

GLUL Antibody (N-term)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP6892a

Specification

GLUL Antibody (N-term) - Product Information

Application	WB, IHC-P,E
Primary Accession	P15104
Other Accession	P46410 , P15105 , Q4R7U3 , P16580 , P15103
Reactivity Predicted	Human, Mouse Bovine, Chicken, Monkey, Pig
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit Ig
Antigen Region	70-96

GLUL Antibody (N-term) - Additional Information

Gene ID 2752

Other Names

Glutamine synthetase, GS, Glutamate decarboxylase, Glutamate--ammonia ligase, GLUL, GLNS

Target/Specificity

This GLUL antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 70-96 amino acids from the N-terminal region of human GLUL.

Dilution

WB~~1:1000

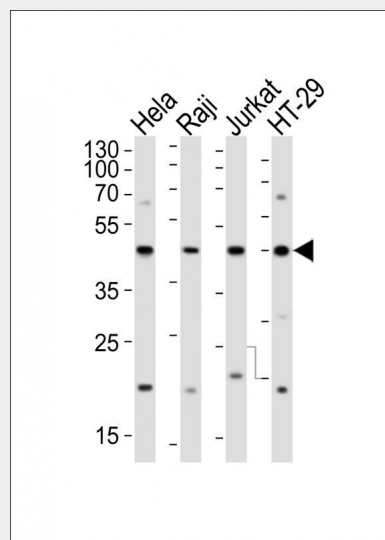
IHC-P~~1:25

Format

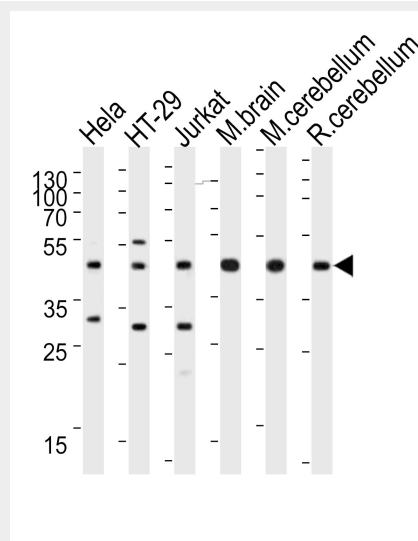
Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.



Western blot analysis of lysates from HeLa, Raji, Jurkat, HT-29 cell line (from left to right), using GLUL Antibody (N-term)(Cat. #AP6892a). AP6892a was diluted at 1:2000 at each lane. A goat anti-rabbit IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20ug per lane.



Western blot analysis of lysates from HeLa, HT-29, Jurkat cell line, mouse brain, mouse cerebellum, rat cerebellum tissue lysate(from

Precautions

GLUL Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

GLUL Antibody (N-term) - Protein Information

Name GLUL

{ECO:0000303|PubMed:30158707,
ECO:0000312|HGNC:HGNC:4341}

Function

Glutamine synthetase that catalyzes the ATP-dependent conversion of glutamate and ammonia to glutamine (PubMed:30158707, PubMed:16267323).

Its role depends on tissue localization: in the brain, it regulates the levels of toxic ammonia and converts neurotoxic glutamate to harmless glutamine, whereas in the liver, it is one of the enzymes responsible for the removal of ammonia (By similarity). Essential for proliferation of fetal skin fibroblasts (PubMed:18662667).

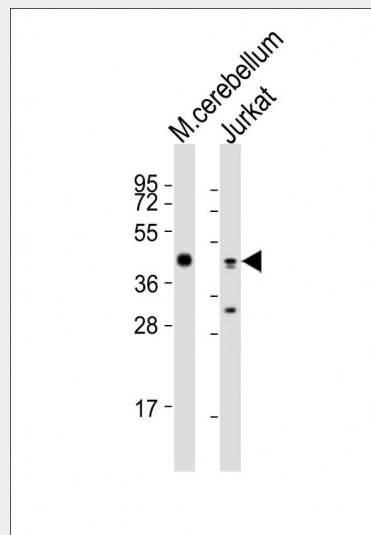
Independently of its glutamine synthetase activity, required for endothelial cell migration during vascular development: acts by regulating membrane localization and activation of the GTPase RHOJ, possibly by promoting RHOJ palmitoylation (PubMed:30158707).

