an ACTH test was performed. Cases with confirmed 21-hydroxylase deficiency were excluded from the study.

Transvaginal scan

A 4–9-MHz transducer (Voluson E8, General Electric Systems, Vipf, Austria; IU22, Philips Healthcare, Bothell, WA, USA; Sonoline Elegra, Siemens Ultrasound Division, Munich, Germany) was used for real-time ultrasound measurements. Ovarian volume was obtained by measuring the greatest diameter in every plane. The formula for a prolate ellipsoid \( V = \pi \times x \times y \times z \times 0.5236/1000 \) was used (19). For determination of the antral follicle count (AFC), small follicles with 2–9 mm diameter were calculated in

Further diagnostic and prognostic aspects (3, 4, 5, 6, 7). Oxidative stress seems to compromise follicle maturation. Surrogate parameters of oxidative stress such as ‘advanced glycosylation end products’ correlated strongly with anovulation, follicle number, and concentrations of anti-Müllerian hormone (AMH) in patients with PCOS (3).

Although insulin resistance (IR) plays an essential role in the pathogenesis of PCOS (8), it is not part of the diagnostic criteria. IR is found in 50% of women suffering from PCOS irrespective of obesity (9). In vitro, insulin enhances effects of luteinizing hormone (LH) in granulosa cells and leads to exaggerated androgen biosynthesis in patients with PCOS in a synergistic manner together with LH (10). An oral glucose tolerance test (OGTT) is recommended in obese patients with PCOS, but further research is required to detect lean patients with PCOS with impaired glucose metabolism (2).

In recent years, research has focused on AMH as a highly reliable diagnostic parameter for PCOS, reflecting the arrested follicle pool by acting as a follicle-stimulating hormone (FSH)-inhibiting parameter (11).

The previously described vitamin E-binding protein afamin may play a role in oxidative stress-related anti-apoptotic cellular processes (12). Vitamin E is an important anti-oxidant and protects from oxidative stress. Afamin was biochemically characterized as a vitamin E-binding protein (13) and was found in follicular fluid (14), indicating a role in the reproductive system, i.e. follicle maturation. Concentrations of afamin remain stable during the menstrual cycle (15); however, its precise role in human reproduction is virtually unknown. The afamin gene resides on chromosome 4 and was first described as the fourth member of the albumin gene family (16).

Overall, only few data exist to support an association between oxidative stress parameters and established diagnostic features of PCOS. Therefore, the purpose of our study was i) to compare serum afamin levels in patients with PCOS and healthy controls, ii) to evaluate the association between afamin and the known diagnostic criteria of PCOS including the AMH HOMA index and iii) to examine a potential diagnostic role of afamin in PCOS.
the longitudinal, transverse, and anterior–posterior cross-sections of each ovary using the most available magnification factor. Women with follicles of diameter >10 mm or any kind of ovarian mass were excluded. The ovary with the most follicles and greatest ovarian volume was used for analysis.

**Glucose tolerance test**

In patients with PCOS, a 3-h OGTT was used to evaluate the parameters of IR and b-cell function. After a 12-h overnight fast, patients ingested 75 g glucose and had their glucose and insulin concentrations determined at baseline and at 30, 60, 90, 120, and 180 min. For this study, we used only fasting insulin and glucose to determine the HOMA index as described previously (20).

**Biochemical analyses**

Blood sample of 27 ml was collected using an S-Monovette tube (Sarstedt AG & Co., Nyembrecht, Germany) from each woman, of which 18 ml were used for hormonal analysis and 9 ml were stored at 4 °C and processed within 4 h to avoid blood cell lysis for AMH and afamin analysis. Serum was obtained by low-speed centrifugation, immediately frozen, and kept at −80 °C until analyses.

Afamin was quantified as described previously (13, 14) by a custom-made double-antibody sandwich ELISA using an affinity-purified biotinylated polyclonal anti-afamin antibody for coating 96-well streptavidin-bound microtiter plates and peroxidase-conjugated MAb N13 for detection (MicroCoat Biotechnologie GmbH, Bernried, Germany). Secondary plasma in serial dilutions initially calibrated with a primary standard served as the assay standard. Afamin purified to homogeneity from human plasma was used as the primary standard; its exact protein concentration was determined by quantitative amino acid compositional analysis. Within-run and total coefficient of variation (CV) values were 3.3 and 6.2%, respectively, at a mean concentration of 73 mg/l (15).

