

assessed by vibration-controlled transient elastography (VCTE). Scores (FIB-4, Agile-3+, Agile-4) were calculated from biochemical and clinical data. Diagnostic accuracy for detecting advanced fibrosis ($\geq F3$) and cirrhosis (F4) was evaluated using ROC curves and Youden index.

Results: Median age was 59 years; 60% were men. Median BMI was 33.3 kg/m²; 69.6% had type 2 diabetes. Median liver stiffness was 9.1 kPa; 29.9% had advanced fibrosis, and 10.5% cirrhosis. Agile-4 outperformed VCTE stiffness in predicting advanced fibrosis (AUROC 0.765, $p=0.037$) and demonstrated superior accuracy for cirrhosis (AUROC 0.875, $p=0.003$) (Figure 1). The optimal cut-offs for Agile-4 were ≥ 0.159 (rule out cirrhosis with 90% sensitivity) and ≥ 0.366 (rule in cirrhosis with 90% specificity).

Conclusions: In this Latin American MASLD cohort, Agile-4 score demonstrated superior noninvasive rule-out performance for advanced fibrosis and cirrhosis. Incorporating these thresholds into VCTE algorithms could reduce unnecessary biopsies and improve streamline MASLD care pathways.

Conflict of interest: Yes, receives support from the Chilean government through the Fondo Nacional de Desarrollo Científico y Tecnológico (FONDECYT 1241450).

Figure 1. Area Under the Curve Performance of VCTE with AGILE 3, and AGILE 4 for predicting Advanced fibrosis and Cirrhosis

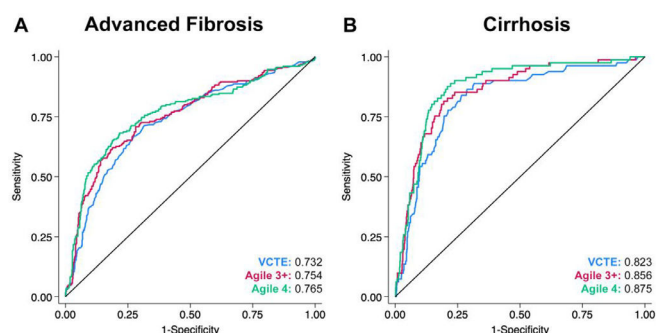


Figure. Receiver-operating characteristic curves comparing the performance of vibration-controlled transient elastography (VCTE) liver stiffness measurements with Agile-3+ and Agile-4 scores in predicting (A) advanced fibrosis and (B) cirrhosis.

<https://doi.org/10.1016/j.aohep.2025.101969>

#139

EXPLORING THE ROLE OF METABOLIC DYSFUNCTION IN ALCOHOL-ASSOCIATED HEPATITIS: A GLOBAL STUDY

María Ignacia Pérez Garayar¹, Vania Cari Gormaz¹, Ignacio Téllez¹, Francisco Idalsoaga², Gene Im³, Bastian Alcayaga¹, Muzzafar Haque⁴, Stephanie Rutledge³, Hanna Blaney⁵, Pojsakorn Danpanichkul⁶, Arun Valsan⁷, Gowripriya Nair⁷, Gustavo Ayares⁸, Renata Farías⁸, Jorge Arnold⁸, Pedro Acuña⁸, Kaanthi Rama⁹, Carlos Esteban Coronel-Castillo¹⁰, María Ayala-Valverde¹¹, Carolina Ramirez-Cadiz¹², Vinay Jahagirdar¹³, Winston Dunn¹⁴, Heer Mehta¹⁴, Maria Poca¹⁵, German Soriano¹⁵, Berta Cuyas¹⁵, Joaquin Cabezas¹⁶, Victor Echavarría¹⁶, Meritxell Ventura Cots¹⁷, Juan G. Abalde¹⁸, Mustafa Al-Karaghoul¹⁸, Lubomir Skladany¹⁹, Daniel J. Havaj¹⁹, Karolina Sulejova¹⁹, Svetlana Adamcova Selcanova¹⁹, Prasun K. Jalal²⁰,

Mohamed A. Elfeki²⁰, Mohamad Ali Ibrahim²¹, Katherine Maldonado²², Juan Pablo Roblero²³, Daniela Simian²³, José Antonio Velarde-Ruiz²⁴, Jacqueline Córdova-Gallardo²⁵, Fátima Higuera-de-la-Tijera²⁶, Rita Silva²⁷, Cristina Melo Rocha²⁸, Roberta C. Araujo²⁹, Gustavo Henrique Pereira³⁰, Fernando Bessone³¹, Mario Tanno³¹, Ayelen Kisch³², Manuel Mendizabal³³, Sebastián Marciano³⁴, Gonzalo Gomez Perdiguerro³⁴, Pedro Montes³⁵, Patricia Guerra Salazar³⁶, Geraldine Ramos³⁶, Enrique Carrera³⁷, Kristina R. Chacko³⁸, Nyingi Kemmer³⁹, Saurabh Agrawal³⁹, Luciana Lofego Goncalves⁴⁰, Oluwatosin Oguntoye⁴¹, Douglas Simonetto⁴², Arun J. Sanyal⁷, Rohit Loomba⁴³, Vijay H. Shah⁴⁴, Patrick S. Kamath⁴⁴, Marco Arrese⁸, Ramon Bataller⁴⁵, Luis Antonio Díaz⁸, Juan Pablo Arab⁸

¹ School of Medicine. Faculty of Medicine. Pontificia Universidad Católica de Chile.

