HIGH SENSITIVE C-REACTIVE PROTEIN AS AN INDICATOR FOR PRO-INFLAMMATORY STATUS IN DIFFERENT DEGREES OF MAJOR DEPRESSION

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ABSTRACT

BACKGROUND

Studies indicate that along with many other factors, low grade inflammation may play an important contributory role in the development of major depressive psychosis. This may add up as a confounding factor for different complications related to this major psychological disorder. The role of hsCRP as a marker of low grade inflammation has been explored and analysed in the present study.

METHODS

HsCRP, Total Cholesterol (TC), Triglyceride (TG) and Fasting Blood Glucose (FBG) were measured in 49 cases of major depression against 40 age and body weight (BMI) matched control subjects. The degree of depression in the case group was assessed by the HAM-D score. Data were analysed for any differences in these parameters between the case and control groups followed by correlation and regression analysis to assess the strength of association between the study parameters and the predictive values of serum cholesterol, triglyceride, FBG and HAM-D score on hsCRP.

RESULTS

HsCRP, TC and TG were significantly higher in the case group. The correlation and regression analyses revealed that hsCRP was strongly associated (r=0.559, P <0.001) and dependent (beta=0.579, P <0.001) on the HAM-D score only.

CONCLUSION

These results not only indicate a definite association of low grade inflammation with major depressive psychosis, but also suggest a close linear relationship between this low grade inflammation and the degree of depression. The researchers propose that hsCRP is a good indicator for monitoring the pro-inflammatory status and the degree of severity of the disease process in major depressive psychosis.

KEYWORDS

HAM-D Score, hsCRP, Low Grade Inflammation, Major Depressive Psychosis.


INTRODUCTION

In terms of its prevalence and sufferings, depression is a disorder of major public health importance along with the huge economic burden for the individual and the society overall. In India, prevalence of depression in community varies from 1.7 to 7.4 per thousand population.[1] Pathogenesis of depression is not fully understood, but studies suggest that among many other precipitating factors low grade systemic inflammation plays an important contributory role in the development of depression.[2] A few studies have suggested that depression may be associated with increased production of pro-inflammatory cytokines such as IL-1, IL-6 and interferon.

C-Reactive Protein (CRP) is a commonly used marker of inflammatory diseases, when used to study the role of low grade inflammation in the future risk for disease development.[3] Elevated CRP levels have been associated with psychological distress and depression.[4-6] Depression may result in elevated CRP level, an inflammatory state may cause depression or both the depression and elevated CRP level may be due to another unmeasured disease process. Data from several large United States and European cohorts indicate that distribution of circulating high-sensitive CRP (hsCRP) level appears comparable among men and women with 50th percentile for gender being about 1.5 mg/dL.[6,7]

A large cross-sectional population studies with 5000–7000 participants reported an association between CRP levels and depression.[6-8] However, in another study this association disappeared when estimates were adjusted for confounding factors, such as chronic illness and body mass index. Similar observations were corroborated in other studies as well.[7,9]

OBJECTIVES

Conflicts exist whether there is any association between the major depression and CRP level. We hypothesized that a low grade inflammation might exist in the pathogenesis of
depression. Accordingly, the present study was designed to assess any possible association of hsCRP level that measures the low grade inflammation with major depression in our study population.

MATERIALS AND METHODS
The study was conducted in the Department of Psychiatry and Department of Biochemistry of a Tertiary Care Medical College and Hospital of Eastern India during the period of one year from January 2013 to January 2014.

Newly diagnosed patients of depression were included in the study following inclusion and exclusion criteria as mentioned below.

Inclusion Criteria
a) All cases of newly diagnosed major depressive disorder.
b) Patients of either sex.
c) Age limit between 15–55 years.
d) Patients who are willing to enrol in the study and who had given written informed consent.

