Correlation of Serum Lipoprotein Ratios with Insulin Resistance in Infertile Women with Polycystic Ovarian Syndrome: A Case Control Study

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Abstract

Background: Dyslipidemia and insulin resistance (IR), occurring in most infertile women with polycystic ovarian syndrome (PCOS), increase the risk of cardiovascular disease (CVD) and type 2 diabetes. This study aimed to assess the relationships between lipoprotein ratios and IR in PCOS women.

Materials and Methods: Thirty six infertile women with PCOS selected based on Androgen Excess Society (AES) criteria and 29 healthy women matched for age were recruited to this case-control study. After physical measurements, fasting serum glucose (Glu), insulin and lipid profile levels [triglycerides (TGs), total cholesterol (TC), low-density lipoprotein-cholesterol (LDL-C) and high-density lipoprotein-cholesterol (HDL-C)] were measured, while lipoprotein ratios (TC/HDL-C, LDL-C/HDL-C, TG/HDL-C) were calculated. IR was also calculated using homeostasis model assessment (HOMA)-IR. The optimal cutoffs of lipoprotein ratios in relation to HOMA-IR were calculated based on the Receiver Operating Characteristics (ROC) curve analysis using the area under curve (AUC).

Results: Waist circumference (WC), insulin levels, HOMA-IR, TG levels, and all lipoprotein ratios were significantly higher, while HDL-C was lower in PCOS group as compared to healthy controls. All lipoprotein ratios, TG levels, and WC are significantly correlated with insulin levels and HOMA-IR. Among lipoprotein ratios, the highest AUC of the ROC belonged to TG/HDL-C ratio with sensitivity of 63.6% and specificity of 84.4% (TG/HDL-C>3.19) as a marker of IR in infertile PCOS women.

Conclusion: Lipoprotein ratios, particularly TG/HDL-C, are directly correlated with insulin levels and can be used as a marker of IR (HOMA-IR) in infertile PCOS patients.

Keywords: Lipoprotein, Infertility, PCOS, Insulin Resistance

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Introduction

Polycystic ovary syndrome (PCOS) is the most common gynecological endocrinopathy disorder and the frequent cause of oligo-ovulatory infertility (1). Abnormalities with ovulation are the cause of infertility in about one third of couples attending infertility clinics that count for 90% of these cases (2). PCOS is generally characterized by chronic anovulation, hyperandrogenism and ovarian polycystic changes that are detected by an ultrasound scan in a clinic (3). The estimated prevalence of PCOS based on the criteria used for diagnosis and recruitment process of the study population has been reported between 2.2 and 26% in different countries (4). Although the etiology of PCOS is still unknown, it has been demonstrated that PCOS is a metabolic disorders rather than a reproductive endocrine disease (3). Insulin is a key component in the pathophysiology of PCOS (1). On average, PCOS patients have higher triglyceride (TG), lower high density lipoprotein-cholesterol (HDL-C) and higher low density lipoprotein-cholesterol (LDL-C) levels than their non-PCOS matched group (5). Insulin resistance (IR), hyperinsulinaemia and dyslipidemia are diagnosed among 50 to 70% of patients with PCOS (6). There is a drastic improvement in PCOS complication when is accompanied by modulation of IR (1). Therefore, PCOS is associated with increased risk of metabolic abnormalities, indicating that the patients are at the risk of developing type 2 diabetes and cardiovascular disease (CVD) (3).

Despite modern treatment options for infertilities and considering economic aspects, it is reasonable to give specific attention to cost effective and easily applied methods for predicting metabolic abnormalities at population level (7). Routine methods for measuring IR are hyperinsulinemic-euglycemic clamp technique (a gold standard to assess insulin sensitivity) (8), homeostasis model assessment (HOMA)-IR, Bennett index, Li Guangwei index, quantitative insulin sensitivity check index (QUICKI), and fasting serum glucose (Glu)/insulin ratio (G/I). Due to being complex, expensive and time-consuming, the latter methods are of limited use in clinical and epidemiological studies (9). Thus for daily clinical practice, it is necessary to use other methods for measuring IR, which are lower in costs and applicable to the general population.

