abcam

Product datasheet

Anti-JunD antibody [EPR17365] ab181615



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Overview

Product name Anti-JunD antibody [EPR17365]

Rabbit monoclonal [EPR17365] to JunD **Description**

Host species Rabbit

Tested applications Suitable for: WB, IHC-P, ICC/IF Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control WB: 293T, Jurkat, HeLa, C6, Raw264.7, PC12, NIH 3T3 cell lysates; Human fetal liver, fetal brain,

fetal heart and fetal kidney lysates. IHC-P: Human mammary gland tissue and lung squamous cell

carcinoma tissue; Mouse and rat cerebral cortex tissues. ICC/IF: HeLa cells.

General notes 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 short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle.

Preservative: 0.01% Sodium azide Storage buffer

Constituents: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA

Purity Protein A purified

Clonality Monoclonal Clone number EPR17365

Isotype lgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab181615 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		1/1000. Detects a band of approximately 39,42 kDa (predicted molecular weight: 35 kDa).
IHC-P		1/1000. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/10000.

Target

Function Transcription factor binding AP-1 sites.

Sequence similarities Belongs to the bZIP family. Jun subfamily.

Contains 1 bZIP domain.

Cellular localization Nucleus.

Images



Western blot - Anti-JunD antibody [EPR17365] (ab181615)

All lanes: Anti-JunD antibody [EPR17365] (ab181615) at 1/1000

dilution

Lane 1: 293T (Human epithelial cells from embryonic kidney)

whole cell lysate at 20 µg

Lane 2: Jurkat (Human T cell leukemia cells from peripheral blood)

whole cell lysate at 20 µg

Lane 3: Human fetal liver lysate at 10 µg

Secondary

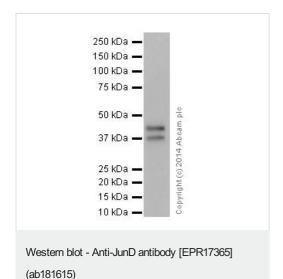
All lanes: Goat Anti-Rabbit IgG, (H+L), Peroxidase conjugated at

1/1000 dilution

Predicted band size: 35 kDa **Observed band size:** 39,42 kDa

Exposure time: 3 minutes

5% NFDM/TBST: Blocking and diluting buffer.



Anti-JunD antibody [EPR17365] (ab181615) at 1/1000 dilution + HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate at 20 µg

Secondary

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

Predicted band size: 35 kDa Observed band size: 39,42 kDa

Exposure time: 5 seconds

1 2 250 kDa -150 kDa -



Western blot - Anti-JunD antibody [EPR17365] (ab181615)

5% NFDM/TBST: Blocking and diluting buffer.

All lanes: Anti-JunD antibody [EPR17365] (ab181615) at 1/1000 dilution

Lane 1: Human fetal brain lysate Lane 2: Human fetal heart lysate Lane 3: Human fetal kidney lysate

Lysates/proteins at 10 µg per lane.

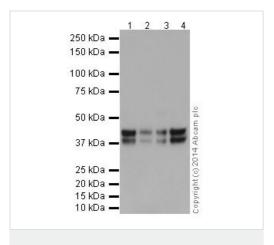
Secondary

All lanes: Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG at 1/1000 dilution

Predicted band size: 35 kDa Observed band size: 39,42 kDa

Exposure time: 1 minute

5% NFDM/TBST: Blocking and diluting buffer.



Western blot - Anti-JunD antibody [EPR17365] (ab181615)

All lanes : Anti-JunD antibody [EPR17365] (ab181615) at 1/1000 dilution

Lane 1: C6 (Rat glial tumor cells) whole cell lysate

Lane 2: Raw264.7 (Mouse macrophage cells transformed with

Abelson murine leukemia virus) whole cell lysate

Lane 3: PC12 (Rat adrenal gland pheochromocytoma) whole cell lysate

Lane 4: NIH 3T3 (Mouse embyro fibroblast cells) whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

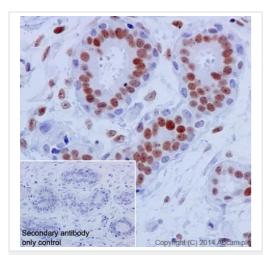
All lanes : Goat Anti-Rabbit lgG, (H+L),Peroxidase conjugated at 1/1000 dilution

Predicted band size: 35 kDa

Observed band size: 39.42 kDa

Exposure time: 30 seconds

5% NFDM/TBST: Blocking and diluting buffer.

