**Baveno VI Recommendation on Avoidance of Screening Endoscopy in Cirrhotic Patients: Are We There Yet?**

Mário Jorge Silva\(^a,\,b\) Carlos Bernardes\(^a\) João Pinto\(^a\) Rafaela Loureiro\(^a\) Pedro Duarte\(^a\) Milena Mendes\(^a\) Filipe Calinas\(^a\)

\(^a\)Gastroenterology Department, Hospital de Santo António dos Capuchos, Centro Hospitalar de Lisboa Central, and \(^b\)Chronic Diseases Research Center (CEDOC) – Integrated Pathophysiological Mechanisms Research Group, Faculdade de Ciências Médicas, NOVA Medical School, Lisbon, Portugal

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**Abstract**

**Introduction:** Recent studies assessed the predictive value of liver transient elastography, combined or not with platelet count, for the presence of esophageal varices in patients with liver cirrhosis, and multiple cutoffs have been proposed. The Baveno VI consensus states that patients with compensated advanced chronic liver disease, liver stiffness <20 kPa, and a platelet count >150,000 have a very low risk of having varices requiring treatment and can avoid screening endoscopy. We aimed to validate this recommendation in a cohort of cirrhotic patients. **Methods:** Retrospective analysis of all patients evaluated at the Gastroenterology Department (Centro Hospitalar de Lisboa Central) between September 2009 and October 2015 with a liver stiffness (FibroScan®) compatible with liver cirrhosis as well as upper endoscopy and blood tests within 12 months from elastography. Patients on propranolol ≥80 mg/day or carvedilol ≥12.5 mg/day, as well as those with previous variceal bleeding, variceal endoscopic treatments, or cirrhosis decompensations were excluded. We validated the new Baveno VI recommendation and explored alternative cutoffs. **Results:** Ninety-seven patients were analyzed, 76.3% (74/97) male, mean age 54.3 ± 11.2 years. Most patients (55.7%) had no varices and 14.4% had varices requiring treatment. Most patients (78.4%) had cirrhosis related to chronic hepatitis C. If the new Baveno VI recommendation had been applied to this cohort, upper endoscopy would have been avoided in 11.3% (11/97) of patients, none of them with esophageal varices requiring treatment: specificity 100%, sensitivity 13.3%, positive predictive value 100%, and negative predictive value 16.3% for absence of varices requiring treatment. If screening endoscopy had been avoided in those patients with liver stiffness <30 kPa and platelet count ≥120,000, endoscopy would have been avoided in 27.8% (27/97) of patients, none of whom with esophageal varices requiring treatment: specificity 100%, sensitivity 32.5%, positive predictive value 100%, and negative predictive value 20% for absence of varices requiring treatment. **Conclusions:** The new Baveno VI criteria identified compensated cirrhotic patients without varices requiring treatment in whom screening endoscopy could have been avoided safely. Further studies are needed to confirm these findings and potentially explore more ambitious but still safe cutoffs for those criteria.
Recomendação de Baveno VI Sobre Evicção da Endoscopia de Rastreio em Doentes com Cirrose: Já Lá Estamos?

Palavras Chave
Cirrose hepática · Consenso · Endoscopia · Hipertensão portal · Varizes esofágicas e gástricas · Técnicas de imagem por elasticidade

