Predictive and Prognostic Value of Thyroid Profile and Lipid profile in Pregnancy Induced Hypertension

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Abstract

Objectives: To evaluate and compare thyroid profile and lipid profile in normotensive pregnant females at 20 weeks or more gestation and in cases of PIH at same gestation.

Methods: The study was conducted on 100 pregnant females, divided into two groups of 50 each: cases of PIH (study group) and normotensive pregnant females (control group) and all these pregnant females were at 20 weeks or more gestation. Blood sample was collected from these patients at the period of gestation when they attended Gynaecology O.P.D at Rajindra Hospital, Patiala and Thyroid profile and Lipid profile (S. Cholesterol and Triglycerides) was evaluated and compared in these patients.

Results: The mean T3 value in the study group was 1.18 ng/ml while it was 1.16 ng/ml in the control group (p=0.8156) and mean T4 value in the study group was 8.19 g/dl compared to 7.69 g/dl in the control group (p=0.2681). The difference did not attain statistical significance. The mean TSH value in the study group was 2.73 mlU/L which was significantly higher statistically than the value of 1.27 mlU/L in the control group (p<0.0001). Lipid profile levels (Serum cholesterol and triglycerides) were significantly higher in the study group compared to the control group (p<0.0001). The mean serum cholesterol value in the study group was 254.64 mg/dl compared to 172.64 mg/dl in the control group. The mean triglycerides value in the study group was 235.86 mg/dl compared to 133.20 mg/dl in the control group.

Conclusions: It was concluded that patients with PIH had a significant increase in TSH levels, a non significant alteration in T3 and T4 levels and a significant increase in lipid profile levels (S.cholesterol and triglycerides) compared to normotensive pregnant females. This suggests role of thyroid hormones and dyslipidemia in the development and pathogenesis of PIH. Therefore, early detection of thyroid abnormalities and dyslipidemia may affect the occurrence and severity of PIH and help in better management of the disease in established pre-eclamptic women.

Keywords: Pregnancy induced hypertension; Thyroid profile; Lipid profile

Introduction

A mother becomes almost a new person during the period of pregnancy. Profound local and systemic changes in maternal physiology are initiated by conception and continued throughout pregnancy [1].

Pre-eclampsia is a multisystem disease unique to human pregnancy characterised by hypertension and organ system derangement. The disease is responsible for considerable morbidity and mortality complicating 5-8% of pregnancies and remains in the top three causes of maternal morbidity and mortality globally. It is the leading cause of fetal growth restriction, intrauterine fetal demise and planned preterm birth [2]. Pre-eclampsia usually occurs after 20 weeks gestation and is classically defined as a triad of hypertension, edema and proteinuria [3].

Pregnancy is usually associated with mild hyperthyroxinemia, but pre-eclamptic women have a high incidence of hypothyroidism that might correlate with the severity of pre-eclampsia [4,5]. In pre-eclampsia, there is decreased estrogen production due to placental dysfunction resulting in lowering of Thyroid binding globulin, total T3 and total T4 along with growth retardation of the fetus [5].

Abnormal lipid metabolism seems important in the pathogenesis of pregnancy induced hypertension (PIH). Endothelial dysfunction is the most important event in the pathogenesis of pre-eclampsia and lipids have a role on this event. Increased TG, found in PIH, is likely to be deposited in predisposed vessels, such as the uterine spiral arteries and contributes to the endothelial dysfunction [6]. Altered lipid synthesis leading to decrease in PGI2:TXA2 ratio is also supposed to be an important way of pathogenesis in PIH [7].

So in the background of altered thyroid and lipid profile being potential risk factors for occurrence of PIH, the present study aims to evaluate and compare the thyroid and lipid (cholesterol and triglycerides) profiles in PIH and normal pregnancy so that early identification of changes in these parameters might be of help in preventing the occurrence of PIH.

