

¹¹ Department of Internal Medicine. Hamilton Medical Center. Medical College, USA.

¹² Stravitz-Sanyal Institute for Liver Disease and Metabolic Health. Division of Gastroenterology, Hepatology, and Nutrition. Department of Internal Medicine. Virginia Commonwealth University School, USA.

¹³ Arizona Liver Health, USA.

¹⁴ Department of Pathological Anatomy. Institute of Gastroenterology. University of Medical Sciences of Havana, Cuba.

¹⁵ Department of Medicine. Schulich School of Medicine. Western University & London Health Sciences Centre. London, UK.

¹⁶ Velocity Clinical Research, USA.

¹⁷ Division of Hepatology. Avera McKennan Hospital & University Health Center, USA.

¹⁸ Department of Medicine. Division of Gastroenterology, Hepatology, and Nutrition. University of Louisville, USA.

¹⁹ Beatty Liver and Obesity Research Program. Inova Health System, USA.

²⁰ Center for Outcomes Research in Liver Disease, USA.

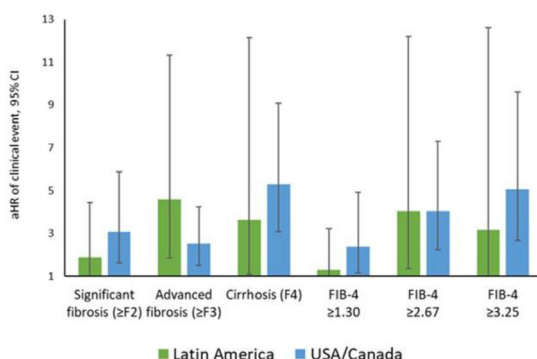
Introduction and Objectives: MASLD is highly prevalent worldwide. We evaluated performance of NITs and predictors of outcomes in patients with MASLD from Latin America (LA) as compared to North America (NA).

Patients and Methods: The Global-MASLD project enrolled MASLD patients with liver biopsies and NITs (FIB-4, liver stiffness measurement (LSM) by transient elastography). NITs' performance to predict advanced fibrosis (AF=F3-F4) and outcomes was assessed.

Results: A total of 3,904 MASLD patients were included [N=892 from 5 LA countries (Argentina, Brazil, Chile, Cuba, Mexico) and N=3012 from NA (USA/Canada)]. MASLD patients from LA were older, had lower BMI (obesity 64% vs. 85%), more lean MASLD (5.6% vs. 2.7%), more T2D (49% vs. 38%) ($p<0.001$) but similar rates of AF ($p=0.56$). Clinico-demographic predictors of AF included older age and T2D ($p<0.05$). The NIT accuracy was lower in LA-MASLD than NA-MASLD: AUC (95% CI) of FIB-4 0.75 (0.71-0.79) vs. 0.81 (0.79-0.83), LSM 0.73 (0.67-0.80) vs. 0.78 (0.75-0.81), Agile-3+ 0.76 (0.70-0.82) for both LA and NA. Sensitivity of 80% (low-risk, screening cutoff) was achieved with FIB-4 ≥ 1.01 in LA vs. FIB-4 ≥ 1.17 in NA; specificity of 95% (high-risk, diagnostic cutoff) with FIB-4 ≥ 2.35 vs. FIB-4 ≥ 2.40 . In adjusted (age, sex, T2D) proportional hazards models, fibrosis severity by histology or NITs was associated with adverse outcomes (death, decompensation, HCC) in both groups (adjusted hazard ratios (aHR) >1.0) (Figure).

Conclusions: MASLD patients from LA have more T2D but less obesity than NA. Common NITs have lower accuracy in LA-MASLD. Histologic and NIT stage of fibrosis are independent predictors of adverse outcomes in both groups.