Resumo
Introdução: Estudos recentes avaliaram o valor preditivo da elastografia hepática transitória, combinada ou não com contagem plaquetária, para a presença de varizes esofágicas em doentes com cirrose hepática, e foram propostos vários valores de corte. O consenso de Baveno VI afirma que doentes com doença hepática crónica avançada compensada, elastografia hepática <20 kPa e >150,000 plaquetas têm muito baixo risco de varizes com necessidade de tratamento, podendo evitar endoscopia de rastreamento. Pretendemos validar esta recomendação numa coorte de doentes cirróticos. Métodos: Análise retrospectiva dos doentes avaliados no Serviço de Gastrenterologia (Centro Hospitalar de Lisboa Central) entre Setembro 2009 e Outubro 2015 com elastografia hepática (FibroScan®) compatível com cirrose hepática e análises num intervalo até 12 meses desde a elastografia. Doentes sob propranolol ≥80 mg/dia ou carvedilol ≥12.5 mg/dia, assim como aqueles com antecedentes de hemorragia variceal, tratamento endoscópico de varizes ou descompensação de cirrose foram excluídos. Validadamos a nova recomendação de Baveno VI e explorámos valores de corte alternativos. Resultados: Noventa e sete doentes foram analisados, 76.3% (74/97) homens, idade média 54.3 ± 11.2 anos. A maioria dos doentes (55.7%) não tinha varizes, 14.4% tinha varizes com indicação para tratamento. A maioria dos doentes (78.4%) tinha cirrose relacionada com hepatite C. Se a nova recomendação de Baveno VI tivesse sido aplicada nesta coorte, a endoscopia de rastreio teria sido evitada em 11.3% (11/97) dos doentes, nenhum com varizes com indicação para tratamento: especificidade 100%, sensibilidade 13.3%, valor preditivo positivo 100% e valor preditivo negativo 16.3% para ausência de varizes com indicação para tratamento. Se a endoscopia de rastreio fosse evitada nos doentes com elastografia hepática <30 kPa e >120,000 plaquetas, teria sido evitada em 27.8% (27/97) destes doentes, nenhum com varizes com indicação para tratamento: especificidade 100%, sensibilidade 32.5%, valor preditivo positivo 100% e valor preditivo negativo 20% para ausência de varizes com indicação para tratamento. Conclusão: Os novos critérios de Baveno VI identificaram doentes com cirrose compensada e sem varizes com indicação para tratamento, em que se podia ter evitado com segurança a endoscopia de rastreio. São necessários mais estudos para confirmar estes resultados e explorar valores de corte mais ambiciosos, mas seguros.

Introduction

Portal hypertension is a progressive complication of liver cirrhosis, and varices and variceal bleeding are its most direct consequences [1]. Gastroesophageal varices are present in approximately 50% of cirrhotic patients, and their existence correlates with severity of disease. The most important complication of varices is variceal bleeding, occurring in 25–40% of patients with cirrhosis [2] and associated with around 15% mortality [3]. The risk of variceal bleeding depends on several factors, including size and appearance of varices and cirrhosis stage [1]. Therefore, patients with varices with high-risk features should be identified, so that treatment (primary prophylaxis of variceal bleeding) may be initiated.

Until recently, all clinical guidelines recommended that every cirrhotic patient should be screened for varices with the gold standard – upper gastrointestinal endoscopy [1, 4], and a recent guideline on noninvasive tests for liver disease highlighted that upper endoscopy cannot be replaced by noninvasive methods [5]. Nevertheless, several studies on the prediction of clinically significant portal hypertension (CSPH) with noninvasive methods have been performed recently, and the report from the Baveno VI Consensus Workshop (held on April 2015) states that patients with compensated advanced chronic liver disease with a liver stiffness <20 kPa and a platelet count >150,000 have a very low risk of having varices requiring treatment and can avoid screening endoscopy [6].

We aimed to validate this new Baveno VI recommendation in a cohort of cirrhotic patients.
Materials and Methods

We performed a retrospective, single-center analysis of all patients evaluated at the Gastroenterology Department of the Centro Hospitalar de Lisboa Central between September 2009 and October 2015 with liver stiffness compatible with liver cirrhosis (>12.5 kPa) and upper endoscopy as well as platelet count performed within 12 months from liver stiffness evaluation. All evaluations were performed at the Centro Hospitalar de Lisboa Central (liver stiffness evaluation and upper gastrointestinal endoscopy at the Gastroenterology Department, platelet count at the Central Laboratory). Liver stiffness was evaluated with transient elastography (FibroScan® model 502, Echosens®, Paris, France), performed by experienced users, and validated according to current recommendations. Upper endoscopy was performed by experienced endoscopists, and esophageal varices were classified as small (comprising grade I/small varices) or large (comprising grade II/medium and grade III/large varices) [1]. Patient records were reviewed for cirrhosis etiology and stage, as well as past and current treatments.

