

Results: A total of 6,991 patients from all 66 dialysis facilities in RS State were enrolled. Most patients (93.3%) were on hemodialysis. All patients completed HCV screening and 454 (6.5%) were anti-HCV positive. So far, nine units have completed the proposed model, with 89 anti-HCV positive patients that resulted in 49 (55.5%) with detectable HCV RNA by PCR. All viremic patients started HCV therapy. Interim analysis showed SVR in 21 (95.5%) of 22 patients.

Conclusions: A collaborative care model increased the rates of diagnosis and treatment for HCV in dialysis facilities to levels near those established by the World Health Organization towards HCV elimination up to 2030.

<https://doi.org/10.1016/j.aohep.2023.101029>

O-20 MOLECULAR AND BIOLOGICAL CHARACTERIZATION OF HEPATITIS B VIRUS SUBGENOTYPE F1b CLUSTERS: UNRAVELING ITS ROLE IN HEPATOCARCINOGENESIS

María Mercedes Elizalde¹, Laura Mojsejczuk², Micaela Speroni¹, Luciana Tadey², Belén Bouzas³, Lilia Mammana³, Rodolfo Héctor Campos², Diego Martín Flichman¹

¹ Institute for Biomedical Research on Retroviruses and AIDS (INBIRS), CONICET, Buenos Aires University, Buenos Aires, Argentina

² Department of Microbiology, Immunology, Biotechnology and Genetics, Chair of Virology, University of Buenos Aires, Autonomous City of Buenos Aires, Argentina

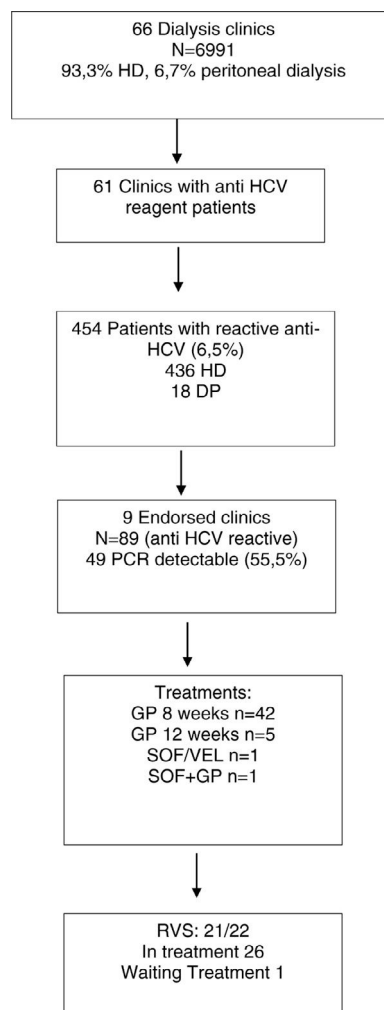
³ Virology Unit, Infectious Diseases Hospital Francisco J. Muñiz, Autonomous City of Buenos Aires, Argentina

Introduction and Objectives: Hepatitis B virus subgenotype F1b infection has been associated with the early occurrence of hepatocellular carcinoma in chronically infected patients from Alaska and Peru. In Argentina, however, despite the high prevalence of subgenotype F1b infection, this relationship has not been described. This study aimed to unravel the observed differences in the progression of the infection, and an in-depth molecular and biological characterization of the subgenotype F1b was performed.

Materials and Methods: 99 subgenotype F1b full-length sequences were obtained, and phylogeny was addressed by the maximum likelihood method. The replicative capacity of the subgenotype F1b clones was assessed by qPCR, Southern and Northern blot analysis. Antigen expression was detected by electrochemiluminescence and Western blot. The analysis of signaling pathways associated with hepatocarcinogenesis was assessed by RT-qPCR.

Results: Phylogenetic analysis of subgenotype F1b genomes revealed the existence of two highly supported clusters. One of the clusters, designated as gtF1b Basal included sequences mostly from Alaska, Peru, and Chile, while the other, called gtF1b Cosmopolitan, contained samples mainly from Argentina and Chile. The clusters were characterized by a differential signature pattern of eight nucleotides distributed throughout the genome. *In vitro* characterization of representative clones from each cluster revealed major differences in viral RNA levels, virion secretion, and antigen expression levels. Interestingly, differential regulation in the expression of genes associated with tumorigenesis was also identified.

