## Synapsin I (phospho Ser62) Polyclonal Antibody

Catalog No: #13499

Package Size: #13499-1 50ul #13499-2 100ul



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## Description

| Decemption            |  |
|-----------------------|--|
| Product Name          | Synapsin I (phospho Ser62) Polyclonal Antibody   |
| Host Species          | Rabbit   |
| Purification          | The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific |
|                       | immunogen.   |
| Applications          | WB,IHC-p,IF(paraffin section),ELISA  |
| Species Reactivity    | Human,Mouse,Rat  |
| Specificity           | Phospho-Synapsin I (S62) Polyclonal Antibody detects endogenous levels of Synapsin I protein only when     |
|                       | phosphorylated at S62.   |
| Immunogen Description | The antiserum was produced against synthesized peptide derived from human Synapsin1 around the             |
|                       | phosphorylation site of Ser62. AA range:26-75  |
| Other Names           | SYN1; Synapsin-1; Brain protein 4.1; Synapsin I  |
| Accession No.         | Swiss Prot:P17600GeneID:6853   |
| Uniprot               | P17600   |
| GenelD                | 6853   |
| SDS-PAGE MW           | 80   |
| Concentration         | 1 mg/ml  |
| Formulation           | Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.                                    |
| Storage               | -20°C/1  |
|                       |  |

## **Application Details**

Western Blot: 1/500 - 1/2000. Immunohistochemistry: 1/100 - 1/300. ELISA: 1/10000. Not yet tested in other applications.

## Background

synapsin I(SYN1) Homo sapiens This gene is a member of the synapsin gene family. Synapsins encode neuronal phosphoproteins which associate with the cytoplasmic surface of synaptic vesicles. Family members are characterized by common protein domains, and they are implicated in synaptogenesis and the modulation of neurotransmitter release, suggesting a potential role in several neuropsychiatric diseases. This member of the synapsin family plays a role in regulation of axonogenesis and synaptogenesis. The protein encoded serves as a substrate for several different protein kinases and phosphorylation may function in the regulation of this protein in the nerve terminal. Mutations in this gene may be associated with X-linked disorders with primary neuronal degeneration such as Rett syndrome. Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq, Jul 2008],

Note: This product is for in vitro research use only