Circulating Markers of Adipocytokines levels and BMI in Insulin Resistance Females with Type2 Diabetes

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Obesity is one of the main factors that lead to development of type 2 diabetes mellitus (T2DM), with the prevalence of both increasing worldwide. Obesity is a modifiable risk factor related with many complications and comorbidities. This study aims to evaluate the level of Resistine, Fasting Blood Glucose and Adiponectin compare with the BMI in Insulin Resistance females with Type2 diabetes. A descriptive cross-sectional - case-control study, the target population is a total 205 Sudanese females were recruited to participate in this study, aged 35-64 years, venous blood samples were collected. Plasma levels of blood glucose were measured using the particle-enhanced immunoturbidimetric assay method Cobas C-311®. While Adiponectin and Resistine estimated by ELISA Kits. Anthropometric measurements, including height and weight, were taken using standard protocols. BMI was calculated as weight (in kilograms) divided by height (in meters squared). Observed from these results significant difference between the means of fasting blood glucose (FBG), adiponectin, esistine and BMI compared with different group classified according to WHO body mass index (BMI) Classification. FBG in underweight (125 ± 15.9), FBG in healthy weight (127 ± 16.7),FBG in overweight (153 ± 19.2), FBG in obesity (169 ± 20.7), FBG in severe obesity(189 ± 23.7), P = 0.037, adiponectin in underweight (12.3 ± 1.1), adiponectin in healthy weight (11.2 ± 1.9), adiponectin in overweight (9.7 ± 1.6), adiponectin in obesity (6.1 ± 1.5), adiponectin in severe obesity (6.8 ± 1.9), P = 0.043, resistine in underweight (11.6 ± 1.1), resistine in healthy weight (12.8 ± 1.1), resistine in overweight (14.2 ± 2.7) resistine in obesity (18.2 ± 2.9), resistine in severe obesity (19.1 ± 3.4), P = 0.021, BMI in underweight(16.5 ± 1.9), BMI in healthy weight (18.3 ± 1.7),BMI in overweight (26.1 ± 2.2),BMI in obesity (30.4 ± 3.7),BMI in severe obesity (40.1 ± 4.0), P = 0.006. In this study, the results show strong negative Correlation between the levels of serum Adiponectin and Fasting Blood Glucose. (P value = 0.013, r = -7.9). Adiponectin and Resistine. (P value = 0.019, r = -6.6). In the other side strong positive Correlation between the levels of serum Resistine and Fasting Blood Glucose. (P value = 0.015, r = 6.0). Observed from this results negative Relationship between the mean of BMI compared with the history of the patient/ years. This study concluded that BMI effected on Adipocytokines levels and Increased risk of insulin resistance in females with Type2 diabetes by increased the level of resistine, fasting blood glucose and reduced of adiponectin in different groups classified according to WHO body mass index (BMI) Classification.

Keywords: Adipocytokines; Adiponectin; BMI; Insulin Resistance; Resistine.

Obesity has emerged as a public health crisis in many populations including Sudanese. Adipose tissue produces several adipokines, one of them is adiponectin which has attracted much attention because of its antidiabetic and antiatherogenic effects. Adipose tissue is a key

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endocrine organ that communicates with brain, muscle, liver, and pancreas, thereby maintaining energy homeostasis. The communication between adipose tissue and other organs is mainly mediated by multiple endocrine substances secreted by adipose tissue, referred to as “adipocytokines". Changes in the levels of adipocytokines are suspected to be indicators of dysfunction in adipose tissue\(^2,3\). Adipocytokines\(^1\) and resistine\(^3\) hormones are thought to link with obesity and MS with cardiovascular risk. Adipose tissue is no longer considered an inactive organ, which only stores lipids and serves as an energy reservoir. These chemical messengers, known as ‘adipocytokines’ or ‘adipokines’, include tumor necrosis factor á (TNF-á), adiponectin, leptin, resistine and virstatin\(^6\). Adiponectin is an adipocyte-secreted polypeptide hormone with molecular weight 30 kDa (244 amino acids) which modulates several metabolic processes, and regulates insulin sensitivity and energy homeostasis, as well as glucose and lipid metabolism\(^7\). The hormone plays a principal role in the suppression of the metabolic derangements that may result in insulin resistance, obesity, MS, and cardiovascular disease\(^5,7,8\). Adiponectin is a protective protein with antidiabetic, anti-inflammatory and anti-atherogenic effects\(^6\). Reduced plasma adiponectin levels have been reported in obese individuals, particularly in those with visceral obesity, and have been negatively correlated with insulin resistance. Recent evidence has also suggested the role of adiponectin in the regulation of insulin action, energy homeostasis, obesity, and insulin resistance. Circulating adiponectin levels and adiponectin gene expression in adipose tissue are reduced in patients with type 2 diabetes\(^9-10\). Available data suggest that adiponectin might reduce hepatic glucose production and increase muscle glucose utilization, perhaps by increasing fat oxidation and thereby reducing circulating NEFA levels and intramyocellular accumulation\(^11\). Studies of adipocytokines in populations with different propensity for obesity, insulin resistance, type 2 diabetes and atherosclerosis are needed, and are interesting in this respect, because they have very high incidence of insulin resistance, central obesity, type 2 diabetes, and cardiovascular disease\(^12-13\). In a previous study by Valsmakis et al\(^14\), South-Asians have lower adiponectin levels compared to Caucasians. It was not clear however, if this feature of the South-Asians is related to aspects of glucose metabolism. No published data have been found related to adipocytokines, resistine and obesity in Sudanese population. This study aims to determine the relationship between body fat distribution and adipocytokines.

