abcam

ab314437

Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free

Recombinant | RabMAb | Advanced Validation

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	EPR27147-176
Purification technique	Affinity purification Protein A
Specificity	Unsuitable for FC-intra human.
Concentration	1.037 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

ChIP-seq

Tested

Species	Human

Dilution info	-
Notes	-

Expected

Species	Mouse, Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Recombinant fragment - Human
Dilution info	-
Notes	-

WB

Tested

Species	Human, Mouse, Rat, Recombinant fragment - Human
Dilution info	-
Notes	-

ICC/IF

Tested

Species	Human, Mouse
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Recombinant fragment - Human
Dilution info	-
Notes	-

IP

Tested

Species	Human, Mouse
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Recombinant fragment - Human
Dilution info	-
Notes	-

Flow Cyt (Intra)

Tested

Species	Mouse
Dilution info	-
Notes	-

Expected

Species	Rat
Dilution info	Use at an assay dependent concentration.
Notes	-

Not recommended

Species	Human, Recombinant fragment - Human
Dilution info	-
Notes	-

IHC-P

Tested

Species	Human
Dilution info	-
Notes	Perform heat-mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Species	Mouse
Dilution info	-
Notes	Perform heat-mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Species	Rat
Dilution info	-
Notes	Perform heat-mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Not recommended

Species	Recombinant fragment - Human
Dilution info	-
Notes	-

Target data

See full target information PHC2

Function	Component of a Polycomb group (PcG) multiprotein PRC1-like complex, a complex class required to maintain the transcriptionally repressive state of many genes, including Hox genes, throughout development. PcG PRC1 complex acts via chromatin remodeling and modification of histones; it mediates monoubiquitination of histone H2A 'Lys-119', rendering chromatin heritably changed in its expressibility.

Storage

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

Notes

ab314437 is the carrier-free version of ab314436.

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	PHC2 also known as Polyhomeotic Homolog 2 is a protein with a molecular mass around 100 kDa. It localizes mainly in the nucleus and expresses widely across different tissue types. The protein is a part of the Polycomb group (PcG) proteins which play key roles in chromatin remodeling and gene silencing. PHC2 contributes to forming stable gene repression essential for regulating development and maintaining cellular identity.
Biological function summary	PHC2 functions as a part of the Polycomb repressive complex 1 (PRC1). This complex is involved in modifying the structure of chromatin which helps maintain the repression of gene transcription. Through interaction with other PRC1 components PHC2 helps establish a compact chromatin state promoting histone modifications that lead to transcriptional silencing. This activity is fundamental for cellular processes like embryonic development and stem cell maintenance.
Pathways	PHC2 plays a role in the regulation of the Wnt signaling and Notch signaling pathways. These pathways are integral to controlling cell proliferation differentiation and apoptosis. Within these pathways PHC2 interacts with other proteins like BMI1 and RING1B which are also part of the PRC1 complex. These interactions ensure the precise control over gene expression needed for cellular homeostasis and proper developmental progression.
Associated diseases and disorders	Aberrant PHC2 function is implicated in a range of conditions particularly cancer and neurodevelopmental disorders. Dysregulation of PHC2 can lead to misexpression of genes that regulate cell cycle and apoptosis contributing to oncogenesis. For instance PHC2 has connections with BMI1 in cancer pathways influencing tumor progression and resistance to apoptosis. Moreover mutations or irregular expression of PHC2 have been linked with neurodevelopmental disorders highlighting its importance in nervous system development and function.

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

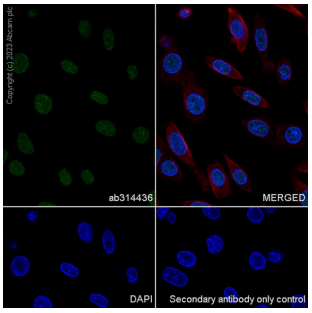
We do not recommend this combination. It is not covered by our product promise.

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.

