

Datasheet for ABIN1607650 anti-GLN1 antibody (Biotin)

Image



Overview

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Quantity:	100 µg
Target:	GLN1
Reactivity:	Brevibacterium
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This GLN1 antibody is conjugated to Biotin
Application:	Western Blotting (WB), Immunoprecipitation (IP), ELISA
Product Details	
Purpose:	Glutamine Synthetase Antibody Biotin Conjugated
Immunogen:	Immunogen: Glutamine Synthetase [Microbial] Immunogen Type: Native Protein
lsotype:	lgG
Cross-Reactivity (Details):	Assay by immunoelectrophoresis resulted in a single precipitin arc against anti-Biotin, anti-Goat Serum as well as purified and partially purified Glutamine Synthetase [Microbial].
Characteristics:	Synonyms: goat anti-Glutamine Synthetase Antibody biotin Conjugation, biotin Conjugated goat anti-Glutamine Synthetase Antibody, Glutamine synthetase
Purification:	Anti-Glutamine Synthetase (microbial) antibody is an IgG fraction antibody purified from monospecific antiserum by a multi-step process which includes delipidation, salt fractionation and ion exchange chromatography followed by extensive dialysis against the buffer stated above.

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Target Details

Target:	GLN1
Alternative Name:	Glutamine Synthetase (GLN1 Products)
Background:	Background: Glutamine Synthetase is a key enzyme in the metabolism of nitrogen. Glutamine
	synthetase catalyzes an ATP-dependent condensation reaction between ammonia and
	glutamate to yield glutamine. Glutamine is a key builder of proteins as well as a vehicle to
	deliver nitrogen atoms to enzymes that build molecules dependent on nitrogen. Glutamine
	Synthetase from a microbial source is composed of twelve subunits that each house an active
	site. During the reaction of glutamine synthetase, the active sites bind ammonia and glutamate
	as well as an ATP molecule to power the reaction. Negative feedback regulation is provided by
	the active sites ability to weakly bind other molecules and once their concentrations rise too
	high, the enzyme shuts off. Glutamine Synthetase has applications in neuroscience due to
	location in astrocytes within the brain and fluctuations in glutamine synthetase can
	detrimentally effect the astrocytes. Anti-Glutamine Synthetase (Microbial) Antibody is ideal for
	investigators in Molecular Biology, Neuroscience, and Enzymology.
Gene ID:	3345165
UniProt:	Q79VE3
Pathways:	Positive Regulation of Peptide Hormone Secretion
Application Details	
Application Notes:	Application Note: Anti-Glutamine Synthetase (microbial) antibody has been assayed against
	1.0 μ g of Glutamine Synthetase in a standard capture ELISA using Peroxidase Conjugated
	Streptavidin #S000-03 and ABTS (2,2'-azino-bis-[3-ethylbenthiazoline-6-sulfonic acid]) code #
	ABTS-100 as a substrate for 30 minutes at room temperature. A working dilution of 1:4,000 to
	1:20,000 of the reconstitution concentration is suggested for this product.
	Western Blot Dilution: 1:500 - 1:2,500
	Immunoprecipitation Dilution: 1:100

ELISA Dilution: 1:2,000 - 1:10,000 Other: User Optimized

Restrictions:

For Research Use only

Handling

Format:

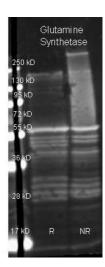
Lyophilized

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Handling

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Reconstitution:	Reconstitution Volume: 100 µL Reconstitution Buffer: Restore with deionized water (or equivalent)
Concentration:	10 mg/mL
Buffer:	Buffer: 0.02 M Potassium Phosphate, 0.15 M Sodium Chloride, pH 7.2 Stabilizer: 10 mg/mL Bovine Serum Albumin (BSA) - Immunoglobulin and Protease free Preservative: 0.01 % (w/v) Sodium Azide
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	4 °C,-20 °C
Storage Comment:	Store vial at 4° C prior to restoration. For extended storage aliquot contents and freeze at -20° C or below. Avoid cycles of freezing and thawing. Centrifuge product if not completely clear after standing at room temperature. This product is stable for several weeks at 4° C as an undiluted liquid. Dilute only prior to immediate use.
Expiry Date:	12 months

Images



Western Blotting

Image 1. Goat anti Glutamine Synthetase antibody was used to detect Glutamine Synthetase under reducing (R) and non-reducing (NR) conditions. Reduced samples of purified target proteins contained 4% BME and were boiled for 5 minutes. Samples of ~1ug of protein per lane were run by SDS-PAGE. Protein was transferred to nitrocellulose and probed with 1:3000 dilution of primary antibody (ON 4 C in ABIN925618). Detection shown was using Dylight 649 conjugated Donkey anti goat (605-743-125 lot 20834 1:10K in TBS/ABIN925618) 1 hr RT. Images were collected using the BioRad VersaDoc System.