Patients on treatment for portal hypertension (propranolol ≥80 mg/day or carvedilol ≥12.5 mg/day), as well as patients with previous variceal bleeding or treatments (such as banding or sclerotherapy) were excluded. Decompensated cirrhotic patients (with ascites, hepatic encephalopathy, Child-Pugh class B, or Child-Pugh class C) were excluded as well.

Esophageal varices requiring treatment were defined, according to Baveno VI, as small varices with red signs or large varices [6]. Statistical analysis was performed with Microsoft Office Excel® 2010 and STATA® v12.1. Descriptive univariate analysis was performed and is reported as means and standard deviations for continuous variables and proportions for categorical variables.

Results

A total of 97 patients were analyzed, 76.3% (74/97) male, with a mean age of 54.3 ± 11.2 years at liver elastography. Most patients (55.7%) had no varices, 29.9% had small varices without red signs, 1.0% had small varices with red signs, and 13.4% had large varices – therefore, 14.4% (14 patients) had varices requiring treatment. All patients were Child-Pugh class A, and most (78.4%) had cirrhosis related to chronic hepatitis C. Population characteristics are detailed in Table 1.

If the new Baveno VI recommendation had been applied to this cohort of cirrhotic patients, upper endoscopy would have been avoided in 11.3% (11/97) of patients, none of them with esophageal varices requiring treatment (2 patients had small varices without red signs and 9 patients had no varices). Upper endoscopy would have been performed in 88.7% (86/97) of patients – 52.3% (56/86) of whom had no varices – and would have identified all patients with varices requiring treatment.

The Baveno VI criteria had a specificity of 100% (95% CI 76.8–100%), a sensitivity of 13.3% (95% CI 6.81–22.5%), a positive likelihood ratio of 0.87 (95% CI 0.80–0.94), a positive predictive value of 100% (95% CI 71.5–100%), and a negative predictive value of 16.3% (95% CI 9.2–25.8%) for absence of varices requiring treatment.

From this cohort, it is possible to generate alternative criteria based on liver elastography and platelet count to identify patients without varices requiring treatment. If screening endoscopy had been avoided in those patients with liver stiffness <30 kPa and platelet count ≥120,000,
endoscopy would have been avoided in 27.8% (27/97) of patients, none of whom with esophageal varices requiring treatment (3 patients had small varices without red signs and 24 patients had no varices). Screening endoscopy would have been performed in 61.9% patients and would have identified all patients in the cohort with varices requiring treatment.

This new criteria (avoiding screening endoscopy when liver stiffness <30 kPa and platelet count ≥120,000) in this derivation cohort had a specificity of 100% (95% CI 76.8–100%), a sensitivity of 32.5% (95% CI 22.6–43.7%), a positive predictive value of 100% (95% CI 87.2–100%), and a negative predictive value of 20.0% (95% CI 11.4–31.3%) for absence of varices requiring treatment.

Discussion

There is a clear potential of noninvasive methods to identify cirrhotic patients with low risk of having varices requiring treatment and that therefore could avoid unpleasant screening endoscopies – relatively invasive procedures with associated risks and costs.

Liver stiffness has been shown to correlate with hepatic venous pressure gradient (HVPG) [7]. Many studies assessed the predictive value of liver elastography (alone [8] or in combination with other noninvasive measures) [9, 10] and platelet count [11–13] for CSHP. Nevertheless, several different surrogate markers of CSHP have been used in those studies (i.e., HVPG, presence of varices, presence of large varices), and their results may not be directly comparable.

