

## Product datasheet

# Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] ab40776

**KO VALIDATED** Recombinant RabMAB<sup>®</sup>

[1 Abreviews](#) [7 References](#) [9 Images](#)

### Overview

<b>Product name</b>	Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y]
<b>Description</b>	Rabbit monoclonal [EP383Y] to PI 3 Kinase catalytic subunit alpha
<b>Tested applications</b>	<b>Suitable for:</b> Flow Cyt, WB, ICC/IF, IP <b>Unsuitable for:</b> IHC-P
<b>Species reactivity</b>	<b>Reacts with:</b> Mouse, Rat, Human
<b>Immunogen</b>	Synthetic peptide (the amino acid sequence is considered to be commercially sensitive) within Human PI 3 Kinase catalytic subunit alpha aa 1000 to the C-terminus (C terminal). The exact sequence is proprietary.
<b>Positive control</b>	WB: Jurkat, MCF-7, Raw264.7 and NIH/3T3 cell lysates. ICC/IF: HeLa and Jurkat cells. IP: Jurkat cell lysate.
<b>General notes</b>	<p>This product is a recombinant rabbit monoclonal antibody.</p> <p>A trial size is available to purchase for this antibody.</p> <p>Rat: We have preliminary internal testing data to indicate this antibody may not react with these species. Please contact us for more information.</p> <p>Alternative versions available:</p> <p><a href="#">Anti-PI 3 Kinase catalytic subunit alpha antibody (Alexa Fluor<sup>®</sup> 488) [EP383Y] (ab202671)</a></p> <p>Our RabMAB<sup>®</sup> technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to <a href="#">RabMab<sup>®</sup> patents</a></p> <p><b>We are constantly working hard to ensure we provide our customers with best in class antibodies. As a result of this work we are pleased to now offer this antibody in purified format. We are in the process of updating our datasheets. The purified format is designated 'PUR' on our product labels. If you have any questions regarding this update, please contact our Scientific Support team.</b></p>

### Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C.

	Avoid freeze / thaw cycle.
<b>Storage buffer</b>	pH: 7.20 Preservative: 0.01% Sodium azide Constituents: 40% Glycerol, 0.05% BSA, 59% PBS
<b>Purity</b>	Protein A purified
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	EP383Y
<b>Isotype</b>	IgG

## Applications

Our [Abpromise guarantee](#) covers the use of **ab40776** 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
Flow Cyt		Use at an assay dependent concentration.
WB		1/1000 - 1/10000. Detects a band of approximately 110 kDa (predicted molecular weight: 110 kDa).
ICC/IF		1/100 - 1/250.
IP		1/20 - 1/30.

**Application notes** Is unsuitable for IHC-P.

## Target

**Function** Phosphorylates PtdIns, PtdIns4P and PtdIns(4,5)P2 with a preference for PtdIns(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 late-stage 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.

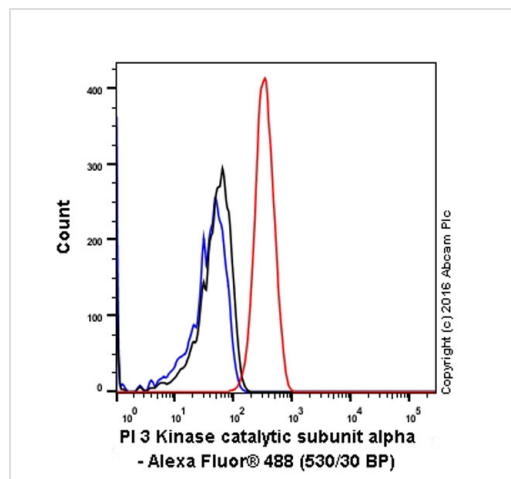
## Sequence similarities

Belongs to the PI3/PI4-kinase family.

Contains 1 C2 domain.

Contains 1 PI3K/PI4K domain.

## Images

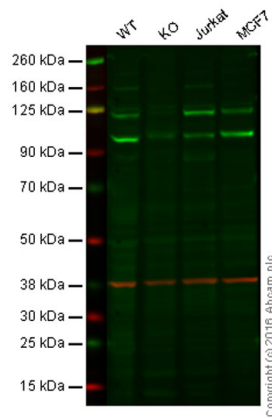


Flow Cytometry - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

ab40776 staining PI 3 Kinase catalytic subunit alpha in the human cell line Jurkat (human acute T cell leukemia) by flow cytometry. Cells were fixed with 4% paraformaldehyde, permeabilised with 90% methanol and the sample was incubated with the primary antibody at a dilution of 1/40. A goat anti rabbit IgG (Alexa Fluor® 488) at a dilution of 1/2000 was used as the secondary antibody.

