Greek hyperinsulinemic women, with or without polycystic ovary syndrome, display altered inositols metabolism

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BACKGROUND: We have shown that American women with polycystic ovary syndrome (PCOS) have decreased glucose-stimulated release of a putative mediator of insulin action, D-chiro-inositol (DCI)-containing inositolphosphoglycan (DCI-IPG), and increased urinary clearance of DCI (uClDCI), which was associated with hyperinsulinemia. METHODS: DCI levels and the release of insulin and DCI-IPG during an oral glucose tolerance test (AUCs) were assessed in 27 Greek PCOS and 10 normal Greek women. RESULTS: PCOS women were heavier than controls (BMI = 28.4 versus 23.7 kg/m², P = 0.05) with higher waist-to-hip ratios (WHR = 0.78 versus 0.71, P = 0.009) and increased free testosterone (P = 0.048) and AUCinsulin (P = 0.04). In PCOS women, incremental AUCDCI-IPG was significantly decreased by 59% (2158 versus 5276 min, P = 0.01), even after correction for BMI and WHR. Finally, increased uClDCI (r = 0.35, P = 0.04) and decreased AUCDCI-IPG; (r = 0.46, P = 0.004) were significantly associated with hyperinsulinemia in all women together, even after correction for BMI and WHR (Ps = 0.02 and 0.007), and regardless of PCOS status. CONCLUSIONS: Greek women, with or without PCOS, display increased uClDCI and decreased AUCDCI-IPG in association with higher insulin levels but independent of adiposity. Increased clearance of inositols might reduce tissue availability of DCI and decrease the release of DCI-IPG mediator, which could contribute to insulin resistance and compensatory hyperinsulinemia in Greek women, as previously described in American women.

Keywords: polycystic ovary syndrome; inositols; inositolphosphoglycans; insulin resistance; hyperinsulinemia

Introduction

The polycystic ovary syndrome (PCOS) is a common but still poorly understood disorder. It is defined by hyperandrogenism, chronic anovulation and/or polycystic ovaries (Zawadzki and Dunaif, 1992; The Rotterdam ESHRE/ASRM-sponsored PCOS consensus workshop group, 2004), and affects 6–10% of women of childbearing age (Knochenhauer et al., 1998; Diamanti-Kandarakis et al., 1999; Asuncion et al., 2000). PCOS is the most common endocrinopathy of women in this age group and is associated with an increased risk of developing hypertension, dyslipidemia, impaired glucose tolerance or Type 2 diabetes (Baillargeon et al., 2003; Cattrall and Healy, 2004), and probably cardiovascular disease (Cibula et al., 2000; Solomon et al., 2002; Cattrall and Healy, 2004).

Increasing evidence supports the central role of insulin resistance and/or increased insulin action in the syndrome’s pathogenesis (Nestler, 1997; De Leo et al., 2003; Baillargeon 2005; Baillargeon and Nestler, 2006; Baillargeon and Carpenter, 2007). Obese and lean women with PCOS manifest insulin resistance independent of fat mass (Dunaif et al., 1989), and administration of insulin-sensitizing drugs, such as metformin (Diamanti-Kandarakis et al., 1998; Nestler et al., 1998; Baillargeon et al., 2004b; Nestler, 2008), thiazolidinediones (Baillargeon et al., 2002, 2004b; Pesant and Baillargeon, 2006) and D-chiro-inositol (DCI) (Nestler et al., 1999; Iuorno et al., 2002; Gerli et al., 2003), to both obese and lean women with the syndrome increases the frequency of ovulation and decreases circulating androgens.

Some actions of insulin may be effected by putative inositol-phosphoglycan (IPG) mediators of insulin action (Saltiel, 1990; Romero and Larner, 1993), and evidence suggests that a deficiency in a specific DCI-containing IPG (DCI-IPG) may...
contribute to insulin resistance in individuals with impaired glucose tolerance or Type 2 diabetes (Kennington et al., 1990; Asplin et al., 1993; Shashkin et al., 1997; Baillargeon et al., 2004a). Furthermore, we have shown that metformin may improve the action of insulin in obese women with PCOS in part by improving insulin-mediated release of the DCI-IPG mediator (Baillargeon et al., 2004a).

Our group has also recently demonstrated that high urinary clearance of DCI (uClDCI) is associated with insulin resistance and high insulin levels in American women, with or without PCOS, and that American PCOS women display reduced insulin-stimulated release of DCI-IPG (Baillargeon et al., 2006). These results are consistent with a defect in tissue availability of DCI in American PCOS that may contribute to the insulin resistance of the syndrome. Moreover, our group (Nestler et al., 1999; Iuorno et al., 2002) and others (Gerli et al., 2003) have shown that oral administration of DCI to women with PCOS improves glucose tolerance while reducing circulating insulin in both obese (Nestler et al., 1999) and lean (Iuorno et al., 2002) women with PCOS, and also decreases serum androgens and improves ovulatory function.

