

Product Data Sheet

Anti-Nectin 4 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5596	Rabbit	H, M, R	WB, IH		
Description	R	Rabbit polyclonal antibody to Nectin 4			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
	r	region of human Nectin 4. The exact sequence is proprietary.			
Purification The antibody was purified by immunogen affinit			immunogen affinity chromatography.		
Specificity	Recognizes endogenous levels of Nectin 4 protein.				
Clonality	Р	Polyclonal			
Conjugation					
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	Р	VRL4			
Alternative Na	i <mark>mes</mark> L	NIR; PRR4; Poliovirus recep	tor-related protein 4; Ig superfamily receptor LNIR;		
	Ν	lectin-4			
Entrez Gene	8	1607 (Human); 71740 (Mo	ise)		
SwissProt	C	Q96NY8 (Human); Q8R007 (Mouse)		
Storage/Stabil	ity S	hipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fı	reeze/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

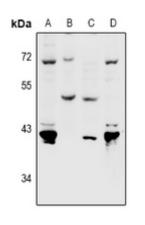
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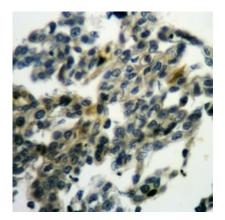


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Western blot analysis of Nectin 4 expression in Hela (A), MCF7 (B), MEF (C), PC12 (D) whole cell lysates. (Predicted band size: 55 kD; Observed band size: 52; 40; 66 kD)



Immunohistochemical analysis of Nectin 4 staining in human ovarian 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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