

Testosterone, obesity, and waist circumference as determinants of metabolic syndrome in Saudi women

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Background: High serum total testosterone is associated with metabolic syndrome (MS). This study aimed to identify possible alterations in total testosterone and their relationship with plasma glucose, blood pressure, and serum lipid profile.

Methods: One hundred forty-two female subjects were selected to participate in this study, and they were recruited by consultant physicians from the Clinic and Medical Out-Patient, King Abdulaziz Hospital, Kingdom of Saudi Arabia. The anthropometric characteristics were obtained from questionnaires by using standard methods. Blood samples were obtained for the determination of glucose, triglycerides, total cholesterol, low-density lipoprotein, and high-density lipoprotein by using enzymatic methods. Total testosterone was determined by enzyme-linked immunosorbent assay for the quantitative measurement of testosterone in human serum.

Results: Significantly higher concentrations of total testosterone, low-density lipoprotein, and glucose, but lower concentrations of high-density lipoprotein, were observed in subjects with MS compared with women without MS ($P < 0.05$).

Conclusion: This study suggests that high levels of total testosterone and disturbance in lipid profile were associated with MS in Saudi women.

Keywords: obesity, hyperlipidemia, testosterone, metabolic syndrome, lipid profile, androgen, insulin resistance

Background

Obesity has reached epidemic proportions globally,¹⁻³ with more than 1 billion adults being overweight – at least 300 million of them being clinically obese.⁴ It leads to adverse metabolic effects, blood pressure,⁵ and lipid and glucose metabolism mediated by insulin resistance.⁴ This constellation of cardiovascular risk factors constitutes the metabolic syndrome (MS).⁶ MS is associated with significant increase in cardiovascular morbidity and mortality,⁷ and it is the most common cause of death in the Western world.⁸ Waist circumference (WC) is the best measure of obesity which is combined with other variables to construct a country-specific definition of MS.⁹

In Saudi Arabia, epidemiological studies reported a potential growth in diabetes with a parallel increase in obesity rates.¹⁰ Currently, 23% of the health care expenditure along with 17% of direct medical costs is spent for diabetes.¹¹ The components of MS, including central obesity, are particularly common among Saudi population.¹²

The largest study, conducted over a 5-year period between 1995 and 2000 and covering different regions of Saudi Arabia, reported the overall MS prevalence in adults aged between 30 and 70 years to be about 40%, with the most common factor being low level of high-density lipoprotein cholesterol (HDL-C).¹³ The prevalence of cardiovascular

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disease (CVD) risk factors is high among women in Saudi Arabia especially in cases of obesity and physical inactivity.^{11,14} Public health authorities must implement solutions from a gender-specific aspect to reverse the trend and decrease the prevalence of CVDs among Saudi women.¹⁴

The first study of MS in Saudi Arabia was published in 2002.¹² Two other smaller studies conducted in the central region of Saudi Arabia reported the prevalence to be 31.4% in men¹⁵ and 35.3% in the whole population.¹⁶ Another study, conducted recently in the city of Jeddah in the western region, reported the MS prevalence to be 21% with low HDL-C as a common component.¹³ The present study was conducted to evaluate the levels of testosterone, obesity, and WC as determinants of MS in Saudi women.

Testosterone is a hormone that plays a key role in the metabolism of carbohydrate, fat, and protein. With a major influence on body fat composition in men, testosterone deficiency was suggested to be associated with reduced insulin sensitivity, impaired glucose tolerance, elevated triglycerides (TGs) and cholesterol, and low HDL-C in previous studies.¹⁷ All these factors are found in the MS and type 2 diabetes. Numerous studies showed associations between serum testosterone and insulin resistance or MS and type 2 diabetes risk in women; however, the conclusion about causal relationship still needs more studies.¹⁸ The present study was aimed to address testosterone levels with the components of MS in Saudi women.

