

# £14971

# TIM-1 (E1R9N) Rabbit mAb



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

<b>Applications:</b> W, IHC-P	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 50, 90-140	<b>Source/Isotype:</b> Rabbit IgG	UniProt ID: #Q96D42	Entrez-Gene Id: 26762
Product Usage Information		<b>Application</b> Western Blotting Immunohistochemistry (Paraffin)			<b>Dilution</b> 1:1000 1:100	
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 #71010.				
Specificity/Sensitivity		TIM-1 (E1R9N) Rabbit mAb recognizes endogenous levels of total TIM-1 protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Val24 of human TIM-1 protein.				
Background		T cell Ig- and mucin-domain-containing molecules (TIMs) are a family of transmembrane proteins expressed by various immune cells. TIM-1 (HAVCR1 (hepatitis A virus cellular receptor 1), KIM-1 (kidney injury molecule-1) was originally identified as a receptor for hepatitis A virus (1). TIM-1 also acts as a costimulatory receptor on T cells and following activation, associates with the TCR complex to upregulate signaling and cytokine production (2-5). Another TIM family member, TIM-4, is expressed by antigen presenting cells and is a ligand for TIM-1 (6). TIM-1 expressed by Th1 and Th17 cells was also recently shown to interact with P-selectin to mediate T cell trafficking during inflammation and autoimmune disease (7). NKT cells also express TIM-1, and engagement of TIM-1 on NKT cells leads to increased production of IL-4, but decreased production of IFN-gamma (8). TIM-1 is also a receptor for phosphatidylserine exposed by cells undergoing apoptosis. Detection of phosphatidylserine by TIM-1 expressed on NKT cells results in activation, proliferation, and cytokine production (9). Expression of TIM-1 on regulatory B cells is required for optimal production of IL-10. Mice lacking the TIM-1 mucin domain have decreased production of IL-10 by regulatory B cells, hyperactive T cells, increased levels of inflammatory cytokines, and enhanced severity of autoimmune disease (10,11). In addition, TIM-1 polymorphisms are associated with susceptibility to atopic diseases including asthma (12,13). Finally, expression of TIM-1 is increased in renal tubular epithelial cells following kidney injury (14).				
Background References		1. Kaplan, G. et al. (1996) <i>EMBO J</i> 15, 4282-96. 2. Umetsu, S.E. et al. (2005) <i>Nat Immunol</i> 6, 447-54. 3. de Souza, A.J. et al. (2005) <i>Proc Natl Acad Sci U S A</i> 102, 17113-8. 4. Binné, L.L. et al. (2007) <i>J Immunol</i> 178, 4342-50. 5. de Souza, A.J. et al. (2008) <i>J Immunol</i> 180, 6518-26. 6. Meyers, J.H. et al. (2005) <i>Nat Immunol</i> 6, 455-64. 7. Angiari, S. et al. (2014) <i>Immunity</i> 40, 542-53. 8. Kim, H.S. et al. (2010) <i>J Immunol</i> 184, 4095-106. 9. Lee, H.H. et al. (2010) <i>J Immunol</i> 185, 5225-35. 10. Xiao, S. et al. (2012) <i>Proc Natl Acad Sci U S A</i> 109, 12105-10. 11. Xiao, S. et al. (2015) <i>J Immunol</i> 194, 1602-8. 12. McIntire, J.J. et al. (2001) <i>Nat Immunol</i> 2, 1109-16. 13. Khademi, M. et al. (2004) <i>J Immunol</i> 172, 7169-76. 14. Ichimura, T. et al. (1998) <i>J Biol Chem</i> 273, 4135-42.				

**Species Reactivity** 

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

**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 IHC-P: Immunohistochemistry (Paraffin)

## **Cross-Reactivity Key**

H: Human

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