

Product Data Sheet

Anti-CRMP2 (Phospho-S522) Antibody

Catalog #	Source	Reactivity	Applications		
CPA3902	Rabbit	H, M, R, B, C, Mk, P	WB, IH		
Description	Rabb	it polyclonal antibody to CRMP2	(Phospho-S522)		
Immunogen	KLH-	conjugated synthetic phosphope	ptide corresponding to residues surrounding		
	S522	of human CRMP2 protein. The	exact sequence is proprietary.		
Purification	The	antibody was purified by immun	ogen affinity chromatography.		
Specificity	Recc	gnizes endogenous levels of CRN	/IP2 protein only when phosphorylated at S522.		
Clonality	Poly	clonal			
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	DPYS	SL2			
Alternative N	ames CRM	P2; ULIP2; Dihydropyrimidinase	related protein 2; DRP-2; Collapsin response		
	med	iator protein 2; CRMP-2; N2A3; l	Inc-33-like phosphoprotein 2; ULIP-2		
Entrez Gene	1808	8 (Human); 12934 (Mouse); 2541	6 (Rat)		
SwissProt	Q16	555 (Human); O08553 (Mouse);	P47942 (Rat)		
Storage/Stabi	lity Ship	oed at 4°C. Upon delivery aliquo	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

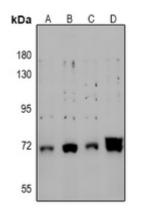
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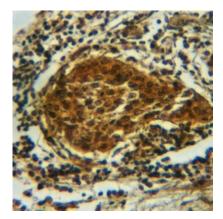


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Western blot analysis of CRMP2 (Phospho-S522) expression in A375 (A), MCF7 (B), PC12 (C), AML12 (D) whole cell lysates. (Predicted band size: 62 kD; Observed band size: 70 kD)



Immunohistochemical analysis of CRMP2 (Phospho-S522) staining in human kidney cancer 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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