Isoytype control: Rabbit monoclonal IgG (Black)

Unlabelled control: Cell without incubation with primary antibody and secondary antibody (Blue)



Western blot - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

**Predicted band size :** 110 kDa

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

**Lane 2:** PI 3 Kinase catalytic subunit alpha knockout HAP1 cell lysate (20 µg)

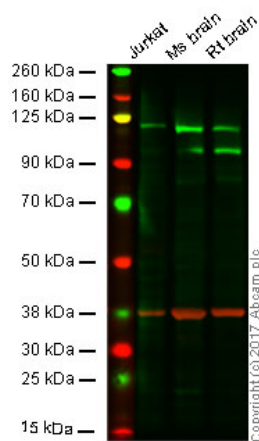
**Lane 3:** Jurkat cell lysate (20 µg)

**Lane 4:** MCF7 cell lysate (20 µg)

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

Green - ab40776 observed at 120 kDa. Red - loading control, ab8245, observed at 37 kDa.

ab40776 was shown to recognize PI 3 Kinase catalytic subunit alpha when PI 3 Kinase catalytic subunit alpha knockout samples were used, along with additional cross-reactive bands. Wild-type and PI 3 Kinase catalytic subunit alpha knockout samples were subjected to SDS-PAGE. ab40776 and ab8245 (loading control to GAPDH) were diluted to 1/1000 and 1/10000 respectively and 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-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

**All lanes** : Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776) at 1/1000 dilution

**Lane 1** : Jurkat Whole Cell Lysate

**Lane 2** : Mouse Brain Tissue Lysate

**Lane 3** : Rat Brain Tissue Lysate

Lysates/proteins at 20 µg per lane.

### Secondary

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

Performed under reducing conditions.

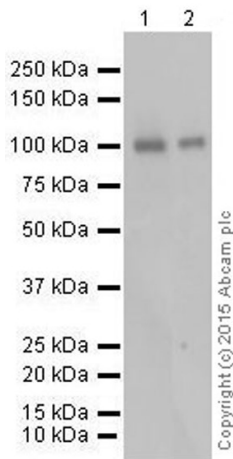
**Predicted band size** : 110 kDa

**Observed band size** : 120 kDa

Lanes 1 - 3: Merged signal (red and green).

Green - ab40776 observed at 120 kDa. Red - loading control, ab8245, observed at 37 kDa.

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using Licor blocking buffer before being incubated with ab40776 and ab8245 (loading control) overnight at 4°C. Antibody binding was detected using Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) at a 1:10000 dilution for 1hr at room temperature and then imaged.



Western blot - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

**All lanes :** Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776) at 1/5000 dilution (purified)

**Lane 1 :** Jurkat whole cell lysate

**Lane 2 :** MCF-7 whole cell lysate

Lysates/proteins at 20 µg per lane.

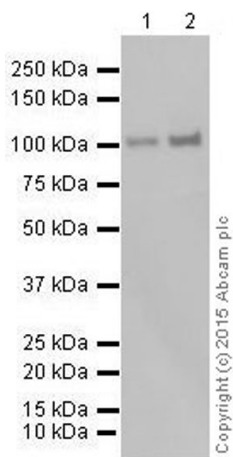
**Secondary**

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

**Predicted band size :** 110 kDa

**Observed band size :** 110 kDa

Blocking and dilution buffer: 5% NFDm/TBST



Western blot - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

**All lanes :** Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776) at 1/5000 dilution (purified)

**Lane 1 :** Raw264.7 whole cell lysate

**Lane 2 :** NIH/3T3 whole cell lysate

Lysates/proteins at 20 µg per lane.

