Fernando Bessone¹⁷, Mario Tanno¹⁷, Gustavo Romero¹⁸, Manuel Mendizabal¹⁹, Sebastián Marciano², Gonzalo Gomez²⁰, Melisa Dirchwolf²¹, Pedro Montes²², Patricia Guerra²³, Geraldine Ramos²³, Juan Carlos Restrepo²⁴, Enrique Carrera²⁵, Mayur Brahmania²⁶, Ashwani Singal²⁷, Ramón Bataller²⁸, Vijay Shah²⁹, Patrick S. Kamath²⁹, Marco Arrese¹, Juan Pablo Arab³⁰

- ¹ Department of Gastroenterology, Pontificia Universidad Católica de Chile, Santiago, Chile
- ² Gastroenterology Unit, Roosevelt Hospital, Ciudad de Guatemala, Guatemala
- ³ Servicio Medicina Interna, Hospital El Pino, Santiago, Chile
- ⁴ Sección Gastroenterología, Hospital Clínico Universidad de Chile, Santiago, Chile
- ⁵ Servicio de Gastroenterología, Hospital Sótero del Río, Santiago, Chile
- ⁶ Servicio de Gastroenterología, Clínica Hospital del Profesor, Santiago, Chile
- ⁷ Hospital Civil Guadalajara, Guadalajara, México
- ⁸ Hospital General Manuel Gea González, Mexico City, México
- ⁹ Servicio de Gastroenterología, Hospital General de México "Dr. Eduardo Liceaga, Mexico City, México 10 Hospital Publica Chibushus México
- ¹⁰ Hospital Dublán, Chihuahua, México
- ¹¹ Hospital Juárez de México, Mexico City, México
- ¹² Unidade de Transplante de Figado e do Hospital de Base, Faculdade de Medicina de São Jose do Rio Preto, São Paulo. Brasil
- ¹³ Fhaj Fundação Hospital Adriano Jorge, Amazonas, Brasil
- ¹⁴ Hospital das Clínicas da Faculdade de Medicina de Ribeirão Preto, Ribeirão Preto, Brasil
- ¹⁵ Hospital Federal de Bonsucesso, Rio de Janeiro, Brasil
- ¹⁶ Hospital das Clinicas Universidade Federal de Minas Gerais, Belo Horizonte, Brasil
- ¹⁷ Hospital Provincial del Centenario, Santa Fe, Argentina
- ¹⁸ Hospital de Gastroenterología " Dr. Carlos Bonorino Udaondo", Buenos Aires, Argentina
- ¹⁹ Hospital Universitario Austral, Buenos Aires, Argentina
- ²⁰ Hospital Italiano Buenos Aires, Buenos Aires, Argentina
- ²¹ Unidad de Trasplante Hepático, Hospital Privado de Rosario, Rosario, Argentina
- ²² Hospital Nacional Daniel Alcides Carrión Callao, Bellavista, Perú
- ²³ Instituto Gastroenterológico Boliviano- Japonés, Cochabamba. Bolivia
- ²⁴ Hospital Pablo Tobon Uribe, Medellin, Colombia
- ²⁵ Hospital Especialidades Eugenio Espejo, Quito, Ecuador
- ²⁶ Univeristy of Calgary, Alberta, Canadá
- ²⁷ Department of Medicine, University of South Dakota Sanford School of Medicine, South Dakota, Estados Unidos
- ²⁸ Liver Unit, Hospital Clinic, Barcelona, España
- ²⁹ Division of Gastroenterology and Hepatology, Mayo Clinic, Minnesota, Estados Unidos
- ³⁰ Department Of Medicine, Division Of Gastroenterology, Western University, London Health Sciences Center, Ontario, Canadá

Introduction and Methods: Severe alcohol-associated hepatitis (AH) is frequently associated with higher infection risk. This study aimed to assess the impact of infections in patients with AH in a multinational cohort in Latin America.

Materials and Methods: Multicenter prospective cohort study including patients with AH (2015-2022). We recorded clinical information, and the impact of infections was assessed using competingrisk models.

Results: We included 511 patients from 24 centers in 8 countries (Argentina, Bolivia, Brazil, Chile, Colombia, Ecuador, Mexico, and Peru). The mean age was 50.1 ± 11.9 years, 426 (83.9%) were men, 264 (58.2%) had a previous diagnosis of cirrhosis, and the median MELD at diagnosis was 24.6 [19.6-30.6] points. Out of the total, 25.9% died, and only 3.7% underwent liver transplantation during follow-up. Also, 44.5% of patients developed an infection. Of them, 50.9% presented with infection at admission, 30.8% developed an infection during hospitalization, and 18.3% presented an infection in both situations. The most common localizations at admission were pulmonary (32.4%), urinary tract (33.1%), spontaneous bacterial peritonitis (15.9%), and cutaneous (9.7%). The main localizations during hospitalization were pulmonary (34.4%), urinary tract (25.8%), spontaneous bacterial peritonitis (14.0%), and bacteremia (8.6%%). The incidence of multidrug-resistant (MDR) organisms was 11.2% at admission and 10.3% during hospitalization, while the incidence of extensively drug-resistant (XDR) organisms was 1.4% and 4.7%, respectively. The presence of infection was associated with higher mortality (sub-distribution hazard ratio [sHR] 1.92, 95%CI:1.56-2.37; p<0.001). In a competing-risk model adjusted by age, sex, MELD, and ACLF grade, the infections were independently associated with mortality (sHR 1.33, 95%CI:1.02-1.75; p=0.037).

