The relationship between fibrosis and nodule structure and esophageal varices

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ABSTRACT

Background/Aims: The aim of the present study was to evaluate the histopathological findings of cirrhosis together with clinical and laboratory parameters, and to investigate their relationship with esophageal varices that are portal hypertension findings.

Materials and Methods: A total of 67 (42 male and 25 female) patients who were diagnosed with cirrhosis were included in the study. The mean age of the patients was 51.6±19.0 (1-81) years. The biopsy specimens of the patients were graded in terms of fibrosis, nodularity, loss of portal area, central venous loss, inflammation, and steatosis. The spleen sizes were graded ultrasonographically, and the esophageal varices were graded endoscopically.

Results: In the multivariate regression analysis, there was a correlation between the advanced disease stage (Child-Pugh score odds ratio (OR): 1.47, 95% confidence interval (CI): 1.018-2.121, p=0.040), presence of micronodularity (OR: 0.318, 95% CI: 0.120-0.842, p=0.021), grade of central venous loss (OR: 5.231, 95% CI: 1.132-24.176, p=0.034), and presence of esophageal varicose veins.

Conclusion: Although thrombocytopenia and splenomegaly may predict the presence of large esophageal varices, cirrhosis histopathology is the main factor in the presence of varices.

Keywords: Cirrhosis, histopathology, portal hypertension, esophageal varices

INTRODUCTION

Cirrhosis is histologically defined by the presence of regenerative nodules surrounded by fibrous tissue. This structural distortion leads to increased intrahepatic resistance and portal hypertension. Esophageal varices and ascites occur after portal pressure exceeds 10–12 mm Hg (1–3). The severity of the functional and structural impairment of the liver determines the course of the disease. Portal hypertension–related esophageal variceal bleeding, ascites, and hepatic encephalopathy generally determine the prognosis of patients with cirrhosis. In clinical practice, some patients with cirrhosis live in an asymptomatic or compensated state, whereas others decompensate over time. Owing to the variability of the clinical course, it is beneficial to predict the clinical course using various laboratory or histological signs. The histological features of cirrhosis do not reflect the clinical severity or the stage of cirrhosis. As certain histological parameters can predict clinical and biochemical decompensation, it is considerably important to be able to determine a relationship between specific histological parameters, such as fibrosis and nodule structure, on liver biopsies of patients with cirrhosis and clinical findings of portal hypertension, such as esophageal varices. The aim of the present study was to evaluate the relationship between some histopathological findings of cirrhosis with clinical, endoscopic, and laboratory parameters.

MATERIALS AND METHODS

The study was approved by the Başkent University Research and Ethics Committee. Data of 194 patients admitted to our hospital between July 1999 and June 2006 who underwent liver biopsies for the evaluation of chronic liver diseases, liver masses, or liver function abnormalities were reviewed retrospectively. Patients with suitable liver biopsy specimens for histopathological reassessment were included in the study.

Histopathological assessment

All liver biopsies were fixed in 10% formalin and routinely stained with hematoxylin and eosin, reticulin, Masson trichrome, Periodic Acid–Schiff, and iron stain. The slides of each biopsy were reviewed by an experienced hepatopathologist blinded to the patient’s clinical status. Liver biopsy samples >10 mm in length were accepted for histopathological examination. In fragmented biopsies, the...
The total length was estimated by combining the maximum dimensions of each individual fragment. Liver fibrosis was staged according to the Batts and Ludwig classification, a modification of the Scheuer classification, in which stage 0 corresponds to no fibrosis, stage 1 to portal fibrosis, stage 2 to periportal fibrosis, stage 3 to bridging fibrosis, and stage 4 to cirrhosis (4). Only biopsies categorized as stage 4 were included in the study.

