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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W, IP	Reactivity: H M R Mk	Sensitivity: Endogenous	MW (kDa): 50	Source/Isotype: Rabbit	UniProt ID: #O95544	Entrez-Gene Id: 65220
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:100	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		NADK Antibody recognizes endogenous levels of total NADK protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the amino terminus of human NADK protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		NADK, also known as NAD kinase, is a cytoplasmic protein responsible for maintaining the pool of available NADP ⁺ and NADPH within the cell (1). Using ATP as a phosphate donor, NADK catalyzes the phosphorylation of NAD ⁺ to NADP ⁺ . This molecule is then reduced to NADPH and utilized in various metabolic and biosynthetic pathways (2). NADK has been suggested to play a role in glucose metabolism due to the effect NADPH production has on both the insulin secretion and survival of pancreatic β -cells (3). NADPH has a vital role in protecting cells from oxidative stress through its neutralizing effect on reactive oxygen species (ROS), which also accumulate during cell growth (2-4). Along with the p53 tumor suppression protein, NADK has been a suggested target in cancer therapy due to its link to NADPH production and its resulting protective role on growing and proliferating cells (2).				
Background References		1. Pollak, N. et al. (2007) <i>J Biol Chem</i> 282, 33562-71. 2. Tedeschi, P.M. et al. (2016) <i>Clin Cancer Res</i> 22, 5189-5195. 3. Gray, J.P. et al. (2012) <i>Am J Physiol Endocrinol Metab</i> 303, E191-9. 4. Grose, J.H. et al. (2006) <i>Proc Natl Acad Sci U S A</i> 103, 7601-6.				
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		W: Western Blotting IP: Immunoprecipitation				
Cross-Reactivity Key		H: Human M: Mouse R: Rat Mk: Monkey				
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