In general, proposed cutoffs for liver stiffness predict presence and size of esophageal varices with high sensitivities but rather unsatisfactory specificities [5]. In fact, it is probable that liver stiffness performs much better at ruling out varices than ruling them in. Correlation between liver stiffness and HVPG is excellent for HVPG values lower than 10–12 mm Hg, but is suboptimal for higher HVPG values [14]. This is explained by the fact that with cirrhosis progression, the mechanisms of portal hypertension become less dependent on the intrahepatic resistance to portal flow due to tissue fibrosis (measured by liver elastography) and progressively more dependent on extrahepatic factors, such as hyperdynamic circulation and splanchnic vasodilation [5]. Therefore, liver stiffness may probably perform better at identifying patients without varices (i.e., with HVPG below 10–12 mm Hg) [1] than at identifying patients with large varices, or varices with red signs, the latter presumably having HVPG values that do not correlate well with liver stiffness values.

The 2015 Baveno VI Consensus Workshop published the first consensus recommendation on avoidance of screening endoscopy in patients with compensated advanced liver disease in an era where evidence in this field was probably not robust enough. The concept of “compensated advanced chronic liver disease” is, per se, innovative, and may be suspected based on liver transient elastography evaluation: values between 10 and 15 kPa are suggestive and values >15 kPa are highly suggestive of compensated advanced chronic liver disease. Furthermore, this new concept and its definition are independent of the etiology of liver disease [6].

In our study, we found that Baveno VI criteria identified correctly patients without varices requiring treatment and would have allowed safe avoidance of screening endoscopy in 11% of this cohort of compensated cirrhotic patients. Nevertheless, robust data on this issue are lacking. It is possible that less stringent criteria will allow to safely avoid a greater proportion of screening endoscopies in low-risk patients in the future, rendering this management strategy even more cost-effective. The new criteria generated from this cohort are speculative and not generalizable, but demonstrate that further cutoffs may be explored.

Our study has several limitations, and our results should be interpreted carefully. This was a single-center, retrospective study, based on liver stiffness measures and endoscopies performed by different operators (although all of them were experienced). Liver stiffness measures were based on single determinations and not confirmed, as recently proposed [6]. A liver stiffness cutoff ≥12.5 kPa [15] was used to define liver cirrhosis irrespective of the etiology of liver disease. Indeed, alternative cutoffs for nonviral liver diseases are insufficiently validated [5], and even for alcohol-related liver disease, presumably associated with higher transient elastography values [5], the proposed cutoff for cirrhosis is 12.5 kPa [16]. Furthermore, the Baveno VI criteria do not recommend differential cutoffs for diagnosis of “compensated advanced chronic liver disease” according to the etiology of liver disease [6].

These Baveno VI criteria performed similarly well (negative predictive value 100%) in some studies [17–19], but failed to identify few patients with varices requiring treatment in other ones (negative predictive value 91.2–98%) [20, 21]. It is important to note that prevalence of varices varied between studies, which were therefore prone to spectrum bias.
In this study, we did not account for the diagnosis of other unsuspected diseases (as gastric antral vascular ectasia, peptic ulcer, or gastric cancer) and surrogate markers of CSPH (as small esophageal varices) in screening endoscopies, which obviously would have a significant impact on patient management and outcomes.

Further studies, with larger sample sizes, less heterogeneous cirrhosis etiologies, and preferentially prospective design, are needed to confirm these findings and potentially explore more robust and cost-effective cutoffs for liver stiffness and platelet count.

### Conclusions

The Baveno VI criteria based on liver stiffness (<20 kPa) and platelet count (>150,000) identified correctly compensated cirrhotic patients without varices requiring treatment in this cohort (in whom screening endoscopy could have been safely avoided), but allowed for avoidance of endoscopy only in a small proportion of patients.

### Statement of Ethics

This study did not require informed consent nor review/approval by the appropriate ethics committee.

### Disclosure Statement

The authors have no conflicts of interest to declare.

### References