Subjects and methods

Ethical approval

This retrospective study was conducted in the Biochemistry Laboratory, Department of Medical Laboratories, Faculty of Medicine, Umm Al-Qura University, Kingdom of Saudi Arabia from January 2015 to March 2017. Written informed consent was obtained from all participants. The overall study methodology was approved by the Biomedical Ethics Committee, Faculty of Medicine, Umm Al-Qura University, Mecca Al-Mukarama, Kingdom of Saudi Arabia.

Subjects

A total of 142 female subjects aged between 20 and 60 years were recruited for this study. Of them, 95 subjects with high WC (≥ 88 cm) and body mass index (BMI ≥ 30 kg/m²) and 47 apparently healthy women in normal weight as controls (BMI ≤ 25 kg/m²) were selected. Those on lipid-lowering drugs and hormonal medications, with CVDs like stroke, and who did not give consent were excluded. Information of age, maternity, menarche, and menstrual cycle disturbance were obtained using a questionnaire. Weight, height, WC,

and systolic and diastolic blood pressures (SBP and DBP, mean of three consecutive measurements) were measured.

Sampling

Following the overnight fasting period, 5 mL of whole venous blood samples were withdrawn from each subject. The sample was divided into two parts (plain and EDTA tubes) as serum and plasma samples, respectively. The samples were used for the estimation of plasma, fasting blood glucose, and total testosterone. Serum was used for the estimation of TGs, total cholesterol (TC), low- and high-density lipoproteins (LDL and HDL), and total testosterone.

Metabolic syndrome

The subjects were recruited by consultant physicians from the Clinic and Medical Out-Patient, King Abdulaziz Hospital. MS is defined by using National Cholesterol Education Program Adult Treatment Panel III guidelines, and it was diagnosed on the basis of five major components: 1) hypertriglyceridemia TG ≥ 150 mg/dL; 2) WC ≥ 102 cm for men and ≥ 88 cm for women; 3) SBP ≥ 130 mmHg, DBP ≥ 85 mmHg; 4) HDL-C ≤ 50 ; and 5) fasting plasma glucose ≥ 110 mg/dL.¹⁹

Biochemical investigation

Serum TC, TGs, HDL-C, and LDL-C were estimated by using direct homogenous method and enzymatic colorimetric test. The kits were provided by HUMAN Gesellschaft für Biochemica und Diagnostica mbH.

Fasting plasma glucose concentration was determined with an automated analyzer (HumaStar 80) by using glucose oxidase-phenol and 4-aminoantipyrine (GOD-PAP) method and an enzymatic colorimetric test kit. The kits were provided by HUMAN Gesellschaft für Biochemica und Diagnostica mbH (REF 10720p).

Hormone assays

The serum level of total testosterone was determined by using the enzyme-linked immunosorbent assay (ELISA) test for the quantitative determination of testosterone in human serum. The kits were provided by HUMAN Gesellschaft für Biochemica und Diagnostica mbH (REF 55010, 96 tests), and the test was performed according to the manufacturer's instructions and with the lowest detection limit of 0.06 ng/mL and a coefficient of variation $\leq 10\%$.

Blood pressure

Blood pressure was measured by a Microlife blood pressure monitor (Microlife A200 AFIB, Taipei, Taiwan). Two values,

the systolic (upper) value and the diastolic (lower) value, are always measured. Any activity was avoided, and the woman was made to sit down for at least 5 minutes before measurement and to relax; close-fitting garments from the upper arm was removed, shirt sleeves were not rolled up, and the cuff was fitted closely and positioned 2 cm above the elbow. The cuff pumped up automatically. When the correct pressure was reached, the pumping was stopped and the pressure fell gradually; the cuff was removed after the measurement. Hypertension was defined as SBP ≥ 130 mmHg and DBP ≥ 85 mmHg.

