

Product Data Sheet

Anti-DCK Antibody

Catalog #	Source	Reactivity	Applications
CQA1079	Rabbit	H, M, R	WB, IH, IF/IC
Description		Rabbit polyclonal antibody	to DCK
Immunogen		Recombinant full length pro	otein of human DCK
Purification		The antibody was purified b	by immunogen affinity chromatography.
Specificity		Recognizes endogenous lev	els of DCK 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 - 1/200)
Gene Symbol		DCK	
Alternative Na	ames	Deoxycytidine kinase; dCK	
Entrez Gene		1633 (Human); 13178 (Mou	ise); 79127 (Rat)
SwissProt		P27707 (Human); P43346 (I	Mouse); P48769 (Rat)
Storage/Stabi	lity	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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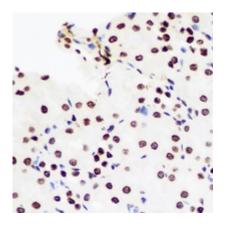
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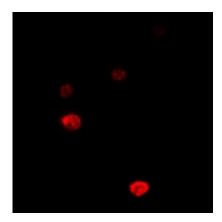
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Western blot analysis of DCK expression in Raji (A), HepG2 (B) whole cell lysates. (Predicted band size: 30 kD; Observed band size: 31 kD)



Immunohistochemical analysis of DCK staining in rat kidney 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 DCK staining in MCF7 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.

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