May act as a palmitoyltransferase for RHOJ: able to autopalmitylate and then transfer the palmitoyl group to RHOJ (PubMed:30158707). Plays a role in ribosomal 40S subunit biogenesis (PubMed:26711351).

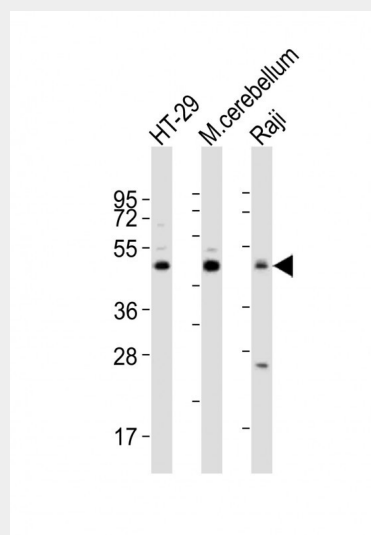
Cellular Location

Cytoplasm, cytosol. Microsome {ECO:0000250|UniProtKB:P09606}. Mitochondrion {ECO:0000250|UniProtKB:P09606}. Cell membrane; Lipid-anchor. Note=Mainly localizes in the cytosol, with a fraction

left to right), using GLUL Antibody (N-term)(Cat. #AP6892a). AP6892a was diluted at 1:1000 at each lane. A goat anti-rabbit IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20ug per lane.



All lanes : Anti-GLUL Antibody (N-term) at 1:2000 dilution Lane 1: mouse cerebellum lysates Lane 2: Jurkat whole cell lysates Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 42 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



All lanes : Anti-GLUL Antibody (N-term) at 1:2000 dilution Lane 1: HT-29 whole cell lysates Lane 2: mouse cerebellum lysates Lane 3: Raji whole cell lysates Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L),

associated with the cell membrane

Tissue Location

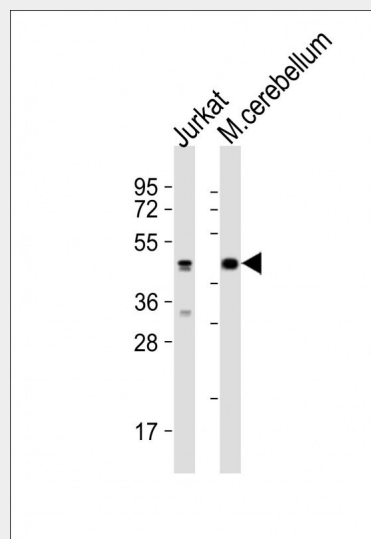
Expressed in endothelial cells.

GLUL Antibody (N-term) - Protocols

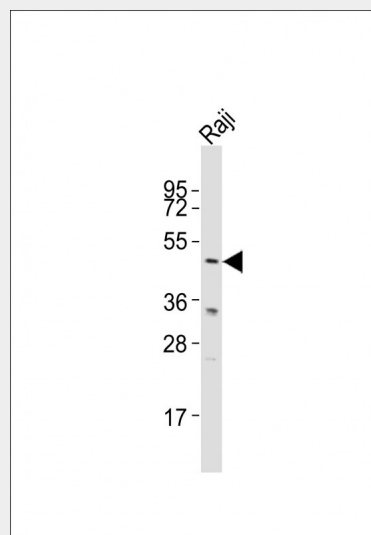
Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

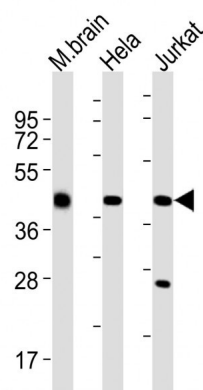
Peroxidase conjugated at 1/10000 dilution
Predicted band size : 42 kDa
Blocking/Dilution buffer: 5% NFDM/TBST.



