

Product Data Sheet

Anti-Netrin 1 Antibody

Catales #	Course	Depativity	Applications		
Catalog #	Source	Reactivity	Applications		
CPA2389	Rabbit	H, M, R, Mk	WB, IH		
Description	Rabb	Rabbit polyclonal antibody to Netrin 1			
Immunogen	KLH-	conjugated synthetic pept	ide encompassing a sequence within the C-term		
	regio	on of human Netrin 1. The	exact sequence is proprietary.		
Purification	The	The antibody was purified by immunogen affinity chromatography.			
Specificity	Reco	gnizes endogenous levels	of Netrin 1 protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/100 -	- 1/200)		
Gene Symbol	NTN	1			
Alternative Names N		NTN1L; Netrin-1; Epididymis tissue protein Li 131P			
Entrez Gene 9423 (H		8 (Human); 18208 (Mouse)	luman); 18208 (Mouse)		
SwissProt	095	O95631 (Human); O09118 (Mouse); Q924Z9 (Rat)			
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	free	ze/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

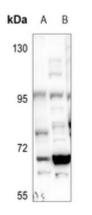
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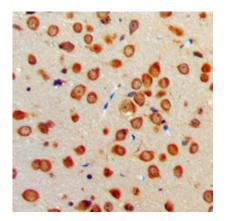
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Western blot analysis of Netrin 1 expression in Hela (A), U87MG (B), CT26 (C) whole cell lysates. (Predicted band size: 67 kD; Observed band size: 66 kD)



Immunohistochemical analysis of Netrin 1 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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