

## Produktinformation



Forschungsprodukte & Biochemikalien



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Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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## Lieferung & Zahlungsart

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## Anti-Alpha Synuclein Antibody (Aggregate-Specific) [2F11]



Mouse Anti-Mouse Alpha Synuclein Aggregate Antibody Monoclonal IgG1 Catalog No. SMC-617

# **Product Name** Alpha Synuclein Antibody (Aggregate-Specific) Description Mouse Anti-Mouse Alpha Synuclein Aggregate Antibody Monoclonal IgG1 **Species Reactivity** Human, Mouse **Applications** WB, DB, IP, IHC, ICC-IF, ImmunoGold labelling **Antibody Dilution** WB (1:1000). Application specific; optimal dilutions for assays should be determined by the user. **Host Species** Mouse **Immunogen Species** Mouse **Immunogen** Mouse alpha synuclein fibrils Concentration 1mg/mL **Conjugates** Alkaline Phosphatase, APC, ATTO 390, ATTO 488, ATTO 594, Biotin, FITC, HRP, PerCP, RPE, Unconjugated Field Of Use

Not for use in humans. Not for use in diagnostics or therapeutics. For in vitro research use only.

## **Properties**

#### **Storage Buffer**

PBS pH 7.4, 50% glycerol, 0.09% sodium azide. \*Storage buffer may change when conjugated

#### **Storage Temperature**

-20°C

#### **Shipping Temperature**

Blue Ice or 4°C

#### **Purification**

Protein G Purified

#### **Clonality**

Monoclonal

#### **Clone Number**

2F11

#### Isotype

IgG1

#### **Specificity**

Binds a 3-dimensional fibrillar epitope including AA101-120, AA61-80

#### **Cite This Product**

Mouse Anti-Mouse Alpha Synuclein Aggregate Antibody Monoclonal (StressMarq Biosciences, Victoria BC, Cat# SMC-617)

#### **Certificate Of Analysis**

A 1:1000 dilution of SMC-617 was sufficient for detection of Alpha Synuclein aggregates in 15  $\mu$ g of human brain cell lysate by ECL immunoblot analysis using goat anti-mouse IgG:HRP as the secondary antibody.

## **Biological Description**

#### **Alternative Names**

Alpha Synuclein Antibody, Non-A beta component of AD amyloid Antibody, Non-A4 component of amyloid precursor Antibody, NACP Antibody, SNCA Antibody, PARK1 Antibody, PARK1 Antibody, alphaSYN Antibody,

PARK 4 antibody, PARK4 antibody, Parkinson disease familial 1 antibody, Parkinson disease (autosomal dominant, Lewy body) 4 antibody, SYN antibody

#### **Research Areas**

Alzheimer's Disease, Neurodegeneration, Neuroscience, Parkinson's Disease, Synuclein, Tangles & Tau, Multiple System Atrophy

#### **Cellular Localization**

Cell Junction, Cytoplasm, Cytosol, Membrane, Nucleus, Synapse

#### **Accession Number**

NP 001035916.1

#### **Gene ID**

20617

#### **Swiss Prot**

055042

#### **Scientific Background**

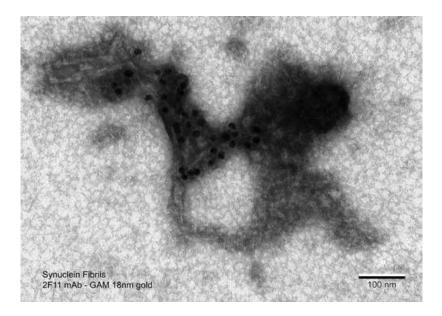
Alpha-Synuclein (SNCA) is expressed predominantly in the brain, where it is concentrated in presynaptic nerve terminals (1). Alpha-synuclein is highly expressed in the mitochondria of the olfactory bulb, hippocampus, striatum and thalamus (2). Functionally, it has been shown to significantly interact with tubulin (3), and may serve as a potential microtubule-associated protein. It has also been found to be essential for normal development of the cognitive functions; inactivation may lead to impaired spatial learning and working memory (4). SNCA fibrillar aggregates represent the major non A-beta component of Alzheimer's disease amyloid plaque, and a major component of Lewy body inclusions, and Parkinson's disease. Parkinson's disease (PD) is a common neurodegenerative disorder characterized by the progressive accumulation in selected neurons of protein inclusions containing alpha-synuclein and ubiquitin (5, 6). Covered by US patent family US11,098,108B2 (7).