Exclusion Criteria for the Case Group were the Following
a) Patients who were on any other medication that affects hsCRP level (e.g. anti-inflammatory drugs, tobacco, oral contraceptives, etc.)
b) Patients with any infection (e.g. Bacterial/viral/fungal/mycobacterial).
c) Patients with any allergic complication of infection, such as rheumatic fever.
d) Patients with any inflammatory disease such as rheumatoid arthritis, juvenile chronic arthritis, ankylosing spondylitis, psoriatic arthritis, systemic vasculitis, polymyalgia rheumatica, Reiter’s disease, Crohn’s disease, familial Mediterranean fever, etc.
e) Patients with myocardial infarction, tumour embolisation, acute pancreatitis.
f) Patients with trauma due to surgery, burns, fracture.
g) Patients with malignancies such as lymphoma, carcinoma, etc.
h) Pregnant women.

Experimental Protocol
Informed written consents were obtained from the participants. The study protocol strictly adhered to the Helsinki Declaration 1975 revised in 2000 for human studies participants. The study protocol strictly adhered to the Helsinki Declaration 1975 revised in 2000 for human studies. Informed written consents were obtained from the study population.

The data were analysed for assessing the significance of differences between the mean values of hsCRP and lipid parameters between the case and control groups with the help of both parametric student t test and non-parametric Mann-Whitney test. Any possible dependence of the hsCRP on the confounding factors was assessed by multiple regression analysis. All statistical analyses were performed by using SPSS software version 17 for Windows.

RESULTS
Data obtained were first analysed for the matching between case and control groups (Results not shown in Tables). There was no significant difference in the mean age of case group (32.24±11.00 years) and control group (33.73±9.07 years), P=0.77. Body weights (Kg) of case and control groups (52.96±7.51 and 53.25±6.9 respectively, P >0.05) and the sex distribution (34 vs 25 males and 16 vs 15 females in the case and control groups respectively, χ²=0.298, P=0.58) revealed that selected case and control subjects were properly matched for body weight and gender.

Further analysis of the data showed that although the FBG concentration showed no significant difference between the case and control groups, serum hsCRP, serum triglyceride and serum cholesterol levels exhibited significant difference between these two groups as evident from both ‘t’ test and Mann-Whitney U test (Table 1 and Table 2).

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Cases (n=50) Mean±SD</th>
<th>Controls (n=40) Mean±SD</th>
<th>Standard Error Mean</th>
<th>t value</th>
<th>P value (Level of Significance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum hsCRP (mg/dL)</td>
<td>2.03±0.93</td>
<td>0.84±0.49</td>
<td>0.13</td>
<td>7.21</td>
<td>P&lt;0.001*</td>
</tr>
<tr>
<td>Serum Total Cholesterol (mg/dL)</td>
<td>174.34±38.32</td>
<td>153.02±27.82</td>
<td>5.47</td>
<td>2.94</td>
<td>P=0.004*</td>
</tr>
<tr>
<td>Serum Triglyceride (mg/dL)</td>
<td>140.93±76.51</td>
<td>99.30±34.28</td>
<td>10.93</td>
<td>3.18</td>
<td>P=0.002*</td>
</tr>
<tr>
<td>Plasma FBG (mg/dL)</td>
<td>103.32±38.18</td>
<td>93.30±12.26</td>
<td>5.45</td>
<td>1.93</td>
<td>P=0.115</td>
</tr>
</tbody>
</table>

Table 1: Group Statistics and Test of Significance (Independent Samples ‘t’ Test) of Different Parameters Between Cases and Control Groups

Level of sig. P <0.05 at 95% CI. *P value significant at 0.05 level.
Results were based on the data analysis and found that hsCRP had a significant association with the degree of depression only (Table 3, Figure 1).

