Product Datasheet

IDH2 Antibody - BSA Free NBP2-22166

Unit Size: 0.1 ml

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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NBP2-22166

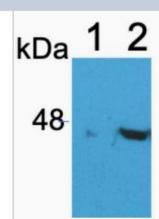
IDH2 Antibody - BSA Free

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Product Information	
Unit Size	0.1 ml
Concentration	1.0 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.02% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	PBS
Product Description	
Host	Rabbit
Gene ID	3418
Gene Symbol	IDH2
Species	Human, Mouse, Rat
Immunogen	A synthetic peptide made to a C-terminal portion of the human IDH2 protein (between residues 375-452) [UniProt P48735].
Product Application Details	
Applications	Western Blot, Immunocytochemistry/ Immunofluorescence,

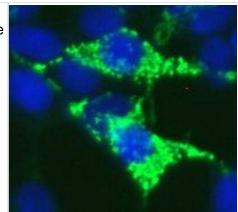
Product Application Details	
Applications	Western Blot, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry, Immunohistochemistry-Paraffin
Recommended Dilutions	Western Blot 1 - 2 ug/mL, Immunohistochemistry 1:100 - 1:500, Immunocytochemistry/ Immunofluorescence 1:250 - 1:500, Immunohistochemistry-Paraffin 1:100 - 1:500
Application Notes	In Western blot a band is observed ~48 kDa. Prior to immunostaining paraffin tissues, antigen retrieval with sodium citrate buffer (pH 6.0) is recommended.

Images

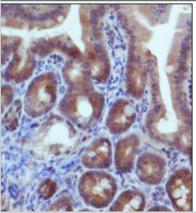
Western Blot: IDH2 Antibody [NBP2-22166] - 1) Non transfected 293T cells. 2) IDH2-flag (c-term) transfected 293T cells. WB image submitted by a verified customer review.



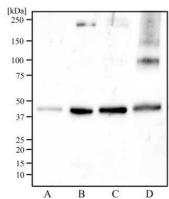
Immunocytochemistry/Immunofluorescence: IDH2 Antibody [NBP2-22166] - 293T cells transfected with IDH2-flag expression construct were fixed and stained with anti-IDH2 antibody followed by Alexa Fluor 488 anti-rabbit secondary antibody. DAPI staining shows nuclei. ICC/IF image submitted by a verified customer review.



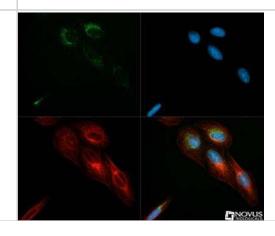
Immunohistochemistry: IDH2 Antibody [NBP2-22166] - Staining of IDH2 in mouse intestine.



Western Blot: IDH2 Antibody [NBP2-22166] - Analysis of Caki-1 cell lysate (A), human skeletal muscle tissue (B), mouse skeletal muscle tissue (C), and rat skeletal muscle tissue (D) using IDH2 antibody at a concentration of 2 ug/mL.



Immunocytochemistry/Immunofluorescence: IDH2 Antibody [NBP2-22166] - IDH2 antibody was tested in U2OS cells with DyLight (green). Nuclei and alpha-tubulin were counterstained with DAPI (blue) and DyLight 550 (red).



Publications

Lin S, Yin S, Shi J et al. Orchestration of energy metabolism and osteogenesis by Mg2+ facilitates low-dose BMP-2-driven regeneration Bioactive Materials 2022-12-01 [PMID: 35387176] (IF/IHC, Mouse)

Oyama Y, Bartman CM, Bonney S et al. Intense Light-Mediated Circadian Cardioprotection via Transcriptional Reprogramming of the Endothelium Cell Rep 2019-08-08 [PMID: 31390562] (WB, Human)

Kim YR, Kim KH, Lee S et al. Expression patterns of members of the isocitrate dehydrogenase gene family in murine inner ear Biotech Histochem 2017-09-19 [PMID: 28925723] (Mouse)

Saha SK, Parachoniak CA, Ghanta KS et al. Mutant IDH inhibits HNF-4alpha to block hepatocyte differentiation and promote biliary cancer. Nature 2014-09-04 [PMID: 25043045] (WB)

Saha SK, Gordan JD, Kleinstiver BP et al. Isocitrate dehydrogenase mutations confer dasatinib hypersensitivity and SRC-dependence in intrahepatic cholangiocarcinoma. Cancer Discov. 2016-05-26 [PMID: 27231123] (WB, Human)



Procedures

Western Blot protocol for IDH2 Antibody (NBP2-22166)

