

World Journal of *Hepatology*

World J Hepatol 2020 March 27; 12(3): 72-115



**REVIEW**

- 72 Polycystic liver disease: Classification, diagnosis, treatment process, and clinical management
Zhang ZY, Wang ZM, Huang Y

ORIGINAL ARTICLE**Basic Study**

- 84 High omega arachidonic acid/docosahexaenoic acid ratio induces mitochondrial dysfunction and altered lipid metabolism in human hepatoma cells
Ghazali R, Mehta KJ, Bligh SA, Tewfik I, Clemens D, Patel VB

Retrospective Study

- 99 Platelet-albumin-bilirubin score - a predictor of outcome of acute variceal bleeding in patients with cirrhosis
Elshaarawy O, Allam N, Abdelsameea E, Gomaa A, Waked I

CASE REPORT

- 108 Successful liver transplantation for acute sickle cell intrahepatic cholestasis: A case report and review of the literature
Alkhayyat M, Saleh MA, Zmaili M, Sanghi V, Singh T, Roupheal C, Simons-Linares CR, Romero-Marrero C, Carey WD, Lindenmeyer CC

ABOUT COVER

Editorial Board Member of *World Journal of Hepatology*, Fernando O Bessone, MD, Associate Professor, Gastroenterology and Hepatology Department, University of Rosario School of Medicine, Rosario 2000, Argentina

AIMS AND SCOPE

The primary aim of *World Journal of Hepatology* (WJH, *World J Hepatol*) is to provide scholars and readers from various fields of hepatology with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJH mainly publishes articles reporting research results and findings obtained in the field of hepatology and covering a wide range of topics including chronic cholestatic liver diseases, cirrhosis and its complications, clinical alcoholic liver disease, drug induced liver disease autoimmune, fatty liver disease, genetic and pediatric liver diseases, hepatocellular carcinoma, hepatic stellate cells and fibrosis, liver immunology, liver regeneration, hepatic surgery, liver transplantation, biliary tract pathophysiology, non-invasive markers of liver fibrosis, viral hepatitis.

INDEXING/ABSTRACTING

The WJH is now abstracted and indexed in PubMed, PubMed Central, Emerging Sources Citation Index (Web of Science), Scopus, China National Knowledge Infrastructure (CNKI), China Science and Technology Journal Database (CSTJ), and Superstar Journals Database.

RESPONSIBLE EDITORS FOR THIS ISSUE

Responsible Electronic Editor: Yun-Xiaojuan Wu

Proofing Production Department Director: Xiang Li

NAME OF JOURNAL

World Journal of Hepatology

ISSN

ISSN 1948-5182 (online)

LAUNCH DATE

October 31, 2009

FREQUENCY

Monthly

EDITORS-IN-CHIEF

Ke-Qin Hu, Koo Jeong Kang, Nikolaos Pylsopoulos

EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/1948-5182/editorialboard.htm>

EDITORIAL OFFICE

Ruo-Yu Ma, Director

PUBLICATION DATE

March 27, 2020

COPYRIGHT

© 2020 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/GerInfo/287>

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

ONLINE SUBMISSION

<https://www.f6publishing.com>

Retrospective Study

Platelet-albumin-bilirubin score - a predictor of outcome of acute variceal bleeding in patients with cirrhosis

Omar Elshaarawy, Naglaa Allam, Eman Abdelsameea, Asmaa Gomaa, Imam Waked

ORCID number: Omar Elshaarawy (0000-0002-6945-6204); Naglaa Allam (0000-0001-7659-8756); Eman Abdelsameea (0000-0003-2674-2427); Asmaa Gomaa (0000-0001-9376-4461); Imam Waked (0000-0002-9857-8972).

Author contributions: Elshaarawy O and Abdelsameea E collected patient's data; Elshaarawy O analyzed the data; all authors wrote the manuscript; Allam N and Waked I revised the manuscript.

Institutional review board statement: The study was approved by the institutional review board (IRB number IRB00003413).

Informed consent statement: Informed consent was obtained from all patients.

Conflict-of-interest statement: Imam Waked received grants/research supports or speaker's honoraria from Abbvie, Marcyrl, MSD, Mylan, Onxio, and Pharco. The authors report no other conflicts of interest in this work.

Data sharing statement: No additional data are available.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build

Omar Elshaarawy, Naglaa Allam, Eman Abdelsameea, Asmaa Gomaa, Imam Waked, Department of Hepatology and Gastroenterology, National Liver Institute, Menoufia University, Shebin Elkom 32511, Egypt

Corresponding author: Eman Abdelsameea, MBChB, MD, MSc, Associate Professor, Doctor, Department of Hepatology and Gastroenterology, National Liver Institute, Menoufia University, Shebin Elkom 32511, Egypt. eabdelsameea@liver-eg.org

Abstract

BACKGROUND

The albumin-bilirubin (ALBI) score was validated as a prognostic indicator in patients with liver disease and hepatocellular carcinoma. Incorporating platelet count in the platelet-albumin-bilirubin (PALBI) score improved validity in predicting outcome of patients undergoing resection and ablation.

AIM

To evaluate the PALBI score in predicting outcome of acute variceal bleeding in patients with cirrhosis.

