abcam

Product datasheet

PD-L1 Antibody Panel (28-8, 73-10, SP142, CAL10) ab239749

リコンピナント

2 References 画像数 5

製品の概要

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PD-L1 Antibody Panel (28-8, 73-10, SP142, CAL10)

PD-L1 Antibody Panel ab239749 is a panel of 4 recombinant rabbit monoclonal antibodies against human PD-L1. They are provided in small sizes to allow you to easily evaluate which is the best PD-L1 antibody for your human PD-L1 research assay needs.

For guidelines on how to use each antibody within the panel, please consult the individual datasheet for each antibody.

Panel contains:

- Rabbit monoclonal [73-10] to PD-L1 (10 µL) ab228415
- Rabbit monoclonal [28-8] to PD-L1 (10 μL) ab205921
- Rabbit monoclonal [SP142] to PD-L1 (25 μL) ab228462
- Rabbit monoclonal [CAL10] to PD-L1 (10 µL) ab237726

See our <u>comparison table of PD-L1 clones 28-8, 73-10, SP142 and CAL10</u> to help you find the optimal recombinant monoclonal PD-L1 antibody clone for your research.

特記事項

All of the antibodies are ideal for use with immunohistochemistry with paraffin-embedded / formalin-fixed sections (as well as with other techniques).

Explore our range of antibody sample panels designed to provide you with a variety of trial-size antibodies in a convenient and cost-effective format.

<u>Carrier-free formulations</u> of our recombinant antibodies are available and ready to use for multiplex IHC analysis including Imaging Mass CytometryTM. Please refer to the 'Associated products' section below.

製品の特性

保存方法

Store at -20°C. Please refer to protocols.

内容	1 kit
ab205921 - Anti-PD-L1 antibody [28-8]	1 x 10µl

1

内容	1 kit
ab228415 - Anti-PD-L1 antibody [73-10]	1 x 10µl
ab237726 - Anti-PD-L1 antibody [CAL10]	1 x 10µl
ab228462 - Anti-PD-L1 antibody [SP142] - C-terminal	1 x 25µl

機能 Involved in the costimulatory signal, essential for T-cell proliferation and production of IL10 and

IFNG, in an IL2-dependent and a PDCD1-independent manner. Interaction with PDCD1 inhibits

T-cell proliferation and cytokine production.

組織特異性 Highly expressed in the heart, skeletal muscle, placenta and lung. Weakly expressed in the

thymus, spleen, kidney and liver. Expressed on activated T- and B-cells, dendritic cells,

keratinocytes and monocytes.

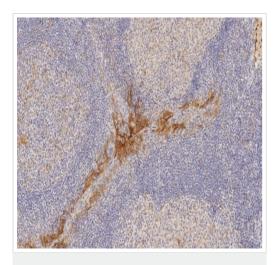
配列類似性 Belongs to the immunoglobulin superfamily. BTN/MOG family.

Contains 1 lg-like C2-type (immunoglobulin-like) domain.

Contains 1 lg-like V-type (immunoglobulin-like) domain.

細胞内局在 Cell membrane and Endomembrane system.

画像

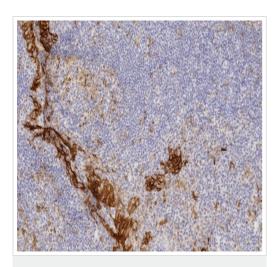


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PD-L1 antibody [28-8]

IHC image of <u>ab205921</u> staining PD-L1 in human tonsil formalin fixed paraffin embedded tissue sections*, performed on a Leica BOND RX (Polymer Refine kit). The section was pre-treated using heat mediated antigen retrieval with EDTA buffer (pH9, epitope retrieval solution 2) for 30 mins at 98°C. The section was then incubated with <u>ab205921</u>, 5µg/ml working concentration, for 60 mins at room temperature and detected using an HRP conjugated compact polymer system for 8 minutes at room temperature. DAB was used as the chromogen for 10 minutes at room temperature. The section was then counterstained with hematoxylin, blued, dehydrated, cleared and mounted with DPX.

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

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

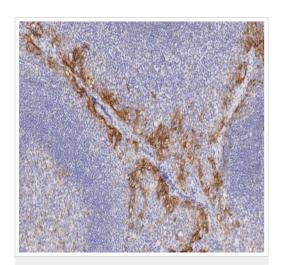


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PD-L1 antibody [SP142]

IHC image of ab228462 staining PD-L1 in human tonsil formalin fixed paraffin embedded tissue sections*, performed on a Leica BOND RX (standard Protocol F, Polymer Refine kit). The section was pre-treated using heat mediated antigen retrieval with EDTA buffer (pH9, epitope retrieval solution 2) for 30 mins at 98°C. The section was then incubated with ab228462, 1/400 working dilution, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system for 8 minutes at room temperature. DAB was used as the chromogen for 10 minutes at room temperature. The section was then counterstained with hematoxylin, blued, dehydrated, cleared and mounted with DPX.

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

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

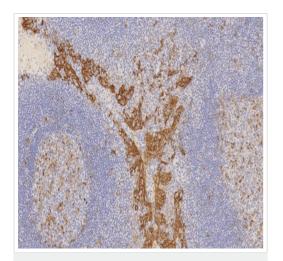


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PD-L1 antibody [CAL10]

IHC image of ab237726 staining PD-L1 in human tonsil formalin fixed paraffin embedded tissue sections*, performed on a Leica BOND RX (standard Protocol F, Polymer Refine kit). The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH6, epitope retrieval solution 1) for 30 mins at 98°C. The section was then incubated with ab237726, 1µg/ml working concentration, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system for 8 minutes at room temperature. DAB was used as the chromogen for 10 minutes at room temperature. The section was then counterstained with hematoxylin, blued, dehydrated, cleared and mounted with DPX.

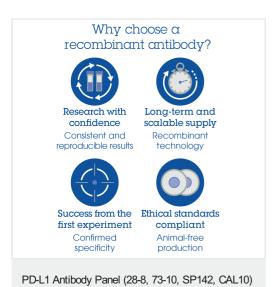
For other IHC staining systems (automated and non-automated) customers should optimize variable parameters such as antigen retrieval conditions, primary antibody concentration and antibody incubation times.

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



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PD-L1 antibody [73-10]

IHC image of <u>ab228415</u> staining PD-L1 in human tonsil formalin fixed paraffin embedded tissue sections*, performed on a Leica BOND RX (standard Protocol F, Polymer Refine kit). The section was pre-treated using heat mediated antigen retrieval with EDTA buffer (pH9, epitope retrieval solution 2) for 30 mins at 98°C. The section was then incubated with <u>ab228415</u>, 0.06µg/ml working concentration, for 15 mins at room temperature and detected using an HRP conjugated compact polymer system for 8 minutes at room temperature. DAB was used as the chromogen for 10 minutes at room temperature. The section was then counterstained with hematoxylin, blued, dehydrated, cleared and mounted with DPX.



(ab239749)

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