

PD-1 (Intracellular Domain) (D7D5W) XP® Rabbit mAb



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Applications: W, IP, IHC-Bond, IHC-P, IF-F, IF-IC, FC-FP	Reactivity: M	Sensitivity: Endogenous	MW (kDa): 40-75	Source/Isotype: Rabbit IgG	UniProt ID: #Q02242	Entrez-Gene Id: 18566
Product Usage Information		Application Western Blotting Immunoprecipitation			Dilution 1:1000 1:200	
		IHC Leica Bond Immunohistochemist Immunofluorescence Immunofluorescence Flow Cytometry (Fixed	(Frozen) (Immunocytochem	nistry)	1:1 1:1 1:1	0 - 1:200 00 - 1:400 00 - 1:400 00 - 1:400 00 - 1:400
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.				
		For a carrier free (BSA and azide free) version of this product see product #55789.				
Specificity/Sensitivity		PD-1 (Intracellular Domain) (D7D5W) $XP^{ ext{@}}$ Rabbit mAb recognizes endogenous levels of total PD-1 protein.				
Species predict based on 100% homology		Rat, Hamster				
Source / Purification Monoclonal antibody is produced by immunizing a residues surrounding Ala242 of mouse PD-1 protei					synthetic peptide co	orresponding to
Background		The programmed cell death 1 protein (PD-1, PDCD1, CD279) is a member of the CD28 family of immunoreceptors that regulate T cell activation and immune responses (1-3). The PD-1 protein contains an extracellular Ig V domain, a transmembrane domain, and a cytoplasmic tail that includes an immunoreceptor tyrosine-based inhibitory motif (ITIM) and an immunoreceptor tyrosine-based switch motif (ITSM). PD-1 is activated by the cell surface ligands PD-L1 and PD-L2 (4). Upon activation, PD-1 ITIM and ITSM phosphorylation leads to the recruitment of the protein tyrosine phosphatases SHP-1 and SHP-2, which suppress TCR signaling (5-7). In addition to activated T cells, PD-1 is expressed in activated B cells and monocytes, although its function in these cell types has not been fully characterized (8). The PD-1 pathway plays an important role in immune tolerance (3); however, research studies show that cancer cells often adopt this pathway to escape immune surveillance (9). Consequently, blockade of PD-1 and its ligands is proving to be a sound strategy for neoplastic intervention (10).				
Background Re	eferences	1. Ishida, Y. et al. (1992) <i>EMBO J</i> 11, 3887-95. 2. Shinohara, T. et al. (1994) <i>Genomics</i> 23, 704-6. 3. Nishimura, H. et al. (1999) <i>Immunity</i> 11, 141-51. 4. Freeman, G.J. et al. (2000) <i>J Exp Med</i> 192, 1027-34. 5. Yokosuka, T. et al. (2012) <i>J Exp Med</i> 209, 1201-17. 6. Sheppard, K.A. et al. (2004) <i>FEBS Lett</i> 574, 37-41. 7. Chemnitz, J.M. et al. (2004) <i>J Immunol</i> 173, 945-54. 8. Thibult, M.L. et al. (2013) <i>Int Immunol</i> 25, 129-37. 9. Dong, H. et al. (2002) <i>Nat Med</i> 8, 793-800. 10. Topalian, S.L. et al. (2012) <i>Curr Opin Immunol</i> 24, 207-12.				

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X

TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key

W: Western Blotting IP: Immunoprecipitation IHC-Bond: IHC Leica Bond IHC-P: Immunohistochemistry

(Paraffin) IF-F: Immunofluorescence (Frozen) IF-IC: Immunofluorescence (Immunocytochemistry) FC-

FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key M: Mouse

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