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Datasheet for ABIN1607648 anti-Beta-Phosphoglucomutase antibody (Biotin)

1 Image



Overview

Quantity:	100 µg
Target:	Beta-Phosphoglucomutase
Reactivity:	Lactococcus lacti
Host:	Goat
Clonality:	Polyclonal
Conjugate:	This Beta-Phosphoglucomutase antibody is conjugated to Biotin
Application:	Western Blotting (WB), ELISA
Product Details	
Product Details Immunogen:	beta Phosphoglucomutase [Lactococcus lacti]
	beta Phosphoglucomutase [Lactococcus lacti] Immunogen Type: Native Protein
Immunogen:	Immunogen Type: Native Protein
Immunogen: Isotype:	Immunogen Type: Native Protein
Immunogen: Isotype: Cross-Reactivity (Details):	Immunogen Type: Native Protein IgG Cross reactivity against Beta-Phosphoglucomutase from other sources is unknown.
Immunogen: Isotype: Cross-Reactivity (Details):	Immunogen Type: Native Protein IgG Cross reactivity against Beta-Phosphoglucomutase from other sources is unknown. Anti-Beta-Phosphoglucomutase is an IgG fraction antibody purified from monospecific
Immunogen: Isotype: Cross-Reactivity (Details):	Immunogen Type: Native Protein IgG Cross reactivity against Beta-Phosphoglucomutase from other sources is unknown. Anti-Beta-Phosphoglucomutase is an IgG fraction antibody purified from monospecific antiserum by a multi-step process which includes delipidation, salt fractionation and ion
Immunogen: Isotype: Cross-Reactivity (Details):	Immunogen Type: Native Protein IgG Cross reactivity against Beta-Phosphoglucomutase from other sources is unknown. Anti-Beta-Phosphoglucomutase 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:	Beta-Phosphoglucomutase
Alternative Name:	Beta-Phosphoglucomutase (Beta-Phosphoglucomutase Products)
Background:	Anti-Beta-Phosphoglucomutase antibody is located in the cytoplasm, working to catalyze the
	transference of a phosphoryl group on a glucose or lactose monomer from the 1' to the 6'
	position in the forward direction or 6' to the 1' position in the reverse direction. Anti-Beta-
	Phosphoglucomutase is essential to regulating the flow of carbohydrate intermediates in
	glycolysis as well as the formation of the sugar nucleotide UDP-glucose. Part of the HAD-like
	hydrolase superfamily, Anti-Beta-Phosphoglucomutase antibody is ideal for researchers
	interested in Metabolism research. Synonyms: Beta PGM antibody, Glucose phosphomutase 1
	antibody, Glucose phosphomutase antibody, PGM antibody, pgmB antibody
Gene ID:	1114041
UniProt:	P71447
Application Details	
Application Notes:	Anti-Beta-Phosphoglucomutase has been assayed against 1.0 μ g of Beta-
	Phosphoglucomutase 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 1:10.000 to 1:50.000 is
	suggested for this product.
	ELISA Dilution: 1:5.000 - 1:25.000
	Western Blot Dilution: 1:500 - 1:3.000
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	10 mg/mL
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
Handling Advice:	Store the vial at -20°C or below after dilution. Avoid cycles of freezing and thawing.
Storage:	-20 °C

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Store vial at -20 °C or below prior to opening. This vial contains a relatively low volume of

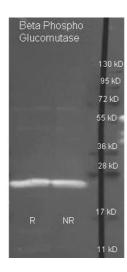
Storage Comment:

reagent (25 μ L). To minimize loss of volume dilute 1:10 by adding 225 μ L of the buffer stated above directly to the vial. Recap, mix thoroughly and briefly centrifuge to collect the volume at the bottom of the vial. Use this intermediate dilution when calculating final dilutions as recommended below.

Expiry Date:

Expiration date is one (1) year from date of opening.

Images



Western Blotting

Image 1. Goat anti antibody was used to detect purified Beta Phospho Glucomutase under reducing (R) and nonreducing (NR) conditions. Reduced samples of protein 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 488 conjugated Donkey anti goat (605-741-125 lot 21094 1:10K in TBS/ABIN925618 1 hr RT). Images were collected using the BioRad VersaDoc System.

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