Conflict of interest: None



<https://doi.org/10.1016/j.aohep.2025.101951>

#127

ACCELERATED PROGRESSION TO CIRRHOSIS AND HEPATIC DECOMPENSATION IN METALD AND ALD COMPARED TO MASLD: A GLOBAL STUDY

Luis Antonio Díaz Piga¹, Xiao-Dong Zhou², Natalia Baeza³, Francisco Idalsoaga⁴, Gustavo Ayares³, Terry Cheuk-Fung Yip⁵, Vincent Wai-Sun Wong⁵, Grace Lai-Hung Wong⁵, Jimmy Che-To Lai⁵, Rakhi Maiwall⁶, Shiv K. Sarin⁶, Yu Jun Wong⁷, Xin En Goh⁷, May Xuan Goh⁸, David Marti-Aguado⁹, Cristiane Villela-Nogueira¹⁰, Ana Carolina Cardoso¹⁰, Natalia Balassiano Wajsbrot¹⁰, Nathalie Leite¹⁰, Gil Fernando Salles¹⁰, Claudia Regina Lopes Cardoso¹⁰, Sebastián Marciano¹¹, Jorge Martínez Morales¹¹, María José Acosta¹¹, Adrian Gadano¹¹, Claudia P. Oliveira¹², Mario G. Pessoa¹², Adelina Lozano¹³, Anand V. Kulkarni¹⁴, Ramagundam Ramyasri¹⁴, Mohamed El-Kassas¹⁵, Mario Reis Alvares-da-Silva¹⁶, Bruno Basso¹⁶, Gabriella Jonko¹⁶, Fátima Higuera de la Tijera¹⁷, Daniza Contreras¹⁸, José Velarde-Ruiz Velasco¹⁹, Alceo Galimberti²⁰, Fernando Bessone²⁰, Claudia Alves Couto²¹, Rafael Theodoro²², Mísia Joyner de Sousa Dias Monteiro²², Francisco Javier Valentin-Cortez²³, Alejandra Mijangos-Trejo²³, Norberto C. Chavez-Tapia²³, Iwona Popiolek²⁴, Michał Kukla²⁴, Ezequiel Ridruejo²⁵, Mirta Peralta²⁶, Ismael Yepes Barreto²⁷, Luis Molina Barrios²⁷, Argemiro Lara Díaz²⁷, Fernando Javier Barreyro²⁸, Juan Pablo Roblero²⁹, Daniela Simian²⁹, Pamela Gil²⁹, Sheel Patel³⁰, Ashwani K. Singal³⁰, Pedro Montes³¹, María Fernanda Saavedra-Chacón³², Alberto Jose García Gonzalez³³, Constanza D. Sabate³⁴, Manuel Mendizabal³⁴, Jordi Gratacós-Ginès³⁵, Elisa Pose³⁵, Johana Acuña³⁶, Angelo Z. Mattos³⁷, Igor Lima Ferraz³⁸, Roberta Chaves Araujo³⁸, Katrina Pekarska³⁹, Richard Parker³⁹, Katherine Marrugo³⁹, Alonso Vera Torres³⁹, Juan Diego Torres³⁹, Valentina Mejía Valdés³⁹, Mohamad Ali Ibrahim⁴⁰, Prasun K. Jalal⁴⁰, Graciela Castro-Narro⁴¹, Mazen Nouredin⁴², Naim Alkhouri⁴³, Winston Dunn⁴⁴, Patrick S. Kamath⁴⁵, Arun Sanyal⁴⁶, Veeral Ajmera⁴⁷, Richard Sterling⁴⁶, Rohit Loomba⁴⁷, Ramon Bataller⁴⁸, Ming-Hua Zheng⁴⁸, Marco Arrese⁴⁹, Juan Pablo Arab⁴⁶

¹ MASLD Research Center. Division of Gastroenterology and Hepatology. University of California San Diego, USA.

² China.

³ Pontificia Universidad Católica de Chile.

⁴ Pontificia Universidad Católica de Chile. Western University.

⁵ Chinese University of Hong Kong.

⁶ Institute of Liver and Biliary Sciences, India.

