

Human CD45RB APC-conjugated Antibody

Monoclonal Mouse IgG₁ Clone # 1068228 Catalog Number: FAB11434A

100 Tests

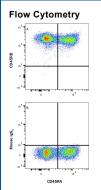
DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human CD45RB in direct ELISA.		
Source	Monoclonal Mouse IgG ₁ Clone # 1068228		
Purification	Protein A or G purified from hybridoma culture supernatant		
Immunogen	Chinese Hamster Ovary cell line, CHO-derived human CD45RB Gln26-Lys463 Accession # P08575		
Conjugate	Allophycocyanin Excitation Wavelength: 620-650 nm Emission Wavelength: 660-670 nm		
Formulation	Supplied in a saline solution containing BSA and Sodium Azide. See Certificate of Analysis for details.		
	*Contains <0.1% Sodium Azide, which is not hazardous at this concentration according to GHS classifications. Refer to the Safety Data Sheet (SDS) for additional information and handling instructions.		

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Flow Cytometry	10 μL/10 ⁶ cells	PBMC lymphocytes

DATA



Detection of CD45RB in PBMC lymphocytes cells by Flow Cytometry. PBMC lymphocytes were stained with either (A) Mouse Anti-Human CD45RB APC-conjugated Monoclonal Antibody (Catalog # FAB11434A) or (B) Mouse IgG₁ Allophycocyanin Isotype Control (Catalog # IC002A). View our protocol for Staining Membrane-associated Proteins.

PREPARATION AND STORAGE

Shipping The product is shipped with polar packs. Upon receipt, store it immediately at the temperature recommended below.

Stability & Storage

Protect from light. Do not freeze.

• 12 months from date of receipt, 2 to 8 °C as supplied.

Rev. 12/19/2023 Page 1 of 2





Human CD45RB APC-conjugated Antibody

Monoclonal Mouse IgG₁ Clone # 1068228 Catalog Number: FAB11434A

100 Tests

BACKGROUND

CD45, previously called LCA (leukocyte common antigen), T200, or Ly5 in mice, is member C of the class 1 (receptor-like) protein tyrosine phosphatase family (PTPRC) (1, 2). It is a variably glycosylated 180-220 kDa transmembrane protein that is abundantly expressed on all nucleated cells of hematopoietic origin (1-3). Multiple splicing isoforms of exon 4 (A), 5 (B), and 6 (C) are expressed according to cell type, developmental stage and antigenic exposure (1-5). The longest form, CD45RABC (called B220 in mouse) is expressed on B lymphocytes and the shortest form, CD45R0 is expressed on memory cells (5). Isoform CD45RB contains exon 5 (B exon) in the extracellular domain, which shares 41% and 40% homology with the ECD of mouse and rat CD45RB. CD45 has been best studied in T cells, where it determines T cell receptor signaling thresholds (3, 6-8). CD45 is moved into or out of the immunological synapse (IS) membrane microdomain depending on the relative influence of interaction with the extracellular galectin lattice or the intracellular actin cytoskeleton (9, 10). Interaction of galaction can be fine-tuned by varying usage of the heavily O-glycosylated spliced regions and sialylation of N-linked carbohydrates of CD45 (4, 9). Within the immunological synapse, CD45 dephosphorylates and negatively regulates the Src family kinase, Lck (8-10). In other leukocytes, CD45 deletion causes in severe immunodeficiency, while point mutations may be associated with autoimmune disorders (6, 7).

References:

- 1. Anderson, J.N. et al. (2004) FASEB J. 18:8.
- 2. Streuli, M. et al. (1987) J. Exp. Med. 166:1548.
- 3. Hermiston, M.L. et al. (2003) Annu. Rev. Immunol. 21:107.
- 4. Earl, L.A. and L.G. Baum (2008) Immunol. Cell Biol. 86:608.
- 5. Ralph, S.J. et al. (1987) EMBO J. 6:1251.
- 6. Falahti, R. and D. Leitenberg (2008) J. Immunol. 181:6082.
- 7. Tchilian, E.Z. and P.C.L. Beverley (2006) Trends Immunol. 27:146.
- 8. McNiell, L. et al. (2007) Immunity 27:425.
- 9. Chen, I-J. et al. (2007) J. Biol. Chem. 282:35361.
- 10. Freiberg, B.A. et al. (2002) Nat. Immunol. 3:911.
- 11. Zhu, J.W. et al. (2008) Immunity 28:183.
- 12. Huntington, N.D. et al. (2006) Nat. Immunol. 7:190.
- 13. Hesslein, D.G. et al. (2006) Proc. Natl. Acad. Sci. USA 103:7012.
- 14. Cross, J.L. et al. (2008) J. Immunol. 180:8020.