abcam

Product datasheet

Alexa Fluor® 568 Anti-HP1 alpha antibody [EPR5777] -Heterochromatin marker ab202533





3 Images

★★★★ 1 Abreviews

Overview

Alexa Fluor® 568 Anti-HP1 alpha antibody [EPR5777] - Heterochromatin marker **Product name**

Description Alexa Fluor® 568 Rabbit monoclonal [EPR5777] to HP1 alpha - Heterochromatin marker

Host species Rabbit

Conjugation Alexa Fluor® 568. Ex: 578nm, Em: 603nm

Tested applications Suitable for: ICC/IF **Species reactivity** Reacts with: Human

Predicted to work with: Mouse, Rat

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control ICC/IF: MCF7 cells, HAP1-CBX5 WT and KO

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit General notes

monoclonal antibodies. For details on our patents, please refer to **RabMAb® patents**.

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outlicensing@thermofisher.com.

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.

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), PBS, 1% BSA

Purity Protein A purified

Clonality Monoclonal
Clone number EPR5777

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab202533 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
ICC/IF	★★★★ (1)	1/200 - 1/500. This product gave a positive signal in MCF7 and HAP1-CBX5 cells fixed with 4% formaldehyde (10 min) and 100% methanol (5 min).

Target

Function Component of heterochromatin that recognizes and binds histone H3 tails methylated at 'Lys-9'

(H3K9me), leading to epigenetic repression. In contrast, it is excluded from chromatin when 'Tyr-41' of histone H3 is phosphorylated (H3Y41ph). Can interact with lamin-B receptor (LBR). This interaction can contribute to the association of the heterochromatin with the inner nuclear membrane. Involved in the formation of functional kinetochore through interaction with MIS12

complex proteins.

Sequence similarities Contains 2 chromo domains.

Post-translational modifications

Phosphorylation of HP1 and LBR may be responsible for some of the alterations in chromatin organization and nuclear structure which occur at various times during the cell cycle (By similarity).

Phosphorylated during interphase and possibly hyper-phosphorylated during mitosis.

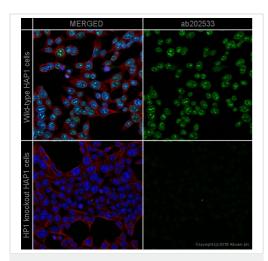
Ubiquitinated.

Cellular localization

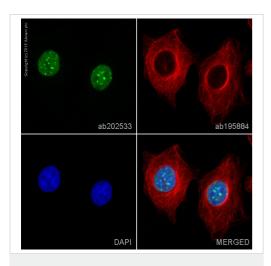
Nucleus. Chromosome. Chromosome > centromere. Component of centromeric and pericentromeric heterochromatin. Associates with chromosomes during mitosis. Associates

specifically with chromatin during metaphase and anaphase.

Images



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 568 Anti-HP1 alpha antibody [EPR5777] -Heterochromatin marker (ab202533)



Immunocytochemistry/ Immunofluorescence - Alexa
Fluor® 568 Anti-HP1 alpha antibody [EPR5777] Heterochromatin marker (ab202533)

ab202533 staining HP1 α in wild-type HAP1 cells (top panel) and HP1 α in wild-type HAP1 cells (bottom panel). The cells were fixed with 100% methanol (5min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated with ab202533 at 1/500 dilution (shown in pseudo colour green) and <u>ab7291</u> at 1 μ g/ml concentration overnight at +4 $^{\circ}$ C, followed by a further incubation at room temperature for 1h with a goat secondary antibody to Mouse μ gG (Alexa Fluor® 647) (<u>ab150119</u>) at 2 μ g/ml (shown in red). Nuclear DNA was labelled in blue with DAPI.

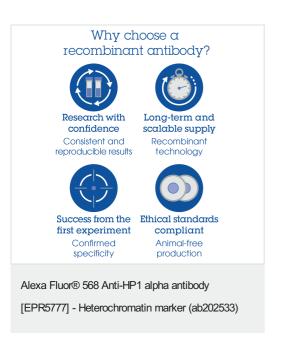
This product also gave a positive signal under the same testing conditions in HAP1 cells fixed with 4% formaldehyde (10 min).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8)

ab202533 staining HP1 alpha in MCF7 cells. The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with ab202533 at 1/200 dilution (shown in green) and ab195884, Rat monoclonal to Tubulin (Alexa Fluor® 647), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

This product also gave a positive signal under the same testing conditions in MCF7 cells fixed with 100% methanol (5min).



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