On the basis of these findings, we hypothesized that elevated uClDCI is associated with insulin resistance not only in American women, with or without PCOS, but also in a population with a different ethnic and geographic origin, such as Greek women. To assess this hypothesis, we studied women with PCOS and normal women from the region of Athens, in Greece, and assessed circulating DCI and 24 h uClDCI, PCOS and normal women from the region of Athens, in Greece, and assessed circulating DCI and 24 h uClDCI, release of insulin and DCI-IPG during an oral glucose tolerance test (OGTT), and insulin sensitivity by the minimal model technique.

Research design and methods

Subjects
Twenty-seven women with PCOS and 10 normal controls were evaluated at the Endocrine unit, first department of Medicine, Laiko Hospital of the University of Athens Medical School, Athens, Greece. PCOS was defined by oligo-amenorrhea (≤8 menstrual periods in the previous year) and hyperandrogenism (hirsutism, acne or elevated serum total or free testosterone concentration); while hyperprolactinemia, thyroid dysfunction and late-onset adrenal hyperplasia were excluded by the appropriate tests. Ovarian ultrasounds were not performed such that our more stringent criteria met both 1990 NIH and 2003 Rotterdam conferences diagnostic criteria (Zawadzki and Dunaif, 1992; The Rotterdam ESHRE/ASRM-sponsored PCOS consensus workshop group, 2004). However, based on these criteria, a woman could have been classified as having PCOS even if her testosterone levels were normal, providing that she had hyperandrogenic signs.

Normal women were recruited from the endocrine outpatient clinic of first department of Medical Laiko Hospital. They were regularly followed-up for thyroid problems, such as simple goiters or thyroid nodules and were euthyroid without medication. They had regular menstrual cycles, absence of acne and hirsutism, normal androgen levels and glucose tolerance, and did not have any history of gestational diabetes or family history of a first-degree relative with diabetes. They were free of diabetes, hypertension, dyslipidemia, or kidney or heart diseases. All women were 18–40 years old and none took oral contraceptives or any medication known to affect insulin sensitivity for at least 2 months prior to study. The study has been approved by the institutional review board of Laiko Hospital and Virginia Commonwealth University, and each woman gave written informed consent.

Study protocol
Because DCI may be ingested as part of a diet high in legumes or fruits, the women were given instructions for a balanced mixed diet to be followed for at least 3 days prior to the start of the study. PCOS women were studied during the equivalent of the follicular phase of the menstrual cycle, as confirmed by a serum progesterone ≤0.5 nmol/l (2 ng/ml). Normal women were studied during the mid-follicular phase of the menstrual cycle (Days 5–9), which most closely approximates the hormonal milieu of anovulatory women with PCOS.

On the first day, fasting baseline laboratories and a 2-h OGTT with 75 g of dextrose were performed. During the OGTT, blood samples were collected every 15 min for determination of serum glucose and insulin concentrations, and serum DCI-IPG bioactivity was determined every 30 min. From the OGTT, the insulinogenic index at 15 min was calculated ([I(0–15) = insulin(0–15) – insulin(0)]/glucose(0–15) – glucose(0)) (Hanson et al., 2000). Women were then instructed to collect all their urine for 24 h before the next visit.

On the second day, after a 12-h overnight fast, insulin sensitivity was measured by the frequent sampling intravenous glucose tolerance test (FSIVGTT) technique as described by Bergman (Bergman, 1989). At zero time, 300 mg/kg of dextrose was administered intravenously and 0.03 U/kg insulin was administered intravenously 20 min later. A total of 27 blood samples for determination of insulin and glucose were collected over the 3-h duration of the protocol. Data were analysed with the Minimal Model Identification Software (MINMOD®, version 6.02, 2004) (Pacini and Bergman, 1986), which yields quantitative determination of glucose-mediated disappearance of glucose (Sg, or glucose effectiveness), tissue insulin sensitivity (S), acute insulin response to glucose (AIRg) and disposition index (DI = S × AIRg).

Laboratory assays
Blood samples were centrifuged immediately, and sera were stored at −70 °C until they were shipped all at once in dry ice to Richmond, and then stored again at −70 °C until assayed. All hormones were assayed as previously described (Nestler et al., 1989, 1991, 1994). Blood glucose and insulin levels were determined by the core laboratory of the General Clinical Research Center of the Virginia Commonwealth University Health System. All other analytes were assayed in Dr Nestler’s laboratory at Virginia Commonwealth University. Serum free testosterone was calculated by the method of Sodergard et al. (1982) using a serum albumin concentration of 40 g/l. To avoid inter-assay variation, all samples were analysed in duplicate in a single assay for each hormone. The intra-assay coefficient of variation for the insulin was 5.5%, and was less than 10% for all steroid hormone assays.

DCI and MYO-inositol analyses
Plasma and urinary inositol concentrations were determined by gas chromatography and mass spectrometry. [1H6]Racemic chiro-inositol and [1H6]myo-inositol were added to plasma or urine as internal standards. The samples were then purified, derivatized with pentafluoropropionic anhydride, separated on a Chirasil-Val capillary column (Alltech, State College, PA, USA), and analysed in negative ion chemical ionization mode on an HP 5973 mass spectrometer (Agilent Technologies, Palo Alto, CA, USA) with methane as the
reagent gas, as previously reported (Ostlund et al., 1993). Twenty-four hour urinary excretion was calculated by dividing 24 h urinary excretion by plasma concentration.

