abcam

Product datasheet

PE/Cy5® Anti-CD45RO antibody [UCHL1] ab95520

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Overview

Product name PE/Cy5® Anti-CD45RO antibody [UCHL1]

Description PE/Cy5® Mouse monoclonal [UCHL1] to CD45RO

Host species Mouse

Conjugation PE/Cy5®. Ex: 496nm, Em: 670nm

Tested applications Suitable for: Flow Cyt

Species reactivity Reacts with: Human

Immunogen Tissue, cells or virus corresponding to Human CD45RO. IL2 dependent T cell line CA1.

Positive control Normal Human peripheral blood cells/leukocytes.

General notes

The 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.

Storage buffer pH: 7.20

Preservative: 0.09% Sodium azide

Constituents: 0.2% BSA, 0.87% Sodium chloride, PBS

Purity Protein G purified

Clonality Monoclonal

Clone number UCHL1

Isotype IgG2a

Light chain type kappa

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Applications

The Abpromise guarantee

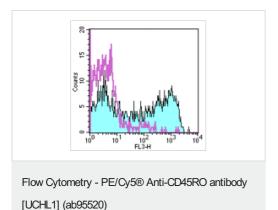
Our <u>Abpromise guarantee</u> covers the use of ab95520 in the following tested applications.

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

Application	Abreviews	Notes
Flow Cyt		Use 5µl for 10 ⁶ cells.

Target		
Function	Protein tyrosine-protein phosphatase required for T-cell activation through the antigen receptor. Acts as a positive regulator of T-cell coactivation upon binding to DPP4. The first PTPase domain has enzymatic activity, while the second one seems to affect the substrate specificity of the first one. Upon T-cell activation, recruits and dephosphorylates SKAP1 and FYN. Dephosphorylates LYN, and thereby modulates LYN activity.	
Involvement in disease	Defects in PTPRC are a cause of severe combined immunodeficiency autosomal recessive T-cell-negative/B-cell-positive/NK-cell-positive (T(-)B(+)NK(+) SCID) [MIM:608971]. A form of severe combined immunodeficiency (SCID), a genetically and clinically heterogeneous group of rare congenital disorders characterized by impairment of both humoral and cell-mediated immunity, leukopenia, and low or absent antibody levels. Patients present in infancy recurrent, persistent infections by opportunistic organisms. The common characteristic of all types of SCID is absence of T-cell-mediated cellular immunity due to a defect in T-cell development. Genetic variations in PTPRC are involved in multiple sclerosis susceptibility (MS) [MIM:126200]. MS is a neurodegenerative disorder characterized by the gradual accumulation of focal plaques of demyelination particularly in the periventricular areas of the brain. Peripheral nerves are not affected. Onset usually in third or fourth decade with intermittent progression over an extended period. The cause is still uncertain.	
Sequence similarities	Belongs to the protein-tyrosine phosphatase family. Receptor class 1/6 subfamily. Contains 2 fibronectin type-Ill domains. Contains 2 tyrosine-protein phosphatase domains.	
Domain	The first PTPase domain interacts with SKAP1.	
Post-translational modifications	Heavily N- and O-glycosylated.	
Cellular localization	Membrane. Membrane raft. Colocalized with DPP4 in membrane rafts.	

Images



Flow cytometric staining of normal Human peripheral blood cells with staining buffer (autofluorescence) (open histogram) or ab95520 (filled histogram). Cells in the lymphocyte gate were used for analysis.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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