Datasheet for ABIN7139160

anti-HIST1H2AG antibody (acLys15)

2 Images

Go to Product page

Overview

Quantity:	100 µL
Target:	HIST1H2AG
Binding Specificity:	acLys15
Reactivity:	Human
Host:	Rabbit
Clonality:	Polyclonal
Conjugate:	This HIST1H2AG antibody is un-conjugated
Application:	ELISA, Immunofluorescence (IF), Immunocytochemistry (ICC)

Product Details

Immunogen:	Peptide sequence around site of Acetyl-Lys (15) derived from Human Histone H2A type 1
Isotype:	IgG
Cross-Reactivity:	Human
Purification:	Antigen Affinity Purified

Target Details

Target:	HIST1H2AG
Alternative Name:	HIST1H2AG (HIST1H2AG Products)
Background:	Background: Core component of nucleosome. Nucleosomes wrap and compact DNA into chromatin, limiting DNA accessibility to the cellular machineries which require DNA as a

Target Details

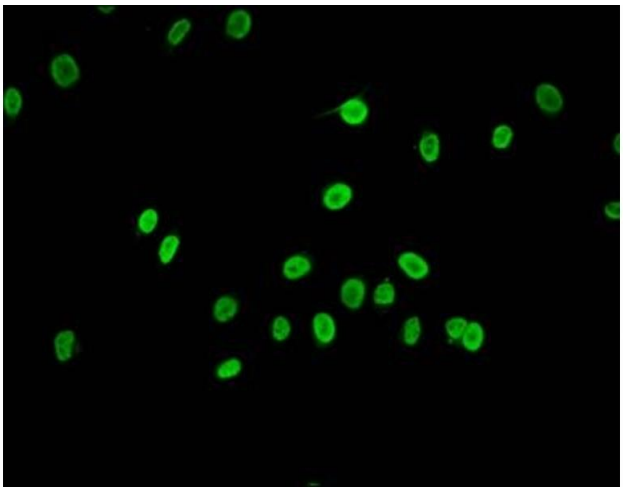
	template. Histones thereby play a central role in transcription regulation, DNA repair, DNA replication and chromosomal stability. DNA accessibility is regulated via a complex set of post-translational modifications of histones, also called histone code, and nucleosome remodeling. Aliases: H2AC11 antibody, H2AFP antibody, HIST1H2AG, antibody, H2AC13 antibody, H2AFC antibody, HIST1H2AI, antibody, H2AC15 antibody, H2AFD antibody, HIST1H2AK, antibody, H2AC16 antibody, H2AFI antibody, HIST1H2AL, antibody, H2AC17 antibody, H2AFN antibody, HIST1H2AMHistone H2A type 1 antibody, H2A.1 antibody, Histone H2A/ptl antibody
UniProt:	P0C0S8

Application Details

Application Notes:	Recommended dilution: ICC:1:1-1:10, IF:1:1-1:10,
Restrictions:	For Research Use only

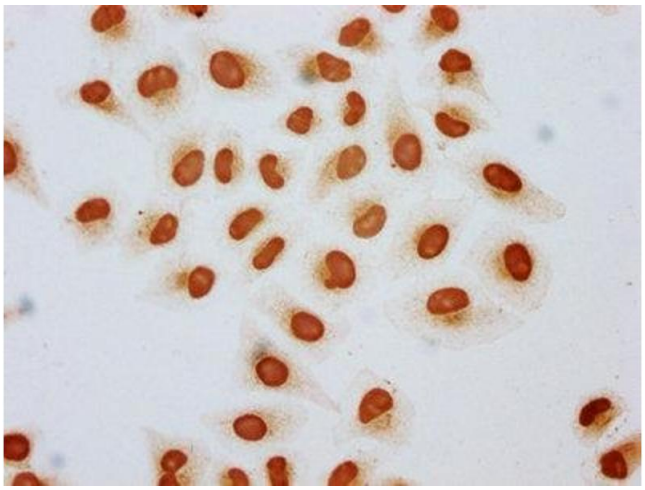
Handling

Format:	Liquid
Buffer:	Preservative: 0.03 % Proclin 300 Constituents: 50 % Glycerol, 0.01M PBS, pH 7.4
Preservative:	ProClin
Precaution of Use:	This product contains ProClin: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.
Storage:	-20 °C,-80 °C
Storage Comment:	Upon receipt, store at -20°C or -80°C. Avoid repeated freeze.



Immunofluorescence

Image 1. Immunofluorescence staining of Hela cells (treated with 30mM sodium butyrate for 4h) with ABIN7139160 at 1:1.5, counter-stained with DAPI. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked in 10% normal Goat Serum. The cells were then incubated with the antibody overnight at 4°C. The secondary antibody was Alexa Fluor 488-congugated AffiniPure Goat Anti-Rabbit IgG(H+L).



Immunocytochemistry

Image 2. Immunocytochemistry analysis of ABIN7139160 diluted at 1:3 and staining in Hela cells (treated with 30mM sodium butyrate for 4h) performed on a Leica BondTM system. The cells were fixed in 4% formaldehyde, permeabilized using 0.2% Triton X-100 and blocked with 10% normal goat serum 30min at RT. Then primary antibody (1% BSA) was incubated at 4°C overnight. The primary is detected by a biotinylated secondary antibody and visualized using an HRP conjugated SP system.

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