

Phospho-Tyrosine Hydroxylase (Ser19) Ab

Cat.#: AF3112 Concn.: 1mg/ml Mol.Wt.: 60 kDa Size: 100ul,200ul Source: Rabbit Clonality: Polyclonal

Application: WB 1:500-1:2000 IHC 1:50-1:200 IF/ICC 1:100-1:500

Reactivity: Human, Mouse, Rat

Purification: The Ab is from purified rabbit serum by affinity purification

via sequential chromatography on phospho- and non-

phospho-peptide affinity columns.

Specificity: Phospho-Tyrosine Hydroxylase (Ser19) Ab detects

endogenous levels of Tyrosine Hydroxylase only when

phosphorylated at Serine 19.

Immunogen: A synthesized peptide derived from human Tyrosine

Hydroxylase around the phosphorylation site of Serine 19.

Uniprot: P07101

Description: Tyrosine hydroxylase (EC 1.14.16.2) is involved in the

conversion of phenylalanine to dopamine. As the ratelimiting enzyme in the synthesis of catecholamines, tyrosine hydroxylase has a key role in the physiology of adrenergic

neurons.

Tissue Specificity: Mainly expressed in the brain and adrenal glands.

Similarity: Belongs to the biopterin-dependent aromatic amino acid

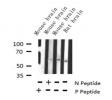
hydroxylase family.

Storage Condition and

Buffer:

Rabbit IgG in phosphate buffered saline, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol. Store at -20

°C.Stable for 12 months from date of receipt.

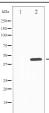


Western blot analysis of Phospho-Tyrosine Hydroxylase

(Ser19) expression in various lysates



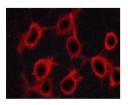
Affinity Biosciences website:www.affbiotech.com order:order@affbiotech.com



Western blot analysis of Tyrosine Hydroxylase phosphorylation expression in PC12 whole cell lysates, The lane on the left is treated with the antigen-specific peptide.



AF3112 at 1/200 staining Rat brain tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the Ab for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit Ab was used as the secondary.



AF3112 staining Hela cells by ICC/IF. Cells were fixed with PFA and permeabilized in 0.1% saponin prior to blocking in 10% serum for 45 minutes at 37°C. The primary Ab was diluted 1/400 and incubated with the sample for 1 hour at 37°C. A Alexa Fluor 594 conjugated goat polyclonal to rabbit IgG (H+L), diluted 1/600 was used as secondary Ab.

<code>IMPORTANT:</code> For western blot, incubate membrane with diluted primary Ab in 5% w/v milk , 1X TBS, 0.1% Tween®20 at 4°C with gentle shaking, overnight.

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