

# **Product Data Sheet**

# **Anti-ERK2** Antibody

Catalog #	Source	e Reactivity	Applications	
CQA8039	Rabbit	н, М, R	WB, IH, IF/IC, IP	
Description		Rabbit monoclonal antibod	y to ERK2	
Immunogen		A synthetic peptide of huma	an ERK2	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous lev	els of ERK2 protein.	
Clonality		Monoclonal		
Conjugation				
Form		Liquid in 50mM Tris-Glycine	e (pH 7.4), 0.15M NaCl, 50% Glycerol, 0.01% Sodium azide	
		and 0.05% BSA.		
Dilution		WB (1/500 - 1/1000), IH (1/50	) - 1/100), IF/IC (1/50 - 1/100), IP (1/10 - 1/50)	
Gene Symbol		MAPK1		
Alternative Names		ERK2; PRKM1; PRKM2; Mitogen-activated protein kinase 1; MAP kinase 1; MAPK 1;		
		ERT1; Extracellular signal-re	gulated kinase 2; ERK-2; MAP kinase isoform p42;	
		p42-MAPK; Mitogen-activat	ed protein kinase 2; MAP kinase 2; MAPK 2	
Entrez Gene		5594 (Human); 26413 (Mouse); 116590 (Rat)		
SwissProt		P28482 (Human); P63085 (I	Mouse); P63086 (Rat)	
Storage/Stabi	ility	Shipped at 4°C. Upon delive	ry 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

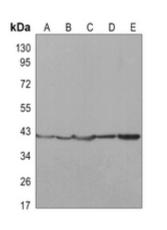
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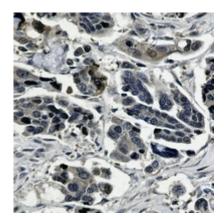


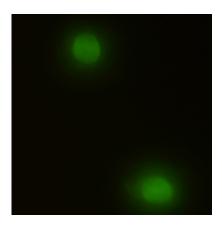
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Western blot analysis of ERK2 expression in K562 (A), rat brain (B), C6 (C), NIH3T3 (D), Hela (E) whole cell lysates. (Predicted band size: 41 kD; Observed band size: 41 kD)





Immunohistochemical analysis of ERK2 staining in human cholangiocarcinoma 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 ERK2 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark.

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