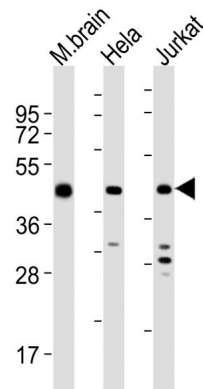
All lanes : Anti-GLUL Antibody (N-term) at 1:2000 dilution Lane 1: Jurkat whole cell lysates Lane 2: mouse cerebellum lysates Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 42 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



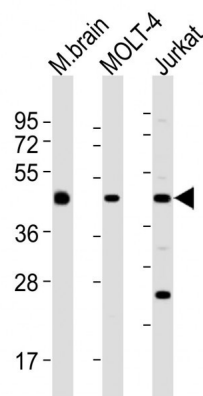
Anti-GLUL Antibody (N-term) at 1:1000 dilution + Raji whole cell lysates Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 42 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



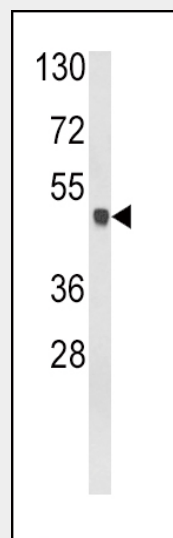
All lanes : Anti-GLUL Antibody (N-term) at 1:2000 dilution Lane 1: mouse brain lysates Lane 2: Hela whole cell lysates Lane 3: Jurkat whole cell lysates Lysates/proteins at 20 μ g per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 42 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



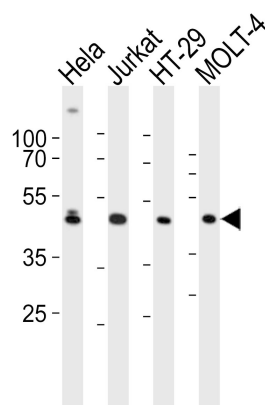
All lanes : Anti-GLUL Antibody (N-term) at 1:2000 dilution Lane 1: mouse brain lysates Lane 2: Hela whole cell lysates Lane 3: Jurkat whole cell lysates Lysates/proteins at 20 μ g per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 42 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



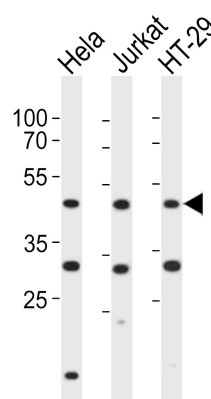
All lanes : Anti-GLUL Antibody (N-term) at 1:2000 dilution Lane 1: mouse brain lysates Lane 2: MOLT-4 whole cell lysates Lane 3: Jurkat whole cell lysates Lysates/proteins at 20 µg per lane. Secondary Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/10000 dilution Predicted band size : 42 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



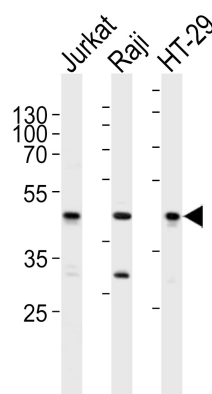
Western blot analysis of GLUL Antibody (N-term) (Cat. #AP6892a) in mouse cerebellum tissue lysates (35ug/lane). GLUL (arrow) was detected using the purified Pab.



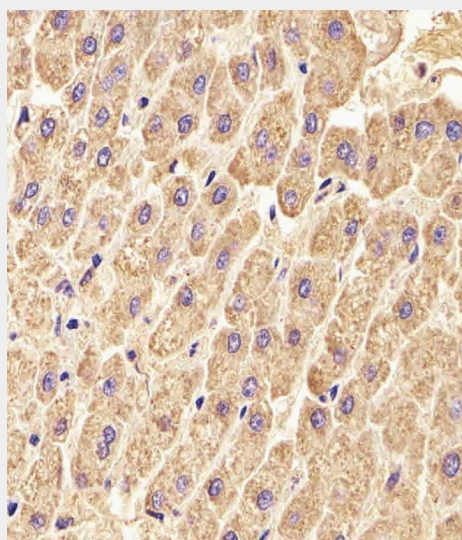
Western blot analysis of lysates from HeLa, Jurkat, HT-29, MOLT-4 cell line (from left to right), using GLUL Antibody (N-term)(Cat. #AP6892a). AP6892a was diluted at 1:1000 at each lane. A goat anti-rabbit IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20ug per lane.



