O-7 PREVALENCE, CHARACTERIZATION AND SURVIVAL OF ACUTE-ON-CHRONIC LIVER FAILURE IN A CHILEAN UNIVERSITY HOSPITAL

Francisco Idalsoaga¹, Francisco Valenzuela¹, Antonio Díaz Luis¹, Franco Manzur², Victor Meza², Joaquin Sotomayor², Maximiliano Schalper², Franco Chianale², Hernan Rodríguez², Juan Arab Pablo¹

Background: Acute-on-chronic liver failure (ACLF) is a serious clinical entity, with no previous reports in Chile.

Aim: To characterize patients with ACLF in a Chilean University Hospital, identifying triggers, organ failure and survival at 30, 90, 180 days, compared to patients with decompensated cirrhosis without ACLF.

Methods: Retrospective cohort study of decompensated cirrhotic patients hospitalized in a chilean University Hospital between 2017-2019.

Results: 334 patients were included, 73 (22%) presented ACLF (33% ACLF-1, 30% ACLF-2, 37% ACLF-3); 16.4% underwent liver transplantation. Patients with ACLF were younger, and had higher MELD-Na and APACHE II on admission. The most common triggers in both groups were infections (42.4%), gastrointestinal bleeding (23.2%) and alcohol intake (31.3%). The main organ failures were kidney (60.2%) and brain (49.3%). All organ failures were more frequent in ACLF-3, except renal failure (greater in ACLF-1). Survival at 180 days was 74% in patients without ACLF and 58.3% in ACLF (p=0.004). Mortality was significantly higher in ACLF-2 and ACLF-3, when compared with patients without ACLF (HR 2.3 and 2.99, respectively; p<0.05). Transplant-free survival in cirrhotics without ACLF was 72.5% versus 43.1% with ACLF (p<0.001). The risk of mortality or transplantation was higher in ACLF-2 and ACLF-3, in contrast to patients without ACLF (HR 2.19 and 4.61, respectively; p<0.05).

Conclusions: ACLF is an entity of younger patients, with lower global and transplantation-free survival at 180 days and multiple organ failure compared to decompensated cirrhotics without ACLF.

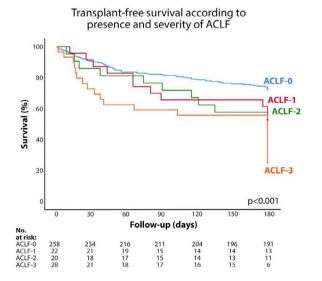


Figure.- Cumulative transplant-free survival according to the presence of ACLF and severity.

O-8 THE IMPORTANCE OF LYSOSOMAL ACID LIPASE DEFICIENCY IN THE ETIOLOGICAL INVESTIGATION OF CRYPTOGENIC LIVER DISEASE IN ADULTS: A MULTICENTER STUDY

Aline Coelho Rocha Candolo¹,
Patricia Momoyo Zitelli¹,
Daniel Ferraz de Campos Mazo^{1,2},
Marlone Cunha-Silva², Raquel Dias Greca²,
Claudia Pinto de Oliveira¹, Roberta Chaves Araújo³,
Amanda Sacha Paulino Tolentino Alustau³,
Claudia Alves Couto⁴, Mateus Jorge Nardelli⁴,
Júlio M. Singer⁵, Roque Gabriel Rezende de Lima¹,
Alberto Queiroz Farias¹, Flair José Carrilho¹,
Mário Guimarães Pessoa¹

 Division of Clinical Gastroenterology and Hepatology, Department of Gastroenterology, University of São Paulo School of Medicine (FMUSP), Sao Paulo, Brazil
 Division of Gastroenterology (Gastrocentro), School of Medical Sciences, University of Campinas (UNICAMP), Campinas. Brazil

³ Division of Gastroenterology, Hospital of the Clinics of the Faculty of Medicine of Ribeirão Preto, University of São Paulo (FMRP-USP), Ribeirao Preto, Brazil ⁴ Department of Internal Medicine, Federal University of Minas Gerais (UFMG), Belo Horizonte, Brazil ⁵ Institute of Mathematics and Statistics, University of Sao Paulo (USP), São Paulo, Brazil

Background: Lysosomal acid lipase deficiency (LAL-D) is a rare genetic disease associated with lipid metabolism deregulation leading to atherosclerosis, dyslipidemia, and hepatic steatosis, with potential progression to cirrhosis. Investigation of LAL-D in patients with chronic liver disease is not routinely performed in most centers.

Aim: The aim of this study was to evaluate whether it is worthwhile to investigate LAL-D in patients with liver disease of unknown etiology, and if there is any particular population that this search should be focused.