Previously, it has been reported that elevated CRP levels were associated with an increased risk of psychological distress and depression. This study, however, stems from the research question whether systemic inflammation precedes the onset of depressive symptoms or occurs as a part of somatic manifestation of the depressive phenotype. Although the CRP level is traditionally only elevated in severe inflammation, newer assays with improved sensitivity are able to measure hsCRP in apparently healthy individuals, permitting an exploration of the postulated association between subclinical systemic inflammation and risk of depression.

Inferential statistical calculation considering both parametric and non-parametric distribution of data in the present study showed that serum hsCRP level was significantly higher in newly diagnosed depressive cases than those of healthy controls (Table 1 and 2). In a recent study it was reported that elevated CRP levels were associated with an increased risk of psychological distress and depression. This association was observed in 73131 individuals in a cross sectional analyses and in prospective analyses for hospitalization with depression.[12] Although previous studies have corroborated the association of elevated CRP levels with psychological distress and depression, the results were conflicting and varied according to differential distribution of race, region or gender.[4,7] A significant positive association between the Centre for Epidemiologic Studies Depression Scale (CES-D) and hsCRP levels among African-American men with depression was found that was absent in their female counterparts.[13] However, no such gender based variations

Table 2: Nonparametric Test: Mann-Whitney U Test for Significance Between Different Study Parameters

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Cases (n=50)</th>
<th>Controls (n=40)</th>
<th>Mann-Whitney U</th>
<th>Wilcoxon W</th>
<th>Z value</th>
<th>Asymp. Sig. (2-tailed)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Serum hsCRP (mg/dL)</td>
<td>1.71</td>
<td>0.87</td>
<td>174.5</td>
<td>994.5</td>
<td>-6.644</td>
<td>P &lt; 0.001*</td>
</tr>
<tr>
<td>Serum Total Cholesterol (mg/dL)</td>
<td>170.0</td>
<td>154.5</td>
<td>652.5</td>
<td>1472.5</td>
<td>-2.704</td>
<td>P = 0.007*</td>
</tr>
<tr>
<td>Serum Triglyceride (mg/dL)</td>
<td>122</td>
<td>100</td>
<td>603.5</td>
<td>1423.5</td>
<td>-3.107</td>
<td>P = 0.002*</td>
</tr>
<tr>
<td>Plasma FBG (mg/dL)</td>
<td>96.2</td>
<td>90.4</td>
<td>750.0</td>
<td>1570.0</td>
<td>-1.898</td>
<td>P = 0.058</td>
</tr>
</tbody>
</table>

Table 3: Bivariate Correlation Analysis among Different Parameters in the Case Group (n = 50)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Pearson’s Correlation Coefficient: r</th>
<th>Level of Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>r12</td>
<td>0.559</td>
<td>P &lt; 0.001*</td>
</tr>
<tr>
<td>r13</td>
<td>0.031</td>
<td>P = 0.831</td>
</tr>
<tr>
<td>r14</td>
<td>0.024</td>
<td>P = 0.868</td>
</tr>
<tr>
<td>r15</td>
<td>0.174</td>
<td>P = 0.233</td>
</tr>
</tbody>
</table>

Based on this finding, multiple linear regression was performed considering hsCRP as the only dependent variable and serum cholesterol, serum triglyceride, FBG and HAM-D score as independent variables taken together. It was found that hsCRP had significant dependence only on the HAM-D score (β = 0.735, t = 6.286, P < 0.001) (Table 4).

Table 4: Multiple Linear Regression showing the Dependence of hsCRP on Selected Study Parameters