Automated chemiluminescence immunoassay systems were used for the determination of LH, FSH, and testosterone (ADVIA Centaur, Siemens Healthcare Diagnostics, Eschborn, Germany), androstenedione and sex hormone-binding globulin (SHBG) (Immulite 2000 XPi, Siemens Healthcare Diagnostics), and insulin (Immuli- rate 2000 XPi, Siemens Healthcare Diagnostics). Glucose was determined photometrically (ADVIA Centaur, Siemens Healthcare Diagnostics). Intra- and inter-assay CV values for these parameters were <5 and <8% respectively.

Serum concentrations of AMH were determined by the enzymatically amplified two-site AMH-Gen-II ELISA (Beckman Coulter, Immunotech, Webster, TX, USA). Concentrations of <0.08 ng/ml were considered undetectable. Intra- and inter-assay CV values were <6%.

**Statistical methods**

Median values and interquartile range (IQR) were calculated for the investigated parameters, stratified for patient and control groups. The respective distributions were compared using the Wilcoxon test. For the evaluation of afamin, Spearman’s correlation coefficients were calculated with each of the parameters of interest in cases with PCOS and controls separately. In addition, a multivariate linear model was applied on afamin including all parameters that were significantly correlated with afamin at the univariate level and age. The association between afamin and PCOS case–control status was tested by means of a univariate as well as a multivariate logistic regression model, adjusted for age, BMI, FTI, and SHBG. In all regression models, skewed variables were log transformed. Statistical analyses were performed with Sigma Plot and R 3.0.

**Results**

The characteristics of patients are given in Table 1. Patients with PCOS were significantly younger than controls. As expected, all tested parameters were significantly different in patients with PCOS when compared with controls. We did not assess an OGTT in the control group and consequently could not compare the parameters of b-cell function between patients with PCOS and controls. In 64 patients with PCOS, homeostatic model assessment-IR (HOMA-IR) was determined: 27 out of 64 patients had a HOMA-IR >2, indicating IR (20, 21). Out of the 85 patients with PCOS, 56 presented with severe phenotype (fulfilling all the three Rotterdam criteria) and 29 with mild phenotype (presenting with two of the three Rotterdam criteria).

Afamin concentrations were significantly higher in patients with PCOS than in controls (P<0.001).

In patients with PCOS, a significant positive correlation was found between afamin and BMI, fasting glucose, HOMA-IR, and FTI and a significant negative correlation between afamin levels and SHBG (P<0.001, Table 2). No significant correlation was observed between afamin and patient age, LH, FSH, LH/FSH ratio, testosterone, androstenedione, AFC, ovarian volume, or AMH. Subgroup analysis of patients with severe phenotype did...
not alter the results (data not shown). In controls, afamin was significantly and positively correlated with the BMI ($P < 0.002$) and a significant inverse correlation with SHBG was also observed ($P = 0.035$, Table 2).

In a multivariate linear model of afamin in patients with PCOS including BMI, age, and log-transformed values of fasting glucose, HOMA-IR, FTI, and SHBG as explanatory variables, only HOMA-IR remained significantly associated with afamin ($P = 0.001$, Table 3).

A logistic regression model of afamin concerning case-control status showed a higher risk for an increase in afamin concentration (per 10 mg/l increase: odds ratio (OR) = 1.307, 95% CI = 1.082–1.579, $P = 0.006$) being a patient with PCOS, which disappeared, however, if adjusted for age, BMI, FTI, and SHBG (per 10 mg/l increase: OR = 1.050, 95% CI = 0.800–1.377, $P = 0.727$).

**Discussion**

To our knowledge, this is the first study evaluating afamin serum concentrations in patients with PCOS and correlating them with diagnostic parameters of PCOS.