² Department of Medicine. Division of Gastroenterology. Western University. London Health Sciences Center. London. Ontario. Canada.

³ Center for Liver Disease and Transplantation. Columbia University Vagelos College of Physicians and Surgeons, USA.

⁴ Department of Internal Medicine. College of Medicine. University of Saskatchewan, Canada.

⁵ MedStar Georgetown University Hospital. Medstar Transplant Hepatology Institute, USA.

⁶ Department of Internal Medicine. Texas Tech University Health Sciences Center, USA.

⁷ Department of Gastroenterology. Hepatology Division. Amrita Institute of Medical Sciences and Research Centre, India.

⁸ Department of Gastroenterology. School of Medicine. Pontificia Universidad Católica de Chile.

⁹ Division of Gastroenterology. Hepatology. and Nutrition. Department of Internal Medicine. Virginia Commonwealth University School of Medicine. Richmond, USA.

¹⁰ Department of Gastroenterology. Hospital General de México "Dr. Eduardo Liceaga", Chile.

¹¹ Internal Medicine Service. Hospital El Pino, Chile.

¹² Department of Anesthesiology. Virginia Commonwealth University School of Medicine, USA.

¹³ Division of Gastroenterology. Hepatology. and Nutrition. Department of Internal Medicine. Virginia Commonwealth University School of Medicine.

¹⁴ Division of Gastroenterology. Department of Medicine. University of Kansas Medical Center, USA.

¹⁵ Department of Gastroenterology. Hospital de la Santa Creu i Sant Pau. Institut de Recerca Sant Pau (IR SANT PAU). Universitat Autònoma de Barcelona. CIBERehd, España.

¹⁶ Gastroenterology and Hepatology Department. University Hospital Marqués de Valdecilla, España.

¹⁷ Liver Unit. Hospital Vall D'Hebron. Universitat Autònoma de Barcelona. CIBERehd, España.

¹⁸ Division of Gastroenterology. Liver Unit. University of Alberta. Edmonton, Canada.

¹⁹ Division of Hepatology. Gastroenterology. and Liver Transplantation. Department of Internal Medicine II. Slovak Medical University. F. D. Roosevelt University Hospital, Slovak Republic.

²⁰ Department of Gastroenterology and Hepatology. Baylor College of Medicine, USA.

²¹ Department of Medicine. University of South Dakota Sanford School of Medicine. Sioux Falls, USA.

²² Division of Gastroenterology. Hepatology and Nutrition. Department of Medicine. University of Louisville School of Medicine. Louisville, USA.

²³ Gastroenterology Section. Clinical Hospital University of Chile. School of Medicine. University of Chile.

²⁴ Civil Hospital of Guadalajara Fray Antonio Alcalde. Guadalajara, México.

²⁵ University Center of Health Sciences. University of Guadalajara. Guadalajara, México.

²⁶ Division of Gastroenterology. Hepatology. and Nutrition. Department of Internal Medicine. Virginia Commonwealth University School of Medicine. Richmond, USA.

²⁷ Faculty of Medicine. National Autonomous University of Mexico.

²⁸ Liver Transplant Unit of the Base Hospital of the Faculty of Medicine of São José do Rio Preto, Brasil.

²⁹ Fhaj Foundation Hospital Adriano Jorge, Brasil.

³⁰ Hospital das Clínicas da Faculdade de Medicina de Ribeirão Preto, Brasil.

³¹ Federal Hospital of Bonsucesso, Brasil.

³² Provincial Hospital del Centenario, Argentina.

³³ Gastroenterology Hospital "Dr. Carlos Bonorino Udaondo", Argentina.

³⁴ Hepatology and Liver Transplant Unit. Austral University Hospital, Argentina.

³⁵ Italian Hospital Buenos Aires, Argentina.

³⁶ National Hospital Daniel Alcides Carrión - Callao, Peru.

³⁷ Bolivian-Japanese Gastroenterology Institute, Bolivia.

³⁸ Eugenio Espejo Specialty Hospital, Ecuador.

³⁹ Division of Hepatology. Montefiore Medical Center. Bronx, USA.

⁴⁰ Division of Gastroenterology. Tampa General Hospital. Tampa, USA.

⁴¹ Gastroenterology Service. University Hospital - Federal University of Espírito Santo, Brasil.

