abcam

ab251303

Anti-PELP1 antibody [EPR15213] - BSA and Azide free

Recombinant | RabMAb

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	EPR15213
Purification technique	Affinity purification Protein A

Reactivity data

ICC/IF

Tested

Species	Human
Dilution info	-
Notes	-

WB

Tested

Species	Human
Dilution info	-
Notes	-

Flow Cyt (Intra)

Tested

Species	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.

Target data

See full target information PELP1

Function

Coactivator of estrogen receptor-mediated transcription and a corepressor of other nuclear hormone receptors and sequence-specific transcription factors (PubMed:14963108). Plays a role in estrogen receptor (ER) genomic activity when present in the nuclear compartment by activating the ER target genes in a hormonal stimulation dependent manner. Can facilitate ER non-genomic signaling via SRC and PI3K interaction in the cytosol. Plays a role in E2-mediated cell cycle progression by interacting with RB1. May have important functional implications in ER/growth factor cross-talk. Interacts with several growth factor signaling components including EGFR and HRS. Functions as the key stabilizing component of the Five Friends of Methylated CHTOP (5FMC) complex; the 5FMC complex is recruited to ZNF148 by methylated CHTOP, leading to desumoylation of ZNF148 and subsequent transactivation of ZNF148 target genes. Component of the PELP1 complex involved in the nucleolar steps of 28S rRNA maturation and the subsequent nucleoplasmic transit of the pre-60S ribosomal subunit. Regulates pre-60S association of the critical remodeling factor MDN1 (PubMed:21326211). May

promote tumorigenesis via its interaction with and modulation of several oncogenes including SRC, PI3K, STAT3 and EGFR. Plays a role in cancer cell metastasis via its ability to modulate E2-mediated cytoskeleton changes and cell migration via its interaction with SRC and PI3K.

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C
Storage information	Do Not Freeze

Notes

ab251303 is the carrier-free version of ab200203.

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	The Proline Glutamic acid and Leucine-rich Protein 1 (PELP1) also known as Modulator of Nongenomic Activity of Estrogen Receptor (MNAR) plays a significant role in chromatin remodeling and transcriptional regulation. PELP1 has a molecular mass of approximately 160 kDa. It is broadly expressed in various tissues with particularly high levels in reproductive tissues and the brain. PELP1 participates in the recruitment of histone-modifying enzymes and acts as a scaffolding protein that interacts with several nuclear receptors and transcription factors.
Biological function summary	PELP1 acts within the nucleus and is central to gene expression regulation. It is an integral part of diverse protein complexes that engage in remodeling chromatin and modifying transcriptional activity. PELP1 interacts with estrogen receptors where it modulates both genomic and non-genomic signaling pathways affecting gene expression programs related to cell growth and differentiation. Additionally PELP1 associates with various other hormone receptors and transcription factors assisting in the intricate regulation of cell functions.
Pathways	Cellular growth and differentiation rely on the signaling roles that PELP1 performs. PELP1 is involved in the estrogen signaling pathway where it closely interacts with estrogen receptor alpha (ERα). This interaction links estrogen signaling to the cell cycle machinery and PI3K/AKT pathway influencing cell proliferation and survival. Another pathway in which PELP1 plays a critical role is the MAPK/ERK pathway where it cooperates with Src kinase highlighting its importance in cell signaling networks that regulate growth and differentiation.
Associated diseases and disorders	PELP1 holds a strong association with hormone-driven cancers particularly breast and prostate cancers. In these contexts PELP1's interaction with estrogen and androgen receptors can dramatically alter tumor progression suggesting its role as an oncogene. Additionally aberrant expression and activity of PELP1 have implications in neurological disorders. In breast cancer PELP1 collaborates with ERα to drive estrogen-mediated growth whereas in prostate cancer it engages with androgen receptor (AR) promoting cell proliferation and resistance to therapy.

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

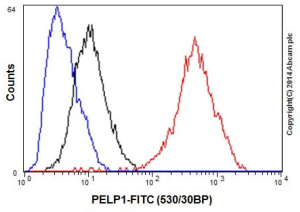
We do not recommend this combination. It is not covered by our product promise.

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.

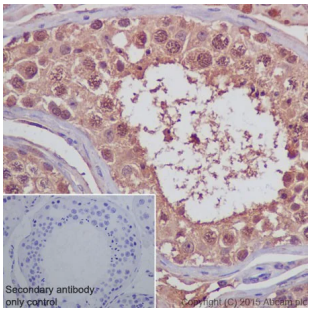
14 product images



Flow Cytometry (Intracellular) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Flow Cytometry (Intracellular) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Intracellular flow cytometric analysis of 2% paraformaldehyde-fixed HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling PELP1 with ab200203 at 1/300 dilution (red) compared with a rabbit monoclonal IgG isotype control (ab172730; black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody; blue). Goat anti rabbit IgG (FITC) at 1/500 dilution was used as the secondary antibody.



