

Product Data Sheet

Anti-LRP11 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2823	Rabbit	Н	WB, IP		
Description	R	Rabbit polyclonal antibody to LRP11			
Immunogen	К	(LH-conjugated synthetic p	eptide encompassing a sequence within the center		
	r	egion of human LRP11. Th	e exact sequence is proprietary.		
Purification	Т	he antibody was purified	by immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous lev	els of LRP11 protein.		
Clonality	Р	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IP (1/1	0 - 1/100)		
Gene Symbol	L	RP11			
Alternative Na	a <mark>mes</mark> L	ow-density lipoprotein re	ceptor-related protein 11; LRP-11		
Entrez Gene		84918 (Human)			
SwissProt	C	Q86VZ4 (Human)			
Storage/Stabi	lity S	hipped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid		
	fi	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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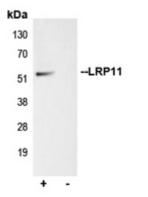
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Western blot analysis of LRP11 expression in A431 (A), HepG2 (B), Jurkat (C) whole cell lysates. (Predicted band size: 53 kD; Observed band size: 49 kD)



Immunoprecipitation of LRP11 from 0.5mg Jurkat whole cell extract lysate, using 5ug of Anti-LRP11 Antibody and 50ul of protein G magnetic beads (+). No antibody was added to the control (-). The antibody was incubated under agitation with Protein G beads for 10min, Jurkat whole cell extract lysate diluted in RIPA buffer was added to each sample and incubated for a further 10min under agitation. Proteins were eluted by addition of 40ul SDS loading buffer and incubated for 10min at 70°C; 10ul of each sample was separated on a SDS PAGE gel, transferred to a nitrocellulose membrane, blocked with 5% BSA and probed with Anti-LRP11 Antibody.

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