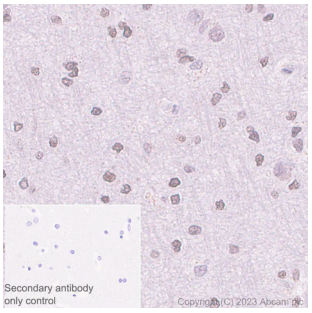
Full details and terms and conditions can be found here:
Terms & Conditions.

18 product images



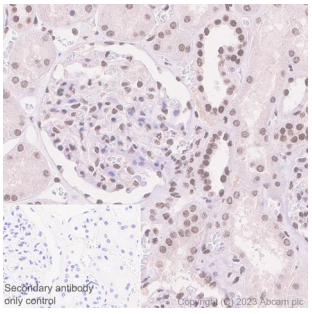
Immunocytochemistry/ Immunofluorescence - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation.Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized SW480 (human colorectal adenocarcinoma epithelial cell) cells labelling PHC2 with ab314436 at 1/50 (10.38 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 SW480 cell line.Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8).ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 (2.5 ug/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-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

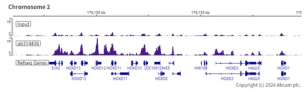
This data was developed using ab314436, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded human cerebrum tissue labeling PHC2 with ab314436 at 1/100 (5.19 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on human cerebrum. The section was incubated with ab314436 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-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

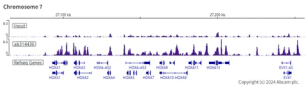
This data was developed using ab314436, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling PHC2 with ab314436 at 1/100 (5.19 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on human kidney.The section was incubated with ab314436 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.



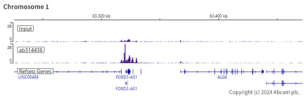
ChIP-sequencing - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

Chromatin was prepared from NCCIT cells. Cells were fixed with 1% formaldehyde for 10 minutes. ChIP was performed with 10^7 cells and 8 µg of ab314436 [EPR27147-176]. ChIP DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 30 million reads. The Input control is also shown.



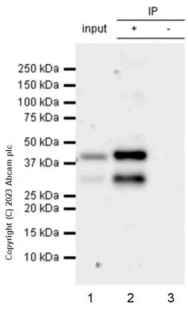
ChIP-sequencing - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

Chromatin was prepared from NCCIT cells. Cells were fixed with 1% formaldehyde for 10 minutes. ChIP was performed with 10^7 cells and 8 µg of ab314436 [EPR27147-176]. ChIP DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 30 million reads. The Input control is also shown. This data was developed using ab314436, the same antibody clone in a different buffer formulation.



ChIP-sequencing - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

Chromatin was prepared from NCCIT cells. Cells were fixed with 1% formaldehyde for 10 minutes. ChIP was performed with 10^7 cells and 8 µg of ab314436 [EPR27147-176]. ChIP DNA was sequenced on the Illumina NovaSeq 6000 to a depth of 30 million reads. The Input control is also shown. This data was developed using ab314436, the same antibody clone in a different buffer formulation.



Immunoprecipitation - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

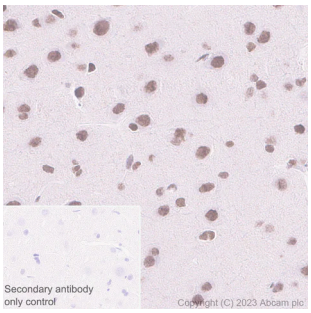
This data was developed using ab314436, the same antibody clone in a different buffer formulation. PHC2 was immunoprecipitated from 0.35 mg SW480 (human colorectal adenocarcinoma epithelial cell) whole cell lysate with ab314436 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab314436 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP)(ab131366) was used at 1/5000 dilution. Lane 1: SW480 (human colorectal adenocarcinoma epithelial cell) whole cell lysate Lane 2: ab314436 IP in SW480 (human colorectal adenocarcinoma epithelial cell) whole cell lysate Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab314436 in SW480 whole cell lysate Blocking and dilution buffer and concentration: 5% NFDM/TBST.
All lanes:
Immunoprecipitation - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade (ab314436) at 1/30 dilution

All lanes:
SW480 (human colorectal adenocarcinoma epithelial cell) whole cell lysate