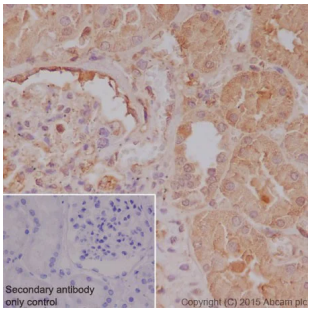
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded Human testis tissue labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and cytoplasm staining on Human testis tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



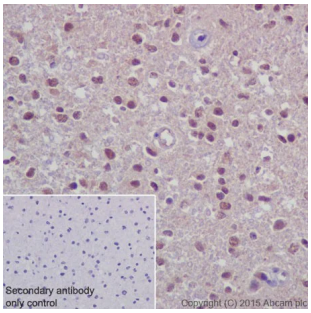
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded human kidney tissue labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and cytoplasm staining on Human kidney tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



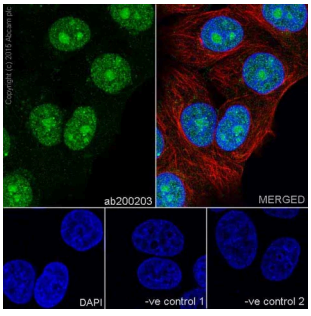
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded Human astrocytoma tissue labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and weakly cytoplasm staining on Human astrocytoma tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

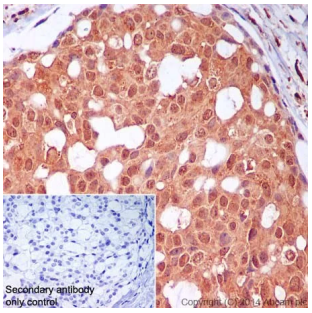


Immunocytochemistry/ Immunofluorescence - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunocytochemistry/ Immunofluorescence staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized MCF7 (Human breast adenocarcinoma cell line) cells labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Nuclear and weakly cytoplasmic staining on MCF7 cell line is observed. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

The negative controls are as follows:
-ve control 1: ab200203 at 1/250 dilution followed by ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution.
-ve control 2: ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution followed by ab150077 (Alexa Fluor®488 Goat Anti-Rabbit IgG H&L) at 1/500 dilution.



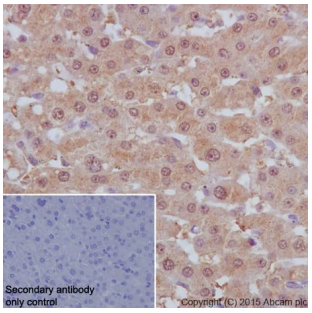
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded Human breast carcinoma tissue labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and cytoplasm staining on Human breast carcinoma tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



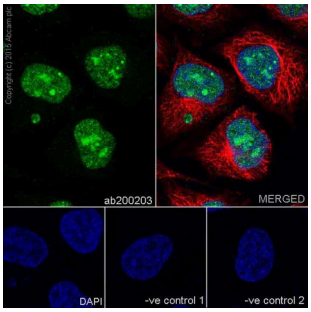
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded Human liver tissue labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and cytoplasm staining on Human liver tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

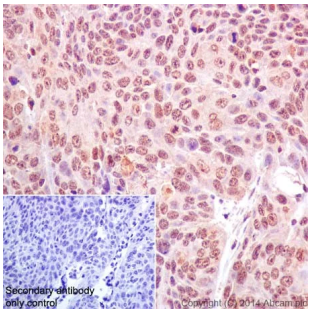


Immunocytochemistry/ Immunofluorescence - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunocytochemistry/ Immunofluorescence staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Nuclear staining on HeLa cell line is observed. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

The negative controls are as follows:
-ve control 1: ab200203 at 1/250 dilution followed by ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution.
-ve control 2: ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution followed by ab150077 (Alexa Fluor®488 Goat Anti-Rabbit IgG H&L) at 1/500 dilution.



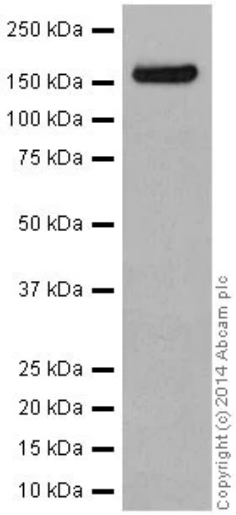
Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) staining using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Immunohistochemical analysis of paraffin-embedded Human cervix carcinoma tissue labeling PELP1 with ab200203 at 1/250 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Nuclear and weakly cytoplasm staining on Human cervix carcinoma tissue is observed. Counter stained with Hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Western blot - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Blocking and dilution buffer: 5% NFDM/TBST.

