

Product datasheet

Anti-HA tag antibody - ChIP Grade ab9110

★★★★★ 51 Abreviews 459 References 6 Images

Overview

Product name	Anti-HA tag antibody - ChIP Grade
Description	Rabbit polyclonal to HA tag - ChIP Grade
Host species	Rabbit
Specificity	ELISA: The anti HA diluted 1:70.000 gave an O.D.=1.0 in a 15 minute reaction against peptide conjugated with a different carrier than used for anti peptide purification. HRP conjugated Goat anti rabbit IgG was used and TMB was the substrate.
Tested applications	Suitable for: ChIP/Chip, IP, ELISA, WB, ICC/IF, ICC, Flow Cyt, ChIP
Immunogen	YPYDVPDYA (influenza hemagglutinin-HA-epitope) conjugated to KLH.
Positive control	WB: 293FT cells transfected with 15kDa HA tagged Vpr (an HIV1 accessory protein). IP: Nuclear lysate of HEK-293T cells transiently expressing HA-tagged protein. ICC/IF: U-2 cells. Mouse olinueu cells. ChIP: Xenopus laevis oocytes were injected with mRNA for HA-tagged human BORIS.

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	Preservative: 0.1% Sodium azide Constituent: PBS
Purity	Immunogen affinity purified
Purification notes	Antibodies were immunoaffinity purified using the peptide conjugated to a solid-phase support.
Clonality	Polyclonal
Isotype	IgG

Applications

Our [Abpromise guarantee](#) covers the use of **ab9110** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

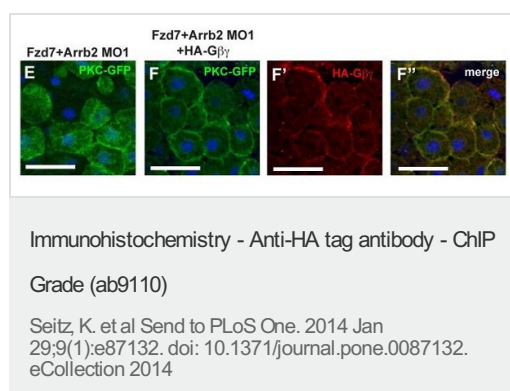
Application	Abreviews	Notes
ChIP/Chip	★★★★★	Use at an assay dependent concentration.
IP	★★★★★	Use at an assay dependent concentration.
ELISA		1/200 - 1/500.
WB	★★★★★	1/4000 - 1/10000.
ICC/IF	★★★★★	Use a concentration of 1 - 4 µg/ml.
ICC	★★★★★	Use at an assay dependent concentration.
Flow Cyt	★★★★★	Use at an assay dependent concentration. ab171870 - Rabbit polyclonal IgG, is suitable for use as an isotype control with this antibody.
ChIP	★★★★★	Use 3 µg for 25 µg of chromatin.

Target

Relevance

Human influenza hemagglutinin (HA) is a surface glycoprotein required for the infectivity of the human virus. The HA tag is derived from the HA molecule corresponding to amino acids 98-106 has been extensively used as a general epitope tag in expression vectors. Many recombinant proteins have been engineered to express the HA tag, which does not appear to interfere with the bioactivity or the biodistribution of the recombinant protein. This tag facilitates the detection, isolation, and purification of the proteins.

Images



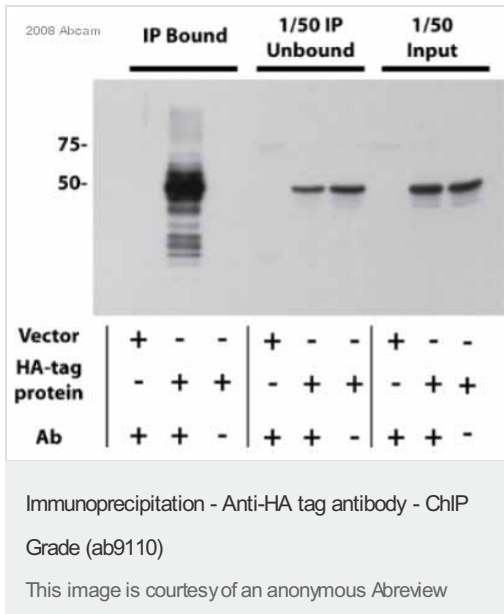
Arrb2 depends on Gβγ to induce membrane translocation of PKCα

Xenopus embryos were injected with 500 pg pkcα-gfp RNA and co-injected as indicated above the images. Animal Caps were prepared at stage 10 and immunostained as indicated. Nuclei were stained with Hoechst 33258 (blue). Images show representative results from at least two independent experiments with a minimum of six Animal Caps per experiment. Scale bars: 50 µm.

