

Product Data Sheet

Anti-BNIP3L Antibody

Catalog #	Source	e Reactivity	Applications	
CQA3171	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody to) BNIP3L	
Immunogen		Recombinant full length prot	ein of human BNIP3L	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of BNIP3L protein.	
Clonality		Polyclonal		
Conjugation				
Form Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7		osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/5	0 - 1/200)	
Gene Symbol		BNIP3L		
Alternative Names		BNIP3A; BNIP3H; NIX; BCL2/adenovirus E1B 19 kDa protein-interacting protein		
		3-like; Adenovirus E1B19K-bi	nding protein B5; BCL2/adenovirus E1B 19 kDa	
		protein-interacting protein 3	A; NIP3-like protein X; NIP3L	
Entrez Gene		665 (Human); 12177 (Mouse)	
SwissProt		O60238 (Human); Q9Z2F7 (N	1ouse)	
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid	
		freeze/thaw cycles.		

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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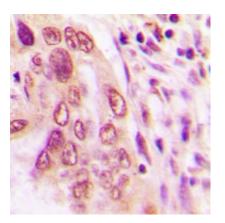
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For research purposes only, not for human use

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Western blot analysis of BNIP3L expression in K562 (A), MCF7 (B) whole cell lysates. (Predicted band size: 19; 23 kD; Observed band size: 35 kD)



Immunohistochemical analysis of BNIP3L 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.

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