abcam

Product datasheet

Anti-PERP antibody [EPR7885] - BSA and Azide free ab248288



יעלאעבני RabMAb

画像数3

製品の概要

特記事項

製品名 Anti-PERP antibody [EPR7885] - BSA and Azide free

製品の詳細 Rabbit monoclonal [EPR7885] to PERP - BSA and Azide free

由来種 Rabbit

アプリケーション 適用あり: ICC/IF, WB

適用なし: Flow Cyt,IHC-P or IP

種交差性 交差種: Mouse, Rat, 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 WB: HeLa and HaCaT cell lysates.

ab248288 is the carrier-free version of ab129083.

Our carrier-free 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 cellbased 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[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar® 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.

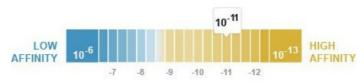
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb patents**.

製品の特性

製品の状態 Liquid

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

解離定数(K_D値) $K_D = 4.10 \times 10^{-11} M$



Learn more about K_D

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 EPR7885

アイソタイプ IgG

アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab248288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。

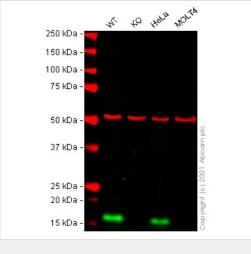
アプリケーション	Abreviews	特記事項
ICC/IF		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 21 kDa.

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

ターゲット情報

機能	Component of intercellular desmosome junctions. Plays a role in stratified epithelial integrity and cell-cell adhesion by promoting desmosome assembly. Plays a role as an effector in the TP53-dependent apoptotic pathway.	
組織特異性	Expressed in skin, heart, placental, liver, pancreas, keratinocytes and dermal fibroblasts.	
配列類似性	Belongs to the TMEM47 family.	
細胞内局在	Cell junction > desmosome. Cell membrane. Associated with desmosomes.	

画像



Western blot - Anti-PERP antibody [EPR7885] - BSA and Azide free (ab248288)

All lanes : Anti-PERP antibody [EPR7885] (**ab129083**) at 1/1000 dilution

Lane 2: PERP knockout A431 cell lysate

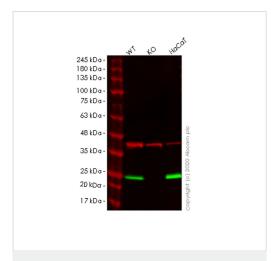
Lane 4: MOLT-4 cell lysate

Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 21 kDa **Observed band size:** 17 kDa

False colour image of Western blot: Anti-PERP antibody [EPR7885] staining at 1/1000 dilution, shown in green; Mouse anti-Alpha Tubulin [DM1A] (ab7291) loading control staining at 1/20000 dilution, shown in red. In Western blot, ab129083 was shown to bind specifically to PERP. A band was observed at 17 kDa in wild-type A431 cell lysates with no signal observed at this size in PERP knockout cell line ab270475 (knockout cell lysate ab270498). To generate this image, wild-type and PERP knockout A431 cell lysates were analysed. First, samples were run on an SDS-PAGE gel then transferred onto a nitrocellulose membrane. Membranes were blocked in 3 % milk in TBS-0.1 % Tween® 20 (TBS-T) before incubation with primary antibodies overnight at 4 °C. Blots were washed four times in TBS-T, incubated with secondary antibodies for 1 h at room temperature, washed again four times then imaged. Secondary antibodies used were Goat anti-Rabbit IgG H&L (IRDye $^{\text{\tiny{(R)}}}$ 800CW) preabsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preabsorbed (ab216776) at 1/20000 dilution.



Western blot - Anti-PERP antibody [EPR7885] - BSA and Azide free (ab248288)

All lanes : Anti-PERP antibody [EPR7885] (ab129083) at 1/1000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: PERP knockout HeLa cell lysate

Lane 3: HaCaT cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) at 1/10000 dilution

Predicted band size: 21 kDa **Observed band size:** 24 kDa

This data was developed using the same antibody clone in a different buffer formulation (ab129083).

Lanes 1-3: Merged signal (red and green). Green - <u>ab129083</u> observed at 24 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

ab129083 Anti-PERP antibody [EPR7885] was shown to specifically react with PERP in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265335 (knockout cell lysate ab258105) was used. Wild-type and PERP knockout samples were subjected to SDS-PAGE. ab129083 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated at room temperature for 2.5 hours at 1 in 1000 dilution and 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.



Anti-PERP antibody [EPR7885] - BSA and Azide free (ab248288)

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