

Product Data Sheet

Anti-ICK Antibody

Catalog #	Source	e Reactivity	Applications		
CPA5327	Rabbit	Н, В	WB, IH, IF/IC		
Description		Rabbit polyclonal antibod	y to ICK		
Immunogen		KLH-conjugated synthetic	peptide encompassing a sequence within the center		
		region of human ICK. The	exact sequence is proprietary.		
Purification		The antibody was purified	by immunogen affinity chromatography.		
Specificity		Recognizes endogenous le	evels of ICK 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 - 1/2000), IH (1/	50 - 1/200), IF/IC (1/50 - 1/100)		
Gene Symbol		ІСК			
Alternative Na	ames	KIAA0936; Serine/threoni	ne-protein kinase ICK; Intestinal cell kinase; hICK;		
		Laryngeal cancer kinase 2	; LCK2; MAK-related kinase; MRK		
Entrez Gene		22858 (Human)			
SwissProt		Q9UPZ9 (Human)			
Storage/Stabi	lity	Shipped at 4°C. Upon deliv	very 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

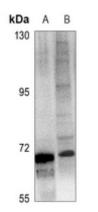
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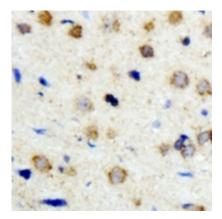
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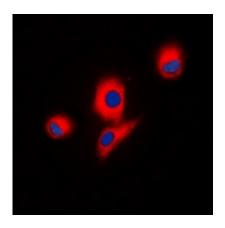
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Western blot analysis of ICK expression in U87MG (A), PC3 (B) whole cell lysates. (Predicted band size: 34; 71 kD; Observed band size: 71 kD)



Immunohistochemical analysis of ICK staining in human brain 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 ICK staining in HEK293T 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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