In order to provide a new idea to evaluate IR in infertilities associated with PCOS, the possibility of establishing the values of total cholesterol (TC)/HDL-C, TG/HDL-C, and LDL-C/HDL-C ratios, waist circumference (WC) as surrogates, as well as LDL-C, TC, and TG levels to estimate insulin levels and IR was investigated. By using Receiver Operating Characteristic (ROC) curves in our subjects, the accuracy of the mentioned parameters was received.

Materials and Methods

Subjects

In this case-control study, subjects were selected among women aged 19 to 35 years who visited a private reproductive medical center, Tabriz, Iran, during the period of February till April 2013, for infertility due to PCOS. Selection was done by the standardized protocol for the initial evaluation. A total of 35 patients were identified as PCOS cases according to the Androgen Excess Society (AES, 2006) criteria (1), while 29 age-matched healthy women (without any infertility and PCOS disorders) were recruited in the study as the control group. Inclusion criteria for case group were as follows: married, clinical and/or biochemical hyperandrogenism, and ovarian dysfunction (oligoanovulation and/or polycystic ovaries detected by ultrasound scans). Exclusion criteria were as follows: congenital adrenal hyperplasia, androgen-secreting tumors, taking androgenic/anabolic medications, Cushing syndrome, severe IR syndrome, thyroid dysfunction, hyperprolactinemia, diabetes, hypertension, CVD, taking vitamins and supplements during the 3 months prior to the study, evidence of recent or recurrent infection, and smoking or drinking alcohol.

Physical measurements

Body weight was measured without shoes with minimal amount of clothing using a digital scale (SECA, Germany) to the nearest 0.1 kg. Height was measured using a non-stretchable stadiometer (SECA, Germany) to the nearest 0.1 cm. Body mass index (BMI) was calculated as weight in kg divided to height in squared me-
ters. WC was measured at the midpoint between
the lowest rib and the top of the lateral border
of iliac crest during minimal respiration. Sys-
tolic blood pressure (SBP) and diastolic blood
pressure (DBP) were measured using Spot Vital
 Signs Device (Welch Allyn, USA). Participants
were asked to lie down and relax for approxi-
mately 8 to 10 minutes, after which three blood
pressure measurements were recorded at five-
minute intervals.

Blood analysis

After 12-hour overnight fast, blood sam-
ple s were collected. Serum and plasma
samples were separated using a centrifuge
(Beckman Coulter Inc., USA) at 1500 rpm
for 15 minutes. Fasting insulin levels were
measured using enzyme-linked immuno-
sorbent assay (ELISA) kits (Monobind Inc.,
USA). Fasting plasma Glu was measured using
enzymatic procedures by an automatic
analyzer (Abbott, USA). IR was estimated
by HOMA using the following formula: HO-
MA-IR=fasting insulin (μU/ml)×fasting Glu
(mg/dl)/405 (10). The concentrations of TC
and TG were measured using enzymatic pro-
cedure with commercial kits (Pars Azmon,
IRI), while HDL-C was measured by a direct
method using polyethylene-glycol-pretreated
enzymes by an automatic analyzer (Abbott,
USA). LDL-C was calculated using Friede-
wald’s formula (11). Lipoprotein ratios (TC/
HDL-C, TG/HDL-C and LDL-C/HDL) were
then calculated.

Ethical considerations

This study was approved by the Medical Ethics
Committee of Ahvaz Jundishapur University and
all participants gave an informed consent before
commencing the study. The code of Ethics Com-
mittee is ETH-702, and registered code of study is
NRC-9110.

