Prevalence of non-alcoholic fatty liver disease on computed tomography in patients with inflammatory bowel disease visiting an emergency department

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\textbf{Introduction} 

Non-alcoholic fatty liver disease (NAFLD) is more common in the western world, with a prevalence of 19-46% in the general US population [1-13]. Risk factors for NAFLD include obesity, insulin resistance, diabetes mellitus (DM), hypertension, and hypertriglyceridemia [14]. NAFLD in patients with inflammatory bowel disease (IBD) has become increasingly recognized, with the prevalence of NAFLD in IBD patients ranging from 8-59% [15-19]. The pathogenesis of NAFLD among the IBD population is not well understood. Other than metabolic syndrome, IBD-specific risk factors, including malnutrition, intestinal inflammation, alterations of gut microbiota, steroid exposure, and drug-induced hepatotoxicity may also contribute to pathogenesis of NAFLD [20].

The aim of this study was to evaluate the prevalence of NAFLD in IBD patients compared with a matched controlled group and to identify the characteristics associated with NAFLD in IBD patients.
Patients and methods

The Institutional Review Board of Saint Luke’s Hospital of Kansas City approved the data collection and methodology for this study. We conducted a retrospective study of IBD patients 18 years of age or older who presented to the emergency department (ED) with a chief complaint of abdominal pain and who completed an abdominal computed tomography (CT) scan in their evaluation. ED encounters within the medical records database between 2009 and 2014 were queried for ICD-9 codes associated with abdominal pain and IBD (either Crohn’s disease or ulcerative colitis). The search was further refined to include only encounters in which abdominal CT scans were performed, based on billing codes associated with the visits. ED provider notes and, when applicable, admission notes were reviewed for possible exclusion criteria, including significant alcohol use, defined as >7 drinks per week in females and >14 drinks per week in males, known liver disease other than NAFLD, and use of corticosteroids within the preceding 12 months. A control group, matched by age (±3 years) and body mass index (BMI; ±2.5), was compiled from patients without IBD who presented to the ED for abdominal pain over the same time period and underwent abdominal CT scans. Medical records were reviewed and the following data were collected: age, BMI, presence of DM, IBD type, levels of aspartate aminotransferase (AST), alanine aminotransferase (ALT), total bilirubin, alkaline phosphatase, albumin, and platelet count. The CT scans of all patients were reviewed specifically for the presence of NAFLD by a specialized abdominal radiologist (SA). Patients were determined to have NAFLD if the liver parenchyma demonstrated a lower density compared with the spleen of >10 Hounsfield units (HU). The primary endpoint was the presence or absence of hepatic steatosis based on CT imaging.

Statistical analysis

The data were presented as mean ± standard deviation for continuous variables and as a total percentage for categorical variables. Student’s t-test and chi-square tests were used in the unadjusted analysis. The predictive factors for NAFLD were assessed with a multivariate logistic regression model adjusted for age, sex, BMI, the presence of DM and transaminase levels. The presence of IBD was independently associated with NAFLD, with an odds ratio of 4.53 (95% confidence interval [CI] 2.00-10.26, P=0.002) (Fig. 1).

Results

Of the 70 patients with IBD, 31 (44%) had hepatic steatosis on abdominal CT images, compared with only 16% (11 of 70) of controls. The difference in NAFLD prevalence between the 2 groups was statistically significant (P<0.001). Patient characteristics, including age, sex, BMI, and the presence of diabetes were similar across the 2 groups (Table 1). There were no significant differences between the 2 groups in transaminase or albumin levels or platelet count (Table 2). The predictive factors associated with NAFLD were examined in a multivariate logistic regression model adjusted for age, sex, BMI, presence of DM, and transaminase levels. The presence of IBD was independently associated with NAFLD, with an odds ratio of 4.53 (95% confidence interval [CI] 2.00-10.26, P=0.002) (Fig. 1).

Table 1 Characteristics of patients enrolled in the cohort

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Total (n=140)</th>
<th>IBD (n=70)</th>
<th>Control* (n=70)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age 38.7±14.9</td>
<td>38.6±15</td>
<td>38.8±15</td>
<td>0.95</td>
<td></td>
</tr>
<tr>
<td>Male sex 58 (41%)</td>
<td>31 (44%)</td>
<td>27 (39%)</td>
<td>0.49</td>
<td></td>
</tr>
<tr>
<td>BMI 26.7±6.7</td>
<td>26.6±6.9</td>
<td>26.7±6.6</td>
<td>0.65</td>
<td></td>
</tr>
<tr>
<td>Crohn’s disease 57 (81%)</td>
<td>57 (81%)</td>
<td>N/A</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diabetes 14 (10%)</td>
<td>8 (11%)</td>
<td>6 (9%)</td>
<td>0.57</td>
<td></td>
</tr>
<tr>
<td>NAFLD 42 (30%)</td>
<td>31 (44%)</td>
<td>11 (16%)</td>
<td>&lt;0.001*</td>
<td></td>
</tr>
</tbody>
</table>