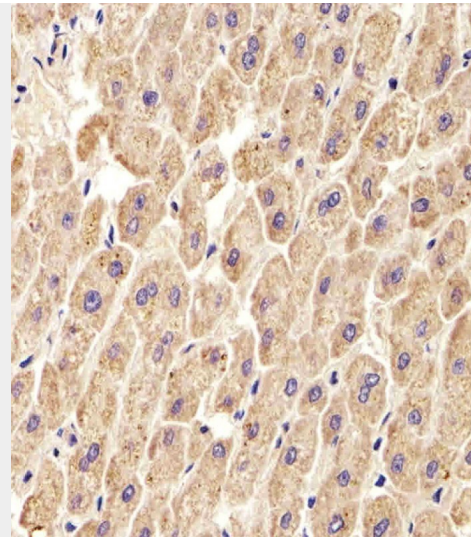
Western blot analysis of lysates from HeLa, Jurkat, HT-29 cell line (from left to right), using GLUL Antibody (N-term)(Cat. #AP6892a). AP6892a was diluted at 1:1000 at each lane. A goat anti-rabbit IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20ug per lane.



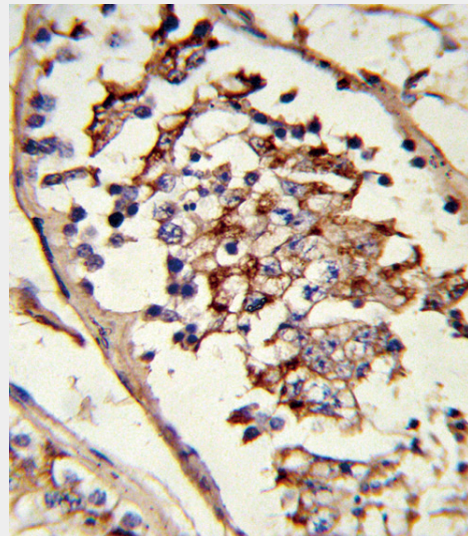
Western blot analysis of lysates from Jurkat, Raji, HT-29 cell line (from left to right), using GLUL Antibody (N-term)(Cat. #AP6892a). AP6892a was diluted at 1:1000 at each lane. A goat anti-rabbit IgG H&L(HRP) at 1:10000 dilution was used as the secondary antibody. Lysates at 20ug per lane.



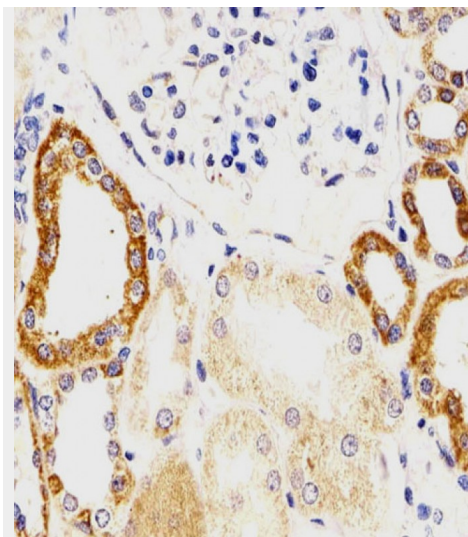
Immunohistochemical analysis of paraffin-embedded H. liver section using GLUL Antibody (N-term)(Cat#AP6892a). AP6892a was diluted at 1:25 dilution. A undiluted biotinylated goat polyvalent antibody was used as the secondary, followed by DAB staining.