Statistical analysis

All statistical analysis was performed using IBM SPSS software version 16.0 (SPSS Inc., Chicago, IL, USA). Descriptive measures were done for each variable in every group. Data in the form of continuous variables were presented as mean $\pm$ SD. Pearson correlation coefficient was used to determine the relationship between variables. Since the test shows that the studied variables are normally distributed for comparing the quantitative variables in the two studied groups, independent sample *t*-test was used. $P < 0.05$ was considered as statistically significant. All results are two-tailed. GraphPad Prism 7 software version 7.03 (GraphPad Software, Inc., La Jolla, CA, USA) was used for graphing.

Results

In this study, we included 142 Saudi women aged between 20 and 60 years. They were divided as cases of MS and controls. Blood pressure, blood glucose, lipid profile, and serum testosterone were measured along with the anthropometric data.

Table 1 provides the age, BMI, WC, blood pressure, and gynecologic data. It has been found that the mean age of the cases was 43.49 years and that of the controls was 39.49 years. The BMI and WC of women with MS were found to be 41.9 and 107.5 cm, respectively. Mean DBP was reported as 134 and 113 mm/Hg among case and controls, respectively.

Table 2 and Figures 1 and 2 show the estimation of blood glucose and lipid profile. The mean fasting blood glucose

was found to be 163.2 mg/dL among women with MS and 66.11 mg/dL among controls. The lipid profile was reported to be as follows: the mean TGs, TC, LDL, and HDL were 194.2, 286.3, 198.0, and 53.73 mg/dL among cases and 68.23, 113.9, 106.1, and 98.49 mg/dL for controls, respectively.

Correlation between lipids profile and BMI among the whole study group (cases and controls) is shown in Figure 2A–D, reflecting that a strong positive correlation exists between BMI and blood glucose, serum TGs, cholesterol, and LDL. A negative correlation was reported between BMI and serum HDL as shown in Figure 3.

Correlation between glucose and lipid profile with WC among the whole study group (cases and controls) is shown in

Table 2 Blood glucose and lipid profile among study groups

Index (mg/dL)	Control n=47	MS n=95	P-value
Glucose	66.11 $\pm$ 3.66	163.2 $\pm$ 6.13	<0.01
Triglyceride	68.23 $\pm$ 4.534	194.2 $\pm$ 11.61	<0.0001
Cholesterol	113.9 $\pm$ 6.347	286.31 $\pm$ 13.9	<0.0001
LDL-C	106.1 $\pm$ 6.86	198.0 $\pm$ 9.4	<0.0001
HDL-C	98.49 $\pm$ 4.77	53.73 $\pm$ 9.82	<0.0001

Note: Values are expressed as mean $\pm$ SD.

Abbreviations: LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; MS, metabolic syndrome.

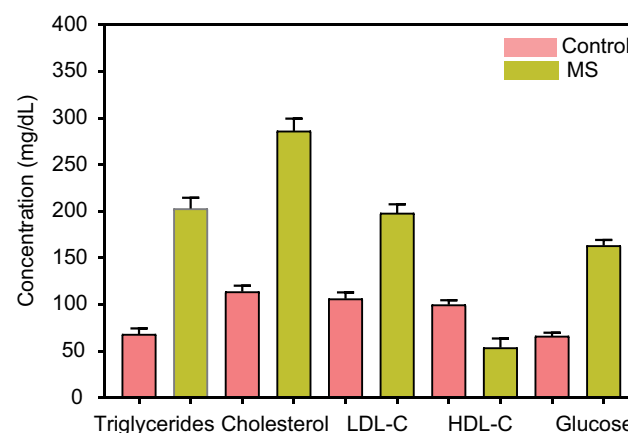


Figure 1 Glucose and lipid profile in controls and cases of MS.

Abbreviations: MS, metabolic syndrome; LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol.

Table 1 Anthropometric data among study female groups

Index	Controls n=47	MS n=95	P-value
Age (years)	39.49 $\pm$ 10.23	43.49 $\pm$ 9.416	<0.05
Body mass index (kg/m ²)	24.53 $\pm$ 2.73	41.905 $\pm$ 6.08	<0.001
Waist circumference (cm)	73.81 $\pm$ 13.93	107.46 $\pm$ 12.06	<0.001
Systolic blood pressure (mmHg)	70.19 $\pm$ 3.66	106.62 $\pm$ 5.67	<0.001
Diastolic blood pressure (mmHg)	113.27 $\pm$ 1.17	134.58 $\pm$ 2.08	<0.001

Note: Values are expressed as mean $\pm$ SD.

