

Product datasheet

Anti-Rb (phospho S807) antibody [EPR17732] ab184796

Recombinant RabMAb

★★★★★ 1 Abreviews 14 References 14 Images

Overview

Product name	Anti-Rb (phospho S807) antibody [EPR17732]
Description	Rabbit monoclonal [EPR17732] to Rb (phospho S807)
Host species	Rabbit
Tested applications	Suitable for: Dot blot, WB, ICC/IF, IHC-P
Species reactivity	Reacts with: Mouse, Rat, Human
Immunogen	Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.
Positive control	WB: K562, MCF7, L-929 and C6 whole cell lysates; Mouse embryo lysate. IHC-P: Human colon, Human squamous cells carcinoma of lung, mouse spleen and rat spleen tissues. ICC/IF: MCF7 cells.
General notes	<p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb[®] patents.</p>

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 long term. Avoid freeze / thaw cycle.
Storage buffer	<p>pH: 7.2</p> <p>Preservative: 0.01% Sodium azide</p> <p>Constituents: 59% PBS, 40% Glycerol, 0.05% BSA</p>
Purity	Protein A purified
Clonality	Monoclonal
Clone number	EPR17732

Isotype

IgG

Applications

The Abpromise guarantee

Our [Abpromise guarantee](#) covers the use of ab184796 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
Dot blot		1/1000.
WB		1/1000. Detects a band of approximately 106 kDa (predicted molecular weight: 106 kDa).
ICC/IF	★★★★★ (1)	1/500.
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

Target

Function

Key regulator of entry into cell division that acts as a tumor suppressor. Promotes G0-G1 transition when phosphorylated by CDK3/cyclin-C. Acts as a transcription repressor of E2F1 target genes. The underphosphorylated, active form of RB1 interacts with E2F1 and represses its transcription activity, leading to cell cycle arrest. Directly involved in heterochromatin formation by maintaining overall chromatin structure and, in particular, that of constitutive heterochromatin by stabilizing histone methylation. Recruits and targets histone methyltransferases SUV39H1, KMT5B and KMT5C, leading to epigenetic transcriptional repression. Controls histone H4 'Lys-20' trimethylation. Inhibits the intrinsic kinase activity of TAF1. Mediates transcriptional repression by SMARCA4/BRG1 by recruiting a histone deacetylase (HDAC) complex to the c-FOS promoter. In resting neurons, transcription of the c-FOS promoter is inhibited by BRG1-dependent recruitment of a phospho-RB1-HDAC1 repressor complex. Upon calcium influx, RB1 is dephosphorylated by calcineurin, which leads to release of the repressor complex (By similarity). In case of viral infections, interactions with SV40 large T antigen, HPV E7 protein or adenovirus E1A protein induce the disassembly of RB1-E2F1 complex thereby disrupting RB1's activity.

Tissue specificity

Expressed in the retina.

Involvement in disease

Childhood cancer retinoblastoma
Bladder cancer
Osteogenic sarcoma

Sequence similarities

Belongs to the retinoblastoma protein (RB) family.

Domain

The Pocket domain binds to the threonine-phosphorylated domain C, thereby preventing interaction with heterodimeric E2F/DP transcription factor complexes.

Post-translational modifications

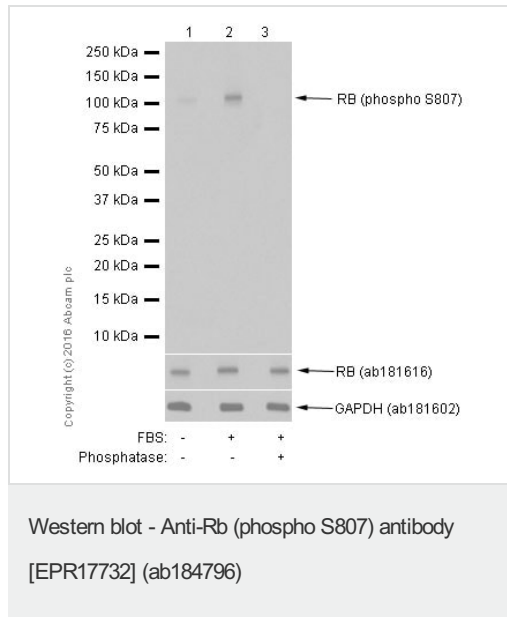
Phosphorylated by CDK6 and CDK4, and subsequently by CDK2 at Ser-567 in G1, thereby releasing E2F1 which is then able to activate cell growth. Dephosphorylated at the late M phase. SV40 large T antigen, HPV E7 and adenovirus E1A bind to the underphosphorylated, active form of pRb. Phosphorylation at Thr-821 and Thr-826 promotes interaction between the C-terminal domain C and the Pocket domain, and thereby inhibits interactions with heterodimeric E2F/DP transcription factor complexes. Dephosphorylated at Ser-795 by calcineurin upon calcium