The biopsies were re-evaluated for two parameters: sinusoidal fibrosis and nodule size. In the definition of nodule size, the “mixed nodule” classification was used when there was at least one small nodule in addition to macro-nodules. For septal thickness, the thickness of the dominant type of septae in each specimen was scored. The histological parameters and scores assessed in the biopsy samples were as follows:

1. Sinusoidal fibrosis: 0, 1 (mild), 2 (moderate), and 3 (severe),
2. Septal thickness: 0, 1 (thin), 2 (medium), and 3 (thick),
3. Nodularity:
   a. Small nodules: nodule size is comparable to the width of the needle biopsy specimen,
   b. Mixed nodules: presence of both small and large nodules,
   c. Large nodules: nodule size is larger than the needle biopsy specimen.

**Endoscopic assessment**

All patients underwent upper gastrointestinal endoscopy for the assessment of esophageal varices as an endoscopic finding of portal hypertension. The Pentax EPM-3500 and Olympus CV-70 endoscopy systems were used to determine the presence of esophageal varices, and two experienced endoscopists performed the procedures. The esophageal varices were graded as I-III according to the Japanese Classification of Esophageal Varices. Patients were classified dichotomously either as Group 1 (large, grade II-III esophageal varices) or Group 2 (none or small, grade I varices).

**Statistical analysis**

All statistical analyses were performed using the SPSS software (version 14 for Windows; SPSS, Chicago, IL, USA). As most variables were not normally distributed, data were presented as median and range (minimum–maximum) values. Owing to the heterogeneity of the distribution, the non-parametric Mann-Whitney U-test, the Kruskal-Wallis test, and the chi-square test were used. Logistic regression analysis was performed for multivariate analysis. A p value of <0.05 was considered statistically significant. A non-parametric statistical analysis was applied to the data, such as esophageal varices and platelet count, which were the clinical and laboratory findings of portal hypertension, the histopathological findings, and the demographic data.

**RESULTS**

The study included 67 (25 (37.3%) female and 42 (62.7%) male) patients with histologically proven cirrhosis. The mean ages of the patients were 52.3±18.6 years for female and 51.2±19.5 years for male. Table 1 shows the etiology, Child-Pugh status, and nodule size distributions of patients.

<table>
<thead>
<tr>
<th>Etiology (N=67)</th>
<th>n (%)</th>
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<tbody>
<tr>
<td>HCV</td>
<td>29 (43.3)</td>
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<tr>
<td>HBV</td>
<td>17 (25.2)</td>
</tr>
<tr>
<td>Cryptogenic cirrhosis</td>
<td>10 (4.9)</td>
</tr>
<tr>
<td>Wilson Disease</td>
<td>3 (4.5)</td>
</tr>
<tr>
<td>Alcoholic Cirrhosis</td>
<td>2 (3.0)</td>
</tr>
<tr>
<td>NASH</td>
<td>2 (3.0)</td>
</tr>
<tr>
<td>HBV+HCV</td>
<td>2 (3.0)</td>
</tr>
<tr>
<td>PBC</td>
<td>1 (1.5)</td>
</tr>
<tr>
<td>AIH</td>
<td>1 (1.5)</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Child-Pugh (N=67)</th>
<th>n (%)</th>
</tr>
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<tbody>
<tr>
<td>A</td>
<td>50 (74.6)</td>
</tr>
<tr>
<td>B</td>
<td>8 (11.9)</td>
</tr>
<tr>
<td>C</td>
<td>9 (13.4)</td>
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<table>
<thead>
<tr>
<th>Nodule size (N=67)</th>
<th>n (%)</th>
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<tbody>
<tr>
<td>Small</td>
<td>12 (17.9)</td>
</tr>
<tr>
<td>Mixed</td>
<td>27 (40.3)</td>
</tr>
<tr>
<td>Large</td>
<td>28 (41.8)</td>
</tr>
</tbody>
</table>

The relationship between the demographic parameters and the presence and group of esophageal varices with liver histopathology of the patients was examined for univariate analysis.

The results are as follows (Table 2):

1. A statistically significant relationship was detected between the nodule size and age (p<0.001), platelet...
count (p=0.04), Child-Pugh score (p=0.001), esophageal varices (p=0.001), and large esophageal varices (p=0.03).

