abcam

ab309963

Anti-PHIP antibody [EPR27150-82] - BSA and Azide free

Recombinant | RabMAb | KO Validated

Key facts

Isotype	IgG
Host species	Rabbit
Storage buffer	pH: 7.2 - 7.4 Constituents: PBS
Form	Liquid
Clonality	Monoclonal
Immunogen	The exact immunogen used to generate this antibody is proprietary information.
Carrier free	Yes
Clone number	EPR27150-82
Purification technique	Affinity purification Protein A
Concentration	0.983 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

ChIP-seq

Not recommended

Species	Human, Mouse, Rat
Dilution info	-

Notes	-

WB

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

IHC-P

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

ICC/IF

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

Flow Cyt (Intra)

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

IP

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

Target data

See full target information PHIP

Function	Probable regulator of the insulin and insulin-like growth factor signaling pathways. Stimulates cell proliferation through regulation of cyclin transcription and has an anti-apoptotic activity through AKT1 phosphorylation and activation. Plays a role in the regulation of cell morphology and cytoskeletal organization.

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C

Notes

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

What are the advantages of a recombinant monoclonal antibody?
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 batch production

For more information, read more on recombinant antibodies.

Conjugation ready
Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. 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 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.

Compatibility
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.

Supplementary info

This supplementary information is collated from multiple sources and compiled automatically.

Activity summary	PHIP also known as pleckstrin homology domain interacting protein is an adapter protein that interacts with insulin receptor substrates. It has a mass of approximately 100 kDa. PHIP is expressed in various tissues including the brain and liver. The protein is involved in intracellular signaling pathways by binding phosphoinositides which are important for cellular communication and growth regulation.
Biological function summary	PHIP plays a significant role in cell proliferation and survival. It contributes to the regulation of the cell cycle by influencing transcription factors and other core components involved in cell growth. PHIP is not only a standalone protein but also part of complexes that modulate chromatin affecting gene expression patterns.
Pathways	PHIP is integrated into insulin signaling and PI3K/AKT pathways. In these pathways PHIP interacts with key proteins like AKT1 supporting cellular responses to insulin and growth factors. The involvement of PHIP in these pathways highlights its role in metabolic regulation and its impact on cell growth and metabolism.
Associated diseases and disorders	PHIP has links to cancer and diabetes. Its dysregulation contributes to abnormal cell proliferation often observed in cancers. PHIP's interaction with insulin signaling components such as IRS1 demonstrates its relevance to diabetes management. These connections show PHIP's potential as a therapeutic target for these conditions.

Product promise

Tested

We have tested this species and application combination and it works. It is covered by our product promise.

Expected

We have not tested this specific species and application combination in-house, but expect it will work. It is covered by our product promise.

Predicted

This species and application combination has not been tested, but we predict it will work based on strong homology. However, this combination is not covered by our product promise.

Not recommended

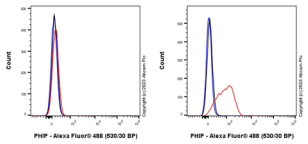
We do not recommend this combination. It is not covered by our product promise.

We are dedicated to supporting your work with high quality reagents and we are here for you every step of the way should you need us.

In the unlikely event of one of our products not working as expected, you are covered by our product promise.

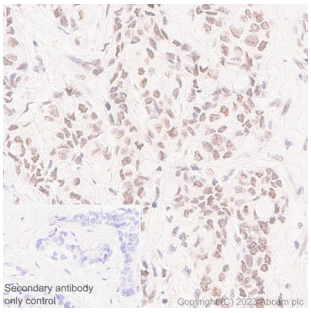
Full details and terms and conditions can be found here:
Terms & Conditions.

11 product images



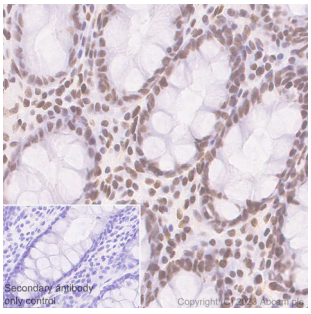
Flow Cytometry (Intracellular) - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Flow cytometric analysis of PHIP KO HeLa (PHIP knockout human cervical adenocarcinoma epithelial cell, Left) / Parental HeLa (Right) cells labelling PHIP with ab309962 at 1/500 dilution (0.1 ug)/Red compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/5000 dilution was used as the secondary antibody.