Material and Methods

The present study was conducted in the Department of Biochemistry, GMC Patiala on 100 pregnant females referred by the Department of Obstetrics and Gynaecology, Rajindra Hospital, Patiala. The ethical committee of GMC, Patiala approved the study protocol. The objectives of the study were explained and written consent was taken from individual subjects. The subjects selected for the present

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study were attending and admitted to Rajindra Hospital, Patiala and were at 20 weeks or more gestation.

**Study group**

50 pregnant females with pregnancy induced hypertension at 20 weeks or more gestation. The diagnosis of Pre-eclampsia was based on the definition of National High Blood Pressure Education Program Working Group, B.P ≥ 140/90 mmHg after 20 weeks gestation and proteinuria ≥ 300 mg/24 hours or ≥ 1+ dipstick.

**Control group**

50 pregnant females at same gestation with normal blood pressure, no proteinuria and without any other systemic or endocrine disorder.

**Exclusion criteria**

All cases of previous history of essential hypertension or chronic hypertension, known cases of thyroid disorders and associated molar pregnancy and multiple pregnancies were excluded from the study.

**Specimen collection**

Blood sample was collected from these patients at the gestational age of 20 weeks or more, when they attended O.P.D at Rajindra Hospital, Patiala. Serum: Under all aseptic conditions, 5 ml blood sample was collected by venipuncture and allowed to clot. The serum was then separated by ultracentrifugation of the sample at room temperature and supernatant (serum) was taken in a separate test tube. The serum was then used for Thyroid and lipid profile analysis in the laboratory.

**Methodology**


**Statistical analysis**

It was done by student's t-test using SPSS version 10.0

**Results**

Table 1 shows demographic characteristics of both the study and control groups. The mean age and period of gestation were comparable in both the groups. The mean BMI was increased in both the groups. The mean SBP and DBP were significantly higher in the study group compared to the control group.

The mean T₄ value in the study group was 1.18 ng/ml while it was 1.16 ng/ml in the control group (p=0.8156) and mean T₃ value in the study group was 8.19 g/dl compared to 7.69 g/dl in the control group (p=0.2681). The difference did not attain statistical significance. The mean TSH value in the study group was 2.73 mIU/L which was significantly higher statistically than the value of 1.27 mIU/L in the control group (p<0.0001) (Table 2) (Figure 1).

Lipid profile levels (Serum cholesterol and triglycerides) were significantly higher in the study group compared to the control group (p<0.0001). The mean serum cholesterol value in the study group was 254.64 mg/dl compared to 172.64 mg/dl in the control group. The mean triglycerides value in the study group was 325.86 mg/dl compared to 133.20 mg/dl in the control group (Table 3) (Figure 2).

**Table 1**: Demographic characteristics of study group and control group.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Study group (n=50)</th>
<th>Control group (n=50)</th>
<th>T</th>
<th>P</th>
<th>S</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean Age (in years)</td>
<td>25.12 ± 3.95</td>
<td>24.14 ± 3.80</td>
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<tr>
<td>Mean Period of gestation (in weeks)</td>
<td>32 ± 3.68</td>
<td>31 ± 3.70</td>
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<tr>
<td>Body Mass Index (kg/m²)</td>
<td>29 ± 5.7</td>
<td>27 ± 3.9</td>
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<td>Mean S.B.P (mmHg)</td>
<td>155.68 ± 14.68</td>
<td>115.68 ± 7.08</td>
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<tr>
<td>Mean D.B.P(mmHg)</td>
<td>105.48 ± 11.02</td>
<td>73.96 ± 6.25</td>
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<tr>
<td>Fasting Plasma Glucose (mg/dl)</td>
<td>81.96 ± 10.59</td>
<td>79.90 ± 8.24</td>
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</tbody>
</table>

