Best Tools for immunology Research

Tyrosine Hydroxylase Polyclonal Antibody

Catalog No :	YT6106
Reactivity :	Human;Mouse;Rat
Applications :	WB;ELISA
Target :	Tyrosine Hydroxylase
Fields :	>>Tyrosine metabolism;>>Folate biosynthesis;>>Metabolic pathways;>>Dopaminergic synapse;>>Prolactin signaling pathway;>>Parkinson disease;>>Cocaine addiction;>>Amphetamine addiction;>>Alcoholism
Gene Name :	TH TYH
Protein Name :	Tyrosine 3-monooxygenase (EC 1.14.16.2) (Tyrosine 3-hydroxylase) (TH),Tyrosine Hydrolase
Human Gene Id :	7054
Human Swiss Prot No :	P07101
Mouse Gene Id :	21823
Mouse Swiss Prot No :	P24529
Rat Gene Id :	25085
Rat Swiss Prot No :	P04177
Immunogen :	Synthesized peptide derived from human Tyrosine Hydroxylase Polyclonal
Specificity :	This antibody detects endogenous levels of Tyrosine Hydroxylase.
Formulation :	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source :	Polyclonal, Rabbit,IgG
Dilution :	WB 1:500-2000, ELISA 1:10000-20000

Purification : The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.

Concentration : 1 mg/ml

Storage Stability : -15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band : 60kD

Cell Pathway : Tyrosine metabolism;Parkinson's disease;

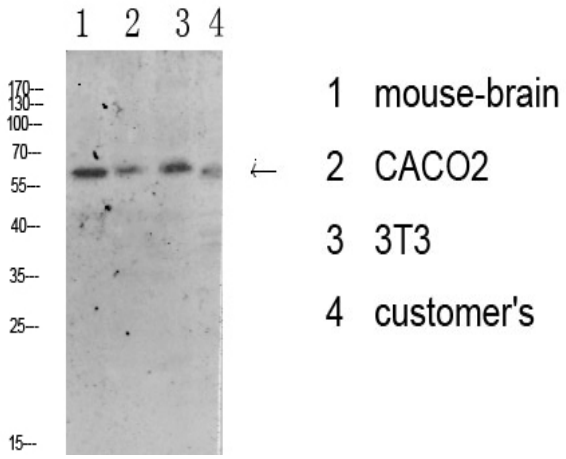
Background : The protein encoded by this gene is involved in the conversion of tyrosine to dopamine. It is the rate-limiting enzyme in the synthesis of catecholamines, hence plays a key role in the physiology of adrenergic neurons. Mutations in this gene have been associated with autosomal recessive Segawa syndrome. Alternatively spliced transcript variants encoding different isoforms have been noted for this gene. [provided by RefSeq, Jul 2008],

Function : catalytic activity:L-tyrosine + tetrahydrobiopterin + O(2) = 3,4-dihydroxy-L-phenylalanine + 4a-hydroxytetrahydrobiopterin.,cofactor:Fe(2+) ion.,disease:Defects in TH are the cause of dystonia DOPA-responsive autosomal recessive (ARDRD) [MIM:605407]; also known as autosomal recessive Segawa syndrome. ARDRD is a form of DOPA-responsive dystonia presenting in infancy or early childhood. Dystonia is defined by the presence of sustained involuntary muscle contractions, often leading to abnormal postures. Some cases of ARDRD present with parkinsonian symptoms in infancy. Unlike all other forms of dystonia, it is an eminently treatable condition, due to a favorable response to L-DOPA.,enzyme regulation:Phosphorylation leads to an increase in the catalytic activity.,function:Plays an important role in the physiology of adrenergic neurons.,online information:Tyrosine hydroxylase entry,pathway:Ca

Subcellular Location : Cytoplasm, perinuclear region . Nucleus . Cell projection, axon . Cytoplasm . Cytoplasmic vesicle, secretory vesicle, synaptic vesicle . When phosphorylated at Ser-19 shows a nuclear distribution and when phosphorylated at Ser-31 as well at Ser-40 shows a cytosolic distribution (By similarity). Expressed in dopaminergic axons and axon terminals. .

Expression : Mainly expressed in the brain and adrenal glands.

Products Images



Western blot analysis of various lysate, antibody was diluted at 1000. Secondary antibody(catalog#:RS0002) was diluted at 1:20000