ASSOCIATION OF VARICEAL BLEED WITH SEVERITY OF LIVER CIRRHOSIS AT A TERTIARY CARE HOSPITAL

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Abstract;
Background: A major cause of cirrhosis-related morbidity and mortality is the development of variceal bleeding, a direct consequence of portal hypertension. Each episode of active variceal bleeding is associated with 30 percent mortality. This study was planned to determine frequency of variceal bleeding in patients with liver cirrhosis and frequency of in-hospital mortality of these patients in our population. Objective: To determine the frequency of variceal bleed in hospitalized patients with cirrhosis of liver and its outcome in terms of in-hospital mortality. Material and Methods: Consecutive 139 Patient diagnosed with cirrhosis of liver were included in this cross-sectional study from department of Medicine, Bahawal Victoria hospital Bahawalpur. Complete history and physical examination was assessed to document duration of duration of liver disease, ascites, Hepatic encephalopathy, Previous GI bleed and systemic coagulopathy. All the patients had undergone diagnostic upper GI endoscopic examination to document varices. Results: Of these 139 study cases, 77 (55.4 %) were male and 62 (44.6 %) were female. Mean age of our study cases was 45.50 ± 10.81 years. Mean duration of disease (liver cirrhosis) was 3.25 ± 2.32 years. Majority of our study cases i.e. 94 (67.6%) were having liver cirrhosis for the duration of less than 5 years. Child-Pugh class C was more prevalent i.e. 77 (55.4%) of our study cases. Variceal bleeding was observed in 100 (71.9 %) of our study cases. Frequency of mortality was 35 (25.2%) in our study cases with liver cirrhosis, while frequency of mortality in patients with variceal bleeding was seen in 31 (31%). Variceal bleeding was significantly associated with disease severity (p<0.001). Conclusion: Very high frequency of variceal bleeding was observed in patients with liver cirrhosis. In-hospital mortality was significantly more prevalent in patients with variceal bleeding than without bleed. Variceal bleeding was significantly more seen in patients with increasing age, duration of disease and with more severe level of disease (Child Pugh class C). There was no statistically significant difference of bleeding with regards to gender. Keywords; Liver Cirrhosis, Variceal bleeding, Mortality.
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Introduction;
Cirrhosis of liver is one of the most frequently encountered entities in clinical setup all over the world especially in our subcontinent. It is defined as a process of diffuse fibrosis and loss of normal architecture with nodule formation. Most common causes of this chronic hepatic condition are viral hepatitis, alcohol and metabolic causes like hemochromatosis and Wilson’s disease. This alteration in hepatic architecture leads to hepatocyte damage and increased resistance to portal blood flow with increase in portal pressure known as portal hypertension. Portal hypertension is most important pathogenic mechanism for development of major complications of cirrhosis like ascites, variceal bleed, Hepatic encephalopathy and Hepatorenal syndrome. Increased portal pressure causes opening up of portosystemic shunts on various sites in body, most important of which are at lower esophagus and stomach. These dilated blood vessels, called Esophageal and gastric varices, are most common cause of upper GIT bleeding in cirrhotic patients. About 65% cases of upper GI bleed (UGIB) in cirrhosis are due to esophageal varices. Esophageal varices are present in 40% cases of cirrhosis at time of hospitalization and are present in 85 % of child class C patients. Variceal bleeding is second most common cause of death in patients with cirrhosis. It carries a 6 week mortality of 10-20% and 1 year mortality of 60%. Similar results were obtained by Hearnshaw and others who found mortality despite of modern methods of treatment to be 10%.
Cirrhosis therefore is one of the most devastating illness affecting the world and particularly in our country. This study will help us to predict the prevalence of esophageal varices and its outcome in terms of in-hospital mortality.

Material and Methods;
Consecutive 139 Patient diagnosed with cirrhosis of liver irrespective of Child Pugh class and duration of disease of either sex less than 70 years were included in this cross-sectional study from department of Medicine, Bahawal Victoria hospital Bahawalpur were included in the study. Patients admitted and expired due to causes other than cirrhosis of liver, acute liver failure, patients with upper GIT bleed with no varices or signs of portal hypertension on Endoscopy and with Coagulopathy were excluded. Informed consent was taken from each patient. Variceal bleed was defined by presence of: Clinical evidence of upper GI bleed in the form of hematemesis and/or melena which is consistent with endoscopic evidence of esophageal and gastric varices. Complete history and physical examination was assessed to document duration of disease, ascites, Hepatic encephalopathy, Previous GI bleed and systemic coagulopathy. All the patients had undergone diagnostic upper GI endoscopic examination to document varices. Data was entered and analyzed using computer program SPSS-16.