Secondary

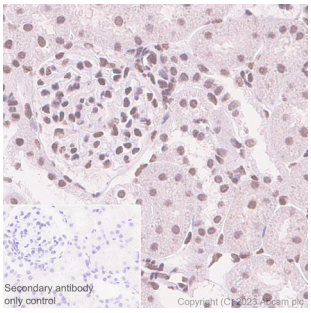
All lanes:
Immunoprecipitation - VeriBlot for IP Detection Reagent (HRP) (ab131366) at 1/5000 dilution

Exposure time: 58s



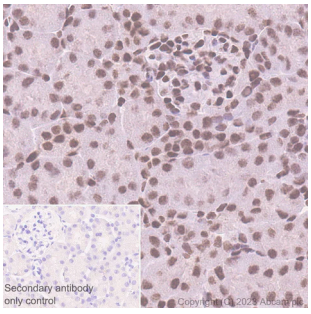
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded rat cerebrum tissue labeling PHC2 with ab314436 at 1/100 (5.19 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on rat cerebrum. The section was incubated with ab314436 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.



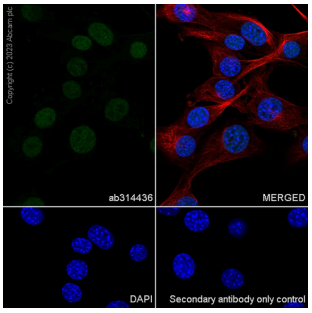
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded rat kidney tissue labeling PHC2 with ab314436 at 1/100 (5.19 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on rat kidney. The section was incubated with ab314436 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.



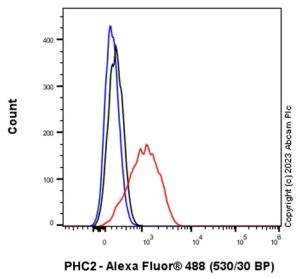
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded mouse kidney tissue labeling PHC2 with ab314436 at 1/100 (5.19 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on mouse kidney. The section was incubated with ab314436 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.



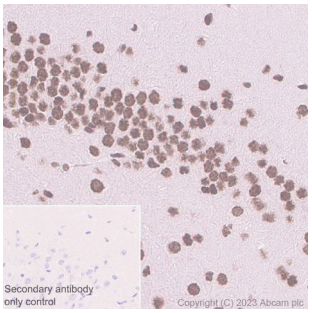
Immunocytochemistry/ Immunofluorescence - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation.Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized NIH/3T3 (mouse embryonic fibroblast) cells labelling PHC2 with ab314436 at 1/50 (10.38 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 NIH/3T3 cell line.Image was taken with a confocal microscope(Leica-Microsystems, TCS SP8). ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 (2.5 ug/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.



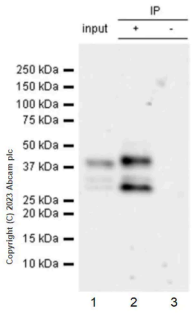
Flow Cytometry (Intracellular) - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation.Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized NIH/3T3 (mouse embryonic fibroblast) cells labelling PHC2 with ab314436 at 1/50 dilution (1 ug)/Red (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). A 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-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Immunohistochemical analysis of paraffin-embedded mouse cerebrum tissue labeling PHC2 with ab314436 at 1/100 (5.19 ug/ml) followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). Nuclear staining on mouse cerebrum. The section was incubated with ab314436 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-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. PHC2 was immunoprecipitated from 0.35 mg NIH/3T3 (mouse embryonic fibroblast) whole cell lysate with ab314436 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab314436 at 1/1000 dilution. VeriBlot for IP secondary antibody(HRP)(ab131366) was used at 1/5000 dilution. Lane 1: NIH/3T3 (mouse embryonic fibroblast) whole cell lysate Lane 2: ab314436 IP in NIH/3T3 (mouse embryonic fibroblast) whole cell lysate Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab314436 NIH/3T3 whole cell lysate Blocking and dilution buffer and concentration: 5% NFDM/TBST.
All lanes:

