Study of high sensitive c-reactive protein in preeclampsia

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Abstract
Introduction: Preeclampsia is a serious complication of the second half of pregnancy and is an important cause of maternal and perinatal morbidity and mortality. Measurement of circulatory inflammatory markers of endothelial cell dysfunction and inflammation may provide an alternative method of detecting women at high risk of delivery. Women with a history of pre eclampsia have an increased susceptibility to lipoprotein oxidation when compared to women who had normal pregnancy.
Aims and objectives: The present study was designed to study hsCRP in association with blood pressure, proteinuria and lipid profile as markers of inflammation and endothelial dysfunction in preeclampsia.
Materials and Methods: The present study was carried out for one year on diagnosed patients of Preeclampsia attending the outpatient and inpatient departments of obstetrics and Gynecology department of Narayana Medical College and Hospital, Nellore. The study was undertaken on 50 Preeclampsia cases and 50 normal pregnant women as controls.
Results: In our study hs-CRP levels were significantly elevated in cases when compared to control subjects. Total cholesterol, triacylglycerol and LDL values were elevated in total cases when compared to controls. However the decrease in HDL cholesterol was not significant when compared to controls.
Conclusion: Elevated levels of LDL undergo oxidation to form oxidized LDL, which is involved in inflammation and free radical generation. Increased inflammation leads to increased levels of hsCRP, which is further associated with dyslipidemia, an important cardiovascular risk factor in preeclampsia.

Keywords: Preeclampsia, hsCRP, Systolic Blood pressure, Dyslipidemia, urine protein creatinine ratio (P/C ratio), Pregnancy induced hypertension.

Introduction

Preeclampsia develops in 4-5% of human pregnancies and is characterised by hypertension, dyslipidemia and increased systemic inflammatory response.1 Pregnancy Induced Hypertension (PIH) is a serious complication of these half of pregnancy with urinary loss of proteins, oedema and activation of hemostatic mechanisms.2

Preeclampsia is characterized by an increased BP equal to or above 140/90 mmHg in the presence of proteinuria that develops after 20 weeks of gestational age. PE results in eclampsia if convulsion develops or manifests as hemolysis, elevated liver enzymes and low platelet count (HELLP) syndrome. Severe complication like cerebral hemorrhage, renal failure, lung edema and liver hemorrhage are associated with eclampsia and HELLP syndrome.3

Endothelial cell dysfunction and inflammation have an important role in the pathophysiological mechanism of preeclampsia. There is Increasing clinical and biochemical evidence which suggests that disturbance of normal endothelial cell function may be a primary cause in the pathogenesis of PE.4 C-reactive protein (CRP) is one of the major acute-phase reactant in humans. It is an important first-line host defence molecule as it activates the complement system and mediates the phagocytic clearance of pathogens and damaged cells.5,6

The term high sensitivity CRP (hsCRP) refers to the lower detection limit of the assay procedures being used and is otherwise similar to routine CRP in structure and function. It has been suggested that hsCRP has provided better sensitivity in establishing inflammation than levels of CRP gives an idea about ongoing inflammation and tissue damage very accurately compared to other laboratory parameters of the acute-phase response.7 The hs-CRP is a sensitive marker of tissue damage and inflammation.It is useful in differentiating acute inflammation as well as assessment of severity of inflammation.8

Mean plasma triglyceride and free fatty acid concentrations undergo near doubling in women with preeclampsia relative to normal pregnancy.9 The hypertriglycerideremia of preeclampsia is also accompanied by increased prevalence of smaller, denser LDL particles and decreased HDL cholesterol. Triacylglycerol levels are a major determinant of cholesterol ester transfer protein (CETP) activity.10 Therefore, in the presence of increased triacylglycerol-rich lipoproteins, CETP activity is increased so that all circulating lipoproteins become enriched in triacylglycerol, in particular HDL and LDL particles.11 LDL particles are triacylglycerol enriched due to the hyper triglycerideremia and increased CETP activity.
They are also converted by the triglyceride lipase activity of hepatic lipase into smaller and denser particles. Large buoyant LDL is cleared rapidly by the LDL receptor pathway and small dense LDL is removed more slowly. Triacylglycerol-rich HDL particles are converted by the triglyceride lipase activity of hepatic lipase into smaller particles which are better substrates for catabolic pathways.