Statistical analysis

Results were expressed as mean ± SD. Lev-
ene’s test for equality of variances was used.
The differences between concerning continu-
ous and categorical variables were analyzed
using unpaired t test (or Mann-Whitney U test
for non-normally distributed data) and χ² test,
respectively. Correlations were determined
by Spearman correlation coefficient method.
ROC curves were used to estimate the sensitiv-
ity and specificity of serum lipoprotein ratios
to diagnose IR. P values less than 0.05 were
considered statistically significant. All statisti-
cal analyses were performed using Statistical
Package for Social Sciences 20.0 (SPSS, SPCC
Inc., USA) software.

Results

The control group was matched with the pa-
tient group for age. Although the values of BMI,
BP, TC, LDL-C, TG and fasting serum Glu were
found to be higher in the infertile PCOS group
than in the control group, indicating that these
differences were not statistically significant. A
higher insulin level and HOMA-IR value were
observed in patients group compared to the con-
trol group (P<0.001 and P=0.024, respectively).
TG levels (P=0.009) as well as the values of TC/
HDL (P=0.002), TG/HDL (P=0.047), LDL/HDL
(P=0.002) and WC (P<0.001) were significantly
higher, while HDL-C levels (P=0.003) were lower
in the cases compared to those of their healthy
counterparts. The results are shown in Table 1.

HOMA-IR value in the patients showed a
positive correlation with TG levels (r=0.56,
P<0.01) as well as the values of TC/HDL-C
(r=0.34, P<0.05), TG/HDL-C (r=0.49, P<0.01),
LDL-C/HDL-C (r=0.33, P<0.05), and WC
(r=0.37, P<0.05). However, HOMA-IR value
showed no significant correlation with TC,
LDL and HDL concentrations. Serum insulin
levels are positively correlated with TG level
(r=0.46, P<0.01), TC level (r=0.33, P<0.05),
and TG/HDL value (r=0.39, P<0.05). We found
no significant correlation between serum insu-
lin levels and LDL-C, HDL-C, TC/HDL, LDL/
HDL and WC values in our patients. The re-
results are shown in Table 2.

According to the ROC curve analysis, all lipid
ratios (TG/HDL-C, TC/HDL-C, and LDL/HDL-
C) showed an area under curve (AUC) greater
than 0.5. Thus, as an effective diagnostic marker for IR
in PCOS patients, the AUC of TG/HDL-C was the
highest with sensitivity of 63.6% and specificity of
84.4% (TG/HDL-C>3.19). The results are shown
in Table 3 and Figure 1.
**Table 1:** Baseline and clinical characteristics of two groups (age range 19-35 years)

| Variables                  | Infertile PCOS (n=36) | Healthy control (n=29) | \(P\) value<sup>a</sup> |
|----------------------------|------------------------|------------------------|--------------------------|
| Age                        | 26.36 ± 4.2            | 27.96 ± 2.47           | 0.107                    |
| BMI (kg/m²)                | 26.72 ± 4.39           | 25.55 ± 4.3            | 0.286                    |
| BMI (%)<sup>b</sup> BMI≥25 | 72.2                   | 48.3                   | 0.049                    |
| WC (cm)                    | 94.77 ± 10.36          | 85.06 ± 8.48           | <0.001                   |
| SBP (mmHg)                 | 118.66 ± 8.98          | 116.89 ± 6.03          | 0.209                    |
| DBP (mmHg)                 | 78.19 ± 6.98           | 76.37 ± 5.15           | 0.274                    |
| Fasting serum Glu (mg/dL)  | 94.47 ± 11.88          | 89.86 ± 8.25           | 0.081                    |
| Insulin (μU/mL)<sup>b</sup> | 21.41 ± 14.14         | 16.24 ± 11.55          | 0.029                    |
| HOMA-IR<sup>b</sup>        | 5.16 ± 3.72            | 3.41 ± 2.53            | 0.024                    |
| TC (mg/dL)                 | 214.83 ± 43.97         | 202.68 ± 46.44         | 0.285                    |
| TG (mg/dL)                 | 139.28 ± 66.98         | 98.17 ± 50.72          | 0.009                    |
| HDL-C (mg/dL)              | 42.88 ± 10.2           | 52.06 ± 13.71          | 0.003                    |
| LDL-C (mg/dL)              | 143.69 ± 36.25         | 129.51 ± 35.70         | 0.119                    |
| TC/HDL-C ratio             | 5.16 ± 1.22            | 4.11 ± 1.36            | 0.002                    |
| TG/HDL-C ratio             | 3.62 ± 2.17            | 2.44 ± 2.52            | 0.047                    |
| LDL-C/HDL-C ratio          | 3.44 ± 0.98            | 2.62 ± 1.02            | 0.002                    |

