abcam

Product datasheet

Anti-PI 3 Kinase catalytic subunit alpha/PIK3CA antibody [EPR19693] - BSA and Azide free ab223532





3 Images

Overview

Product name Anti-PI3 Kinase catalytic subunit alpha/PIK3CA antibody [EPR19693] - BSA and Azide free

Description Rabbit monoclonal [EPR19693] to PI 3 Kinase catalytic subunit alpha/PIK3CA - BSA and Azide

free

Host species Rabbit

Suitable for: IP, WB **Tested applications**

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: F9 and M1 whole cell lysates; Mouse hypothalamus and fetal brain lysates; P0 Rat brain

lysate. IP: Mouse hypothalamus lysate.

General notes ab223532 is the carrier-free version of ab183957.

> 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**.

Properties

Form Liquid

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

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

ClonalityMonoclonalClone numberEPR19693

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab223532 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 |
|-------------|-----------|---|
| IP | | Use at an assay dependent concentration. |
| WB | | Use at an assay dependent concentration. Detects a band of approximately 110 kDa (predicted molecular weight: 124 kDa). |

Target

Function

Phosphorylates Ptdlns, Ptdlns4P and Ptdlns(4,5)P2 with a preference for Ptdlns(4,5)P2.

Involvement in disease

Defects in PIK3CA are associated with colorectal cancer (CRC) [MIM:114500].

Defects in PIK3CA are a cause of susceptibility to breast cancer (BC) [MIM:114480]. A common

malignancy originating from breast epithelial tissue. Breast neoplasms can be distinguished by their histologic pattern. Invasive ductal carcinoma is by far the most common type. Breast cancer is etiologically and genetically heterogeneous. Important genetic factors have been indicated by familial occurrence and bilateral involvement. Mutations at more than one locus can be involved in different families or even in the same case.

Defects in PIK3CA are a cause of susceptibility to ovarian cancer (OC) [MIM:167000]. Ovarian cancer common malignancy originating from ovarian tissue. Although many histologic types of ovarian neoplasms have been described, epithelial ovarian carcinoma is the most common form. Ovarian cancers are often asymptomatic and the recognized signs and symptoms, even of latestage disease, are vague. Consequently, most patients are diagnosed with advanced disease. Defects in PIK3CA may underlie hepatocellular carcinoma (HCC) [MIM:114550].

Defects in PIK3CA are a cause of keratosis seborrheic (KERSEB) [MIM:182000]. A common benign skin tumor. Seborrheic keratoses usually begin with the appearance of one or more sharply defined, light brown, flat macules. The lesions may be sparse or numerous. As they initially

grow, they develop a velvety to finely verrucous surface, followed by an uneven warty surface with multiple plugged follicles and a dull or lackluster appearance.

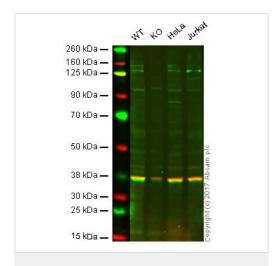
Sequence similarities

Belongs to the PI3/PI4-kinase family.

Contains 1 C2 domain.

Contains 1 PI3K/PI4K domain.

Images



Western blot - Anti-PI 3 Kinase catalytic subunit alpha/PIK3CA antibody [EPR19693] - BSA and Azide free (ab223532)

This WB data was generated using the same anti-PI3 Kinase catalytic subunit alpha/PIK3CA antibody clone [EPR19693] in a different buffer formulation (cat# <u>ab183957</u>).

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

Lane 2: PI3 Kinase catalytic subunit alpha/PIK3CA knockout

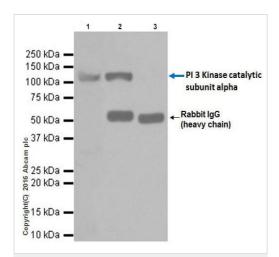
HAP1 whole cell lysate (20 µg)

Lane 3: HeLa whole cell lysate (20 µg)

Lane 4: Jurkat whole cell lysate (20 µg)

Lanes 1 - 4: Merged signal (red and green). Green - <u>ab183957</u> observed at 125 kDa. Red - loading control, <u>ab9484</u>, observed at 37 kDa.

ab183957 was shown to recognize PI 3 Kinase catalytic subunit alpha/PIK3CA in wild type cells as signal was lost at the expected MW in PI 3 Kinase catalytic subunit alpha/PIK3CA knockout cells. Additional cross-reactive bands were observed in the wild-type and knockout cells. Wild-type and PI 3 Kinase catalytic subunit alpha/PIK3CA knockout samples were subjected to SDS-PAGE. ab183957 and ab9484 (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.



Immunoprecipitation - Anti-PI 3 Kinase catalytic subunit alpha/PIK3CA antibody [EPR19693] - BSA and Azide free (ab223532)

This IP data was generated using the same anti-PI3 Kinase catalytic sunbunit alpha antibody clone [EPR19693] in a different buffer formulation (cat# <u>ab183957</u>).

PI 3 Kinase catalytic subunit alpha was immunoprecipitated from 0.35 mg of mouse hypothalamus lysate with <u>ab183957</u> at 1/30 dilution.

Western blot was performed from the immunoprecipitate using **ab183957** at 1/500 dilution.

VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>), was used for detection at 1/1000 dilution.

Lane 1: Mouse hypothalamus lysate, 10µg (Input).

Lane 2: ab183957 IP in mouse hypothalamus lysate.

Lane 3: Rabbit monoclonal $\lg G$ ($\underline{ab172730}$) instead of $\underline{ab183957}$ in mouse hypothalamus lysate.

Blocking and dilution buffer and concentration: 5% NFDM/TBST.

Exposure time: 3 minutes.



Anti-Pl 3 Kinase catalytic subunit alpha/PlK3CA antibody [EPR19693] - BSA and Azide free (ab223532)

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