Abbreviation: MS, metabolic syndrome.

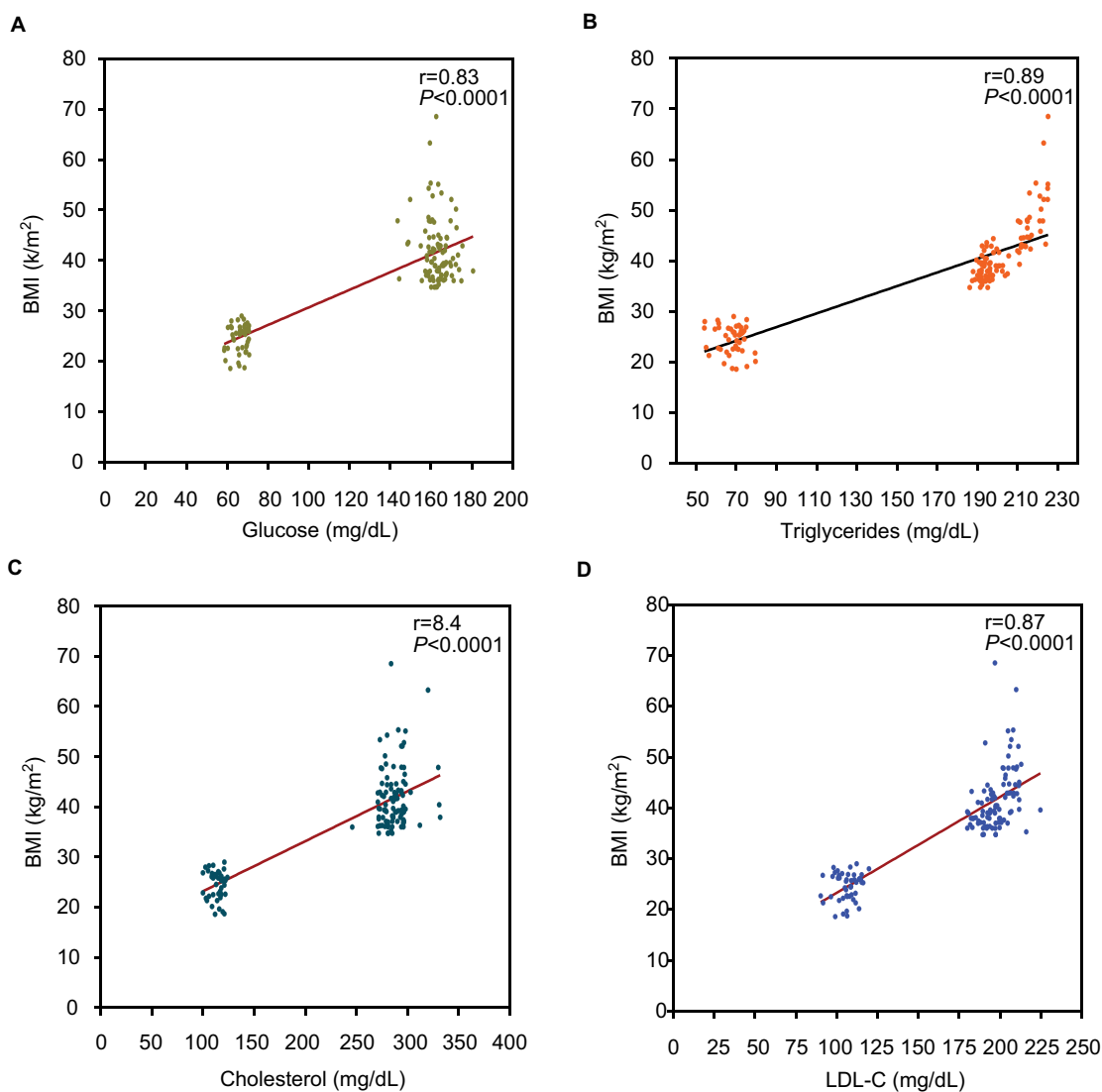


Figure 2 Correlation between glucose (A), triglycerides (B), cholesterol (C), LDL-C (D), and BMI among the whole study group.

