



presence of any one of: varices on endoscopy, portal hypertensive bleeding, or porto-systemic collaterals on imaging. Univariate analysis was used to examine variables that can predict varices and CSPH. Multivariate model was constructed using a logistic regression analysis of statistically significant variables in univariate analysis. A sequential-testing algorithm was developed using the best performing model. **Results:** Of the 44 patients, 28 (64%) had varices and 35 (79.5%) had CSPH. LSM was higher in patients with varices (12.7 kPa vs 6 kPa, $p < 0.01$) and CSPH (12 kPa vs 5.4 kPa, $p < 0.01$). Platelet count was lower in patients with varices (71 vs 151, $p < 0.01$) and CSPH (84 vs 144, $p = 0.03$). Multivariate analysis combining LSM, and platelet count predicts varices (AUROC 0.82 ± 0.07 , $p < 0.01$) and CSPH (AUROC 0.86 ± 0.07 , $p < 0.01$). A step-wise algorithm combining LSM of 10 kPa and platelet of $80 \times 10^9 /L$ was developed. (Figure) This algorithm performed with specificity of 81%, negative predictive value (NPV) of 81% and positive predictive value (PPV) of 89% to detect varices ($p < 0.001$). The same model performed with sensitivity of 77%, specificity of 89% and PPV of 96% to detect CSPH ($p < 0.001$). Accuracy of the model for predicting varices and CSPH was 93% and 80%, respectively. **Conclusion:** We developed a simple model by combining LSM with platelet count that can be used to identify CSPH in PSVD patients. This noninvasive model can predict the risk of varices and aid in deciding the need for endoscopic surveillance or initiating therapy for portal hypertension. However, this model requires further validation in a larger independent cohort.

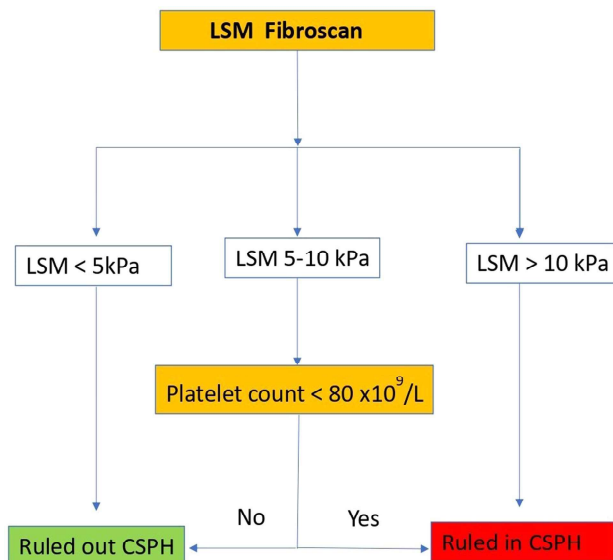


Figure: Step-wise decision algorithm

Disclosures: The following people have nothing to disclose: Harish Gopalakrishna, Maria Mironova, Nehna Abdul Majeed, Asif Ali Hitawala, Shani Scott, Jaha Norman-Wheeler, David E Kleiner, Christopher Koh, Theo Heller

3133-A | ACCURACY OF A DEDICATED 100 HZ VIBRATION-CONTROLLED SPLEEN STIFFNESS MEASUREMENT VERSUS BAVENO CRITERIA FOR THE DETECTION OF VARICES IN PATIENTS WITH COMPENSATED CIRRHOSIS

Angelo Armandi^{1,2}, Tiziana Sanavia², Emma Vanderschueren^{3,4}, Georg Semmler⁵, Antonio Liguori⁶, Salvatore Petta⁷, Maurice Michel¹, Merle Marie Werner¹, Talal Merzian¹, Christian Labenz¹, Mathias Jachs⁸, Mattias Mandorfer⁵, Wim Laleman^{3,9}, Luca Miele⁶, Thomas Reiberger¹⁰ and Jörn M. Schattenberg¹, (1)University of Mainz, (2)University of Turin, (3)University Hospitals Leuven, (4)Catholic University of Leuven, Department of Chronic Diseases, Metabolism and Aging (CHROMETA), Leuven, Belgium, (5)Division of Gastroenterology and Hepatology, Department of Medicine III, Medical University of Vienna, Vienna, Austria, (6)Università Cattolica Di Roma, Department of Internal Medicine, Fondazione Policlinico a. Gemelli, (7)Sezione Di Gastroenterologia, Dipartimento Promozione Della Salute, Materno-Infantile, Di Medicina Interna e Specialistica Di Eccellenza "G. D'alessandro", Università Di Palermo, Palermo, Italy, (8)Medical University of Vienna, (9)Catholic University of Leuven, Leuven, Belgium, (10)Division of Gastroenterology and Hepatology, Department of Internal Medicine III, Medical University of Vienna

Background: Clinically significant portal hypertension (CSPH) marks a critical step in the natural history of compensated advanced chronic liver disease (cACLD) and may lead to esophageal varices (EV). The Baveno VI criteria suggest liver stiffness measurement (LSM) and platelet count (PLT) for non-invasive identification of cACLD patients not requiring screening gastroscopy. We investigated the accuracy of the novel 100 Hz vibration-controlled transient elastography-based spleen stiffness measurement (SSM) exam for the identification of EV in cACLD patients. **Methods:** Retrospective study of Mainz, Vienna, Leuven, Rome and Palermo. Patients with cACLD of any etiology ($LSM \geq 10kPa$ or histological F4 fibrosis), but without previous decompensation (bleeding, encephalopathy, ascites) were included. SSM and LSM were obtained using Fibroscan F630 ≤ 1 month within screening gastroscopy. Prediction performance between different SSM cut-offs with respect to the Baveno criteria ($LSM > 20kPa$ and/or $PLT < 150 G/L$) were compared by logistic regression with 10-fold cross-validation, adjusted for age, gender, BMI, transaminases, INR, albumin, and bilirubin. Backward feature selection based on likelihood ratio test was applied to identify significant confounders. Performance was calculated by

