

Product Datasheet

Cathepsin B Antibody - BSA Free NBP1-19797

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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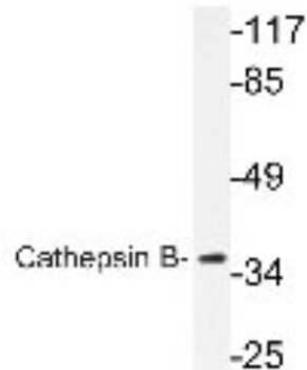
NBP1-19797

Cathepsin B Antibody - BSA Free

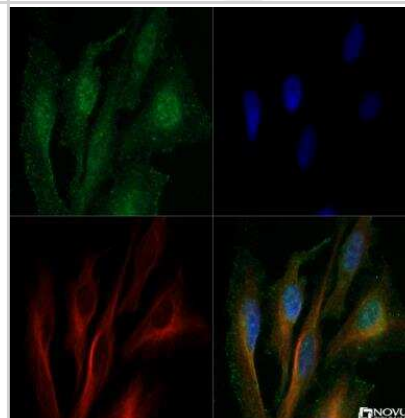
Product Information	
Unit Size	0.1 ml
Concentration	1.0 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Polyclonal
Preservative	0.05% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	PBS
Product Description	
Host	Rabbit
Gene ID	1508
Gene Symbol	CTSB
Species	Human, Mouse, Rat
Reactivity Notes	Immunogen displays the following percentage of sequence identity for non-tested species: bovine (84%) and chicken (84%).
Immunogen	Synthetic peptide made to an internal portion of the Cathepsin B human protein (between residues 200-270) [UniProt P07858]
Product Application Details	
Applications	Western Blot, Immunohistochemistry-Paraffin, Immunocytochemistry/Immunofluorescence, Immunohistochemistry
Recommended Dilutions	Western Blot 1.0 - 2.0 ug/ml, Immunohistochemistry 1:200, Immunocytochemistry/ Immunofluorescence 1:50 - 1:100, Immunohistochemistry-Paraffin 1:200

Images

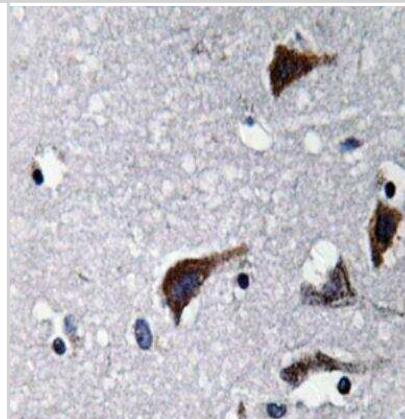
Western Blot: Cathepsin B Antibody [NBP1-19797] - extracts from COLO cells.



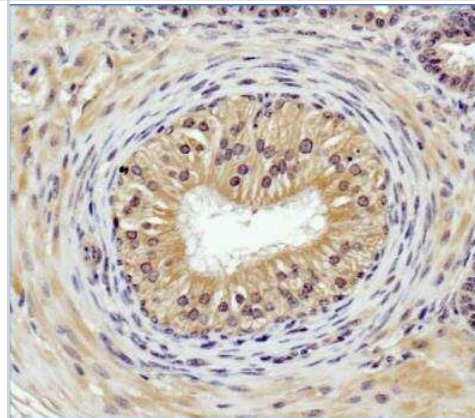
Immunocytochemistry/Immunofluorescence: Cathepsin B Antibody [NBP1-19797] - ICC/IF analysis of HeLa cells using 1:50 dilution of Cathepsin B antibody followed by detection using Dylight 488 labelled secondary antibody (green). Nuclei and alpha-tubulin were counterstained using DAPI (blue) and Dylight 550 (red) respectively. Punctate staining was observed for Cathepsin B indicating its presence in the cellular lysosomes, endosomes and melanosomes. Image objective 40x.



Immunohistochemistry-Paraffin: Cathepsin B Antibody [NBP1-19797] - Human brain tissue.



Immunohistochemistry-Paraffin: Cathepsin B Antibody [NBP1-19797] - Staining for Cathepsin B in mouse ureter.



Publications

Felix Humphries, Bridget Chang-McDonald, Josie Patel, Nicholas Bockett, Erin Paterson, Paul F. Davis, Swee T. Tan
Cathepsins B, D, and G Are Expressed in Metastatic Head and Neck Cutaneous Squamous Cell Carcinoma *Frontiers in Oncology* 2021-09-21 [PMID: 34621666]

Murray ER, Menezes S, Henry JC et al. Disruption of pancreatic stellate cell myofibroblast phenotype promotes pancreatic tumor invasion *Cell reports* 2022-01-25 [PMID: 35081338] (IF/IHC, Mouse)

Hansen L, Brasch HD, Paterson E Et al. Expression of Cathepsins B, D, and G in Extracranial Arterio-Venous Malformation *Frontiers in surgery* 2021-08-02 [PMID: 34409065] (IHC-P, Human)

Maniati E, Berlato C, Gopinathan G et al. Mouse Ovarian Cancer Models Recapitulate the Human Tumor Microenvironment and Patient Response to Treatment *Cell Rep* 2020-01-14 [PMID: 31940494] (IF/IHC, Mouse)

Farizatto K. Study of hyperphosphorylated Tau protein degradation by proteasome-independent pathways, in an experimental model of neurodegeneration Thesis. 2014-04-28 (WB, Rat)



Procedures

Western Blot protocol for Cathepsin B Antibody (NBP1-19797)

Cathepsin B Antibody:

Western Blot Protocol for Cathepsin B antibody

1. Perform SDS-PAGE on samples to be analyzed, loading 40 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 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 Protocol for Cathepsin B antibody

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.
2. 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 Protocol for Cathepsin B antibody

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 1:25,000 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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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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