Methods: This was a multicenter cross-sectional study in 295 patients followed with presumed cryptogenic liver disease from four tertiary centers in Brazil. Clinical, demographic and laboratory data from participants were assessed, with the exclusion of all known causes of liver disease. All patients were submitted to the investigation of LAL enzyme activity. The exams were collected on dried blood spot (DBS).

Results: A total of 135 patients were included in the study. Three patients (2.22%) presented values of LAL below the reference limit, compatible with LAL-D. The mean age of these patients was 43.9 ± 10.1 years, of which 2 were females. The mean BMI was 24.3 ± 0.7 and mean serum glycemia was 89.7 ± 3.2 mg/dL. The mean serum HDL and triglycerides were 21.7 ± 3.2 mg/dL and 206.7 ± 25.5 mg/dL, respectively.

Conclusion: Despite being a rare disease, also in our study population, LAL-D investigation may be considered in those individuals without overweight with reduced serum HDL and elevated triglycerides levels and chronic liver disease of unknown etiology.

https://doi.org/10.1016/j.aohep.2021.100495

O-9 COMPARISON OF THE PERFORMANCE OF DIFFERENT SCORES FOR THE PREDICTION OF IN-HOSPITAL MORTALITY IN PATIENTS WITH CIRRHOSIS AND BACTERIAL INFECTIONS

Agustina Martinez Garmendia¹, Maria Nelly Gutierrez Acevedo², Sabrina Barbero³,

¹ Departamento of Gastroenterología, Escuela de Medicina, Pontificia Universidad Católica de Chile, Santiago. Chile

² Escuela de Medicina, Pontificia Universidad Católica de Chile, Santiago, Chile

Abstracts Annals of Hepatology 24 (2021) 100366

Lorena del Carmen Notari³, Marina Agozino⁴, Iose Luis Fernandez⁴, Maria Margarita Anders⁵, Nadia Grigera⁵. Florencia Antinucci⁵. Orlando Orozco Ganem⁵, Maria Dolores Murga⁶, Daniea Perez⁶. Ana Palazzo⁶. Liria Martinez Rejtman⁷, Ivonne Giselle Duarte⁸, Julio Vorobioff⁹, Victoria Trevizan⁹, Sofía Bulaty⁹, Fernando Bessone⁹, José Daniel Bosia⁹, Silvia Mabel Borzi¹⁰, Teodoro E. Stieben¹¹, Adriano Masola¹¹, Sebastian Eduardo Ferretti¹², Diego Arufe¹³, Ezequiel Demirdjian¹³, Maria Pia Raffa¹³, Cintia Elizabet Vazquez¹⁴, Pablo Ruiz¹⁴, José Emanuel Martínez¹⁵, Hugo Fainboim¹⁶, Mirta Peralta¹⁶, Leandro Alfredo Heffner¹⁷, Andrea Odzak¹⁷, Melisa Dirchwolf¹⁸, Astrid Smud¹⁹, Manuel Mendizabal²⁰, Carla Bellizzi²¹, Diego Giunta²², Marcelo Valverde^{23,24}, Martin Elizondo^{23,24}, Ezequiel Mauro¹, Ana Martinez²¹, Jesica Tomatis¹⁸, Andres Bruno¹⁷, Agñel Ramos¹², Josefina Pages²⁰, Silvina Tevez⁴, Salvatore Piano²⁵, Adrian Gadano^{1,22}, Sebastián Marciano^{1,22}

- ¹ Hospital Italiano de Buenos Aires, Liver Unit, CABA, Argentina
- ² Clinica Chapelco, San Martin de los Andes, Argentina
- ³ Hospital Churruca Visca, CABA, Argentina, San Martin de los Andes, Argentina
- ⁴ Sanatorio Güemes, CABA, Argentina
- ⁵ Hospital Aleman, CABA, Argentina
- ⁶ Hospital A.C. Padilla, San Miguel de Tucuman, Argentina
- ⁷ Hospital T J Schestakow, Mendoza, Argentina
- ⁸ Hospital 4 de Junio, Saenz Peña, Argentina
- ⁹ Hospital provincial del Centenario, Rosario, Argentina
- ¹⁰ Hospital Rossi, La Plata, Argentina
- ¹¹ Hospital San Martín, Paraná, Argentina
- ¹² Sanatorio Parque, Rosario, Argentina
- ¹³ Sanatorio Sagrado Corazon, CABA, Argentina
- ¹⁴ Hospital Regional de Rio Gallegos, Río Gallegos, Argentina
- ¹⁵ Sanatorio Boratti, Posadas, Argentina
- ¹⁶ Hospital Muñiz, CABA, Argentina
- ¹⁷ Hospital Argerich, CABA, Argentina
- ¹⁸ Hospital Privado de Rosario, Rosario, Argentina
- ¹⁹ Hospital Italiano de Buenos Aires, Infectious Diseases Section, ABH, CABA, Argentina
- ²⁰ Hospital Universitario Austral, Pilar, Argentina
- ²¹ Hospital Fernández, CABA, Argentina
- ²² Hospital Italiano de Buenos Aires, Department of Research, CABA, Argentina
- ²³ Hospital de Clinicas, Montevideo, Uruguay
- ²⁴ Hospital Militar, Montevideo, Uruguay
- ²⁵ Unit of Internal Medicine and Hepatology, Department of Medicine, University of Padova, Padova, Italy

