

Product Datasheet

PUMA Antibody NB500-261SS

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: 1

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Updated 6/15/2014 v.20.1

NB500-261SS

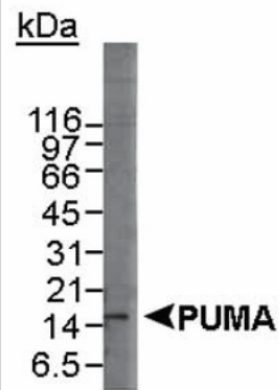
PUMA Antibody

Product Information	
Unit Size	0.025 ml
Concentration	5.65 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
Purity	Immunogen affinity purified
Buffer	Tris-glycine, 150 mM NaCl
Target Molecular Weight	16 kDa
Product Description	
Host	Rabbit
Gene ID	27113
Gene Symbol	BBC3
Species	Human
Species Reactivity	Human.
Immunogen	A synthetic peptide made to a region within the C-terminus of the human PUMA protein sequence (between residues 150-193). [UniProt# Q9BXH1]
Product Application Details	
Applications	Western Blot, Immunocytochemistry/Immunofluorescence
Recommended Dilutions	Immunocytochemistry/Immunofluorescence 1:20-1:100, Western Blot 2-4 ug/ml
Application Notes	This PUMA antibody is useful for Western blot and Immunocytochemistry/Immunofluorescence. By WB a band at ~16 kDa is seen, representing the beta form of PUMA. Reactivity against the alpha form of PUMA is unknown.

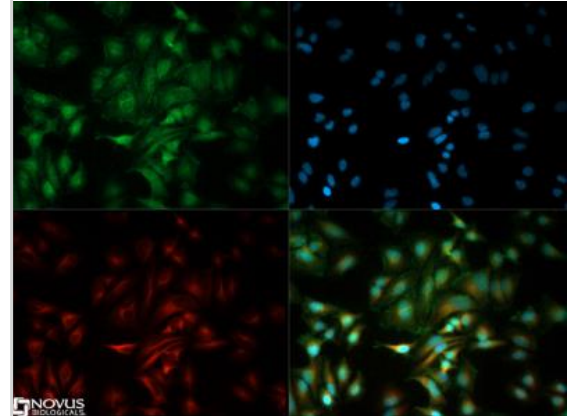


Images

Western Blot: PUMA Antibody [NB500-261] - Detection of the beta isoform of PUMA in HL-60 whole cell lysate using NB 500-261(4 ug/ml). ECL detection at a 30 minute exposure.



Immunocytochemistry/Immunofluorescence: PUMA Antibody [NB500-261] - The PUMA antibody was tested at a 1:20 dilution in HeLa cells against Dylight 488 (Green). Alpha-tubulin and nuclei were counterstained against Dylight 550 (Red) and DAPI (Blue).



Publications

Stylianou, S, Clarke, R B, Brennan, K. Aberrant activation of notch signaling in human breast cancer. Cancer Res;66 (3):1517-25. 2006 Feb 1. [PMID: 16452208] (WB, Human)

Procedures

Protocol specific for PUMA Antibody (NB500-261)

Western Blot Protocol

1. Perform SDS-PAGE (4-12%) on samples to be analyzed, loading 35 ug of total protein per lane.
2. Transfer proteins to Nitrocellulose according to the instructions provided by the manufacturer of the transfer apparatus.
3. 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% non-fat dry milk + 0.5% BSA in TBS for 1 hour.
6. Dilute the rabbit anti-PUMA primary antibody (NB 500-261) in blocking buffer and incubate 2 hours at room temperature.
7. Wash the membrane in water for 5 minutes and apply the diluted rabbit-IgG HRP-conjugated secondary antibody in blocking buffer (as per manufacturer's instructions) and incubate 1 hour at room temperature.
8. Wash the blot in TBS containing 0.05-0.1% Tween-20 for 10-20 minutes.
9. Wash the blot in type I water for an additional 10-20 minutes (this step can be repeated as required to reduce background).
10. Apply the detection reagent of choice in accordance with the manufacturer's instructions (Amersham's ECL is the standard reagent used at Novus Biologicals).

Note: Tween-20 can be added to the blocking 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.

