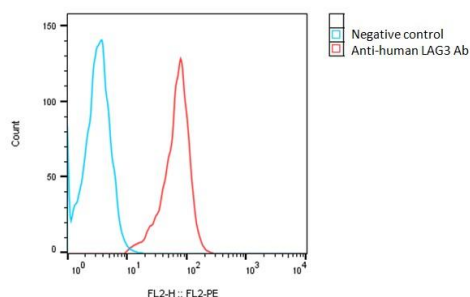


**SPECIFICATIONS**

<b>Catalog Number</b>	C3071
<b>Cell Line Name</b>	Cynomolgus LAG-3-CHO-K1 stable cell line
<b>Accession Number</b>	XP_045220577.1
<b>Host Cell</b>	Adherent CHO-K1
<b>Quantity</b>	Two vials of frozen cells (2x10 <sup>6</sup> per vial)
<b>Culture Medium</b>	DMEM with 10% FBS, 4µg/ml puromycin
<b>Freezing Medium</b>	90% FBS and 10% DMSO
<b>Storage</b>	Liquid nitrogen

**DATA**

Detection of human LAG-3 expression on cynomolgus LAG-3-CHO-K1 stable cells using a PE anti-human LAG-3 antibody (BioLegend, Cat. #369305).


**BACKGROUND**

LAG-3 (Lymphocyte-activation gene 3) is a protein receptor in the immunoglobulin superfamily expressed on various immune cells. LAG-3 interacts with major histocompatibility complex class II (MHC-II) molecules on antigen-presenting cells (APCs) to regulate T cell activation and tolerance. By binding to MHC-II, LAG-3 can negatively regulate the activation and proliferation of T cells, promoting immune tolerance and preventing excessive immune responses. LAG-3 is involved in maintaining immune homeostasis and preventing autoimmunity. The expression of LAG-3 in cancer is often upregulated in tumor-infiltrating lymphocytes (TILs) and exhausted T cells within the tumor microenvironment leading to immune dysfunction and impaired antitumor immune responses. Tumor cells may exploit the LAG-3 pathway to evade immune surveillance and promote immune tolerance, thereby facilitating tumor growth and metastasis. The unique property of LAG-3 as an immune checkpoint molecule has led to its exploration as a potential therapeutic target in cancer immunotherapy.

**References**

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