abcam

Product datasheet

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

画像数1

製品の概要

製品名 PE/Cy5® Anti-CD45RO antibody [UCHL1]

製品の詳細 PE/Cy5® Mouse monoclonal [UCHL1] to CD45RO

由来種 Mouse

標識 PE/Cy5®. Ex: 496nm, Em: 670nm

アプリケーション 適用あり: Flow Cyt

種交差性 交差種: Human

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

ポジティブ・コントロール Normal Human peripheral blood cells/leukocytes.

特記事項

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

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C.

バッファー pH: 7.20

Preservative: 0.09% Sodium azide

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

精製度 Protein G purified

ポリ/モノ モノクローナル

クローン名 UCHL1 **アイソタイプ** lgG2a

軽鎖の種類 kappa

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The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab95520の使用に適用されますアプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
Flow Cyt		Use 5µl for 10 ⁶ cells.

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機能

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.

関連疾患

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.

配列類似性

Belongs to the protein-tyrosine phosphatase family. Receptor class 1/6 subfamily. Contains 2 fibronectin type-III domains.

Contains 2 tyrosine-protein phosphatase domains.

ドメイン

The first PTPase domain interacts with SKAP1.

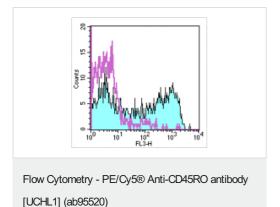
翻訳後修飾

Heavily N- and O-glycosylated.

細胞内局在

Membrane. Membrane raft. Colocalized with DPP4 in membrane rafts.

画像



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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