abcam

Product datasheet

Rabbit IgG, polyclonal - Isotype Control (ChIP Grade) ab171870

245 References 4 Images

Overview

Product name Rabbit IgG, polyclonal - Isotype Control (ChIP Grade)

Specificity The sera for this product is un-immunized, naive sera.

Tested applications Suitable for: WB, ChIP, Flow Cyt (Intra)

General notes For more information regarding the isotype control selection, please

see https://www.abcam.com/primary-antibodies/your-guide-to-selecting-an-isotype-

control

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

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

pH: 7.40

Preservative: 0.02% Sodium azide

Constituent: PBS

Batches with a concentration less than 1 mg/ml will have 1% BSA

Purity Protein G purified

Clonality Polyclonal

Isotype IgG

1

Applications

The Abpromise guarantee

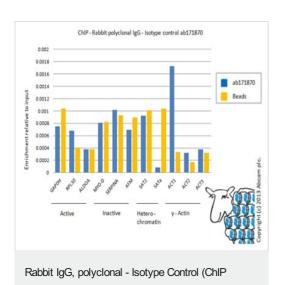
Our <u>Abpromise guarantee</u> covers the use of ab171870 in the following tested applications.

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

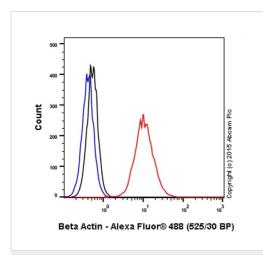
Application	Abreviews	Notes
WB		Use a concentration of 1 µg/ml.
ChIP		Use a concentration of 1 µg/ml.
Flow Cyt (Intra)		Use at an assay dependent concentration. Please note: This product should be diluted to the same concentration (not dilution) of the primary antibody to be used.

Images

Grade) (ab171870)



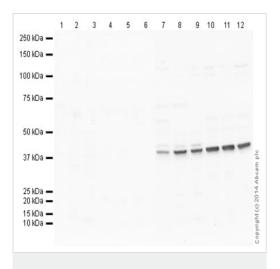
Chromatin was prepared from HeLa cells according to the Abcam X-ChIP protocol. Cells were fixed with formaldehyde for 10 minutes. The ChIP was performed with 25µg of chromatin, 2µg of ab171870 (blue), and 20µl of Protein A/G sepharose beads. No antibody was added to the beads control (yellow). The immunoprecipitated DNA was quantified by real time PCR (Taqman approach for active and inactive loci, Sybr green approach for heterochromatic loci). Primers and probes are located in the first kb of the transcribed region.



Flow Cytometry (Intracellular) - Rabbit IgG, polyclonal - Isotype Control (ChIP Grade) (ab171870) Overlay histogram showing HeLa cells stained with <u>ab75186</u> (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Triton X-100 for 20 min. The cells were then incubated in 1x PBS / 10% normal goat serum / 0.3M glycine to block non-specific protein-protein interactions followed by the antibody (<u>ab75186</u>, 0.05µg/1x10⁶ cells) for 30 min at 22°C. The secondary antibody used was Alexa Fluor® 488 goat anti-rabbit lgG (H&L) (<u>ab150081</u>) at 1/4000 dilution for 30 min at 22°C.

lsotype control antibody (black line) was rabbit lgG (polyclonal) ($\underline{ab171870}$, $0.05\mu g/1x10^6$ cells) used under the same conditions. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 20mW Argon ion laser (488nm) and 525/30 bandpass filter.



Western blot - Rabbit IgG, polyclonal - Isotype Control (ChIP Grade) (ab171870)

Lanes 1-6 : Rabbit lgG, polyclonal - Isotype Control (ChIP Grade) (ab171870) at 1 μg/ml

Lanes 7-12: Anti-beta Actin antibody (ab8227) at 1 µg/ml

Lanes 1 & 7: Human liver tissue lysate - total protein (ab29889)

Lanes 2 & 8: Liver (Mouse) Tissue Lysate

Lanes 3 & 9: Liver (Rat) Tissue Lysate

Lanes 4 & 10: HeLa (Human epithelial carcinoma cell line) Whole

Cell Lysate

Lanes 5 & 11: Jurkat (Human T cell lymphoblast-like cell line)

Whole Cell Lysate

Lanes 6 & 12: HEK293 (Human embryonic kidney cell line) Whole

Cell Lysate

Lysates/proteins at 10 µg per lane.

Secondary

All lanes : Goat Anti-Rabbit lgG H&L (HRP) (ab97051) at 1/10000 dilution

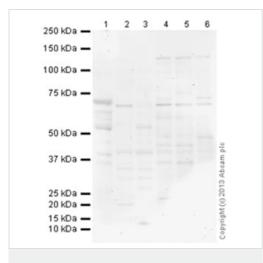
Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 2 minutes

Please note that ab171870 in lanes 1-6 represents a negative control for Beta Actin, positively seen in lanes 7-12.

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab171870 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution.



Western blot - Rabbit IgG, polyclonal - Isotype Control (ChIP Grade) (ab171870)

All lanes : Rabbit lgG, polyclonal - Isotype Control (ChIP Grade) (ab171870) at 1 µg/ml

Lane 1: Liver (Human) Tissue Lysate - adult normal tissue

Lane 2: Brain (Mouse) Tissue Lysate

Lane 3: Rat Kidney Tissue Lysate

Lane 4 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 5 : HepG2 (Human hepatocellular liver carcinoma cell line)

Lane 6: MCF7 (Human breast adenocarcinoma cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

Secondary

Whole Cell Lysate

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/10000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 20 minutes

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Bovine Serum Albumin before being incubated with ab171870 overnight at 4°C. Antibody binding was detected using an anti-rabbit antibody conjugated to HRP, and visualised using ECL development solution.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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