METHODS

The data of 1517 patients with cirrhosis presenting with variceal bleeding were analyzed. Child Turcotte Pugh (CTP) class, Model of End-stage Liver Disease (MELD), ALBI and PALBI scores were calculated on admission, and were correlated to the outcome of variceal bleeding. Areas under the receiving-operator characteristic curve (AUROC) were calculated for survival and rebleeding.

RESULTS

Mean age was 52.6 years; 1176 were male (77.5%), 69 CTP-A (4.5%), 434 CTP-B (29.2%), 1014 CTP-C (66.8%); 306 PALBI-1 (20.2%), 285 PALBI-2 (18.8%), and 926 PALBI-3 (61.1%). Three hundred and thirty-two patients died during hospitalization (21.9%). Bleeding-related mortality occurred in 11% of CTP-B, 28% of CTP-C, in 21.8% of PALBI-2 and 34.4% of PALBI-3 patients. The AUROC for predicting survival of acute variceal bleeding was 0.668, 0.689, 0.803 and 0.871 for CTP, MELD, ALBI and PALBI scores, respectively. For predicting rebleeding the AUROC was 0.681, 0.74, 0.766 and 0.794 for CTP, MELD, ALBI and PALBI scores, respectively.

CONCLUSION

PALBI score on admission is a good prognostic indicator for patients with acute

upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See:

<http://creativecommons.org/licenses/by-nc/4.0/>

Manuscript source: Unsolicited manuscript

Received: September 19, 2019

Peer-review started: September 19, 2019

First decision: October 14, 2019

Revised: January 7, 2020

Accepted: January 14, 2020

Article in press: January 14, 2020

Published online: March 27, 2020

P-Reviewer: Garbuzenko DV, Mura VL

S-Editor: Dou Y

L-Editor: Webster JR

E-Editor: Wu YXJ



variceal bleeding and predicts early mortality and rebleeding.

Key words: Variceal bleeding; Platelet-albumin-bilirubin score; Albumin-bilirubin score; Rebleeding

©The Author(s) 2020. Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: This study describes a comparative analysis of the performance of different scoring systems in a large number of patients with acute variceal bleeding. The platelet-albumin-bilirubin score performed better in predicting short-term outcome and the incidence of rebleeding compared with the other 4 scoring systems, the Child Pugh score, albumin-bilirubin score and Model of End-stage Liver Disease and its modification for acute variceal bleeding.

Citation: Elshaarawy O, Allam N, Abdelsameea E, Gomaa A, Waked I. Platelet-albumin-bilirubin score - a predictor of outcome of acute variceal bleeding in patients with cirrhosis. *World J Hepatol* 2020; 12(3): 99-107

URL: <https://www.wjgnet.com/1948-5182/full/v12/i3/99.htm>

DOI: <https://dx.doi.org/10.4254/wjh.v12.i3.99>

INTRODUCTION

Acute variceal bleeding is a frequent, ominous complication of liver cirrhosis and portal hypertension and is responsible for high morbidity and mortality^[1].

The Child-Turcotte-Pugh (CTP) and Model of End-stage Liver Disease (MELD) scores are two of the most important models for predicting the survival of upper gastrointestinal bleeding. The CTP score was originally developed to be a prognostic score in patients with cirrhosis and portal hypertension undergoing surgery for variceal bleeding^[2]. The limitations of the CTP score in assessing liver functions in patients with upper gastro-intestinal tract (GIT) bleeding are the subjective assessment of encephalopathy and amount of ascites, together with the interrelated variables such as ascites and serum albumin, besides the parameters that are scored based on arbitrarily defined cutoff points^[3,4].

The MELD score was initially developed to determine prognosis following a trans jugular intra-hepatic porto-systemic shunt procedure, and is widely used in liver transplant settings to prioritize donor liver allocation^[5,6]. Johnson *et al*^[7] developed the albumin-bilirubin (ALBI) score, which depends on two laboratory variables; bilirubin and albumin, omits the subjective CTP variables ascites and hepatic encephalopathy, and results are expressed as three grades with three different cutoff points^[7]. Roayaie *et al*^[8] proposed modifying the ALBI score by including the platelet count as an indicator of portal hypertension.

In this study, we aimed to determine the value of the platelet-albumin-bilirubin (PALBI) score in predicting the outcome of patients with cirrhosis presenting with acute variceal bleeding.

MATERIALS AND METHODS

This retrospective study included 1517 patients with cirrhosis presenting with acute variceal bleeding who were admitted to the National Liver Institute Hospital. The study was approved by the institutional review board (IRB number IRB00003413).

Inclusion criteria

This study included patients with acute variceal bleeding. Acute upper gastrointestinal bleeding was considered in patients with liver cirrhosis presenting with hematemesis, defined as either one or more than one episode of vomiting either fresh blood or a coffee ground-like material, or reported or observed melena, with a drop in hemoglobin, and blood in the nasogastric tube.