**DCI containing inositolphosphoglycans (DCI-IPG) insulin mediator bioactivity assay**

DCI-IPG mediator was isolated from serum as previously described (Baillargeon et al., 2004a). To date, it has not been possible to measure the content of extracted DCI-IPG because its structure and exact mass are unknown, and no specific antibody suitable for an immunosassay has been developed. Therefore, DCI-IPG mediator bioactivity was determined using the specific activation of pyruvate dehydrogenase (PDH) phosphatase, as previously validated in women with PCOS and described in detail (Baillargeon et al., 2004a). The inter-assay coefficient of variation of the bioassay was 17.4%. The PDH activity intra-assay coefficient of variation was 6.7%. Coefficients of variation of the entire method (extraction and assay) were 10.7 and 8.5%, respectively, for the absolute values of basal and peak DCI-IPG bioactivity.

In order to adjust for variation in basal PDH activity from one assay to the other, and therefore from subject to subject, the water-blank activity was subtracted from the bioactivity of DCI-IPG released into serum during OGTT, which was then expressed as the percentage of its bioactivity at baseline (0 min).

**Statistical analyses**

We analysed the response of serum insulin concentrations and relative bioactivities of DCI-IPG to the oral administration of glucose by calculating the areas under the respective response curves (AUC) by the trapezoidal rule. Results not normally distributed, based on the Normal Quintile Plot, were log-transformed for all statistical analyses and calculated using the specific activation of pyruvate dehydrogenase (PDH) phosphatase, as previously validated in women with PCOS and described in detail (Baillargeon et al., 2004a). The inter-assay coefficient of variation of the bioassay was 17.4%. The PDH activity intra-assay coefficient of variation was 6.7%. Coefficients of variation of the entire method (extraction and assay) were 10.7 and 8.5%, respectively, for the absolute values of basal and peak DCI-IPG bioactivity.

In order to adjust for variation in basal PDH activity from one assay to the other, and therefore from subject to subject, the water-blank activity was subtracted from the bioactivity of DCI-IPG released into serum during OGTT, which was then expressed as the percentage of its bioactivity at baseline (0 min).

The primary variables of interest for this study were 24 h uCl_{DCI} and insulin-stimulated release of the DCI-IPG insulin mediator (AUC_{DCI-IPG}). Variable comparisons between groups were made with Student’s two-tailed t-test, and equalities of variances were tested with the Brown–Forsythe test. For comparisons with unequal variances, Ps of Welch ANOVA tests were reported, as indicated. Correlation analysis was performed using Pearson’s correlation test. With 27 PCOS and 10 control women, this study has 70% power to detect the same group difference in uCl_{DCI} than the difference we previously observed in American PCOS women (1.75 on loge scale) (Baillargeon et al., 2006), considering the same common standard deviation (1.85 on loge scale).

Since BMI and waist-to-hip ratio (WHR) differed between groups (see Table I), all comparisons and correlations were corrected for both variables using multiple linear regression analyses. Possible interactions between BMI and PCOS status for DCI or DCI-IPG parameters were also tested because a significant interaction with uCl_{DCI} was found in American women (Baillargeon et al., 2006). However, no interaction was found in the present study.

In order to determine the best models to predict our primary variables of interest, manual stepwise multiple linear regression analyses were performed. All variables reported in Table II that were associated with uCl_{DCI} or AUC_{DCI-IPG} (P ≤ 0.10) were entered successively in the model based on the next best P-value (forward method); except for 24 h urinary excretions, which were collinear with clearances, AUC_{DCI-IPG}–AUC_{insulin} ratio, which was collinear with AUC_{insulin, as well as DI and IGI_{15}, which were redundant with AIR_{g}. All possible interactions with variables previously kept in the model were also included at each step. Thereafter, parameters that did not contribute significantly to the model were excluded (partial P > 0.01 for interactions). We also performed a stepwise analysis to determine the best independent predictors of AUC_{insulin} using the same method.

**Results**

**Clinical and biochemical characteristics**

Women with PCOS did not differ significantly from normal control women with respect to age, systolic blood pressure and diastolic blood pressure (Table I). However, their BMI tended to be higher \( (P = 0.05) \) and their WHR was significantly higher \( (P = 0.009) \).

