

Datasheet for JOT0002-5 Anti-PD-L1 VHH Antibody

BASIC INFORMATION

Name Anti-PD-L1 VHH Antibody

Specificity Programmed Death-Ligand 1 (PD-L1)

Human, Cynomolgus macaques **Species Reactivity**

Description Alpaca derived anti-PD-L1 VHH single domain antibody (molecular

weight: 14.8 kDa), expressed in E. coli under conditions free from animal

derived components

ELISA, WB, IHC, IF

Conjugate and tags Un-conjugated; with a 6*His tag at its C-terminal

Tested

Applications

Purification Affinity chromatography purified via Ni-charged resin

Format Liquid

Product buffer

25 mM TAPS pH8.5, 500 mM NaCl, 5 mM EDTA, 0.1% Proclin 300

solution

Solution

4°C; Do not freeze. **Storage**

Analyte description

ligand 1. It is a type I transmembrane protein that is thought to repress immune responses by binding to its receptor (PD1), thus inhibiting T-cell activation, proliferation, and cytokine production. It contains V-like and Clike immunoglobulin domains. PD-L1 expression is regulated by various cytokines, such as TNF-α or LPS (ISSN: 1848-7718). Increased expression of this protein in certain types of cancers, e.g., renal cell carcinoma or colon

PD-L1, also known as CD274 or B7H1, stands for programmed cell death

cancer, correlates with poor prognosis.

PARAMETERS TESTED ON EACH LOT

Product concentration 1 mg/mL by Nanodrop

QUALITY - TRACEABILITY - INTEGRITY

Life Science Group

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Purity ≥95% (4-12% gradient SDS-PAGE)

Affinity constant KD of 0.363 nM by Biacore T200

Positive controls Positive ELISA detected in: recombinant full-length human and

Cynomolgus macaques PD-L1 proteins

Positive WB detected in: PD-L1 stable expressing PLC/PRF/5 cells Positive FC detected in: PD-L1 stable expressing PLC/PRF/5 cells

Positive IHC detected in: human tonsillitis tissue, human lung cancer tissue

Positive IF detected in: MCF-7 cells

Recommended dilutions

ELISA 1:10000-1:100000

WB 1:1000-1:2000

FC: 1:50-1:100 IHC 1:200-1:800 IF 1:200-1:800

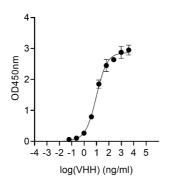
Material safety note

This product is sold **for research use only**. Standard Laboratory Practices

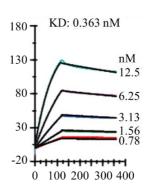
should be followed when handling this material.

SELECTED VALIDATION DATA

ELISA



SPR



Indirect ELISA showing anti-PD-L1 VHH antibody (JOT0002-5) binding to purified PD-L1. Plates were coated with 200ng/well-purified protein and the binding of JOT0002-5 was assessed in serial dilution from 0.06ng/ml primary antibody in triplicate.

Captured human PD-L1 on CM5 Chip can bind anti-PD-L1 VHH antibody (JOT0002-5) with an affinity constant of 0.363 nM as determined in SPR assay (Biacore T200).

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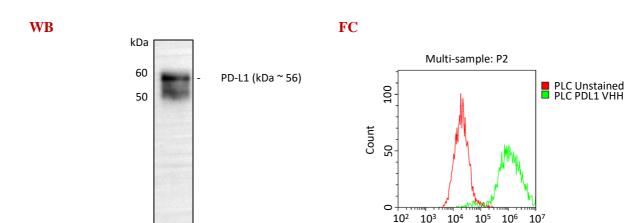
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Lysates (30 µg) were subjected to SDS-PAGE followed by western blot with anti-PDL1 VHH antibody (JOT0002-5) at dilution of 1:1000 incubated overnight at 4°C.

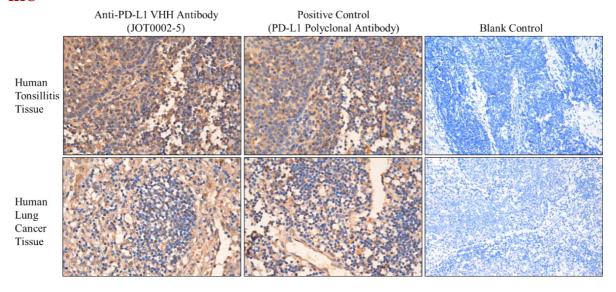
PD-L1 stable expressing PLC/PRF/5 cells were stained with anti-PDL1 VHH antibody (JOT0002-5) at a dilution of 1:50 followed by Rabbit Anti-Camelid VHH Cocktail secondary antibody [iFluor 488] at 1:1000 dilution.

10⁵ 10^{6}

 10^{3}

 10^{4}

IHC



Immunohistochemical analysis of paraffin-embedded human tonsillitis tissue and human lung cancer tissue slides using anti-PD-L1 VHH antibody (JOT0002-5) at 2.5ug/ml and positive control (PD-L1 polyclonal antibody, a competitor product) at 1:500 dilution (under 40x lens), respectively. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).

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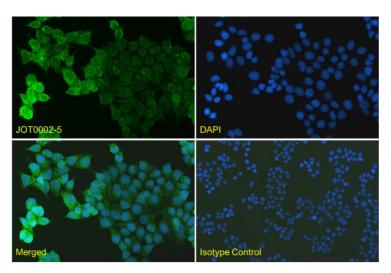


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IF



Immunofluorescence analysis of paraformaldehyde fixed MCF-7 cells stained with anti-PD-L1 VHH antibody (JOT0002-5) at 2.5 μ g/ml followed by CoraLite® 488 secondary antibody at 1:200 dilution, showing cytoplasmic staining (under 40x lens). The nuclear stain is DAPI (blue). The isotype control was stained with anti-unknown antibody followed by CoraLite® 488 secondary antibody (under 20x lens).

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