

Product Data Sheet

Anti-ERK5 Antibody

Catalog #	Source	Reactivity	Applications		
CQA2069	Rabbit	H, M, R	WB, IH, IF/IC		
Description	R	abbit polyclonal antibody	to ERK5		
Immunogen	R	ecombinant full length pro	tein of human ERK5		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of ERK5 protein.			
Clonality	Р	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	vb (1/500 - 1/2000), IH (1/50) - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	N	ЛАРК7			
Alternative N	ames B	MK1; ERK5; PRKM7; Mito	en-activated protein kinase 7; MAP kinase 7; MAPK 7;		
	В	ig MAP kinase 1; BMK-1; E	xtracellular signal-regulated kinase 5; ERK-5		
Entrez Gene		5598 (Human); 23939 (Mouse)			
SwissProt	Q	213164 (Human); Q9WVS8	(Mouse); P0C865 (Rat)		
Storage/Stabi	i lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fr	reeze/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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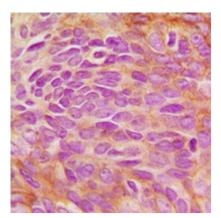
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Western blot analysis of ERK5 expression in K562 (A), Ramos (B), mouse brain (C), mouse lung (D) whole cell lysates. (Predicted band size: 50; 59; 73; 88 kD; Observed band size: 115 kD)



Immunohistochemical analysis of ERK5 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.



Immunofluorescent analysis of ERK5 staining in K562 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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