Elabscience®

Glutamine Synthetase Polyclonal Antibody

catalog number: E-AB-93350

Note: Centrifuge before opening to ensure complete recovery of vial contents.

Description	
Reactivity	Human;Mouse;Rat
Immunogen	Recombinant fusion protein of human Glutamine Synthetase
Host	Rabbit
Isotype	IgG
Purification	Affinity purification
Buffer	Phosphate buffered solution, pH 7.4, containing 0.05% stabilizer and 50% glycerol.
Applications	Recommended Dilution

WB	1:500-1:2000
IF	1:50-1:200

Data





Western blot analysis of various lysates using Glutamine Synthetase Polyclonal Antibody at 1:400 dilution.

> Observed-MW:42 kDa Calculated-MW:42 kDa



Immunofluorescence analysis of HeLa cells using [KO Validated] Glutamine Synthetase Polyclonal Antibodyat dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.



Immunofluorescence analysis of NIH/3T3 cells using [KO Validated] Glutamine Synthetase Polyclonal Antibody at dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining. dilution of 1:100 (40x lens). Blue: DAPI for nuclear staining.

Immunofluorescence analysis of PC-12 cells using [KO Validated] Glutamine Synthetase Polyclonal Antibody at

Preparation & Storage	
Storage	Store at -20°C Valid for 12 months. Avoid freeze / thaw cycles.
Shipping	The product is shipped with ice pack, upon receipt, store it immediately at the temperature recommended.

Background

For Research Use Only

Toll-free: 1-888-852-8623 Web:www.elabscience.com

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The protein encoded by this gene belongs to the glutamine synthetase family. It catalyzes the synthesis of glutamine from glutamate and ammonia in an ATP-dependent reaction. This protein plays a role in ammonia and glutamate detoxification, acid-base homeostasis, cell signaling, and cell proliferation. Glutamine is an abundant amino acid, and is important to the biosynthesis of several amino acids, pyrimidines, and purines. Mutations in this gene are associated with congenital glutamine deficiency, and overexpression of this gene was observed in some primary liver cancer samples. There are six pseudogenes of this gene found on chromosomes 2, 5, 9, 11, and 12. Alternative splicing results in multiple transcript variants.

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