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anti-BNIP1 antibody (C-Term)

3 Images



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Overview

Quantity:	100 μL
Target:	BNIP1
Binding Specificity:	C-Term
Reactivity:	Human, Mouse, Rat, Cow
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This BNIP1 antibody is un-conjugated
Application:	Western Blotting (WB), Immunohistochemistry (IHC), Immunofluorescence (IF), Immunoprecipitation (IP), Immunochromatography (IC)
Product Details	
Immunogen:	KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of human NIP1.
Specificity:	Recognizes endogenous levels of NIP1 protein.
Characteristics:	Rabbit polyclonal antibody to NIP1
Purification:	The antibody was purified by immunogen affinity chromatography.
Target Details	
Target:	BNIP1
Alternative Name:	NIP1 (BNIP1 Products)

Target Details

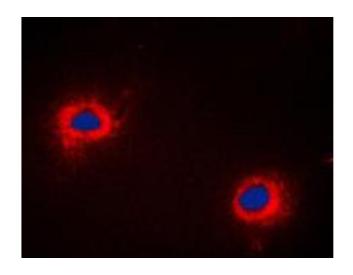
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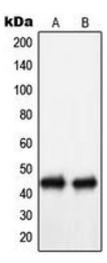
12 months

Expiry Date:

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Background:	APBA2BP, NIP1, SYTIP2, XB51, N-terminal EF-hand calcium-binding protein 3, Amyloid beta A4 protein-binding family A member 2-binding protein, Nek2-interacting protein 1, Neuronal calcium-binding protein 3, X11L-binding protein 51
Gene ID:	63941, 56846
UniProt:	Q96P71, Q9D6J4
Application Details	
Application Notes:	WB (1:500 - 1:1000), IH (1:100 - 1:200), IF/IC (1:100 - 1:500), IP (1:10 - 1:100)
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	Liquid in 0.42 % Potassium phosphate, 0.87 % Sodium chloride, pH 7.3, 30 % glycerol, and 0.01 % sodium azide.
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C

Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid freeze/thaw cycles.





Immunofluorescence

Image 1. Immunofluorescent analysis of NIP1 staining in COLO205 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

Immunohistochemistry

Image 2. Immunohistochemical analysis of NIP1 staining in human lung cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

Western Blotting

Image 3. Western blot analysis of NIP1 expression in Ramos (A), COLO205 (B) whole cell lysates.