

Product datasheet for TA809809BM

OriGene Technologies, Inc.

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PD-L1 (CD274) Mouse Monoclonal Antibody (HRP conjugated) [Clone ID: OTI13G7]

Product data:

Product Type: Primary Antibodies

Clone Name: OTI13G7

Applications: ELISA, FC, IF

Recommended Dilution: IF 1:5000, FLOW 1:100

Reactivity: Human (Does not react with: Mouse)

Host: Mouse Isotype: IgG1

Clonality: Monoclonal

Immunogen: Human recombinant protein fragment corresponding to amino acids 19-239 of human PD-

L1/CD274(NP_054862) produced in HEK293 cells.

Formulation: PBS (pH 7.3) containing 1% BSA, 50% glycerol.

Concentration: 0.5 mg/ml

Purification: Purified from mouse ascites fluids or tissue culture supernatant by affinity chromatography

(protein A/G)

Conjugation: HRP

Storage: Store at -20°C as received.

Stability: Stable for 12 months from date of receipt.

Predicted Protein Size: 31 kDa

Gene Name: CD274 molecule

Database Link: NP 054862

Entrez Gene 60533 MouseEntrez Gene 29126 Human

Q9NZQ7

Synonyms: B7-H; B7H1; PD-L1; PDCD1L1; PDCD1LG1; PDL1

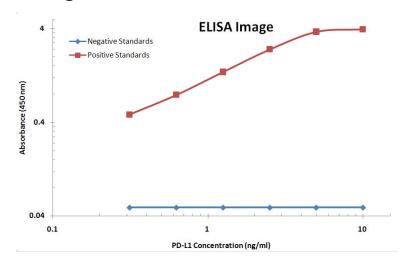
Protein Families: Druggable Genome, Transmembrane

Protein Pathways: Cell adhesion molecules (CAMs)

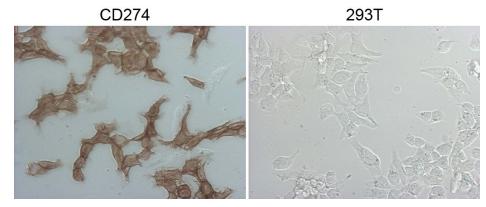




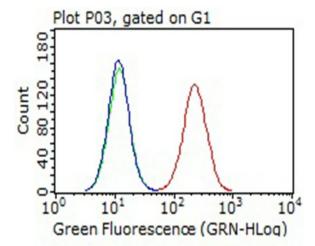
Product images:



PD-L1 ELISA with 13G7 Capture ([TA809809]) and 9E12 Detection ([TA808771]) Antibodies. Substrate used: Recombinant Human PD-L1 ([TP700201])

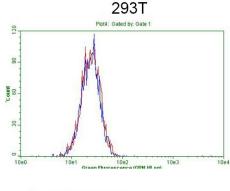


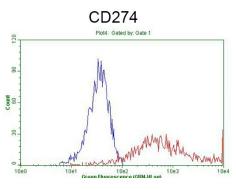
Immunocytochemistry staining of CD274 stable expression cells using anti-CD274 mouse monoclonal antibody ([TA507087]) (Left). The right is negative control (1:5000).



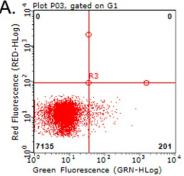
Flow cytometric Analysis of HCC78 cells, using anti-PDL1 antibody ([TA809809]), (Red), compared to isotype control, (green), and negative control (PBS), (Blue) (1:100).

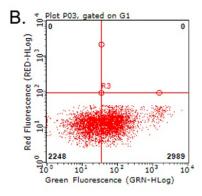




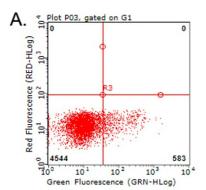


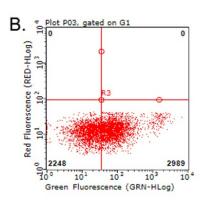
Flow cytometric Analysis of stable expression CD274 cells using anti-CD274 antibody ([TA809809]) (Red) compared to a nonspecific negative control antibody (Blue). The left is 293T as negative control (1:100).



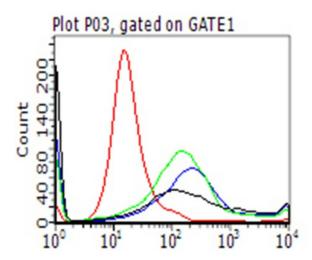


Flow cytometric analysis of living PBMCs treated with 10ug/ml PHA for 72h (Right)/untreated (Left) using anti-PDL1 antibody ([TA809809]) (1:100).





Flow cytometric analysis of living PBMCs treated with 10ug/ml PHA for 72h (Right) using anti-PDL1 antibody ([TA809809]). Cells incubated with a non-specific antibody (Left) were used as isotype control (1:100).



Detection of PDL1 neutralizing antibody using MACS column. GFP+/PDL1+ 293T cells (cotransfected with PDL1 and GFP plasmid ([RC213071], PS10010) were incubated with either PDL1 antibody [TA809809] (red), nonspecific antibody (green), isotype control (blue) or PBS (black) and then mixed with PD1+ 293T cells ([RC210364]) linked with magnetic-beads. The mixed cells were pulled down using MACS column (Miltenyi Biotec) and analysed by Flow Cytometry. GFP+/PDL1+ cells would not be collected if PD1/PDL1 interaction is neutralized by the tested antibody (1:50).