

Product Data Sheet

Anti-RPS19BP1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4216	Rabbit	H, M, R, Mk	WB, IH		
Description	Ra	bbit polyclonal antibody to	o RPS19BP1		
Immunogen	KL	H-conjugated synthetic pe	ptide encompassing a sequence within the center		
	re	gion of human RPS19BP1.	The exact sequence is proprietary.		
Purification	Th	e antibody was purified by	/ immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous leve	ls of RPS19BP1 protein.		
Clonality	onality Polyclonal				
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/100	0 - 1/200)		
Gene Symbol	RP	PS19BP1			
Alternative Na	ames AR	OS; Active regulator of SIR	RT1; 40S ribosomal protein S19-binding protein 1;		
	RP	S19-binding protein 1; S19)BP		
Entrez Gene	91	.582 (Human); 66538 (Mou	use)		
SwissProt	Q	36WX3 (Human); Q8C6B9 ((Mouse)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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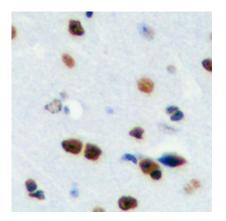
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Western blot analysis of RPS19BP1 expression in H446 (A) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 20 kD)



Immunohistochemical analysis of RPS19BP1 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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