## Product Name: Choactase Rabbit Polyclonal Antibody Catalog #: APRab08771



### **Summary**

Production Name Choactase Rabbit Polyclonal Antibody

**Description** Rabbit Polyclonal Antibody

Host Rabbit
Application WB,ELISA

**Reactivity** Human, Mouse, Rat

### **Performance**

ConjugationUnconjugatedModificationUnmodified

**Isotype** IgG

ClonalityPolyclonalFormLiquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw

cycles.

**Buffer** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.

**Purification** Affinity purification

### **Immunogen**

Storage

Gene Name CHAT

Alternative Names CHAT; Choline O-acetyltransferase; CHOACTase; ChAT; Choline acetylase

**Gene ID** 1103.0

P28329. The antiserum was produced against synthesized peptide derived from human

Choactase. AA range:334-383

### **Application**

**SwissProt ID** 

**Dilution Ratio** WB 1:500 - 1:2000. ELISA: 1:10000

Molecular Weight 82,70kD

### **Background**

## Product Name: Choactase Rabbit Polyclonal Antibody Catalog #: APRab08771

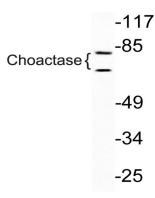


This gene encodes an enzyme which catalyzes the biosynthesis of the neurotransmitter acetylcholine. This gene product is a characteristic feature of cholinergic neurons, and changes in these neurons may explain some of the symptoms of Alzheimer's disease. Polymorphisms in this gene have been associated with Alzheimer's disease and mild cognitive impairment. Mutations in this gene are associated with congenital myasthenic syndrome associated with episodic apnea. Multiple transcript variants encoding different isoforms have been found for this gene, and some of these variants have been shown to encode more than one isoform. [provided by RefSeq, May 2010],catalytic activity:Acetyl-CoA + choline = CoA + O-acetylcholine, disease:Defects in CHAT are the cause of congenital myasthenic syndrome with episodic apnea (CMSEA) [MIM:254210]; formerly known as familial infantile myasthenia gravis 2 (FIMG2). CMSEA is an autosomal recessive congenital myasthenic syndrome. Patients have myasthenic symptoms since birth or early infancy, negative tests for anti-AChR antibodies, and abrupt episodic crises with increased weakness, bulbar paralysis, and apnea precipitated by undue exertion, fever, or excitement, function:Catalyzes the reversible synthesis of acetylcholine (ACh) from acetyl CoA and choline at cholinergic synapses, online information:Choline acetyltransferase entry, similarity:Belongs to the carnitine/choline acetyltransferase family.

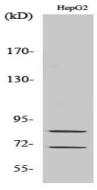
#### **Research Area**

Glycerophospholipid metabolism;

### **Image Data**



Western blot analysis of lysate from HepG2 cells, using Choactase antibody.



# Product Name: Choactase Rabbit Polyclonal Antibody Catalog #: APRab08771



Western Blot analysis of various cells using Choactase Polyclonal Antibody diluted at 1: 1000

### Note

For research use only.

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838