abcam

Product datasheet

Anti-SIAH1 antibody ab2237

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Overview

Product name Anti-SIAH1 antibody

Description Goat polyclonal to SIAH1

Host species Goat

Tested applications Suitable for: WB, ICC/IF

Species reactivity Reacts with: Rat, Human

Predicted to work with: Mouse, Cow, Dog

Immunogen Synthetic peptide corresponding to Human SIAH1 aa 2-16 (N terminal).

Sequence:

SRQTATALPTGTSKC

Database link: Q8IUQ4

(Peptide available as ab22876)

Run BLAST with
Run BLAST with

Positive control WB: Rat liver lysate, human liver lysate. ICC/IF- U2OS cells and HepG2 cells.

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. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer pH: 7.30

Preservative: 0.02% Sodium azide Constituents: 0.05% Tris, 0.5% BSA

Purity Immunogen affinity purified

Purification notes Purified from goat serum by ammonium sulphate precipitation followed by antigen affinity

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chromatography using the immunizing peptide.

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise guarantee covers the use of ab2237 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	*****(1)	Use a concentration of 1 - 3 µg/ml. Predicted molecular weight: 31 kDa. 1 hour primary incubation at room temperature is recommended for this product.
ICC/IF		Use a concentration of 10 µg/ml.

Target

Function

E3 ubiquitin-protein ligase that mediates ubiquitination and subsequent proteasomal degradation of target proteins. E3 ubiquitin ligases accept ubiquitin from an E2 ubiquitin-conjugating enzyme in the form of a thioester and then directly transfers the ubiquitin to targeted substrates. Mediates E3 ubiquitin ligase activity either through direct binding to substrates or by functioning as the essential RING domain subunit of larger E3 complexes. Triggers the ubiquitin-mediated degradation of many substrates, including proteins involved in transcription regulation (MYB, POU2AF1, PML and RBBP8), a cell surface receptor (DCC), the cell-surface receptor-type tyrosine kinase FLT3, the cytoplasmic signal transduction molecules (KLF10/TIEG1 and NUMB), an antiapoptotic protein (BAG1), a microtubule motor protein (KIF22), a protein involved in synaptic vesicle function in neurons (SYP), a structural protein (CTNNB1) and SNCAIP. Confers constitutive instability to HIPK2 through proteasomal degradation. It is thereby involved in many cellular processes such as apoptosis, tumor suppression, cell cycle, axon guidance, transcription regulation, spermatogenesis and TNF-alpha signaling. Has some overlapping function with SIAH2. Induces apoptosis in cooperation with PEG3. Upon nitric oxid (NO) generation that follows apoptotic stimulation, interacts with S-nitrosylated GAPDH, mediating the translocation of GAPDH to the nucleus. GAPDH acts as a stabilizer of SIAH1, facilitating the degradation of nuclear proteins.

Tissue specificity

Widely expressed at a low level. Down-regulated in advanced hepatocellular carcinomas.

Pathway

Protein modification; protein ubiquitination.

Sequence similarities

Belongs to the SINA (Seven in absentia) family.

Contains 1 RING-type zinc finger. Contains 1 SIAH-type zinc finger.

Domain

The RING-type zinc finger domain is essential for ubiquitin ligase activity.

The SBD domain (substrate-binding domain) mediates the homodimerization and the interaction

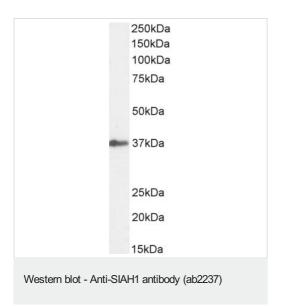
with substrate proteins. It is related to the TRAF family.

Post-translational modifications

Phosphorylated on Ser-19 by ATM and ATR. This phosphorylation disrupts SIAH1 interaction with

HIPK2, and subsequent proteasomal degradation of HIPK2.

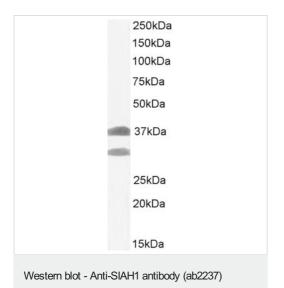
Images



Anti-SIAH1 antibody (ab2237) at 1 mg/ml + Human liver lysate at 35 μg

Predicted band size: 31 kDa **Observed band size:** 37 kDa

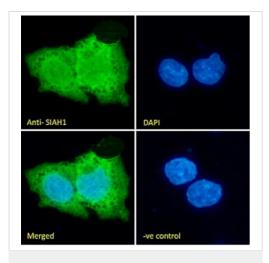
Primary incubation 1 hour at room temperature. Detected by chemiluminescence. RIPA buffer was used.



Anti-SIAH1 antibody (ab2237) at 1 μg/ml + Rat liver lysate at 35 μg

Predicted band size: 31 kDa **Observed band size:** 30, 37 kDa

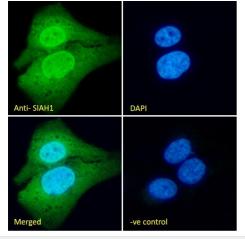
Primary incubation 1 hour at room temperature. Detected by chemiluminescence. RIPA buffer was used.



Immunocytochemistry/ Immunofluorescence - Anti-SIAH1 antibody (ab2237)

Immunofluorescence analysis of paraformaldehyde fixed, 0.15% Triton permeabilized HepG2 (human liver hepatocellular carcinoma cell) cells labelling SIAH1 with primary anti-SIAH1 antibody (ab2237) at 10ug/ml for 1 hour, followed by Alexa Fluor 488 secondary antibody at 2ug/ml. Image showing nuclear and cytoplasmic staining. The nuclear counterstain is DAPI (blue).

Negative control: Unimmunized goat IgG at 10ug/ml followed by Alexa Fluor 488 secondary antibody.



Immunocytochemistry/ Immunofluorescence - Anti-SIAH1 antibody (ab2237)

Immunofluorescence analysis of paraformaldehyde fixed, 0.15% Triton permeabilized U2OS (human bone osteosarcoma epithelial cell) cells labelling SIAH1 with primary anti-SIAH1 antibody (ab2237) at 10ug/ml for 1 hour, followed by Alexa Fluor 488 secondary antibody at 2ug/ml. Image showing nuclear and cytoplasmic staining. The nuclear counterstain is DAPI (blue).

Negative control: Unimmunized goat IgG at 10ug/ml followed by Alexa Fluor 488 secondary antibody.

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