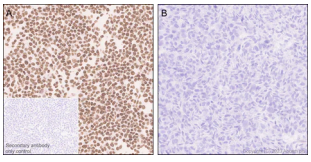
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human breast carcino tissue labeling PHIP with ab309962 at 1/100 (4.92 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on human breast carcinoma.The section was incubated with ab309962 for 30 mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

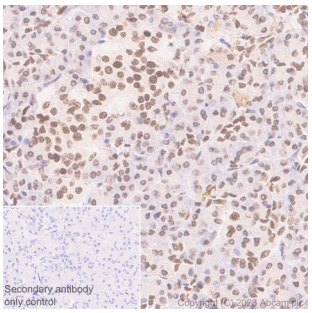
This data was developed using ab309962, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human colon tissue labeling PHIP with ab309962 at 1/100 (4.92 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on human colon.The section was incubated with ab309962 for 30 mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

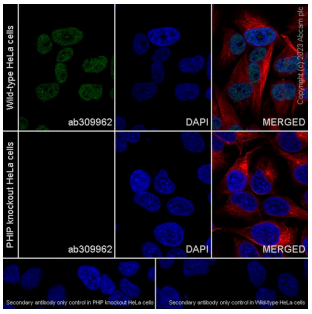
This data was developed using ab309962, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded (A) Wild-type Hela (tissue labeling PHIP with ab309962 at 1/2000 (0.246 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on (A) Wild-type Hela cell pellet, no staining on (B) PHIP knockout Hela (ab265403) cell pellet.The section was incubated with ab309962 for 30

mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



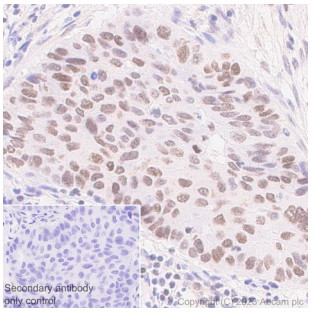
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human pancreas tissue labeling PHIP with ab309962 at 1/100 (4.92 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on human pancreas.The section was incubated with ab309962 for 30 mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



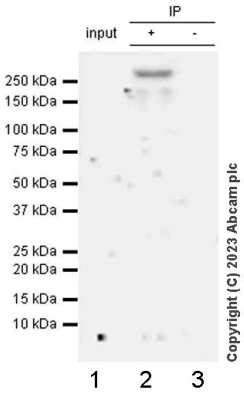
Immunocytochemistry/ Immunofluorescence - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized PHIP KO HeLa (human cervical adenocarcinoma epithelial cell) (ab265403) cells labelling PHIP with ab309962 at 1/100 (4.92 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 2 ug/mL dilution (Green). Confocal image showing nuclear staining in parental HeLa cells.Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8). is observed. ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 2.5ug/ml dilution (Red). The Nuclear counterstain was DAPI (Blue). Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 2 ug/mL dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded Human lung adenocarc tissue labeling PHIP with ab309962 at 1/100 (4.92 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on human lung adenocarcinoma.The section was incubated with ab309962 for 30 mins at room temperature.The immunostaining was performed on a Leica Biosystems BOND® RX instrument Counterstained with Hematoxylin. Secondary antibody only control: Secondary antibody is a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Heat mediated antigen retrieval was performed with Tris-EDTA buffer (pH 9.0, Epitope Retrieval Solution2) for 20 mins



