

产品名称: Tyrosine Hydroxylase Rabbit Polyclonal Antibody
产品货号: APRab19473



产品概述 (Summary)

产品名称 (Production Name)	Tyrosine Hydroxylase Rabbit Polyclonal Antibody
描述 (Description)	Rabbit polyclonal Antibody
宿主 (Host)	Rabbit
应用 (Application)	WB,ELISA
种属反应性 (Reactivity)	Human,Mouse,Rat

产品性能 (Performance)

偶联物 (Conjugation)	Unconjugated
修饰 (Modification)	Unmodified
同种型 (Isotype)	IgG
克隆 (Clonality)	Polyclonal
形式 (Form)	Liquid
存放说明 (Storage)	Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.
储存溶液 (Buffer)	Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% New type preservative N.
纯化方式 (Purification)	Affinity purification

免疫原信息 (Immunogen)

基因名 (Gene Name)	TH TYH
别名 (Alternative Names)	Tyrosine 3-monooxygenase (EC 1.14.16.2) (Tyrosine 3-hydroxylase) (TH)
基因 ID (Gene ID)	7054.0
蛋白 ID (SwissProt ID)	P07101.Synthesized peptide derived from human Tyrosine Hydroxylase Polyclonal

产品应用 (Application)

稀释比 (Dilution Ratio)	WB 1:500-1:2000,ELISA 1:10000-1:20000
蛋白分子量 (Molecular Weight)	60kDa

研究背景 (Background)

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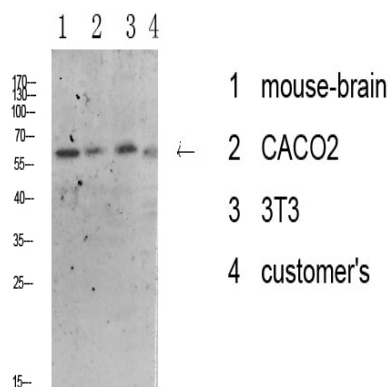


The protein encoded by this gene is involved in the conversion of tyrosine to dopamine. It is the rate-limiting enzyme in the synthesis of catecholamines, hence plays a key role in the physiology of adrenergic neurons. Mutations in this gene have been associated with autosomal recessive Segawa syndrome. Alternatively spliced transcript variants encoding different isoforms have been noted for this gene. [provided by RefSeq, Jul 2008], 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: Catecholamine biosynthesis; dopamine biosynthesis; dopamine from L-tyrosine: step 1/2., similarity: Belongs to the biopterin-dependent aromatic amino acid hydroxylase family., tissue specificity: Mainly expressed in the brain and adrenal glands.,

研究领域 (Research Area)

Tyrosine metabolism; Parkinson's disease;

图片 (Image Data)



Western blot analysis of various lysate, antibody was diluted at 1000. Secondary antibody was diluted at 1:20000

注意事项 (Note)

For research use only .