

## Product Data Sheet

## RANKL/TNFSF11 Protein, Cynomolgus (HEK293, Fc)

Cat. No.:	HY-P75999
Synonyms:	Tumor necrosis factor ligand superfamily member 11; RANKL; CD254; ODF; OPGL; TNFSF11; TRANCE
Species:	Cynomolgus
Source:	HEK293
Accession:	A0A7N9DBU4 (G136-D317)
Gene ID:	/
Molecular Weight:	Approximately 56 kDa

PROPERTIES	
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AA Sequence	GSQHIRAEKA MVDGSWLDLA KRSKLEAQPF AHLTINATNI PTGSHKVSLS SWYHDRGWAK ISNMTFSNGK LIVNQDGFYY LYANICFRHH ETSGDLATEY LQLMVYVTKT SIKIPSSHTL MKGGSTKYWS GNSEFHFYSI NVGGFFKLRS GEEISVEVSN PSLLDPDQDA TYFGAFKVRD ID
Biological Activity	The bioactivity was determined by measuring the ability of RANKL to induce TRAP activity in RAW 264.7 cells. The $ED_{50}$ for this effect is $\leq 10$ ng/mL, corresponding to a specific activity is $\geq 1 \times 10^5$ U/mg
Appearance	Lyophilized powder
Formulation	Lyophilized from a 0.2 $\mu m$ filtered solution of PBS, pH 7.4.
Endotoxin Level	<1 EU/µg, determined by LAL method.
Reconsititution	It is not recommended to reconstitute to a concentration less than 100 μg/mL in ddH <sub>2</sub> O. For long term storage it is recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).
Storage & Stability	Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is recommended to freeze aliquots at -20°C or -80°C for extended storage.
Shipping	Room temperature in continental US; may vary elsewhere.

DESCRIPTION	
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Background	RANKL (TNFSF11) belongs to TNF family. RANKL is a type II transmembrane protein and is a receptor activator of NF-κB (RANK) ligand. RANKL is an activator of RANK. RANKL binds to RANK and induces the differentiation of monocyte/macrophage-lineage cells into osteoclasts and leads to osteoclast precursor maturation. In bone tissue, RANKL is expressed by osteoblasts, osteocytes and immune cells, especially in osteoblasts and osteocytes <sup>[1]</sup> . RANKL is also expressed

by T cells and increases proliferation and survival of dendritic cells<sup>[2]</sup>. RANKL consists of cytoplasmic domain (1-47), helical domain (48-68), and extracellular domain (69-317). The soluble chain (140-317) is released when cleaved by enzymes such as matrix metalloproteinases (MMP3 or 7) and ADAM<sup>[1][3]</sup>. RANKL is critical for osteoclasts maturation, bone modeling, and bone remodeling, as well as the development of lymph nodes (LNs)<sup>[1]</sup>.

## REFERENCES

[1]. Ono T, et al. RANKL biology: bone metabolism, the immune system, and beyond. Inflamm Regen. 2020 Feb 7;40:2.

[2]. Li B, et al. Roles of the RANKL-RANK Axis in Immunity-Implications for Pathogenesis and Treatment of Bone Metastasis. Front Immunol. 2022 Mar 21;13:824117.

[3]. Tobeiha M, et al. RANKL/RANK/OPG Pathway: A Mechanism Involved in Exercise-Induced Bone Remodeling. Biomed Res Int. 2020 Feb 19;2020:6910312.

[4]. Mikami S, et al. Increased RANKL expression is related to tumour migration and metastasis of renal cell carcinomas. J Pathol. 2009 Aug;218(4):530-9.

[5]. Peng X, et al. Differential expression of the RANKL/RANK/OPG system is associated with bone metastasis in human non-small cell lung cancer. PLoS One. 2013;8(3):e58361.

[6]. Lloyd SA, et al. Soluble RANKL induces high bone turnover and decreases bone volume, density, and strength in mice. Calcif Tissue Int. 2008 May;82(5):361-72.

Caution: Product has not been fully validated for medical applications. For research use only.

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