abcam

Product datasheet

FITC Anti-CD105 antibody [MEM-229] ab53318

20 References 画像数 1

製品の概要

製品名 FITC Anti-CD105 antibody [MEM-229]

製品の詳細 FITC Mouse monoclonal [MEM-229] to CD105

由来種 Mouse

標識 FITC. Ex: 493nm, Em: 528nm

特異性 The antibody MEM-229 recognizes an extracellular epitope of CD105 (Endoglin), a 90 kDa type I

integral membrane homodimer glycoprotein expressed on vascular endothelial cells (small and large vessels), activated monocytes and tissue macrophages, stromal cells of certain tissues

including bone marrow, pre-B

lymphocytes in fetal marrow and erythroid precursors in fetal and adult bone marrow; it is also

present on syncytiotrophoblast on placenta throughout pregnancy.

アプリケーション 適用あり: IHC-Fr

種交差性 交差種: Pig

免疫原 Tissue, cells or virus corresponding to Human CD105.

Database link: P17813

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: 7.4

Preservative: 0.097% Sodium azide

Constituents: 0.2% BSA, PBS

精製度 Size exclusion

特記事項(精製) Purified by size-exclusion chromatography.

1

ポリÆノ モノクローナル **クローン名** MEM-229 アイソタイプ lgG2a

アプリケーション

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ab53318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
IHC-Fr		1/200. Fix with acetone.

ターゲット情報

機能 Major glycoprotein of vascular endothelium. May play a critical role in the binding of endothelial

cells to integrins and/or other RGD receptors.

組織特異性 Endoglin is restricted to endothelial cells in all tissues except bone marrow.

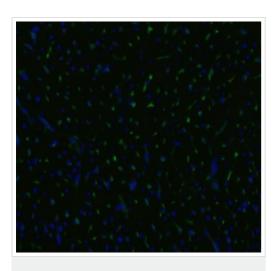
関連疾患 Defects in ENG are the cause of hereditary hemorrhagic telangiectasia type 1 (HHT1)

[MIM:187300, 108010]; also known as Osler-Rendu-Weber syndrome 1 (ORW1). HHT1 is an autosomal dominant multisystemic vascular dysplasia, characterized by recurrent epistaxis, muco-cutaneous telangiectases, gastro-intestinal hemorrhage, and pulmonary (PAVM), cerebral (CAVM) and hepatic arteriovenous malformations; all secondary manifestations of the underlying vascular dysplasia. Although the first symptom of HHT1 in children is generally nose bleed, there

is an important clinical heterogeneity.

細胞内局在 Membrane.

画像



Immunohistochemistry (Frozen sections) - FITC
Anti-CD105 antibody [MEM-229] (ab53318)

ab53318 staining CD105 in infarcted porcine heart by Immunohistochemistry (Frozen sections). Cell nuclei were counterstained blue with DAPI.

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