Conclusions: Infections during an AH episode are frequent and independently associated with mortality in Latin America. However, the incidence of multidrug-resistant organisms was lower than in other regions. Efforts should be made to prevent, diagnose, and adequately treat infections in AH.

https://doi.org/10.1016/j.aohep.2023.101256

O-7 CURRENT PRACTICE OF LIVER TRANSPLANTATION IN LATIN AMERICAN COUNTRIES: AN ALEH INTEREST GROUP SURVEY 2023

Paulo Bittencourt¹, Liana Codes¹, Adrian Gadano², Alejandra Villamil², Alfeu de Medeiros Fleck Jr³, Álvaro Urzua⁴, Debora Raquel Terrabuio⁵, Eira Cerda⁶, Graciela Elia Castro Narro⁷, Ignacio Roca⁸, John Abad González⁹, Josefina Pages¹⁰, Juan Carlos Restrepo Gutierrez¹¹, Leonardo de Lucca Schiavon¹², Mario Uribe¹³, Martin Padilla¹⁴, Norma Marlene Perez Figueroa¹⁵, Pablo Coste Murillo¹⁶, Raquel Stucchi¹⁷, Ricardo Chong¹⁸, Rodrigo Wolff¹⁹, Victoria Mainardi²⁰, Rodrigo Zapata²¹

Misericórdia, Porto Alegre, Rio Grande do Sul, Brasil

¹ Portuguese Hospital, Salvador, Bahia, Brasil

² Italian Hospital, Buenos Aires, Argentina

³ Adult Liver Transplant Group, Santa Casa

⁴ Clinical Hospital, University of Chile, Santiago, Chile

⁵ Clinical Hospital, University of São Paulo, São Paulo, Brasil

⁶ Central Military Hospital, Mexico City, México

⁷ National Institute of Medical Sciences and Nutrition Salvador Zubirán, Mexico City, México

⁸ El Cruce Hospital, Buenos Aires, Argentina

Abstracts Annals of Hepatology 29 (2024) 101187

- ⁹ Carlos Andrade Marín Hospital, Quito, Ecuador
- ¹⁰ Austral University Hospital, Buenos Aires, Argentina
- ¹¹ Pablo Tobon Uribe Hospital, Medellin, Colombia
- ¹² Federal University of Santa Catarina, Florianópolis, Santa Catarina, Brasil
- ¹³ Del Salvador Hospital, Santiago, Chile
- ¹⁴ Guillermo Almenara National Hospital, Lima, Perú
- ¹⁵ General Plaza de la Salud Hospital, S. Domingos, República Dominicana
- ¹⁶ RA Calderón Guardia Hospital, San Jose, Costa Rica
- ¹⁷ State University of Campinas, Campinas, Brasil
- ¹⁸ Francisco Clinic, Quito, Ecuador
- ¹⁹ Red de Salud UC Christus, Santiago, Chile
- ²⁰ National Liver Transplant Program, Montevideo, Uruguay
- ²¹ Alemana Clinic, Santiago, Chile

Introduction and Objectives: Little is known about current practice of liver transplantation (LT) in Latin American countries (LATAM). This study aimed to describe LT activity, immunosuppression protocols and policies regarding prophylaxis of cytomegalovirus (CMV) infection and hepatitis B virus (HBV) recurrence in different active LATAM centers.

Materials and Methods: A web-based survey with 20 questions regarding LT practice was sent to all members of ALEH LT SIG in December 2022.

Results: 22 centers performing 35 [5-160] LT per year from Brazil (n=5), Argentina (n=4), Chile (n=4), Ecuador (n-2), Mexico (n=2), Colombia (n=1), Costa Rica (n=1), Peru (n=1), Dominican Republic (n=1) and Uruguay (n=1) answered the survey. Tacrolimus, mycophenolate and prednisone was the main immunosuppressive regimen employed by most (72%) centers and 81% of them referred basiliximab use for induction therapy in selected patients. Tailoring of immunosuppression was universally accepted, particularly in autoimmune hepatitis (AIH) (59%), hepatocellular carcinoma (54%) kidney dysfunction (77%) and primary biliary cirrhosis (33%). Weaning of corticosteroids at three, six and 12 months after LT was reported, respectively, by 41%, 36% and 23% of the centers, but policy for lifelong corticosteroid use in AIH-transplanted subjects was commonly observed (90%). Just four centers are currently performing protocol liver biopsies, while 18 of them are considering liver biopsy prior to steroid pulse therapy. HBIG and nucleos(t)ide analogs are used in most instances (73%) for HBV recurrence prevention, whereas CMV infection prophylaxis was shown to vary sharply across centers. Of note, all but two of them referred major changes in LT practice over the years due to economical restraints.