All patients were managed in the emergency unit and subjected to the following: (1) Resuscitation to maintain hemodynamic stability (systolic blood pressure above 80-90 mmHg); (2) Blood transfusion using type and cross-matched packed red cells

and plasma as required to increase hematocrit to 25%-30% or hemoglobin to above 9 g/dL; (3) Vaso-active drugs as required and according to requirement (octreotide or terlipressin); and (4) Antibiotic prophylaxis was given in the form of I.V. cefotaxime 1 g/12 h for up to 5 d, unless acute infection, especially spontaneous bacterial peritonitis, was diagnosed, which was treated accordingly.

Urgent endoscopy was performed within the first six hours. The diagnosis of variceal hemorrhage was considered when active bleeding from an esophageal or gastric varix was observed or when a sign of recent bleeding, such as a "white nipple", was observed. Variceal hemorrhage was inferred when varices were the only pathology found, with blood present in the stomach. Patients with other causes of acute upper gastrointestinal bleeding were not considered for this analysis.

Esophageal varices were managed by band ligation or injection sclerotherapy if banding was not feasible. Bleeding from gastric varices was managed by injection of tissue adhesive (histoacryl) injection.

Variceal bleeding was considered to have been controlled if the following criteria were met: Stable blood pressure (no reduction in systolic pressure exceeding 20 mmHg, once the blood pressure had stabilized); a stable hemoglobin concentration (> 9 g/dL), measured twice daily; and a hematocrit above 30% (measured hourly during the first 12 h and every 2 to 6 h thereafter, depending on the patient's hemodynamic status), with a transfusion requirement of no more than two units in a 2-h period and fewer than four units within the first 4 h after endoscopic therapy.

Patients were followed for 5 d after the control of acute bleeding for occurrence of rebleeding. Rebleeding was defined as the occurrence of new hematemesis or melena after a period of 24 h or more from the 24-h point of stable vital signs and hematocrit/hemoglobin following an episode of acute bleeding^[9]. In patients with significant rebleeding, defined as frank hematemesis, new onset of melena, fresh blood in nasogastric tube aspirate, or hemodynamic compromise, with a decrease in hemoglobin level of > 2 g/dL, re-endoscopy was performed.

The CTP class, MELD, ALBI, MELD-Acute variceal bleeding (MELD-AVB) and PALBI scores were calculated from admission labs, and were correlated with control of bleeding, rebleeding, and in-hospital mortality. (1) The CTP score was calculated numerically as previously described including bilirubin, albumin, international normalized ratio (INR), and presence and grade of ascites and encephalopathy. CTP class was A if the score was 5-6, B if the score was 7-9, and C if the score was 10 or higher^[2]; (2) The MELD score was calculated as: $0.957 \times \log_e(\text{creatinine mg/dL}) + 0.378 \times \log_e(\text{bilirubin mg/dL}) + 1.120 \times \log_e(\text{INR}) + 0.643$ ^[5]; (3) The ALBI score was calculated as: $=(\log_{10} \text{bilirubin} \times 0.66) + (\text{albumin} \times -0.085)$ where bilirubin is in $\mu\text{mol/L}$ and albumin in g/L. ALBI was categorized into three grades: ALBI-1 (≤ -2.60), ALBI-2 (> 2.60 to -1.39), ALBI-3 (> -1.39)^[6]; (4) MELD-AVB was adapted from the basic MELD score through this equation $\text{logit}, -5.312 + 0.207 \times \text{MELD}$; bootstrapped R^2 , 0.3295; and (5) PALBI score was calculated as: $(2.02 \times \text{Log}_{10} \text{bilirubin}) + [-0.37 \times (\text{Log}_{10} \text{bilirubin})^2] + (-0.04 \times \text{albumin}) + (-3.48 \times \text{Log}_{10} \text{platelets}) + [1.01 (\text{Log}_{10} \text{platelets})^2]$ where bilirubin is in $\mu\text{mol/L}$ and albumin in g/L, and platelet count in $1000/\mu\text{L}$. PALBI was categorized as: PALBI 1 (score ≤ 2.53), PALBI 2 (score > 2.53 and ≤ 2.09), and PALBI 3 (score > 2.09)^[7].

The areas under the receiving-operator characteristic curve (AUROC) were calculated for survival and rebleeding.

Statistical analysis

Data were collected and entered into the computer using SPSS program version 23 for statistical analysis (IBM Corp., Armonk, NY, United States). Continuous data were expressed as the mean \pm SD and the median with minimum and maximum. Categorical data were expressed as the frequency (percentage). A comparison between the chosen variables and control of bleeding, rebleeding, and mortality was performed. Univariate analysis included the Fisher's exact test or Chi-square test for categorical variables and the analysis of variance for continuous variables. Multivariate logistic regression using the stepwise selection method was performed starting from the variables with $P < 0.01$ in the univariate analysis. ROC curve analyses were performed to determine the value of CTP class, MELD, ALBI, MELD-AVB and PALBI scores in predicting the in-hospital mortality and the control of bleeding. AUROCs with 95% confidence intervals were calculated and compared. Also, the AUROCs were tested for significance using DeLong test. The sensitivity, specificity, positive likelihood ratio, negative likelihood ratio, positive predictive value, and negative predictive value with 95% confidence intervals were reported at the best cutoff value. $P < 0.05$ was considered statistically significant.

RESULTS

This analysis contains the data of 1517 patients who presented with acute variceal bleeding. There were 1176 males (77.5%), and mean patient age was 52.6 ± 10.3 years.

