# abcam

# Product datasheet

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

1 References 3 Images

Overview

Product name Anti-SNF2H antibody [3.25(2)]

**Description** Mouse monoclonal [3.25(2)] to SNF2H

Host species Mouse

Tested applications Suitable for: WB, Flow Cyt (Intra), ICC/IF

Species reactivity Reacts with: Mouse, Human

**Immunogen** Fusion protein. This information is proprietary to Abcam and/or its suppliers.

Positive control 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

**General notes** We can conjugate this antibody to FITC for you (please see <u>ab150233</u> 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°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

Purity Protein G purified

**Clonality** Monoclonal

1

Clone number 3.25(2)

**Isotype** IgG

#### **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 µg/ml. Detects a band of approximately 122 kDa (predicted molecular weight: 122 kDa).
Flow Cyt (Intra)		Use 1µl for 10 <sup>6</sup> cells.
ICC/IF		Use a concentration of 5 µg/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 H4 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 histone-modifying 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. Contains 1 helicase C-terminal domain.

Contains 2 SANT domains.

**Developmental stage** 

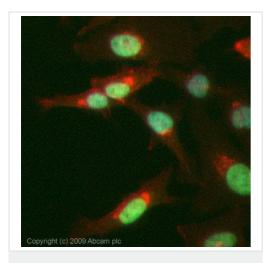
Overexpressed in CD34-positive erythrocyte progenitor cells in acute myeloid leukemia. Down-

regulation correlates with hematologic remission following chemotherapy.

**Cellular localization** 

Nucleus.

### **Images**



Immunocytochemistry/ Immunofluorescence - Anti-SNF2H antibody [3.25(2)] (ab33747)

1 2 3 4 5

460kDa —

268kDa —
238kDa —

171kDa —

117kDa —

71kDa —

41kDa —

41kDa —

31kDa —

Western blot - Anti-SNF2H antibody [3.25(2)]

Western blot - Anti-SNF2H 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 1h to permeabilise the cells and block non-specific protein-protein interactions. The cells were then incubated with the antibody (ab33747, 5µg/ml) overnight at +4°C. The secondary antibody (green) was Alexa Fluor® 488 Goat anti-Mouse IgG (H+L) used at a 1/1000 dilution for 1h. Alexa Fluor® 594 WGA was used to label plasma membranes (red) at a 1/200 dilution for 1h. 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 5µg/ml and in 100% Methanol fixed (5 min) HeLa cells, HEK293 cells, HepG2 cells, and MCF-7 cells at 5µg/ml.

**All lanes :** Anti-SNF2H antibody [3.25(2)] (ab33747) at 10 μg/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 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

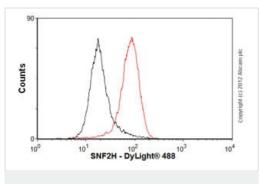
Lane 5 : MCF7 (Human breast adenocarcinoma cell line) Whole Cell Lysate

Lysates/proteins at 20 µg per lane.

## Secondary

 $\begin{tabular}{ll} \textbf{All lanes:} Goat polyclonal to Mouse $\lg G$ - H\&L - Pre-Adsorbed (HRP) at 1/3000 dilution \end{tabular}$ 

**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 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (ab33747, 1µg/1x10<sup>6</sup> cells) for 30 min at 22°C. The secondary antibody used was DyLight® 488 goat anti-mouse IgG (H+L) (ab96879) at 1/500 dilution for 30 min at 22°C. Isotype control antibody (black line) was mouse IgG (1µg/1x10<sup>6</sup> 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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