

BACKGROUND

The enzyme tyrosine hydroxylase (TH), also designated tyrosine 3-monooxygenase (TY3H), catalyzes the conversion of tyrosine to L-DOPA, which is the rate limiting step in the biosynthesis of catecholamines such as dopamine, adrenalin and noradrenalin. TH is thought to play a role in the pathogenesis of Parkinson's disease, which is associated with reduced dopamine levels. Two transcription factor binding sites in the proximal region of the TH gene, the TPA-responsive element (TRE) and the c-AMP responsive element (CRE), have been implicated in the complex regulation of the TH gene. TH is also known to be upregulated by the glia maturation factor (GMF), a Cdc 10/SWI6 motif-containing protein called V-1, and a variety of additional compounds.

REFERENCES

1. Stull, N.D., et al. 1996. Acidic fibroblast growth factor and catecholamines synergistically up-regulate tyrosine hydroxylase activity in developing and damaged dopamine neurons in culture. *J. Neurochem.* 67: 1519-1524.
2. Nagatsu, T., et al. 1998. Catecholamine synthesis and release. Overview *Adv. Pharmacol.* 42: 1-14.

CHROMOSOMAL LOCATION

Genetic locus: TH (human) mapping to 11p15.5; Th (mouse) mapping to 7 F5.

SOURCE

TH (H-196) is a rabbit polyclonal antibody raised against amino acids 1-196 of TH of human origin.

PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

APPLICATIONS

TH (H-196) is recommended for detection of TH of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunoprecipitation [1-2 µg per 100-500 µg of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for TH siRNA (h): sc-36662, TH siRNA (m): sc-36661, TH shRNA Plasmid (h): sc-36662-SH, TH shRNA Plasmid (m): sc-36661-SH, TH shRNA (h) Lentiviral Particles: sc-36662-V and TH shRNA (m) Lentiviral Particles: sc-36661-V.

Molecular Weight of TH: 60 kDa.

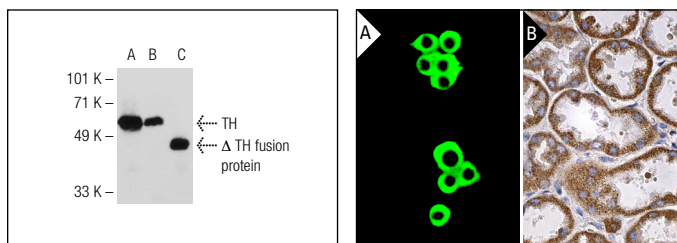
Positive Controls: mouse brain extract: sc-2253, rat adrenal gland extract: sc-364802 or PC-12 cell lysate: sc-2250.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

DATA

TH (H-196): sc-14007. Western blot analysis of TH expression in rat adrenal gland (A) and mouse brain (B) tissue extracts and truncated human recombinant TH fusion protein (C).

TH (H-196): sc-14007. Immunofluorescence staining of methanol-fixed PC-12 cells showing cytoplasmic localization (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human kidney tissue showing granular cytoplasmic staining of cells in tubules (B).

SELECT PRODUCT CITATIONS

1. Chao, H.J., et al. 2005. The conditioned enhancement of neutrophil activity is catecholamine dependent. *J. Neuroimmunol.* 158: 159-169.
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3. Costa, C., et al. 2008. Electrophysiology and pharmacology of striatal neuronal dysfunction induced by mitochondrial complex I inhibition. *J. Neurosci.* 28: 8040-8052.
4. van Roon-Mom, W.M., et al. 2008. Mutant huntingtin activates Nrf2-responsive genes and impairs dopamine synthesis in a PC12 model of Huntington's disease. *BMC Mol. Biol.* 9: 84.
5. Gauthier, M.A., et al. 2008. Covalent arylation of metallothionein by oxidized dopamine products: a possible mechanism for zinc-mediated enhancement of dopaminergic neuron survival. *Neurotox. Res.* 14: 317-328.
6. Rana, O.R., et al. 2009. Regulation of nerve growth factor in the heart: the role of the calcineurin-NFAT pathway. *J. Mol. Cell. Cardiol.* 46: 568-578.
7. Rana, O.R., et al. 2010. Mechanical stretch induces nerve sprouting in rat sympathetic neurocytes. *Auton. Neurosci.* 155: 25-32.
8. Paillé, V., et al. 2010. Distinct levels of dopamine denervation differentially alter striatal synaptic plasticity and NMDA receptor subunit composition. *J. Neurosci.* 30: 14182-14193.
9. Clewes, O., et al. 2011. Human epidermal neural crest stem cells (hEPI-NCSC)-characterization and directed differentiation into osteocytes and melanocytes. *Stem Cell Rev.* 7: 799-814.
10. Collo, G., et al. 2012. Pre-synaptic dopamine D₃ receptor mediates cocaine-induced structural plasticity in mesencephalic dopaminergic neurons via ERK and Akt pathways. *J. Neurochem.* 120: 765-778.