**Table 2**: Comparison of mean thyroid profile (T₃, T₄ and TSH) values in study and control group.

**Table 3**: Comparison of lipid profile in study group and control group.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Study group (n=50)</th>
<th>Control group(n=50)</th>
<th>T</th>
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<th>S</th>
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</thead>
<tbody>
<tr>
<td>S.Cholesterol (mg/dl)</td>
<td>254.64 ± 50.56</td>
<td>172.64 ± 39.15</td>
<td>9.0661</td>
<td>&lt;0.0001</td>
<td>Highly Significant</td>
</tr>
<tr>
<td>S.Triglycerides (mg/dl)</td>
<td>235.86 ± 64.90</td>
<td>133.20 ± 48.58</td>
<td>8.9537</td>
<td>&lt;0.0001</td>
<td>Highly Significant</td>
</tr>
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</table>
Discussion

Between 5% and 15% of pregnant women experience thyroid abnormalities, a fact which justifies screening by means of clinical laboratory testing. Maternal thyroid dysfunction during pregnancy has been shown to be associated with a number of adverse outcomes. It has been suggested that there may be an existence of mutual influences between pre-eclampsia and thyroid function [12,13].

The two groups were comparable in terms of fasting plasma glucose and this rule out the influence of systemic illness like Diabetes Mellitus on thyroid gland and on the lipid profile of the patients. In the present study, though T₃ and T₄ level were higher in the study group, the values did not attain statistical significance while the mean TSH level was significantly higher than the control group. Many researchers have shown that women with pre-eclampsia had high level of TSH [14]. The finding of increased TSH levels in pre-eclamptic cases can be explained by excessive release of antiangiogenic proteins-most notably sFlt-1 (soluble Fms-like tyrosine kinase-1) from the placenta into maternal blood, resulting in an antiangiogenic state with low levels of free placentat growth factor and free vascular endothelial growth factor which leads to endothelial dysfunction [15].

The explanation of non-significantly increased T₃ and T₄ levels in the present study in PIH cases may be attributed to pathological changes in pre-eclampsia which leads to endocrinological disturbances and consequently to thyroid gland dysfunction that may affect synthesis and secretion of T₃ and T₄. Other studies have shown decreased T₃ associated with higher T₄ in pre-eclampsia attributed to reduced conversion of T₄ to T₃ in the liver and kidneys [16].

Disorders of lipoprotein metabolism are a major cause of endothelial dysfunction that may result in hypertension and proteinuria, the clinical hall-marks of pre-eclampsia [17].

In the present study, the mean serum cholesterol level was significantly higher in the study group compared to the control group. Physiological insulin resistance is exaggerated in pre-eclampsia. Gestational insulin resistance may accentuate the suppression of lipoprotein lipase activity and increase mobilization of free fatty acids from visceral adipocytes. This may explain the hypercholesterolemia in pre-eclampsia [18,19].

The mean triglycerides level was significantly higher in the study group compared to the control group. Hypertriglyceridaemia may be modulated by hyperinsulinemia found in pregnancy [20]. In pregnancies complicated by hypertension, there appears to be an exaggeration of insulin resistance and associated metabolic changes. Pre-eclamptic women are more insulin resistant than normotensive controls, so in pre-eclampsia triglyceride levels further increase due to the exaggeration of insulin resistance [21]. Also our findings of greater increase in mean BMI in pregnant females with PIH could explain the significant increase in triglycerides.

Conclusion

In the present study, an effort was made to explore the role of thyroid hormones and lipid profile in the pathogenesis of PIH and it was found that patients with PIH had altered levels of thyroid hormones and dyslipidemia. This suggests role of thyroid hormones and dyslipidemia in the development and pathogenesis of PIH. Therefore, early detection of thyroid abnormalities and dyslipidemia may affect the occurrence and severity of PIH and help in better management of the disease in established pre-eclamptic women.

Keeping in view the target for the Fifth Millennium Development Goal (MDG 5) which aims to improve maternal health and reduce maternal mortality rate, by three - quarters by 2015, and PIH being one of the top three leading causes of maternal mortality, it is recommended that each and every pregnant women should have thyroid profile and lipid profile done as soon as she reports to the antenatal clinic and then again preferably in the second trimester to detect any abnormality that might put her at risk for PIH.

References