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## TIM-3 (D5D5R™) XP® Rabbit mAb (Alexa Fluor® 647 Conjugate)

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

Applications:	Reactivity:	Sensitivity:	Source/Isotype:	UniProt ID:	Entrez-Gene Id:
FC-L	H	Endogenous	Rabbit IgG	#Q8TDQ0	84868
<b>Product Usage Information</b>	<b>Application</b>			<b>Dilution</b>	
	Flow Cytometry (Live)			1:50	
<b>Storage</b>	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.				
<b>Specificity/Sensitivity</b>	TIM-3 (D5D5R™) XP® Rabbit mAb (Alexa Fluor® 647 Conjugate) recognizes endogenous levels of total TIM-3 protein.				
<b>Source / Purification</b>	Monoclonal antibody is produced by immunizing animals with recombinant protein specific to the extracellular domain of human TIM-3 protein.				
<b>Description</b>	This Cell Signaling Technology antibody is conjugated to Alexa Fluor® 647 fluorescent dye and tested in-house for direct flow cytometric analysis in human cells. This antibody is expected to exhibit the same species cross-reactivity as the unconjugated TIM-3 (D5D5R™) XP® Rabbit mAb #45208.				
<b>Background</b>	T cell Ig- and mucin-domain-containing molecules (TIMs) are a family of transmembrane proteins expressed by various immune cells. TIM-3 is an inhibitory molecule that is induced following T cell activation (1-3). TIM-3 is expressed by exhausted T cells in the settings of chronic infection and cancer (4,5), and tumor-infiltrating T cells that coexpress PD-1 and TIM-3 exhibit the most severe exhausted phenotype (5). Tumor-infiltrating dendritic cells (DCs) also express TIM-3. TIM-3 expression on DCs was found to suppress innate immunity by reducing the immunogenicity of nucleic acids released by dying tumor cells (6). Research studies show that heterodimerization of TIM-3 with CEACAM-1 is critical for the inhibitory function of TIM-3, and co-blockade of TIM-3 and CEACAM-1 enhanced anti-tumor responses in a mouse model of colorectal cancer (7). In addition, blockade of TIM-3 in mouse models of autoimmunity enhanced the severity of disease (1). Finally, binding of Galectin-9 to TIM-3 expressed by Th1 cells induces T cell death (8).				
<b>Background References</b>	<ol style="list-style-type: none"> <li>1. Monney, L. et al. (2002) <i>Nature</i> 415, 536-41.</li> <li>2. Sánchez-Fueyo, A. et al. (2003) <i>Nat Immunol</i> 4, 1093-101.</li> <li>3. Sabatos, C.A. et al. (2003) <i>Nat Immunol</i> 4, 1102-10.</li> <li>4. Jones, R.B. et al. (2008) <i>J Exp Med</i> 205, 2763-79.</li> <li>5. Sakuishi, K. et al. (2010) <i>J Exp Med</i> 207, 2187-94.</li> <li>6. Chiba, S. et al. (2012) <i>Nat Immunol</i> 13, 832-42.</li> <li>7. Huang, Y.H. et al. (2015) <i>Nature</i> 517, 386-90.</li> <li>8. Zhu, C. et al. (2005) <i>Nat Immunol</i> 6, 1245-52.</li> </ol>				

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

**Applications Key** **FC-L:** Flow Cytometry (Live)

**Cross-Reactivity Key** **H:** Human

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