

## 20574

## TIGIT (E6L7H) Rabbit mAb



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

<b>Applications:</b> W, IP	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 18, 30-40	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #Q495A1	Entrez-Gene Id: 201633
Product Usage Information	2	<b>Application</b> Western Blotting Immunoprecipitation			<b>Dilution</b> 1:1000 1:50	
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		TIGIT (E6L7H) Rabbit mAb recognizes endogenous levels of total TIGIT protein. This antibody cross-reacts with an unidentified protein of 50 kDa and 75 kDa in some cell extracts.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val228 of human TIGIT protein.				
Background		T-cell immunoreceptor with Ig and ITIM domains (TIGIT), also known as VSIG9, VSTM3, and WUCAM, is a member of the poliovirus receptor family of immunoglobulin proteins (1-3). TIGIT is expressed at low levels on subsets of T cells and NK cells, and is upregulated at the protein level following activation of these cells (1-4). TIGIT marks exhausted T cells in the tumor microenvironment (5) and during human immunodeficiency virus (HIV) infection (6). Research has shown TIGIT interacts with several receptors expressed on antigen presenting cells, such as dendritic cells and macrophages, as well as tumor cells and cells of the microenvironment. TIGIT binds with high affinity to PVR/CD155, and with low affinity to Nectin-2/CD112 and Nectin-3/CD113 (2,4,7). Upon binding to its ligands, TIGIT suppresses T cell activation, and inhibits T and NK cell cytotoxicity. This inhibition can be blocked using monoclonal antibodies directed at the extracellular domain of TIGIT, resulting in rejuvenated antigen-specific CD8 <sup>+</sup> T cell responses in tumors and during HIV infection (5,6,8). Three potential isoforms of TIGIT have been computationally mapped (9).				
Background References		<ol> <li>Yu, X. et al. (2009) Nat Immunol 10, 48-57.</li> <li>Levin, S.D. et al. (2011) Eur J Immunol 41, 902-15.</li> <li>Boles, K.S. et al. (2009) Eur J Immunol 39, 695-703.</li> <li>Stanietsky, N. et al. (2009) Proc Natl Acad Sci U S A 106, 17858-63.</li> <li>Chauvin, J.M. et al. (2015) J Clin Invest 125, 2046-58.</li> <li>Chew, G.M. et al. (2016) PLoS Pathog 12, e1005349.</li> <li>Stengel, K.F. et al. (2012) Proc Natl Acad Sci U S A 109, 5399-404.</li> <li>Johnston, R.J. et al. (2014) Cancer Cell 26, 923-37.</li> <li>Bechtel, S. et al. (2007) BMC Genomics 8, 399.</li> </ol>				
Species Reacti	vity	Species reactivity is de	etermined by testin	g in at least one approve	ed application (e.g.,	western blot).

**Western Blot Buffer** 

**Applications Key** 

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

W: Western Blotting IP: Immunoprecipitation

**Cross-Reactivity Key** 

**H:** Human

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