abcam

Product datasheet

Anti-Phospholipase C gamma 1/PLC-gamma-1 antibody [EPR5358] ab109501



יוטבעבער RabMAb

画像数 5

製品の概要

製品名 Anti-Phospholipase C gamma 1/PLC-gamma-1 antibody [EPR5358]

製品の詳細 Rabbit monoclonal [EPR5358] to Phospholipase C gamma 1/PLC-gamma-1

由来種 Rabbit

アプリケーション 適用あり: Flow Cyt (Intra), WB

適用なし: ICC/IF,IHC-P or IP

種交差性 交差種: Mouse, Human

交差が予測される動物種: Rat 🔷

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 WB: HEK293-T, HepG2, Jurkat, and MCF-7 cell lysates. Flow Cyt (intra): Jurkat 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**.

製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

バッファー pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.5% BSA

精製度 Protein A purified

ポリモノ モノクローナル クローン名 **EPR5358**

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ab109501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
Flow Cyt (Intra)		1/10 - 1/100. ab172730 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
WB		1/1000 - 1/10000. Predicted molecular weight: 149 kDa.

追加情報

Is unsuitable for ICC/IF,IHC-P or IP.

ターゲット情報

機能 Plays a role in actin reorganization and cell migration. The production of the second messenger molecules diacylglycerol (DAG) and inositol 1,4,5-trisphosphate (IP3) is mediated by activated

phosphatidylinositol-specific phospholipase C enzymes. Major substrate for heparin-binding

growth factor 1 (acidic fibroblast growth factor)-activated tyrosine kinase.

配列類似性 Contains 1 C2 domain.

Contains 1 EF-hand domain. Contains 2 PH domains.

Contains 1 PI-PLC X-box domain.
Contains 1 PI-PLC Y-box domain.

Contains 2 SH2 domains. Contains 1 SH3 domain.

ドメイン The SH3 domain mediates interaction with CLNK (By similarity). The SH3 domain also mediates

interaction with RALGPS1.

翻訳後修飾 The receptor-mediated activation of PLC-gamma-1 and PLC-gamma-2 involves their

phosphorylation by tyrosine kinases in response to ligation of a variety of growth factor receptors

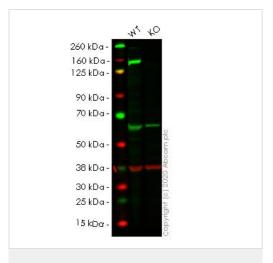
and immune system receptors. May be dephosphorylated by PTPRJ.

Ubiquitinated by CBLB in activated T-cells.

細胞内局在 Cell projection > lamellipodium. Cell projection > ruffle. Rapidly redistributed to ruffles and

lamellipodia structures in response to epidermal growth factor (EGF) treatment.

画像



Western blot - Anti-Phospholipase C gamma 1/PLCgamma-1 antibody [EPR5358] (ab109501)

All lanes : Anti-Phospholipase C gamma 1/PLC-gamma-1 antibody [EPR5358] (ab109501) at 1/1000 dilution

Lane 1: Wild-type HEK-293T cell lysate

Lane 2: PLCG1 knockout HEK-293T cell lysate

Lysates/proteins at 40 µg/ml per lane.

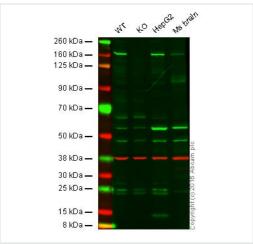
Secondary

All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) at 1/20000 dilution

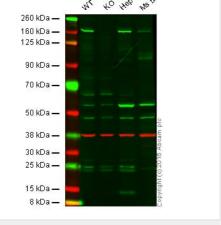
Predicted band size: 149 kDa

Lanes 1-2: Merged signal (red and green). Green - ab109501 observed at 160 kDa. Red - Anti-GAPDH antibody [6C5] - Loading Control (ab8245) observed at 37 kDa.

ab109501 was shown to react with Phospholipase C gamma 1/PLC-gamma-1 in wild-type HEK-293T cells in western blot. Loss of signal was observed when knockout cell line ab266530 (knockout cell lysate ab266530 (knockout cell lysate ab266530 (knockout cell lysate ab266530 (knockout HEK-293T cell lysates were subjected to SDS-PAGE. Membrane was blocked for 1 hour at room temperature in 0.1% TBST with 3% non-fat dried milk. ab109501 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at a 1 in 1000 dilution and a 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-Phospholipase C gamma 1/PLCgamma-1 antibody [EPR5358] (ab109501)



knockout samples were used, along with additional cross-reactive bands. Wild-type and Phospholipase C gamma 1 / PLC-gamma-1 knockout samples were subjected to SDS-PAGE. ab109501 at a

Lane 1: Wild-type HAP1 cell lysate (20 µg)

Lane 4: Mouse brain tissue lysate (20 µg)

HAP1 cell lysate (20 µg)

at 37 kDa.

Lane 3: HepG2 cell lysate (20 µg)

Lane 2: Phospholipase C gamma 1 / PLC-gamma-1 knockout

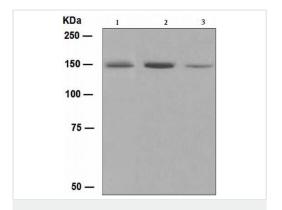
Lanes 1 - 4: Merged signal (red and green). Green - ab109501

observed at 160 kDa. Red - loading control, ab8245, observed

ab109501 was shown to recognize Phospholipase C gamma 1 /

PLC-gamma-1 when Phospholipase C gamma 1 / PLC-gamma-1

dilution of 1/1000 and ab8245 (loading control to GAPDH) at a dilution of 1/10000 were incubated overnight at 4°C. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed ab216776 secondary antibodies at 1/10000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-Phospholipase C gamma 1/PLCgamma-1 antibody [EPR5358] (ab109501)

All lanes: Anti-Phospholipase C gamma 1/PLC-gamma-1 antibody [EPR5358] (ab109501) at 1/1000 dilution

Lane 1: HepG2 cell lysate

Lane 2: Jurkat cell lysate

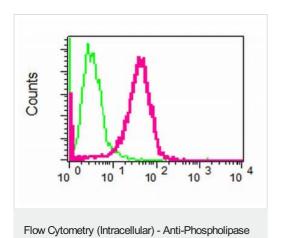
Lane 3: MCF-7 cell lysates

Lysates/proteins at 10 µg per lane.

Secondary

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

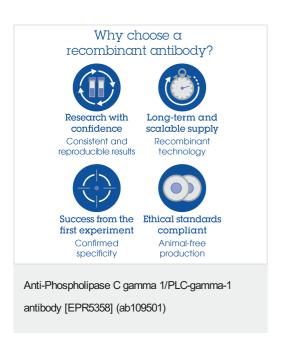
Predicted band size: 149 kDa



C gamma 1/PLC-gamma-1 antibody [EPR5358]

(ab109501)

Intracellular flow cytometric analysis of permeabilized Jurkat cells using ab109501 at 1/10 dilution (red)or a rabbit lgG (negative) (green).



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