Technical Data Sheet

Purified Mouse Anti-Glutamine Synthetase

Product Information

Material Number: 610517

Alternate Name: Glutamate-Ammonia Ligase; GLUL; GLNS

 Size:
 50 μg

 Concentration:
 250 μg/ml

Clone: 6/Glutamine Synthetase

Immunogen: Human Glutamine Synthetase aa. 1-373

Isotype:Mouse IgG2aReactivity:QC Testing: Rat

Tested in Development: Human, Mouse

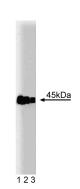
Target MW: 45 kDa

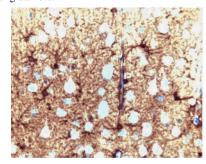
Storage Buffer: Aqueous buffered solution containing BSA, glycerol, and ≤0.09% sodium

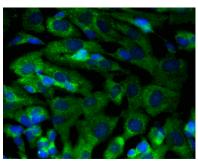
azide.

Description

Glutamine synthetase catalyzes the amination of glutamic acid to form glutamine. It is found in mammals as an octamer of identical 45 kDa subunits. Glutamine synthetase activity is a useful marker for astrocytes and an important differentiation feature in retina. It is also considered to be a key enzyme in the recycling of the neurotransmitter glutamate.







Western blot analysis of glutamine synthetase on a rat cerebrum lysate (left). Lane 1: 1:5000, lane 2: 1:10,000, lane 3: 1:20,000 dilution of the anti-glutamine synthetase antibody.

Glutamine synthetase staining on a rat cerebrum section (center). Section prepared during antibody development was formalin fixed and paraffin embedded without citrate buffer pretreatment. Note visible staining of astrocytes in the section. Magnification: 40X.

Immunofluorescent staining of SK-N-SH cells (right). Cells were seeded in a 384 well collagen coated Microplates (Material # 353962) at ~ 8,000 cells per well. After overnight incubation, cells were stained using the methanol fix/perm protocol (see Recommended Assay Procedure; Bioimaging protocol link) and the anti- Glutamine Synthetase antibody. The second step reagent was Alexa Fluor® 488 goat anti mouse Ig (Invitrogen)(pseudo colored green). Cell nuclei were counter stained with Hoechst 33342 (pseudo colored blue). The image was taken on a BD Pathway™ 855 or 435 Bioimager System using a 20x objective and merged using the BD AttoVison ™ software. This antibody also stained SH-SY5Y, C6, U87 and U373 cells using both the Triton™ X-100 and methanol fix/perm protocols (see Recommended Assay Procedure; Bioimaging protocol link).

Preparation and Storage

The monoclonal antibody was purified from tissue culture supernatant or ascites by affinity chromatography. Store undiluted at -20°C.

Application Notes

Application

Аррисации			
Western blot	Routinely Tested		
Immunohistochemistry	Tested During Development		
Bioimaging	Tested During Development	Tested During Development	
Immunoprecipitation	Not Recommended		
Immunofluorescence	Not Recommended		

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Recommended Assay Procedure:

For Western blot and IHC: Please refer to http://www.bdbiosciences.com/resources/cellbiology/index.jsp For Bioimaging: Please refer to http://www.bdbiosciences.com/resources/cellularimaging/index.jsp

Suggested Companion Products

Catalog Number	Name	Size	Clone
611463	Rat Cerebrum Lysate	500 μg	(none)
554002	HRP Goat Anti-Mouse Ig	1.0 ml	(none)
554001	FITC Goat Anti-Mouse Ig	0.5 mg	Polyclonal
353219	BD Falcon™ 96-well Imaging Plate		(none)
554655	Fixation Buffer	100 ml	(none)
558050	Perm Buffer III	125 ml	(none)

Product Notices

- Since applications vary, each investigator should titrate the reagent to obtain optimal results.
- Please refer to www.bdbiosciences.com/pharmingen/protocols for technical protocols.
- Caution: Sodium azide yields highly toxic hydrazoic acid under acidic conditions. Dilute azide compounds in running water before discarding to avoid accumulation of potentially explosive deposits in plumbing.
- 4. Source of all serum proteins is from USDA inspected abattoirs located in the United States.
- 5. Triton is a trademark of the Dow Chemical Company.

References

Fei ZL, D'Ambrosio C, Li S, Surmacz E, Baserga R. Association of insulin receptor substrate 1 with simian virus 40 large T antigen. Mol Cell Biol. 1995; 15(8):4232-4239. (Biology: Immunoprecipitation)

Kentroti S, Baker R, Lee K, Bruce C, Vernadakis A. Platelet-activating factor increases glutamine synthetase activity in early and late passage C-6 glioma cells. J Neurosci Res. 1991; 28(4):497-506. (Biology)

Kronfeld I, Kazimirsky G, Lorenzo PS, Garfield SH, Blumberg PM, Brodie C. Phosphorylation of protein kinase Cdelta on distinct tyrosine residues regulates specific cellular functions. J Biol Chem. 2000; 275(45):35491-35498. (Biology: Western blot)

Labow BI, Souba WW, Abcouwer SF. Glutamine synthetase expression in muscle is regulated by transcriptional and posttranscriptional mechanisms. Am J Physiol. 1999; 276(6):E1136-E1145. (Biology: Western blot)

Vardimon L, Fox LE, Cohen-Kupiec R, Degenstein L, Moscona AA. Expression of v-src in embryonic neural retina alters cell adhesion, inhibits histogenesis, and prevents induction of glutamine synthetase. Mol Cell Biol. 1991; 11(10):5275-5284. (Biology)

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