

Alexa Fluor® 488 Anti-PD1 antibody [NAT105] ab220300

2 Images

Overview

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|----------------------------|---|
| Product name | Alexa Fluor® 488 Anti-PD1 antibody [NAT105] |
| Description | Alexa Fluor® 488 Mouse monoclonal [NAT105] to PD1 |
| Host species | Mouse |
| Conjugation | Alexa Fluor® 488. Ex: 495nm, Em: 519nm |
| Specificity | This antibody recognizes human PD-1, a checkpoint protein expressed by T cells that is involved in the control of immune cell responses. |
| Tested applications | Suitable for: IHC-Fr, Flow Cyt (Intra) |
| Species reactivity | Reacts with: Human |
| Immunogen | Tissue, cells or virus corresponding to Human PD1. T cells (human T/NK cell Leukemia) Database link: Q15116 |
| Positive control | IHC-Fr: normal human tonsil tissue sections Flow Cyt (intra): MOLT4 cells. |
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The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be found below, along with publications, customer reviews and Q&As

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: PBS, 30% Glycerol (glycerin, glycerine), 1% BSA |
| Purity | Immunogen affinity purified |
| Clonality | Monoclonal |
| Clone number | NAT105 |
| Isotype | IgG1 |
| Light chain type | kappa |

Applications

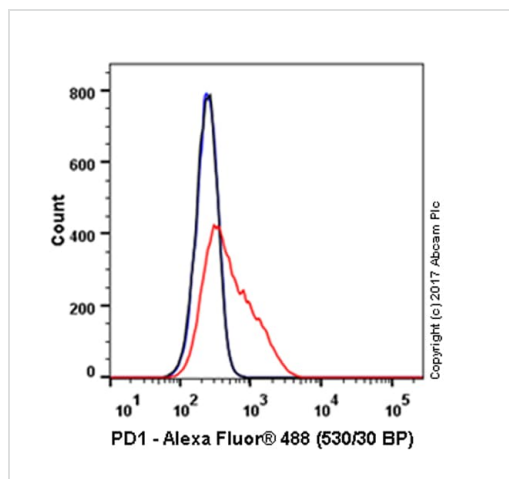
The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab220300 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| Application | Abreviews | Notes |
|------------------|-----------|--|
| IHC-Fr | | 1/100. This product gave a positive signal in frozen human tonsil tissue fixed with 10% formaldehyde (10 min) |
| Flow Cyt (Intra) | | 1/500. |

Target

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| Function | Possible cell death inducer, in association with other factors. |
| Involvement in disease | Genetic variation in PDCD1 is associated with susceptibility to systemic lupus erythematosus type 2 (SLEB2) [MIM:605218]. Systemic lupus erythematosus is a chronic, inflammatory and often febrile multisystemic disorder of connective tissue. It affects principally the skin, joints, kidneys and serosal membranes. It is thought to represent a failure of the regulatory mechanisms of the autoimmune system. |
| Sequence similarities | Contains 1 Ig-like V-type (immunoglobulin-like) domain. |
| Developmental stage | Induced at programmed cell death. |
| Cellular localization | Membrane. |

Images

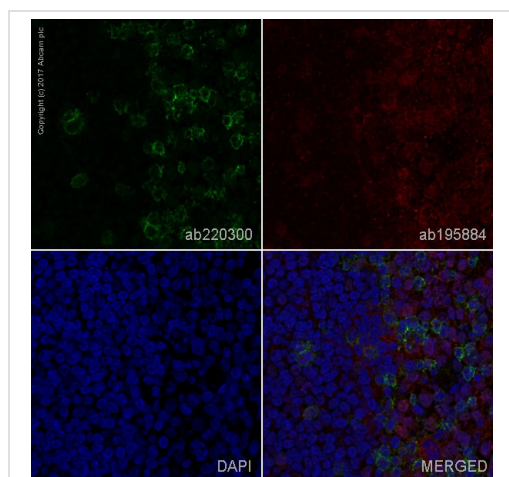


Flow Cytometry (Intracellular) - Alexa Fluor® 488
Anti-PD1 antibody [NAT105] (ab220300)

Overlay histogram showing MOLT4 cells stained with ab220300 (red line). The cells were incubated in 1x PBS / 10% normal goat serum to block non-specific protein-protein interactions followed by the antibody (ab220300, 1/500 dilution) for 30 min at 22°C.

Isotype control antibody (black line) was Mouse IgG1 (monoclonal) Alexa Fluor® 488 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 50 mW Blue laser (488nm) and 530/30 bandpass filter.



Immunohistochemistry (Frozen sections) - Alexa Fluor® 488 Anti-PD1 antibody [NAT105] (ab220300)

IHC image of PD1 staining in a section of frozen normal human tonsil*.

The section was fixed using 10% formaldehyde in 1XPBS for 10 minutes. No antigen retrieval step was performed prior to staining. Non-specific protein-protein interactions were then blocked in TBS containing 0.025% (v/v) Triton X-100, 0.3M (w/v) glycine and 1% (w/v) BSA for 1h at room temperature. The section was then incubated overnight at +4°C in TBS containing 0.025% (v/v) Triton X-100 and 1% (w/v) BSA with ab220300 at 1/100 dilution (shown in green) and counterstained using **ab195884**, Rat monoclonal to Tubulin (Alexa Fluor® 647), at 1/250 dilution (shown in red). Nuclear DNA was labelled with DAPI (shown in blue). The section was then mounted using Fluoromount®.

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

For other IHC staining systems (automated and non-automated), customers should optimize variable parameters such as antigen retrieval conditions, antibody concentrations and incubation times.

*Tissue obtained from the Human Research Tissue Bank, supported by the NIHR Cambridge Biomedical Research Centre.

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