PCOS; Polycystic ovarian syndrome, BMI; Body mass index, WC; Waist circumference, SBP; Systolic blood pressure, DBP; Diastolic blood pressure, Glu; Glucose, HOMA-IR; Homeostasis model assessment of insulin resistance, TC; Total cholesterol, TG; Triglyceride, HDL-C; High density lipoprotein-cholesterol, LDL-C; Low density lipoprotein-cholesterol,<sup>a</sup>; Statistical analyses performed by unpaired t test for comparison,<sup>b</sup>; Statistical analyses performed by Mann-Whitney U test and<sup>c</sup>; Statistical analyses performed by Chi-squared test. Data are the mean ± SD.

**Table 2:** Spearman’s correlations of lipid profile, lipoprotein ratios and WC values with serum insulin level and IR in infertile women with PCOS

| Variables                  | Serum insulin levels | IR          |
|----------------------------|----------------------|-------------|
| TGs                        | 0.46<sup>b</sup>     | 0.56<sup>b</sup>| |
| TC                         | 0.33<sup>a</sup>     | 0.316       | |
| LDL-C                      | 0.29                 | 0.28        | |
| HDL-C                      | 0.14                 | 0.08        | |
| TC/HDL                     | 0.3                  | 0.34<sup>a</sup>| |
| TG/HDL                     | 0.39<sup>a</sup>     | 0.49        | |
| LDL/HDL                    | 0.28                 | 0.33<sup>a</sup>| |
| WC                         | 0.32                 | 0.37        | |

IR; Insulin resistance, PCOS; Polycystic ovary syndrome, WC; Waist circumference, TC; Total cholesterol, TG; Triglyceride, HDL-C; High density lipoprotein-cholesterol, LDL-C; Low density lipoprotein-cholesterol,<sup>a</sup>; Statistical analyses performed by unpaired t test for comparison,<sup>b</sup>; Statistical analyses performed by Mann-Whitney U test and<sup>c</sup>; Statistical analyses performed by Chi-squared test. Data are the mean ± SD.

**Table 3:** Serum lipoprotein ratios, AUC, cut-off points and sensitivity and specificity calculated from ROC curves for the detection of PCOS with IR

| Serum lipoprotein ratios | AUC± SE | 95% CI | Cut-off point | Infertile PCOS patients | P value |
|-------------------------|---------|-------|---------------|-------------------------|---------|
|                         | Sensitivity (%) | Specificity (%)| |
| TG/HDL                  | 0.743 ± 0.062 | 0.622-0.864 | 3.19 | 63.6 | 84.4 | 0.001 |
| TC/HDL                  | 0.651 ± 0.069 | 0.515-0.786 | 4.37 | 69.7 | 65.6 | 0.037 |
| LDL/HDL                 | 0.638 ± 0.070 | 0.502-0.775 | 2.84 | 69.7 | 63.5 | 0.055 |

CI; Confidence interval, IR; Insulin resistance, AUC; Area under curve area, ROC; Receiver operating characteristic, PCOS; Polycystic ovarian syndrome, TC; Total cholesterol, TG; Triglyceride, HDL; High density lipoprotein and LDL; Low density lipoprotein.
Discussion

