abcam

Product datasheet

Anti-Phospho Serine/Threonine Chk1/Chk2 substrate antibody ab131023

1 References 1 Image

Overview

Product name Anti-Phospho Serine/Threonine Chk1/Chk2 substrate antibody

Description Rabbit polyclonal to Phospho Serine/Threonine Chk1/Chk2 substrate

Host species Rabbit

Tested applications Suitable for: WB

Species reactivity Reacts with: Human

Immunogen Synthetic peptide within Human Phospho Serine/Threonine Chk1/Chk2 substrate aa 1-100

conjugated to keyhole limpet haemocyanin. The exact sequence is proprietary.

Database link: P04637

Positive controlThis antibody gave a positive signal in HeLa and HeLa UV irradiated whole cell lysates.

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches of this product that have a concentration < 1mg/ml may have BSA added as a stabilising

agent. If you would like information about the formulation of a specific lot, please contact our

scientific support team who will be happy to help.

Purity Immunogen affinity purified

1

Clonality

Polyclonal

Isotype

lgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab131023 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

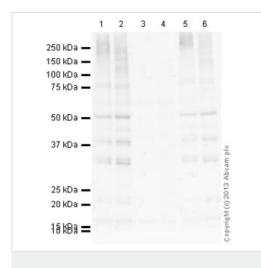
Application	Abreviews	Notes
WB		Use a concentration of 1 µg/ml. Detects a band of approximately 53 kDa (predicted molecular weight: 53 kDa).

Target

Relevance

Several protein kinases mediate cellular responses to DNA damage. This includes the serine/threonine kinases Chk1 and Chk2. In response to DNA damage Chk1 and Chk2 phosphorylate a large range of proteins on Serine (S) or Threonine (T) residues on RxxS/T degenerated consensus sites.

Images



Western blot - Anti-Phospho Serine/Threonine Chk1/Chk2 substrate antibody (ab131023) **All lanes :** Anti-Phospho Serine/Threonine Chk1/Chk2 substrate antibody (ab131023) at 1 μg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2: Hela Whole Cell Lysate - UV Irradiated

Lane 3: HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate with Immunising peptide at 1 µg/ml

Lane 4: Hela Whole Cell Lysate - UV Irradiated with Immunising peptide at 1 µg/ml

Lane 5: HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate with Anti-HIF-1 alpha antibody [H1alpha67] (ab1)

Lane 6 : Hela Whole Cell Lysate - UV Irradiated with Anti-HIF-1 alpha antibody [H1alpha67] (**ab1**)

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/10000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Predicted band size: 53 kDa Observed band size: 53 kDa

Exposure time: 4 minutes

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab131023 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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