PD-1 (D3W4U) Rabbit mAb (Alexa Fluor® 488 Conjugate)



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Applications: FC-FP	Reactivity: H	Sensitivity: Endogenous	Source/Isotype: Rabbit IgG	UniProt ID: #Q15116	Entrez-Gene Id: 5133
Product Usage Information		Application Flow Cytometry (Fixed/P	ermeabilized)		Dilution 1:50
Storage		Supplied in PBS (pH 7.2), less than 0.1% sodium azide and 2 mg/ml BSA. Store at 4° C. Do not aliquot the antibody. Protect from light. Do not freeze.			
Specificity/Sensitivity		PD-1 (D3W4U) Rabbit mAb (Alexa Fluor [®] 488 Conjugate) recognizes endogenous levels of total PD-1 protein. This antibody binds the intracellular domain of human PD-1 protein.			
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala249 of human PD-1 protein.			
Description		This Cell Signaling Technology antibody is conjugated to Alexa Fluor [®] 488 fluorescent dye and tested in-house for direct flow cytometry analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated PD-1 (D3W4U) Rabbit mAb #15121.			
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 References		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.			

Species Reactivity

Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Applications Key FC-FP: Flow Cytometry (Fixed/Permeabilized)

Cross-Reactivity Key H: Human

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