abcam

Product datasheet

Anti-HMGA2 antibody [EPR18114] - BSA and Azide free ab251468



7 Images

Overview

Product name Anti-HMGA2 antibody [EPR18114] - BSA and Azide free

Description Rabbit monoclonal [EPR18114] to HMGA2 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: WB, IHC-P, ICC/IF

Species reactivity Reacts with: Human

Immunogen Recombinant full length protein. This information is proprietary to Abcam and/or its suppliers.

General notes ab251468 is the carrier-free version of ab207301.

> Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility
- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® patents.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal
Clone number EPR18114

Isotype IgG

Applications

The Abpromise quarantee Our Abpromise quarantee covers the use of ab251468 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 at an assay dependent concentration. Predicted molecular weight: 12 kDa.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		Use at an assay dependent concentration.

Target

Function Functions as a transcriptional regulator. Functions in cell cycle regulation through CCNA2.

Involvement in disease

Note=A chromosomal aberration involving HMGA2 is associated with a subclass of benign mesenchymal tumors known as lipomas. Translocation t(3;12)(q27-q28;q13-q15) with LPP is

The serior ly mar turnors who wire as inportions. Translocation (0, 12)(q27-q20,q10-q10) with El

shown in lipomas. HMGA2 is also fused with a number of other genes in lipomas.

Note=A chromosomal aberration involving HMGA2 is associated with pulmonary chondroid hamartomas. Translocation t(3;12)(q27-q28;q14-q15) with LPP is detected in pulmonary

chondroid hamartomas.

Note=A chromosomal aberration involving HMGA2 is associated with parosteal lipomas.

Translocation t(3;12)(q28;q14) with LPP is also shown in one parosteal lipoma.

Note=A chromosomal aberration involving HMGA2 is found in uterine leiomyoma. Translocation t(12;14)(q15;q23-24) with RAD51L1. Chromosomal rearrangements involving HMGA2 do not

seem to be the principle pathobiological mechanism in uterine leiomyoma.

Sequence similarities Belongs to the HMGA family.

Contains 3 A.T hook DNA-binding domains.

Developmental stage Expressed predominantly during embryogenesis.

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Post-translational Regulated by cell cycle-dependent phosphorylation which alters its DNA binding affinity. modifications

Images



Western blot - Anti-HMGA2 antibody [EPR18114] - BSA and Azide free (ab251468)

All lanes : Anti-HMGA2 antibody [EPR18114] (**ab207301**) at 1/1000 dilution

Lane 1: HEK-293 (Human embryonic kidney epithelial cell) prepared in RIPA lysis method

Lane 2: HepG2 (Human hepatocellular carcinoma epithelial cell) prepared in RIPA lysis method

Lane 3: NCCIT (Human pluripotent embryonic carcinoma epithelial cell) prepared in 1% SDS Hot lysis method

Lysates/proteins at 20 µg per lane.

Secondary

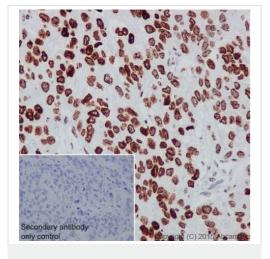
All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 12 kDa Observed band size: 18 kDa

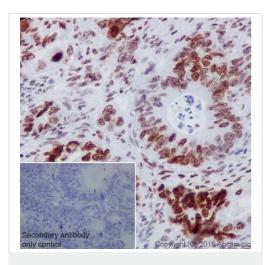
This data was developed using <u>ab207301</u>, the same antibody clone in a different buffer formulation.

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Immunohistochemical analysis of paraffin-embedded human gastric cancer tissue labeling HMGA2 with <u>ab207301</u> at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Nucleus staining on Human gastric cancer is observed. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

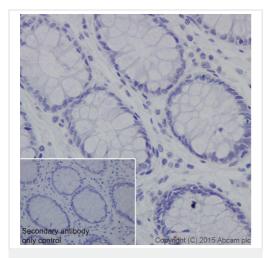


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HMGA2 antibody
[EPR18114] - BSA and Azide free (ab251468)



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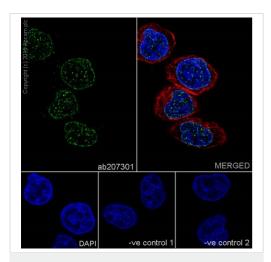
This data was developed using <u>ab207301</u>, the same antibody clone in a different buffer formulation.lmmunohistochemical analysis of paraffin-embedded human colon cancer tissue labeling HMGA2 with <u>ab207301</u> at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Nucleus staining on Human colon cancer is observed. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



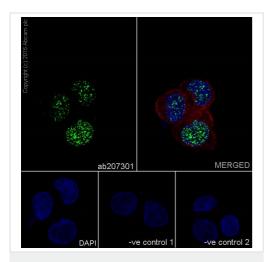
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-HMGA2 antibody

[EPR18114] - BSA and Azide free (ab251468)

This data was developed using <u>ab207301</u>, the same antibody clone in a different buffer formulation.lmmunohistochemical analysis of paraffin-embedded Human colon tissue labeling HMGA2 with <u>ab207301</u> at 1/1000 dilution, followed by Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Negative staining on Human colon tissue. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (HRP) (<u>ab97051</u>) at 1/500 dilution. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Immunocytochemistry/ Immunofluorescence - Anti-HIMGA2 antibody [EPR18114] - BSA and Azide free (ab251468)



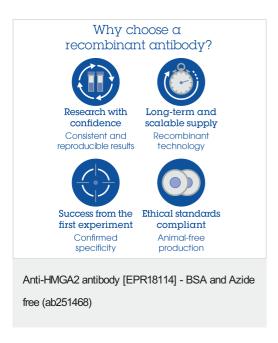
Immunocytochemistry/ Immunofluorescence - Anti-HMGA2 antibody [EPR18114] - BSA and Azide free (ab251468)

This data was developed using <u>ab207301</u>, the same antibody clone in a different buffer formulation.lmmunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells labeling HMGA2 with <u>ab207301</u> at 1/100 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (<u>ab150077</u>) secondary antibody at 1/1000 dilution (green). Confocal image showing nuclear staining on HeLa cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution and <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution (red). The negative controls are as follows:

-ve control 1: <u>ab207301</u> at 1/100 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution. -ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit IgG H&L) at 1/1000 dilution.

This data was developed using <u>ab207301</u>, the same antibody clone in a different buffer formulation.Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HepG2 (Human liver hepatocellular carcinoma) cells labeling HMGA2 with <u>ab207301</u> at 1/100 dilution, followed by Goat antirabbit lgG (Alexa Fluor® 488) (<u>ab150077</u>) secondary antibody at 1/1000 dilution (green). Confocal image showing nuclear staining on HepG2 cell line. The nuclear counter stain is DAPI (blue). Tubulin is detected with <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution and <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/1000 dilution (red).The negative controls are as follows:

-ve control 1: <u>ab207301</u> at 1/100 dilution followed by <u>ab150120</u> (Alexa Fluor[®] 594 Goat anti-Mouse secondary) at 1/1000 dilution.
-ve control 2: <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor[®] 488 Goat Anti-Rabbit IgG H&L) at 1/1000 dilution.



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