BOSTER BIOLOGICAL TECHNOLOGY Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

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antibody and FLISA

Basic Information		
Product Name	Anti-CHAT Antibody	
Gene Name	СНАТ	
Source	Rabbit	
Clonality	Polyclonal	
lsotype	lgG	
Species Reactivity	mouse, rat	
Tested Application	WB, IHC, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	E.coli-derived rat Choline Acetyltransferase/Chat recombinant protein (Position: E19-D612).	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	83 kDa	
Dilution Ratios	Western blot (WB): Immunohistochemistry (IHC): ELISA: (Boiling the paraffin sections in 10mM citrate buffer, mins is required for the staining of formalin/paraffin s determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied.

Background Information

Choline acetyltransferase (commonly abbreviated as°ChAT,but sometimes°CAT) is a transferase enzyme responsible for the synthesis of the neurotransmitter acetylcholine. In humans,the choline acetyltransferase enzyme is encoded by the°CHAT gene. 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.

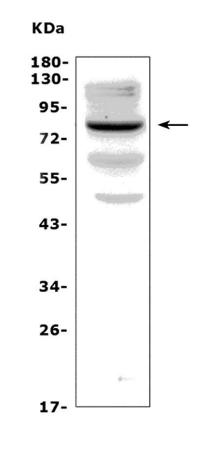
antibody and ELISA experts BOSTER BIOLOGICAL TECHNOLOGY Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

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Reference

Anti-CHAT Antibody 被引用在14文献中。

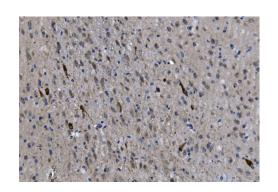
Selected Validation Data



Western blot analysis of CHAT using anti-CHAT antibody (A01192-5). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: rat brain tissue lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-CHAT antigen affinity purified polyclonal antibody (A01192-5) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for CHAT at approximately 83 kDa. The expected band size for CHAT is at 83 kDa.



IHC analysis of CHAT using anti-CHAT antibody (A01192-5). CHAT was detected in a paraffin-embedded section of mouse brain tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-CHAT Antibody (A01192-5) at a dilution of 1:200 and developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.

Antibody | ELISA Kits | IHC | WB | Cell Culture



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