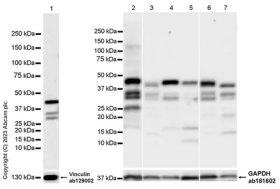
Immunoprecipitation - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade (ab314436) at 1/30 dilution

All lanes:
NIH/3T3 (mouse embryonic fibroblast) whole cell lysate

Secondary

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

Exposure time: 76s



Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

PHC2 Western blot staining using rabbit Anti-PHC2 antibody

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST The expression profile/molecular weight observed is consistent with what has been described in the literature (PMID: 16024804). The identity of the bands below 31 kDa is unknown. In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution. In Western blot, Anti-Vinculin antibody [EPR8185] (ab129002) staining at 1/10000 dilution. Exposure time: Lane1: 180 seconds, lane2-7: 37 seconds
All lanes:
Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade (ab314436) at 1/1000 dilution

Lane 1:
NCCIT (human pluripotent embryonic carcinoma epithelial cell) whole cell lysate at 20 µg

Lanes 2 - 3:
Human testis tissue lysate at 20 µg

Lane 4:
Mouse testis tissue lysate at 20 µg

Lane 5:
Mouse kidney tissue lysate at 20 µg

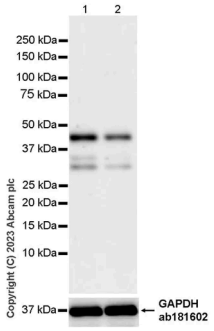
Lane 6:
Rat testis tissue lysate at 20 µg

Lane 7:
Rat kidney tissue lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 31 kDa, 36 kDa, 40 kDa



Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST The expression profile/molecular weight observed is consistent with what has been described in the literature (PMID: 16024804). In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution.
All lanes:
Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade (ab314436) at 1/1000 dilution

Lane 1:
SW480 (human colorectal adenocarcinoma epithelial cell) transfected with scrambled siRNA control whole cell lysate at 20 µg

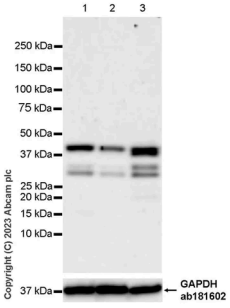
Lane 2:
SW480 transfected with siRNA specifically targeti PHC2 whole cell lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 31 kDa, 36 kDa, 40 kDa

Exposure time: 80s



Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST Low expression: Raji (PMID:9121482) The expression profile/ molecular weight observed is consistent with what has been described in the literature (PMID: 16024804). In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/200000 dilution.
All lanes:
Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade (ab314436) at 1/1000 dilution

Lane 1:
SW480 (human colorectal adenocarcinoma epithelial cell) whole cell lysate at 20 µg

Lane 2:
Raji (human burkitts lymphoma b lymphocyte) whole cell lysate at 20 µg

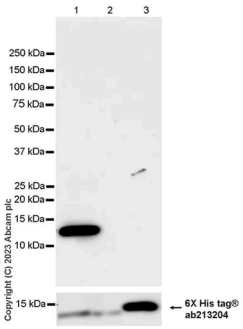
Lane 3:
NIH/3T3 (mouse embryonic fibroblast) whole cell lysate at 20 µg

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 31 kDa, 36 kDa, 40 kDa

Exposure time: 92s



Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade - BSA and Azide free (ab314437)

This data was developed using ab314436, the same antibody clone in a different buffer formulation. Blocking and diluting buffer and concentration: 5% NFDM/TBST This antibody does not react with human PHC1 and PHC3. In Western blot, Anti-6X His tag® antibody [EPR20547] - ChIP Grade (ab213204) staining at 1/5000 dilution
All lanes:
Western blot - Anti-PHC2 antibody [EPR27147-176] - ChIP Grade (ab314436) at 1/1000 dilution

Lane 1:
His-tagged human PHC2 recombinant protein at 10 ng

Lane 2:
His-tagged human PHC1 recombinant protein at 10 ng

Lane 3:
His-tagged human PHC3 recombinant protein at 10 ng

Secondary

All lanes:
Western blot - Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/100000 dilution

Observed band size: 14 kDa

Exposure time: 48s

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