Results;
A total of 139 patients with liver cirrhosis meeting inclusion and exclusion criteria of this study were registered. Of these 139 study cases, 77 (55.4%) were male and 62 (44.6%) were female. Mean age of our study cases was 45.50 ± 10.81 years (minimum age was 19 years while maximum was 60 years). Study results have also indicated that majority of our cases were in the range of age groups 41 to 60 years i.e. 94 (67.6%) of our study cases. Mean duration of disease (liver cirrhosis) was 3.25 ± 2.32 years (minimum duration of disease was 1 year while maximum duration of disease was 9 years). Majority of our study cases i.e. 94 (67.6%) were having liver cirrhosis for the duration of less than 5 years. Child-Pugh class C was more prevalent i.e. 77 (55.4%) of our study cases. Variceal bleeding was observed in 100 (71.9%) of our study cases. Frequency of mortality was 35 (25.2%) in our study cases with liver cirrhosis while frequency of mortality in patients with variceal bleeding was seen in 31 (31%) of patients i.e. 31/100. Frequencies of variceal bleeding in different child pugh classes has been described in Table No. 1.

Table-1
Stratification of Variceal bleed with Severity of disease.
(n=139)

| Disease Severity | Yes (n=100) | No (n=39) | P-Value |
|------------------|------------|-----------|---------|
| Child Pugh Class A (n=31) | 12 (39%) | 19 (61%) | |
| Child Pugh Class B (n=31) | 15 (48.3%) | 16 (51.7%) | |
| Child Pugh Class C (n=77) | 73 (94.8%) | 04 (5.2%) | < 0.001 |
| Total | 139 | | |
Discussion;

Cirrhosis affects 3.6 out of every 1000 adults in North America, and is responsible for over one million days of work loss and 32,000 deaths annually. A major cause of cirrhosis-related morbidity and mortality is the development of variceal bleeding, a direct consequence of portal hypertension\textsuperscript{12}. Each episode of active variceal bleeding is associated with 30 percent mortality \textsuperscript{13, 14}. In addition, survivors of an episode of active bleeding have a 70 percent risk of recurrent hemorrhage within one year of the bleeding episode \textsuperscript{15}. Variceal bleeding occurs in 25 to 40 percent of patients with cirrhosis \textsuperscript{16}. Our study registered 139 study cases, 77 (55.4 \%) were male and 62 (44.6 \%) were female patients having liver cirrhosis. Different authors have reported liver cirrhosis being more common among males than those of females. A study conducted by Sohail et al.\textsuperscript{17} reported 56.7 \% male patients of cirrhosis in their study and 43.3\% among females, our study results are close to that of Sohail et al.\textsuperscript{17}. Devrajani et al.\textsuperscript{18} reported 59.3\% male patients with liver cirrhosis and 40.7\% female patients which are close to our study results. Two other studies very high frequencies of male patients than females.\textsuperscript{19, 20} Mean age of our study cases was 45.50 ± 10.81 years (minimum age was 19 years while maximum was 60 years). Study results have also indicated that majority of our cases were in the range of age groups 41 to 60 years i.e. 94 (67.62 \%).

Variceal bleeding was observed in 100 (71.9 \%) of our study cases. Variceal bleeding was observed in 77 (55.4 \%) of our study cases. Variceal bleeding was observed in 35 (25.2 \%) in our study cases with liver cirrhosis. Our overall frequency of mortality was 35 (25.2 \%) in our study cases with liver cirrhosis. Kirge et al.\textsuperscript{20} reported 20.8 \% of mortality among cirrhotic patients, these findings are close to that of our study results. Pichilinque et al.\textsuperscript{21} reported 18.03 \% in-hospital mortality in patients having liver cirrhosis. Hearnshaw et al.\textsuperscript{11} reported 10\% in-hospital mortality in patients having liver cirrhosis. This frequency is lesser than our study results. Sarwar et al.\textsuperscript{19} reported 6.7 \% in-hospital mortality among Cirrhotic patients, which is quite less than our study results.

Variceal bleeding was associated with 30 percent mortality rates \textsuperscript{13, 14} and our study results are also in accordance of these figures as frequency of mortality in patients with variceal bleeding was seen in 31 (31\%). In our study variceal bleeding was not significantly related to gender but was significantly more common with increasing age, duration of disease and Child Pugh class C (severity of disease).

Conclusion;

Very high frequency of variceal bleeding was observed in patients with liver cirrhosis. In-hospital mortality was significantly more prevalent in patients with variceal bleeding than without bleed. Variceal bleeding was significantly more seen in patients with increasing age, duration of disease and with more severe level of disease (Child Pugh class C). There was no statistically significant difference of bleeding with regards to gender.

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