Conclusions: This study provides new insights into the molecular and biological characteristics of the subgenotype F1b clusters and contributes to unraveling the different clinical outcomes of subgenotype F1b chronic infections.



<https://doi.org/10.1016/j.aohep.2023.101030>

O-21 EVIDENCE OF SUBOPTIMAL PUBLIC HEALTH POLICIES ON HEPATOCELLULAR CARCINOMA IN THE AMERICAS: A HUGE DEBT OF OUR REGION

Luis Antonio Díaz¹, Gustavo Ayares¹, Francisco Idalsoaga¹, Jorge Arnold¹, Blanca Norero^{1,2}, Oscar Corsi¹, Gonzalo Pizarro³, Sergio García⁴, Eduardo Fuentes-López⁵, Edmundo Martínez², Patricia Guerra Salazar⁶, Roberta C. Araújo⁷, Mario Reis Alvares-Da-Silva⁸, Florencia D. Pollarsky⁹, Nelia Hernandez¹⁰, Juan Carlos Restrepo¹¹, Mirtha Infante¹², Enrique Carrera¹³, Abel Sanchez¹⁴, Marcos Giralá¹⁵, Martín Padilla¹⁶, Javier Díaz¹⁷, Martín Tagle¹⁸, Melisa Dirchwolf¹⁹, Manuel Mendizabal²⁰, Mariana Lazo²¹, Catterina Ferreccio²², Thomas G. Cotter²³, Mayur Brahmanian²⁴, Nahum Méndez-Sánchez²⁵, Juan Pablo Roblero²⁶, Winston Dunn²⁷, Patrick S. Kamath²⁸, Ashwani K. Singal²⁹, Ramón Bataller³⁰, Marco Arrese¹, Juan Pablo Arab¹

¹ Department of Gastroenterology, Medical School, Pontifical Catholic University of Chile, Santiago, Chile

² Gastroenterology Service, Dr. Sótero del Río Hospital, Santiago, Chile

³ Department of Hematology-Oncology, Medical School, Pontifical Catholic University of Chile, Santiago, Chile

⁴ Medical School, Pontifical Catholic University of Chile, Santiago, Chile

⁵ Department of Health Sciences, Speech-Language Pathology Department, Medical School, Pontifical Catholic University of Chile, Santiago, Chile

⁶ Gastroenterology Institute Bolivian-Japanese, Cochabamba, Bolivia

⁷ Gastroenterology Division, Ribeirão Preto Medical School, University of São Paulo, 14048-900 Ribeirão Preto, SP, Brazil

⁸ Clinic Hospital of Porto Alegre, Porto Alegre, Brazil

⁹ Hepatology Section, Gastroenterology Hospital of Dr. Carlos Bonorino Udaondo, Buenos Aires, Argentina

¹⁰ Gastroenterology Clinic, Clinic Hospital Medical School, University of República Uruguay, Montevideo, Uruguay

¹¹ Hepatology Unit of Pablo Tobon Uribe Hospital, Gastrohepatology Group of Antioquia University, Medellín, Colombia

¹² Medical Science University of La Habana, Havana, Cuba

¹³ Eugenio Espejo Hospital, Quito, Ecuador

¹⁴ Gastroenterology Digestive Endoscopy Hepatology, Roosevelt Hospital, Guatemala City, Guatemala

¹⁵ Gastroenterology Department, Clinic Hospital, National University of Asunción, Asunción, Paraguay

¹⁶ Mayor Nacional University of San Marcos. Guillermo Almenara National Hospital, Lima, Perú

¹⁷ Nacional Hospital Edgardo Rebagliati, Jesús María, Perú

¹⁸ Medical School Alberto Hurtado, Peruvian University Cayetano Heredia, Lima, Perú

¹⁹ Liver Unit, Private Hospital of Rosario, Rosario, Argentina

²⁰ Hepatology and Liver Transplant Unit, Austral University Hospital, Buenos Aires, Argentina

²¹ Department of Community Health and Prevention, Dornsife School of Public Health, Drexel University, Philadelphia, Pennsylvania; Urban Health Collaborative, Dornsife School of Public Health, Drexel University, Philadelphia, Pennsylvania; Division of General Internal Medicine, Johns Hopkins University School of Medicine, Baltimore, Maryland, USA

