

Product Data Sheet

Anti-MRCK alpha Antibody

Catalog #	Source	Reactivity	Applications			
CPA2282	Rabbit	H, M, R	WB, IF/IC			
Description	Ra	Rabbit polyclonal antibody to MRCK alpha				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center				
	re	region of human MRCK alpha. The exact sequence is proprietary.				
Purification		The antibody was purified by immunogen affinity chromatography.				
Specificity		Recognizes endogenous levels of MRCK alpha protein.				
Clonality	Po	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	ar	nd 0.01% sodium azide.				
Dilution	W	/B (1/500 - 1/1000), IF/IC (1	/50 - 1/200)			
Gene Symbol	CI	DC42BPA				
Alternative Na	ames Kl	IAA0451; Serine/threonine	e-protein kinase MRCK alpha; CDC42-binding protein			
	ki	nase alpha; DMPK-like alp	ha; Myotonic dystrophy kinase-related CDC42-binding			
	ki	nase alpha; MRCK alpha; i	Myotonic dystrophy protein kinase-like alpha			
Entrez Gene	84	476 (Human); 226751 (Mo	use); 114116 (Rat)			
SwissProt	Q	5VT25 (Human); Q3UU96	(Mouse); O54874 (Rat)			
Storage/Stabi	lity Sł	nipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	fr	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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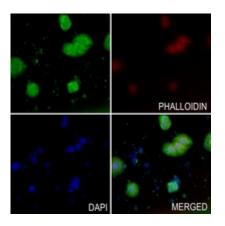
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KDA A B C D E

Western blot analysis of MRCK alpha expression in C6 (A), BV2 (B), HEK293T (C), Panc1 (D), U87MG (E) whole cell lysates. (Predicted band size: 197 kD; Observed band size: 186 kD)



Immunofluorescent analysis of MRCK alpha staining in Hela cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin - AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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