abcam

Product datasheet

Alexa Fluor® 488 Anti-TRIM25/EFP antibody [EPR7315] ab200781

Recombinant RabMAb

3 Images

Overview

Product name Alexa Fluor® 488 Anti-TRIM25/EFP antibody [EPR7315]

Description Alexa Fluor® 488 Rabbit monoclonal [EPR7315] to TRIM25/EFP

Host species Rabbit

Conjugation Alexa Fluor® 488, Ex: 495nm, Em: 519nm

Tested applications Suitable for: Flow Cyt (Intra), ICC/IF

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

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

Positive control ICC/IF: HeLa cells Flow Cyt (intra): HeLa cells.

This product is a recombinant monoclonal antibody, which offers several advantages including: **General notes**

- 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**.

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outlicensing@thermofisher.com.

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.

Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 30% Glycerol (glycerin, glycerine), 1% BSA, PBS

Purity Protein A purified

ClonalityMonoclonalClone numberEPR7315

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab200781 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)		1/50.
ICC/IF		1/100.

Target

Function Functions as an ubiquitin E3 ligase and as an ISG15 E3 ligase. Involved in innate immune

defense against viruses by mediating ubiquitination of DDX58. Mediates 'Lys-63'-linked polyubiquitination of the DDX58 N-terminal CARD-like region which is crucial for triggering the cytosolic signal transduction that leads to the production of interferons in response to viral infection. Promotes ISGylation of 14-3-3 sigma (SFN), an adapter protein implicated in the regulation of a large spectrum signaling pathway. Mediates estrogen action in various target

organs.

Tissue specificity Ubiquitous.

Pathway Protein modification; protein ubiquitination.

Sequence similarities Contains 1 B30.2/SPRY domain.

Contains 1 RING-type zinc finger.

Domain The RING-type zinc finger is important for ISG15 E3 ligase activity and autoISGylation.

AutoISGylation negatively regulates ISG15 E3 ligase activity.

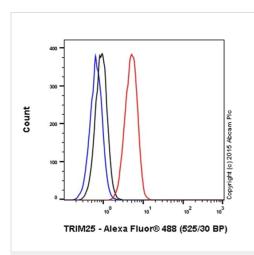
The C-terminal B30.2/SPRY domain interacts with the first N-terminal CARD domain of DDX58.

Post-translational

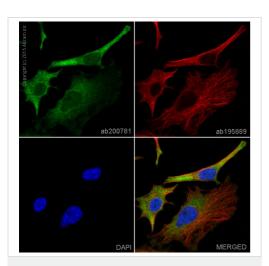
modifications

Auto-ISGylated.

Images



Flow Cytometry (Intracellular) - Alexa Fluor® 488 Anti-TRIM25/EFP antibody [EPR7315] (ab200781)

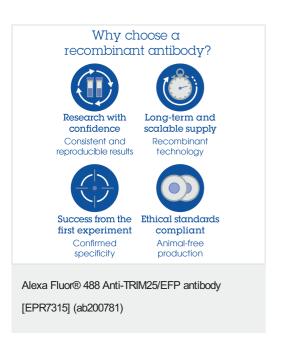


Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-TRIM25/EFP antibody [EPR7315] (ab200781)

Overlay histogram showing HeLa cells stained with ab200781 (red line). The cells were fixed with 80% methanol (5 min) and then permeabilized with 0.1% PBS-Tween 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 (ab200781, 1/50 dilution) for 30 min at 22°C. Isotype control antibody (black line) was rabbit monoclonal IgG [EPR25A] Alexa Fluor® 488 (ab199091) 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 20mW Argon ion laser (488nm) and 525/30 bandpass filter. This antibody gave a positive signal in HeLa cells fixed with 4% formaldehyde (10 min)/permeabilized with 0.1% PBS-Tween for 20 min used under the same conditions.

ab200781 staining TRIM25/EFP in HeLa cells. The cells were fixed with 100% methanol (5min), 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 ab200781 at 1/100 dilution (shown in green) and <u>ab195889</u>, Mouse monoclonal to alpha Tubulin (Alexa Fluor[®] 594), at $2\mu g/ml$ (shown in red). Nuclear DNA was labelled with DAPI (shown in blue).

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



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