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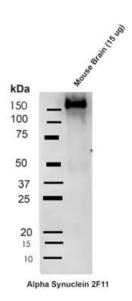
#### References

- 1. "Genetics Home Reference: SNCA". US National Library of Medicine. (2013).
- 2. Zhang L., et al. (2008) Brain Res. 1244: 40-52.
- 3. Alim M.A., et al. (2002) J Biol Chem. 277(3): 2112-2117.
- 4. Kokhan V.S., Afanasyeva M.A., Van'kin G. (2012) Behav. Brain. Res. 231(1): 226-230.
- 5. Spillantini M.G., et al. (1997) Nature. 388(6645): 839-840.
- 6. Mezey E., et al. (1998) Nat Med. 4(7): 755-757.
- 7. Louwrier, A. US Patent family US11,098,108 B2

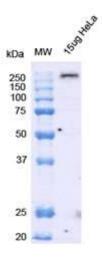
#### **Product Images**



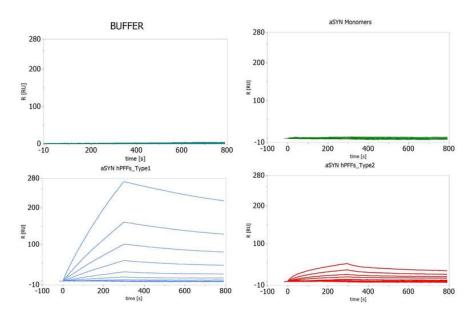
Immuno-gold electron micrograph of sonicated  $\alpha$ -Syn Fibrils (SPR-322) stained with 2F11 antibody (1:400 dilution) and visualized using goat anti-mouse IgG conjugated with 18 nm gold particles. The sample was then placed on a transmission electron microscopy metallic grid and stained with uranyl acetate.



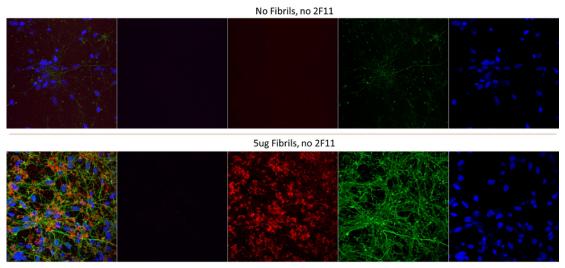
Western blot on 15ug mouse brain lysate from Fmr1 KO mice (FXS mice) with C57Bl/6J genetic background. Block: 5% Skim Milk in 1X TBST. Primary Antibody: Mouse Anti-Alpha Synuclein Monoclonal Antibody clone 2F11 at 1:1000 for 2 hours at RT. Secondary Antibody: Goat Anti-Mouse HRP:lgG at 1:3000 for 1 hour at RT. Color Development: ECL solution (Super Signal West Pico) for 5 min in RT. Band clearly identifies large molecular weight oligomeric band (monomeric band predicted at 14kD).



Western blot on 15ug Hela lysate. Block: 5% Skim Milk in 1X TBST. Primary Antibody: Mouse Anti-Alpha Synuclein Monoclonal Antibody clone 2F11 at 1:1000 for 2 hours at RT. Secondary Antibody: Goat Anti-Mouse HRP:lgG at 1:3000 for 1 hour at RT. Color Development: ECL solution (Super Signal West Pico) for 5 min in RT. Band clearly identifies large molecular weight oligomeric band (monomeric band predicted at 14kD).

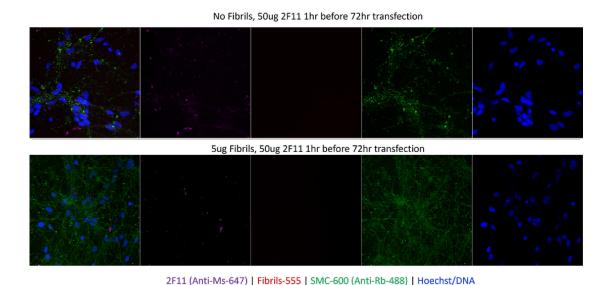


SPR binding analysis of 2F11 antibody and human alpha synuclein type 1 fibrils (SPR-322), type 2 (SPR-317) fibrils and monomeric protein (SPR-321). Buffer (top left), alpha synuclein monomers (top right), alpha synuclein fibrils type1 (bottom left), and alpha synuclein fibrils type 2 (bottom right). Data: KD[nM] for SPR-322 = 42.5, SPR-317 = 365.3; 2F11 binds SPR-322 with 8.6 x greater affinity than SPR-317. 2F11 does not bind monomeric alpha synuclein (SPR-321).



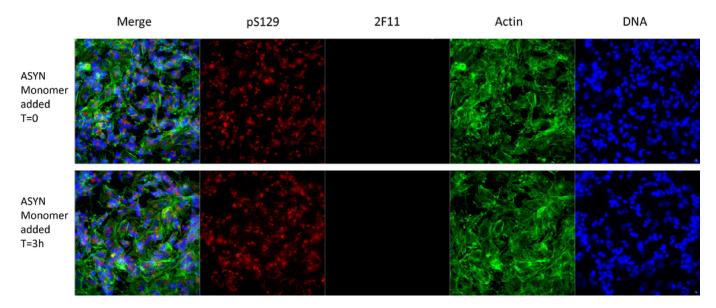
2F11 (Anti-Ms-647) | Fibrils-555 | SMC-600 (Anti-Rb-488) | Hoechst/DNA

8 Day differentiated iPSCs with added alpha synuclein fibrils (SPR-322), conjugated to Alexa Fluor 555 with or without 1h prior 2F11 addition. SPR-322 Alexa Fluor 555 added at 5µg/well for 48h, 2F11 (50µg/well added 1h before fibrils where applicable). Cells were subsequently washed 48h after addition and imaged a following 24h later. Imaging and channels: 2F11 antibody: 647nm (using an anti-mouse-647 conjugate); Fibrils: 555nm, phosph-ser129 (SPR-600) 488nm, and Hoechst stain for DNA. Image 4a top, no fibrils, no 2F11 added. Bottom, fibrils added, no 2F11 antibody. Fibrils are clearly visible in the cells as is enhanced phosph-ser129 signal. Image 4b top, no fibrils, 2F11 antibody added. Bottom, fibrils and 2F11 antibody added. The image clearly shows 2F11 antibody addition stops cell entry of the 555-labelled fibrils.



