abcam

Product datasheet

Anti-IKZF3 antibody [EPR9342(B)] - BSA and Azide free ab192678



1 References 9 Images

Overview

Product name Anti-IKZF3 antibody [EPR9342(B)] - BSA and Azide free

Description Rabbit monoclonal [EPR9342(B)] to IKZF3 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: Flow Cyt (Intra), ICC/IF, WB, IHC-P

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control Human thymus, Jurkat, Raji, and Ramos cell lysates; Human spleen tissue.

General notes ab192678 is the carrier-free version of <u>ab139408</u>.

Our <u>carrier-free</u> 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 cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our <u>conjugation kits</u> 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**.

1

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

ClonalityMonoclonalClone numberEPR9342(B)

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab192678 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
Flow Cyt (Intra)		Use at an assay dependent concentration. ab199376 - Rabbit monoclonal lgG, is suitable for use as an isotype control with this antibody.
ICC/IF		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 58 kDa. Can be blocked with IKZF3 peptide (ab184024).
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. See IHC antigen retrieval protocols.

Target

Function Transcription factor that plays an important role in the regulation of lymphocyte differentiation.

Plays an essential role in regulation of B-cell differentiation, proliferation and maturation to an effector state. Involved in regulating BCL2 expression and controlling apoptosis in T-cells in an

L2-dependent manner.

Tissue specificity Expressed most strongly in peripheral blood leukocytes, the spleen, and the thymus.

Sequence similaritiesBelongs to the lkaros C2H2-type zinc-finger protein family.

Contains 6 C2H2-type zinc fingers.

Post-translational

modifications

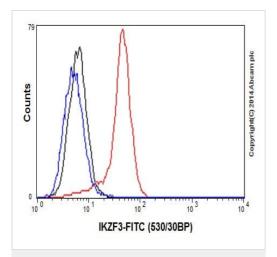
Phosphorylation on tyrosine residues induced by IL2 is required for dissociation from HRAS and

nuclear translocation of IKZF3 in T-cells. Phosphorylation on tyrosine residues induced by IL4 is

required for dissociation from Bcl-X(L) in T-cells.

Cellular localization Nucleus. Cytoplasm.

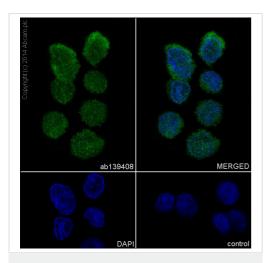
Images



Flow Cytometry (Intracellular) - Anti-IKZF3 antibody [EPR9342(B)] - BSA and Azide free (ab192678)

Intracellular Flow Cytometry analysis of Ramos cells labelling IKZF3 with purified **ab139408** at 1/90 (red). Cells were fixed with 2% paraformaldehyde. A FITC-conjugated goat anti-rabbit IgG (1/150) was used as the secondary antibody. Black - Isotype control, rabbit monoclonal IgG. Blue - Unlabelled control, cells without incubation with primary and secondary antibodies.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab139408).

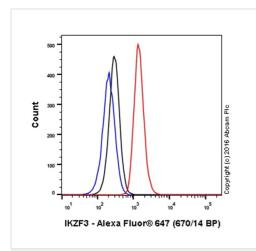


Immunocytochemistry/ Immunofluorescence - Anti-IKZF3 antibody [EPR9342(B)] - BSA and Azide free (ab192678)

Immunocytochemistry/Immunofluorescence analysis of Jurkat cells labelling IKZF3 (green) with purified <u>ab139408</u> at 1/100. Cells were fixed with 4% paraformaldehyde and permeabilized with 0.1% Triton X-100. <u>ab150077</u>, an Alexa Fluor[®] 488-conjugated goat antirabbit IgG (1/500) was used as the secondary antibody. DAPI (blue) was used as the nuclear counterstain.

Control: primary antibody (1/100) and secondary antibody **ab150120**, an Alexa Fluor[®] 594-conjugated goat anti-mouse lgG (1/500).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab139408</u>).



Flow Cytometry (Intracellular) - Anti-IKZF3 antibody [EPR9342(B)] - BSA and Azide free (ab192678)

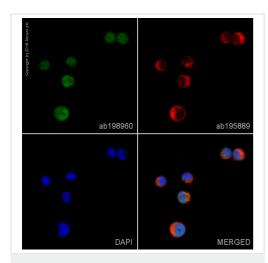
Clone EPR9342 (B) (ab192678) has been successfully conjugated by Abcam. This image was generated using Anti-IKZF3 antibody [EPR9342 (B)] (Alexa Fluor[®] 647). Please refer to **ab198962** for protocol details.

