# abcam

# Product datasheet

# Anti-CD105 antibody [EPR10145-12] - BSA and Azide free ab271922



リコンピナント

RabMAb

★★★★★ 1 Abreviews

画像数7

#### 製品の概要

製品名 Anti-CD105 antibody [EPR10145-12] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR10145-12] to CD105 - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: WB, IHC-P

種交差性 交差種: Human

免疫原 Recombinant fragment corresponding to Human CD105 aa 1-200.

Database link: P17813

ポジティブ・コントロール WB: ECV-304 and HUVEC cell lysates, human tonsil tissue lysate and immunoprecipitation pellet

from ECV-304 cell lysate. IHC-P: Human glioma, clear cell carcinoma, tonsil and kidney tissues.

特記事項 ab271922 is the carrier-free version of ab169545.

Our <u>carrier-free</u> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for

increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our **conjugation kits** for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar $^{(\!0\!)}$  Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar $^{(\!0\!)}$  is a trademark of Fluidigm Canada Inc.

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit

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#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C. Do Not Freeze.

**バッファー** pH: 7.2

Constituent: PBS

キャリア・フリー はい

精製度 Protein A purified

**ポリ/モノ** モノクローナル

**クローン名** EPR10145-12

アイソタイプ lgG

#### 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ab271922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
WB		Use at an assay dependent concentration. Predicted molecular weight: 70 kDa.  For unpurified use at 1/50.
IHC-P	**** (1)	Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.  See IHC antigen retrieval protocols. For unpurified use at 1/30.

## 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.

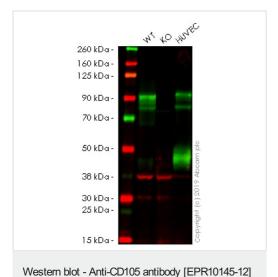
組織特異性 Endoglin is restricted to endothelial cells in all tissues except bone marrow.

**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.



- BSA and Azide free (ab271922)

**All lanes :** Anti-CD105 antibody [EPR10145-12] (**ab169545**) at 1/1000 dilution

Lane 1: Wild-type HeLa whole cell lysate

Lane 2: CD105 knockout HeLa whole cell lysate

Lane 3: HUVEC whole cell lysate

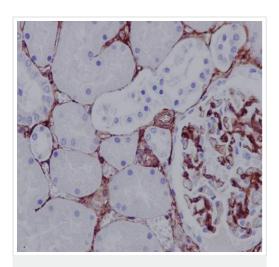
Lysates/proteins at 20 µg per lane.

Predicted band size: 70 kDa

**Lanes 1 - 3:** Merged signal (red and green). Green - <u>ab169545</u> observed at 70 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

<u>ab169545</u> was shown to recognize CD105 in wild-type HeLa cells as signal was lost at the expected MW in CD105 knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and CD105 knockout samples were subjected to SDS-PAGE. The membrane was blocked with 3% milk. Ab169545 and <u>ab8245</u> (Mouse anti-GAPDH loading control) were incubated overnight at 4°C at 1/1000 dilution and 1/20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed <u>ab216773</u> and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed <u>ab216776</u> secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab169545</u>).

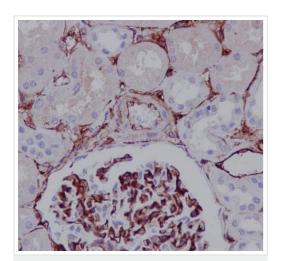


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD105 antibody

[EPR10145-12] - BSA and Azide free (ab271922)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue labelling CD105 with purified <a href="mailto:ab169545">ab169545</a> at 1/900. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (**ab169545**).

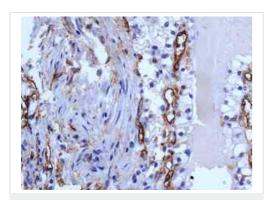


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD105 antibody

[EPR10145-12] - BSA and Azide free (ab271922)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human kidney tissue labelling CD105 with unpurified <a href="mailto:ab169545">ab169545</a> at 1/30. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. A prediluted HRP-polymer conjugated anti-rabbit lgG was used as the secondary antibody. Counterstained with hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab169545</u>).

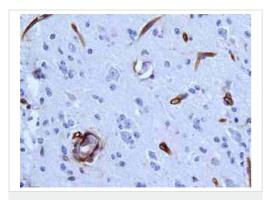


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD105 antibody

[EPR10145-12] - BSA and Azide free (ab271922)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis human clear cell carcinoma tissue labelling CD105 with unpurified <u>ab169545</u> at 1/250.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab169545</u>).

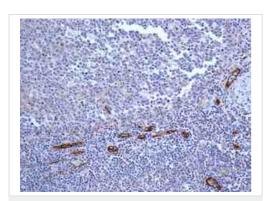


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD105 antibody

[EPR10145-12] - BSA and Azide free (ab271922)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human glioma tissue labelling CD105 with unpurified <u>ab169545</u> at 1/250.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab169545).

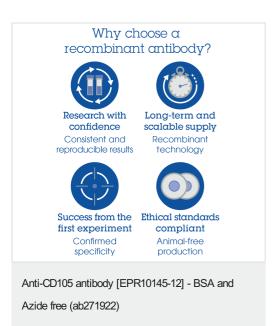


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD105 antibody

[EPR10145-12] - BSA and Azide free (ab271922)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human tonsil tissue labelling CD105 with unpurified <u>ab169545</u> at 1/250.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab169545</u>).



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