EAAT1 (A-3): sc-515839



The Power to Question

BACKGROUND

Excitatory amino acid transporter 1 (EAAT1) is one of the two glial glutamate transporters that clear the extracellular glutamate generated during neuronal signal transmission. Excitatory amino acid transporters (EAATs) are membrane-bound proteins that are localized in glial cells and pre-synaptic glutamatergic nerve endings. EAATs transport the excitatory neurotransmitters L-glutamate and D-aspartate, a process that is essential for terminating the postsynaptic acction of glutamate. The reuptake of amino acid neurotransmitters by EAAT proteins has been shown to protect neurons from excitotoxicity, which is caused by the accumulation of amino acid neurotransmitters. Three glutamate transporters have been identified in human brain, designated EAAT1–3. EAAT1 and EAAT3 are also expressed in various non-nervous tissues, while EAAT2 expression appears to be restricted to the brain. Surface expression of the glial glutamate transporter EAAT1 is stimulated by Insulin-like growth factor 1 through activation of phosphatidylinositol-3-kinase.

CHROMOSOMAL LOCATION

Genetic locus: SLC1A3 (human) mapping to 5p13.2; Slc1a3 (mouse) mapping to 15 A1.

SOURCE

EAAT1 (A-3) is a mouse monoclonal antibody raised against amino acids 1-50 mapping near the N-terminus of EAAT1 of human origin.

PRODUCT

Each vial contains 200 μg lgG_{2a} kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

EAAT1 (A-3) is available conjugated to agarose (sc-515839 AC), 500 μg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-515839 HRP), 200 μg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-515839 PE), fluorescein (sc-515839 FITC), Alexa Fluor $^{\circ}$ 488 (sc-515839 AF488), Alexa Fluor $^{\circ}$ 546 (sc-515839 AF546), Alexa Fluor $^{\circ}$ 594 (sc-515839 AF594) or Alexa Fluor $^{\circ}$ 647 (sc-515839 AF647), 200 μg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor $^{\circ}$ 680 (sc-515839 AF680) or Alexa Fluor $^{\circ}$ 790 (sc-515839 AF790), 200 μg/ml, for Near-Infrared (NIR) WB, IF and FCM.

APPLICATIONS

EAAT1 (A-3) is recommended for detection of EAAT1 of mouse, rat and human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2 μ g per 100-500 μ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for EAAT1 siRNA (h): sc-35253, EAAT1 siRNA (m): sc-35254, EAAT1 shRNA Plasmid (h): sc-35253-SH, EAAT1 shRNA Plasmid (m): sc-35254-SH, EAAT1 shRNA (h) Lentiviral Particles: sc-35253-V and EAAT1 shRNA (m) Lentiviral Particles: sc-35254-V.

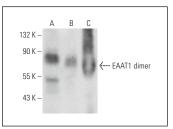
Molecular Weight of EAAT1: 65 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200, IMR-32 cell lysate: sc-2409 or human cerebellum extract: sc-516706.

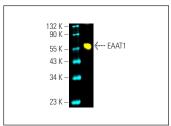
STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

DATA







EAAT1 (A-3) Alexa Fluor® 488: sc-515839 AF488. Direct fluorescent western blot analysis of EAAT1 expression in human cerebellum tissue extract. Blocked with UltraCruz® Blocking Reagent: sc-516214. Cruz Marker™ Molecular Weight Standards detected with Cruz Marker MW Tag-Alexa Fluor® 647: sc-516791.

SELECT PRODUCT CITATIONS

- Martín-Hernández, D., et al. 2019. Chronic mild stress alters kynurenine pathways changing the glutamate neurotransmission in frontal cortex of rats. Mol. Neurobiol. 56: 490-501.
- 2. Saba, J., et al. 2020. Astrocytes from cortex and striatum show differential responses to mitochondrial toxin and BDNF: implications for protection of striatal neurons expressing mutant huntingtin. J. Neuroinflammation 17: 290
- 3. Lacaille, H., et al. 2021. Preterm birth alters the maturation of the GABAergic system in the human prefrontal cortex. Front. Mol. Neurosci. 14: 827370.
- Marchese, N.A., et al. 2022. The intrinsic blue light responses of avian müller glial cells imply calcium release from internal stores. ASN Neuro 14: 17590914221076698.
- He, W., et al. 2022. Adenosine A_{2A} receptor antagonist improves cognitive impairment by inhibiting neuroinflammation and excitatory neurotoxicity in chronic periodontitis mice. Molecules 27: 6267.
- Haferkamp, U., et al. 2023. Human isogenic cells of the neurovascular unit exert transcriptomic cell type-specific effects on a blood-brain barrier in vitro model of late-onset Alzheimer disease. Fluids Barriers CNS. 20: 78.
- Mätlik, K., et al. 2024. Cell-type-specific CAG repeat expansions and toxicity of mutant Huntingtin in human striatum and cerebellum. Nat. Genet. 56: 383-394.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

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