

Datasheet for ABIN933109

anti-ASH2L antibody

3 Images



Overview

Overview	
Quantity:	100 μg
Target:	ASH2L
Reactivity:	Human
Host:	Mouse
Clonality:	Monoclonal
Conjugate:	This ASH2L antibody is un-conjugated
Application:	Western Blotting (WB), Immunocytochemistry (ICC), Flow Cytometry (FACS), Dot Blot (DB)
Product Details	
Immunogen:	ASH2 L antibody was raised in mouse using recombinant Human Ash2 (Absent, Small, Or
	Homeotic)-Like (Drosophila) (Ash2)
Clone:	2046D2a
Isotype:	lgG1
Cross-Reactivity:	Human, Mouse (Murine), Rat (Rattus)
Cross-Reactivity (Details):	Other species not studied.
Purification:	Protein G affinity chromatography
Target Details	
Target:	ASH2L
Alternative Name:	ASH2L (ASH2L Products)

Target Details

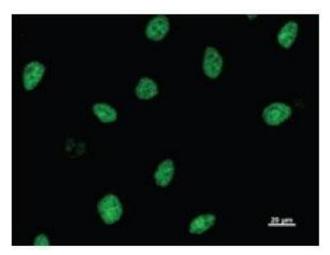
Storage Comment:

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Component of the Set1/Ash2 histone methyltransferase (HMT) complex, a complex that specifically methylates 'Lys-4' of histone H3, but not if the neighboring 'Lys-9' residue is already methylated. May function as a transcriptional regulator. May play a role in hematopoiesis. Synonyms: Monoclonal ASH2L antibody, Anti-ASH2L antibody, Ash2 histone methyltransferase complex subunit antibody, ASH2 antibody, ASH2L1 antibody, ASH2L2 antibody.

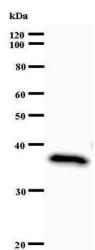
Application Details	
Application Notes:	WB: 0.2-2 μg/mL, FC: 0.5-2 μg/sample, ICC: 2-100 μg/mL Optimal conditions should be determined by the investigator.
Restrictions:	For Research Use only
Handling	
Concentration:	Lot specific
Buffer:	ASH2 L antibody in PBS (3.0 mM KCl, 1.5 mM KH2 PO4, 140 mM NaCl, 8.0 mM Na2 HPO4 (pH 7.4)) containing 1 % bovine serum albumin (BSA) and 0.05 % sodium azide (NaN3).
Preservative:	Sodium azide
Precaution of Use:	This product contains Sodium Azide: a POISONOUS AND HAZARDOUS SUBSTANCE, which should be handled by trained staff only.
Handling Advice:	Avoid repeated freeze/thaw cycles. Dilute only prior to immediate use.
Storage:	4 °C/-20 °C

Store at 2-8 °C for up to one year. We recommend long term storage at -20 °C.



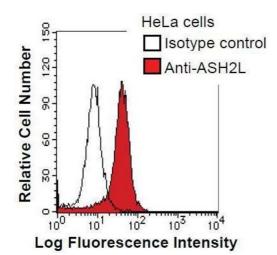
Immunofluorescence

Image 1. Immunostaining analysis in HeLa cells. HeLa cells were fixed with 4% paraformaldehyde and permeabilized with 0.01% Triton-X100 in PBS. The cells were immunostained with anti-ASH2L antibody.



Western Blotting

Image 2.



Flow Cytometry

Image 3. HeLa cells were fixed in 2% paraformaldehyde/PBS and then permeabilized in 90% methanol. Cells were stained with anti-ASH2L antibody (shaded) or isotype control (unshaded) followed by Alexa Fluor 488 conjugated goat anti-mouse IgG.