The main findings of our study are as follows:

i) Afamin concentrations as well as conventional diagnostic PCOS parameters were significantly higher in patients with PCOS than in the control group.

### Table 1 Patient characteristics, showing median (IQR) and $P$ values of the Wilcoxon test.

| Parameters          | Patients with PCOS | Controls | $P$ value |
|---------------------|--------------------|----------|-----------|
| $n$                 | 85                 | 76       | <0.001    |
| Age (years)         | 26 (23–32)         | 35 (30–38)| <0.001    |
| BMI (kg/m$^2$)      | 27.9 (23.5–35.8)   | 24.1 (22.0–28.4)| <0.001    |
| Ovarian volume (ml) | 13.3 (10.6–16.8)   | 7.7 (6.6–10.0)| <0.001    |
| AFC (10–15)         | 12 (10–15)         | 7 (5–8)  | <0.001    |
| Total testosterone (nmol/l) | 2.04 (1.54–2.75)   | 1.32 (0.97–1.86)| <0.001    |
| Androstenedione (ng/ml) | 2.76 (2.01–3.96)   | 1.63 (1.17–2.03)| <0.001    |
| FTI (%)             | 5.80 (3.18–9.17)   | 2.34 (1.54–4.32)| <0.001    |
| SHBG (nmol/l)       | 41.6 (24.8–62.1)   | 52.7 (36.1–71.1)| <0.001    |
| LH (IU/l)           | 7.4 (5.0–11.0)     | 4.6 (3.6–6.3) | <0.001    |
| FSH (IU/l)          | 5.6 (4.2–6.9)      | 7.1 (6.3–8.0) | <0.001    |
| LH/FSH ratio        | 1.38 (0.95–2.12)   | 0.64 (0.50–0.80)| <0.001    |
| Fasting glucose (mg/dl) | 87.0 (83.0–96.5)   | 2.26 (1.05–3.00)| <0.001    |
| HOMA-IR             | 1.32 (0.57–3.56)   | –        | –         |
| AMH (ng/ml)         | 4.39 (2.86–7.91)   | 74.9 (67.3–89.0)| <0.001    |
| Afamin (mg/l)       | 83.1 (72.1–101.2)  | 74.9 (67.3–89.0)| <0.001    |

AFC, anter follicle count (number of follicles with a diameter between 2 and 9 mm; FTI, free testosterone index; SHBG, sex hormone-binding globulin; LH, luteinizing hormone; FSH, follicle-stimulating hormone; HOMA-IR, homeostasis model assessment-insulin resistance; AMH, anti-Müllerian hormone.

### Table 2 Correlation between afamin and patients’ characteristics using Spearman’s rank correlation test, sorted by the absolute value of the correlation coefficient for cases with PCOS.

| Parameters          | Patients with PCOS | Controls | $P$ value |
|---------------------|--------------------|----------|-----------|
| $n$                 | 85                 | 76       | <0.001    |
| HOMA-IR             | 0.65               | 0.34     | 0.002     |
| BMI                 | 0.54               | 0.06     | 0.693     |
| Fasting glucose     | 0.47               | 0.00     | 0.001     |
| FTI                 | 0.45               | 0.06     | 0.001     |
| SHBG                | −0.44              | −0.29    | 0.035     |
| Ovarian volume      | −0.18              | −0.01    | 0.941     |
| Androstenedione     | 0.18               | 0.00     | 0.969     |
| Testosterone        | 0.17               | 0.10     | 0.387     |
| LH/FSH ratio        | −0.09              | −0.10    | 0.763     |
| FSH                 | 0.09               | −0.13    | 0.267     |
| AFC                 | −0.08              | −0.13    | 0.281     |
| AMH                 | −0.07              | −0.18    | 0.127     |
| LH                  | −0.03              | −0.12    | 0.286     |
| Age                 | 0.03               | −0.06    | 0.589     |
Until now, OGTT was recommended only in obese patients with PCOS with a BMI ≥27 in light of the high lifetime risk for developing diabetes mellitus type 2 (2). As an OGTT with the measurement of insulin is expensive, time consuming, and restricted to experienced personnel, screening tools for lean women are strongly needed. In this context, afamin determination could detect those patients who would benefit from an OGTT. While designing the study, we did not focus mainly on biomarker search that could replace an OGTT in patients with PCOS. Therefore, our data can only be interpreted as a preliminary result possibly indicating an important biomarker of disturbed β-cell function in several, partly unknown conditions as well as in patients with PCOS.