Calculated free testosterone levels were significantly increased by more than 75% in women with PCOS as compared to control women \( (P = 0.048) \). This difference was no longer significant after correction for BMI and WHR. Although fasting insulin levels were 50% higher in PCOS women compared with control women, the difference was not significant \( (P = 0.21) \), whereas areas under the insulin curves during OGTT (AUC_{insulin}) were 65% higher \( (P = 0.04) \) in PCOS women compared with normal controls. AUC_{insulin} was no longer significantly different between groups after correction for BMI and WHR. IGI_{15} was not calculable (negative result) in two PCOS women and was not significantly different between groups. Regarding the fitting of FSIVGTT data, the MINMOD program did not generate S_{g} result for one PCOS woman and AIR_{g} for three PCOS women. Accordingly, DI \( (S_{g} \times AIR_{g}) \) was unavailable for four PCOS women, which reduced the power of this analysis. Although \( S_{g} \) was reduced by 44% in PCOS women, this difference was only borderline significant \( (P = 0.07) \). \( S_{g} \), AIR_{g} and DI were not significantly different between groups, although there was a trend for lower mean DI in PCOS women (decreased by 30%, \( P = 0.43) \).

**DCI and MYO-inositol (MYO) metabolism**

Plasma concentrations of DCI and 24 h urinary excretion and clearance of DCI were comparable between PCOS women and normal controls (Table I). Corresponding parameters were also similar between groups for MYO. However, AUC_{DCI-IPG} was significantly reduced in PCOS women as compared with controls \( (P = 0.01) \), even after correction for BMI and WHR \( (P = 0.01) \). Incremental AUC_{DCI-IPG}, i.e. AUC_{DCI-IPG} above baseline, was decreased by almost 60% in PCOS women compared with controls. Finally, the ratio of AUC_{DCI-IPG} to AUC_{insulin} was reduced by half in women with PCOS \( (P = 0.02) \), which remained significant after correction for BMI and WHR \( (P = 0.047) \).
Correlation between hyperinsulinemia (AUC_{insulin}) and uCl_{DCI} or insulin-stimulated release of DCI-IPG (AUC_{DCI-IPG})

When all women were included, there was a direct correlation between AUC_{insulin} and AUC_{DCI-IPG} that was statistically significant (Fig. 1A; r = 0.35, P = 0.04). The negative correlation between AUC_{insulin} and AUC_{DCI-IPG} was even more robust and significant (Fig. 1B; r = 0.46, P = 0.004). These associations persisted after adjustment for differences both in BMI and WHR (P = 0.02 and 0.007, respectively). Moreover, no interaction was found between PCOS status and AUC_{insulin} (P = 0.12 for uCl_{DCI} and P = 0.34 for AUC_{DCI-IPG}), such that these correlations were not driven by the effects in only one of the groups. When assessing these correlations within each group, AUC_{insulin} was positively associated with uCl_{DCI} both in PCOS (r = 0.40, P = 0.04) and normal women (r = 0.64, P = 0.04). There were also negative correlations between AUC_{insulin} and AUC_{DCI-IPG} within each group (r = 0.33 for PCOS and r = 0.57 for controls), but they were not significant in these sub-group analyses (P = 0.08 and 0.09, respectively).

After combining the data from the present population with the previously published data (Baillargeon et al., 2006) from Richmond, Virginia, a total of 86 women were analysed, i.e. 50 women with PCOS (27 from Greece and 23 from USA) and 36 normal controls (10 from Greece and 26 from USA). Using this database and analysing all the women together (n = 86), a significant positive correlation persisted between AUC_{insulin} and uCl_{DCI} (Fig. 1C: r = 0.32, P = 0.003) and a negative correlation persisted between AUC_{insulin} and AUC_{DCI-IPG} (Fig. 1D: r = −0.31, P = 0.004). The association between hyperinsulinemia and increased uCl_{DCI} remained significant after adjustment for BMI and WHR (P = 0.002), but this was not the case for AUC_{DCI-IPG} (P = 0.06). There was no interaction between AUC_{insulin} and PCOS status for either analysis, as well as between AUC_{insulin} and Greek or American origin. Therefore, these significant correlations are valid regardless of women’s origin and PCOS status.

Stepwise multivariate analyses

As described in the Statistical analyses section, the best models predicting our primary variable of interest, i.e. uCl_{DCI} and AUC_{DCI-IPG}, were determined using manual stepwise multivariate analyses. Variables associated with uCl_{DCI} with P ≤ 0.10 were pDCI (P = 0.004), uCl_{MYO} (P = 0.009), AUC_{insulin} (P = 0.04), pMYO (P = 0.09) and AUC_{DCI-IPG} (P = 0.10). Using these independent variables, the stepwise analysis (partial P = 0.003) and MYO urinary clearance (partial P = 0.006), which explained 37.6% of the variance of uCl_{DCI} (R^2 = 0.376, P < 0.001). Variables associated with AUC_{DCI-IPG} with P ≤ 0.10 were AUC_{insulin} (P = 0.004), PCOS status (P = 0.01), S_g (P = 0.07), pDCI (P = 0.08) and uCl_{DCI} (P = 0.10). After stepwise analysis, the only independent variable significantly associated with AUC_{DCI-IPG} was AUC_{insulin}, which explained 21.6% of the variability of AUC_{DCI-IPG} (R^2 = 0.216, P = 0.004).

Table 1. Clinical and biochemical characteristics of Greek women with the PCOS and normal control women.

