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*DECENIO DE LA IGUALDAD DE OPORTUNIDADES PARA MUJERES Y HOMBRES
" AÑO DEL FORTALECIMIENTO DE LA SOBERANÍA NACIONAL "*

INFECTOLOGÍA

➤ **Outcomes of HTLV-1 Carriers with Diffuse Large B-Cell Lymphoma: A Single-Center Retrospective Matched Cohort Study**

INVESTIGADORES: Bryan Valcarcel, Gustavo Sandival Ampuero, Gabriel de la Cruz-Ku, Daniel J Enriquez, Luis Malpica.

REVISTA: Clin Lymphoma Myeloma Leuk 2021 Sep 29;S2152-2650(21)02069-3. doi: 10.1016/j.clml.2021.09.017.

ABSTRACTO: Background: The human T-cell lymphotropic virus type 1 (HTLV-1) is associated with aggressive diseases, such as adult T-cell leukemia/lymphoma (ATLL). However, less is known on the impact of HTLV-1 infection in non-ATLL hematologic malignancies. We aimed to investigate if HTLV-1 carriers with diffuse large B-cell lymphoma (DLBCL) have worse survival outcomes than non-HTLV-1 carriers. Materials and methods: We performed a single-center retrospective cohort study by matching HTLV-1 carriers to non-carriers based on age, sex, Ann Arbor stage, and year of diagnosis. Our outcomes of interest were overall survival (OS) and progression-free survival (PFS). The Kaplan-Meier method was used to estimate OS and PFS between carriers and non-carriers. We fitted multivariate Cox regression models to assess the mortality and recurrence/disease progression risk of HTLV-1 infection. Results: A total of 188 patients, 66 with HTLV-1 infection and 122 without HTLV-1, were included in the study. HTLV-1 carriers had higher extranodal involvement than non-carriers (47% vs. 27%, $P = .010$). With a median follow-up of 78 months (95% CI: 41-90 months), HTLV-1 carriers had a similar 5 year OS (41% vs. 42%, $P = .940$) and PFS (34% vs. 32%, $P = .691$) compared to non-carriers. In the multivariate Cox analysis, HTLV-1 infection was not associated with worse OS (aHR: 0.98, 95% CI: 0.64-1.50) or PFS (aHR: 0.90, 95% CI: 0.60-1.34). Conclusion: HTLV-1 carriers with DLBCL did not have worse survival outcomes compared to non-carriers. Our results suggest that clinicians should follow standard guidelines for DLBCL management on HTLV-1 seropositive patients.

➤ **Association between convalescent plasma treatment and mortality in COVID-19: a collaborative systematic review and meta-analysis of randomized clinical trials**

INVESTIGADORES: Cathrine Axfors, Perrine Janiaud, Andreas M Schmitt, Janneke Van't Hooft, Emily R Smith, Noah A Haber, Akin Abayomi, Manal Abduljalil, Abdulkarim Abdulrahman, Yeny Acosta-Ampudia, Manuela Aguilar-Guisado, Farah Al-Beidh, Marissa M Alejandria, Rachelle N Alfonso, Mohammad Ali, Manaf AlQahtani, Alaa AlZamrooni, Juan-Manuel Anaya, Mark Angelo C Ang, Ismael F Aomar, Luis E Argumanis, Alexander Averyanov, Vladimir P Baklaushev, Olga Balionis, Thomas Benfield, Scott Berry, Nadia Birocco, Lynn B Bonifacio, Asha C Bowen, Abbie Bown, Carlos Cabello-Gutierrez, Bernardo Camacho, Adrian Camacho-Ortiz, Sally Campbell-Lee, Damon H Cao, Ana Cardesa, Jose M Carnate, German Jr J Castillo, Rossana Cavallo, Fazle R Chowdhury, Forhad U H Chowdhury, Giovannino Ciccone, Antonella Cingolani, Fresthel Monica M Climacosa, Veerle Compennolle, Carlo Francisco N Cortez, Abel Costa Neto, Sergio D'Antico, James Daly, Franca Danielle, Joshua S Davis, Francesco Giuseppe De Rosa, Justin T Denholm, Claudia M Denkinge, Daniel Desmecht, Juan C Díaz-Coronado, Juan A Díaz Ponce-Medrano, Anne-Françoise Donneau, Teresita E Dumagay, Susanna Dunachie, Cecile C Dungog, Olufemi Erinoso, Ivy Mae S Escasa, Lise J Estcourt, Amy Evans, Agnes L

