abcam

Product datasheet

Anti-HPRT antibody ab10479



★★★★ 4 Abreviews 48 References 6 Images

Overview

Product name Anti-HPRT antibody

Description Rabbit polyclonal to HPRT

Host species Rabbit

Specificity Replenishment batches of our polyclonal antibody, ab10479 are tested in WB. Previous batches

were additionally validated in ICC/IF. This application is still expected to work and is covered by our Abpromise guarantee. You may also be interested in our alternative recombinant antibody,

ab133242.

Tested applications Suitable for: WB, ICC/IF

Species reactivity Reacts with: Mouse, Rat, Human

Predicted to work with: Chicken, Gerbil, Chinese hamster

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: MEF1, NIH/3T3, HeLa, MCF-7, HEK-293, A-431, Wild-type HAP1 and PC12 whole cell

lysates, mouse brain tissue lysate and Recombinant Human HPRT protein (ab117153). ICC/IF:

HepG2 cells.

General notesThe Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

1

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising agent. If you would like information about the formulation of a specific lot, please contact our scientific support team who will be happy to help.

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee co

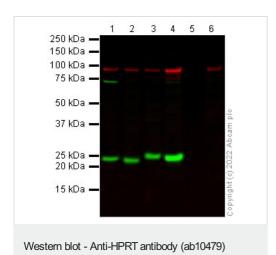
Our $\underline{\textbf{Abpromise guarantee}}$ covers the use of ab10479 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB	**** <u>(4)</u>	Use a concentration of 1 µg/ml. Predicted molecular weight: 24 kDa.
ICC/IF		Use a concentration of 5 µg/ml.

Target		
Function	Converts guanine to guanosine monophosphate, and hypoxanthine to inosine monophosphate. Transfers the 5-phosphoribosyl group from 5-phosphoribosylpyrophosphate onto the purine. Plays a central role in the generation of purine nucleotides through the purine salvage pathway.	
Pathway	Purine metabolism; IMP biosynthesis via salvage pathway; IMP from hypoxanthine: step 1/1.	
Involvement in disease	Defects in HPRT1 are the cause of Lesch-Nyhan syndrome (LNS) [MIM:300322]. LNS is characterized by complete lack of enzymatic activity that results in hyperuricemia, choreoathetosis, mental retardation, and compulsive self-mutilation. Defects in HPRT1 are the cause of gout HPRT-related (GOUT-HPRT) [MIM:300323]; also known as HPRT-related gout or Kelley-Seegmiller syndrome. Gout is characterized by partial enzyme activity and hyperuricemia.	
Sequence similarities	Belongs to the purine/pyrimidine phosphoribosyltransferase family.	
Cellular localization	Cytoplasm.	

Images



Lane 1: HeLa whole cell lysate (20 µg)

Lane 2: NIH/3T3 whole cell lysate (20 µg)

Lane 3: PC12 whole cell lysate (20 µg)

Lane 4: Wild-type HAP1 whole cell lysate (20 µg)

Lane 5: Empty Lane

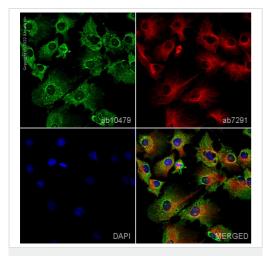
Lane 6: HPRT1 knockout HAP1 whole cell lysate (20 µg)

Lanes 1 - 6: Merged signal (red and green). Green - ab10479 observed at 24 kDa. Red - loading control <u>ab6301</u> observed at 95 kDa.

Gel type: MES

Blocking buffer: LiCOR blocking buffer

ab10479 was shown to specifically react with HPRT1 in wild-type HAP1 cells. No band was observed when HPRT1 knockout samples were examined. Wild-type and HPRT1 knockout samples were subjected to SDS-PAGE. Ab10479 and Ab6301 (Beta catenin - Loading control) were incubated overnight at 4°C at 1 µg/ml and 1/10,000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800RD) preadsorbed and Goat anti-Mouse lgG H&L (IRDye® 680CW) preadsorbed secondary antibodies at 1/10,000 dilution for 1 hour at room temperature before imaging.