Three hundred and thirty-two (21.9%) patients died during hospitalization, and 364 (23.9%) developed hepatic encephalopathy during admission.

All patients received resuscitation, blood as needed, antibiotics, and vasoactive drugs. Endoscopy was performed as soon as feasible, and the mean time between admission and endoscopy was 5.6 ± 1.9 h (median 4 h) (Table 1). The mean number of blood units transfused was 2.8 ± 1.1 units, and the mean duration of hospitalization was 5 d. The baseline characteristics of the patients on admission are shown in Table 1.

Endoscopy revealed that 1283 patients (84.57%) had esophageal varices as the source of bleeding. Seven hundred and eighty-five patients (51.74%) had band ligation alone; 301 patients (19.84%) had sclerotherapy as a single treatment modality, while 197 (12.9%) had combined treatment modalities. Gastric varices were present in 366 patients, and were the only source of bleeding in 165. Histoacryl injection was conducted in 271 patients, in 165 as the only therapeutic intervention and with band ligation of esophageal varices in 106. Sixty-nine patients (4.54%) received conservative medical treatment (Table 2).

Sixty-nine patients were CTP class A (4.5%), 434 were CTP class B (29.2 %) and 1014 were CTP class C (66.8%). 692 (46%) had ascites and 364 (24%) presented with hepatic encephalopathy. Twenty-six patients were ALBI grade 1 (1.6%); 669 were ALBI grade 2 (44.1%), and 822 were ALBI grade 3 (54.1%). Using the PALBI score to classify patients, 306 patients were PALBI grade 1 (20.2%), 285 patients were PALBI grade 2 (18.8%), and 926 patients were PALBI grade 3 (61%). Table 3 shows the patient classification using the different scores.

The incidence of early rebleeding (within 1 wk) was 3.12% and the incidence of recurrent bleeding (after 1 wk) was 8.9%. The predictors of rebleeding are shown in Table 4.

In-hospital mortality among CTP classes A, B and C were 0%, 11% and 28%, among ALBI grades 1, 2 and 3 were 0%, 7.2% and 34.5%, and among PALBI grade 1, 2, and 3 were 0%, 21.8% and 34.4%, respectively (Table 3).

Table 5 shows the predictors of mortality, where PALBI grade showed the highest odds ratio.

The AUROC for predicting survival following acute variceal bleeding was 0.668, 0.689, 0.803, 0.849 and 0.871 for CTP, MELD, ALBI, MELD-AVB and PALBI scores, respectively (Figure 1). PALBI score showed a significantly higher performance than MELD, ALBI and CTP (Table 6). For predicting rebleeding, the AUROC was 0.681, 0.74, 0.766, 0.769 and 0.794 for CTP, MELD, ALBI, MELD-AVB and PALBI scores, respectively (Figure 2). The performance of PALBI was significantly higher than that of MELD-AVB, MELD, ALBI and CTP (Table 6).

DISCUSSION

The present study found a mortality rate of 21.9% due to variceal bleeding. Randomized, controlled trials have shown that mortality due to variceal bleeding in cirrhosis has decreased over the past 3 decades from about 50% to 20%-30%, but this figure is still remarkably high. Hence, stratifying the risk for mortality is paramount^[10]. The best method to stratify risk is not clear. In this study we report the important predictive value of the PALBI score.

The PALBI score, proposed by Roayaie *et al*^[8] was initially used for the assessment of patients with HCC undergoing resection or ablation. It was later validated in several studies in predicting the outcome of interventions for the management of HCC^[11,12]. This report, however, evaluates the predictive power of the PALBI score in a large cohort of patients with cirrhosis presenting with acute variceal bleeding, in comparison to ALBI, MELD and CTP scores. In the overall analysis, PALBI score performed significantly better than ALBI, MELD score and CTP classification.

The performance of CTP and MELD scores in predicting survival of patients with upper gastrointestinal bleeding is comparable^[13,14]. The better performance of ALBI in the current study was most probably due to omitting the subjective factors (ascites and encephalopathy) in the CTP score. The MELD score also includes creatinine that may be affected temporarily during gastrointestinal tract bleeding. Additionally, the INR results, as noted by Lisman *et al*^[15] showed great variation among seven different European laboratories and Trotter *et al*^[16] also confirmed such a variation in INR levels among 14 different laboratories in the United States. However, the adapted version

Table 1 Characteristics of all studied participants

Variable	Number/mean (%)	Median	SD
Age (yr)	52.6	52	10.3
Male:Female	1176 (77.5):341 (22.5)	-	-
In-hospital mortality	332 (21.9)	-	-
Serum albumin (g/dL)	2.4	2.4	0.6
Serum total bilirubin (μmol/L)	64.98	30.78	99.18
Serum creatinine (mg/dL)	1.3	1	0.97
International normalized ratio (INR)	1.7	1.6	0.53
Hemoglobin level (g/dL)	9.5	9.4	2.35
Hematocrit	28.8	29	6.25
Platelet count ($\times 10^3/\text{mm}^3$)	116	104	56
Blood units transfused	2.8	2	1.1
Time to first endoscopy (h)	5.6	4	1.9
Duration of admission (d)	5	4	3.3
MELD score	15.2	14.8	3.2

SD: Standard deviation; MELD: Model of End-stage Liver Disease.

for variceal bleeding showed significantly better performance compared with the basic version.

