

Applications: W, IP	Reactivity: H M	Sensitivity: Endogenous	MW (kDa): 90	Source/Isotype: Rabbit	UniProt ID: #Q13627	Entrez-Gene Id: 1859	
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:50		
Storage		Supplied in 10 mM sod 20°C. Do not aliquot th	ium HEPES (pH 7.5 e antibody.), 150 mM NaCl, 100 μg/	ml BSA and 50% gly)	/cerol. Store at –	
Specificity/Sensitivity		DYRK1A Antibody detects endogenous levels of total DYRK1A protein.					
Species predicted to react based on 100% sequence homology		Rat					
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to human DYRK1A. Antibodies are purified by protein A and peptide affinity chromatography.					
Background		The DYRK family includ capable of phosphoryla identified based on hor Seven mammalian isofu DYRK4, and DYRK4B. D seen across the DYRK fa activation loop; phosph kinase activity. DYRKs to phosphorylate substrate DYRK1A phosphorylate Substrates include tran such as NFAT (7,8). DYR multiple copies of DYRK1A g and may contribute to	es several d ual-spe ating proteins at be mology to the yeas orms have been di ifferences in subst amily (4,5). All DYR norylation of the se ypically autophosp tes at Ser and Thr s serine and threo iscription factors se K1A is ubiquitously K1A exhibit learnin gene localizes to ch pathological traits	ecificity t y rosine-phosph oth Tyr and Ser/Thr resid it Yak1 (2) and the <i>Drosc</i> scovered, including DYR rate specificity, expressi K proteins have a Tyr-X- cond Tyr residue (e.g. Ty shorylate the Tyr residue residues (1,6). nine residues within a R uch as FoxO1 and cAMP y expressed in fetal and g and motor defects sug romosome 21q22.2, a re observed in chromoson	orylated and regula dues (1). The DYRK f <i>pphila</i> minibrain (mr K1A, DYRK1B, DYRK on, and subcellular Tyr motif in the cata yr312 of DYRK1A) is within their activat PX(S/T)P consensus response element- adult tissues. Trans ggesting that it is a egion implicated in ne 21 trisomy (10).	ated kinases amily was ab) kinases (3). (1C, DYRK2, DYRK3, localization are alytic domain necessary for ion loop, but sequence. binding proteins genic mice with dosage-sensitive Down syndrome,	
Background Ref	erences	1. Becker, W. and Joost, 2. Garrett, S. and Broac 3. Tejedor, F. et al. (1995 4. Kentrup, H. et al. (1995 5. Becker, W. et al. (1997 6. Lochhead, P.A. et al. (1997 6. Lochhead, P.A. et al. (1997 8. Gwack, Y. et al. (2006 9. Altafaj, X. et al. (2001 10. Guimera, J. et al. (1997)	H.G. (1999) <i>Prog.</i> h, J. (1989) <i>Genes</i> 5) <i>Neuron</i> 14, 287- 96) <i>J. Biol. Chem.</i> 2 8) <i>J. Biol. Chem.</i> 27 (2005) <i>Cell</i> 121, 929 haier, F. et al. (2003) 5) <i>Nature</i> 441, 646- 1) <i>Hum. Mol. Genet</i> 999) <i>Genomics</i> 57,	Nucleic Acid Res. Mol. Bi Dev. 3, 1336-1348. 301. 71, 3488-3495. 3, 25893-25902. 5-936.) Biochem. Biophys. Res. 650. : 10, 1915-1923. 407-418.	ol. 62, 1-17. . Commun. 300, 764	-769.	
Species Reactivi	tv	Species reactivity is det	ermined by testing	n in at least one approve	ed application (e.g.,	western blot).	
Western Blot Bu	iffer	IMPORTANT: For weste TBS, 0.1% Tween® 20 a	rn blots, incubate at 4°C with gentle s	membrane with diluted haking, overnight.	primary antibody ir	n 5% w/v BSA, 1X	
Applications Key	y	W: Western Blotting IP	: Immunoprecipita	tion			
Cross-Reactivity	' Key	H: Human M: Mouse					

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