Symbols: ♦, Poster of Distinction; ★, Foundation Award Recipient

balanced accuracy (BA), specificity (SP) and sensitivity (SE). Wilcoxon test was used to evaluate significant performance improvement of SSM cut-offs with respect to Baveno criteria, or to significant confounders. **Results:** 343 cACLD patients with a median age of 59 years (60.3% male) and NAFLD as the main etiology (51.3%) were included. 137 had EV with 49 high-risk EV (HR-EV), while median SSM, LSM and PLT were 40.5kPa, 21kPa and 139 G/L, respectively. The figure shows BAs at different SSM cut-offs, compared to Baveno (red line). The best overall performance with all-type EV was at SSM=60 kPa (BA=0.72, SP=0.86, SE=0.58); Baveno: BA=0.66, SP=0.86, SE=0.39. Comparing HR-EV vs. absence of EV, the best cut-off was at 50kPa (BA=0.71, SP=0.95, SE=0.47; Baveno: BA=0.56, SP=0.92, SE=0.29). These SSM thresholds significantly improved BA when significant confounders were considered. **Conclusion:** The novel spleen-dedicated 100 Hz SSM is associated with presence of EV in cACLD patients. In both all-type EV and HR-EV, SSM showed better accuracy than the Baveno LSM-PLT criteria, achieving a better trade-off between SP and SE.

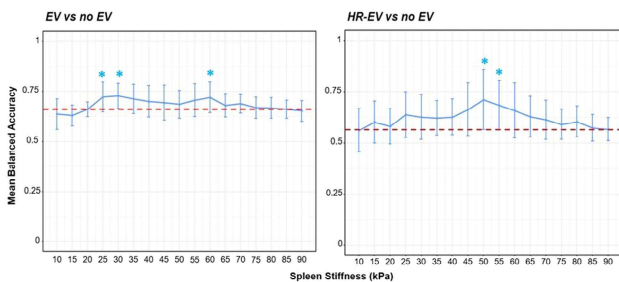


Figure 1. Mean Balanced Accuracy at different Spleen Stiffness cut-offs. Dashed red line: Baveno performance. The asterisk indicates that the SSM models showed a balanced accuracy significantly higher (i.e. Wilcoxon test p-value >5%) with respect to both the same model including only the selected confounders and Baveno.

Disclosures: Wim Laleman – Cook Medical, CSL Behring, Norgine: Speaking and Teaching, No, No; Cook Medical, Boston Scientific, CSL Behring: Consultant, No, No; Boston Scientific: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), No, No; Thomas Reiberger – AbbVie: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), Yes, Yes; Boehringer Ingelheim: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), Yes, No; Gilead: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), Yes, No; Intercept: Grant/Research Support (research funding from ineligible

companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), No, Yes; MSD: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), No, Yes; Myr Pharmaceuticals: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), Yes, Yes; Philips Healthcare: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), No, Yes; Pliant: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), No, Yes; Siemens: Grant/Research Support (research funding from ineligible companies should be disclosed by the principal or named investigator even if that individual's institution receives the research grant and manages the funds), Yes, No; AbbVie: Consultant, Yes, Yes; Boehringer Ingelheim: Consultant, Yes, No; Gilead: Consultant, Yes, Yes;

The following people have nothing to disclose: Angelo Armandi, Tiziana Sanavia, Emma Vanderschueren, Georg Semmler, Antonio Liguori, Salvatore Petta, Maurice Michel, Merle Marie Werner, Talal Merizian, Christian Labenz, Mathias Jachs, Mattias Mandorfer, Luca Miele, Jörn M. Schattenberg

3134-A | AN INTERNATIONAL SURVEY ON PRACTICE PATTERNS FOR THE MANAGEMENT OF GASTRIC VARICES

Lolwa Al-Obaid¹, Mohammad Bilal², Katarzyna M. Pawlak³, Nadeem Tehami⁴, Diogo De Moura⁵, Rashid Ns Luj⁶, Jayanta Samanta⁷, Aymen Almuhaideb⁸, Andres Rodriguez Parra⁹, Andres Cardenas¹⁰, Marvin Ryou¹¹ and Ahmad Najdat Bazarbashi¹, (1)Washington University in St. Louis, (2)University of Minnesota, (3) Samodzielny Publiczny Zaklad Opieki Zdrowotnej Ministerstwa Spraw Wewnętrznych i Administracji w Szczecinie, (4)University Hospital Southampton NHS Foundation Trust, (5)Universidade De Sao Paulo Hospital Das Clinicas Da Faculdade De Medicina De Ribeirao Preto, (6)The Chinese University of Hong Kong, (7)Postgraduate Institute of Medical and Educational Research, (8)King Faisal Specialist Hospital and Research Center, (9)Hospital General Dr Manuel Gea Gonzalez, (10)Barcelona Clinic, Barcelona, Spain, (11)Brigham and Women's Hospital

Symbols: ♦, Poster of Distinction; ★, Foundation Award Recipient