The performance of PALBI, an updated version of ALBI, in predicting mortality was significantly better than the ALBI score possibly because the PALBI score includes platelet count which reflects the effect of portal hypertension; the main cause of acute variceal bleeding. In other scenarios as testing the prognostic performance after radiofrequency ablation, ALBI has been proved to be superior to PALBI^[17]. Thus, it appears that ALBI has a better prognostic power in patients with minimal liver dysfunction, whereas PALBI appears to be of more value in stratifying risk for portal hypertension.

The present study is the first to evaluate the performance of the PALBI score in predicting in-hospital mortality after variceal bleeding. The performance of the ALBI score in predicting the in-hospital mortality of acute upper gastrointestinal bleeding was previously tested by Zou *et al*^[18]. Their estimated AUROC result for the ALBI score was in agreement with the current study. However, they found the performance of ALBI comparable to the CTP and MELD scores. However, it is worth noting that in their study the cohort was much smaller than the present cohort (631 *vs* 1517) and unlike the current study not all patients underwent endoscopy and hence the source of bleeding was not restricted to variceal bleeding.

The incidence of early rebleeding during hospitalization (within 1 wk) among our patients was 3.12%. Several studies reported overall rebleeding rates of 2.56%, 3.2%, 6.1%, 3.9%^[19-23]. PALBI was significantly better than MELD, CTP and ALBI in predicting rebleeding. If the PALBI score could replace the CTP score and MELD, it would be easier and quicker to identify candidates for an early transjugular intra-hepatic porto-systemic shunt procedure. Furthermore, the use of PALBI score allows for better survival stratification within a CTP class. PALBI is a purely objective score incorporating both liver function and portal hypertension and is not subject to the inconsistencies of CTP resulting from the inclusion of ascites and encephalopathy.

Despite the limitation of being a retrospective single-center study, the large number of patients with all patients undergoing endoscopy and bleeding confirmed to be from varices, provides further validity to our findings.

In conclusion, the PALBI score is a simple, objective score that may be a good option for predicting in-hospital rebleeding and mortality in patients with acute variceal bleeding. We recommend future prospective studies to further validate the PALBI score and its value in predicting long-term prognosis.

Table 2 Treatment modalities

Treatment modality	n (%)
Band ligation	785 (51.8)
Sclerotherapy	301 (19.9)
Fundal varix (histoacryl injection alone)	165 (10.8)
Combined treatment	197 (13)
Conservative treatment	69 (4.5)

Table 3 Patients distribution according to Child Turcotte Pugh, Albumin-bilirubin, Model of End-stage Liver Disease and Platelet-albumin-bilirubin scores

Class	No. of patients (%)	Bleeding mortality (%)
CTP A	69 (4.5)	0
CTP B	434 (29.2)	11
CTP C	1014 (66.8)	28
MELD < 12	186 (12)	0
MELD 12-20	353 (23.5)	16.3
MELD > 20	978 (64.5)	31.9
ALBI grade 1	26 (1.7)	0
ALBI grade 2	669 (44.1)	7.2
ALBI grade 3	822(54.1)	34.5
MELD-AVB < 11	159 (10.5)	0
MELD-AVB 11-19	370 (24.4)	14.3
MELD-AVB > 19	988 (65.1)	36.3
PALBI grade 1	306 (20.2)	0
PALBI grade 2	285(18.8)	21.8
PALBI grade 3	926 (61)	34.4

PALBI: Platelet-albumin-bilirubin score; ALBI: Albumin-bilirubin; MELD: Model of End-stage Liver Disease; CTP: Child Turcotte Pugh; MELD-AVB: MELD-acute variceal bleeding.

Table 4 Predictors of rebleeding

Parameter	OR	95%CI	P value
PALBI	3.987	1.994-7.775	< 0.001
Conservative therapy	2.473	1.237-4.822	< 0.001
Previous beta blockers	0.387	0.194-0.755	< 0.001
ALBI	3.214	1.607-6.267	< 0.01
Time to endoscopy	1.572	1.086-3.065	< 0.01
Ascites	2.319	1.160-4.522	< 0.05
Spleen size	1.961	1.181-3.824	< 0.05
Diabetes	1.631	1.116-3.180	< 0.05
Platelet count	0.445	0.223-0.868	< 0.05
INR	1.618	1.109-3.155	< 0.05
Bilirubin	1.981	1.291-3.863	< 0.05
Albumin	0.613	0.457-0.897	< 0.05

ALBI: Albumin-bilirubin; PALBI: Platelet-albumin-bilirubin score; INR: International normalized ratio.

