abcam

Product datasheet

Anti-ErbB3 / HER3 (phospho Y1289) antibody [EPR2325] ab76469

יובעדער RabMAb

5 References 画像数3

製品の概要

製品名 Anti-ErbB3 / HER3 (phospho Y1289) antibody [EPR2325]

製品の詳細 Rabbit monoclonal [EPR2325] to ErbB3 / HER3 (phospho Y1289)

由来種 Rabbit

特異性 This antibody only detects ErbB 3 phosphorylated on Tyrosine 1289.

アプリケーション 適用あり: WB. ICC/IF

適用なし: Flow Cyt,IHC-P or IP

交差種: Human 種交差性

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 SKBR3 cell lysate treated with Neuregulin. MCF7 cells.

特記事項 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 monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

Rat: We have preliminary internal testing data to indicate this antibody may not react with this

species. Please contact us for more information.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

バッファー

Preservative: 0.01% Sodium azide

Constituents: 9% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA, 50% Tissue culture

supernatant

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精製度 Protein A purified

ポリ/モノ モノクローナル

クローン名 EPR2325

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ab76469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		1/2500 - 1/5000. Detects a band of approximately 185 kDa (predicted molecular weight: 148 kDa).
ICC/IF		1/100.

追加情報 Is unsuitable for Flow Cyt,IHC-P or IP.

ターゲット情報

機能 Binds and is activated by neuregulins and NTAK.

組織特異性 Epithelial tissues and brain.

関連疾患 Defects in ERBB3 are the cause of lethal congenital contracture syndrome type 2 (LCCS2)

[MIM:607598]; also called Israeli Bedouin multiple contracture syndrome type A. LCCS2 is an autosomal recessive neurogenic form of a neonatally lethal arthrogryposis that is associated with atrophy of the anterior horn of the spinal cord. The LCCS2 syndrome is characterized by multiple joint contractures, anterior horn atrophy in the spinal cord, and a unique feature of a markedly distended urinary bladder. The phenotype suggests a spinal cord neuropathic etiology.

配列類似性 Belongs to the protein kinase superfamily. Tyr protein kinase family. EGF receptor subfamily.

Contains 1 protein kinase domain.

発生段階 Overexpressed in a subset of human mammary tumors.

ドメイン The cytoplasmic part of the receptor may interact with the SH2 or SH3 domains of many signal-

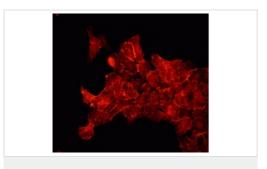
transducing proteins.

翻訳後修飾 Ligand-binding increases phosphorylation on tyrosine residues and promotes its association with

the p85 subunit of phosphatidylinositol 3-kinase.

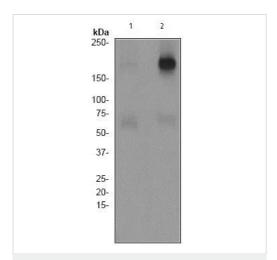
細胞内局在 Secreted and Cell membrane.

画像



Immunocytochemistry/ Immunofluorescence - Anti-ErbB3 / HER3 (phospho Y1289) antibody [EPR2325] (ab76469)

Immunofluorescent staining of MCF7 cells using ab76469 at 1/100 dilution.



Western blot - Anti-ErbB3 / HER3 (phospho Y1289) antibody [EPR2325] (ab76469)

All lanes: Anti-ErbB3 / HER3 (phospho Y1289) antibody [EPR2325] (ab76469) at 1/5000 dilution

Lane 1: SKBR3 cell lysate untreated

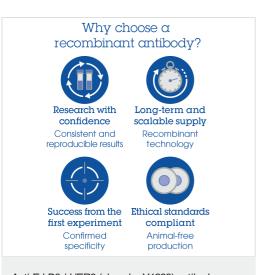
Lane 2: SKBR3 cell lysate treated with Neuregulin

Lysates/proteins at 10 µg per lane.

Secondary

All lanes: HRP labelled goat anti-rabbit at 1/2000 dilution

Predicted band size: 148 kDa **Observed band size:** 185 kDa



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