**Table 1. Number of diabetic female’s patients according to WHO body mass index (BMI) Classification**

<table>
<thead>
<tr>
<th>Categories Based on BMI</th>
<th>Body Mass Index (BMI-Kg/m(^2))</th>
<th>Number of Diabetic Female</th>
<th>% Of Diabetic Female</th>
</tr>
</thead>
<tbody>
<tr>
<td>Under weight</td>
<td>&lt; 18.5</td>
<td>9</td>
<td>4%</td>
</tr>
<tr>
<td>Healthy weight</td>
<td>18.5 – 24.9</td>
<td>53</td>
<td>25%</td>
</tr>
<tr>
<td>Overweight</td>
<td>25 – 29.9</td>
<td>107</td>
<td>52%</td>
</tr>
<tr>
<td>Obesity</td>
<td>30 – 39.9</td>
<td>31</td>
<td>17%</td>
</tr>
<tr>
<td>Sever obesity</td>
<td>&gt; 40</td>
<td>5</td>
<td>2%</td>
</tr>
</tbody>
</table>

**Table 2. Sociodemographic of diabetic female’s patients’ participants**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Age / Year Range (mean ± SD)</th>
<th>Body Mass Index -Kg/m(^2) Range (mean ± SD)</th>
<th>Duration of diabetic / years Range (mean ± SD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Under weight</td>
<td>38– 60 (48.7 ± 5.8)</td>
<td>17.1 – 18.3 (16.5 ± 1.9)</td>
<td>8 – 11 (9.8 ± 1.7)</td>
</tr>
<tr>
<td>Healthy weight</td>
<td>40 – 62 (49.6 ± 6.2)</td>
<td>19 – 24.9 (18.3 ± 1.7)</td>
<td>10 – 15 (13.6 ± 3.5)</td>
</tr>
<tr>
<td>Overweight</td>
<td>35 – 61 (51 ± 6.6)</td>
<td>25.2 – 28.8 (26.1 ± 2.2)</td>
<td>11 – 14 (11.1 ± 2.4)</td>
</tr>
<tr>
<td>Obesity</td>
<td>43 – 62 (50 ± 6.4)</td>
<td>30.1 – 33.7 (30.4 ± 3.7)</td>
<td>7 – 13 (12.0 ± 2.9)</td>
</tr>
<tr>
<td>Sever obesity</td>
<td>40– 64 (49.4 ± 6.1)</td>
<td>40.7 – 40.9 (40.1 ± 4.0)</td>
<td>6 – 11 (10.1 ± 1.8)</td>
</tr>
</tbody>
</table>
MATERIALS AND METHODS

In this is a descriptive cross-sectional and analytical case-control study. The study was done in Khartoum state. The target population is a total 205 Sudanese females were recruited to participate in this study, classified according to WHO body mass index (BMI) Classification, aged 20-50 years. The sample size calculations were based on the formula for unmatched case-control studies. Open EPI-INFO statistical package version 7 was used with 99 % two-sided confidence level, 80% power and 40 % of controls exposed, 57 % of cases exposed and odds ratio > 2. The ratio of controls to cases is 1:1. Venous blood sample (5 ml) was drawn by a well-trained medical technologist into vacutainer tubes, blood collected in anticoagulant fluoride oxalate and then plasma samples were obtained by centrifugation at room temperature at 3000 rpm/10 minutes. BMI was obtained by anthropometric measurements, including height and weight, were taken using standard protocols. BMI was calculated as weight (in kilograms) divided by height (in meters squared).

Estimation of Fasting Blood Glucose

Plasma levels of blood glucose were measured using the particle-enhanced immunoturbidimetric assay method Cobas C-311®. Human glucose agglutinates with latex particles coated with monoclonal anti-glucose antibodies, and then, the precipitate was determined turbidimetrically.

