



Abstract LAG-3 Role in Infection *

Luisa Chocarro ^{1,*}, Ester Blanco ¹, Hugo Arasanz ^{1,2}, Ana Bocanegra ¹, Leticia Fernández Rubio ¹, Miriam Echaide ¹, Maider Garnica ¹, Pablo Ramos ¹, Grazyna Kochan ^{1,*}, and David Escors ^{1,*}

- ¹ Oncoimmunology Group, Navarrabiomed-Fundacion Miguel Servet, Universidad Pública de Navarra, IdISNA, Irunlarrea 3, 31008 Pamplona, Navarra, Spain
- ² Department of Medical Oncology, Complejo Hospitalario de Navarra CHN-IdISNA, 31008 Pamplona, Navarra, Spain
- * Correspondence: <u>luisa.chocarro.deerauso@navarra.es</u> (L.C.); <u>grkochan@navarra.es</u> (G.K.); <u>descorsm@na-varra.es</u> (D.E.)
- + Presented at the 1st International Electronic Conference on Molecular Sciences: Druggable Targets of Emerging Infectious Diseases, online, 01-14 September 2021.

Academic Editor: Clemente Capasso

Published: 31 August 2021

Abstract: Lymphocyte activation gene 3 (LAG-3) is a cell surface inhibitory receptor with multiple biological activities over T cell activation and effector functions. LAG-3 plays a regulatory role in immunity and emerged some time ago as an inhibitory immune checkpoint molecule. A systematic research was performed using the PubMed and ClinicalTrial.gov databases. Articles published up to 2021 meeting the inclusion criteria were investigated. LAG-3 expression has been linked to increased pathology in certain infections, such as the ones caused by Salmonella, Plasmodium parasites, Mycobacterium tuberculosis, human immunodeficiency virus (HIV), non-pathogenic simian immunodeficiency virus (SIV), in hepatitis B virus (HBV), human papillomavirus (HPV), chronic hepatitis C virus (HCV), lymphocytic choriomeningitis virus (LCMV) and herpes simplex virus 1 (HSV-1). Here, we will discuss the impaired control of cell-mediated immunity associated with high accumulation of LAG-3 after infection, in most cases associated with a high bacterial/viral load, a reduced survival rate or persisting metabolic and inflammation disorders. Interestingly, the in vitro blockade of PD-1/LAG-3 interactions enhanced cytokine production in response to some of these infections.

Keywords: LAG-3; Immune Checkpoint