abcam

Product datasheet

Anti-CD45 antibody [YTH24.5] ab30446

7 References 1 图像

概述

产品名称 Anti-CD45抗体[YTH24.5]

宿主 Rat

经测试应用适用于: Flow Cyt种属反应性与反应: Human

免疫原 Tissue, cells or virus corresponding to Human CD45. Human T lymphocytes.

常规说明

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

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

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

性能

形式 Liquid

存放说明 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

存储溶液 Preservative: 0.09% Sodium azide

Constituent: PBS

纯**度** Protein A purified

纯**化**说明 Immunoglobulin fraction prepared by ammonium sulphate precipitation.

 克隆
 单克隆

 克隆编号
 YTH24.5

 骨髓瘤
 Y3/Ag1.2.3

 同种型
 IgG2b

应用

The Abpromise guarantee

Abpromise™承诺保证使用ab30446于以下的经测试应用

"应用说明"部分 下显示的仅为推荐的起始稀释度;实际最佳的稀释度/浓度应由使用者检定。

应用	Ab评论	说明
Flow Cyt		Use 1µg for 10 ⁶ cells. ab18536 - Rat monoclonal lgG2b, is suitable for use as an isotype control with this antibody.

靶标

功能

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.

疾病相关

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.

序列相似性

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

Contains 2 tyrosine-protein phosphatase domains.

结**构域**

The first PTPase domain interacts with SKAP1.

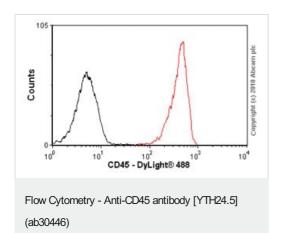
翻译后修饰

Heavily N- and O-glycosylated.

细胞定位

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

图片



Overlay histogram showing peripheral blood lymphocytes stained with ab30446 (red line). The cells were incubated with the antibody (ab30446, 1µg/1x10⁶ cells) for 30 min at 4°C. The secondary antibody used was DyLight® 488 goat anti-rat lgG (Fc) (ab96971) at 1/250 dilution for 30 min at 22°C. Isotype control antibody (black line) was rat lgG2b [RTK4530] (ab18541, 2µg/1x10⁶ cells) used under the same conditions. Acquisition of >5,000 events was performed gating on peripheral blood lymphocytes.

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

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