abcam

Product datasheet

Recombinant Human Alpha-synuclein protein ab51189

6 References 2 Images

Description

Product name Recombinant Human Alpha-synuclein protein

Purity > 95 % SDS-PAGE.

Endotoxin level < 1.000 Eu/μg
Expression system Escherichia coli

Accession P37840

Protein length Full length protein

Animal free No

Nature Recombinant

Species Human

Sequence MDVFMKGLSKAKEGVVAAAEKTKQGVAEAAGKTKEGVL

YVGSKTKEGVVH

GVATVAEKTKEQVTNVGGAVVTGVTAVAQKTVEGAGSIA

AATGFVKKDQL

GKNEEGAPQEGILEDMPVDPDNEAYEMPSEEGYQDYEP

EΑ

Predicted molecular weight 14 kDa

Amino acids 1 to 140

Specifications

Our Abpromise guarantee covers the use of ab51189 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Applications Western blot

SDS-PAGE

Form Liquid

Preparation and Storage

Stability and Storage Shipped on Dry Ice. Upon delivery aliquot. Store at -80°C. Avoid freeze / thaw cycle.

pH: 7.50

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General Info

Function May be involved in the regulation of dopamine release and transport. Induces fibrillization of

microtubule-associated protein tau. Reduces neuronal responsiveness to various apoptotic

stimuli, leading to a decreased caspase-3 activation.

Tissue specificity Expressed principally in brain but is also expressed in low concentrations in all tissues examined

except in liver. Concentrated in presynaptic nerve terminals.

Involvement in disease Genetic alterations of SNCA resulting in aberrant polymerization into fibrils, are associated with

several neurodegenerative diseases (synucleinopathies). SNCA fibrillar aggregates represent the major non A-beta component of Alzheimer disease amyloid plaque, and a major component of Lewy body inclusions. They are also found within Lewy body (LB)-like intraneuronal inclusions, glial inclusions and axonal spheroids in neurodegeneration with brain iron accumulation type 1.

Parkinson disease 1 Parkinson disease 4 Dementia Lewy body

Sequence similarities Belongs to the synuclein family.

Domain The 'non A-beta component of Alzheimer disease amyloid plaque' domain (NAC domain) is

involved in fibrils formation. The middle hydrophobic region forms the core of the filaments. The C-

terminus may regulate aggregation and determine the diameter of the filaments.

Post-translational modifications

Phosphorylated, predominantly on serine residues. Phosphorylation by CK1 appears to occur on residues distinct from the residue phosphorylated by other kinases. Phosphorylation of Ser-129 is selective and extensive in synucleinopathy lesions. In vitro, phosphorylation at Ser-129 promoted

insoluble fibril formation. Phosphorylated on Tyr-125 by a PTK2B-dependent pathway upon

osmotic stress.

Hallmark lesions of neurodegenerative synucleinopathies contain alpha-synuclein that is modified by nitration of tyrosine residues and possibly by dityrosine cross-linking to generated stable

oligomers.

Ubiquitinated. The predominant conjugate is the diubiquitinated form.

Acetylation at Met-1 seems to be important for proper folding and native oligomeric structure.

Cellular localization Cytoplasm, cytosol. Membrane. Nucleus. Cell junction, synapse. Secreted. Membrane-bound in

dopaminergic neurons.

Images



Western blot - Recombinant Human Alpha-synuclein protein (ab51189)



Anti-Alpha-synuclein antibody [LB 509] ($\underline{ab27766}$) at 1/1000 dilution + Recombinant Human Alpha-synuclein protein (ab51189) at 0.1 μg

Secondary

Goat Anti-Mouse IgG H&L (HRP) preadsorbed (ab97040) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 8 minutes

ab51189 (3 µg) in 14% SDS-PAGE.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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