

HuPro®

TNFRSF21 (Human) Recombinant Protein

Catalog # H00027242-H06 Size 25 ug

Specification	
Product Description	Purified TNFRSF21 (AAH17730.1 47 a.a 214 a.a.) human recombinant protein with His-Flag-Stre pll tag at N-terminus expressed in human cells.
Transfected Cell Line	Human HEK293H cells
Sequence	ASNLIGTYRHVDRATGQVLTCDKCPAGTYVSEHCTNTSLRVCSSCPVGTFTRHENGIEKCHDCSQ PCPWPMIEKLPCAALTDRECTCPPGMFQSNATCAPHTVCPVGWGVRKKGTETEDVRCKQCAR GTFSDVPSSVMKCKAYTDCLSQNLVVIKPGTKETDNVCGTL
Host	Human
Theoretical MW (kDa)	23.76
Interspecies Antigen Sequence	Mouse (88); Rat (89)
Form	Liquid
Preparation Method	Transfection of pSuper-TNFRSF21 plasmid into HEK293H cell, and the expressed protein was purified by <i>Strep</i> -Tactin affinity column.
Purification	Strep-Tactin affinity columns
Concentration	≥ 10 ug/ml
Storage Buffer	100 mM Tris-HCl pH 8.0, 150 mM NaCl, 1 mM EDTA, and 5 mM desthiobiotin.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot
- Enzyme-linked Immunoabsorbent Assay



- SDS-PAGE
- Protein Interaction

Gene Info — TNFRSF21	
Entrez GenelD	27242
GeneBank Accession#	BC017730.1
Protein Accession#	AAH17730.1
Gene Name	TNFRSF21
Gene Alias	BM-018, DR6, MGC31965
Gene Description	tumor necrosis factor receptor superfamily, member 21
Omim ID	605732
Gene Ontology	<u>Hyperlink</u>
Gene Summary	The protein encoded by this gene is a member of the TNF-receptor superfamily. This receptor has been shown to activate NF-kappaB and MAPK8/JNK, and induce cell apoptosis. Through its de ath domain, this receptor interacts with TRADD protein, which is known to serve as an adaptor the at mediates signal transduction of TNF-receptors. Knockout studies in mice suggested that this gene plays a role in T-helper cell activation, and may be involved in inflammation and immune regulation. [provided by RefSeq
Other Designations	OTTHUMP00000016561 OTTHUMP00000039915 TNFR-related death receptor 6 death receptor 6

Pathway

• Cytokine-cytokine receptor interaction

Disease

- Asthma
- Chromosome Aberrations
- Epilepsy



- Genetic Predisposition to Disease
- Head and Neck Neoplasms
- Migraine Disorders
- Neoplasm Recurrence
- Neoplasms