stimulation. CDK3/cyclin-C-mediated phosphorylation at Ser-807 and Ser-811 is required for G0-G1 transition. Phosphorylated by CDK1 and CDK2 upon TGFB1-mediated apoptosis. N-terminus is methylated by METTL11A/NTM1 (By similarity). Monomethylation at Lys-810 by SMYD2 enhances phosphorylation at Ser-807 and Ser-811, and promotes cell cycle progression. Monomethylation at Lys-860 by SMYD2 promotes interaction with L3MBTL1. Acetylation at Lys-873 and Lys-874 regulates subcellular localization, at least during keratinocytes differentiation.

Cellular localization

Nucleus.

Images



All lanes : Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/20000 dilution

Lane 1 : Jurkat (human T cell leukemia cell line from peripheral blood) serum starved for 44 h. Whole cell lysates

Lane 2 : Jurkat (human T cell leukemia cell line from peripheral blood) serum starved for 24 h. Then 10% FBS incubated for 20 h. Whole cell lysates

Lane 3 : Jurkat (human T cell leukemia cell line from peripheral blood) serum starved for 24 h. Then 10% FBS incubated for 20 h. Whole cell lysates. Then the membrane was incubated with phosphatase.

Lysates/proteins at 15 µg per lane.

Secondary

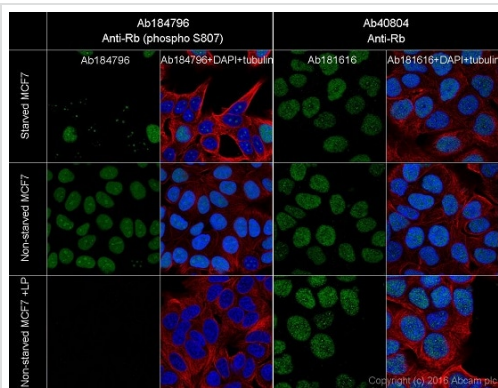
All lanes : Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 10 seconds

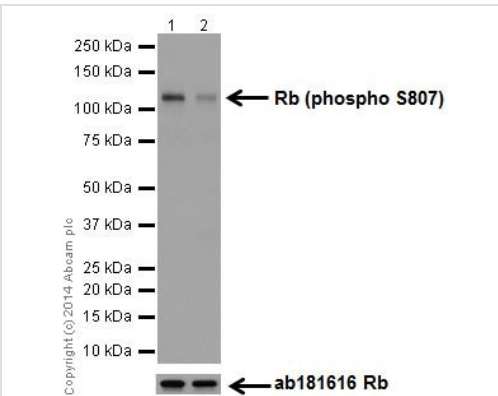
Blocking and diluting buffer and concentration: 5% NFDm/TBST.



Immunocytochemistry/ Immunofluorescence - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% tritonX-100 permeabilized MCF7 (human breast adenocarcinoma cell line) cells, Serum starved and non-starved, labeling Rb (phospho S807) with Ab184796 at 1/200 dilution followed by Goat anti-Rabbit secondary IgG AlexaFluor@488 (ab150077) secondary antibody at 1/1000 dilution (green).

Confocal image showing positive staining on MCF7 cells. The number of positive cells increased after treatment with FBS (fetal bovine serum) for 48 hours, then decreased after Lambda Protein Phosphatase treatment (311 for 2 hours).



Western blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

All lanes : Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/20000 dilution

Lane 1 : K562 (human chronic myelogenous leukemia cells from bone marrow) whole cell lysates

Lane 2 : K562 (human chronic myelogenous leukemia cells from bone marrow) whole cell lysates serum starved for 2 days

Lysates/proteins at 10 µg per lane.

