

Product Data Sheet

Anti-p116 Rip Antibody

Catalog #	Source	Reactivity	Applications		
CPA3396	Rabbit	H, M, R, B, P	WB, IH		
Description		Rabbit polyclonal antibody to p116 Rip			
Immunogen		KLH-conjugated synthetic pep	tide encompassing a sequence within the center		
		region of human p116 Rip. Th	e exact sequence is proprietary.		
Purification		The antibody was purified by	mmunogen affinity chromatography.		
Specificity		Recognizes endogenous levels	of p116 Rip protein.		
Clonality		Polyclonal			
Conjugation					
Form		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 (1/100	- 1/200)		
Gene Symbol		MPRIP			
Alternative N	ames	KIAA0864; MRIP; RHOIP3; My	osin phosphatase Rho-interacting protein; M-RIP;		
		Rho-interacting protein 3; RIP	3; p116Rip		
Entrez Gene		23164 (Human); 26936 (Mouse); 116504 (Rat)			
SwissProt		Q6WCQ1 (Human); P97434 (Mouse); Q9ERE6 (Rat)			
Storage/Stabi	ility	Shipped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
		freeze/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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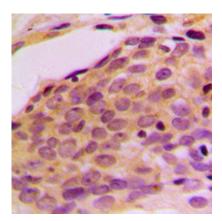
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KDa A B C D 180 130 95 72 Western blot analysis of p116 Rip expression in PC12 (A), MEF (B), A549 (C), A2780 (D) whole cell lysates. (Predicted band size: 116 kD; Observed band size: 116 kD)



Immunohistochemical analysis of p116 Rip staining in human breast 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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