Abbreviations: LDL-C, low-density lipoprotein cholesterol; BMI, body mass index.

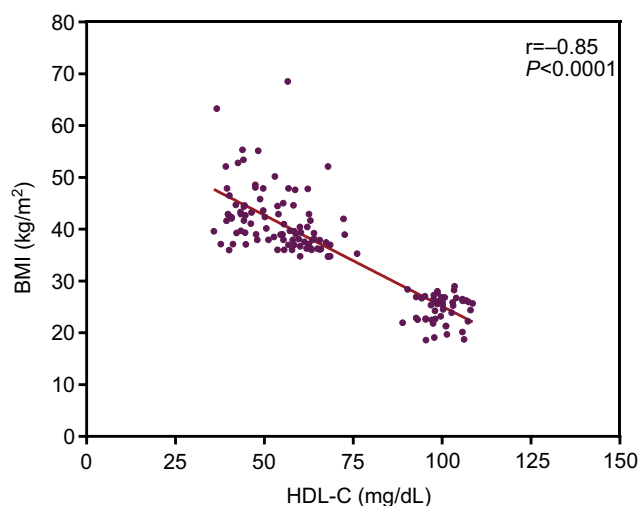


Figure 3 Correlation between HDL-C and BMI among the whole study group.

Abbreviations: HDL-C, high-density lipoprotein cholesterol; BMI, body mass index.

Figure 4A–D, reflecting a strong positive correlation between WC and blood glucose, serum TGs, cholesterol, and LDL. A negative correlation was reported between WC and serum HDL as shown in Figure 5.

The levels of testosterone were also found to be higher in women with MS than in women without MS as shown in Figure 6.

Discussion

This study aimed to detect the relationship between obesity, BMI, and WC as determinants of MS among Saudi women. The age of the study group ranged between 20 and 60 years. The main findings of the current study showed a strong significant relationship between BMI and WC with MS. The mean age was higher in women with MS compared to controls (43 and 39 years, respectively).