Estimation Adiponectin and Resistine

For Adiponectin and Resistin used ELISA Kits-(ab222508)/ (ab222403) is a single-wash 90 min sandwich ELISA designed for the quantitative measurement of Adiponectin and Resistine in plasma. Simple step ELISA® technology employs capture antibodies conjugated to an affinity tag that is recognized by the monoclonal antibody used to coat our Simple step ELISA® plates. (15)

Statistical analysis

Finally, the result analyzed by SPSS version 24. The mean and SD were obtained, and one way ANOVA are used for comparison. Linear regression was also use for correlation. P. value was obtained to assess the significance of the results (P value of < 0.05 was significant).
RESULTS

The study done on 105 Females Diabetic Patient Type2 with different body mass index BMI (Underweight, Healthy weight Overweight, Obesity, Sever obesity) table (1,2). Used statistical analysis test one away ANOVA to determine the mean difference between different groups (Underweight, Healthy weight Overweight, Obesity, Sever obesity), observed from this means significant difference between the means of fasting blood glucose (FBG), adiponectin, resistine and

Figure 1. Shows strong negative Correlation between the levels of serum Adiponectin and Fasting Blood Glucose. (P.value = 0.013, r = -7.9)

Figure 2. Shows moderate positive Correlation between the levels of serum Resistine and Fasting Blood Glucose. (P.valu = 0.015, r = 6.0)
BMI compare with underweight, healthy weight, overweight, obesity and sever obesity respectively. FBG in underweight (125 ± 15.9), FBG in healthy weight (127 ± 16.7), FBG in overweight (153 ± 19.2), FBG in obesity (169 ± 20.7), FBG in sever obesity (189 ± 23.7), P = 0.037, adiponectin in underweight (12.3 ± 1.1), adiponectin in healthy weight (11.2 ± 1.9), adiponectin in overweight (9.7 ± 1.6), adiponectin in obesity (6.1 ± 1.5), adiponectin in sever obesity (6.8 ± 1.9), P = 0.043, resistine

**Figure 3.** Shows strong negative Correlation between the levels of serum Resistine and Adiponectin. (P.value = 0.019, r = -6.6)

**Figure 4.** Shows negative Relationship between the mean of BMI compare with History of the patients
in underweight (11.6 ± 1.9), resistine in healthy weight (12.8 ± 1.1), resistine in overweight (14.2 ± 2.7), resistine in obesity (18.2 ± 2.9), resistine in sever obesity (19.1 ± 3.4), P = 0.021), BMI in underweight (16.5 ± 1.9), BMI in healthy weight (18.3 ± 1.7), BMI in overweight (26.1 ± 2.2), BMI in obesity (30.4 ± 3.7), BMI in sever obesity (40.1 ± 4.0), P = 0.006) Table (3). Person T test was used for correlation, the statistical analysis shows strong negative Correlation between the levels of serum Adiponectin versus Fasting Blood Glucose and Resistine. (Adiponectin versus FBG: P.value = 0.013, r = -7.9). (Adiponectin versus Resistine: P.value = 0.019, r = -6.6) figure (1) and figure (3).

In the other side of hand shows moderate positive Correlation between the levels of serum Resistine and Fasting Blood Glucose. (P.value = 0.015, r = 6.0) figure (2). Observed from this results negative Relationship between the mean of BMI compare with the history of the patient/ year’s figure (4).

**DISCUSSION**

Obesity is a modifiable risk factor in the development of type 2 diabetes mellitus (T2DM), with the prevalence of both increasing worldwide and the risks of many complications and comorbidities (e.g., cardiovascular disease and chronic kidney disease) are considerably increased in patients with T2DM who have concomitant obesity. Through this study, we have shown that circulating adipocytokine concentrations differed based on the degree of obesity in females’ patients with diagnosed T2DM, we demonstrated that higher plasma concentrations of FBG, Resistine and lower plasma concentrations of adiponectin were present in patients diagnosed T2DM who were overweight, obese and sever obesity than in patients diagnosed T2DM who had underweight and healthy weight. According to results of this study observed significant difference in resistine concentration was identified between patients with T2DM who had underweight, healthy weight and those who were overweight, obese and sever obesity. Foula WH et al found that resistine concentrations were higher in obese females patients with diabetes than non-obese females participants; moreover, they were higher in obese patients with diabetes than in non-obese healthy controls. Observed from this study significant difference in adiponectin concentration was identified between patients with T2DM who had underweight, healthy weight and those who were overweight, obese and sever obesity. Serum adiponectin concentrations have been shown to be inversely correlated with the severity of insulin resistance in patients with T2DM consistent with the findings of previous studies the present study showed that a lower level of adiponectin was present in patients with diagnosed T2DM who were obese than in those who had underweight and healthy weight BMI. Adiponectin is considered to have anti-diabetic and anti-inflammatory effects; therefore, it is reasonable to presume that patients with T2DM who obese exhibit more severe insulin resistance status are than patients with T2DM who have underweight and healthy weight BMI. Some of the studies have suggested that adiponectin/resistine ratios are more closely related to the severity of insulin resistance. It became clear through this study positive correlation between the levels of serum resistine and Fasting Blood Glucose. While on the other side of the study it was observed negatively correlation with these same parameters. In conclusion, serum resistin and adiponectin levels are correlated with the occurrence of insulin Resistance females T2DM.

**CONCLUSION**

Our findings suggest that, in patients diagnosed T2DM, adipocytokine concentrations (resistine, and adiponectin) differed between patients who had normal BMI and those who were obese. This study concluded that BMI effected on Adipocytokines levels and Increased risk of insulin resistance in females with Type2 diabetes by increased the level of Resistine, Fasting Blood Glucose and reduced of Adiponectin in different groups of BMI.

**ACKNOWLEDGMENTS**

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REFERENCES


