

# **Product Data Sheet**

## Anti-LRP10 Antibody

Catalog #	Source	Reactivity	Applications			
CPA5586	Rabbit	H, M, R	WB, IH			
Description	Rabb	Rabbit polyclonal antibody to LRP10				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center				
	regio	region of human LRP10. The exact sequence is proprietary.				
Purification	Thea	The antibody was purified by immunogen affinity chromatography.				
Specificity	Reco	gnizes endogenous lev	zes endogenous levels of LRP10 protein.			
Clonality	Poly	Polyclonal				
Conjugation						
Form	Liqui	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	and	0.01% sodium azide.				
Dilution	WB (	1/500 - 1/1000), IH (1/50	) - 1/100)			
Gene Symbol	LRP1	0				
Alternative Na	ames Low-	Low-density lipoprotein receptor-related protein 10; LRP-10				
Entrez Gene 26020		0 (Human); 65107 (Mo	(Human); 65107 (Mouse)			
SwissProt	Q7Z4	Q7Z4F1 (Human); Q7TQH7 (Mouse)				
Storage/Stabi	lity Ship	oed at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	freez	e/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

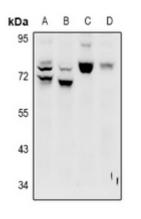
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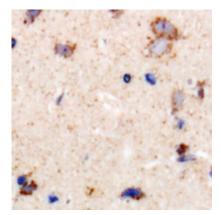


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Western blot analysis of LRP10 expression in HepG2 (A), A549 (B), PC12 (C), CT26 (D) whole cell lysates. (Predicted band size: 76 kD; Observed band size: 76 kD)



Immunohistochemical analysis of LRP10 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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