Conclusions: Compliance with standard of care recommendations for management of LT was reported by most centers. Heterogeneity in practices regarding HBV infection recurrence and CMV prophylaxis may reflect local financial restraints and point to the importance of developing ALEH guidelines to encourage LT activity in LATAM.

https://doi.org/10.1016/j.aohep.2023.101257

O- 8 PNPLA3 RS738409 C>G POLYMORPHISM IMPACT ON HCV-RELATED HCC DEVELOPMENT IN THE BRAZILIAN POPULATION: PRELIMINARY RESULTS

Claudia Maccali, Aline L Chagas, Lisa Rc Saud, Regiane Ssm Alencar, Michele Sg Gouvea, Joyce Mks Etto, Isabel V Pereira, Arthur In Oliveira, Jose T Stefano, Rafael Sn Pinheiro, Wellington Andraus, Paulo Herman, Luiz Ac D'albuquerque, Mario G Pessoa, Claudia P Oliveira Department of Gastroenterology, University of São Paulo School of Medicine, São Paulo, Brasil

Introduction and Objectives: The PNPLA3 rs738409 C>G polymorphism has been associated with hepatocellular carcinoma (HCC) and liver cirrhosis regardless of the etiology, although the association was stronger with non-viral etiologies. However, the influence of PNPLA3 polymorphism on Hepatitis C Virus (HCV) and whether this polymorphism could be a risk factor for HCV-related HCC is not well defined. We aimed to evaluate the influence of the PNPLA3 rs738409 C>G polymorphism on the risk of HCC occurrence in HCV patients in Brazil.

Materials and Methods: This study included 90 patients with HCV-related cirrhosis and HCC who underwent liver transplantation or resection at a tertiary center in Brazil and 111 patients non-HCC with HCV-related cirrhosis, as the control group. The rs738409 polymorphism was detected in the DNA extracted from patients' blood samples using the TaqMan assay. All clinical data were collected using the Research Electronic Data Capture (REDCap) tool. The statistical analyses were performed using Jamovi software version 2.3.23.

Results: In the HCV+HCC group there was a higher proportion of male gender (79.1% vs. 45.9%, p<0.001), history of alcoholism (80.5% vs. 22.5%, p<0.001) and smoking (68.9% vs. 25.2%, p<0.001), however there was no statistical difference in age (p=0.519) and BMI (p=0.403) between both groups. The genotype frequencies of the rs738409 polymorphism in the HCV+HCC group was CC 41,2% CC and CG/GG 58,8% vs. controls CC 49,5% and CG/GG 50,5%. The presence of the G allele was not an independent factor associated with the risk of HCC occurrence (r=0,199, p=0.53).

Conclusions: Even in an admixed population such as the Brazilian, there was no association between the PNPLA3 rs738409 C>G polymorphism and the risk of developing HCV-related HCC, as previously shown in published studies in caucasian and oriental population

https://doi.org/10.1016/j.aohep.2023.101258

O-9 PRESENCE OF METABOLIC ASSOCIATED LIVER DISEASE IN AUTOIMMUNE HEPATITIS IS ASSOCIATED WITH ADVANCED LIVER FIBROSIS

Giselle Arévalo¹, Daniela Simian¹, Jorge Chiong², Consuelo Palomo¹, Laura Carreño¹, Jaime Poniachik¹, Álvaro Urzúa¹

 Sección de Gastroenterología, Hospital Clínico Universidad de Chile, Santiago, Chile
Facultad de Medicina, Universidad de Chile, Santiago, Chile

Introduction and Objectives: Metabolic associated liver disease (MAFLD) is one of the most frequent causes of chronic liver disease worldwide. Steatohepatitis is a major risk factor for fibrosis and its progression. MAFLD might be present in other causes of chronic liver damage, such as autoimmune hepatitis (AIH). There is scarce information of the role of MAFLD in fibrosis in patients with AIH. This study aimed to describe frequency of steatosis, steatohepatitis and fibrosis in AIH liver biopsies and the impact of steatosis and steatohepatitis on fibrosis severity and survival.

Materials and Methods: Observational, retrospective study of liver biopsies performed prior to initiation of immunosuppressive therapy, between 2014 and 2019. Presence of steatosis, steatohepatitis and fibrosis was recorded. Alcohol and other etiologies of liver disease were excluded. Clinical data was obtained from electronic charts and outcome variables by telephone contact. Chi2 and exact Fisher test and Odds Ratio (OR) were performed (p < 0.05 considered statistically significant).