Phospho-Tyrosine Hydroxylase (Ser31) Antibody



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For Research Use Only. Not for Use in Diagnostic Procedures.							
Applications: WB, IP	Reactivity: R	Sensitivity: Endogenous	MW (kDa): 55-60	Source: Rabbit	UniProt ID: #P04177	Entrez-Gene Id: 25085	
Product Usage	Ар	plication			Dilution		
Information	We	Western Blotting			1:1000		
	Imr	munoprecipitation			1:50		
Storage	•	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA and 50% glycerol. Store at $-$ 20°C. Do not aliquot the antibody.					
Specificity / Sens		Phospho-Tyrosine Hydroxylase (Ser31) Antibody detects endogenous levels of tyrosine hydroxylase only when phosphorylated at Ser31.					
Species predicted react based on 10 sequence homological contracts and contracts are contracted by the contract of the contrac	0%	ise					
Source / Purificat	to th	Polyclonal antibodies are produced by immunizing animals with a synthetic phosphope to the sequence surrounding Ser31 of mouse tyrosine hydroxylase. Antibodies are puril peptide affinity chromatography.					
Background	dop regu emb	Tyrosine hydroxylase (TH) catalyzes the rate-limiting step in the synthesis of the neurotransmitter dopamine and other catecholamines. TH functions as a tetramer, with each subunit composed of a regulatory and catalytic domain, and exists in several different isoforms (1,2). This enzyme is required for embryonic development since TH knockout mice die before or at birth (3). Levels of transcription, translation and post-translational modification regulate TH activity. The amino-terminal regulatory domain					

translation and post-translational modification regulate TH activity. The amino-terminal regulatory domain contains three serine residues: Ser9, Ser31, and Ser40. Phosphorylation at Ser40 by PKA positively regulates the catalytic activity of TH (4-6). Phosphorylation at Ser31 by CDK5 also increases the catalytic

activity of TH through stabilization of TH protein levels (7-9).

1. Kumer, S.C. and Vrana, K.E. (1996) *J Neurochem* 67, 443-62. **Background References**

2. Bodeau-Péan, S. et al. (1999) J Biol Chem 274, 3469-75.

3. Kobayashi, K. et al. (1995) J Biol Chem 270, 27235-43.

4. Lew, J.Y. et al. (1999) Mol Pharmacol 55, 202-9.

5. Vié, A. et al. (1999) J Biol Chem 274, 16788-95.

6. Lindgren, N. et al. (2000) J Neurochem 74, 2470-7.

7. Moy, L.Y. and Tsai, L.H. (2004) J Biol Chem 279, 54487-93.

8. Lehmann, I.T. et al. (2006) J Biol Chem 281, 17644-51.

9. Saraf, A. et al. (2007) J Biol Chem 282, 573-80.

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 WB: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: human M: mouse R: rat Hm: hamster Mk: monkey Vir: virus Mi: mink C: chicken Dm: D. melanogaster

X: Xenopus Z: zebrafish B: bovine Dg: dog Pg: pig Sc: S. cerevisiae Ce: C. elegans Hr: horse

GP: Guinea Pig Rab: rabbit All: all species expected

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