antibodies

Datasheet for ABIN361757 anti-Phosphothreonine antibody

3 Images



Overview

Quantity:	400 µL
Target:	Phosphothreonine
Reactivity:	Please inquire
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This Phosphothreonine antibody is un-conjugated
Application:	Western Blotting (WB), ELISA, Immunoprecipitation (IP), Immunocytochemistry (ICC), Immunofluorescence (IF)

Product Details

Immunogen:	Phosphothreonine conjugated to KLH
Specificity:	Detects proteins phosphorylated on threonine residues. Does not cross-react with phosphotyrosine.
Purification:	Peptide Affinity Purified

Target Details

Target:	Phosphothreonine
Abstract:	Phosphothreonine Products
Target Type:	Amino Acid
Background:	Protein phosphorylation is an important posttranslational modification that serves many key
	functions to regulate a protein's activity, localization, and protein-protein interactions.

Order at www.antibodies-online.com | www.antikoerper-online.de | www.anticorps-enligne.fr | www.antibodies-online.cn International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 1/4 | Product datasheet for ABIN361757 | 09/12/2023 | Copyright antibodies-online. All rights reserved. Phosphorylation is catalyzed by various specific protein kinases, which involves removing a phosphate group from ATP and covalently attaching it to to a recipient protein that acts as a substrate. Most kinases act on both serine and threonine, others act on tyrosine, and a number (dual specificity kinases) act on all three. Because phosphorylation can occur at multiple sites on any given protein, it can therefore change the function or localization of that protein at any time (1). Changing the function of these proteins has been linked to a number of diseases, including cancer, diabetes, heart disease, inflammation and neurological disorders (2-4).

Application Details

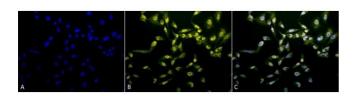
Application Notes:	 WB (1:500) ICC/IF (1:60) ELISA (1:2000) IP (1:100) optimal dilutions for assays should be determined by the user.
Comment:	2 µg/ml of ABIN361757 was sufficient for detection of phosphorylation signal in western blot analysis using mouse spleen extract treated with Vanadium.
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Concentration:	0.25 mg/mL
Buffer:	PBS, 50 % glycerol, 0.01 % sodium azide, Storage buffer may change when conjugated
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 Comment: -20°C

Storage:

-20 °C

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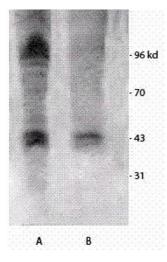


Immunocytochemistry

1. Immunocytochemistry/Immunofluorescence Image analysis using Rabbit Anti-Phosphothreonine Polyclonal Antibody (ABIN361757). Tissue: Cervical cancer cell line (HeLa). Species: Human. Fixation: 2 % Formaldehyde for 20 min at RT. Primary Antibody: Rabbit Anti-Phosphothreonine Polyclonal Antibody (ABIN361757) at 1:60 for 12 hours at 4 °C. Secondary Antibody: R-PE Goat Anti-Rabbit (yellow) at 1:200 for 2 hours at RT. Counterstain: DAPI (blue) nuclear stain at 1:40000 for 2 hours at RT. Localization: Cytoplasm. Nucleus. Magnification: 20x. (A) DAPI (blue) nuclear stain. (B) Anti-Phosphothreonine Antibody. (C) Composite.

Western Blotting

Image 2. Phospho Threonine, mouse brain extract.





Immunocytochemistry

Image 3. Immunocytochemistry/Immunofluorescence analysis using Rabbit Anti-Phosphothreonine Polyclonal Antibody (ABIN361757). Tissue: Cervical cancer cell line (HeLa). Species: Human. Fixation: 2 % Formaldehyde for Antibody: 20 min at RT. Primary Rabbit Anti-Phosphothreonine Polyclonal Antibody (ABIN361757) at 1:60 for 12 hours at 4 °C. Secondary Antibody: FITC Goat Anti-Rabbit (green) at 1:200 for 2 hours at RT. Counterstain: DAPI (blue) nuclear stain at 1:40000 for 2 hours at RT. Localization: Cytoplasm. Nucleus. Magnification: 100x. (A)

DAPI (blue) nuclear stain. (B) Anti-Phosphothreonine Antibody. (C) Composite.

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