

Recombinant Mouse TREML4/TLT-4 Fc Chimera

Catalog Number: 10172-TL

DESCRIPTION				
Source	Mouse myeloma cell line, NS0-derived mouse TREML4/TLT-4 protein			
	Mouse TREML4/TLT-4 (Ser29-Leu197) Accession # Q3LRV9	IEGRMD	Human IgG ₁ (Pro100-Lys330)	
	N-terminus		C-terminus	
N-terminal Sequence Analysis	Ser29			
Structure / Form	Disulfide-linked homodimer			
Predicted Molecular Mass	45 kDa			

SPECIFICATIONS		
SDS-PAGE	64-72 kDa, under reducing conditions	
Activity	Measured by its ability to bind apoptotic EL-4 mouse lymphoblasts in a flow cytometry assay. The ED ₅₀ for this effect is 0.45-4.5 μg/mL.	
Endotoxin Level	<0.10 EU per 1 µg of the protein by the LAL method.	
Purity	>95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.	
Formulation	Supplied as a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.	

PREPARATION AND S	TORAGE		
Shipping	The product is shipped with dry ice or equivalent. Upon receipt, store it immediately at the temperature recommended below.		
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.		
	 12 months from date of receipt, -70 °C as supplied. 		
	• 1 month 0 to 0 °C under starile conditions often encoder		

- 1 month, 2 to 8 °C under sterile conditions after opening.
- 3 months, -20 to -70 °C under sterile conditions after opening.



BACKGROUND

TREML4 (Triggering Receptor Expressed on Myeloid cells-like 4), also known as TLT-4, is a type I transmembrane member of the TREM family and Ig superfamily (1, 2). Mature mouse TREML4 consists of a 172 amino acid (aa) extracellular domain (ECD) with one Ig-like domain, a 21 aa transmembrane segment, and a 42 aa cytoplasmic domain (1, 2). Within the ECD, mouse TREML4 shares 51% and 76% aa sequence identity with human and rat TREM-2, respectively. TREML4 is

expressed in the spleen, $CD8\alpha^+$ dendritic cells, and macrophages, and it is capable of recognizing necrotic cells by different phagocytes within the spleen (1). TREML4 is characterized by its Ig-like extracellular domain and short cytoplasmic tail that associates with adaptor DAP12 (2). TREML4 has been shown to play a role in coronary calcification and artery disease (3), and it is also a positive regulator of TLR7 signaling in autoimmunity and antiviral responses (4).

References:

- 1. Hemmi, H. et al. (2009) J. Immunol. 182:1278.
- 2. Hemmi, H. et al. (2012) J. Immunol. 188:1147.
- 3. Sen, S.K. et al. (2014) Am. J. Hum. Genet. 95:66
- 4. Ramirez-Ortiz, Z. G. et al. (2015) Nat. Immunol. 16:495.

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