Table 5 Multivariate analysis: Predictors of in-hospital mortality

Parameter	OR	95%CI	P value
PALBI	4.187	2.093-8.164	< 0.001
MELD-AVB	3.842	2.983-7.431	< 0.01
ALBI	3.153	1.576-6.148	< 0.001
MELD	2.981	1.490-5.812	< 0.001
CTP	2.144	1.072-4.180	< 0.01
Bilirubin	1.981	1.015-3.862	< 0.01
Albumin	0.269	0.134-0.524	< 0.01
Age	1.718	1.034-3.350	< 0.05
Ascites	1.547	1.173-3.016	< 0.05
Encephalopathy	1.869	1.124-3.644	< 0.05
Hematocrit	0.191	0.095-0.372	< 0.05
Platelet count	0.543	0.321-0.853	< 0.05
INR	1.923	1.061-3.749	< 0.05
Creatinine	1.561	1.180-3.043	< 0.05

PALBI: Platelet-albumin-bilirubin score; ALBI: Albumin-bilirubin; MELD: Model of End-stage Liver Disease; CTP: Child Turcotte Pugh; MELD-AVB: MELD-acute variceal bleeding; INR: International normalized ratio.

Table 6 Area under the receiver operating characteristic curve values of different scores to predict the outcome of acute variceal bleeding and the incidence of rebleeding

Predicting outcome of acute variceal bleeding					
Score	CTP	MELD	ALBI	MELD-AVB	PALBI
AUROC	0.668	0.689	0.803	0.849	0.871
P value ¹	< 0.01	< 0.01	< 0.01	0.043	-
Predicting rebleeding					
Score	CTP	MELD	ALBI	MELD-AVB	PALBI
AUROC	0.681	0.74	0.766	0.769	0.794
P value ¹	< 0.01	< 0.05	0.052	0.032	-

¹P values are tested with Delong test between each consecutive test. PALBI: Platelet-albumin-bilirubin score; ALBI: Albumin-bilirubin; MELD: Model of End-stage Liver Disease; CTP: Child Turcotte Pugh; MELD-AVB: MELD-acute variceal bleeding; INR: International normalized ratio.

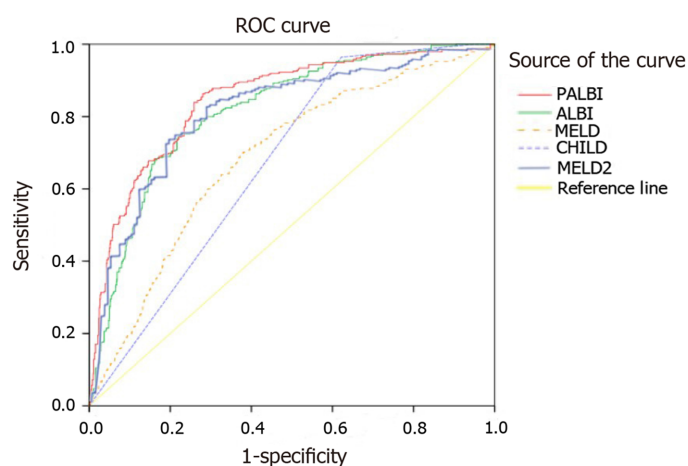


Figure 1 The AUROCs of the platelet-albumin-bilirubin score, albumin-bilirubin, Model of End-stage Liver Disease, Child Turcotte Pugh and Model of End-stage Liver Disease-acute variceal bleeding scores for predicting outcome of acute variceal bleeding. PALBI: Platelet-albumin-bilirubin score; ALBI: Albumin-bilirubin; MELD: Model of End-stage Liver Disease; CTP: Child Turcotte Pugh; MELD-AVB: MELD-acute variceal bleeding.

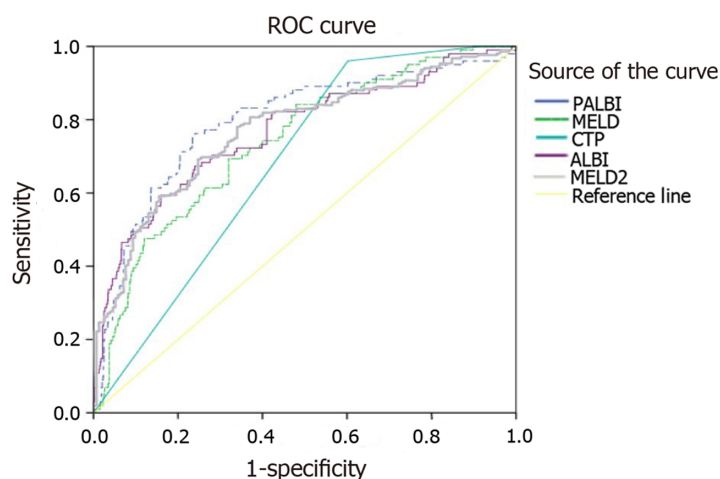


Figure 2 The AUROCs of the platelet-albumin-bilirubin score, albumin-bilirubin, Model of End-stage Liver Disease, Child Turcotte Pugh and Model of End-stage Liver Disease-acute variceal bleeding scores for predicting rebleeding. PALBI: Platelet-albumin-bilirubin score; ALBI: Albumin-bilirubin; MELD: Model of End-stage Liver Disease; CTP: Child Turcotte Pugh; MELD-AVB: MELD-acute variceal bleeding.