<table>
<thead>
<tr>
<th>Model</th>
<th>Unstandardized Coefficients</th>
<th>Standardized Coefficients</th>
<th>t</th>
<th>Sig.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>B</td>
<td>Std. Error</td>
<td>Beta</td>
<td></td>
</tr>
<tr>
<td>(Constant)</td>
<td>1.024</td>
<td>0.625</td>
<td>1.640</td>
<td>P=1.08</td>
</tr>
<tr>
<td>FBG</td>
<td>2.16E-03</td>
<td>0.002</td>
<td>0.123</td>
<td>1.06</td>
</tr>
<tr>
<td>Serum Cholesterol (mg/dL)</td>
<td>-5.68E-03</td>
<td>0.003</td>
<td>-0.233</td>
<td>-1.913</td>
</tr>
<tr>
<td>Serum Triglyceride (mg/dL)</td>
<td>-1.13E-03</td>
<td>0.001</td>
<td>-0.092</td>
<td>-0.762</td>
</tr>
<tr>
<td>HAM-D Score</td>
<td>9.84E-02</td>
<td>0.019</td>
<td>0.579</td>
<td>5.070</td>
</tr>
</tbody>
</table>

Level of sig. P < 0.05 at 95% CI.
*P-value significant at 0.05 level.

Precision of the tests was monitored assessing the coefficient of variation (CV). CV for the FBG, serum cholesterol and triglyceride remained under 9 percent, while that for the hsCRP was below 6 percent on average throughout the study period.

DISCUSSION

For nearly two decades, it has been recognized that immune system plays a major role in depression. Role of systemic immune activation has been documented in major depression with respect to changes in acute phase response, notably enhancement of positive and diminution of negative acute phase proteins that mark the systemic inflammation.[12-19] CRP is an acute phase protein that suggests the systemic inflammation. This study, however, stems from the research question whether systemic inflammation precedes the onset of depressive symptoms or occurs as a part of somatic manifestation of the depressive phenotype. Although the CRP level is traditionally only elevated in severe inflammation, newer assays with improved sensitivity are able to measure hsCRP in apparently healthy individuals, permitting an exploration of the postulated association between subclinical systemic inflammation and risk of depression.

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Regarding the association of hsCRP with depression was observed in the present study.

Some earlier studies have shown the probable mechanism of CRP rise in depression. Depression might promote an inflammatory response by activating immune response. A higher transcription of both Chemokine Ligands (CCL24) as well as Chemokine Receptors (CCR6) have been reported recently in major depressive disorders. These chemokines are found to attract increasing numbers of T cells and eosinophils, whereas their receptors potentiate the immunological functions of macrophages, B cells and helper T cells. Alternatively, the effects of depression on inflammation might be due to its link to psychological stress. For a long time, depressive illness is known to be associated with an increase in several phases of psychological stress as evident by various biochemical tests including the cortisol response test to psychological stressors. The latter has also been associated with excessive production of interleukin-6 (IL-6). IL-6 is also the main pro-inflammatory cytokine inducing the synthesis of Type I acute phase protein such as CRP. Also psychological stress has been shown to increase oxidative stress, which in turn through modified lipids and lipoproteins is thought to initiate an inflammatory response in arterial wall. It is recognized that depressive symptoms are associated with heightened stress-induced norepinephrine responses and norepinephrine dysregulation. Preliminary evidence suggests that nor-epinephrine dependent adrenal stimulation results in activation of nuclear factor κB (NF-κB), a transcription factor known to promote IL-6 gene expression. Severity of the depressive symptoms, anger and hostility, alone and in combination is associated with increased gene expression of pro-inflammatory cytokines and chemokines, and elevated levels of IL-6. Thus, individuals who show elevated levels of stress may respond to daily life stressors with excessive stress induced sympathetic activation that triggers an NF-κB-dependent cascade of pro-inflammatory events that contribute to increase in hsCRP.

CONCLUSION
Thus, the present study strongly suggests that the depressive illness is closely linked to a pro-inflammatory condition and therefore heralds an early onset of all such related complications like premature atherosclerosis, cardiovascular disorders and cerebrovascular accidents.

RECOMMENDATION
The present study, hence, suggests a close monitoring and surveillance of all depressive patients for pro-inflammatory markers, particularly the hsCRP to enable an early intervention for preventing low grade inflammation related disorders and atherosclerotic diseases in these patients.

REFERENCES