

## **Product Data Sheet**

### **Anti-MARK1** Antibody

Catalog #	Source	Reactivity	Applications
CQA3457	Rabbit	Н, М	WB, IH
Description	Rabl	oit polyclonal antibod	y to MARK1
Immunogen	Reco	ombinant full length p	rotein of human MARK1
Purification	The	antibody was purified	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous le	vels of MARK1 protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqu	id 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)
Gene Symbol	MAF	RK1	
Alternative Na	ames KIAA	1477; MARK; Serine/	hreonine-protein kinase MARK1; MAP/microtubule
	affin	ity-regulating kinase 1	l; PAR1 homolog c; Par-1c; Par1c
Entrez Gene	4139	) (Human); 226778 (N	louse)
SwissProt	Q9P	0L2 (Human); Q8VHJ5	(Mouse)
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	very aliquot and store at -20°C for one year. Avoid
	free	ze/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

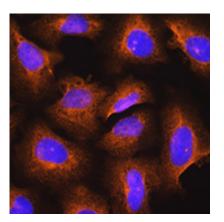
200

140

For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of MARK1 expression in Hela (A) whole cell lysates. (Predicted band size: 72; 84; 89 kD; Observed band size: 90; 72 kD)



Immunohistochemical analysis of MARK1 staining in human lung cancer 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.

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