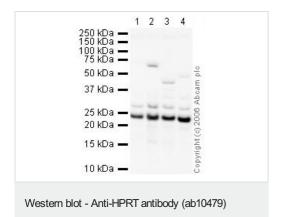


Immunocytochemistry/ Immunofluorescence - Anti-HPRT antibody (ab10479)

ab10479 staining HPRT in HepG2 cells. The cells were fixed with 100% methanol (5 min), permeabilized with 0.1% PBS-Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1%PBS-Tween for 1h. The cells were then incubated overnight at 4°C with ab10479 at 5µg/ml and ab7291, Mouse monoclonal [DM1A] to alpha Tubulin - Loading Control. Cells were then incubated with ab150081, Goat polyclonal Secondary Antibody to Rabbit IgG - H&L (Alexa Fluor® 488), pre-adsorbed at 1/1000 dilution (shown in green) and ab150120, Goat polyclonal Secondary Antibody to Mouse IgG - H&L (Alexa Fluor® 594), pre-adsorbed at 1/1000 dilution (shown in pseudocolour red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was acquired with a high-content analyser (Operetta CLS, Perkin Elmer) and a maximum intensity projection of confocal

sections is shown.



All lanes: Anti-HPRT antibody (ab10479) at 1 µg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2: A-431 whole cell lysate (ab7909)

Lane 3: MCF-7 (Human breast adenocarcinoma cell line) Whole Cell Lysate

Lane 4: HEK-293 whole cell lysate (ab7902)

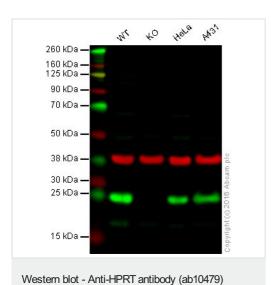
Lysates/proteins at 20 µg per lane.

Secondary

All lanes : Goat polyclonal to Rabbit lgG H&L (HRP) Pre-Adsorbed at 1/10000 dilution

Performed under reducing conditions.

Predicted band size: 24 kDa Observed band size: 24 kDa



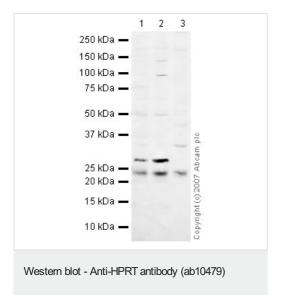
Lane 1: Wild-type HAP1 whole cell lysate (20 µg)

Lane 2: HPRT1 knockout HAP1 whole cell lysate (20 µg)

Lane 3: HeLa whole cell lysate (20 μg) **Lane 4:** A431 whole cell lysate (20 μg)

Lanes 1 - 4: Merged signal (red and green). Green - ab10479 observed at 25 kDa. Red - loading control, **ab8245**, observed at 37 kDa.

ab10479 was shown to specifically react with HPRT1 in wild-type HAP1 cells. No band was observed when HPRT1 knockout samples were examined. Wild-type and HPRT1 knockout samples were subjected to SDS-PAGE. Ab10479 and ab8245 (Mouse anti GAPDH loading control) were incubated overnight at 4°C at 1/500 dilution and 1/10,000 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 1/10,000 dilution for 1 hour at room temperature before imaging.



All lanes: Anti-HPRT antibody (ab10479) at 1 µg/ml

Lane 1: NIH/3T3 whole cell lysate (ab7179)

Lane 2: MEF1 (Mouse embryonic fibroblast cell line) Whole Cell

Lysate

Lane 3: Brain (Mouse) Tissue Lysate

Lysates/proteins at 10 µg per lane.

Secondary

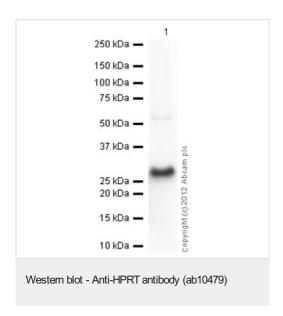
All lanes: IRDye 680 Conjugated Goat Anti-Rabbit IgG (H+L) at

1/10000 dilution

Performed under reducing conditions.

Predicted band size: 24 kDa Observed band size: 24 kDa

Additional bands at: 28 kDa (possible glycosylated form)



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