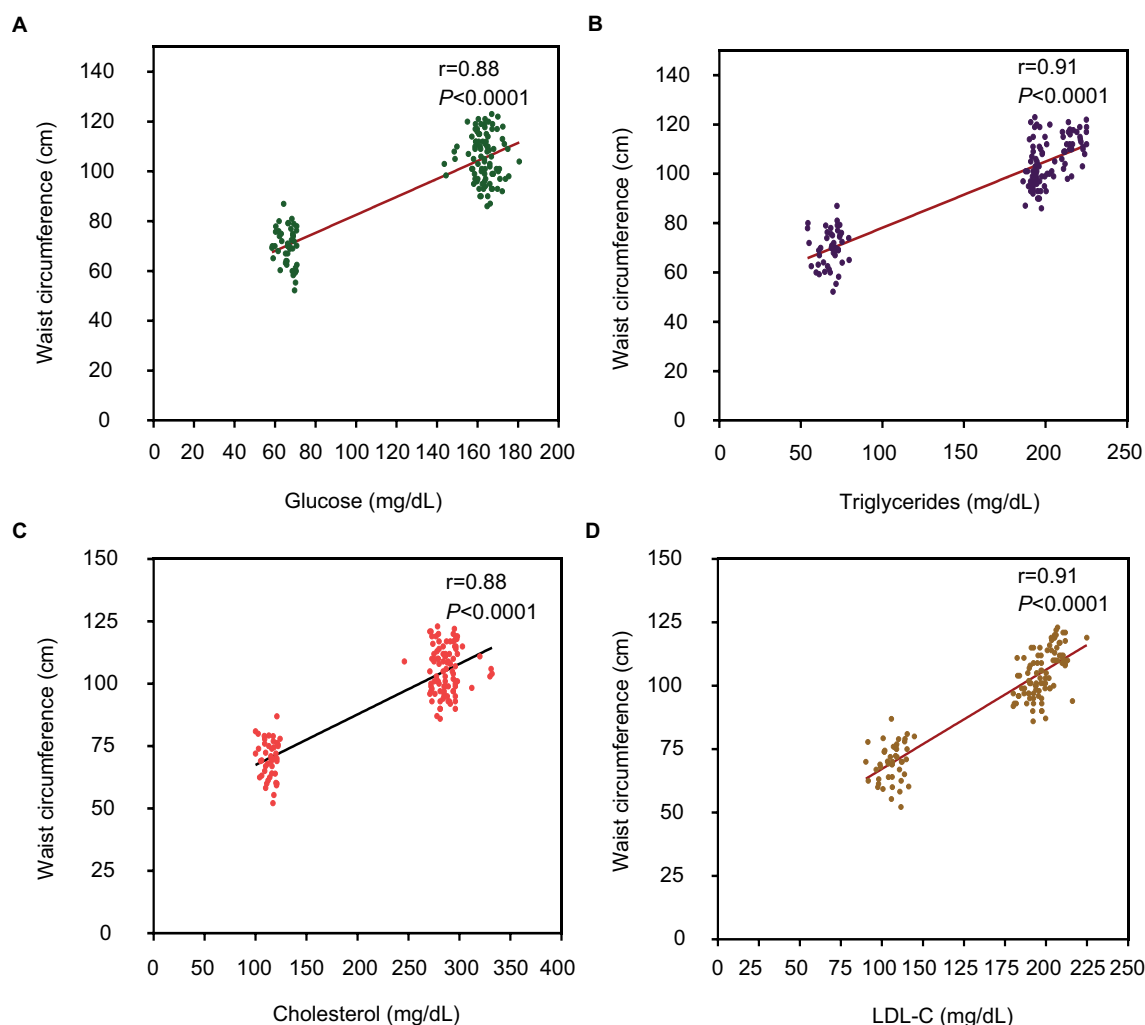


Figure 4 Correlation between glucose, cholesterol, triglycerides, LDL-C, and waist circumference among the whole study group. (A) Fasting blood sugar levels and BMI; (B) triglycerides and BMI; (C) cholesterol and BMI; (D) LDL-C and BMI.

Abbreviation: LDL-C, low-density lipoprotein cholesterol.

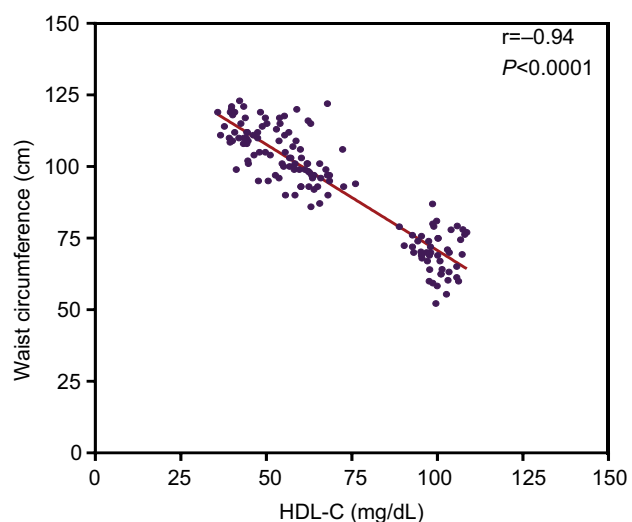


Figure 5 Correlation between HDL-C and waist circumference among the whole study group.