⁷ Changi General Hospital / NUS, Singapore.

⁸ National University of Singapore (NUS).

⁹ Clinic University Hospital. INCLIVA Health Research Institute, España.

¹⁰ Universidade Federal do Rio de Janeiro, Brasil.

¹¹ Hospital Italiano de Buenos Aires, Argentina.

- ¹² Universidade de São Paulo, Brasil.
¹³ Clínica Avendaño, Peru.
¹⁴ AIG Hospitals, India.
¹⁵ Helwan University, Egypt.
¹⁶ Hospital de Clínicas de Porto Alegre, Brasil.
¹⁷ Hospital General de México.
¹⁸ Instituto Nacional de Diabetes. Endocrinología y Nutrición (INDEN), República Dominicana.
¹⁹ Hospital Civil de Guadalajara, México.
²⁰ Hospital Centenario de Rosario, Argentina.
²¹ Universidade Federal de Minas Gerais, Brasil.
²² Hospital das Clínicas da Universidade Federal de Minas Gerais, Brasil.
²³ Medica Sur Hospital, México.
²⁴ Jagiellonian University Medical College, Poland.
²⁵ Centro de Educación Médica e Investigaciones Clínicas Norberto Quirno "CEMIC", Argentina.
²⁶ Hospital Francisco J. Muñiz, Argentina.
²⁷ Universidad de Cartagena, Colombia.
²⁸ Universidad Nacional de Misiones, Argentina.
²⁹ Universidad de Chile.
³⁰ University of Louisville, USA.
³¹ Hospital Daniel Alcides Carrión, Perú.
³² Clínica CES, Colombia.
³³ Universidad Central de Venezuela.
³⁴ Hospital Universitario Austral, Argentina.
³⁵ Hospital Clínic de Barcelona, España.
³⁶ Fundación Santa Fe de Bogotá, Colombia.
³⁷ Santa Casa de Porto Alegre, Brasil.
³⁸ Universidade de São Paulo. Ribeirão Preto, Brasil.
³⁹ Leeds Teaching Hospitals NHS Trust, UK.
⁴⁰ Baylor College of Medicine, USA.
⁴¹ INCMNSZ, México.
⁴² Houston Methodist Hospital, USA.
⁴³ Summit Clinical Research, USA.
⁴⁴ University of Kansas Medical Center, USA.
⁴⁵ Mayo Clinic, USA.
⁴⁶ Virginia Commonwealth University, USA.
⁴⁷ University of California San Diego, USA.
⁴⁸ MAFLD Research Center. Department of Hepatology. The First Affiliated Hospital of Wenzhou Medical University, China.

Introduction and Objectives: The natural history of MetALD remains poorly characterized. In a large global cohort, we compared the natural history of the main steatotic liver disease (SLD) subtypes in terms of liver fibrosis progression and hepatic decompensation.

Materials and Methods: Retrospective cohort study of adult participants with SLD (2003–2025), including MASLD, MetALD, and ALD according to the 2023 SLD criteria. Sociodemographic and clinical data, vibration-controlled transient elastography (VCTE) parameters, and liver biopsy (when available) were recorded. Other causes of liver disease were excluded. The primary outcome was progression to cirrhosis in those without cirrhosis at baseline (defined by 1. liver biopsy, or 2. VCTE ≥ 13.6 kPa, or Fibrosis-4 [FIB-4] > 3.25 if other techniques were missing). Secondary outcomes included incidence of hepatic decompensation (ascites, hepatic encephalopathy, variceal bleeding, or hepatorenal syndrome). A multivariable Cox regression adjusted by age, sex, race, body mass index, diabetes, hypertension, hyperlipidemia, and smoking (for HCC) was performed.