One of the reasons for this dyslipidemia is due to hormonal imbalance in preeclampsia. Preeclampsia is a state of hypoestrogenemia. There is impairment in the formation of Dehydroepiandrosterone (DHEA) by fetal adrenal glands due to decreased uteroplacental blood flow there by leading to decreased uptake of lipids by the fetus. DHEA is the important source of estrogen in pregnancy. The decrease in utero-placental blood flow leads to relative stasis of maternal blood in the intervillous space which leads to redistribution of steroids formed in syncytium back to the fetus instead of entering into maternal circulation which also results in a state of hypoestrogenemia.

Oxidative stress has been implicated in the pathogenesis of several complication of human pregnancy including preeclampsia. In preeclampsia dyslipidemia pattern of increased concentration of triglyceride, cholesterol and LDL and decreased concentration of HDL have been noticed. Preeclampsia is associated with elevated lipid peroxidation and reduced antioxidant status. Preeclampsia is an endothelial disease with a major involvement of lipid mediated oxidative damage.

The present study was designed to establish an association among blood pressure, proteinuria, hsCRP and lipid profile as markers of inflammation and endothelial dysfunction in pregnancy induced hypertension.

Materials and Methods

The present study was carried out for one year on diagnosed patients of preeclampsia attending the outpatient and inpatient departments of Obstetrics and Gynecology department of Narayana Medical College and Hospital, Nellore. The study was undertaken on 50 preeclampsia cases and 50 normal pregnant women.

Inclusion criteria: 50 normal healthy pregnant women of > 20 weeks of gestational age were selected as controls. 50 diagnosed preeclampsia cases > 20 weeks of gestational age and with urine protein creatinine ratio (P/C ratio) more than 0.3 were selected as cases for the study.

Exclusion criteria: Pregnant women with diabetes mellitus, anemia and other chronic illnesses were excluded from the study. Women with the following conditions were also excluded: a known kidney disease, heavy exercise (more than 1 hour of vigorous exercise on the day of urine collection), bacteriuria, bed rest longer than 24 hours, and gestational diabetes mellitus.

In addition, women who delivered their babies during the urine collection day were excluded.

6ml venous blood was collected from the cases and controls under aseptic precautions and it was transferred to the Plain tube and serum was separated by centrifugation and stored at -20°C to measure the analytes. Haemolysed and lipemic samples were avoided. 5ml of the collected urine sample was centrifuged, supernatant separated and stored at -20°C to measure the analytes.

hsCRP was determined by turbidimetric immunoassay using commercially available (Euro diagnostics) kit method. Total cholesterol, triacylglycerol, HDL, LDL, urine microprotein and urine creatinine were estimated by commercially available kits in a semi auto analyser.

The mean and standard deviation were calculated for all the Biochemical parameters. The significance between the groups were determined using Student t-test for Equality of means. The p-value of < 0.05 was considered significant.

Results

In the present study the mean ± SD of SBP in controls (n=50) and pre-eclamptic cases (n=50) were 110.8 ± 8.0 and 159.2 ± 16.87 mm of Hg respectively. This increase was statistically significant (p<0.0001) as shown in Table 1.

The mean ± SD of DBP in controls (n=50) and pre-eclamptic cases (n=50) were 73.6 ± 5.6 and 103.56 ± 8.95 mm of Hg respectively. This increase was statistically significant (p<0.0001) as shown in (Table 1, Fig. 1). The mean ± SD of urine P/C ratio in controls (n=50) and pre-eclamptic cases (n=50) were 0.13 ± 0.02 and 0.64 ± 0.26 respectively. This increase was statistically significant (p<0.0001) as shown in Table 1.

**Fig. 1: Comparison of mean SBP and DBP levels in controls and PE cases**
Study of high sensitive c-reactive protein in preeclampsia

In our present study total cholesterol, triacylglycerol and LDL values were elevated in total cases when compared to controls (P-value<0.05, Table 1, Fig. 2). The decrease in HDL cholesterol was not significant when compared to controls (Table 1). The mean ± SD of hsCRP in controls (n=50) and pre-eclamptic cases (n=50) were 1.71 ± 0.85 and 8.07 ± 2.09 mg/L respectively. hs-CRP levels in our study were significantly elevated in cases when compared to control subjects(Table 1, P-value<0.0001, Fig. 3).