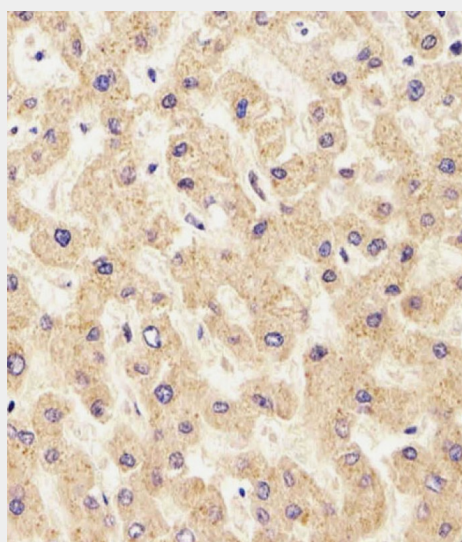
Immunohistochemical analysis of paraffin-embedded H. liver section using GLUL Antibody (N-term)(Cat#AP6892a). AP6892a was diluted at 1:25 dilution. A undiluted biotinylated goat polyvalent antibody was used as the secondary, followed by DAB staining.



Formalin-fixed and paraffin-embedded human testis tissue reacted with GLUL Antibody (N-term), which was peroxidase-conjugated to the secondary antibody, followed by DAB staining. This data demonstrates the use of this antibody for immunohistochemistry; clinical relevance has not been evaluated.

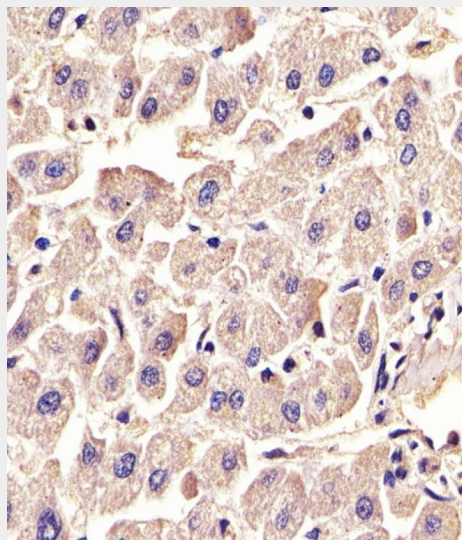


AP6892a staining GLUL in Human kidney tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with formaldehyde and blocked with 3% BSA for 0.5 hour at room temperature; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody (1/25) for 1 hour at 37°C. A undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.

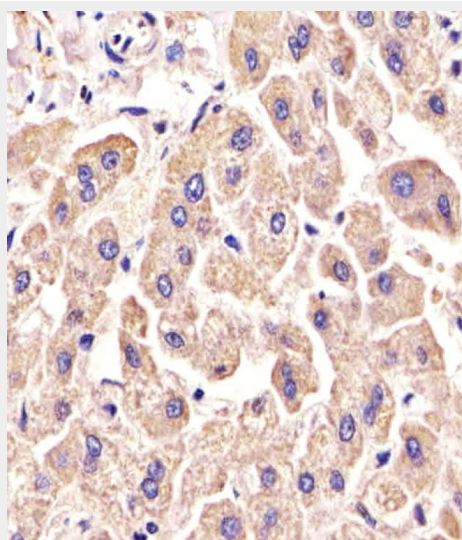


AP6892a staining GLUL in Human liver tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with formaldehyde and blocked with 3% BSA for 0.5 hour at room temperature; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody (1/25) for 1 hour at 37°C. A

undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.



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AP6892a staining GLUL in Human liver tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with formaldehyde and blocked with 3% BSA for 0.

5 hour at room temperature; antigen retrieval was by heat mediation with a citrate buffer (pH6). Samples were incubated with primary antibody (1/25) for 1 hours at 37°C. A undiluted biotinylated goat polyvalent antibody was used as the secondary antibody.

GLUL Antibody (N-term) - Background

GLUL belongs to the glutamine synthetase family. It catalyzes the synthesis of glutamine from glutamate and ammonia. Glutamine is a main source of energy and is involved in cell proliferation, inhibition of apoptosis, and cell signaling.

GLUL Antibody (N-term) - References

Di Tommaso, L., et.al., J. Hepatol. 50 (4), 746-754 (2009)