## Phospho-NPM1 (Thr95) Antibody





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Applications:ReactW, IP, IF-IC, FC-FPH		<b>MW (kDa):</b> 38	Source/Isotype: Rabbit	<b>UniProt ID:</b> #P06748	Entrez-Gene Id: 4869			
Product Usage Information	Immunofluorescence	••			<b>Dilution</b> 1:1000 1:50 1:50 1:50 - 1:200			
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	Phospho-NPM1 (Thr9 Thr95.	Phospho-NPM1 (Thr95) Antibody detects endogenous levels of NPM1 only when phosphorylated at Thr95.						
Species predicted to re based on 100% sequen homology	eact Mouse, Rat, Monkey Ce							
Source / Purification	corresponding to resi	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphopeptide corresponding to residues around Thr95 of human NPM1. Antibodies are purified by protein A and peptide affinity chromatography.						
Background	primarily found in nuc assembly and transpo activity, centrosome d known for its fusion w portion contributes to	Nucleophosmin (NPM; also known as B23, numatrin, or NO38) is an abundant phosphoprotein primarily found in nucleoli. It has been implicated in several distinct cellular functions, including assembly and transport of ribosomes, cytoplasmic/nuclear trafficking, regulation of DNA polymerase α activity, centrosome duplication, and molecular chaperoning activities (1,2). The <i>NPM</i> gene is also known for its fusion with the anaplastic lymphoma kinase (ALK) receptor tyrosine kinase. The NPM portion contributes to transformation by providing a dimerization domain, which results in activation of the fused kinase (3,4).						
Background Reference	2. Takemura, M. et al. 3. Morris, S.W. et al. (1	1. Okuda, M. et al. (2000) <i>Cell</i> 103, 127-140. 2. Takemura, M. et al. (1999) <i>J. Biochem. (Tokyo)</i> 125, 904-909. 3. Morris, S.W. et al. (1994) <i>Science</i> 263, 1281-1284. 4. Bischof, D. et al. (1997) <i>Mol. Cell. Biol.</i> 17, 2312-2325.						
Species Reactivity	Species reactivity is de	etermined by testing	g in at least one approve	ed 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 IF-IC: Immunofluorescence (Immunocytochemistry) FC- FP: Flow Cytometry (Fixed/Permeabilized)						
Cross-Reactivity Key	H: Human	H: Human						
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