Further research in larger cohorts is recommended to explore afamin cutoff values indicating a pathological diagnosis.

Afamin was biochemically characterized as a vitamin E-binding protein (13). Vitamin E belongs to the group of non-enzymatic antioxidants and it is plausible that its binding protein afamin indicates oxidative stress with vitamin E presence or requirement. Furthermore, there is some evidence to support an association among oxidative stress, hyperinsulinemia, and PCOS (4, 22).

Oxidative stress, defined as an imbalance between pro- and antioxidants (23), is known to be present in patients with PCOS with follicle maturation failure (3, 24). Kuroglu et al. (7) demonstrated this imbalance independent of obesity and hyperinsulinemia in patients with PCOS. However, there is some evidence of an association among pathways of hyperinsulinemia, oxidative stress, and inflammation in PCOS (4, 22, 25). The authors observed a hyperglycemia-induced increase in reactive oxygen species (ROS) and activated NF-κB in patients with PCOS (4, 22), thus leading to transcription of tumor necrosis factor α (TNFα), most pronounced in obese patients with PCOS (22). Additionally, TNFα is a key marker of chronic inflammation directly inducing IR (26, 27) and strongly associated with PCOS (28). The investigation of oxidative stress status including features of metabolic syndrome in patients with PCOS and BMI-matched controls is thus highly recommended for future research.

A limitation in our study is the significant age difference between patients and controls. We have therefore adjusted our results for age although there was no significant correlation between afamin and age in our study population. This is in agreement with a previous study reporting no age dependency of afamin concentrations in a large group of healthy blood donors (15).

### Table 3 Results of a multivariate linear model of selected variables performed for afamin serum concentrations.

| Covariate            | \( \beta \) estimate | 95% CI          | \( P \) value |
|----------------------|-----------------------|-----------------|--------------|
| HOMA-IR              | 2.86                  | 0.63 to 5.09    | 0.001        |
| BMI                  | 0.49                  | −0.25 to 1.23   | 0.622        |
| Fasting glucose      | 0.04                  | −0.23 to 0.30   | 0.971        |
| FTI                  | 0.16                  | −0.58 to 0.89   | 0.350        |
| SHBG                 | −0.11                 | −0.28 to 0.06   | 0.679        |
| Age                  | −0.25                 | −0.97 to 0.48   | 0.349        |

\( ^a \)Skewed variables were log transformed (HOMA-IR, fasting glucose, FTI, and SHBG) to hold model assumptions.

\( ^b \)Skewed variables were log transformed (HOMA-IR, fasting glucose, FTI, and SHBG) to ensure interpretability of estimates.
Our control group had median afamin values of 74.9 mg/l (IQR 67.3–89.0), comparable to previously published data showing a median value of 70.7 mg/l in a healthy cohort of 177 participants without gynecological diseases (29).

Declaration of interest
H Dieplinger is the owner and a shareholder of Vitateq Biotechnology GmbH, Innsbruck, Austria, a spin-off biotech company from Innsbruck Medical University, holding several patents related to research described in this article. All other coauthors report no conflict of interest.

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Author contribution statement
A Königger, S Kasimir-Bauer, and H Dieplinger made substantial contributions to the study’s conception and design. A Königger, P Edimiris, L Koch, A Enekev, R Kimmig, and H Dieplinger made substantial contributions to acquisition and interpretation of data. A Königger, S Kasimir-Bauer, and H Dieplinger made substantial contributions to drafting the article and the final approval of the version to be published. P Edimiris, L Koch, A Enekev, C Lamina, and R Kimmig made substantial contributions to revising the article critically for important intellectual content and the final approval of the version to be published. C Lamina gave substantial contributions to analysis and interpretation of data.