The expression profile observed is consistent with what has been described in the literature, PMID: 16141397.

All lanes:
Western blot - Anti-PELP1 antibody [EPR15213] (ab200203) at 1/1000 dilution

All lanes:
HEK-293 (Human epithelial cells from embryonic kidney) cell lysate at 20 µg

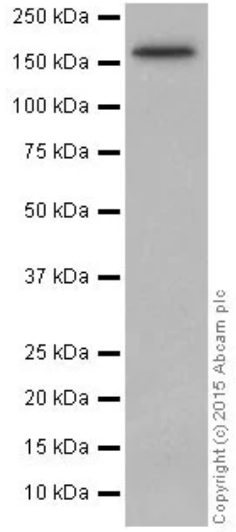
Secondary

All lanes:
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 120 kDa

Observed band size: 160 kDa

Exposure time: 15s



Western blot - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Western blot staining of Human breast cancer lysate using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Blocking and dilution buffer: 5% NFDM/TBST.

The expression profile observed is consistent with what has been described in the literature (PMID: 16141397).

All lanes:
Western blot - Anti-PELP1 antibody [EPR15213] (ab200203) at 1/20000 dilution

All lanes:
Human breast cancer lysate at 10 µg

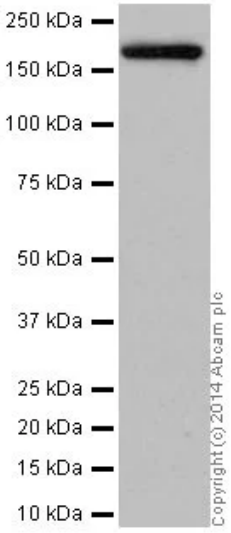
Secondary

All lanes:
Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 120 kDa

Observed band size: 160 kDa

Exposure time: 3min



Western blot - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Western blot staining of Human fetal brain lysate using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Blocking and dilution buffer: 5% NFDM/TBST.

The expression profile observed is consistent with what has been described in the literature, PMID: 16141397.

All lanes:
Western blot - Anti-PELP1 antibody [EPR15213] (ab200203) at 1/1000 dilution

All lanes:
Human fetal brain lysate at 20 µg

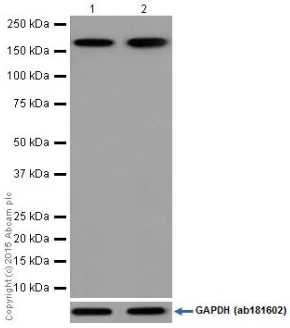
Secondary

All lanes:
Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 120 kDa

Observed band size: 160 kDa

Exposure time: 3min



Western blot - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Blocking and dilution buffer: 5% NFDM/TBST.

The expression profile observed is consistent with what has been described in the literature (PMID: 16141397).

All lanes:
Western blot - Anti-PELP1 antibody [EPR15213] (ab200203) at 1/20000 dilution

Lane 1:
T-47D (Human ductal breast epithelial tumor cell line) cell lysate at 10 µg

Lane 2:
MCF7 (Human breast adenocarcinoma cell line) cell lysate at 10 µg

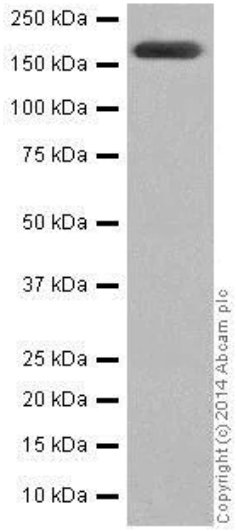
Secondary

All lanes:
Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 120 kDa

Observed band size: 160 kDa

Exposure time: 10s



Western blot - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303)

PELP1 Western blot staining of HeLa (Human epithelial cells from cervix adenocarcinoma) cell lysate using rabbit Anti-PELP1 antibody

This data was developed using ab200203, the same antibody clone in a different buffer formulation.
Blocking and dilution buffer: 5% NFDM/TBST.

The expression profile observed is consistent with what has been described in the literature (PMID: 16141397).

All lanes:
Western blot - Anti-PELP1 antibody [EPR15213] - BSA and Azide free (ab251303) at 1/10000 dilution

All lanes:
HeLa (Human epithelial cells from cervix adenocarcinoma) cell lysate

Secondary

All lanes:
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 120 kDa

Observed band size: 160 kDa

Exposure time: 30s

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