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REVISTA: BMC Infect Dis 2021 Nov 20;21(1):1170. doi: 10.1186/s12879-021-06829-7.

ABSTRACTO: Background: Convalescent plasma has been widely used to treat COVID-19 and is under investigation in numerous randomized clinical trials, but results are publicly available only for a small number of trials. The objective of this study was to assess the benefits of convalescent plasma treatment compared to placebo or no treatment and all-cause mortality in patients with COVID-19, using data from all available randomized clinical trials, including unpublished and ongoing trials. Methods: In this collaborative systematic review and meta-analysis, clinical trial registries (ClinicalTrials.gov, WHO International Clinical Trials Registry Platform), the Cochrane COVID-19 register, the LOVE database, and PubMed were searched until April 8, 2021. Investigators of trials registered by March 1, 2021, without published results were contacted via email. Eligible were ongoing, discontinued and completed randomized clinical trials that compared convalescent plasma with placebo or no treatment in COVID-19 patients, regardless of setting or treatment schedule. Aggregated mortality data were extracted from publications or provided by investigators of unpublished trials and combined using the Hartung-Knapp-Sidik-Jonkman random effects model. We investigated the contribution of unpublished trials to the overall evidence. Results: A total of 16,477 patients were included in 33 trials

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(20 unpublished with 3190 patients, 13 published with 13,287 patients). 32 trials enrolled only hospitalized patients (including 3 with only intensive care unit patients). Risk of bias was low for 29/33 trials. Of 8495 patients who received convalescent plasma, 1997 died (23%), and of 7982 control patients, 1952 died (24%). The combined risk ratio for all-cause mortality was 0.97 (95% confidence interval: 0.92; 1.02) with between-study heterogeneity not beyond chance ($I^2 = 0\%$). The RECOVERY trial had 69.8% and the unpublished evidence 25.3% of the weight in the meta-analysis. Conclusions: Convalescent plasma treatment of patients with COVID-19 did not reduce all-cause mortality. These results provide strong evidence that convalescent plasma treatment for patients with COVID-19 should not be used outside of randomized trials. Evidence synthesis from collaborations among trial investigators can inform both evidence generation and evidence application in patient care.

➤ **Prevalence of viral hepatitis type C in blood donors in Peru 2016 - 2017**

INVESTIGADORES: Alvaro Bellido, Enrique Argumanis, Patricia Segura, Martin Tagle.

REVISTA: Rev Gastroenterol Peru Jul-Sep 2021;41(3):164-168.

ABSTRACTO: Objectives: Knowing the current prevalence of hepatitis C virus infection has taken great importance nowadays due to the new advances in the treatment of hepatitis C virus infection in which the new direct-acting antivirals have demonstrated a high rate of sustained viral response. Materials and methods: This is a descriptive study based on the detection of anti VHC in the blood banks of "Ministerio de Salud del Perú" according to the information obtained from "Programa Nacional de Hemoterapia y Bancos de Sangre" (PRONAHEBAS). Results: In our study, the prevalence of anti-hepatitis C antibodies in blood donors nationwide was 0.428% in 2016 and 0.301% in 2017. Conclusion: To know updated data about the prevalence of hepatitis C virus infection in our population. It is the first step to be able to treat the disease and drastically reduce the infection rate according to the WHO target for 2030.