

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

<b>MW (kDa):</b> 19	<b>UniProt ID:</b> #035235	Entrez-Gene Id: 21943
Background		RANKL, also known as TRANCE, TNFSF11, or OPGL, is a member of the TNF superfamily of ligands. T cells, mammary epithelial cells, and endothelial cells can produce RANKL (1). RANKL is expressed as a type II transmembrane protein or cleaved into a soluble form by extracellular proteases, such as TACE, ADAM10, and matrix metalloproteases (1). Alternative splicing also results in the production of soluble RANKL (1). RANKL signaling is antagonized by osteoprotegerin, which functions as a soluble decoy receptor (2). RANKL plays key roles in mammary gland development and dendritic cell survival and is required for osteoclast differentiation and survival (3-6). Research studies have shown that RANKL deficiencies in both mice and humans are associated with abnormally increased bone density and defects in lymphoid organogenesis (5,6).
Endotoxin		Endotoxin levels are less than or equal to 1 EU / 1 $\mu g$ mRANKL.
Purity		A greater than or equal to 95% purity was determined by SDS-PAGE.
Source / Purifica	ation	Recombinant mouse RANKL was expressed in <i>E. coli</i> and is supplied in a lyophilized form. This product was produced without any FBS or animal-derived ingredients.
Bioactivity		The bioactivity of recombinant mRANKL was determined in a secreted embryonic alkaline phosphatase (SEAP) induction assay using RAW-Blue™ cells. The ED <sub>50</sub> of each lot is less than or equal to 50 ng/ml. RAW-Blue™ Cells were purchased from InvivoGen Inc.
Background Ref	erences	1. O'Brien, C.A. (2010) <i>Bone</i> 46, 911-9. 2. Lacey, D.L. et al. (1998) <i>Cell</i> 93, 165-76. 3. Wong, B.R. et al. (1997) <i>J Exp Med</i> 186, 2075-80. 4. Fata, J.E. et al. (2000) <i>Cell</i> 103, 41-50. 5. Kong, Y.Y. et al. (1999) <i>Nature</i> 397, 315-23. 6. Conklin, J.L. et al. (1991) <i>Gastroenterology</i> 101, 657-63.
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