

Product Data Sheet

Anti-NLRX1 Antibody

Catalog #	Source	e Reactivity	Applications	
CPA7502	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody t	D NLRX1	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the center	
		region of human NLRX1. The	exact sequence is proprietary.	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of NLRX1 protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium pl	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/2000), IH (1/50	- 1/200)	
Gene Symbol		NLRX1		
Alternative Names		NOD26; NOD5; NOD9; NLR family member X1; Caterpiller protein 11.3; CLR11.3;		
		Nucleotide-binding oligomer	ization domain protein 26; Nucleotide-binding	
		oligomerization domain prot	ein 5; Nucleotide-binding oligomerization domain	
		protein 9		
Entrez Gene		79671 (Human); 270151 (Mo	ouse); 315599 (Rat)	
SwissProt		Q86UT6 (Human); Q3TL44 (I	Mouse); Q5FVQ8 (Rat)	
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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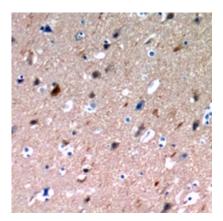
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Western blot analysis of NLRX1 expression in HepG2 (A), Hela (B) whole cell lysates. (Predicted band size: 107 kD; Observed band size: 107 kD)



Immunohistochemical analysis of NLRX1 staining in human brain 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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