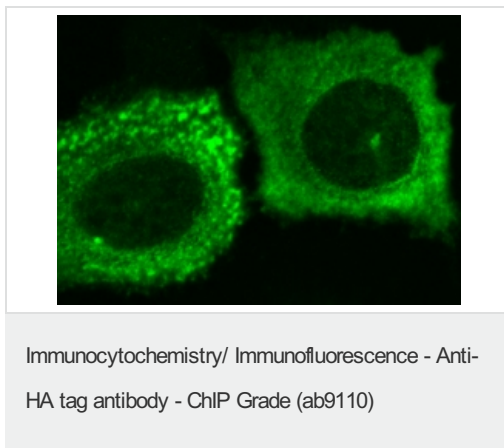
The inhibitory effect of Arrb2 MO1 (E) on PKCα-GFP membrane translocation was rescued by (F) co-injection of HA-Gβ and HA-Gγ mRNA (anti-HA (red): F', merge: F").

HA was detected with ab9110.

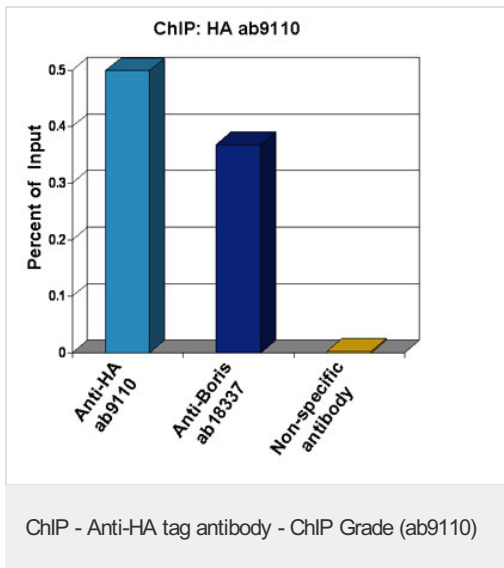
(After Figure 2 of Seitz et al)



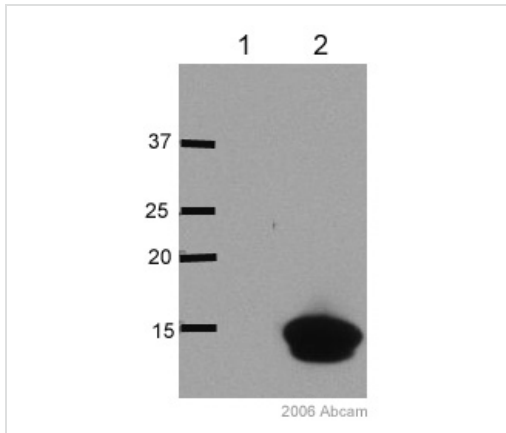
ab9110 was diluted to 4 µg/mg lysate and incubated with a nuclear lysate of HEK293T cells transiently expressing HA-tagged protein and a Protein A matrix for 2 hours at 23°C to achieve immunoprecipitation. 1000 µg of lysate was present in the input. A HRP-conjugated anti-rabbit HA monoclonal antibody diluted 1/1000 was used for the Western Blot step.



This image was kindly supplied as part of the review submitted by Kasper Fugger. Immunofluorescence staining of U-2 cells expressing HA-tagged protein with ab9110.



Xenopus laevis oocytes were injected with mRNA for HA-tagged human BORIS. Chromatin was prepared according to the Abcam X-ChIP protocol. Oocytes were fixed with formaldehyde for 10min. The ChIP was performed with 25µg of chromatin, 20µl of Protein A/G sepharose beads, and 3µg of ab9110 (anti-HA, light blue) or, 3µg of ab18337 (anti-Boris, dark blue). A non-specific antibody was used as a control (yellow). The immunoprecipitated DNA was quantified by real time PCR (Taqman approach).



Western blot - Anti-HA tag antibody - ChIP Grade (ab9110)

All lanes : Anti-HA tag antibody - ChIP Grade (ab9110) at 1/4000 dilution

Lane 1 : 15ug untransfected wcl lysate

Lane 2 : 293FT cells transfected with 15kDa HA tagged Vpr (an HIV1 accessory protein)

Secondary

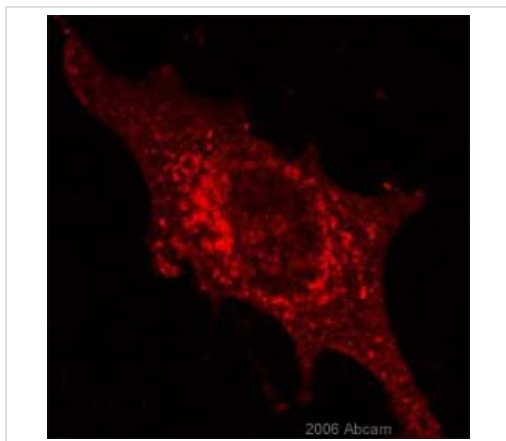
All lanes : HRP conjugated Goat anti-Rabbit

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 5 seconds

This image is courtesy of an Abreview



Immunocytochemistry/ Immunofluorescence - Anti-HA tag antibody - ChIP Grade (ab9110)

ab 9110 at a 1/200 dilution staining the mouse olinueu cell line (oligodendrocyte precursor cell) by immunocytochemistry. The antibody was incubated with the cells for 30 minutes and then detected using a Cy5 conjugated goat anti-rabbit antibody.

This image is courtesy of an Abreview submitted by **Katarina Trajkovic** on **15 March 2006**

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