

# **Product Data Sheet**

## Anti-AK4 Antibody

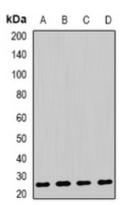
Catalog #	Source	Reactivity	Applications			
CQA1113	Rabbit	Н, М	WB, IH, IF/IC			
Description	Rab	bit polyclonal antibody	v to AK4			
Immunogen	Rec	ombinant full length pr	otein of human AK4			
Purification	The	antibody was purified	by immunogen affinity chromatography.			
Specificity	Rec	ognizes endogenous le	vels of AK4 protein.			
Clonality	Poly	yclonal				
Conjugation						
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	and	l 0.01% sodium azide.				
Dilution	WB	(1/500 - 1/2000), IH (1/5	50 - 1/200), IF/IC (1/50 - 1/200)			
Gene Symbol	AK4	Ļ				
Alternative Na	ames AK3	; AK3L1; Adenylate kin	ase 4 mitochondrial; AK 4; Adenylate kinase 3-like;			
	GTF	P:AMP phosphotransfer	rase AK4			
Entrez Gene	100	100507855, 205 (Human); 11639 (Mouse)				
SwissProt	P27	'144 (Human); Q9WUR	9 (Mouse)			
Storage/Stabi	lity Ship	oped at 4 $^\circ$ C. Upon del	ivery aliquot and store at -20 $^\circ~$ C for one year. Avoid			
	free	eze/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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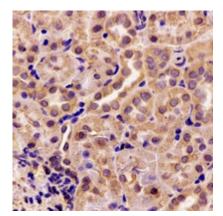




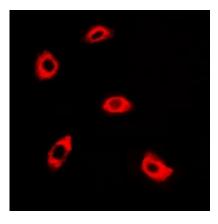
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Western blot analysis of AK4 expression in HepG2 (A), Raji (B), mouse liver (C), mouse kidney (D) whole cell lysates. (Predicted band size: 25 kD; Observed band size: 27 kD)



Immunohistochemical analysis of AK4 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 AK4 staining in A549 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  $^{\circ}$  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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