

# CD45RO Monoclonal Antibody (UCHL1), Super Bright™ 600, eBioscience™

Product Details	
Size	100 Tests
Species Reactivity	Human
Host/Isotype	Mouse / IgG2a, kappa
Recommended Isotype Control	Mouse IgG2a kappa Isotype Control (eBM2a), Super Bright™ 600, eBioscience™
Class	Monoclonal
Type	Antibody
Clone	UCHL1
Conjugate	Super Bright™ 600
Excitation/Emission Max	414/601 nm
Form	Liquid
Concentration	5 µL/Test
Purification	Affinity chromatography
Storage buffer	PBS with BSA
Contains	0.09% sodium azide
Storage conditions	4° C, store in dark, DO NOT FREEZE!
RRID	AB_2662466

Applications	Tested Dilution	Publications
Flow Cytometry (Flow)	5 µL (0.5 µg)/Test	2 Publications

## Product Specific Information

The UCHL1 monoclonal antibody reacts with human CD45RO, a 180 kDa isoform of CD45. CD45RO is expressed by most thymocytes, activated memory T cells, granulocytes and monocytes. CD22 is a ligand for CD45RO. Expression of CD45RO and CD45RA is used commonly to discriminate subsets of peripheral T cells.

This UCHL1 antibody has been pre-titrated and tested by flow cytometric analysis of normal human peripheral blood cells. This can be used at 5 µL (0.5 µg) per test. A test is defined as the amount (µg) of antibody that will stain a cell sample in a final volume of 100 µL. Cell number should be determined empirically but can range from 10<sup>5</sup> to 10<sup>8</sup> cells/test.

Super Bright 600 is a tandem dye that can be excited with the violet laser line (405 nm) and emits at 600 nm. We recommend using a 610/20 bandpass filter. Please make sure that your instrument is capable of detecting this fluorochrome.

When using two or more Super Bright dye-conjugated antibodies in a staining panel, it is recommended to use Super Bright Complete Staining Buffer (Product # SB-4401) to minimize any non-specific polymer interactions. Please refer to the datasheet for Super Bright Staining Buffer for more information.

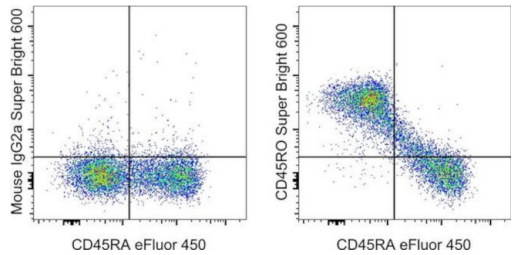
Light sensitivity: This tandem dye is sensitive to photo-induced oxidation. Protect this vial and stained samples from light.

Fixation: Samples can be stored in IC Fixation Buffer (cat. 00-8222) (100 L of cell sample + 100 L of IC Fixation Buffer) or 1-step Fix/Lyse Solution (cat. 00-5333) for up to 3 days in the dark at 2-8°C with minimal impact on brightness and FRET efficiency/compensation. Some generalizations regarding fluorochrome performance after fixation can be made, but clone specific performance should be determined empirically.

Excitation: 405 nm; Emission: 600 nm; Laser: Violet Laser

Super Bright Polymer Dyes are sold under license from Becton, Dickinson and Company.

Product Images For CD45RO Monoclonal Antibody (UCHL1), Super Bright™ 600, eBioscience™

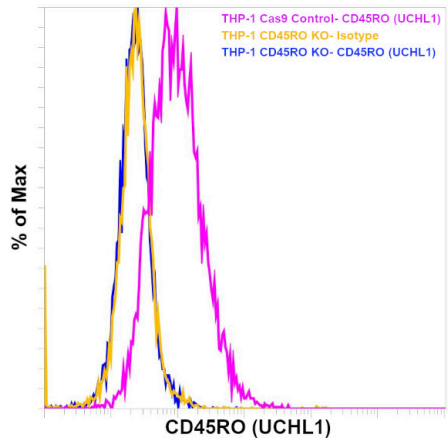


CD45RO Antibody (63-0457-42) in Flow

Staining of normal human peripheral blood cells with Anti-Human CD45RA eFluor® 450 (Product # 48-0458-42) and Mouse IgG2a K Isotype Control Super Bright 600 (Product # 63-4724-80) (left) or Anti-Human CD45RO Super Bright 600 (right). Cells in the lymphocyte gate were used for analysis.

CD45RO Antibody (63-0457-42)

Antibody clone (UCHL1) specificity was demonstrated by CRISPR-Cas9 mediated knockout of target protein. Loss of signal was observed for target protein in UCHL1 KO cells (blue histogram) compared to the control Cas9 cells (pink histogram) using CD45RO antibody (UCHL1). Yellow histogram represents staining with the isotype control. {KO}



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### Flow Cytometry (2)

#### Oncoimmunology

#### Development of CAR-T cell therapy for B-ALL using a point-of-care approach.

"Published figure using CD45RO monoclonal antibody (Product # 63-0457-42) in Flow Cytometry"

Authors: de Macedo Abdo L,Barros LRC,Saldanha Viegas M,Vieira Codeço Marques L,de Sousa Ferreira P,Chicaybam L,Bonamino MH

Year  
2021

#### Retrovirology

#### Longitudinal analysis of subtype C envelope tropism for memory CD4<sup>+</sup> T cell subsets over the first 3 years of untreated HIV-1 infection.

"Published figure using CD45RO monoclonal antibody (Product # 63-0457-42) in Flow Cytometry"

Authors: Gartner MJ,Gorry PR,Tumpach C,Zhou J,Dantanarayana A,Chang JJ,Angelovich TA,Ellenberg P,Laumaea AE,Nonyane M,Moore PL,Lewin SR,Churchill MJ,Flynn JK,Roche M

Year  
2020

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