Background: Predicting short-term mortality in patients with cirrhosis and bacterial infections is challenging.

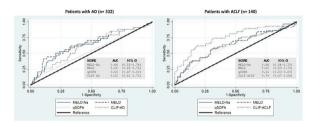
Aims: To compare the performance of various scores in predicting in-hospital mortality in this population.

Methods: We performed an analysis of the multicenter prospective cohort study of patients with cirrhosis with bacterial infections throughout Argentina and Uruguay (clinicatrials.gov.NCT03919032). Patients were classified according to the CLIF criteria as having ACLF or mere acute decompensation (AD). We evaluated the performance of scores of liver disease and infection severity in predicting in-

hospital mortality. MELD, MELD-Na, and Quick SOFA (qSOFA) were computed in all patients. CLIF-AD was only computed in patients without ACLF, and CLIF-ACLF only in patients with ACLF. We plotted ROC curves and estimated their area under the curve (AUROC).

Results: We included 472 patients: 66% male, mean age 57 \pm 12 years. Most frequent infections: SBP (30%) and urinary tract infection (25%). Overall, 332 (70%) patients had acute decompensation, and 140 (30%) ACLF. In-hospital mortality rate was 19%: 41% in patients with ACLF vs 10% in patients with AD (p<0,001). When we evaluated the AUROC of the entire cohort, MELD and MELD-Na performed similarly: 0.74 (95% CI 0.68-0.81) and 0.74 (95% CI 0.67-0.80), respectively; whereas qSOFA showed the lowest performance: 0.62 (95% CI 0.57-0.68). When evaluating only patients with ACLF, CLIFACLF performed significantly better than the other ones: AUROC 0.76 (95% CI 0.67-0.84, p =0.01). All scores performed poorly in patients with AD (Figure).

Conclusion: The best tool to predict in-hospital mortality in patients with infection-related ACLF was the CLIF-ACLF score. In patients with infection-related AD, all scores performed poorly. Evaluation of the scores performance is of paramount importance in different regions and for each complication of cirrhosis separately.



https://doi.org/10.1016/j.aohep.2021.100496

O-10 PRIMARY BILIARY CHOLANGITIS PATIENTS DIAGNOSED BY DIFFERENT COMBINATIONS OF THE DIAGNOSTIC CRITERIA PRESENT CLINICAL AND LABORATORY PECULIARITIES

Guilherme G.L. Cançado^{1,2}, Eduardo L.R. Cançado³, Maria L.G. Ferraz⁴, Cristiane A. Villela-Nogueira⁵, Debora R.B. Terrabuio³, Michelle H. Braga³, Mateus J. Nardelli¹, Luciana C. Faria¹, Nathalia M.F. GOMES⁴, ELZE M.G. Oliveira⁴, Vivian Rotman⁵, Maria Beatriz Oliveira⁶, Simone M.C.F. Cunha⁷, Daniel F. Mazo⁸, Liliana S.C. Mendes⁹, Claudia A.P. Ivantes¹⁰, Valéria Faborges^{11,12}, Fabio H.L. Pace¹³, Mario G. Pessoa³, Izabelle V. Signorelli¹⁴, Gabriela P. Coral¹⁵, Paulo L. Bittencourt^{16,17}, Cynthia Levy¹⁸, Cláudia A. Couto¹, Members of the Brazilian Cholestasis Study Group Consortium¹

¹ Hospital das Clínicas da Universidade Federal de Minas Gerais, Belo Horizonte, Brazil

² Hospital da Polícia Militar de Minas Gerais, Belo Horizonte, Brazil

³ Departamento de Gastroenterologia, Faculdade de Medicina da Universidade de São Paulo, São Paulo, Brazil

⁴ Disciplina de Gastroenterologia, Universidade Federal de São Paulo, São Paulo, Brazil

⁵ Hospital Universitário Clementino Fraga Filho e Departamento de Clínica Médica da Faculdade de