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>PCOS (n = 27)</th>
<th>Normals (n = 10)</th>
<th>P (t-test)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>26 (24–29)</td>
<td>28 (23–33)</td>
<td>0.42</td>
</tr>
<tr>
<td>BMI (kg/m^2)</td>
<td>28.4 (25.7–31.4)</td>
<td>23.7 (19.9–28.2)</td>
<td>0.05</td>
</tr>
<tr>
<td>WHR†</td>
<td>0.78 (0.75–0.81)</td>
<td>0.71 (0.67–0.76)</td>
<td>0.009</td>
</tr>
<tr>
<td>S_g (l/min)</td>
<td>117 (112–122)</td>
<td>114 (106–121)</td>
<td>0.41</td>
</tr>
<tr>
<td>Di (mU/min)†</td>
<td>80 (76–83)</td>
<td>76 (71–80)</td>
<td>0.17</td>
</tr>
<tr>
<td>Calculated free testosterone (pmol/l)‡</td>
<td>20.5 (15.4–27.1)</td>
<td>11.6 (6.4–21.2)</td>
<td>0.048</td>
</tr>
<tr>
<td>Fasting insulin (pmol/l)†</td>
<td>29 (21–40)</td>
<td>19 (10–39)</td>
<td>0.21</td>
</tr>
<tr>
<td>AUC_{insulin} (mmol-min/l)†</td>
<td>24.8 (19.0–32.3)</td>
<td>15.0 (10.5–21.5)</td>
<td>0.04</td>
</tr>
<tr>
<td>IGI_g (pmol/nmol)</td>
<td>69 (51–94) (n = 25)</td>
<td>61 (37–100)</td>
<td>0.66</td>
</tr>
<tr>
<td>S_g (l/min)</td>
<td>0.023 (0.019–0.027)</td>
<td>0.028 (0.019–0.038)</td>
<td>0.17</td>
</tr>
<tr>
<td>AUC_{DCI-IPG} (%-min)‡</td>
<td>142 (93–216) (n = 24)</td>
<td>113 (63–201)</td>
<td>0.53</td>
</tr>
<tr>
<td>Urinary clearance of DCI (ml/min)⁰</td>
<td>1070 (629–1821) (n = 23)</td>
<td>1539 (682–3471)</td>
<td>0.43</td>
</tr>
<tr>
<td>Plasma MYO (µmol/l)</td>
<td>0.04 (0.02–0.09)</td>
<td>0.03 (0.01–0.08)</td>
<td>0.54*</td>
</tr>
<tr>
<td>24 h urinary DCI (µmol/d)</td>
<td>0.8 (0.3–1.9)</td>
<td>1.7 (0.3–11.1)</td>
<td>0.39</td>
</tr>
<tr>
<td>24 h urinary MYO (µmol/d)</td>
<td>12.7 (5.3–31.1) (n = 26)</td>
<td>36.9 (4.2–325)</td>
<td>0.26</td>
</tr>
<tr>
<td>Urinary clearance of MYO (ml/min)⁰</td>
<td>414 (93–216) (n = 24)</td>
<td>113 (63–201)</td>
<td>0.53</td>
</tr>
<tr>
<td>24 h urinary DCI (µmol/d)</td>
<td>81 (66–98)</td>
<td>70 (49–100)</td>
<td>0.43</td>
</tr>
<tr>
<td>24 h urinary MYO (µmol/d)</td>
<td>27.2 (2.2–3.3)</td>
<td>2.4 (1.7–3.4)</td>
<td>0.56</td>
</tr>
<tr>
<td>Incremental AUC_{DCI-IPG} (%-min)</td>
<td>2.7 (13.063–15 345)</td>
<td>17 276 (14 907–20 021)</td>
<td>0.01‡</td>
</tr>
<tr>
<td>AUC_{DCI-IPG}/AUC_{insulin}</td>
<td>4.0 (2.9–5.4)</td>
<td>8.0 (5.0–12.6)</td>
<td>0.02‡</td>
</tr>
</tbody>
</table>