ARTICLE HIGHLIGHTS

Research background

In 1964, the CTP score was proposed to assess patients with portal hypertension related to gastrointestinal tract bleeding and was used later to assess patients with cirrhosis in general. Many doubts have been raised recently regarding the performance of the CTP score as a subjective scoring system with abrupt points and overlapping parameters. Recently, new scoring systems such as the ALBI and PALBI scores were proposed which use linear predictive equations to overcome the disadvantages of the CTP score.

Research motivation

To identify a reliable prognostic score to predict the short-term outcome of patients with acute variceal bleeding.

Research objectives

Scoring systems with more specific parameters and using linear predictive equations showed better performance than subjective point-based scoring systems.

Research methods

We retrospectively analyzed the data of a large number of patients with acute variceal bleeding and their short-term outcome.

Research results

The PALBI score is a simple, objective score that is considered a good option for predicting in-hospital rebleeding and mortality in patients with acute variceal bleeding in comparison to other scoring systems. However, we still recommend performing a prospective study to better analyze the performance of these scoring systems.

Research conclusions

The PALBI score could be used to predict the short-term outcome and the incidence of rebleeding in patients with acute variceal bleeding.

Research perspectives

Performing a prospective large-scale multicenter study to test the performance of these scores in different management settings.

REFERENCES

- 1 Lebrech D. Life, death, and varices. *Gut* 2001; **49**: 607-608 [PMID: 11600456 DOI: 10.1136/gut.49.5.607]
- 2 Pugh RN, Murray-Lyon IM, Dawson JL, Pietroni MC, Williams R. Transection of the oesophagus for bleeding oesophageal varices. *Br J Surg* 1973; **60**: 646-649 [PMID: 4541913 DOI: 10.1002/bjs.1800600817]
- 3 van Dam GM, Gips CH, Reisman Y, Maas KW, Purmer IM, Huizenga JR, Verbaan BW. Major clinical events, signs and severity assessment scores related to actual survival in patients who died from primary biliary cirrhosis. A long-term historical cohort study. *Hepatology* 1999; **46**: 108-115 [PMID: 10228773]

