Status of high sensitive C-reactive protein (hs-CRP) in NAFLD obese type-2 diabetic subjects

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Abstract
Introduction: Non alcoholic fatty liver disease (NAFLD) is a disease occurring in patients without any significant alcohol consumption. It includes a broad spectrum and advanced forms of liver diseases, from simple steatosis to nonalcoholic steatohepatitis, fibrosis, and cirrhosis and is associated with diabetes mellitus, insulin resistance, obesity, dyslipidemia and metabolic syndrome. The aim of this study is to evaluate the metabolic significance of NAFLD in obese and type-2 diabetic subjects compare to healthy group of individuals and those independent factors associated with NAFLD.

Materials and Methods: Our study was conducted in Govt. Medical College, Banda (UP) as well as G. R. Medical College, Gwalior (MP). A total of 200 subjects were involved in our study, diagnosis being based on ultrasonography. Out of these, 100 were obese with type-2 diabetic patients and 100 were healthy subjects. They were evaluated by measurement of BMI and several biochemical blood parameters such as fasting blood glucose, lipid profile, liver function tests (ALT, ALP and GGT) and hs-CRP level.

Results: An increase in the BMI and levels of FBS, total cholesterol, triglycerides, LDL, VLDL, GGT, ALT, ALP and hs-CRP level and a decrease in HDL was observed in NAFLD group of individuals. The values of these biochemical markers were found higher in NAFLD patients and the variations were found to be statically significant. BMI and triglycerides were positively correlated with fatty liver.

Conclusion: Obesity, hyperglycemia, dyslipidemia and increased liver enzymes and hs-CRP level are seen more commonly in non alcoholic fatty liver in type-2 diabetic obese patients.

Keywords: NAFLD, BMI, High sensitive C-reactive protein.

Introduction
Non alcoholic fatty liver disease (NAFLD) describes a clinicopathological condition that is described by significant lipid deposition in the hepatocytes of the liver parenchyma in patients with no history of excessive intake of alcohol. The spectrum of NAFLD is broad, ranging from a simple steatosis to nonalcoholic steatohepatitis, fibrosis and cirrhosis. Obesity, dyslipidemia, insulin resistance and diabetes mellitus are well known risk factors for the development of a fatty liver disease.1,2 NAFLD includes patients with simple steatosis and non-alcoholic steatohepatitis, which can lead to cirrhosis and hepatocellular carcinoma.3 Another inflammatory marker, hs-CRP has given mixed results as a biomarker while interleukin-6 (IL-6) has been shown to distinguish differentiante between disease states and independently correlates with fibrosis.4,5

Materials and Methods
Subjects were randomly taken from those attending the medical outpatient department (OPD) of Government Medical College, Banda (UP) as well as G.R. Medical College, Gwalior (MP). A total number of 200 cases were included in our study, diagnosis being based on ultrasonography. Out of these, 100 were non obese and non diabetic healthy controls and 100 were obese with type-2 diabetic patients of both sexes male and female included in our observation, except based on significant alcohol consumption (>20 g/day). A written consent was obtained from the patients. The approval was taken from the institutional ethics committee for conducting the study. Subject was instructed to stand still on the platform, with the body weight evenly distributed between both the feet, for the measurement of weight. Weight (Seca 803, digital scale, Germany) was measured to the nearest of 0.1 kg. Stadiometer (Seca 206, Germany) was used for the measurement of height with head held in Frankfort plane to the nearest of 0.1 cm. BMI was calculated by the formula i.e. weight (kg)/height (m²).

The patients were then evaluated by the measurement of Body Mass Index (BMI), FBS (Fasting Blood Sugar), TC (Total Cholesterol), TG (Triglycerides), LDL (Low Density Lipoprotein), HDL (High Density Lipoprotein), VLDL (Very Low Density Lipoprotein), TB (Total Bilirubin), ALT (Alanine Aminotransferase), GGT (Gamma Glutamyl Transpeptidase), ALP (Alkaline Phosphatase), and hs-CRP. The data was analysed by Statistical packages for
social science (SPSS version 21.0). Mean and standard deviation were analysed for quantitative variables like BMI, FBS, TC, TG, LDL, HDL, VLDL, Total Bilirubin, ALT, GGT, ALP and hs-CRP. To compare mean of all the quantitative variables between the two groups the independent sample t-test of patients were 0.001 level was considered significant. Correlation coefficient analysis was performed for risk factors of NAFLD. P-value less than 0.05 was considered statistically significant.

### Results

In our study, 100 patients of type-2 diabetes mellitus obese subjects and 100 healthy controls were included. The average age of the patients was 52 ± 8 years (Ranging from 36 to 74). Comparison of means of serum biochemical markers between fatty liver and healthy groups is presented in Table. The values of all these biochemical parameters except HDL were increased in fatty liver disease patients in comparison to healthy group and the variations were found to be statistically significant (P value <0.01 and <0.001). BMI and triglycerides were positively correlated with nonalcoholic fatty liver disease.