Correlation analysis of study variables was done among cases. P value less than 0.05 was considered significant. In our study hsCRP correlated positively with SBP, DBP and P/C ratio. SBP levels in cases correlated positively with P/C ratio (Figure No.4). Total cholesterol, LDL and hsCRP.

P/C ratio in our study correlated positively with Total cholesterol, triacyl glycerol, LDL and hscRP. (P-value <0.05, Table 2).

Table 1: Biochemical Parameters in Cases and Controls

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Parameter</th>
<th>Controls</th>
<th>Cases</th>
<th>P value</th>
<th>t value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>SBP mmHg</td>
<td>110.8 ± 8.0</td>
<td>159.2 ± 16.87</td>
<td>&lt;0.0001</td>
<td>18.33</td>
</tr>
<tr>
<td>2.</td>
<td>DBP mmHg</td>
<td>73.6 ± 5.62</td>
<td>103.56 ± 8.95</td>
<td>&lt;0.0001</td>
<td>20.04</td>
</tr>
<tr>
<td>3.</td>
<td>P/C ratio</td>
<td>0.13 ± 0.02</td>
<td>0.64 ± 0.26</td>
<td>&lt;0.0001</td>
<td>13.82</td>
</tr>
<tr>
<td>4.</td>
<td>T. Cholesterol mg/dl</td>
<td>156.32 ± 32.04</td>
<td>183.26 ± 40.39</td>
<td>0.0004</td>
<td>3.695</td>
</tr>
<tr>
<td>5.</td>
<td>TGL mg/dl</td>
<td>107.28 ± 37.2</td>
<td>142.38 ± 32.49</td>
<td>&lt;0.0001</td>
<td>5.02</td>
</tr>
<tr>
<td>6.</td>
<td>HDL-C mg/dl</td>
<td>46.92 ± 12.2</td>
<td>44.42 ± 10.39</td>
<td>0.27</td>
<td>1.10</td>
</tr>
<tr>
<td>7.</td>
<td>LDL-C mg/dl</td>
<td>85.6 ± 18.86</td>
<td>112.64 ± 27.24</td>
<td>&lt;0.0001</td>
<td>5.77</td>
</tr>
<tr>
<td>8.</td>
<td>hsCRP mg/L</td>
<td>1.71 ± 0.85</td>
<td>8.07 ± 2.09</td>
<td>&lt;0.0001</td>
<td>19.93</td>
</tr>
</tbody>
</table>

Table 2: Correlation analysis of study variables

<table>
<thead>
<tr>
<th>hsCRP Vs</th>
<th>SBP Vs</th>
<th>P/C ratio Vs</th>
</tr>
</thead>
<tbody>
<tr>
<td>r-value</td>
<td>p-value</td>
<td>r-value</td>
</tr>
<tr>
<td>SBP</td>
<td>0.55</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>DBP</td>
<td>0.62</td>
<td>&lt;0.05</td>
</tr>
<tr>
<td>P/C ratio</td>
<td>0.49</td>
<td>&lt;0.05</td>
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</table>

Discussion

The present study was conducted on fifty normal healthy pregnant women as controls and fifty preeclamptic women as cases. Pregnancy Induced Hypertension (PIH) is a pregnancy specific disorder that adversely affects the mother by vascular dysfunction and the fetus by intrauterine growth
Albuminuria serves as a surrogate marker of increased glomerular capillary pressure and a predictor of progressive renal injury. There might be lower eGFR when there is proteinuria and produces significant renal injury in association with hypertension, which further leads to cardiovascular and renal complications.²⁴

Conclusion

Preeclampsia is one of the leading causes of maternal mortality and morbidity. In preeclampsia there is an exaggeration of systemic inflammatory response that might induce reactive oxygen species which further induces endothelial dysfunction. This leads to clinical symptoms of hypertension and proteinuria in preeclampsia. The hs-CRP is sensitive marker of tissue damage and inflammation. The present study showed that hsCRP was elevated in preeclampsia. Dyslipidemia was associated with preeclampsia, which acts as a precursor for sub clinical atherosclerotic progression in PIH. In addition, hypertension in pregnancy is considered as a marker for the later development of cardiovascular and renal morbidity.

References