Results are expressed as means with 95% confidence intervals, unless indicated otherwise. BMI, body mass index; WHR, ratio of waist circumference to hip circumference; BP, blood pressure; AUC_{insulin}, area under the insulin levels curve during OGTT; IGI_g, insulinogenic index at 15 min during the OGTT; S_g, glucose-mediated disappearance of glucose; S_g/(by FSIVGTT), sensitivity to insulin measure by the frequent sampling intra-venous glucose tolerance test; AUC_{DCI-IPG}, area under the DCI-containing inositolphosphoglycan bioactivity curve during OGTT; incremental AUC_{DCI-IPG}, area under the DCI-IPG curve above baseline during OGTT (i.e. AUC_{DCI-IPG} minus area under baseline curve that is by definition 100% × 120 min); AUC_{DCI-IPG}/AUC_{insulin}, ratio of AUC_{DCI-IPG} to AUC_{insulin} measures. To convert values for free testosterone to ng/dl, divide by 34.7; and to convert values for insulin to mU/ml, divide by 6.945. *Unequal variance Welch ANOVA test. †Geometric means. ‡P = 0.01 or §P = 0.047 when adjusted for BMI and WHR.
Since we determined that our primary variables of interest are significantly associated with hyperinsulinemia, we sought to determine if parameters of inositols metabolism were significant predictors of AUC_{insulin}, independently from classical determinants of hyperinsulinemia that were assessed in our study. Thus, a third stepwise analysis was performed. Variables associated with AUC_{insulin} during univariate analyses ($P < 0.10$) are reported in Table II, and the final model in Table III. The total number of subjects included in the final model is 36 due to unavailable Si in one PCOS women. As shown, the best parameters significantly associated with AUC_{insulin} are insulin sensitivity (S_i, partial $P < 0.001$), plasmonic MYO levels (partial $P = 0.008$), MYO urinary clearance (partial $P = 0.01$) and AUC_{DCI-IPG} (partial $P = 0.02$), which explained 71.4% of the variance of AUC_{insulin} ($R^2 = 0.714$, $P < 0.001$). Of note, inositols and DCI-IPG parameters increased the predictive precision of the model from 34.9%, when considering S_i alone, to 71.4%, i.e. an absolute increase of 36.5%.

Discussion

This study was designed to test the hypothesis that Greek women with or without PCOS would exhibit alterations in the metabolism of DCI and insulin-stimulated release of the putative insulin mediator DCI-IPG similar to those previously observed in American PCOS women and controls (Baillargeon et al., 2006). Direct comparisons of both populations are possible because identical protocols and laboratory techniques were used. In Greek PCOS women, glucose-stimulated insulin levels (AUC_{insulin}) were significantly increased by 65% compared with normal Greek women, which reflects mainly compensatory hyperinsulinemia. Concurrently, insulin-stimulated release of bioactive DCI-IPG (incremental AUC_{DCI-IPG}) was reduced by 60%. Moreover, this last analysis remained significant after correction for BMI and WHR. Thus, these findings support a role for defective DCI-IPG insulin mediator activity in the pathophysiology of PCOS, which is independent of adiposity and most likely related to compensatory hyperinsulinemia.
AUCDCI-IPG, even after correction for adiposity. After combination of insulin sensitivity alone; and only 29% of the contribution of insulin sensitivity alone; and only 29% of the compensatory hyperinsulinemia in Greek women. Another important finding of our study is that parameters of PCOS status, measured by pelvic ultrasonography and fasting plasma insulin concentrations, were related to compensatory hyperinsulinemia in all women, regardless of adiposity, origin and PCOS status. However, PCOS and normal women with BMI < 30 kg/m². Furthermore, Greek PCOS women were substantially less insulin-resistant, less hyperinsulinemic and less hyperandrogenic than American PCOS women. Thus, a milder clinical syndrome in Greek women might explain the lack of difference in the development of insulin resistance and compensatory hyperinsulinemia, but alteration in MYO metabolism may be more apparent simply because the amounts of MYO in plasma and tissues largely exceed the amounts of DCI. It is however unknown why MYO and DCI clearance correlate in Greek but not in American women. This might be due to differences in diet content of inositols, although this variable was not assessed in our study.

Nonetheless, it remains possible that the metabolism of MYO also plays a role in the hyperinsulinemia of Greek women, resulting in less availability of MYO to tissues. Since it has been shown in tissues that [3H]MYO is converted to [3H]DCI and subsequently to [3H]DCI-IPG (Pak et al., 1993), decreased MYO availability would result in less DCI substrate for the generation of the DCI-IPG insulin mediator. A possible role of MYO in the pathophysiology of PCOS in women of Mediterranean origin is further supported by the results of a recent randomized-controlled trial in 40 lean Italian PCOS patients undergoing ovulation induction for ICSI (Papaleo et al., 2007b). This trial demonstrated that oral administration of MYO reduced r-FSH dose and duration, estradiol levels at hCG injection and the number of oocytes of poor quality in PCOS women. Another uncontrolled prospective study in 25 infertile Italian PCOS women found that oral MYO increases spontaneous ovarian activity and fertility (Papaleo et al., 2007a). Surprisingly, 24 h uClDCI and other inositols did not differ in Greek PCOS women compared with Greek controls, which contrasts with our previous findings in American PCOS women (Baillargeon et al., 2006). One possible explanation is that American women were substantially more obese, i.e. the mean BMI for American women was 33.9 versus 28.4 kg/m² for Greek women. Indeed, sub-group analyses in American PCOS women revealed that DCI urinary clearance was increased only in PCOS women whose BMI was ≥ 30 kg/m² (Baillargeon et al., 2006), and was similar in PCOS and normal women with BMI < 30 kg/m². Furthermore, Greek PCOS women were substantially less insulin-resistant, less hyperinsulinemic and less hyperandrogenic than American PCOS women. Thus, a milder clinical syndrome in Greek women might explain the lack of difference in urinary DCI clearance between Greek PCOS and normal women. Finally, this discrepancy might just result from the small number of Greek control women.

