

GAD-65 (A-3): sc-377145

BACKGROUND

There are two forms of glutamic acid decarboxylases (GADs) that are found in the brain: GAD-65 (also known as GAD2) and GAD-67 (also known as GAD1, GAD or SCP). GAD-65 and GAD-67 are members of the group II decarboxylase family of proteins and are responsible for catalyzing the rate limiting step in the production of GABA (γ -aminobutyric acid) from L-glutamic acid. Although both GADs are found in the brain, GAD-65 localizes to synaptic vesicle membranes in nerve terminals, while GAD-67 is distributed throughout the cell. GAD-67 is responsible for the basal levels of GABA synthesis. In the case of a heightened demand for GABA in neurotransmission, GAD-65 will transiently activate to assist in GABA production. The loss of GAD-65 is detrimental and can impair GABA neurotransmission, however the loss of GAD-67 is lethal. Due to alternative splicing, two isoforms exist for GAD-67, the predominant GAD-67 form and the minor GAD-25 form. GAD-25 is not expressed in brain but can be found in a variety of endocrine tissues.

CHROMOSOMAL LOCATION

Genetic locus: GAD2 (human) mapping to 10p12.1; Gad2 (mouse) mapping to 2 A3.

SOURCE

GAD-65 (A-3) is a mouse monoclonal antibody raised against amino acids 1-95 of GAD-65 of human origin.

PRODUCT

Each vial contains 200 μ g IgG₁ kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

GAD-65 (A-3) is available conjugated to agarose (sc-377145 AC), 500 μ g/0.25 ml agarose in 1 ml, for IP; to HRP (sc-377145 HRP), 200 μ g/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-377145 PE), fluorescein (sc-377145 FITC), Alexa Fluor[®] 488 (sc-377145 AF488), Alexa Fluor[®] 546 (sc-377145 AF546), Alexa Fluor[®] 594 (sc-377145 AF594) or Alexa Fluor[®] 647 (sc-377145 AF647), 200 μ g/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor[®] 680 (sc-377145 AF680) or Alexa Fluor[®] 790 (sc-377145 AF790), 200 μ g/ml, for Near-Infrared (NIR) WB, IF and FCM.

APPLICATIONS

GAD-65 (A-3) is recommended for detection of GAD-65 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), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

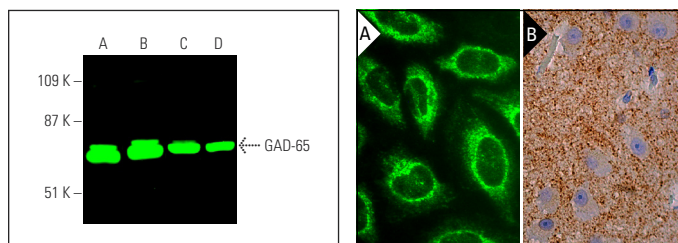
Suitable for use as control antibody for GAD-65 siRNA (h): sc-41964, GAD-65 siRNA (m): sc-41965, GAD-65 siRNA (r): sc-61888, GAD-65 shRNA Plasmid (h): sc-41964-SH, GAD-65 shRNA Plasmid (m): sc-41965-SH, GAD-65 shRNA Plasmid (r): sc-61888-SH, GAD-65 shRNA (h) Lentiviral Particles: sc-41964-V, GAD-65 shRNA (m) Lentiviral Particles: sc-41965-V and GAD-65 shRNA (r) Lentiviral Particles: sc-61888-V.

Molecular Weight of GAD-65: 65 kDa.

STORAGE

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

DATA



GAD-65 (A-3): sc-377145. Near-infrared western blot analysis of GAD-65 expression in rat cerebellum (A) and rat brain (B) tissue extracts and U-87 MG (C) and EOC 20 (D) whole cell lysates. Blocked with UltraCruz[®] Blocking Reagent: sc-516214. Detection reagent used: m-IgG κ BP-CFL 680: sc-516180.

GAD-65 (A-3): sc-377145. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization (A). Immunoperoxidase staining of formalin fixed, paraffin-embedded human cerebral cortex tissue showing neuropil staining (B).

SELECT PRODUCT CITATIONS

- Hernández-Cerón, M., et al. 2017. Participation of the dentate-rubral pathway in the kindling model of epilepsy. *J. Neurosci. Res.* 95: 1495-1502.
- Lacaille, H., et al. 2019. Impaired interneuron development in a novel model of neonatal brain injury. *eNeuro* 6: ENEURO.0300-18.2019.
- Frandsen, J., et al. 2020. Neural glyoxalase pathway enhancement by morin derivatives in an Alzheimer's disease model. *ACS Chem. Neurosci.* 11: 356-366.
- Branscome, H., et al. 2022. Retroviral infection of human neurospheres and use of stem cell EVs to repair cellular damage. *Sci. Rep.* 12: 2019.
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- Singh, S., et al. 2022. Somatostatin-mediated regulation of retinoic acid-induced differentiation of SH-SY5Y cells: neurotransmitters phenotype characterization. *Biomedicines* 10: 337.
- Tadokoro, T., et al. 2022. Precision spinal gene delivery-induced functional switch in nociceptive neurons reverses neuropathic pain. *Mol. Ther.* 30: 2722-2745.
- Liang, D., et al. 2024. A GABAergic system in atrioventricular node pacemaker cells controls electrical conduction between the atria and ventricles. *Cell Res.* 34: 556-571.
- Hayes, S.H., et al. 2024. Neurophysiological, structural, and molecular alterations in the prefrontal and auditory cortices following noise-induced hearing loss. *Neurobiol. Dis.* 200: 106619.

RESEARCH USE

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

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