⁴² Division of Gastroenterology and Hepatology. Mayo Clinic, USA.

⁴³ MASLD Research Center. Division of Gastroenterology and Hepatology. University of California San Diego, USA.

⁴⁴ Division of Gastroenterology and Hepatology. Mayo Clinic. Rochester, USA.

⁴⁵ Liver Unit. Hospital Clinic, España.

Introduction and Objectives: Severe alcohol-associated hepatitis (AH) carries high mortality. Although the role of cardiometabolic risk factors (CMRF)—including obesity, type 2 diabetes mellitus (T2DM), hypertension (HTN), and dyslipidemia (DLP)—has been characterized in steatotic liver disease, their role in the severity of AH remains unclear.

To evaluate the impact of CMRF on mortality and infection risk in AH.

Materials and Methods: Multinational prospective cohort study (2015–2024) including hospitalized patients with severe AH across 24 centers in 14 countries (Global AlCHep Network). Diagnosis of AH was done using NIAAA criteria. Analyses included competing-risk models, with liver transplantation as a competing risk. Models were adjusted by age, sex, ethnicity, history of cirrhosis, CMRF, corticosteroids use, MELD, and ACLF grade.

Results: 935 participants were included. Median BMI was 24.2kg/m², prevalence of T2DM was 21%, HTN 17%, DLP 7%. In adjusted competing-risk models, age (sHR 1.02, 95%CI: 1.01–1.04; p<0.001), MELD

(sHR 1.04, 95%CI: 1.01–1.06; p<0.001), infections (sHR 1.76, 95%CI: 1.28–2.41; p<0.001), and ACLF grade 2 (sHR 1.67, 95%CI: 1.05–2.69; p<0.032) and 3 (sHR 3.06, 95%CI: 1.88–4.99; p<0.001) were associated with higher risk of mortality, while obesity (sHR 0.67, 95%CI: 0.48–0.93; p=0.016) and corticosteroids use (sHR 0.67, 95%CI: 0.49–0.92; p=0.014) were associated with lower mortality. T2DM, HTN and DLP weren't associated with higher mortality.

Conclusions: Metabolic dysfunction was not associated with increased mortality in AH. Although obesity may be a protective factor, these findings could be explained by a better nutritional status than the lean population.

Conflict of interest: None

<https://doi.org/10.1016/j.aohep.2025.101970>

#143

CHARACTERIZATION OF THE UNRECORDED ALCOHOL USE WORLDWIDE: A SYSTEMATIC REVIEW AND SURVEY-BASED STUDY

Katherine Emilia Maldonado Cardona¹, Hanna Blaney², Deepika Devuni³, Muzzafer Haque⁴, Pojsakorn Danpanichkul⁵, Francisco Idalosoaga⁶, Vinay Jahagirdar⁷, Kaanthi Rama⁷, Natalia Baeza⁶, Frank Murray⁸, Patrick Kamath⁹, Ramon Bataller¹⁰, Marco Arrese⁶, Jeffrey Lazarus¹¹, Luis Antonio Díaz¹², Juan Pablo Arab⁷

¹ Clínica de gastroenterología Intera, Guatemala.

² MedStar Georgetown University Hospital. Medstar Transplant Hepatology Institute, USA.

³ Division of Gastroenterology and Hepatology. UMass Chan Medical School, USA.

⁴ Department of Internal Medicine. College of Medicine. University of Saskatchewan, Canada.

⁵ Department of Medicine. Texas Tech University Health Sciences Center, USA.

⁶ Departamento de Gastroenterología. Escuela de Medicina. Pontificia Universidad Católica de Chile.

⁷ Division of Gastroenterology. Hepatology. and Nutrition. Department of Internal Medicine. Virginia Commonwealth University School of Medicine, USA.

⁸ Department of Medicine. Royal College of Surgeons in Ireland. Beaumont Hospital, Ireland.

⁹ Division of Gastroenterology and Hepatology. Mayo Clinic College of Medicine and Science, USA.

¹⁰ Liver Unit. Hospital Clinic, Barcelona.

¹¹ Barcelona Institute for Global Health (ISGlobal), España.

¹² MASLD Research Center. Division of Gastroenterology and Hepatology. University of California San Diego, USA.

Introduction and Objectives: Unrecorded alcohol - products that escape taxation, regulation, and safety checks - represents up to one quarter of world alcohol intake and is strongly linked to hazardous drinking and alcohol-related liver disease. Knowledge gaps regarding unrecorded alcohol worldwide need to be addressed to better inform region-specific harm reduction strategies.

To characterize the population, contemporary consumption patterns, and physicians' interest in unrecorded alcohol worldwide.

Materials and Methods: Cross-sectional survey-based study. Data was collected between August and November 2024, distributing a 19 item electronic questionnaire to hepatology-focused physicians worldwide. Responses were categorized into 15 geographic regions and were analyzed by descriptive statistics.