Resource Summary Report

Generated by FDI Lab - SciCrunch.org on Apr 13, 2025

Brilliant Violet 785(TM) anti-mouse CD4

RRID:AB_2565843 Type: Antibody

Proper Citation

(BioLegend Cat# 100453, RRID:AB_2565843)

Antibody Information

URL: http://antibodyregistry.org/AB_2565843

Proper Citation: (BioLegend Cat# 100453, RRID:AB_2565843)

Target Antigen: CD4

Host Organism: rat

Clonality: monoclonal

Comments: Applications: FC

Antibody Name: Brilliant Violet 785(TM) anti-mouse CD4

Description: This monoclonal targets CD4

Target Organism: mouse

Clone ID: Clone GK1.5

Antibody ID: AB_2565843

Vendor: BioLegend

Catalog Number: 100453

Record Creation Time: 20231110T035157+0000

Record Last Update: 20240725T062033+0000

Ratings and Alerts

No rating or validation information has been found for Brilliant Violet 785(TM) anti-mouse CD4.

No alerts have been found for Brilliant Violet 785(TM) anti-mouse CD4.

Data and Source Information

Source: Antibody Registry

Usage and Citation Metrics

We found 24 mentions in open access literature.

Listed below are recent publications. The full list is available at FDI Lab - SciCrunch.org.

Zhou C, et al. (2024) Anti-tumor efficacy of HRS-4642 and its potential combination with proteasome inhibition in KRAS G12D-mutant cancer. Cancer cell, 42(7), 1286.

Bonetti L, et al. (2024) A Th17 cell-intrinsic glutathione/mitochondrial-IL-22 axis protects against intestinal inflammation. Cell metabolism, 36(8), 1726.

Li Z, et al. (2024) Therapeutic application of human type 2 innate lymphoid cells via induction of granzyme B-mediated tumor cell death. Cell, 187(3), 624.

Kenney LL, et al. (2024) mRNA-delivery of IDO1 suppresses T cell-mediated autoimmunity. Cell reports. Medicine, 5(9), 101717.

Joshi S, et al. (2024) Tim4 enables large peritoneal macrophages to cross-present tumor antigens at early stages of tumorigenesis. Cell reports, 43(4), 114096.

Osorio JC, et al. (2023) The antitumor activities of anti-CD47 antibodies require Fc-Fc?R interactions. Cancer cell, 41(12), 2051.

First NJ, et al. (2023) Bordetella spp. block eosinophil recruitment to suppress the generation of early mucosal protection. Cell reports, 42(11), 113294.

Huseni MA, et al. (2023) CD8+ T cell-intrinsic IL-6 signaling promotes resistance to anti-PD-L1 immunotherapy. Cell reports. Medicine, 4(1), 100878.

Pankhurst TE, et al. (2023) MAIT cells activate dendritic cells to promote TFH cell differentiation and induce humoral immunity. Cell reports, 42(4), 112310.

Guo M, et al. (2023) Molecular, metabolic, and functional CD4 T cell paralysis in the lymph node impedes tumor control. Cell reports, 42(9), 113047.

Briukhovetska D, et al. (2023) T cell-derived interleukin-22 drives the expression of CD155 by cancer cells to suppress NK cell function and promote metastasis. Immunity, 56(1), 143.

Soriano-Baguet L, et al. (2023) Pyruvate dehydrogenase fuels a critical citrate pool that is essential for Th17 cell effector functions. Cell reports, 42(3), 112153.

Blomberg OS, et al. (2023) IL-5-producing CD4+ T cells and eosinophils cooperate to enhance response to immune checkpoint blockade in breast cancer. Cancer cell, 41(1), 106.

Gonçalves R, et al. (2023) SARS-CoV-2 variants induce distinct disease and impact in the bone marrow and thymus of mice. iScience, 26(2), 105972.

Earley ZM, et al. (2023) GATA4 controls regionalization of tissue immunity and commensaldriven immunopathology. Immunity, 56(1), 43.

Hailemichael Y, et al. (2022) Interleukin-6 blockade abrogates immunotherapy toxicity and promotes tumor immunity. Cancer cell, 40(5), 509.

Mastandrea I, et al. (2022) Isolation and characterization of the immune cell fraction from murine brain tumor microenvironment. STAR protocols, 3(1), 101106.

Cardoso A, et al. (2021) Interleukin-10 induces interferon-?-dependent emergency myelopoiesis. Cell reports, 37(4), 109887.

Magod P, et al. (2021) Exploring the longitudinal glioma microenvironment landscape uncovers reprogrammed pro-tumorigenic neutrophils in the bone marrow. Cell reports, 36(5), 109480.

Monteiro de Oliveira Novaes JA, et al. (2021) Targeting of CD40 and PD-L1 Pathways Inhibits Progression of Oral Premalignant Lesions in a Carcinogen-induced Model of Oral Squamous Cell Carcinoma. Cancer prevention research (Philadelphia, Pa.), 14(3), 313.