abcam

Product datasheet

Anti-CLPTM1 antibody [EPR8800] - BSA and Azide free ab248641





RabMAb

3 Images

Overview

Product name Anti-CLPTM1 antibody [EPR8800] - BSA and Azide free

Description Rabbit monoclonal [EPR8800] to CLPTM1 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: ICC/IF, WB, IHC-P, Flow Cyt (Intra)

Unsuitable for: IP

Species reactivity Reacts with: Human

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

Positive control WB: HAP1, A549 and HeLa cell lysates. Flow Cyt: A549 cells

General notes ab248641 is the carrier-free version of **ab133756**.

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 <u>conjugation kits</u> 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**.

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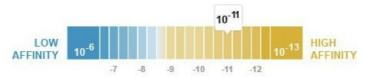
Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with these species. Please contact us for more information.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C. Do Not Freeze.

Dissociation constant (K_D) $K_D = 3.45 \times 10^{-11} M$



Learn more about K_D

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

ClonalityMonoclonalClone numberEPR8800

Isotype IgG

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab248641 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
ICC/IF		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Detects a band of approximately 90 kDa (predicted molecular weight: 76 kDa).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
Flow Cyt (Intra)		Use at an assay dependent concentration.

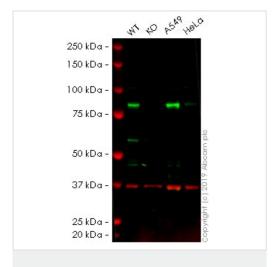
Application notes Is unsuitable for IP.

Target

Function May play a role in T-cell development.

Tissue specificity Widely expressed.

Images



Western blot - Anti-CLPTM1 antibody [EPR8800] - BSA and Azide free (ab248641)

All lanes : Anti-CLPTM1 antibody [EPR8800] (<u>ab133756</u>) at 1/1000 dilution

Lane 1: Wild-type HAP1 whole cell lysate

Lane 2: CLPTM1 knockout HAP1 whole cell lysate

Lane 3: A549 whole cell lysate
Lane 4: HeLa whole cell lysate

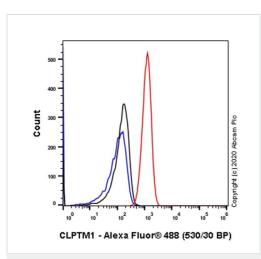
Lysates/proteins at 20 µg per lane.

Predicted band size: 76 kDa **Observed band size:** 76 kDa

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab133756</u> observed at 76 kDa. Red - loading control, <u>ab8245</u>, observed at 37 kDa.

ab133756 was shown to recognize in wild-type HAP1 cells as signal was lost at the expected MW in CLPTM1 knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and CLPTM1 knockout samples were subjected to SDS-PAGE. The membrane was blocked with 3pc Milk. Ab133756 and ab8245 (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 ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.

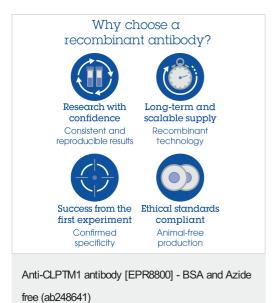
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab133756</u>).



Flow Cytometry (Intracellular) - Anti-CLPTM1 antibody [EPR8800] - BSA and Azide free (ab248641)

This data was developed using <u>ab133756</u>, the same antibody clone in a different buffer formulation.

Flow Cytometry analysis of A549 (Human lung carcinoma epithelial cell) cells labeling CLPTM1 with purified <u>ab133756</u> at 1/200 dilution (1 µg/mL) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit lgG (Alexa Fluor[®] 488, <u>ab150077</u>) secondary antibody was used at 1/2000. Isotype control - Rabbit monoclonal lgG (Black). Unlabeled control - Cell without incubation with primary antibody and secondary antibody (Blue).



Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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