Results: The total cohort included 150,306 participants from 15 countries; 87.5% MASLD, 7.9% MetALD, and 4.6% ALD. Overall, the median age was 61 years [IQR 51–70]; 52.8% men, and 97.6% Asian. At baseline, 12.2% of the cohort had a liver biopsy with F4 or a VCTE/FIB-4 suggestive of cirrhosis. During a median follow-up of 2.1 years [IQR 0.6–4.7], 0.9% of participants progressed to cirrhosis and 0.4%

had hepatic decompensations. Individuals with MetALD and ALD exhibited a higher risk of progression to cirrhosis (MetALD aHR 1.34, 95%CI: 1.06–1.68, $p=0.013$; ALD aHR 1.82, 95%CI: 1.41–2.35, $p<0.0001$; MASLD: reference) and of hepatic decompensation (MetALD aHR 9.33, 95%CI: 6.32–13.75, $p<0.0001$; ALD aHR 20.56, 95%CI: 13.86–30.48, $p<0.0001$).

Conclusions: In this multi-ethnic global cohort, MetALD and ALD were associated with more rapid cirrhosis progression and greater decompensation rates than MASLD, independent of cardiometabolic factors.

Conflict of interest: None

Figure
Cumulative Incidence of Cirrhosis by SLD Subtype

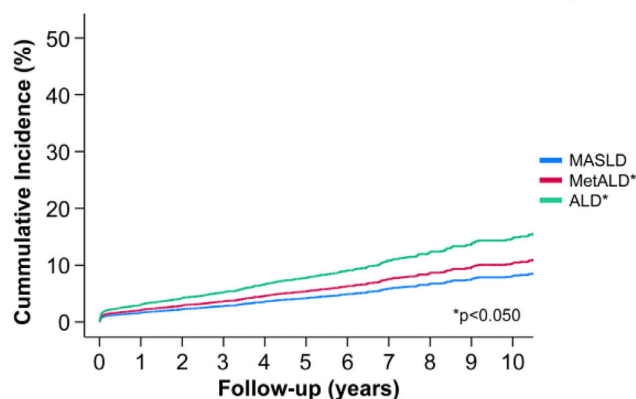


Figure. Cumulative incidence of incident cirrhosis by steatotic liver disease subtype. Participants with baseline cirrhosis—defined by (1) histologic stage F4, (2) liver stiffness ≥ 13.6 kPa on vibration-controlled transient elastography, or (3) Fibrosis-4 index > 3.25 —were excluded. Cox models were adjusted for age, sex, race, body mass index, diabetes, hypertension, and hyperlipidemia.

<https://doi.org/10.1016/j.aohep.2025.101952>

#14

FACTORS ASSOCIATED WITH HEALTH LITERACY IN PATIENTS DIAGNOSED WITH LIVER CIRRHOSIS IN COLOMBIA

Ismael de Jesus Yepes Barreto¹, Carlos Martelo², Nicole Chamorro²

¹ Universidad de Cartagena - Asociación Colombiana de Hepatología, Colombia.

² Universidad de Cartagena, Colombia.

Introduction and Objectives: Health literacy (HL) refers to a patient's ability to obtain, process, and understand medical information needed to make informed health decisions. Low HL is associated with increased healthcare costs, higher hospitalization rates, reduced access to transplantation, and increased mortality—especially among vulnerable populations. In patients with cirrhosis, HL has been linked to sex, education, employment, and disease etiology in other countries, but data are scarce in Latin America. This study aimed to identify factors associated with HL in patients with liver cirrhosis in Cartagena, Colombia.

Patients and Methods: We conducted a cross-sectional, analytical study between September and December 2024. Adults with a confirmed diagnosis of cirrhosis completed the validated Spanish version of the Short Assessment of Health Literacy (SAHL-S). Scores below 14 indicated inadequate HL. A separate, validated questionnaire assessed disease-specific knowledge across four domains: diagnosis, signs/symptoms, treatment, and medication.

Results: A total of 93 patients were analyzed (61.2% female; mean age: 63.9 $\pm$ 11.7 years). MASLD and cryptogenic cirrhosis were the