

PD-L1 (E1L3N[®]) XP[®] Rabbit mAb (Biotinylated)



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For Research Use Only. Not for Use in Diagnostic Procedures.

Applications: W	Reactivity: H	Sensitivity: Endogenous	MW (kDa): 40-50	Source/Isotype: Rabbit IgG	UniProt ID: #Q9NZQ7	Entrez-Gene Id: 29126
Product Usage Information		Application Western Blotting			Dilution 1:1000	
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		PD-L1 (E1L3N [®]) XP [®] Rabbit mAb (Biotinylated) recognizes endogenous levels of total PD-L1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human PD-L1 protein.				
Description		This Cell Signaling Technology antibody is conjugated to biotin under optimal conditions. The biotinylated antibody is expected to exhibit the same species cross-reactivity as the unconjugated PD-L1 (E1L3N [®]) XP [®] Rabbit mAb #13684.				
Background		Programmed cell death 1 ligand 1 (PD-L1, B7-H1, CD274) is a member of the B7 family of cell surface ligands that regulate T cell activation and immune responses. The PD-L1 ligand binds the PD-1 transmembrane receptor and inhibits T cell activation. PD-L1 was discovered following a search for novel B7 protein homologs and was later shown to be expressed by antigen presenting cells, activated T cells, and tissues including placenta, heart, and lung (1-3). Similar in structure to related B7 family members, PD-L1 protein contains extracellular IgV and IgC domains and a short, cytoplasmic region. Research studies demonstrate that PD-L1 is expressed in several tumor types, including melanoma, ovary, colon, lung, breast, and renal cell carcinomas (4-6). Expression of PD-L1 in cancer is associated with tumor-infiltrating lymphocytes, which mediate PD-L1 expression through the release of interferon gamma (7). Additional research links PD-L1 expression to cancers associated with viral infections (8,9).				
Background References		1. Dong, H. et al. (1999) <i>Nat Med</i> 5, 1365-9. 2. Freeman, G.J. et al. (2000) <i>J Exp Med</i> 192, 1027-34. 3. Liang, S.C. et al. (2003) <i>Eur J Immunol</i> 33, 2706-16. 4. Dong, H. et al. (2002) <i>Nat Med</i> 8, 793-800. 5. Thompson, R.H. et al. (2006) <i>Cancer Res</i> 66, 3381-5. 6. Pardoll, D.M. (2012) <i>Nat Rev Cancer</i> 12, 252-64. 7. Taube, J.M. et al. (2012) <i>Sci Transl Med</i> 4, 127ra37. 8. Lyford-Pike, S. et al. (2013) <i>Cancer Res</i> 73, 1733-41. 9. Chen, B.J. et al. (2013) <i>Clin Cancer Res</i> 19, 3462-73.				
Species Reactiv	ity	Species reactivity is d	etermined by testin	n in at least one approve	ad application (o.g.	western blot\

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Western Blot Buffer

IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat dry milk, 1X TBS, 0.1% Tween® 20 at 4° C with gentle shaking, overnight.

Applications Key W: Western Blotting

Cross-Reactivity Key H: Human

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