IDH2 Antibody:

Western Blot Protocol

- 1. Perform SDS-PAGE on samples to be analyzed, loading 25 ug of total protein per lane.
- 2. Transfer proteins to membrane according to the instructions provided by the manufacturer of the membrane and transfer apparatus.
- 3. Stain according to standard Ponceau S procedure (or similar product) to assess transfer success, and mark molecular weight standards where appropriate.
- 4. Rinse the blot.
- 5. Block the membrane using standard blocking buffer for at least 1 hour.
- 6. Wash the membrane in wash buffer three times for 10 minutes each.
- 7. Dilute anti-IDH2 primary antibody in blocking buffer and incubate 1 hour at room temperature.
- 8. Wash the membrane in wash buffer three times for 10 minutes each.
- 9. Apply the diluted HRP conjugated secondary antibody in blocking buffer (as per manufacturers instructions) and incubate 1 hour at room temperature.
- 10. Wash the blot in wash buffer three times for 10 minutes each (this step can be repeated as required to reduce background).
- 11. Apply the detection reagent of choice in accordance with the manufacturers instructions.

Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%.

Immunohistochemistry-Paraffin protocol for IDH2 Antibody (NBP2-22166)

IDH2 Antibody:

Immunohistochemistry-Paraffin Embedded Sections

Antigen Unmasking:

Bring slides to a boil in 10 mM sodium citrate buffer (pH 6.0) then maintain at a sub-boiling temperature for 10 minutes. Cool slides on bench-top for 30 minutes.

Staining:

- 1. Wash sections in deionized water three times for 5 minutes each.
- Wash sections in wash buffer for 5 minutes.
- 3. Block each section with 100-400 ul blocking solution for 1 hour at room temperature.
- 4. Remove blocking solution and add 100-400 ul diluted primary antibody. Incubate overnight at 4 C.
- 5. Remove antibody solution and wash sections in wash buffer three times for 5 minutes each.
- 6. Add 100-400 ul biotinylated diluted secondary antibody. Incubate 30 minutes at room temperature.
- 7. Remove secondary antibody solution and wash sections three times with wash buffer for 5 minutes each.
- 8. Add 100-400 ul Streptavidin-HRP reagent to each section and incubate for 30 minutes at room temperature.
- 9. Wash sections three times in wash buffer for 5 minutes each.
- 10. Add 100-400 ul DAB substrate to each section and monitor staining closely.
- 11. As soon as the sections develop, immerse slides in deionized water.
- 12. Counterstain sections in hematoxylin.
- 13. Wash sections in deionized water two times for 5 minutes each.
- 14. Dehydrate sections.
- 15. Mount coverslips.



Immunocytochemistry/Immunofluorescence protocol for IDH2 Antibody (NBP2-22166)

IDH2 Antibody:

Immunocytochemistry Protocol

Culture cells to appropriate density in 35 mm culture dishes or 6-well plates.

- 1. Remove culture medium and add 10% formalin to the dish. Fix at room temperature for 30 minutes.
- 2. Remove the formalin and add ice cold methanol. Incubate for 5-10 minutes.
- 3. Remove methanol and add washing solution (i.e. PBS). Be sure to not let the specimen dry out. Wash three times for 10 minutes.
- 4. To block nonspecific antibody binding incubate in 10% normal goat serum from 1 hour to overnight at room temperature.
- 5. Add primary antibody at appropriate dilution and incubate at room temperature from 2 hours to overnight at room temperature.
- 6. Remove primary antibody and replace with washing solution. Wash three times for 10 minutes.
- 7. Add secondary antibody at appropriate dilution. Incubate for 1 hour at room temperature.
- 8. Remove antibody and replace with wash solution, then wash for 10 minutes. Add Hoechst 33258 to wash solution at 1:25,0000 and incubate for 10 minutes. Wash a third time for 10 minutes.
- 9. Cells can be viewed directly after washing. The plates can also be stored in PBS containing Azide covered in Parafilm (TM). Cells can also be cover-slipped using Fluoromount, with appropriate sealing.

*The above information is only intended as a guide. The researcher should determine what protocol best meets their needs. Please follow safe laboratory procedures.





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Products Related to NBP2-22166

NB820-59253 Human Skeletal Muscle Whole Tissue Lysate (Adult Whole Normal)

NBP2-22166PEP IDH2 Antibody Blocking Peptide

HAF008 Goat anti-Rabbit IgG Secondary Antibody [HRP]

NB7160 Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]

NBP2-24891 Rabbit IgG Isotype Control

Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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