²² Public Health Department, School of Medicine, Pontifical Catholic University of Chile, Santiago, Chile. Advanced Center for Chronic Diseases, ACCDis, Santiago, Chile

²³ Division of Digestive and Liver Diseases, UT Southwestern Medical Center, Dallas, Texas, USA

²⁴ Department of Medicine, Division of Gastroenterology, Western University, London Health Sciences Center, London, Ontario, Canada

²⁵ Liver Research Unit, Medica Sur Clinic & Foundation, Faculty of Medicine, National Autonomous University of Mexico, Mexico City, 14050, Mexico

²⁶ Gastroenterology Section, Clinical Hospital of University of Chile, Medical School, University of Chile, Santiago, Chile

²⁷ University of Kansas Medical Center, KS, USA

²⁸ Division of Gastroenterology and Hepatology, Mayo Clinic, Rochester, MN, USA

²⁹ Department of Medicine, University of South Dakota Sanford School of Medicine, Division of Transplant

Hepatology, Avera Transplant Institute, Sioux Falls, SD, United States

³⁰ Center for Liver Diseases, Division of Gastroenterology, Hepatology and Nutrition, University of Pittsburgh Medical Center, PA, USA

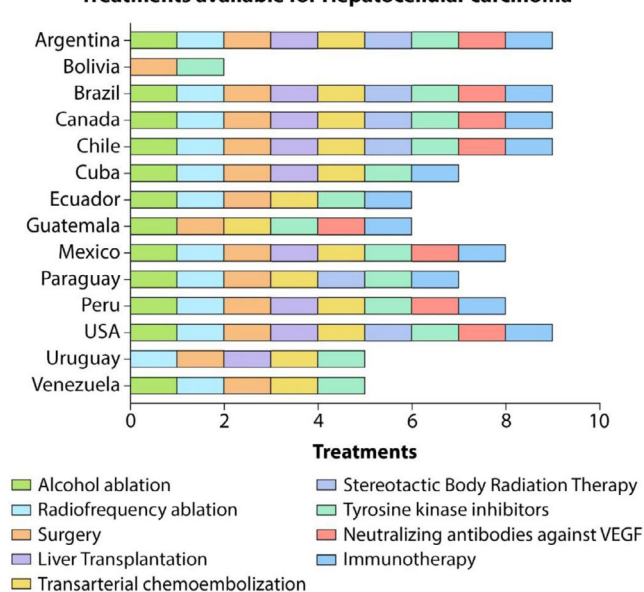
Introduction and Objectives: Although most cases of HCC occur in patients with advanced liver disease and there are effective screening methods, health policies aimed at preventing and detecting HCC are not often on the agenda of government initiatives and policies. This study aimed to explore HCC-related population-wide public health policies, treatment availability, epidemiological surveillance, and awareness campaigns in the Americas.

Materials and Methods: We conducted a 43-item survey about HCC among experts from 12 countries. Questions were classified into four categories: policies and civil society (18 questions), clinical guidelines (5 questions), epidemiology (7 questions), and care management (13 questions). The survey was administered using an electronic form in May 2022. Data was collected in a spreadsheet, revised by two independent reviewers, and contrasted with governmental institutions.

Results: We obtained 22 responses from 15 out of the 18 countries targeted. A total of 7 (47%) countries had a written national cancer strategy or action plan. Only 4 (27%) countries had a specific written national HCC strategy or action plan, including Argentina, Brazil, Mexico, and the United States. These same four countries also had national clinical guidelines on HCC. HCC is managed by various providers, including Hepatologists (80%), Oncologists (80%), Gastroenterologists (60%), Surgeons (47%), and Palliative medicine (20%). There were important differences in the availability of treatments among countries in the Americas (Figure). Of note, 60% of countries had liver transplantation available for HCC, but only 67% of them had this therapy outside the capital city. Nine (60%) countries had a national disease registry that included HCC. However, only Brazil (7%) had governmental funded awareness campaigns on HCC prevention or screening.

Conclusions: Implementation of public health policies on HCC is scarce in the Americas. Important differences in treatments were observed across countries, especially in curative therapies. Our results strongly encourage developing public health policies on HCC in the Americas.

Treatments available for Hepatocellular carcinoma



<https://doi.org/10.1016/j.aohep.2023.101031>