Secondary

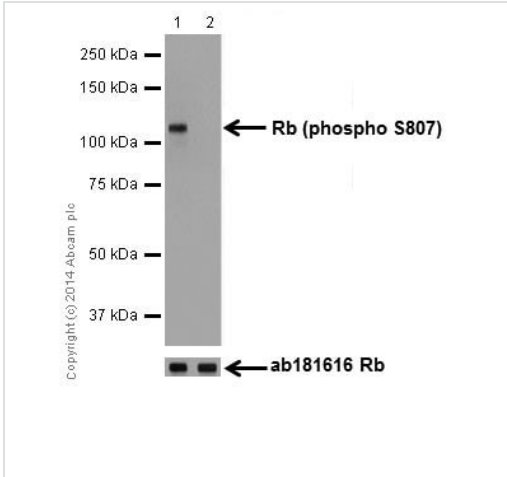
All lanes : Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDm/TBST.



Western blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

All lanes : Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/5000 dilution

Lane 1 : K562 (human chronic myelogenous leukemia cells from bone marrow) whole cell lysates

Lane 2 : K562 (human chronic myelogenous leukemia cells from bone marrow) whole cell lysates treated with Lambda Phosphatase.

Lysates/proteins at 10 µg per lane.

Secondary

Lane 1 : Goat Anti-Rabbit IgG (H+L), Peroxidase conjugated at 1/1000 dilution

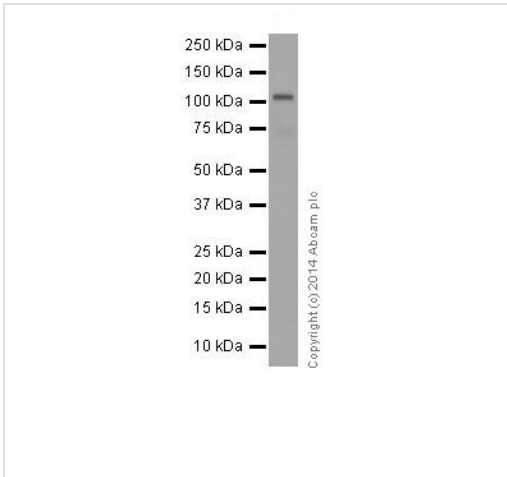
Lane 2 : Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDm/TBST.



Western blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/1000 dilution + L-929 (mouse connective tissue fibroblast cell line) whole cell lysate at 10 µg

Secondary

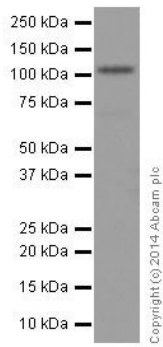
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDm/TBST.



Western blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/2000 dilution + Mouse embryo lysate at 10 μ g

Secondary

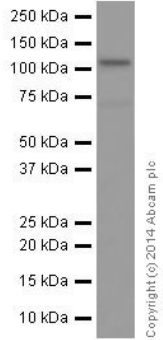
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/2000 dilution + C6 (rat glial tumor cell line) whole cell lysate at 10 μ g

Secondary

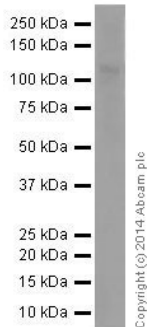
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 30 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Anti-Rb (phospho S807) antibody [EPR17732] (ab184796) at 1/1000 dilution + MCF7 (human breast adenocarcinoma cell line) whole cell lysate at 10 µg

Secondary

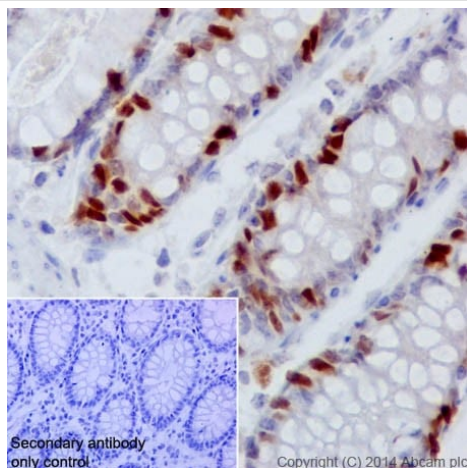
Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at 1/1000 dilution

Predicted band size: 106 kDa

Observed band size: 106 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.

