abcam

Product datasheet

Anti-CD45RB antibody [16A] (FITC) ab112464

画像数1

製品の概要

製品名 Anti-CD45RB antibody [16A] (FITC)

製品の詳細 Rat monoclonal [16A] to CD45RB (FITC)

由来種 Rat

標識 FITC. Ex: 493nm, Em: 528nm

アプリケーション 適用あり: Flow Cyt 種交差性 交差種: Mouse

免疫原 Mouse TH₂ cell clones, final boost with TH₂ clone D10.

ポジティブ・コントロール Mouse spleen cells (BALB/c strain).

特記事項 Abcam is committed to meeting high standards of ethical manufacturing and as such, we will be

discontinuing this product, which has been generated by the ascites method, within the next year. We are sorry for any inconvenience this may cause. If you would like help finding an alternative

product, please do not hesitate to contact our scientific support team.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C.

ארע"ד Preservative: 0.1% Sodium azide

Constituents: 0.5% BSA, PBS

精製度 Protein G purified

ポリ/モノ モノクローナル

クローン名 16A

₹IП-▼ Ag8.653

アイソタイプ lgG2a

アプリケーション

Our Abpromise guarantee covers the use of ab112464 in the following tested applications.

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

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アプリケーション	Abreviews	特記事項
Flow Cyt		Use 1µg for 10 ⁶ cells. ab18446 - Rat monoclonal lgG2a, is suitable for use as an isotype control with this antibody.

ターゲット情報

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.

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.

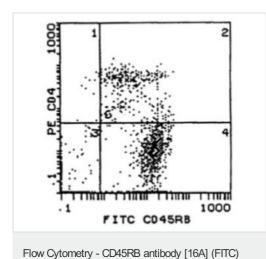
Contains 2 tyrosine-protein phosphatase domains.

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 raft. Colocalized with DPP4 in membrane rafts.

画像



ab112464 at $1\mu g/10^6$ cells staining CD45RB in Mouse spleen cells (BALB/c strain) by Flow Cytometry.

Percentage of cells stained above control (FITC Rat lgG2a): 79.8%.

(ab112464)

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