8 Day differentiated iPSCs with added alpha synuclein fibrils (SPR-322), conjugated to Alexa Fluor 555 with or without 1h prior 2F11 addition. SPR-322 Alexa Fluor 555 added at 5µg/well for 48h, 2F11 (50µg/well added 1h before fibrils where applicable). Cells were subsequently washed 48h after addition and imaged a following 24h later. Imaging and channels: 2F11 antibody: 647nm (using an anti-mouse-647 conjugate); Fibrils: 555nm, phosph-ser129 (SPR-600) 488nm, and Hoechst stain for DNA. Image 4a top, no fibrils, no 2F11 added. Bottom, fibrils added, no 2F11 antibody. Fibrils are clearly visible in the cells as is enhanced phosph-ser129 signal. Image 4b top, no fibrils, 2F11 antibody added. Bottom, fibrils and 2F11 antibody

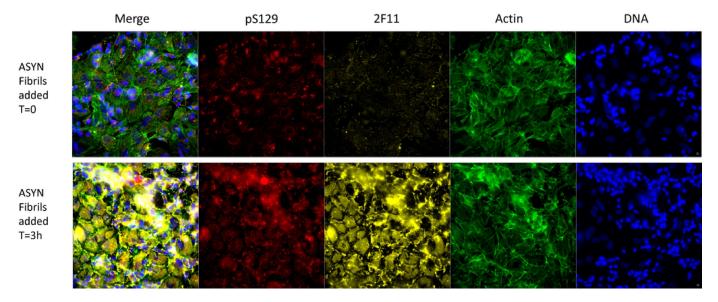
added. The image clearly shows 2F11 antibody addition stops cell entry of the 555-labelled fibrils.



8 Day differentiated SH-SY5Y cells with added alpha synuclein monomers (SPR-321; 5µg) or fibrils (SPR-322; 5µg) at time zero and 3h time points. Staining and channels phospho-ser129 (ab51253) and secondary DaRb-647; 2F11 (1:1000 dilution, DaMs-555); Actin stain (Phalloidin dye: 488nm); DNA (Hoechst dye, 405nm). Image 5a, top T=0, bottom T=3h, no signal change from 2F11 showing that alpha synuclein monomers do not bind the antibody. Image 5b, top t=0 and bottom, t=3h. 2F11 channel clearly shows a large increase in fibril signal entering the cells by 3h, with a concomitant increase in phspho-Ser129 signal.

Mouse Ileum 40x	Mouse Ascending Colon 40x	Mouse Transcending Colon 40x	Mouse Descending Colon 40x
Mouse Ileum 40x	Mouse Ascending Colon 40x	Mouse Transcending Colon 40x	Mouse Descending Colon 40x

Mouse gut samples stained with 2F11 at 1:1000 dilution with overnight incubation at 4°C in PBS with 0.1% BSA, Secondary biotinylated anti-mouse antibody for 1 hour at RT also in 0.1% BSA PBS. VECTASTAIN ABC reagent kit was added after secondary incubation. Peroxidase HRP substrate was left on the gut tissue for 6 minutes. H&E staining of all tissue was performed at 1:1 with dH20, rinsed, then stained with blueing solution prior to dehydration. Imaging at 40 x magnification. Mouse tissue used from a mouse Parkinson's disease (PD) seeding model based on C57BLJ/6 mice with brain injections of Human PD patient alphasynuclein fractions. Mouse gut sections were used at a thickness of 3µm. Top panel shows (in biological order of mouse gut from ileum to descending colon) control secondary staining. Bottom panel shows gut samples from mice injected with human PD patient brain-derived alpha synuclein, also in biological order (ileum to descending colon). There is clear aggregate staining visible in the lower panel.



8 Day differentiated SH-SY5Y cells with added alpha synuclein monomers (SPR-321; 5µg) or fibrils (SPR-322; 5µg) at time zero and 3h time points. Staining and channels phospho-ser129 (ab51253) and secondary DaRb-647; 2F11 (1:1000 dilution, DaMs-555); Actin stain (Phalloidin dye: 488nm); DNA (Hoechst dye, 405nm). Image 5a, top T=0, bottom T=3h, no signal change from 2F11 showing that alpha synuclein monomers do not bind the antibody. Image 5b, top t=0 and bottom, t=3h. 2F11 channel clearly shows a large increase in fibril signal entering the cells by 3h, with a concomitant increase in phspho-Ser129 signal.

## **Product Citations (0)**

Currently there are no citations for this product.

#### Reviews

There are no reviews yet.