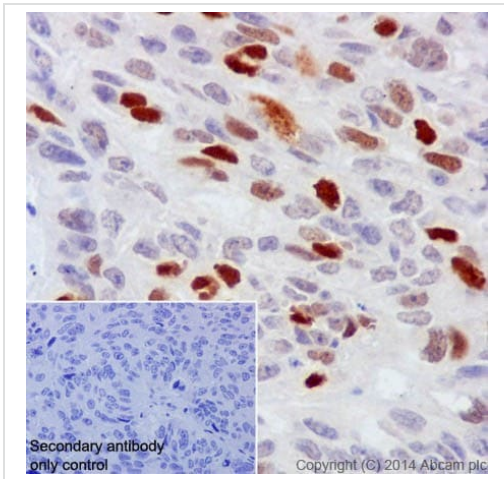


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Immunohistochemical analysis of paraffin-embedded Human colon tissue labeling Rb (phospho S807) with ab184796 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) secondary antibody at 1/500 dilution. Nucleus staining on epithelial cells of Human colon 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) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

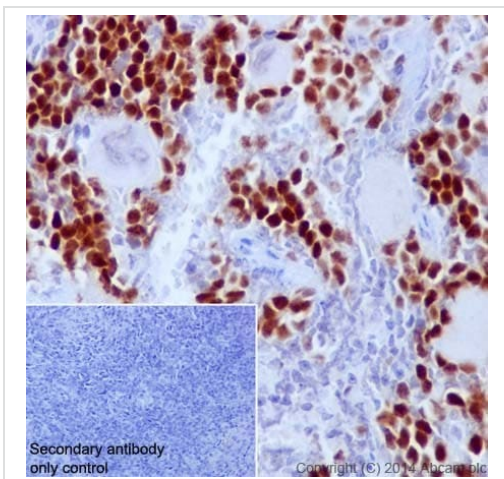


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Immunohistochemical analysis of paraffin-embedded human Squamous cells carcinoma of lung tissue labeling Rb (phospho S807) with ab184796 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) secondary antibody at 1/500 dilution. Nucleus staining on squamous cells carcinoma of lung 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) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

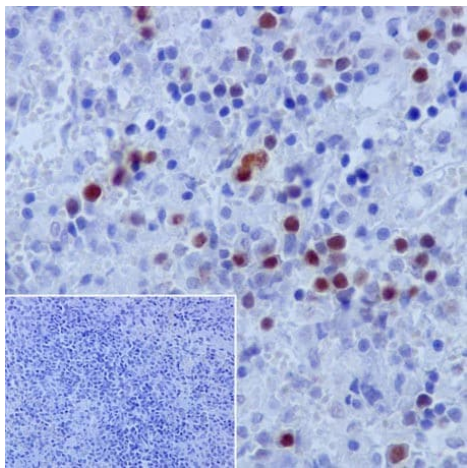


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Immunohistochemical analysis of paraffin-embedded mouse spleen tissue labeling Rb (phospho S807) with ab184796 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) secondary antibody at 1/500 dilution. Nucleus staining on lymphocytes of mouse spleen 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) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

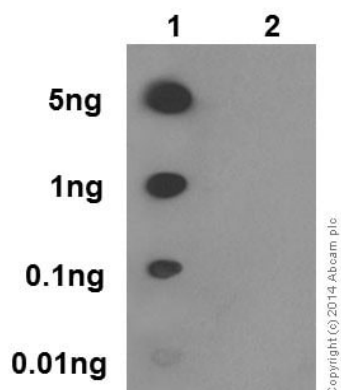


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Immunohistochemical analysis of paraffin-embedded rat spleen tissue labeling Rb (phospho S807) with ab184796 at 1/500 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) secondary antibody at 1/500 dilution. Nucleus staining on lymphocytes of rat spleen is observed. Counter stained with Hematoxylin.

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



Dot Blot - Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

Dot blot analysis of Rb (phospho S807) peptide (Lane 1), and non-phospho peptide (Lane 2), labeled using ab184796 at 1/1000 dilution, followed by Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated secondary antibody at 1/1000 dilution.

Blocking/Dilution buffer: 5% NFDm/TBST.

Why choose a recombinant antibody?



Research with confidence
Consistent and reproducible results



Long-term and scalable supply
Recombinant technology



Success from the first experiment
Confirmed specificity



Ethical standards compliant
Animal-free production

Anti-Rb (phospho S807) antibody [EPR17732] (ab184796)

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

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