

# **Human TRANCE/TNFSF11/RANK L Antibody**

Polyclonal Goat IgG Catalog Number: AB626

DESCRIPTION		
Species Reactivity	Human	
Specificity	Detects TRANCE/TNFSF11/RANK L in direct ELISAs and Western blots. In direct ELISAs, approximately 30% cross-reactivity with recombinant mouse TRANCE is observed.	
Source	Polyclonal Goat IgG	
Purification	Protein A or G purified	
Immunogen	S. frugiperda insect ovarian cell line Sf 21-derived recombinant human TRANCE/TNFSF11/RANK L Gln73-Asp317 (Ala194Gly) Accession # O14788	
Endotoxin Level	<0.10 EU per 1 µg of the antibody by the LAL method.	
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.	

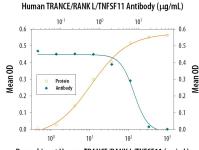
#### **APPLICATIONS**

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Western Blot	1 μg/mL	Recombinant Human TRANCE/TNFSF11/RANK L (Truncated) (Catalog # 390-TN)
Neutralization	splenocytes. Nakagav	v to neutralize TRANCE/TNFSF11/RANK L-induced osteoclast-like cell formation in mouse va, N. et al. (1998) Biochim. Biophys. Res. Commun. <b>253</b> :395. The Neutralization Dose 5 µg/mL in the presence of 30 ng/mL Recombinant Human TRANCE/TNFSF11/RANK L.

## DATA

## Neutralization



Recombinant Human TRANCE/RANK L/TNFSF11 (ng/mL)

Osteoclast-like Cell Formation Induced by TRANCE/ TNFSF11/RANK L and **Neutralization by Human** TRANCE/TNFSF11/RANK L Antibody. In the presence of Recombinant Mouse M-CSF (20 ng/mL, Catalog # 416-ML), Recombinant Human TRANCE/TNFSF11/RANK L (Catalog # 390-TN) induces osteoclast-like cell formation in mouse splenocytes in a dosedependent manner (orange line), as measured by TRAP (tartrateresistant acid phosphatase) solution assay. Under these conditions, osteoclast-like cell formation elicited by Recombinant Human TRANCE/TNFSF11/ RANK L (30 ng/mL) is neutralized (green line) by increasing concentrations of Goat Anti-Human TRANCE/ TNFSF11/RANK L Polyclonal Antibody (Catalog # AB626). The ND<sub>50</sub> is typically 5-15 µg/mL.

#### PREPARATION AND STORAGE Reconstitute at 1 mg/mL in sterile PBS. Reconstitution

The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below Shipping

### Stability & Storage

### Use a manual defrost freezer and avoid repeated freeze-thaw cycles.

- 12 months from date of receipt, -20 to -70 °C as supplied.
- 1 month, 2 to 8 °C under sterile conditions after reconstitution.
- 6 months, -20 to -70 °C under sterile conditions after reconstitution.





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#### BACKGROUND

RANK Ligand (receptor activator of NF-kB ligand [RANKL], also called TNF-related activation-induced cytokines (TRANCE), osteoprotegerin ligand [OPGL], and osteoclast differentiation factor [ODF]), is a member of the tumor necrosis factor (TNF) family. RANK Ligand was originally identified as an immediate early gene upregulated by T cell receptor stimulation. The human RANK Ligand cDNA encodes a type II transmembrane protein of 317 amino acids with a predicted cytoplasmic domain of 47 amino acids, a 21 amino acids transmembrane region, and an extracellular domain of 249 amino acids. The extracellular domain contains two potential N-linked glycosylation sites. Mouse and human RANK Ligand share 85% amino acid identity. RANK Ligand is primarily expressed in T cells and T cell rich organs, such as thymus and lymph nodes. The multi-functions of RANK Ligand include induction of activation of the c-jun N-terminal kinase, enhancement of T cell growth and dendritic cell function, induction of osteoclastogenesis, and lymph node organogenesis. RANK is the cell surface signaling receptor of RANK Ligand. RANK has been shown to undergo receptor clustering during signal transduction. Osteoprotegerin, a soluble member of the TNF receptor family which binds RANK Ligand, is a naturally occurring decoy receptor that counterbalances the effects of RANK Ligand.

#### References:

- 1. Wong, B.R. et al. (1997) J. Biol. Chem. 272:25190.
- 2. Anderson, D.M. et al. (1997) Nature 390:175.
- 3. Nakagawa, N. et al. (1998) Biochem. Biophys. Res. Commun. 245:382.
- 4. Kong, Y-Y. et al. (1999) Nature 397:315.