On the basis of these results, we propose that higher renal clearance of inositols tends to decrease plasma inositols levels and, ultimately, intra-tissue generation of DCI-IPG. Decreased

### Table II Variables associated with hyperinsulinemia (AUC_{insulin}) during univariate analyses.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Direction of association</th>
<th>P*</th>
</tr>
</thead>
<tbody>
<tr>
<td>PCOS status, n (%)</td>
<td>Positive</td>
<td>0.04</td>
</tr>
<tr>
<td>WHR</td>
<td>Positive</td>
<td>0.07</td>
</tr>
<tr>
<td>Diastolic BP (mmHg) (n = 36)</td>
<td>Positive</td>
<td>0.02</td>
</tr>
<tr>
<td>S_0 (by FSIVGTT)</td>
<td>Negative</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Plasma DCI (µmol/l)</td>
<td>Negative</td>
<td>0.09</td>
</tr>
<tr>
<td>Urinary clearance of DCI (ml/min)</td>
<td>Positive</td>
<td>0.04</td>
</tr>
<tr>
<td>Plasma MYO (µmol/l)</td>
<td>Negative</td>
<td>0.004</td>
</tr>
<tr>
<td>24 h urinary MYO (µmol/d)</td>
<td>Positive</td>
<td>0.04</td>
</tr>
<tr>
<td>Urinary clearance of MYO (ml/min)</td>
<td>Positive</td>
<td>0.001</td>
</tr>
<tr>
<td>AUC_{DCI-IPG} (%-min)</td>
<td>Negative</td>
<td>0.004</td>
</tr>
</tbody>
</table>

All variables from Table I, except fasting insulin. AUC_{DCI-IPG}-to-AUC_{insulin} ratio, DI and IGI15, were tested but only those associated to AUC_{insulin} with a P ≤ 0.10 are reported in this table. *By Pearson’s correlation tests, except for comparison of AUC_{insulin} by PCOS status that was tested using two-tailed unpaired Student’s t-tests. †Variable was log-transformed for analysis. ‡P ≤ 0.007 or §P < 0.03 when adjusted for BMI and WHR.

### Table III Best independent predictors of AUC_{insulin} (nmol/min-1) in Greek women with or without PCOS (n = 36).

<table>
<thead>
<tr>
<th>Variables</th>
<th>Partial P</th>
<th>Partial model R²*</th>
<th>Total model P</th>
</tr>
</thead>
<tbody>
<tr>
<td>S_0 (by FSIVGTT)</td>
<td>&lt;0.001</td>
<td>27.8%</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>AUC_{DCI-IPG} (%-min)</td>
<td>0.002</td>
<td>46.0%</td>
<td></td>
</tr>
<tr>
<td>Urinary clearance of MYO (ml/min)</td>
<td>0.003</td>
<td>62.5%</td>
<td></td>
</tr>
<tr>
<td>Plasma MYO (µmol/l)</td>
<td>0.005</td>
<td>71.0%</td>
<td></td>
</tr>
</tbody>
</table>

*R² for the model including all the variables that are listed up to this point (exclude the variables below this line). †Variable was log-transformed for analysis.

Importantly, this study also demonstrated that, when all Greek women were analysed together, hyperinsulinemia was significantly associated with increased 24 h uClDCI and reduced AUC_{DCI-IPG}, even after correction for adiposity. After combining primary data from this study with our previously published study in American women (Baillargeon et al., 2006), these associations remained highly significant. Finally, there was no interaction with origin or PCOS status. Therefore, these findings suggest that alterations in the metabolism of DCI and DCI-IPG are related to compensatory hyperinsulinemia in all women, regardless of adiposity, origin and PCOS status. However, DCI and DCI-IPG metabolism were not related to acute insulin response (IGI13 and AIRg) and beta-cell function (DI). Another important finding of our study is that parameters of inositols metabolism were the best independent predictors of high insulin levels, together with insulin resistance. Moreover, inositols and measures of DCI-IPG release increased the precision of predicting AUC_{insulin} by an additional 43% over the contribution of insulin sensitivity alone; and only 29% of the variability of AUC_{insulin} was explained by other unknown factors. This supports an important role of inositols metabolism and the DCI-IPG insulin mediator in the expression of compensatory hyperinsulinemia in Greek women.

However, it was unexpected that MYO plasma levels and urinary clearance (uClMYO) would be more strongly associated with higher insulin concentrations than corresponding parameters of DCI. Indeed, MYO metabolism was not altered in American PCOS women and was not correlated with insulin sensitivity or insulin levels (Baillargeon et al., 2006). This new finding may be explained in part by high correlations between MYO and DCI parameters in Greek women, such that the best predictors of uClDCI were uClMYO and plasmatic DCI levels. Thus, it is possible that the same renal defect affects urinary clearance of both inositols in Greek hyperinsulinemic women. Accordingly, DCI may still be the inositol implicated in the development of insulin resistance and compensatory hyperinsulinemia, but alteration in MYO metabolism may be more apparent simply because the amounts of MYO in plasma and tissues largely exceed the amounts of DCI. It is however unknown why MYO and DCI clearance correlate in Greek but not in American women. This might be due to differences in diet content of inositols, although this variable was not assessed in our study.

