abcam

Product datasheet

Anti-SLAM / CD150 antibody [9D1] - Low endotoxin, Azide free ab174599

1 Image

Overview

Product name Anti-SLAM / CD150 antibody [9D1] - Low endotoxin, Azide free

Description Rat monoclonal [9D1] to SLAM / CD150 - Low endotoxin, Azide free

Host species Rat

Tested applications
Suitable for: Flow Cyt
Species reactivity
Reacts with: Mouse

Immunogen The details of the immunogen for this antibody are not available.

Positive control Mouse thymic and splenic cell suspensions (C57Bl/6 splenocytes).

General notesThe Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Carrier free Yes

Purity Protein G purified

Clonality Monoclonal

Clone number 9D1 lsotype lgG1

Applications

1

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab174599 in the following tested applications.

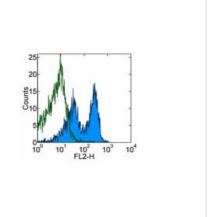
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
Flow Cyt		Use $0.5\text{-}1\mu\text{g}$ for $10^{5\text{-}8}$ cells. (Final test volume = $100~\mu\text{L}$). $\underline{ab18407}$ - Rat monoclonal lgG1, is suitable for use as an isotype control with this antibody.

Target

Function	High-affinity self-ligand important in bidirectional T-cell to B-cell stimulation. SLAM-induced signal-transduction events in T-lymphocytes are different from those in B-cells. Two modes of SLAM signaling are likely to exist: one in which the inhibitor SH2D1A acts as a negative regulator and another in which protein-tyrosine phosphatase 2C (PTPN11)-dependent signal transduction operates.	
Tissue specificity	Constitutively expressed on peripheral blood memory T-cells, T-cell clones, immature thymocytes and a proportion of B-cells, and is rapidly induced on naive T-cells after activation.	
Sequence similarities	Contains 1 lg-like C2-type (immunoglobulin-like) domain. Contains 1 lg-like V-type (immunoglobulin-like) domain.	
Domain	The most membrane-proximal SH2-binding motif interacts with SH2 domain of SH2D1A and does not need to be phosphorylated on tyrosine residues.	
Post-translational modifications	Phosphorylated by FYN.	
Cellular localization	Cell membrane. Present on the surface of B-cells and T-cells.	

Images



Flow Cytometry - Anti-SLAM / CD150 antibody [9D1] - Low endotoxin, Azide free (ab174599)

Flow cytometry analysis of C57Bl/6 splenocytes labeling SLAM / CD150 using ab174599 (blue histogram) at 0.5 μ g, or purified Rat lgG1 kappa Isotype Control (open histogram) at 0.5 μ g; followed by an anti-Rat lgG Biotin and Streptavidin PE. Cells in the lymphocyte gate were used for analysis.

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