

Product Datasheet

Sulfatase-2/SULF2 Antibody

NBP1-36727SS

Unit Size: 0.025 ml

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

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Publications: 4

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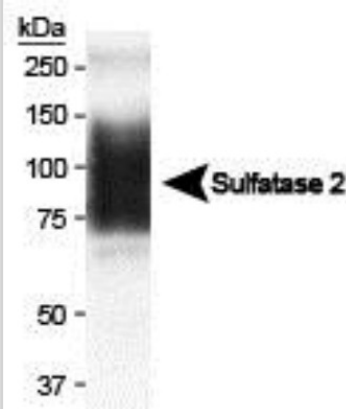
NBP1-36727SS**Sulfatase-2/SULF2 Antibody (2B4)**

Product Information	
Unit Size	0.025 ml
Concentration	1 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Monoclonal
Clone	2B4
Preservative	0.05% Sodium Azide
Isotype	IgG1
Purity	Protein G purified
Buffer	PBS
Product Description	
Host	Mouse
Gene ID	55959
Gene Symbol	SULF2
Species	Human, Mouse
Species Reactivity	Human and mouse.
Specificity/Sensitivity	Does not recognize human or mouse Sulfatase 1 or other family members.
Immunogen	Partial recombinant human Sulfatase 2 isoform b [Accession# NP_940998].
Product Application Details	
Applications	Western Blot, ELISA, Flow Cytometry, Immunohistochemistry, Immunohistochemistry-Paraffin, Immunoprecipitation
Recommended Dilutions	ELISA 1:100-1:2000, Flow Cytometry, Immunohistochemistry 1:250-1:500, Immunohistochemistry-Paraffin 1:250-1:500, Immunoprecipitation 1:10-1:100, Western Blot 1:1000
Application Notes	This Sulfatase 2 (2B4) antibody is useful for Western Blot, Immunoprecipitation, Immunohistochemistry on paraffin-embedded sections and ELISA. For WB a band can be seen at 50 kDa. Other bands may be seen at 135 kDa (unprocessed protein) and 37 kDa (fragment of 50 kDa subunit). For ELISA, use this antibody at 1-10 ug/ml for capture and 1-5 ug/ml for detection. It has been reported that NBP1-36727 picks up a 130 kDa band that is not Sulfatase 2 in human plasma. Because of this we do not recommend NBP1-36727 for capture of the human protein. Flow data from customer review on FITC-conjugated antibody.

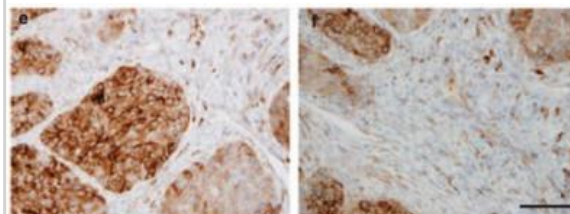


Images

Western Blot: Sulfatase 2 Antibody (2B4) [NBP1-36727] - Analysis showing Sulfatase 2 expression in MCF7 conditioned media (CM). In CM, the 75 kDa band is dominant (Tang R, Rosen SD. J Biol Chem. 2009 Aug 7;284(32):21505-14). Image courtesy of Mark Singer at UCSF.



Immunohistochemistry-Paraffin: Sulfatase 2 Antibody (2B4) [NBP1-36727] - Immunohistochemical analysis of Sulfatase 2 in non-small-cell lung carcinoma. (left) lung squamous cell carcinoma stained with 2B4 antibody and (right) tumor-associated stromal cells stained with 2B4 antibody. Image courtesy of Mark Singer at UCSF.



Publications

Solari V, Borriello L, Turcatel G et al. MYCN-dependent Expression of Sulfatase-2 Regulates Neuroblastoma Cell Survival Cancer Res. 2014 Aug 27 [PMID: 25164011]

Yue X, Lu J, Auduong L et al. Overexpression of Sulf2 in idiopathic pulmonary fibrosis. Glycobiology 2013 Feb 14 [PMID: 23418199] (WB, Human)

Lui NS, van Zante A, Rosen SD et al. SULF2 expression by immunohistochemistry and overall survival in oesophageal cancer: a cohort study BMJ Open 2012 Nov 23 [PMID: 23180455] (IHC, Human)

Lemjabbar-Alaoui H, van Zante A, Singer MS, Xue Q, Wang YQ, Tsay D, He B, Jablons DM, Rosen SD. Sulf-2, a heparan sulfate endosulfatase, promotes human lung carcinogenesis. Oncogene;29(5):635-46. 2010 Feb 4. [PMID: 19855436] (WB, IHC-P, Human)

Procedures

Protocol specific for Sulfatase 2 Antibody (NBP1-36727)

Western Blot Protocol

1. Perform SDS-PAGE (4-12% MOPS) on samples to be analyzed, loading 25 ug of total protein per lane.
 2. Transfer proteins to Nitrocellulose according to the instructions provided by the manufacturer of the transfer apparatus.
 3. Rinse membrane with dH₂O and then stain the blot using Ponceau S for 1-2 minutes to access the transfer of proteins onto the nitrocellulose membrane. Rinse the blot in water to remove excess stain and mark the lane locations and locations of molecular weight markers using a pencil.
 4. Rinse the blot in TBS for approximately 5 minutes.
 5. Block the membrane using 5% NFDM + 1% BSA in TBS + Tween, 1 hour at RT.
 6. Rinse the membrane in dH₂O and then wash the membrane in wash buffer [TBS + 0.1% Tween] 3 times for 10 minutes each.
 7. Dilute the anti-Sulfatase 2 primary antibody (NBP1-36727) in blocking buffer and incubate 1 hour at room temperature.
 8. Rinse the membrane in dH₂O and then wash the membrane in wash buffer [TBS + 0.1% Tween] 3 times for 10 minutes each.
 9. Apply the diluted mouse-IgG 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 [TBS + 0.1% Tween] 3 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 (Pierce ECL).
- Note: Tween-20 can be added to the blocking or antibody dilution buffer at a final concentration of 0.05-0.2%, provided it does not interfere with antibody-antigen binding.





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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.

For more information on our guarantee, please visit www.novusbio.com/guarantee.

