

Product Data Sheet

Anti-RPL26L1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2658	Rabbit	H, M, R	WB, IH		
Description	Rat	Rabbit polyclonal antibody to RPL26L1			
Immunogen	KLł	H-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
	reg	gion of human RPL26L1. T	ne exact sequence is proprietary.		
Purification	The	e antibody was purified by	immunogen affinity chromatography.		
Specificity	Red	cognizes endogenous leve	ls of RPL26L1 protein.		
Clonality	Pol	Polyclonal			
Conjugation					
Form	Liq	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	d 0.01% sodium azide.			
Dilution	WE	3 (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	RPI	L26L1			
Alternative Na	ames RPI	L26P1; 60S ribosomal pro	ein L26-like 1		
Entrez Gene 51		51121 (Human)			
SwissProt	Q9	Q9UNX3 (Human)			
Storage/Stabi	lity Shi	ipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fre	eze/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

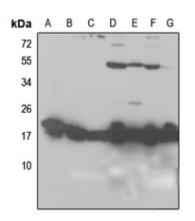
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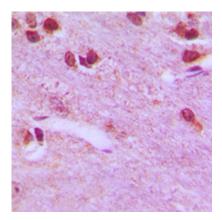


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Western blot analysis of RPL26L1 expression in HEK293T (A), Hela (B), H1688 (C), mouse liver (D), mouse kidney (E), rat liver (F), rat kidney (G) whole cell lysates. (Predicted band size: 17 kD; Observed band size: 17 kD)



Immunohistochemical analysis of RPL26L1 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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