- 4 **Shetty K**, Rybicki L, Carey WD. The Child-Pugh classification as a prognostic indicator for survival in primary sclerosing cholangitis. *Hepatology* 1997; **25**: 1049-1053 [PMID: [9141416](#) DOI: [10.1002/hep.510250501](#)]
- 5 **Malinchoc M**, Kamath PS, Gordon FD, Peine CJ, Rank J, ter Borg PC. A model to predict poor survival in patients undergoing transjugular intrahepatic portosystemic shunts. *Hepatology* 2000; **31**: 864-871 [PMID: [10733541](#) DOI: [10.1053/he.2000.5852](#)]
- 6 **Kamath PS**, Wiesner RH, Malinchoc M, Kremers W, Therneau TM, Kosberg CL, D'Amico G, Dickson ER, Kim WR. A model to predict survival in patients with end-stage liver disease. *Hepatology* 2001; **33**: 464-470 [PMID: [11172350](#) DOI: [10.1053/jhep.2001.22172](#)]
- 7 **Johnson PJ**, Berhane S, Kagebayashi C, Satomura S, Teng M, Reeves HL, O'Beirne J, Fox R, Skowronska A, Palmer D, Yeo W, Mo F, Lai P, Iñarrairaegui M, Chan SL, Sangro B, Miksad R, Tada T, Kumada T, Toyoda H. Assessment of liver function in patients with hepatocellular carcinoma: a new evidence-based approach-the ALBI grade. *J Clin Oncol* 2015; **33**: 550-558 [PMID: [25512453](#) DOI: [10.1200/JCO.2014.57.9151](#)]
- 8 **Roayaie S**, Jibara G, Berhane S, Tabriz-ian P, Park JW, Yang J, Yan L, Han G, Izzo F, Chen M, Blanc JF, Kudo M, Roberts LR, Sherman M, Johnson P. 851 PALBI-An Objective Score Based on Platelets, Albumin Bilirubin Stratifies HCC Patients Undergoing Resection & Ablation Better than Child's Classification. *Hepatology* 2015; **62**: 624A-690A [DOI: [10.1002/hep.28219](#)]
- 9 **Tripathi D**, Stanley AJ, Hayes PC, Patch D, Millson C, Mehrzad H, Austin A, Ferguson JW, Olliff SP, Hudson M, Christie JM; Clinical Services and Standards Committee of the British Society of Gastroenterology. U.K. guidelines on the management of variceal haemorrhage in cirrhotic patients. *Gut* 2015; **64**: 1680-1704 [PMID: [25887380](#) DOI: [10.1136/gutjnl-2015-309262](#)]
- 10 **Fortune BE**, Garcia-Tsao G, Ciarleglio M, Deng Y, Fallon MB, Sigal S, Chalasani NP, Lim JK, Reuben A, Vargas HE, Abrams G, Lewis MD, Hassanein T, Trotter JF, Sanyal AJ, Beavers KL, Ganger D, Thuluvath PJ, Grace ND, Grossmann RJ; Vapreotide Study Group. Child-Turcotte-Pugh Class is Best at Stratifying Risk in Variceal Hemorrhage: Analysis of a US Multicenter Prospective Study. *J Clin Gastroenterol* 2017; **51**: 446-453 [PMID: [27779613](#) DOI: [10.1097/MCG.0000000000000733](#)]
- 11 **Hansmann J**, Evers MJ, Bui JT, Lokken RP, Lipnik AJ, Gaba RC, Ray CE. Albumin-Bilirubin and Platelet-Albumin-Bilirubin Grades Accurately Predict Overall Survival in High-Risk Patients Undergoing Conventional Transarterial Chemoembolization for Hepatocellular Carcinoma. *J Vasc Interv Radiol* 2017; **28**: 1224-1231.e2 [PMID: [28688815](#) DOI: [10.1016/j.jvir.2017.05.020](#)]
- 12 **Liu PH**, Hsu CY, Hsia CY, Lee YH, Chiou YY, Huang YH, Lee FY, Lin HC, Hou MC, Huo TI. ALBI and PALBI grade predict survival for HCC across treatment modalities and BCLC stages in the MELD Era. *J Gastroenterol Hepatol* 2017; **32**: 879-886 [PMID: [27696519](#) DOI: [10.1111/jgh.13608](#)]
- 13 **Peng Y**, Qi X, Dai J, Li H, Guo X. Child-Pugh versus MELD score for predicting the in-hospital mortality of acute upper gastrointestinal bleeding in liver cirrhosis. *Int J Clin Exp Med* 2015; **8**: 751-757 [PMID: [25785053](#)]
- 14 **Chalasani N**, Kahi C, Francois F, Pinto A, Marathe A, Bini EJ, Pandya P, Sitaraman S, Shen J. Model for end-stage liver disease (MELD) for predicting mortality in patients with acute variceal bleeding. *Hepatology* 2002; **35**: 1282-1284 [PMID: [11981782](#) DOI: [10.1053/jhep.2002.32532](#)]
- 15 **Lisman T**, van Leeuwen Y, Adelmeijer J, Pereboom IT, Haagsma EB, van den Berg AP, Porte RJ. Interlaboratory variability in assessment of the model of end-stage liver disease score. *Liver Int* 2008; **28**: 1344-1351 [PMID: [18482269](#) DOI: [10.1111/j.1478-3231.2008.01783.x](#)]
- 16 **Trotter JF**, Olson J, Lefkowitz J, Smith AD, Arjal R, Kenison J. Changes in international normalized ratio (INR) and model for endstage liver disease (MELD) based on selection of clinical laboratory. *Am J Transplant* 2007; **7**: 1624-1628 [PMID: [17511686](#) DOI: [10.1111/j.1600-6143.2007.01822.x](#)]
- 17 **Ho SY**, Liu PH, Hsu CY, Chiou YY, Su CW, Lee YH, Huang YH, Lee FY, Hou MC, Huo TI. Prognostic Performance of Ten Liver Function Models in Patients with Hepatocellular Carcinoma Undergoing Radiofrequency Ablation. *Sci Rep* 2018; **8**: 843 [PMID: [29339752](#) DOI: [10.1038/s41598-018-19251-y](#)]
- 18 **Zou D**, Qi X, Zhu C, Ning Z, Hou F, Zhao J, Peng Y, Li J, Deng H, Guo X. Albumin-bilirubin score for predicting the in-hospital mortality of acute upper gastrointestinal bleeding in liver cirrhosis: A retrospective study. *Turk J Gastroenterol* 2016; **27**: 180-186 [PMID: [27015623](#) DOI: [10.5152/tjg.2016.15502](#)]
- 19 **Khan A**, Mannan F, Din Ru. Outcome of endoscopic band ligation for oesophageal variceal bleed in patients with chronic liver disease. *Gomal J Med Sci* 2013; **11**: 84-87
- 20 **Kuran S**, Oğuz D, Parlak E, Asil M, Çiçek B, Kiliç M, Dişibeyaz S, Sahin B. Secondary prophylaxis of esophageal variceal treatment: Endoscopic sclerotherapy, band ligation and combined therapy--long-term results. *Turk J Gastroenterol* 2006; **17**: 103-109 [PMID: [16830291](#)]
- 21 **Petrascu F**, Grothaus J, Mössner J, Schiefke I, Hoffmeister A. Differences in bleeding behavior after endoscopic band ligation: a retrospective analysis. *BMC Gastroenterol* 2010; **10**: 5 [PMID: [20074379](#) DOI: [10.1186/1471-230X-10-5](#)]
- 22 **Hameed K**, Saeed A, Khan Y, Khan IM, Javed M. Endoscopic band ligation: an ideal procedure to manage bleeding oesophageal varices. *Ann Pak Inst Med Sci* 2007; **3**: 234-237
- 23 **de Franchis R**; Baveno VI Faculty. Expanding consensus in portal hypertension: Report of the Baveno VI Consensus Workshop: Stratifying risk and individualizing care for portal hypertension. *J Hepatol* 2015; **63**: 743-752 [PMID: [26047908](#) DOI: [10.1016/j.jhep.2015.05.022](#)]



Published By Baishideng Publishing Group Inc
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA
Telephone: +1-925-3991568
E-mail: bpgoffice@wjgnet.com
Help Desk: <https://www.f6publishing.com/helpdesk>
<https://www.wjgnet.com>

