

## CDK5 (D1F7M) Rabbit mAb



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<b>Applications:</b> W, W-S, IP, IF-F, IF- IC	Reactivity: H M R Mk	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 30	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q00535	Entrez-Gene Id: 1020
Product Usage Information		Application			Dilution	
		Western Blotting			1:1000	
		Simple Western™			1:1	0 - 1:50
		Immunoprecipitation			1:5	0
		Immunofluorescence (Frozen)			1:50	
		Immunofluorescence (Immunocytochemistry)			1:200 - 1:400	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. <i>Do not aliquot the antibody.</i>				
Specificity/Sensitivity		CDK5 (D1F7M) Rabbit mAb recognizes endogenous levels of total CDK5 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Pro253 of human CDK5 protein.				
Cyclin-dependent kinases (CDKs) are serine/threonine kinases that are activated by cyclins and eukaryotic cell cycle progression. While CDK5 shares high sequence homology with its family it is thought mainly to function in postmitotic neurons to regulate the cytoarchitecture of thes Analogous to cyclins, the regulatory subunits p35 and p39 associate with and activate CDK5 delack of sequence homology. CDK5 is ubiquitously expressed, with high levels of kinase activity primarily in the nervous system due to the narrow expression pattern of p35 and p39 in post-neurons. A large number of CDK5 substrates have been identified although no substrates have specifically attributed to p35 or p39. Substrates of CDK5 include p35, PAK1, Src, β-catenin, tau, neurofilament-H, neurofilament-M, synapsin-1, APP, DARPP32, PP1-inhibitor, and Rb. p35 is rap degraded ( $T_{1/2} < 20$ min) by the ubiquitin-proteasome pathway (1). However, p35 stability incressing CDK5 kinase activity decreases, likely as a result of decreased phosphorylation of p35 at Thr13 CDK5 (2). Proteolytic cleavage of p35 by calpain produces p25 upon neurotoxic insult, resulting prolonged activation of CDK5 by p25. Research studies have shown accumulation of p25 in neurodegenerative diseases, such as Alzheimer's disease and amyotrophic lateral sclerosis (Alzheimer's disease and amyotrophic lateral sclerosis (Alzheimer's disease and amyotrophic lateral sclerosis).						ts family members, are of these cells. The CDK5 despite the se activity detected in post-mitotic trates have been tenin, tau, p35 is rapidly polity increases as at Thr138 by the resulting in p25 in
Background References		<ol> <li>Dhavan, R. and Tsai, L.H. (2001) Nat Rev Mol Cell Biol 2, 749-59.</li> <li>Patrick, G.N. et al. (1998) J Biol Chem 273, 24057-64.</li> <li>Lee, M.S. et al. (2000) Nature 405, 360-4.</li> <li>Kusakawa, G. et al. (2000) J Biol Chem 275, 17166-72.</li> </ol>				
Species Reactiv	ity	Species reactivity is d	etermined by testin	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 W-S: Simple Western™ IP: Immunoprecipitation IF-F: Immunofluorescence (Frozen) IF-IC: Immunofluorescence (Immunocytochemistry)

**Cross-Reactivity Key** 

H: Human M: Mouse R: Rat Mk: Monkey

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