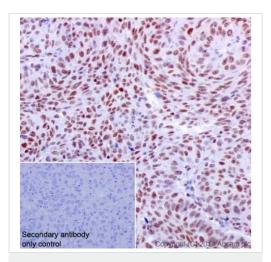


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-JunD antibody
[EPR17365] (ab181615)

Immunohistochemical analysis of paraffin-embedded Human mammary gland tissue labeling JunD using ab181615 at 1/1000 dilution. A Goat Anti-Rabbit IgG H&L (HRP) (ab97051) was used as secondary at 1/500 dilution. Counterstain: Hematoxylin. Inset image: negative control obtained using PBS instead of ab181615 and secondary antibody only.

Note: Nuclear staining on the epithelial cells of Human mammary gland was observed.

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

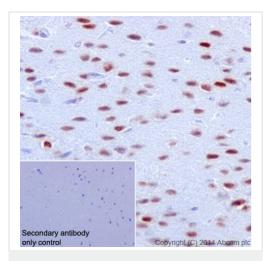


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-JunD antibody
[EPR17365] (ab181615)

Immunohistochemical analysis of paraffin-embedded Human lung squamous cell carcinoma tissue labeling JunD using ab181615 at 1/1000 dilution. A Goat Anti-Rabbit IgG H&L (HRP) (ab97051) was used as secondary at 1/500 dilution. Counterstain: Hematoxylin. Inset image: negative control obtained using PBS instead of ab181615 and secondary antibody only.

Note: Nucleus staining on the cancer cells of lung squamous cell carcinoma was observed.

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

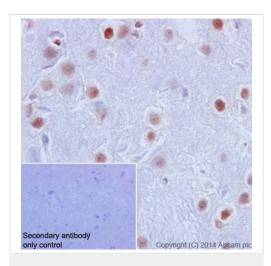


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-JunD antibody
[EPR17365] (ab181615)

Immunohistochemical analysis of paraffin-embedded mouse cerebral cortex tissue labeling JunD using ab181615 at 1/1000 dilution. A Goat Anti-Rabbit IgG H&L (HRP) (ab97051) was used as secondary at 1/500 dilution. Counterstain: Hematoxylin. Inset image: negative control obtained using PBS instead of ab181615 and secondary antibody only.

Note: Nuclear staining on neurons of the mouse cerebral cortex was observed.

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

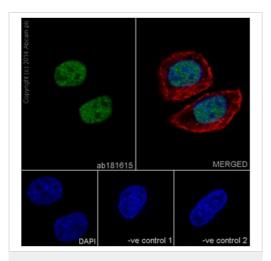


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-JunD antibody
[EPR17365] (ab181615)

Immunohistochemical analysis of paraffin-embedded Rat cerebral cortex tissue labeling JunD using ab181615 at 1/1000 dilution. A Goat Anti-Rabbit IgG H&L (HRP) (ab97051) was used as secondary at 1/500 dilution. Counterstain: Hematoxylin. Inset image: negative control obtained using PBS instead of ab181615 and secondary antibody only.

Note: Nuclear staining on neurons of the rat cerebral cortex was observed.

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

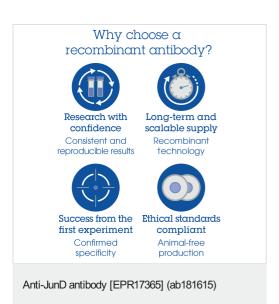


Immunocytochemistry/ Immunofluorescence - Anti-JunD antibody [EPR17365] (ab181615)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human epithelial cells from cervix adenocarcinoma) cells, labeling JunD with ab181615 at 1/10000 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Confocal image shows nuclear staining on the HeLa cell line. The nuclear counter stain is DAPI (blue) . Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

The negative controls are as follows:

- 1. ab181615 at 1/10000 dilution followed by ab150120
 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution.
- 2. <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/500 dilution.



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