Immunoprecipitation - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

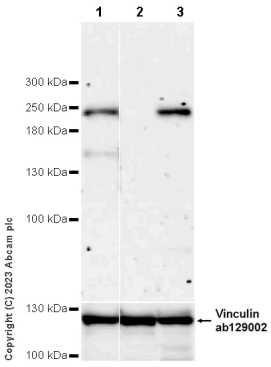
This data was developed using ab309962, the same antibody clone in a different buffer formulation. PHIP was immunoprecipitated from 0.35 mg HeLa (human cervical adenocarcinoma epithelial cell) whole cell lysate with ab309962 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab309962 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP)(ab131366) was used at 1/5000 dilution. Lane 1: HeLa (human cervical adenocarcinoma epithelial cell) whole cell lysate Lane 2: ab309962 IP in HeLa (human cervical adenocarcinoma epithelial cell) whole cell lysate Lane 3:Rabbit monoclonal IgG (ab172730) instead of ab309962 in HeLa whole cell lysate Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 3 minutes
All lanes:
Immunoprecipitation - Anti-PHIP antibody [EPR27150-82] (ab309962) at 1/30 dilution

All lanes:
HeLa (human cervical adenocarcinoma epithelial cell) whole cell lysate

Secondary

All lanes:
Immunoprecipitation - VeriBlot for IP Detection Reagent (HRP) (ab131366) at 1/5000 dilution

Exposure time: 3min



Western blot - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

PHIP Western blot staining using rabbit Anti-PHIP antibody

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST Performed under reducing conditions. In Western blot, ab309962 was shown to bind specifically to PHIP. A band was observed at 250 kDa in wild-type Hela cell lysates whereas no signal observed at this size in PHIP knockout cell line ab265403 (knockout cell lysate ab259039). The band beneath the target band (250 kDa) is likely to be degraded target fragment. In Western blot, anti- Vinculin antibody (ab129002) loading control staining at 1/10000 dilution. Exposure time: 3 minutes
All lanes:
Western blot - Anti-PHIP antibody [EPR27150-82] (ab309962) at 1/1000 dilution

Lane 1:
Wild-type HeLa(human cervical adenocarcinoma epithelial cell) whole cell lysate at 40 µg

Lane 2:
Western blot - Human PHIP knockout HeLa cell lysate (ab259039) at 40 µg

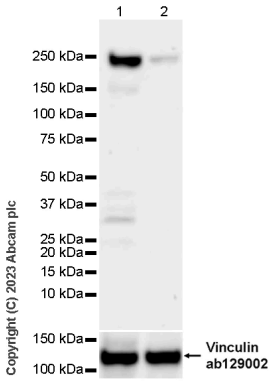
Lane 3:
K-562 (human chronic myelogenous leukemia lymphoblast) whole cell lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Observed band size: 250 kDa

Exposure time: 3min



Western blot - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

PHIP Western blot staining using rabbit Anti-PHIP antibody

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST The band beneath the target band (250 kDa) is likely to be degraded target fragment. In Western blot, anti-Vinculin antibody (ab129002) loading control staining at 1/10000 dilution. Exposure time: 48 seconds
All lanes:
Western blot - Anti-PHIP antibody [EPR27150-82] (ab309962) at 1/1000 dilution

Lane 1:
HeLa (human cervical adenocarcinoma epithelial cell) transfected with scrambled siRNA control whole cell lysate at 20 µg

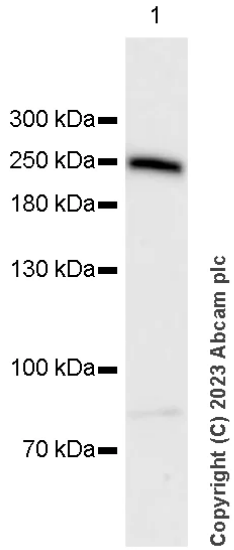
Lane 2:
HeLa transfected with siRNA specifically targeti PHIP whole cell lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Observed band size: 250 kDa

Exposure time: 48s



Western blot - Anti-PHIP antibody [EPR27150-82] - BSA and Azide free (ab309963)

This data was developed using ab309962, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST Lysates were freshly made and used for Western blotting immediately to minimize protein degradation. Exposure time: 59 seconds
All lanes:
Western blot - Anti-PHIP antibody [EPR27150-82] (ab309962) at 1/1000 dilution

All lanes:
293T (human embryonic kidney epithelial cell) whole cell lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Observed band size: 250 kDa

Exposure time: 59s

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.