

## Monoclonal Antibody to CD49d / ITGA4 - FITC

<b>Alternate names:</b>	CD49 antigen-like family member D, Integrin alpha-4, Integrin alpha-IV, VLA-4, VLA4
<b>Catalog No.:</b>	CL030FX
<b>Quantity:</b>	0.3 mg
<b>Concentration:</b>	0.1 mg/ml
<b>Background:</b>	<p>Integrin alpha 4 (also called CD49d) is a 150 kDa protein that possesses a large extracellular domain involved in ligand binding, a single transmembrane domain, and an intracellular regulatory domain possessing multiple sites for phosphorylation. Integrin alpha 4 forms heterodimers with integrins beta 1 and beta 7. Integrin alpha 4 is expressed on leukocytes and leukocyte precursors, neural crest cells, and developing skeletal muscles and is essential for embryogenesis, hematopoiesis, and immune responses. The presence of integrin alpha 4 promotes cell migration and inhibits cell spreading and contractility. Integrin alpha 4 function has been implicated in the pathogenesis of multiple diseases including asthma, rheumatoid arthritis, Crohn's disease, ulcerative colitis, hepatitis C, and multiple sclerosis, and therefore, modulation of integrin alpha 4 function has become an important target for drug discovery.</p>
<b>Uniprot ID:</b>	<a href="#">Q00651</a>
<b>NCBI:</b>	<a href="#">NP_034706.3</a>
<b>GeneID:</b>	<a href="#">16401</a>
<b>Host / Isotype:</b>	Rat / IgG2b
<b>Clone:</b>	R1-2
<b>Immunogen:</b>	Peyers Patch HEV binding lymphoma line (TK1)
<b>Format:</b>	<p><b>State:</b> Liquid <b>Purification:</b> Purified from ascitic fluid via Protein G Chromatography <b>Buffer System:</b> PBS, 0.02% NaN<sub>3</sub> and EIA grade BSA as a stabilizing protein to bring total protein concentration to 4-5 mg/ml. <b>Label:</b> FITC</p>
<b>Applications:</b>	<p>Flow cytometry (see protocol). Immunoprecipitation. Immunohistochemistry. (1,2,3) Other applications not tested. Optimal dilutions are dependent on conditions and should be determined by the user.</p>
<b>Specificity:</b>	<p>Antibody CL030FX reacts with α4 integrin (mouse CD49d), which helps to mediate cell-cell and cell-matrix interactions. <b>Species:</b> mouse. Other species not tested.</p>

**For research and in vitro use only. Not for diagnostic or therapeutic work.**

Material Safety Datasheets are available at [www.acris-antibodies.com](http://www.acris-antibodies.com) or on request.

Antibody Hotline - Technical Questions - Antibody Location Service  
Free Call: 0800-2274746 (Germany only) - [www.acris-antibodies.com](http://www.acris-antibodies.com)

**Storage:** Store at 4°C. For long term storage, aliquot and freeze unused portion at -20°C in volumes appropriate for single usage. Avoid prolonged exposure to light. Shelf life: one year from despatch.

**General References:** 1) Berlin, C., E. L. Berg, M. J. Briskin, D. P. Andrew, P. J. Kilshaw, B. Holzmann, I. L. Weissman, A. Hamann, E.C. Butcher 1993. a4b7 integrin mediates lymphocyte binding to the mucosal vascular addressin MAdCam-1. *Cell* 704:185-195

2) Holzmann, B., I. L. Weissman 1989. Peyer's patch-specific lymphocyte homing receptors consist of a VLA-4 like a chain associated with either of two integrin b chains, one of which is novel. *EMBO* 8:1736-1741

3) Holzmann, B., B. W. McIntyre, I. W. Weissman 1989. Identification of a murine Peyer's patch-specific lymphocyte homing receptor as an integrin molecule with an a chain homologous to human VLA-4a. *Cell* 56:37-46

**Protocols:** FLOW CYTOMETRY ANALYSIS:

Method:

1. Prepare a cell suspension in media A. For cell preparations, deplete the red blood cell population with Lympholyte®-M cell separation medium.
2. Wash 2 times.
3. Resuspend the cells to a concentration of 2x10<sup>7</sup> cells/ml in media A. Add 50 µl of this suspension to each tube (each tube will then contain 1 x 10<sup>6</sup> cells, representing 1 test).
4. To each tube, add 1.0 µg\* of CL030F per 10<sup>6</sup> cells.
5. Vortex the tubes to ensure thorough mixing of antibody and cells.
6. Incubate the tubes for 30 minutes at 4°C.  
(It is recommended that the tubes are protected from light, since most fluorochemicals are light sensitive.)
7. Wash 2 times at 4°C.
8. Resuspend the cell pellet in 50 µl ice cold media B.
9. Transfer to suitable tubes for flow cytometric analysis containing 15 µl of propidium iodide at 0.5 mg/ml in PBS. This stains dead cells by intercalating in DNA.

Media:

- A. Phosphate buffered saline (pH 7.2) + 5% normal serum of host species + sodium azide (100 µl of 2M sodium azide in 100 mls).
- B. Phosphate buffered saline (pH 7.2) + 0.5% Bovine serum albumin + sodium azide (100 µl of 2M sodium azide in 100 mls).

Results:

Tissue Distribution by Flow Cytometry Analysis:

Mouse Strain: BALB/c

Cell Concentration: 1x10<sup>6</sup> cells per tests

Antibody Concentration Used: 1.0 µg/10<sup>6</sup> cells

Isotypic Control: FITC Rat IgG2b

Cell Source Percentage of cells stained above control:

TK1 cell line 96.8%

Thymus 45.6%

Spleen 88.0%

Bone Marrow 84.7%

Strain Distribution by Flow Cytometry Analysis:

Cell Concentration: 1x10<sup>6</sup> cells per tests

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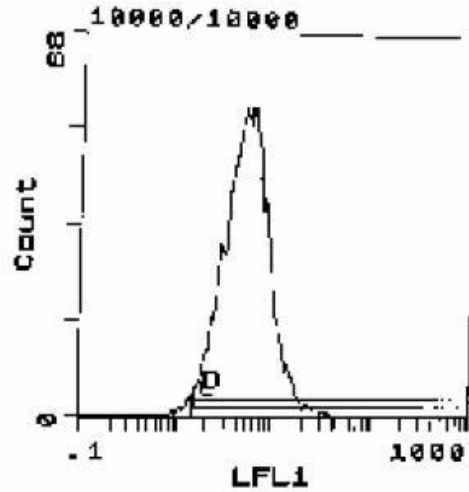
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Antibody Concentration Used: 1.0 µg / 10<sup>6</sup> cells  
Strains Tested: BALB/c, C57BL/6, C3H/He, CBA/J, AKR  
Positive: BALB/c, C57BL/6, C3H/He, CBA/J, AKR  
Negative: non

Pictures:



Cell Source: TK1 cell line  
Percentage of cells stained above control: 96.8%