

Product Datasheet

Bcl-xL Antibody NB110-83982

Unit Size: 0.1 ml

Store at 4C. Do not freeze.

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NB110-83982

Bcl-xL Antibody

Product Information

Unit Size	0.1 ml
Concentration	1 mg/ml
Storage	Store at 4C. Do not freeze.
Clonality	Polyclonal
Preservative	0.1% Sodium Azide
Isotype	IgG
Purity	Immunogen affinity purified
Buffer	Tris-Citrate/Phosphate (pH 7.0 - 8.0)
Target Molecular Weight	26 kDa

Product Description

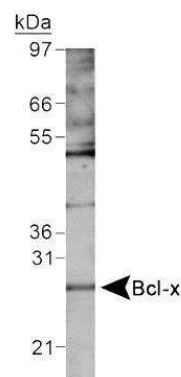
Host	Rabbit
Gene Symbol	BCL2L1
Species	Human, Canine, Feline, Sheep
Reactivity Notes	Immunogen displays the following percentage of sequence identity for non-tested species: Mouse (87%), Rat (81%)
Specificity/Sensitivity	There are three known isoforms of Bcl-X; Bcl-X(S), Bcl-X(beta), and Bcl-X(L). The immunogen used to create this has 100% identity to all three isoforms.
Immunogen	Synthetic peptide made to an N-terminal portion of human BCL2L1 (within residues 30-70). [Swiss-Prot# Q07817]

Product Application Details

Applications	Western Blot, ICC/IF (Negative)
Recommended Dilutions	Western Blot 2 ug/ml, ICC/IF (Negative)
Application Notes	This BCL2L1 antibody is useful for Western blot, where a band is seen ~26 kDa. This antibody is not useful for ICC/IF. The observed molecular weight of the protein may vary from the listed predicted molecular weight due to post translational modifications, post translation cleavages, relative charges, and other experimental factors.

Images

Western Blot: Bcl-XL Antibody [NB110-83982] - Detection of BCL2L1 in whole cell extract of A431 cell line using NB110-83982. The band at ~26kDa is the specific target band, whereas the band above ~50 kDa position is potentially a homodimer of BCL2L1.



Procedures

Protocol specific for BCL2L1 Antibody (NB110-83982)

Bcl-xL Antibody:

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% 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 rabbit anti-Bcl-X primary antibody (NB 110-83982) 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 rabbit-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.

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