Nonetheless, it remains possible that the metabolism of MYO also plays a role in the hyperinsulinemia of Greek women, resulting in less availability of MYO to tissues. Since it has been shown in tissues that [3H]MYO is converted to [3H]DCI and subsequently to [3H]DCI-IPG (Pak et al., 1993), decreased MYO availability would result in less DCI substrate for the generation of the DCI-IPG insulin mediator. A possible role of MYO in the pathophysiology of PCOS in women of Mediterranean origin is further supported by the results of a recent randomized-controlled trial in 40 lean Italian PCOS patients undergoing ovulation induction for ICSI (Papaleo et al., 2007b). This trial demonstrated that oral administration of MYO reduced r-FSH dose and duration, estradiol levels at hCG injection and the number of oocytes of poor quality in PCOS women. Another uncontrolled prospective study in 25 infertile Italian PCOS women found that oral MYO increases spontaneous ovarian activity and fertility (Papaleo et al., 2007a). Surprisingly, 24 h uClDCI and other inositols did not differ in Greek PCOS women compared with Greek controls, which contrasts with our previous findings in American PCOS women (Baillargeon et al., 2006). One possible explanation is that American women were substantially more obese, i.e. the mean BMI for American women was 33.9 versus 28.4 kg/m² for Greek women. Indeed, sub-group analyses in American PCOS women revealed that DCI urinary clearance was increased only in PCOS women whose BMI was ≥ 30 kg/m² (Baillargeon et al., 2006), and was similar in PCOS and normal women with BMI < 30 kg/m². Furthermore, Greek PCOS women were substantially less insulin-resistant, less hyperinsulinemic and less hyperandrogenic than American PCOS women. Thus, a milder clinical syndrome in Greek women might explain the lack of difference in urinary DCI clearance between Greek PCOS and normal women. Finally, this discrepancy might just result from the small number of Greek control women.

On the basis of these results, we propose that higher renal clearance of inositols tends to decrease plasma inositols levels and, ultimately, intra-tissue generation of DCI-IPG. Decreased
insulin-stimulated release of DCI-IPG, in turn, contributes to lower insulin sensitivity and a resulting compensatory hyperinsulinemia. Variability in urinary clearance of inositols might be due to obesity, lifestyle, or PCOS-related genetic and/or other genetic factors. Thus, renal clearance of inositols probably contributes to the variability of insulin sensitivity in women. Although it might not be a major contributor to insulin resistance in the general population, this contribution might be more important in some individuals, such as PCOS women.

Another possible scenario is that compensatory hyperinsulinemia itself induces a defect that increases renal clearance of inositols. It is also possible that both scenarios are valid and present in hyperinsulinemic women, causing a ‘vicious cycle’ whereby insulin resistance and compensatory hyperinsulinemia are amplified in these women through the induction of a defect in renal clearance of inositols.

Despite interesting and highly significant results, even after correction for adiposity, there are limitations to our study. First, the number of Greek normal controls is relatively small. Therefore, it was not possible to ascertain whether non-significant differences between Greek PCOS and control were due to random distribution instead of true physiological differences. This also holds true for the absence of a modifying effect of PCOS status on reported correlations in Greek women alone. Second, since only PCOS and normal women were studied, inference to other populations would be speculative. Furthermore, because of the cross-sectional design of this study, it is unknown if significant correlations reflect causative links or only associations. Also, cases and controls were not matched for adiposity (e.g. BMI), such that an effect of obesity cannot be completely excluded. Indeed, adjusting for BMI, as we did, assumes linear relationships, but it may not be the case. Finally, our control group was recruited from an endocrine clinic and might not be entirely representative of the community.

In conclusion, results of the present study demonstrated that hyperinsulinemic Greek women with or without PCOS displayed increased urinary clearance of inositols and decreased insulin-stimulated release of bioactive DCI-IPG, independently of adiposity. Thus, increased inositols clearance and reduced DCI-IPG generation might contribute to insulin resistance and compensatory hyperinsulinemia in Greek as well as American women. These findings offer potential mechanisms for previously reported benefits of DCI or MYO oral administration on metabolic and clinical derangements in PCOS women (Nestler et al., 1999; Iuorno et al., 2002; Gerl et al., 2003; Papaleo et al., 2007a,b). Whether abnormalities of DCI metabolism and DCI-IPG release are specific to PCOS or relevant to other disorders characterized by insulin resistance is still unknown.

**Funding**

Funding for this work was provided by the National Institutes of Health (R01HD35 629 to J.E.N., K24HD40 237 to J.E.N., R01DK58 698 to R.E.O.) and the Fonds de Recherche en Santé du Québec (Award #3158 to J-P.B.).

**Reference**


Submitted on November 13, 2007; resubmitted on February 24, 2008; accepted on March 4, 2008