# Anti-Glutamine Synthetase, clone GS-6

Monoclonal Antibody Cat. # MAB302 Lot # 2318760

FOR RESEARCH USE ONLY NOT FOR USE IN HUMANS

pack size: 100 µg

Store at -20°C



## Certificate of Analysis

Applications WB, ELISA, IH	Species Cross-Reactivity H, M, R, Sh	Antibody Isotype IgG2a	Epitope/ Region	Host Species	Molecular Weight	Accession #
Background				IVI	45 kDa	NM_005178.3

Glutamine Synthetase catalyzes the conversion of ammonia and glutamate to glutamine. It is found in astrocytes as an octamer of identical 45 kDa subunits. The function of Glutamine Synthetase is the detoxification of brain ammonia. It also has an important role in the metabolic regulation of neurotransmitter glutamate. Because of the multiple functions and importance Glutamine Synthetase in cellular metabolism, both catalytic activities and synthesis are highly regulated. The activity of Glutamine Synthetase is controlled by adenylylation. Its activity is decreased in the cerebral cortex of brains affected by Alzheimer's disease, particularly in the vicinity of senile plaques. It is also decreased under conditions of glucose deprivation. The level of expression Glutamine Synthetase is increased during ischemia in vivo or hypoxia in culture.

### Presentation

Liquid in PBS containing 50% glycerol, BSA and < 0.09% sodium azide.

### Concentration

1 mg/mL

### Specificity

Reacts strongly with a single band on Western blots of sheep brain and rat brain.

### Immunogen

Glutamine synthetase purified from sheep brain.

### Molecular Weight

45 kDa

### **Method of Purification**

Protein A Purified

### Storage and Handling

Stable for 1 year at -20°C in undiluted aliquots from date of receipt.

Handling Recommendations: Upon receipt, and prior to removing the cap, centrifuge the vial and gently mix the solution. Aliquot into microcentrifuge tubes and store at -20°C. Avoid repeated freeze/thaw cycles, which may damage IgG and affect product performance. Note: Variability in freezer temperatures below -20°C may cause glycerol containing solutions to become frozen during storage.

#### Control

Rat brain tissue, rat brain cytosolic fraction extract, human brain lysate

### **Quality Control Testing**

Evaluated by Western Blot on Human brain lysates.

Western Blot Analysis: 1:500 dilution of this antibody detected Glutamine Synthetase on 10 μg of Human brain lysates.

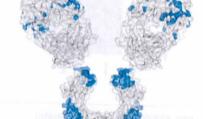
# 130 KD 95 KD 72 KD-55 KD-36 KD-28 KD 17 KD-

### Western Blot Analysis:

Representative lot data. Human brain lysate was resolved by electrophoresis, transferred to PVDF membrane and probed with anti-Glutamine Synthetase (1:500 dilution).

Proteins were visualized using a goat anti-mouse secondary antibody conjugated to HRP and a chemiluminescence detection

Arrow indicates
GLUTAMINE SY SYNTHETASE (~44 kDa).



### References

- 1. Oh, Edwin C T, et al. (2007). Proc. Natl. Acad.
- Sci. U.S.A. 104: 1679-84. Suarez, I, et al. (2005). Neuropathol. Appl. Neurobiol. 31:163-9.
- 3. Wilhelmsson, U., et al. (2004). J. Neuroscience. 24:5016-5021
- 4. Ward, M., et al. (2004). Cell Tissue Res. 315:305-310.
- 5. Weiner, I.D., et al. (2003). Gastroenterol. 124:1432-1440.

# Additional Research Applications

Immunohistochemistry: A previous lot of this antibody was used in immunohistochemistry on rat brain tissue.

ELISA: A previous lot of this antibody was used on ELISA.

Optimal working dilutions must be determined by end user.

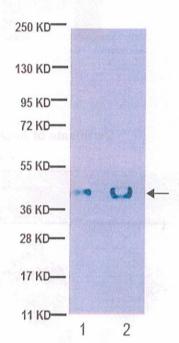
APPLICATION LEGEND: WB Western Blotting ELISA Enzyme-linked Immunosorbent Assay IP Immunoprecipitation IC Immunocytochemistry IF Immunofluorescence IH Immunohistochemistry (Tissue) SPECIES LEGEND: H Human M Mouse R Rat Rb Rabbit Sh Sheep WR Most Common Vertebrates

11 KD

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### Additional Research Applications



### Western Blot Analysis:

Representative image from a previous

lot.
Human brain lysate (Lane 1) and
Mouse brain membrane lysate (Lane
2) were resolved by electrophoresis,
transferred to PVDF membrane and
probed with ahti-GLUTAMINE SYNTHETASE (1:500 dilution of a previous lot).

Proteins were visualized using a goat anti-mouse secondary antibody secondary to HRP to conjugated to HRP and a chemiluminescence detection system.

Arrow indicates protein GLUTAMINE SYNTHETASE (~44 kDa).

### PROTOCOL

### Western Blotting

- Perform SDS-polyacrylamide gel electrophoresis (SDS-PAGE) on cell lysate and transfer the proteins to a PVDF membrane. Wash the PVDF membrane twice with water.
- Block the blotted PVDF membrane in freshly prepared 5% BSA with 0.05% Tween®-20 for 1 hour at room temperature with constant agitation.
- Incubate the PVDF with the recommended dilution of Anti-Glutamine Synthetase, clone GS-6 diluted in freshly prepared 5% BSA for 1 hour at room temperature or overnight with agitation at 2-8°C.
- Wash the PVDF 3 times with TBST. 4.
- Incubate the PVDF in the secondary reagent of choice (a goat anti-rabbit HRP conjugated IgG, Catalog # 12-348 1:1000 dilution was used) in 5% milk for 1 hour with agitation at room temperature.
- 6. Wash the PVDF 3-5 times with TBST.
- Use Spray and Glow Catalog # 17-373 to visualize results. Use as directed.

### **RELATED PRODUCTS (specific)**

cat#

### description

AP124P

Goat anti-Mouse IgG, Peroxidase Conjugated, H+L

### RELATED PRODUCTS (non-specific)

cat#		description
IPVH00010		Immobilon-P 26.5 cm x 3.75 m Roll PVDF 0.45 µm
IPFL00010	1	Immobilon-FL 26.5 cm x 3.75 m Roll PVDF 0.45 µm
IPVH07850		Immobilon-P 7 x 8.4 cm PVDF 0.45 mm (sheet) 50/pk
ISEQ00010	100	Immobilon-P SQ 26.5 cm x 3.75 m 1 roll PVDF 0.2 µm
ISEQ07850	1871	Immobilon-P 7 x 8.4 cm PVDF 0.2 mm (sheet) 50/pk
IPFL07810	180	Immobilon-FL 7 x 8.4 cm PVDF 0.45 mm (sheet) 10/pk
WBKLS0100	100	Immobilon Western Chemilum HRP Substrate 100 mL
17-373		Spray & Glow™ ECL WB Detection System 1 ea
2060		Re-Blot Western Blot Recycling Kit
2500	100	Re-Blot Plus Western Blot Recycling Kit
B2080- 175GM	100	Blot Quick Blocker Membrane Blocking Agent 175G

💹 antibodies 📕 Multiplex products 💹 biotools 💹 cell culture 📕 enzymes 📕 kits 📓 proteins/peptides 📒 siRNA/cDNA products

Please visit www.millipore.com for additional product information, test data and references