

## **Product Data Sheet**

### Anti-LIMK1/2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3256	Rabbit	H, M, R, B, C	WB, IF/IC		
Description	Rab	Rabbit polyclonal antibody to LIMK1/2			
Immunogen	KLH	I-conjugated synthetic pep	ide encompassing a sequence within the C-term		
	regi	region of human LIMK1/2. The exact sequence is proprietary.			
Purification	The	e antibody was purified by i	mmunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous levels	of LIMK1/2 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 - 1/1000), IF/IC (1/50	) - 1/200)		
Gene Symbol	LIM	IK1; LIMK2			
Alternative N	ames LIM	IK; LIM domain kinase 1; LI	ИК-1		
Entrez Gene	398	34 (Human); 16885 (Mouse	)		
SwissProt	P53	667 (Human); P53668 (Mc	use); P53669 (Rat)		
Storage/Stabi	<b>lity</b> Ship	oped at 4°C. Upon delivery	aliquot and store at -20°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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**kDa** A B 95 72 55 43 Western blot analysis of LIMK1/2 expression in Hela (A), DLD (B) whole cell lysates. (Predicted band size: 72 kD; Observed band size: 72 kD)



Immunofluorescent analysis of LIMK1/2 staining in SGC7901 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. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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