

## 65957

## Non-phospho-STEP (Ser221) (D74H3) XP® Rabbit mAb



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

W, IP, IF-F	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 46, 61	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #P54829	Entrez-Gene Id: 84867
Product Usage Information		Application			Dilution	
		Western Blotting			1:1000	
		Immunoprecipitation			1:50	
		Immunofluorescence	(Frozen)		1:50	)
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. Do not aliquot the antibody.				
Specificity/Sensitivity		Non-phospho-STEP (Ser221) (D74H3) XP <sup>®</sup> Rabbit mAb detects endogenous levels of STEP61 protein only when dephosphorylated at Ser221 and of STEP46 protein when dephosphorylated at Ser49.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic nonphosphopeptide corresponding to residues surrounding Ser221 of human STEP61 protein.				
Background		Striatal enriched phosphatase (STEP, also known as PTPN5), is a protein tyrosine phosphatase expressed in dopaminoceptive neurons of the central nervous system (1). Alternative splicing produces the cytosolic STEP46 and the membrane-associated STEP61 isoforms of STEP. Dopamine activates D1 receptors and PKA, which in turn phosphorylate both isoforms of STEP. Phosphorylation of STEP61 occurs at Ser160 and Ser221, while STEP46 is phosphorylated at Ser49 (equivalent to Ser221 of STEP61) (2). NMDA-mediated activation of STEP is an important mechanism for regulation of Erk activity in neurons (3). Furthermore, STEP is involved in the regulation of both NMDAR and AMPAR trafficking (4,5). Due to its importance in cognitive function, STEP may play a role in Alzheimer's disease (1). Activity of STEP61 is reduced upon phosphorylation of Ser221 (Ser49 of STEP46) due to lower affinity for its substrates (2).				
		,	educed upon phosp		r49 of STEP46) due	disease (1).
Background R	eferences	,	al. (2006) <i>Trends Ne</i> <i>J Neurosci</i> 20, 5630 <i>Nat Neurosci</i> 6, 34- al. (2006) <i>Eur J Neur</i>	horylation of Ser221 (Se <i>urosci</i> 29, 452-8. -8. 42. <i>osci</i> 23, 2847-56.	r49 of STEP46) due	disease (1).
Background R		its substrates (2).  1. Braithwaite, S.P. et 2. Paul, S. et al. (2000) 3. Paul, S. et al. (2003) 4. Braithwaite, S.P. et 5. Zhang, Y. et al. (200	al. (2006) <i>Trends Ne</i> <i>J Neurosci</i> 20, 5630 <i>Nat Neurosci</i> 6, 34- al. (2006) <i>Eur J Neur</i> 8) <i>J Neurosci</i> 28, 109	horylation of Ser221 (Se <i>urosci</i> 29, 452-8. -8. 42. <i>osci</i> 23, 2847-56.	,	s disease (1). to lower affinity for
	vity	its substrates (2).  1. Braithwaite, S.P. et . 2. Paul, S. et al. (2000) 3. Paul, S. et al. (2003) 4. Braithwaite, S.P. et . 5. Zhang, Y. et al. (200	al. (2006) <i>Trends Ne J Neurosci</i> 20, 5630 <i>Nat Neurosci</i> 6, 34- al. (2006) <i>Eur J Neur</i> (8) <i>J Neurosci</i> 28, 109	horylation of Ser221 (Se urosci 29, 452-8. -8. 42. osci 23, 2847-56. 561-6.	ed application (e.g.,	s disease (1). to lower affinity for

Applications Key W: Western Blotting IP: Immunoprecipitation IF-F: Immunofluorescence (Frozen)

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

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