2. A statistically significant relationship was detected between a high Child-Pugh score (p=0.047) and the thickness of septal fibrosis.

Table 3 presents the relationship between the presence of large esophageal varices (FII or FIII) and the other clinical and laboratory parameters of portal hypertension and demographic data. Table 4 presents the relationship between the presence of esophageal varices and the similar parameters and data. A positive and statistically significant relationship was detected between a high Child-Pugh score (p<0.001) and the presence of both large esophageal varices and esophageal varices. Multivariate logistic regression analysis revealed a statistically significant relationship between the Child-Pugh score (p=0.040), micronodularity (p=0.021), and presence of varices (Table 5).

**Discussion**
The consequences of portal hypertension, such as variceal bleeding, are important factors in the morbidity and mortality of patients with cirrhosis (5,6). At the
Third Baveno Conference on portal hypertension and its treatment, all patients with cirrhosis are suggested to be screened for the presence of esophageal varices regardless of predictor factors. Endoscopic scanning every 2 to 3 years for patients who do not show any presence of varices endoscopically and every 1 to 2 years for those who have ambiguous varices has been recommended (7,8). However, an endoscopy is an invasive and relatively expensive procedure that is not tolerated by some patients. Moreover, the definition of variceal sizes depends on the endoscopist. Therefore, researchers have turned to alternative non-invasive methods for variceal screening. Several clinical and laboratory parameters have been examined in terms of their ability to predict the presence of varices. In the literature, >10 studies have specifically investigated this issue (9-20). Chalasani et al. argued that the presence of thrombocytopenia and splenomegaly is an independent predictor for the presence of esophageal varices (15). Madhotra showed that the presence of thrombocytopenia and splenomegaly is an independent predictor for the presence of large esophageal varices. However, in the present study, the presence of splenomegaly could not be found in 32% of the cases where platelet counts were below the cut-off value (68,000/mm\(^3\)). The sensitivity and specificity of splenomegaly and thrombocytopenia (<68,000/mm\(^3\)) were 75% and 58% and 71% and 73%, respectively (16). Therefore, the presence of splenomegaly per se does not explain the thrombocytopenia seen in patients with cirrhosis. Other possible reasons may include “decreased thrombopoietin synthesis” and “altered immune mechanisms” (21).

Other than the clinical and laboratory findings of portal hypertension, other remaining studies have focused on the histopathological characteristics of the liver and the development of esophageal varices.

For example, the Fibro Test is an alternative non-invasive method for analyzing liver fibrosis instead of histopathology. Its relationship with varices prediction was examined by Thabut et al. (14), who investigated the relationship of platelet count, prothrombin time, presence of ascites, Child-Pugh score, and Fibro Test results with large esophageal varices. In their study, Fibro Test was found to be the parameter with the highest differentiation power. Nagula et al. (22) investigated portal vein pressure in patients with liver cirrhosis and its relationship with histopathological parameters. A statistically significant relationship was found between cirrhotic nodule size and septal thickness and the presence of portal hypertension. In the univariate analysis of data, Nagula et al. (22) found a statistically significant relationship between the presence of micronodules and platelet count, Child-Pugh score, presence of esophageal varices, presence of large varices, and age.