Abbreviation: HDL-C, high-density lipoprotein cholesterol.

The fact that an increase in TG levels is observed along with the increment of the WC and BMI is similar to the findings in a previous study.²⁰ The study conducted by Du et al²¹ has demonstrated that the HDL-C level is affected by weight. Furthermore, in this study, the average of the HDL-C levels in normal-weight and obese women with MS is higher in comparison with our results.²¹ Another finding of our study that should be emphasized is the fact that the HDL-C level decreases along with the increase of total testosterone concentration. There was a significant change in the LDL-C level parallel to the increment of WC, BMI, and testosterone level.

With regard to the effect on metabolic abnormal values which were in our study for TC, HDL-C, and LDL-C, similar strong association was found between their abnormal levels and abnormal total testosterone concentration, for high TGs

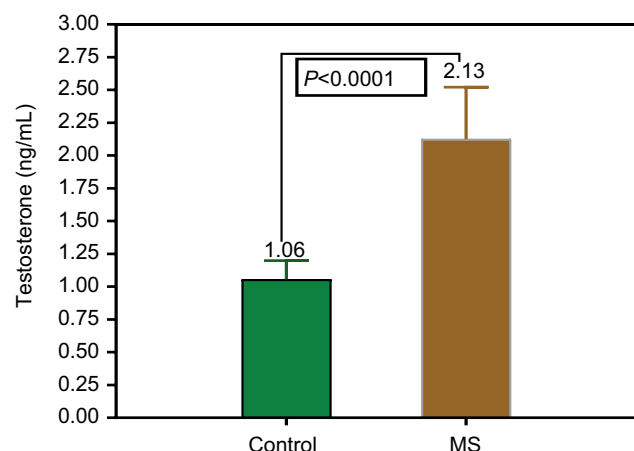


Figure 6 The levels of testosterone in controls and women with MS.
Abbreviation: MS, metabolic syndrome.

($P=0.0001$), TC ($P=0.0001$), LDL-C ($P=0.0001$), and low levels of HDL ($P=0.0001$).

With regard to hormonal effects, significant difference between the fasting blood levels of testosterone ($P=0.0001$) was found between the two groups. Higher levels of testosterone in women with MS was suggested in previous studies.¹⁸ Excess of androgen could potentially contribute to the pathogenesis of insulin resistance and MS in women with polycystic ovary syndrome.¹⁸ The data in the present study show that women with MS have twofold increase in testosterone levels. The causal relationship needs more investigation and larger studies.

Recently, obesity is reaching epidemic proportions with worldwide figures estimated at 1.4 billion and rising every year.²² Obesity affects all socioeconomic backgrounds and different ethnicities and is considered as a prerequisite for MS. MS is a clustering of risk factors that increase the risk of type 2 diabetes mellitus and CVDs. The analysis of the distribution of MS suggested that what follows under the common definition of this pathologic condition is not a unique entity, it is rather a constellation of cluster of components that may differ across different countries.²³ As these pathologic conditions are among the leading causes of deaths worldwide, it is of critical importance to continuously evaluate the factors associated with increased prevalence of MS especially in communities of modern and sedentary lifestyles. Further studies are recommended to validate the risk factors and biochemical parameters associated with the incidence of MS.

Conclusion

In summary, the data of this study showed a significant positive correlation between BMI, WC, and disturbance in lipid

profile and MS, and a high total testosterone concentrations was found in Saudi women with MS.

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Author contributions

SHF contributed to design of the study, interpretation of biochemical laboratory investigations, and editing, styling, writing, and revising of the main manuscript text. HSA and MMM were involved in design of the study, interpretation of biochemical laboratory investigations, preparation of tables and figures, and editing, styling, writing, and revising of the main manuscript text. NAA and ATB contributed to biochemical laboratory investigations, statistical analysis, preparation of tables and figures, and editing, styling, writing, and revising of the main manuscript text. All the authors read and approved the final manuscript.

Disclosure

The authors report no conflicts of interest in this work.

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