#92215

## Tryptase (E5F3Q) Rabbit mAb



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

Applications: W	Reactivity: H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 25-40	Source/Isotype: Rabbit IgG	UniProt ID: #Q15661	Entrez-Gene Id: 7177	
Product Usage Information		<b>Application</b> Western Blotting			Dilution 1:1000		
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. <i>Do not aliquot the antibody.</i>					
Specificity/Sens	<b>ficity/Sensitivity</b> Tryptase (E5F3Q) Rabbit mAb recognizes endogenous levels of total Tryptase protein. This antibod does not cross-react with murine Tryptase proteins.				. This antibody		
Source / Purifica	<b>fication</b> Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues surrounding Ala120 of human Tryptase protein.					rresponding to	
Background	d Tryptase is the most abundant neutral serine protease expressed in the secretory granules of all human mast cells (1). Tryptase is secreted upon the coupled activation-degranulation response of mas cells in peripheral tissues to physical factors such as trauma, toxins, venoms, endogenous mediators, and immune mechanisms (IgE-dependent and IgE-independent) (2). Tryptase has distinct enzymatic functions that depend on the monomeric or homotetrameric state of this protein, the pH of the environment, and the presence or absence of heparin (3-5). Tryptase has the ability to cleave extracellular substrates such as vasoactive intestinal peptides (6), calcitonin gene-related peptides (7), fibronectins (8), fibrinogens (3), and kininogens (9). Tryptase is also a potent growth factor for epithelia cells, airway smooth muscle cells, and fibroblasts (10-13).						
Background Ref	erences	<ol> <li>Vanderslice, P. et al. (1990) <i>Proc Natl Acad Sci U S A</i> 87, 3811-5.</li> <li>Krishnaswamy, G. et al. (2001) <i>Front Biosci</i> 6, D1109-27.</li> <li>Fukuoka, Y. and Schwartz, L.B. (2007) <i>Int Immunopharmacol</i> 7, 1900-8.</li> <li>Ren, S. et al. (1998) <i>J Immunol</i> 160, 4561-9.</li> <li>Sakai, K. et al. (1996) <i>J Clin Invest</i> 97, 988-95.</li> <li>Tam, E.K. and Caughey, G.H. (1990) <i>Am J Respir Cell Mol Biol</i> 3, 27-32.</li> <li>Walls, A.F. et al. (1992) <i>Biochem Pharmacol</i> 43, 1243-8.</li> <li>Lohi, J. et al. (1992) <i>J Cell Biochem</i> 50, 337-49.</li> <li>Walls, A.F. et al. (1992) <i>Biochem Soc Trans</i> 20, 260S.</li> <li>Cairns, J.A. and Walls, A.F. (1996) <i>J Immunol</i> 156, 275-83.</li> <li>Ruoss, S.J. et al. (1991) <i>J Clin Invest</i> 88, 493-9.</li> <li>Brown, J.K. et al. (1995) <i>Am J Respir Cell Mol Biol</i> 13, 227-36.</li> <li>Gruber, B.L. et al. (1997) <i>J Immunol</i> 158, 2310-7.</li> </ol>					
Species Reactivi	ty	Species reactivity is de	termined by testing	g in at least one approve	d application (e.g.,	western blot).	
Western Blot Bu	ıffer	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 Ke	y	W: Western Blotting					
Cross-Reactivity	y Key	H: Human					
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