### Discussion

The prevalence of NAFLD is highly associated with insulin resistance, type-2 diabetes mellitus, obesity, dyslipidemia and metabolic syndrome. The presence of NAFLD correlates significantly with BMI. In our study only BMI was taken as a marker for obesity, raised BMI indicated strong correlation with presence of nonalcoholic fatty liver. In obese type-2 diabetic fatty liver group, the mean BMI was 32.85±3.89 but in non-fatty liver group it was 22.6±1.88 (P value < 0.01). In literature, the prevalence of NAFLD has been found to be 100 %, among severe obese patients with type-2 diabetes mellitus patients. Visceral obesity is frequently associated with NAFLD and their coexistence in the same individual subject increases the chances of having more advanced forms of liver disease. NAFLD is found in 60%–95% of people with obesity. The correlation between fatty liver, impaired glucose tolerance, diabetes mellitus and hyperlipidemia is well recognized. It has been established that insulin resistance can lead to increased free fatty acid in the liver, subsequently higher triglyceride synthesis and increased secretion of triglyceride rich VLDL from the liver. Hypertriglyceridemia have been also strongly associated with liver fat accumulation. Our study showed FBS levels in fatty liver disease group (mean 148.0±37.7) were higher than healthy control group (mean 87.46±13.46), which confirmed the obvious dysglycemia in these patients (P value <0.001). We also observed that increased triglyceride levels (mean 212.7±55.05) in diabetic fatty liver group in comparison to healthy group (triglycerides mean 128.09±19.49) and the results were statistically significant (P value <0.001). In a correlation coefficient analysis triglycerides were also found to be increase in obese type 2 diabetic obese nonalcoholic fatty liver population. The study was done in China also found that fatty liver positively correlated with plasma triglyceride levels and negatively with plasma HDL-C level. In our study also, the elevated total cholesterol positively correlated with fatty liver disease. We also found that increased levels of ALT, ALP and GGT in obese type 2 diabetic NAFLD subjects in comparison to healthy group and the results were statistically significant (P value <0.001). This is also reported in other studies as well. In this study, though raised ALT levels are taken as the first marker of fatty infiltration of the liver. The mechanism behind the increased secretion of liver enzymes in NAFLD patients is that dyslipidemia and insulin resistance

<table>
<thead>
<tr>
<th>Study Variables</th>
<th>Healthy controls n=100</th>
<th>NAFLD n=100</th>
<th>t values</th>
<th>P values</th>
</tr>
</thead>
<tbody>
<tr>
<td>BMI</td>
<td>22.6±1.88</td>
<td>32.85±3.89</td>
<td>4.048</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>FBS</td>
<td>87.46±13.46</td>
<td>148.0±37.7</td>
<td>6.426</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>Total Cholesterol (mg/dl)</td>
<td>186.3±21.09</td>
<td>248.4±41.9</td>
<td>4.234</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>128.09±19.49</td>
<td>212.7±55.05</td>
<td>8.549</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>HDL (mg/dl)</td>
<td>47.45±6.14</td>
<td>33.27±7.02</td>
<td>3.267</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>LDL (mg/dl)</td>
<td>114.1±20.20</td>
<td>161.7±39.3</td>
<td>7.384</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>VLDL (mg/dl)</td>
<td>23.58±4.09</td>
<td>52.89±11.3</td>
<td>6.725</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>ALT (IU/L)</td>
<td>27.3±6.44</td>
<td>54.16±16.1</td>
<td>5.362</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>ALP (IU/L)</td>
<td>179.7±336.75</td>
<td>292.6±21.0</td>
<td>7.180</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>GGT (IU/L)</td>
<td>25.05±5.24</td>
<td>49.39±9.15</td>
<td>6.752</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>hs-CRP (mg/L)</td>
<td>1.03±0.46</td>
<td>3.66±1.20</td>
<td>7.902</td>
<td>P&lt;0.001**</td>
</tr>
<tr>
<td>Total bilirubin (mg/dl)</td>
<td>0.72±0.26</td>
<td>1.22±0.50</td>
<td>2.567</td>
<td>P&lt;0.01*</td>
</tr>
</tbody>
</table>

* Significant at P<0.01, ** significant at P<0.001
leading to significant lipid deposition in hepatocytes causes initiation of mitochondrial swelling, increased lysosomal fragility and impaired membrane integrity, resulting release of hepatic enzymes from the damaged hepatocytes. Furthermore, it is concluded that fasting blood glucose level was an independent predictor of nonalcoholic fatty liver disease. This deviation between our data to that of Marchesini et al implies that the clinicopathological profile of Indian NAFLD patients are different from that seen in other ethnic group of individuals. The present study also, revealed significant higher levels of serum high sensitive C-reactive protein in NAFLD female and male patients (mean 3.66±1.20) in comparison to those of their corresponding healthy controls (mean 1.03±0.46) with significant higher levels in female patients compared to those levels of male patients. These findings similar with those of previous studies. hs-CRP levels have been shown to be closely related to obesity, in particular central or visceral fat deposition. The study of Koruk et al. found that increased levels of hs-CRP could be helpful in the diagnostic work-up of patients with fatty liver disease.

Moreover added that hs-CRP could be a clinical feature that not only distinguishes (Non-alcoholic steatohepatitis) NASH from simple non-progressive steatosis but also indicates the severity of hepatic fibrosis. Bilirubin levels may also be linked to NAFLD via fatty acid metabolism. Higher de novo lipogenesis and peripheral fatty acids mainly derived from lipolysis of adipose tissue contribute to the accumulation of hepatic fat in NAFLD.

Conclusion
The occurrence of NAFLD is high in type-2 diabetic patients, insulin resistance and obesity, dyslipidemia, dysglycemia, increased secretion of liver enzymes (ALT, ALP and GGT) and hs-CRP is seen more frequently in fatty liver than in non-fatty liver subjects. The associated risk factors for diabetic fatty liver are the raised BMI and increased levels of triglycerides.

Limitation
Our study also came up with few limits. The diagnosis of NAFLD in the study was based on ultrasonography (USG) and exception of the known causes of chronic liver disease, but this was not confirmed by liver biopsy, in our study. Some clinical trials regarding use of other inflammatory markers also prove significant.

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Conflict of Interest: There has been no conflict of interest at any stage of the study.

References


