

Resource Summary Report

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SynGAP Polyclonal Antibody

RRID:AB_2287112

Type: Antibody

Proper Citation

(Thermo Fisher Scientific Cat# PA1-046, RRID:AB_2287112)

Antibody Information

URL: http://antibodyregistry.org/AB_2287112

Proper Citation: (Thermo Fisher Scientific Cat# PA1-046, RRID:AB_2287112)

Target Antigen: SynGAP

Host Organism: rabbit

Clonality: polyclonal

Comments: Applications: WB (1 µg/mL), ICC/IF (2 µg/mL), IHC (P) (2 µg/mL)

Antibody Name: SynGAP Polyclonal Antibody

Description: This polyclonal targets SynGAP

Target Organism: rat, mouse, human

Defining Citation: [PMID:27194588](#), [PMID:17904250](#), [PMID:12040032](#), [PMID:25158904](#), [PMID:27623146](#), [PMID:20410461](#), [PMID:26041915](#), [PMID:16041714](#), [PMID:9620694](#), [PMID:14970204](#), [PMID:20554866](#), [PMID:25533468](#), [PMID:22072671](#), [PMID:14715953](#), [PMID:23268962](#), [PMID:9581761](#)

Antibody ID: AB_2287112

Vendor: Thermo Fisher Scientific

Catalog Number: PA1-046

Record Creation Time: 20250416T091819+0000

Record Last Update: 20250416T094241+0000

Ratings and Alerts

No rating or validation information has been found for SynGAP Polyclonal Antibody.

No alerts have been found for SynGAP Polyclonal Antibody.

Data and Source Information

Source: [Antibody Registry](#)

Usage and Citation Metrics

We found 11 mentions in open access literature.

Listed below are recent publications. The full list is available at [FDI Lab - SciCrunch.org](#).

Robinson K, et al. (2024) Mapping proteomic composition of excitatory postsynaptic sites in the cerebellar cortex. *Frontiers in molecular neuroscience*, 17, 1381534.

Daniel JA, et al. (2023) An intellectual-disability-associated mutation of the transcriptional regulator NACC1 impairs glutamatergic neurotransmission. *Frontiers in molecular neuroscience*, 16, 1115880.

Yang R, et al. (2023) Upregulation of SYNGAP1 expression in mice and human neurons by redirecting alternative splicing. *Neuron*, 111(10), 1637.

Yokoi S, et al. (2022) The SYNGAP1 3'UTR Variant in ALS Patients Causes Aberrant SYNGAP1 Splicing and Dendritic Spine Loss by Recruiting HNRNPK. *The Journal of neuroscience : the official journal of the Society for Neuroscience*, 42(47), 8881.

Murtaza N, et al. (2022) Neuron-specific protein network mapping of autism risk genes identifies shared biological mechanisms and disease-relevant pathologies. *Cell reports*, 41(8), 111678.

Gou G, et al. (2020) SynGAP splice variants display heterogeneous spatio-temporal expression and subcellular distribution in the developing mammalian brain. *Journal of neurochemistry*, 154(6), 618.

McEachern EP, et al. (2020) PSD-95 deficiency alters GABAergic inhibition in the prefrontal cortex. *Neuropharmacology*, 179, 108277.

Muheim CM, et al. (2019) Dynamic- and Frequency-Specific Regulation of Sleep Oscillations by Cortical Potassium Channels. *Current biology : CB*, 29(18), 2983.

Dejanovic B, et al. (2018) Changes in the Synaptic Proteome in Tauopathy and Rescue of Tau-Induced Synapse Loss by C1q Antibodies. *Neuron*, 100(6), 1322.

Walkup WG, et al. (2016) A model for regulation by SynGAP- γ 1 of binding of synaptic proteins to PDZ-domain 'Slots' in the postsynaptic density. *eLife*, 5.

Walkup WG, et al. (2015) Phosphorylation of synaptic GTPase-activating protein (synGAP) by Ca^{2+} /calmodulin-dependent protein kinase II (CaMKII) and cyclin-dependent kinase 5 (CDK5) alters the ratio of its GAP activity toward Ras and Rap GTPases. *The Journal of biological chemistry*, 290(8), 4908.