Common causes of micronodular cirrhosis include chronic alcohol usage, hemochromatosis, biliary tract obstruction, chronic venous flow obstruction, and childhood...
metabolic diseases, and one of the rare causes is cirrhosis secondary to chronic active hepatitis. In the current study group, the micronodular cirrhosis rate was almost 18%, whereas alcoholic cirrhosis was relatively infrequent at 3% (Table 1). The etiology of cirrhosis was hepatitis B or C infections or hepatitis B and C co-infection in 72% of the patients. The mixed nodularity rate in this group at almost 40% was classed as relatively high, in contrast with the conventional explanation that “the illnesses that cause macronodular cirrhosis from the beginning are chronically viral and autoimmune hepatitis” (23). In the present study, although the rate of alcoholic etiology in the patient population was very low, and most of the patients showed viral etiology, micronodule formation in the pathology specimens was a very frequent finding when mixed nodularity was added (Tables 1 and 6). This can be attributed to the transformation of the cirrhosis morphology, which was initially micronodular and became macronodular during the course of the disease. Other possible explanations could include other etiological reasons that accompany the viral etiology but are not seen as dominant or sufficient enough to determine this morphological classification in most biopsy materials (23). In the current study, a statistically significant relationship was determined between micronodularity and the average high level of the Child-Pugh score (Table 7). Accordingly, it may be advisable that patients with micronodular cirrhosis are closely monitored. The relationship between the presence of large varices and micronodularity was not statistically significant in the multivariate analysis, possibly due to the low number of patients with micronodular cirrhosis. The majority of patients with micronodular cirrhosis in the current study group were viral hepatitis B, C, or D positive. The results of the study showed a statistically significant relationship between the Child-Pugh score and the presence of varices in multivariate analysis. With respect to liver structure, compared with macronodules, micronodules can cause greater structural damage, leading to the development of portal hypertension and thereby contributing to the development of esophageal varices. Previous studies have shown the relationship between thrombocytopenia and liver fibrosis (14,24). However, as in alcoholic hepatitis without manifest fibrosis, varices and other portal hypertension findings can occur in the early stage of chronic liver disease. In the current study, a statistically significant relationship was also found between the thickness of the fibrous septae and the Child-Pugh score (p=0.047). This relationship supports a possible structural reason that negatively affects liver synthesis functions (Table 2). Micronodule formation was frequent in younger patients in the current study group (p<0.001), as seen in the alcoholic cirrhosis and hemochromatosis cases. This situation can be explained by an increase in the age of onset of cirrhosis or of discontinuing alcohol usage and by the transformative ability of cirrhosis, which was initially micronodular to macronodular cirrhosis following treatments, such as phlebotomy.

A liver biopsy is still the gold standard in the diagnosis of liver diseases. When the cirrhosis diagnosis is proven by biopsy, reporting additional findings, such as nodule size and septal thickness, can serve as a guide for clinicians in predicting the development and course of portal hypertension. In the present study, which aimed to analyze the relationship between the findings of histopathological cirrhosis and various parameters related to portal hypertension, some results that can be transferred to clinical practice were obtained, although the study was retrospective, and the volume of data was relatively low. It can be recommended that all patients with cirrhosis are scanned in daily practice, in terms of the presence of esophageal varices, independently of the predicting factors. The most reliable method of variceal screening, which allows for possible intervention if needed, is endoscopic assessment. Endoscopic variceal screening is indicated and is cost-effective particularly in cases of compensated cirrhosis. Several alternative non-invasive methods have been defined. However, in case of negative results, it is important to keep in mind that the possibility of small varices cannot be excluded, in which case, the use of combined multiple parameters is recommended.

In conclusion, the presence of micronodularity can be accepted as a predictor of varices. This is useful for patients who do not have clear clinical cirrhosis and who are diagnosed after biopsy. It is possible to say that patients with micronodular cirrhosis have (1) a lower platelet count, (2) a higher Child-Pugh score, (3) a high probability of the presence of esophageal varices, (4) a high probability of the presence of large esophageal varices, and (5) lower age.

**Ethics Committee Approval:** Ethics committee approval was obtained from the local institutional ethical committee.

**Informed Consent:** N/A.

**Peer-review:** Externally peer-reviewed.

**Author Contributions:** Concept - A.M.C, E.S; Design - A.M.C, T.Y; Supervision - E.S, B.O; Resources - A.M.C, F.K; Materials - A.M.C,
Conflict of Interest: The authors have no conflict of interest to declare.

Financial Disclosure: The authors declared that this study has received no financial support.

REFERENCES