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References
1 Aziz R. PCOS: a diagnostic challenge. Reproductive Biomedicine Online 2004 8 644–648. (doi:10.1684/15147-6438(10)6144-6)
2 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). Human Reproduction 2004 19 41–47. (doi:10.1093/humrep/deh098)
3 Diamanti-Kandarakis E, Plouka A, Livadas S, Piperi C, Katsikis I, Papavassiliou AG & Panidis D. Anti-Müllerian hormone is associated with advanced glycosylated end products in lean women with polycystic ovary syndrome. European Journal of Endocrinology 2009 160 847–853. (doi:10.1530/EJE-08-0510)
4 Gonzalez F, Rote NS, Minium J & Kirwan JP. Reactive oxygen species-induced oxidative stress in the development of insulin resistance and hyperandrogenism in polycystic ovary syndrome. Journal of Clinical Endocrinology and Metabolism 2006 91 336–340. (doi:10.1210/jc.2005-1696)
5 Victor VM, Rocha M, Banuls C, Alvarez A, de Pablo C, Sanchez-Serrano M, Gomez M & Hernandez-Mijares A. Induction of oxidative stress and human leukocyte/endothelial cell interactions in polycystic ovary syndrome patients with insulin resistance. Journal of Clinical Endocrinology and Metabolism 2011 96 3115–3122. (doi:10.1210/jc.2011-06651)
6 Murri M, Luque-Ramirez M, Insenser M, Ojeda-Ojeda M & Escobar-Morreale HF. Circulating markers of oxidative stress and polycystic ovary syndrome (PCOS): a systematic review and meta-analysis. Human Reproduction Update 2013 19 268–288. (doi:10.1093/humupd/dms059)
7 Kurodoga Z, Oasis H, Tuluce Y & Koyuncu I. Oxidative status and its relation with insulin resistance in young non-obese women with polycystic ovary syndrome. Journal of Endocrinological Investigation 2012 35 317–321. (doi:10.3275/7682)
8 Baillargeon JP & Nestler JE. Commentary: polycystic ovary syndrome: a syndrome of ovarian hypersensitivity to insulin? Journal of Clinical Endocrinology and Metabolism 2006 91 22–24. (doi:10.1210/jc.2005-1804)
9 Dunài F, Segal KR, Futterweit C & Franks S. Modulation by insulin of follicle-stimulating hormone and luteinizing hormone actions in human granulosa cells of normal and polycystic ovaries. Journal of Clinical Endocrinology and Metabolism 1996 81 302–309. (doi:10.1210/ jc.81.3.550768)
10 Jonard S & Dewailly D. The follicular excess in polycystic ovaries, due to intra-ovarian hyperandrogenism, may be the main culprit for the follicular arrest. Human Reproduction Update 2004 10 107–117. (doi:10.1093/humupd/dmh010)
11 Heiser M, Hutter-Paier B, Jerkovic L, Pfagnier R, Windisch M, Becker-Andre M & Dieplinger H. Vitamin E binding protein afamin protects neuronal cells in vitro. Journal of Neural Transmission. Suplementum 2002 62 337–345.
12 Voegele AF, Jerkovic L, Wellenzohn R, Eller P, Kronenberg F, Liedl KR & Dieplinger H. Characterization of the vitamin E-binding properties of human plasma afamin. Biochemistry 2002 41 14532–14538. (doi:10.1021/bi026513v)
13 Jerkovic L, Voegele AF, Chwatal S, Kronenberg F, Radcliffe CM, Wormald MR, Lobentanz EM, Ezeh B, Eller P, Dejori N et al. Afamin is a novel human vitamin E-binding glycoprotein characterization and in vitro expression. Journal of Proteome Research 2005 4 889–899. (doi:10.1021/pr0400105)
14 Dieplinger B, Egger M, Gabriel C, Poelz W, Morandell E, Seeber B, Kronenberg F, Haltmayer M, Mueller T & Dieplinger H. Analytical characterization and clinical evaluation of an enzyme-linked immunosorbent assay for measurement of afamin in human plasma. Clinica Chimica Acta 2013 425 236–241. (doi:10.1016/j.cca.2013.08.016)
15 Lichenstein HS, Lyons DE, Wurfel MM, Johnson DA, McGinley MD, Leidl JC, Trollinger DR, Mayer JP, Wright SD & Zukowski MM. Afamin is a new member of the albumin, α-fetoprotein, and vitamin D-binding protein gene family. Journal of Biological Chemistry 1994 269 18149–18154.
16 Ferriman D & Gallwey JD. Clinical assessment of body hair growth in women. Journal of Clinical Endocrinology and Metabolism 1961 21 1440–1447. (doi:10.1210/jcem-21-11-1440)
17 Ludwig E. Classification of the types of androgenetic alopecia (common baldness) occurring in the female sex. British Journal of Dermatology 1977 97 247–254. (doi:10.1111/j.1365-2133.1977.tb15179.x)
18 Balen AH, Laven JS, Tan SL & Devalia D. Ultrasound assessment of the polycystic ovary: international consensus definitions. Human Reproduction Update 2003 9 505–514. (doi:10.1093/humupd/dmg044)

