

# ChAT Recombinant Rabbit mAb

ChAT Recombinant Rabbit mAb Catalog # AP94815

#### **Product Information**

**Application** WB, IHC-P, IHC-F, IF

Host Rabbit Clonality Recombinant **Calculated MW** 82 KDa **Physical State** Liquid

KLH conjugated synthetic peptide derived from human CHAT **Immunogen** 

Isotype

**Purity** affinity purified by Protein A

**Buffer** 0.01M TBS(pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.

SUBCELLULAR LOCATION Cytosol; Nucleus

**SIMILARITY** Belongs to the carnitine/choline acetyltransferase family.

**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.

This product as supplied is intended for research use only, not for use in **Important Note** 

human, therapeutic or diagnostic applications.

**Background Descriptions** 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].

### **Additional Information**

**Dilution** WB=1:500-2000,IHC-P=1:100-500,IHC-F=1:100-500,IF=1:100-500

**Format** 0.01M TBS(pH7.4) with 1% BSA, 0.09% (W/V) sodium azide and 50% Glyce

**Storage** Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. When

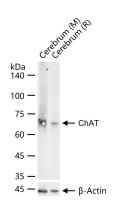
reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody

is stable for at least two weeks at 2-4 °C.

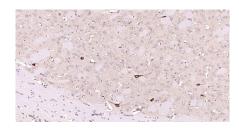
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# **Images**



25 ug total protein per lane of various lysates (see on figure) probed with ChAT monoclonal antibody, unconjugated (AP94815) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



Paraformaldehyde-fixed, paraffin embedded Mouse Cerebrum; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with ChAT Monoclonal Antibody, Unconjugated(AP94815) at 1:200 overnight at 4°C, followed by conjugation to the SP Kit (Rabbit, SP-0023) and DAB (C-0010) staining.



Paraformaldehyde-fixed, paraffin embedded Rat Cerebrum; Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15 min; Antibody incubation with ChAT Monoclonal Antibody, Unconjugated(AP94815) at 1:200 overnight at 4°C, followed by conjugation to the SP Kit (Rabbit, SP-0023) and DAB (C-0010) staining.

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