*Age- and BMI-matched control group

Table 2 Laboratory data of patients enrolled in the cohort

<table>
<thead>
<tr>
<th>Lab data</th>
<th>Total (n=140)</th>
<th>IBD (n=70)</th>
<th>Control* (n=70)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>AST 35±38</td>
<td>32±39</td>
<td>38±36</td>
<td>0.7</td>
<td></td>
</tr>
<tr>
<td>ALT 38±56</td>
<td>34±34</td>
<td>42±71</td>
<td>0.19</td>
<td></td>
</tr>
<tr>
<td>Albumin 4±0.6</td>
<td>3.8±0.6</td>
<td>4.3±0.5</td>
<td>0.24</td>
<td></td>
</tr>
<tr>
<td>Platelet 267±93</td>
<td>286±101</td>
<td>250±82</td>
<td>0.07</td>
<td></td>
</tr>
</tbody>
</table>

*Age- and BMI-matched control group

Figure 1 Predictive factors associated with the presence of NAFLD from the study cohort, adjusted for age, male sex, BMI, presence of diabetes, and transaminase levels (n=140)

NAFLD, non-alcoholic fatty liver disease; IBD, inflammatory bowel disease; BMI, body mass index; CI, confidence interval
Discussion

Because of the obesity epidemic and the increasing prevalence of metabolic syndrome, the global prevalence of NAFLD has been rising and is now estimated to be 24% [13]. The reported prevalence of NAFLD in IBD patients has varied from 8-59%, depending on the diagnostic criteria: 59% based on liver histology, 34% based on hepatic steatosis index score, 8% based on combined multiple imaging modalities, and 26-40% based on ultrasonography [15-19]. In this population, NAFLD was diagnosed in 44% of IBD patients on the basis of abdominal CT results, compared with 16% in non-IBD patients. The diagnostic value of abdominal CT for NAFLD has been investigated and a liver-to-spleen difference in attenuation of >9 HU had a specificity of 100% and a sensitivity of 82% in the detection of moderate to severe hepatic steatosis [20]. In our study, patients were determined to have NAFLD if the liver-to-spleen difference in attenuation was >10 HU. Hypertension, obesity and metabolic syndrome were associated with the presence of NAFLD in IBD patients [16,19]. We compared our study population of IBD patients with an age- and BMI-matched controlled group; there were no significant differences in metabolic profile between the 2 groups.

Of our IBD study population, 81% were diagnosed with Crohn’s disease, whereas 19% had ulcerative colitis. The sample sizes were too small to perform a subgroup analysis to identify the distinction between Crohn’s disease and ulcerative colitis in relation to NAFLD prevalence.

Some studies have reported that IBD disease activity or severity and prior small bowel surgery were associated with NAFLD [17,21] However, other studies reported no such association between NAFLD and IBD [19,22]. In this study group, the presence of IBD was very strongly associated with NAFLD, with an odds ratio of 4.53 (95%CI 2.00-10.26), P=0.002.

A limitation of this study was the incomplete data recording of our ED cohort. A detailed smoking history, IBD disease severity, duration of diagnosis, prior IBD-related surgery, and IBD medications or treatment were not completely noted. As the study evaluated an ED population, there may have been an inherent selection bias towards patients with more severe disease activity and less medical compliance. Since glucocorticoids have been implicated in the pathogenesis of NAFLD, we excluded patients who had taken corticosteroids within the previous 12 months. Thus, the NAFLD prevalence reported in our study does not apply to IBD patients exposed to corticosteroid therapy within a 1-year period.

In conclusion, data concerning the association between IBD disease activity and NAFLD are conflicting. Our results from this controlled study demonstrate that the presence of IBD in patients who visited the ED was strongly associated with NAFLD. Systemic inflammation and alteration of the intestinal microbiome have been proposed as potential causative factors, but further studies will be needed to elucidate the pathophysiology.

Summary Box

What is already known:

• The global prevalence of non-alcoholic fatty liver disease (NAFLD) has been rising because of the epidemic of obesity and metabolic syndrome
• The prevalence of NAFLD in patients with inflammatory bowel disease (IBD) reportedly varies from 8-59%, depending on the diagnostic criteria
• Hypertension, obesity and metabolic syndrome are associated with NAFLD in IBD patients
• Data regarding IBD disease severity and prior small bowel surgery in association with NAFLD in IBD patients are conflicting

What the new findings are:

• Based on abdominal computed tomography, NAFLD was also more common (44%) in IBD patients who visited the emergency department, compared with a control group without IBD
• NAFLD is independently associated with the presence of IBD
• Further studies are still needed to elucidate the pathophysiology of NAFLD development in IBD populations

References

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