## NME1/NDKA (G19) Antibody





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Applications: W	<b>Reactivity:</b> H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 18	Source/Isotype: Rabbit	<b>UniProt ID:</b> #P15531	<b>Entrez-Gene Id:</b> 4830
Product Usage Information		<b>Application</b> Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM soc 20°C. Do not aliquot th		), 150 mM NaCl, 100 μg/	ml BSA and 50% gly	/cerol. Store at –
Specificity/Sen	sitivity	NM23-H1/H2 (G19) Antibody detects endogenous levels of total NM23-H1/H2 protein.			۱.	
Source / Purific	cation	Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to Gly19 of human NM23-H1/H2. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		The NDK/NME/NM23 kinase family (encoded by the <i>NME</i> gene family) consists of at least eight distinct proteins that exhibit different cellular localization (1). Members of this group inhibit metastasis in a variety of tumor cell types (2). All NDK/NME/NM23 proteins possess nucleoside diphosphatase kinase (NDK) activity and catalyze the phosphorylation of nucleoside diphosphate to the corresponding nucleoside triphosphate to regulate a diverse array of cellular events (3). At least four classes of NDK biochemical activities have been described, including protein-protein interactions (4-6), regulation of GTP-binding protein function (7-9), DNA-associated activities (10,11), and histidine-dependent protein phosphotransferase activity (12). NDK/NME proteins participate in the regulation of a broad spectrum of cellular responses, including development, differentiation, proliferation, endocytosis, and apoptosis (13). Because of its role in metastasis suppression and oncogenesis, NDKA (NME1/NM23-H1) has been widely studied (14). NDKA (NM23-H1) and NDKB (NM23-H2) are encoded by adjacent <i>NME1</i> and <i>NME2</i> genes and share 90% sequence identity. Two serine residues (Ser122 and Ser144) on NDKA/NM23-H1 can be phosphorylated by AMPKα1, but only phosphorylation at Ser122 determines whether NDKA channels ATP to AMPKα1. This regulates AMPKα1 activity towards ACC1, an important regulator of fatty acid metabolism (15). Mutation of NDKB/NM23-H2 at Ser122 (S122P) in melanoma cells results in altered phosphoryl transfer activity (16).				
Background Re	eferences	<ol> <li>Lacombe, M.L. et al. (2000) <i>J Bioenerg Biomembr</i> 32, 247-58.</li> <li>Tee, Y.T. et al. (2006) <i>Taiwan J Obstet Gynecol</i> 45, 107-13.</li> <li>Ishikawa, N. et al. (2003) <i>J Bioenerg Biomembr</i> 35, 7-18.</li> <li>Paravicini, G. et al. (1996) <i>Biochem Biophys Res Commun</i> 227, 82-7.</li> <li>Reymond, A. et al. (2001) <i>Nat Med</i> 7, 350-5.</li> <li>Zhu, J. et al. (2001) <i>Proc Natl Acad Sci USA</i> 96, 14911-8.</li> <li>Otsuki, Y. et al. (2001) <i>Proc Natl Acad Sci USA</i> 98, 4385-90.</li> <li>Palacios, F. et al. (2002) <i>Nat Cell Biol</i> 4, 929-36.</li> <li>Fan, Z. et al. (2003) <i>J Bioenerg Biomembr</i> 35, 31-40.</li> <li>Wagner, P.D. and Vu, N.D. (2000) <i>Biochem J</i> 346 Pt 3, 623-30.</li> <li>Kimura, N. et al. (2004) <i>J Natl Cancer Inst</i> 96, E4.</li> <li>Crawford, R.M. et al. (2006) <i>Mol Cell Biol</i> 26, 5921-31.</li> <li>Schaertl, S. et al. (1999) <i>J Biol Chem</i> 274, 20159-64.</li> </ol>				
Species Reactiv	/ity	Species reactivity is de	termined by testing	g in at least one approve	ed application (e.g.,	western blot).
Western Blot B	uffer	IMPORTANT: For weste TBS, 0.1% Tween® 20		membrane with diluted haking, overnight.	primary antibody ir	1 5% w/v BSA, 1X
Applications Ke	ey	W: Western Blotting				

Cross-Reactivity Key	H: Human M: Mouse R: Rat Mk: Monkey	
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