http://www.endocrinologypages.org
DOI: 10.1530/EC-14-0053
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20 Matthews DR, Hosker JP, Rudenski AS, Naylor BA, Treacher DF & Turner RC. Homeostasis model assessment: insulin resistance and β-cell function from fasting plasma glucose and insulin concentrations in man. Diabetologia 1985 28 412–419. (doi:10.1007/BF00280883)

21 Jensterle M, Weber M, Pfeifer M, Prezelj J, Pfutzner A & Janez A. Assessment of insulin resistance in young women with polycystic ovary syndrome. International Journal of Gynaecology and Obstetrics 2008 102 137–140. (doi:10.1016/j.ijigo.2008.03.017)

22 Gonzalez F, Rote NS, Minium J & Kirwan JP. Increased activation of nuclear factor κB triggers inflammation and insulin resistance in polycystic ovary syndrome. Journal of Clinical Endocrinology and Metabolism 2006 91 1508–1512. (doi:10.1210/jc.2005-2327)

23 Agarwal A, Aponte-Mellado A, Premkumar BJ, Shaman A & Gupta S. The effects of oxidative stress on female reproduction: a review. Reproductive Biology and Endocrinology 2012 10 49. (doi:10.1186/1477-7827-10-49)

24 Blair SA, Kyaw-Tun T, Young IS, Phelan NA, Gibney J & McEneny J. Oxidative stress and inflammation in lean and obese subjects with polycystic ovary syndrome. Journal of Reproductive Medicine 2013 58 107–114.

25 Gonzalez F, Minium J, Rote NS & Kirwan JP. Hyperglycemia alters tumor necrosis factor-α release from mononuclear cells in women with polycystic ovary syndrome. Journal of Clinical Endocrinology and Metabolism 2005 90 5336–5342. (doi:10.1210/jc.2005-0694)

26 Hotamisligil GS, Murray DL, Choy LN & Spiegelman BM. Tumor necrosis factor α inhibits signaling from the insulin receptor. PNAS 1994 91 4854–4858.

27 Hotamisligil GS, Shargill NS & Spiegelman BM. Adipose expression of tumor necrosis factor-α: direct role in obesity-linked insulin resistance. Science 1993 259 87–91. (doi:10.1126/science.7678183)

28 Escobar-Morreale HF, Luque-Ramirez M & Gonzalez F. Circulating inflammatory markers in polycystic ovary syndrome: a systematic review and metaanalysis. Fertility and Sterility 2011 95 1048–1058.e1041–1042. (doi:10.1016/j.fertnstert.2010.11.036)

29 Dieplinger H, Ankerst DP, Burges A, Lenhard M, Lingenhel A, Fieder L, Buchner H & Stieber P. Afamin and apolipoprotein A-IV: novel protein markers for ovarian cancer. Cancer Epidemiology, Biomarkers & Prevention 2009 18 1127–1133. (doi:10.1158/1055-9965.EPI-08-0653)

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