Overlay histogram showing K562 cells stained with <u>ab198962</u> (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Triton X-100 for 15 min. The cells were then incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (<u>ab198962</u>, 1/500 dilution) for 30 min at 22°C.

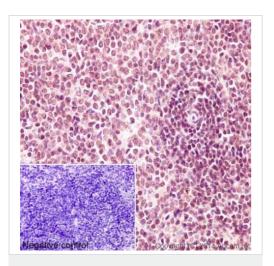
Isotype control antibody (black line) was Rabbit IgG (monoclonal) Alexa Fluor[®] 647 (<u>ab199093</u>) used at the same concentration and conditions as the primary antibody. Unlabelled sample (blue line) was also used as a control.

Acquisition of >5,000 events were collected using a 40 mW Red laser (640nm) and 670/14 bandpass filter.

This antibody gave a positive signal in K562 cells fixed with 4% formaldehyde (10 min)/permeabilized with 0.1% PBS-Triton X-100 for 15 min used under the same conditions.



Immunocytochemistry/ Immunofluorescence - Anti-IKZF3 antibody [EPR9342(B)] - BSA and Azide free (ab192678)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IKZF3 antibody

[EPR9342(B)] - BSA and Azide free (ab192678)

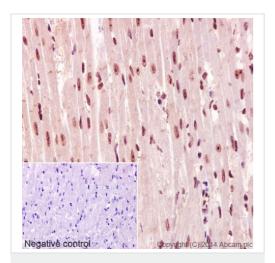
Clone EPR9342(B) (ab192678) has been successfully conjugated by Abcam. This image was generated using Anti-IKZF3 antibody [EPR9342(B)] (Alexa Fluor® 488). Please refer to **ab198960** for protocol details.

<u>ab198960</u> staining IKZF3 in Jurkat cells. The cells were fixed with 4% formaldehyde (10 min), permeabilized with 0.1% Triton X-100 for 5 minutes and then blocked with 1% BSA/10% normal goat serum/0.3M glycine in 0.1% PBS-Tween for 1h. The cells were then incubated overnight at +4°C with <u>ab198960</u> at 1/100 dilution (shown in green) and <u>ab195889</u>, Mouse monoclonal to alpha Tubulin (Alexa Fluor[®] 594), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of rat spleen tissue labelling IKZF3 with purified **ab139408** at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. **ab97051**, a HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab139408).

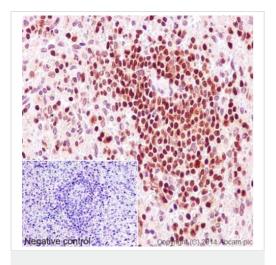


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IKZF3 antibody

[EPR9342(B)] - BSA and Azide free (ab192678)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of mouse cardiac muscle tissue labelling IKZF3 with purified ab139408 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. ab97051, a HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

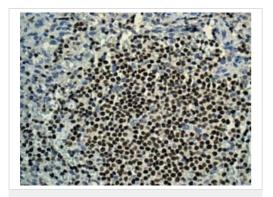
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab139408).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IKZF3 antibody
[EPR9342(B)] - BSA and Azide free (ab192678)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human spleen tissue labelling IKZF3 with purified ab139408 at 1/100. Heat mediated antigen retrieval was performed using Tris/EDTA buffer pH 9. ab97051, a HRP-conjugated goat anti-rabbit IgG (H+L) was used as the secondary antibody (1/500). Negative control using PBS instead of primary antibody. Counterstained with hematoxylin.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab139408).



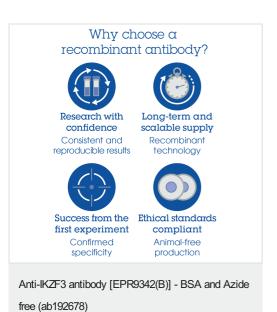
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IKZF3 antibody

[EPR9342(B)] - BSA and Azide free (ab192678)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of human spleen tissue labelling IKZF3 with unpurified <u>ab139408</u> at 1/250 dilution.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab139408</u>).

Heat mediated antigen retrieval was performed with citrate buffer pH 6 before commencing with IHC staining protocol.



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

Our Abpromise to you: Quality guaranteed and expert technical support

- Replacement or refund for products not performing as stated on the datasheet
- · Valid for 12 months from date of delivery
- Response to your inquiry within 24 hours
- We provide support in Chinese, English, French, German, Japanese and Spanish
- Extensive multi-media technical resources to help you
- We investigate all quality concerns to ensure our products perform to the highest standards

If the product does not perform as described on this datasheet, we will offer a refund or replacement. For full details of the Abpromise, please visit https://www.abcam.com/abpromise or contact our technical team.

Terms and conditions

8