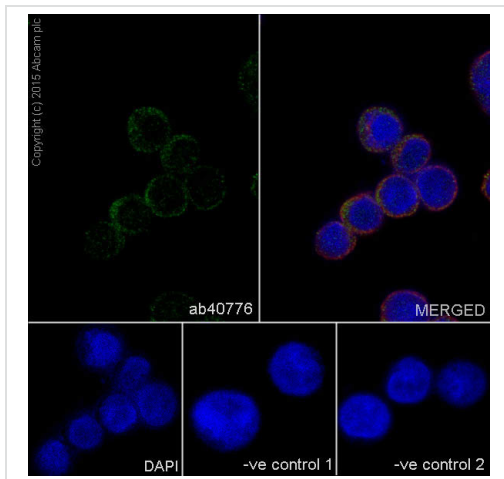
**Secondary**

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

**Predicted band size :** 110 kDa

**Observed band size :** 110 kDa

Blocking and dilution buffer: 5% NFDm/TBST

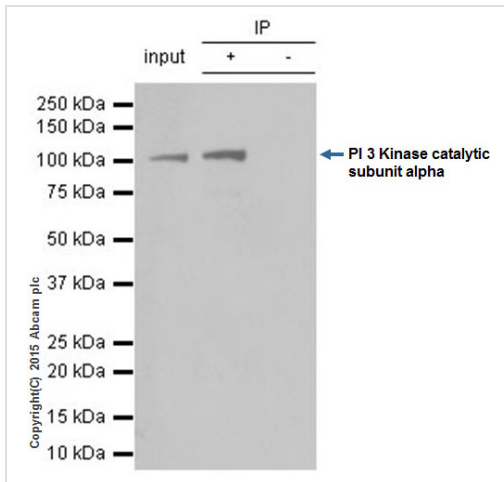


Immunocytochemistry/ Immunofluorescence - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

Immunocytochemistry/Immunofluorescence analysis of Jurkat cells labelling PI 3 Kinase catalytic subunit alpha with purified ab40776 at a dilution of 1/100. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. ab150077, an Alexa Fluor® 488-conjugated goat anti-rabbit IgG (1/1000) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain. ab7291, a mouse anti-tubulin (1/1000) and ab150120, an Alexa Fluor® 594-conjugated goat anti-mouse IgG (1/1000) were also used.

Control 1: primary antibody (1/100) and secondary antibody, ab150120, an Alexa Fluor® 594-conjugated goat anti-mouse IgG (1/1000).

Control 2: ab7291 (1/1000) and secondary antibody, ab150077, an Alexa Fluor® 488-conjugated goat anti-rabbit IgG (1/1000).



Immunoprecipitation - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

ab40776 (purified) at a dilution of 1/20 immunoprecipitating PI 3 Kinase catalytic subunit alpha in Jurkat whole cell lysate.

Lane 1 (input): Jurkat whole cell lysate (10µg)

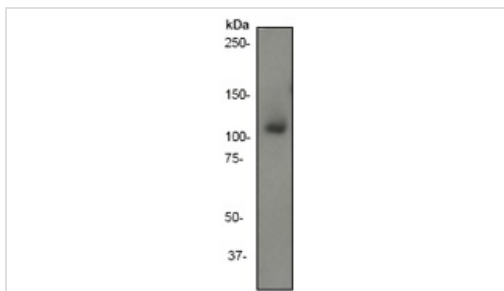
Lane 2 (+): ab40776 + Jurkat whole cell lysate.

Lane 3 (-): Rabbit monoclonal IgG ([ab172730](#)) instead of ab40776 in Jurkat whole cell lysate.

For western blotting, [ab131366](#) VeriBlot for IP (HRP) was used as the secondary antibody (1/1000).

Blocking buffer and concentration: 5% NFDm/TBST.

Diluting buffer and concentration: 5% NFDm /TBST.

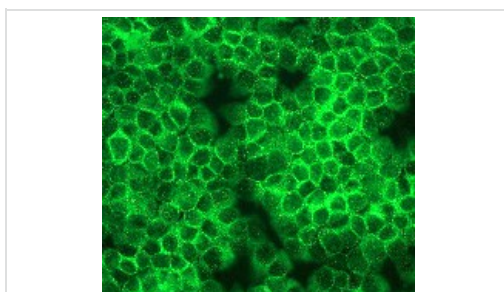


Western blot - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776) at 1/5000 dilution (unpurified) + Jurkat cell lysate at 10 µg

**Predicted band size** : 110 kDa

**Observed band size** : 110 kDa



Immunocytochemistry/ Immunofluorescence - Anti-PI 3 Kinase catalytic subunit alpha antibody [EP383Y] (ab40776)

Immunocytochemistry/Immunofluorescence analysis of HeLa cells labelling PI 3 Kinase catalytic subunit alpha with unpurified ab40776 at a dilution of 1/100.



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