## abcam

## Product datasheet

## Anti-SNF2H antibody [3.25(2)] ab33747

1 References 3 Images

## Overview

Product name
Description
Host species
Tested applications
Species reactivity
Immunogen
Positive control

## General notes

Anti-SNF2H antibody [3.25(2)]
Mouse monoclonal [3.25(2)] to SNF2H
Mouse
Suitable for: WB, Flow Cyt (Intra), ICC/IF
Reacts with: Mouse, Human
Fusion protein. This information is proprietary to Abcam and/or its suppliers.
This antibody gave a positive signal in the following Human whole cell lysates: HeLa, Jurkat, HepG2, MCF7 This antibody gave a negative signal in the following Mouse whole cell lysate: NIH3T3

We can conjugate this antibody to FITC for you (please see ab150233 for details).

This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or conjugation for your experiments, please contact orders@abcam.com.

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 $4^{\circ} \mathrm{C}$. Store at $+4^{\circ} \mathrm{C}$ short term (1-2 weeks). Upon delivery aliquot. Store at $-20^{\circ} \mathrm{C}$ or - |
|  | $80^{\circ} \mathrm{C}$. Avoid freeze / thaw cycle. |
| Storage buffer | $\mathrm{pH}: 7.40$ |
|  | Preservative: $0.02 \%$ Sodium azide |
|  | Constituent: PBS |
| Purity | Protein G purified |
| Clonality | Monoclonal |


| Clone number | $3.25(2)$ |
| :--- | :--- |
| Isotype | $\operatorname{lgG}$ |

## Applications

The Abpromise guarantee
Our Abpromise guarantee covers the use of ab33747 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 $10 \mu \mathrm{~g} / \mathrm{ml}$. Detects a band of approximately <br> 122 kDa (predicted molecular weight: 122 kDa ). |
| Flow Cyt (Intra) |  | Use $1 \mu \mathrm{l}$ for $10^{6}$ cells. |
| ICC/IF |  | Use a concentration of $5 \mu \mathrm{~g} / \mathrm{ml}$. |


| Target |  |
| :---: | :---: |
| Function | Helicase that possesses intrinsic ATP-dependent nucleosome-remodeling activity. Complexes containing SMARCA5 are capable of forming ordered nucleosome arrays on chromatin; this may require intact histone H 4 tails. Also required for replication of pericentric heterochromatin in S phase specifically in conjunction with BAZ1A. Probably plays a role in repression of poll dependent transcription of the rDNA locus, through the recruitment of the SIN3/HDAC1 corepressor complex to the rDNA promoter. Essential component of the WICH complex, a chromatin remodeling complex that mobilizes nucleosomes and reconfigures irregular chromatin to a regular nucleosomal array structure. The WICH complex regulates the transcription of various genes, has a role in RNA polymerase I and RNA polymerase III transcription, mediates the histone H2AX phosphorylation at 'Tyr-142', and is involved in the maintenance of chromatin structures during DNA replication processes. Essential component of the NoRC (nucleolar remodeling complex) complex, a complex that mediates silencing of a fraction of rDNA by recruiting histonemodifying enzymes and DNA methyltransferases, leading to heterochromatin formation and transcriptional silencing. |
| Tissue specificity | Ubiquitously expressed. |
| Sequence similarities | Belongs to the SNF2/RAD54 helicase family. ISWI subfamily. Contains 1 helicase ATP-binding domain. <br> Contains 1 helicase C-terminal domain. <br> Contains 2 SANT domains. |
| Developmental stage | Overexpressed in CD34-positive erythrocyte progenitor cells in acute myeloid leukemia. Downregulation correlates with hematologic remission following chemotherapy. |
| Cellular localization | Nucleus. |

Images


Immunocytochemistry/ Immunofluorescence - AntiSNF2H antibody [3.25(2)] (ab33747)


ICC/IF image of ab33747 stained HeLa cells. The cells were 4\% PFA fixed ( 10 min ) and then incubated in $1 \%$ BSA / $10 \%$ normal Goat serum / 0.3M glycine in $0.1 \%$ PBS-Tween for 1 h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab33747, $5 \mu \mathrm{~g} / \mathrm{ml}$ ) overnight at $+4^{\circ} \mathrm{C}$. The secondary antibody (green) was Alexa Fluor® 488 Goat anti-Mouse lgG (H+L) used at a 1/1000 dilution for 1 h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1 h . DAPI was used to stain the cell nuclei (blue). This antibody also gave a positive result in 4\% PFA fixed (10min) HEK293 cells, HepG2 cells at $5 \mathrm{ug} / \mathrm{ml}$ and in $100 \%$ Methanol fixed ( 5 min ) HeLa cells, HEK293 cells, HepG2 cells, and MCF-7 cells at $5 \mu \mathrm{~g} / \mathrm{ml}$.

All lanes : Anti-SNF2H antibody [3.25(2)] (ab33747) at $10 \mu \mathrm{~g} / \mathrm{ml}$

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

Lane 2 : Jurkat (Human T cell lymphoblast-like cell line) Whole Cell Lysate

Lane 3 : HepG2 (Human hepatocellular liver carcinoma cell line) Whole Cell Lysate

Lane 4 : NIH 3 T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

Lane 5 : MCF7 (Human breast adenocarcinoma cell line) Whole Cell Lysate

Lysates/proteins at $20 \mu \mathrm{~g}$ per lane.

## Secondary

All lanes: Goat polyclonal to Mouse lgG - H\&L - Pre-Adsorbed (HRP) at 1/3000 dilution

Predicted band size: 122 kDa
Observed band size: 122 kDa


Flow Cytometry (Intracellular) - Anti-SNF2H antibody [3.25(2)] (ab33747)

Overlay histogram showing HeLa cells stained with ab33747 (red line). The cells were fixed with $80 \%$ methanol ( 5 min ) and then permeabilized with $0.1 \%$ PBS-Tween for 20 min . The cells were then incubated in $1 \times$ PBS / 10\% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab33747, $1 \mu \mathrm{~g} / 1 \times 10^{6}$ cells) for 30 min at $22^{\circ} \mathrm{C}$. The secondary antibody used was DyLight® 488 goat anti-mouse lgG $(\mathrm{H}+\mathrm{L})(\mathbf{a b 9 6 8 7 9})$ at $1 / 500$ dilution for 30 min at $22^{\circ} \mathrm{C}$. Isotype control antibody (black line) was mouse $\operatorname{lgG}\left(1 \mu \mathrm{~g} / 1 \times 10^{6}\right.$ cells) used under the same conditions. Acquisition of $>5,000$ events was performed.

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

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