abcam

Product datasheet

Alexa Fluor® 647 Anti-Von Willebrand Factor antibody [EPSISR15] ab195029

ייבעבעדער RabMAb

画像数 2

製品の概要

製品名 Alexa Fluor® 647 Anti-Von Willebrand Factor antibody [EPSISR15]

製品の詳細 Alexa Fluor® 647 Rabbit monoclonal [EPSISR15] to Von Willebrand Factor

由来種 Rabbit

標識 Alexa Fluor® 647. Ex: 652nm. Em: 668nm

アプリケーション 適用あり: ICC/IF 種交差性 交差種: Human

免疫原 Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

ポジティブ・コントロール ICC/IF: HepG2 cells

特記事項 Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb® patents**.

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outlicensing@thermofisher.com.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -20°C. Avoid freeze / thaw cycle.

Store In the Dark.

バッファー pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

精製度 Protein A purified

ポリ/モノ モノクローナル **クローン名** EPSISR15

アイソタイプ lgG

アプリケーション

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アプリケーション	Abreviews	特記事項
ICC/IF		1/50.

ターゲット情報

機能 Important in the maintenance of hemostasis, it promotes adhesion of platelets to the sites of

vascular injury by forming a molecular bridge between sub-endothelial collagen matrix and platelet-surface receptor complex GPlb-IX-V. Also acts as a chaperone for coagulation factor VIII, delivering it to the site of injury, stabilizing its heterodimeric structure and protecting it from

premature clearance from plasma.

組織特異性 Plasma.

関連疾患 Defects in VWF are the cause of von Willebrand disease (VWD) [MIM:277480]. VWD defines a

group of hemorrhagic disorders in which the von Willebrand factor is either quantitatively or qualitatively abnormal resulting in altered platelet function. Symptoms vary depending on severity and disease type but may include prolonged bleeding time, deficiency of factor VIII and impaired platelet adhesion. Type I von Willebrand disease is the most common form and is characterized by partial quantitative plasmatic deficiency of an otherwise structurally and functionally normal Willebrand factor; type II is associated with a qualitative deficiency and functional anomalies of the

Willebrand factor; type III is the most severe form and is characterized by total or near-total absence of Willebrand factor in the plasma and cellular compartments, also leading to a profound

deficiency of plasmatic factor VIII.

配列類似性 Contains 1 CTCK (C-terminal cystine knot-like) domain.

Contains 4 TIL (trypsin inhibitory-like) domains.

Contains 3 VWFA domains. Contains 3 VWFC domains. Contains 4 VWFD domains.

ドメイン The von Willebrand antigen 2 is required for multimerization of vWF and for its targeting to

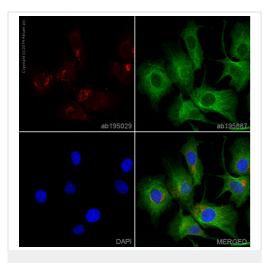
storage granules.

翻訳後修飾 All cysteine residues are involved in intrachain or interchain disulfide bonds.

N- and O-glycosylated.

細胞内局在 Secreted. Secreted > extracellular space > extracellular matrix. Localized to storage granules.

画像



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 647 Anti-Von Willebrand Factor antibody [EPSISR15] (ab195029) ab195029 staining Von Willebrand Factor in HepG2 cells. The cells were fixed with 100% methanol (5 min), permeabilised in 0.1% Triton X-100 for 5 minutes and then blocked in 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated with ab195029 at 1/50 dilution (shown in red) and ab195887, Mouse monoclonal [DM1A] to alpha Tubulin (Alexa Fluor® 488, shown in green) at 2µg/ml overnight at +4°C. Nuclear DNA was labelled in blue with DAPI.

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).



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