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MARK1 Antibody



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or Research Use Only. Not for Use in Diagnostic Procedures.							
Applications: WB	Reactivity: H M R	Sensitivity: Endogenous	MW (kDa): 85	Source: Rabbit	UniProt ID: #Q9P0L2	Entrez-Gene ld: 4139	
Product Usage Information	Application			Dilution			
	We	Western Blotting			1:1000		
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at $-$ 20°C. Do not aliquot the antibody.					
Specificity / Sensitivity		MARK1 Antibody detects endogenous levels of total MARK1 protein.					
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala547 of MARK1. Antibodies were purified by protein A and peptide affinity chromatography.					
Background	pola (MA base MAF diss pho	Microtubule associated proteins regulate the stability of microtubules and control processes such as cell polarity/differentiation, neurite outgrowth, cell division and organelle trafficking (1). The MARK (MAP/microtubule affinity-regulating kinases) family (MARK1-4) of serine/threonine kinases was identified based on their ability to phosphorylate microtubule-associated proteins (MAPs) including tau, MAP2 and MAP4 (2-6). MARK proteins phosphorylate MAPs within their microtubule binding domains, causing dissociation of MAPs from microtubules and increased microtubule dynamics (2-4). In the case of tau, phosphorylation has been hypothesized to contribute to the formation of neurofibrillary tangles observed in Alzheimer's disease. Overexpression of MARK leads to hyperphosphorylation of MAPs, morphological					

related AMP-kinases within their T-loops, leading to increased activity (7).

1. Drubin, D.G. and Nelson, W.J. (1996) Cell 84, 335-44. **Background References**

2. Illenberger, S. et al. (1996) J Biol Chem 271, 10834-43.

3. Drewes, G. et al. (1995) J Biol Chem 270, 7679-88.

4. Drewes, G. et al. (1997) Cell 89, 297-308.

5. Kato, T. et al. (2001) Neoplasia 3, 4-9.

6. Trinczek, B. et al. (2004) J Biol Chem 279, 5915-23.

7. Lizcano, J.M. et al. (2004) EMBO J 23, 833-43.

Species Reactivity Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS,

0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key WB: Western Blotting

Cross-Reactivity Key H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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changes and cell death (4). The tumor suppressor kinase LKB1 phosphorylates MARK and the closely

information.

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MARK1 Antibody (#3319) Datasheet Without Images Cell Signaling Technology

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