Among the factors responsible for a reduction in fecundity and successful pregnancy, the hormonal changes associated with various factors are considered as an important cause for interrupting normal ovulatory menstrual cycles (12). Among these factors, visceral adiposity is a common finding in PCOS patient, even when the subjects are not classified as overweight (25<BMI<29.9) (13). According to our findings, we observed a significant different in WC between groups. Although the difference in BMI index between cases and controls was not significant, but in case group, percentage of overweight BMI was higher than that of controls. Also there was a significantly positive correlation between WC and IR. Our Findings are in agreement with those of some previous studies (13-15). However, the latter results differ from those of the study conducted by Iuhas et al. (16). In their study, visceral fat area showed no significant difference in PCOS and healthy subjects, which might be due to the difference in method of measuring visceral fat and larger sample size. Pathophysiology of PCOS is unknown. It is regarded as an endocrinol disorder due to IR, which presents in about 70% of PCOS patients (17, 18). In PCOS patients, IR is mostly associated with dyslipidemia. Methods used for measuring IR are mostly sophisticated and expensive that are not applicable for epidemiological studies. Hence, more reasonable methods for IR measurements have been investigated in several studies, of which lipoprotein ratios were proposed for the identification of IR as an alternative method. Our investigation was carried out in order to provide evidences for the application of lipoprotein ratios as an indicator of IR in infertile PCOS women. In this investigation, PCOS was diagnosed by AES criteria, while for the first time, subjects were selected among infertile PCOS women.

This study showed the case group had higher TG levels and lower HDL-C levels compared to control group. While no significant difference was detected in TC and LDL-C levels between groups. Most studies have shown low levels of HDL-C in women with PCOS, but composition of HDL in PCOS is still unknown. There is still a need for further studies in order to determination of the HDL-C composition in these patients. One of the mechanisms that could explain the observed difference is the activity of hepatic lipase (HL) enzyme induced by IR and hyperan-
dyslipidemia, which removes lipid from HDL and plays as a key role of the lipid-depleted HDL particles in PCOS patients. Also insulin-resistant states along with low HDL levels are frequently associated with hypertriglyceridemia. However, another possible mechanism of dyslipidemia in PCOS could be a reduction in clearance of triglyceride-rich proteins (19).

Result of this study demonstrated a significant association in TC/HDL-C, TG/HDL-C and LDL-C/HDL-C ratios and TG with IR (HOMA-IR) in PCOS patients. In a study on women with PCOS, Xiang et al. (20) also suggested that serum lipoprotein ratios could be used as a marker of IR due to the significant positive correlation of the indices with IR. However, in their study, Rotterdam criteria were used for diagnosing PCOS, so there was a significant difference in terms of BMI between case and control groups, which could be a confounding factor. Hence the present study was designed more specifically by use of updated criteria (AES) on infertile women for diagnosing PCOS. Moreover in our study BMI was not significantly different between case and control groups, which could justify the confounding impact of BMI on results. Serum lipoprotein ratios were also reported to be significantly correlated to IR in type 2 diabetes patients (21). Furthermore TG/HDL could be considered as a simple reliable indicator to determine IR in healthy (22) and severely obese non-diabetic individuals (23). Our results on women with ovulatory disorder infertility also confirmed these findings. ROC curve analysis showed that TG/HDL-C, TC/HDL-C, and LDL-C/HDL-C with an AUC greater than 0.5 were effective and useful diagnostic markers for IR in infertile PCOS women. AUC of TG/HDL-C was the highest with sensitivity of 63.6% and specificity of 84.4% (TG/HDL-C>3.19). Xiang et al. (20) have also shown that AUC of TC/HDL-C had the highest sensitivity and specificity (TC/HDL-C>3.6). This discrepancy could be partially due to lesser sample size in our study or possible racial differences.

Future studies with higher sample size and more specific markers are needed to show the correlation between lipid ratios and IR in an extended level.

Conclusion

Our investigation demonstrated that despite the routine methods used for measuring IR, TC/HDL-C, TG/HDL-C, and LDL/HDL ratios could be regarded as simple, reliable and economic indicators of IR in PCOS infertile women. Moreover the combination of higher serum lipoprotein ratios and TG levels with abdominal obesity may predispose a group of patients to more marked risks for IR.

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