abcam

Product datasheet

Anti-IRF8 antibody [EPR20441] - BSA and Azide free ab251475



6 Images

Overview

Product name Anti-IRF8 antibody [EPR20441] - BSA and Azide free

Description Rabbit monoclonal [EPR20441] to IRF8 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: WB, IP, IHC-P

Reacts with: Human Species reactivity

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

General notes ab251475 is the carrier-free version of ab207418.

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

Use our conjugation kits 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.

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

Clonality Monoclonal
Clone number EPR20441

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab251475 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 at an assay dependent concentration. Detects a band of approximately 48 kDa (predicted molecular weight: 48 kDa).
IP		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

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Function Specifically binds to the upstream regulatory region of type I IFN and IFN-inducible MHC class I

genes (the interferon consensus sequence (ICS)). Plays a negative regulatory role in cells of the

immune system.

Tissue specificity Predominantly in lymphoid tissues.

Sequence similarities Belongs to the IRF family.

Contains 1 IRF tryptophan pentad repeat DNA-binding domain.

Post-translational

modifications

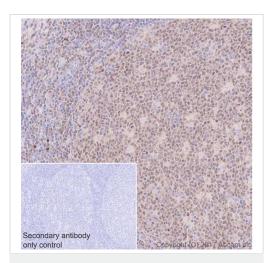
Ubiquitinated. Ubiquitination by TRIM21 in macrophages, a process that is strongly increased

upon interferon gamma stimulation, leds to the enhanced transcriptional activity of target cytokine

genes.

Cellular localization Nucleus.

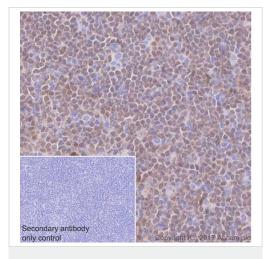
Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-IRF8 antibody

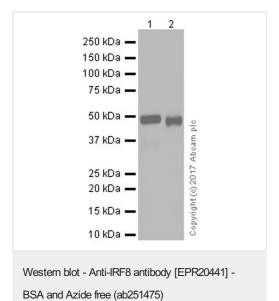
[EPR20441] - BSA and Azide free (ab251475)

This data was developed using <u>ab207418</u>, the same antibody clone in a different buffer formulation.Immunohistochemical analysis of paraffin-embedded human tonsil tissue labeling IRF8 with <u>ab207418</u> at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use. Nuclear staining on human tonsil (×200) [PMID: 18663414] [PMID: 16380510]. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) Ready to use. 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-IRF8 antibody
[EPR20441] - BSA and Azide free (ab251475)

This data was developed using <u>ab207418</u>, the same antibody clone in a different buffer formulation.lmmunohistochemical analysis of paraffin-embedded human non-hodgkin's lymphoma tissue labeling IRF8 with <u>ab207418</u> at 1/1000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) ready to use. Nuclear staining on human non-hodgkin's lymphoma. Counter stained with Hematoxylin. Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ready to use. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.



All lanes : Anti-IRF8 antibody [EPR20441] (**ab207418**) at 1/1000 dilution

Lane 1 : THP-1 (Human monocytic leukemia cell line) whole cell lysate

Lane 2 : Raji (Human Burkitt's lymphoma cell line) whole cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

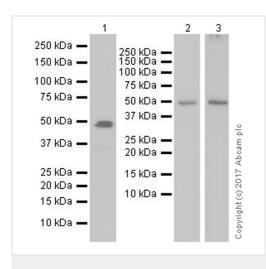
All lanes : Goat Anti-Rabbit $\lg G \ H\&L \ (HRP) \ (\underline{ab97051})$ at 1/100000 dilution

Predicted band size: 48 kDa **Observed band size:** 48 kDa

Exposure time: 1 minute

This data was developed using <u>ab207418</u>, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.



Western blot - Anti-IRF8 antibody [EPR20441] - BSA and Azide free (ab251475)

All lanes : Anti-IRF8 antibody [EPR20441] (<u>ab207418</u>) at 1/1000 dilution

Lane 1: Human fetal spleen lysate

Lane 2 : Human tonsil lysate
Lane 3 : Human colon lysate

Lysates/proteins at 10 µg per lane.

Secondary

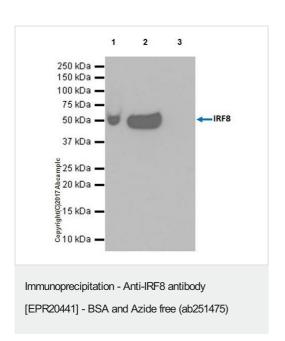
All lanes : VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) at 1/1000 dilution

Predicted band size: 48 kDa Observed band size: 48 kDa

This data was developed using <u>ab207418</u>, the same antibody clone in a different buffer formulation.

Blocking and dilution buffer: 5% NFDM/TBST.

Exposure time: Lane 1: 30 seconds; Lane 2: 2 seconds; Lane 3: 15 seconds.



This data was developed using <u>ab207418</u>, the same antibody clone in a different buffer formulation.IRF8 was immunoprecipitated from 0.35 mg of Raji (Human Burkitt's lymphoma cell line) whole cell lysate with <u>ab207418</u> at 1/40 dilution. Western blot was performed from the immunoprecipitate using <u>ab207418</u> at 1/500 dilution. VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>), was used for detection at 1/1000 dilution. Lane 1: Raji whole cell lysate 10 μ g (Input). Lane 2: <u>ab207418</u> IP in Raji whole cell lysate. Lane 3: Rabbit monoclonal IgG (<u>ab172730</u>) instead of <u>ab207418</u> in Raji whole cell lysate. Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 1 second.



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