

# Product datasheet for MR228773

## Trem2 (NM\_001272078) Mouse Tagged ORF Clone

### **Product data:**

#### OriGene Technologies, Inc.

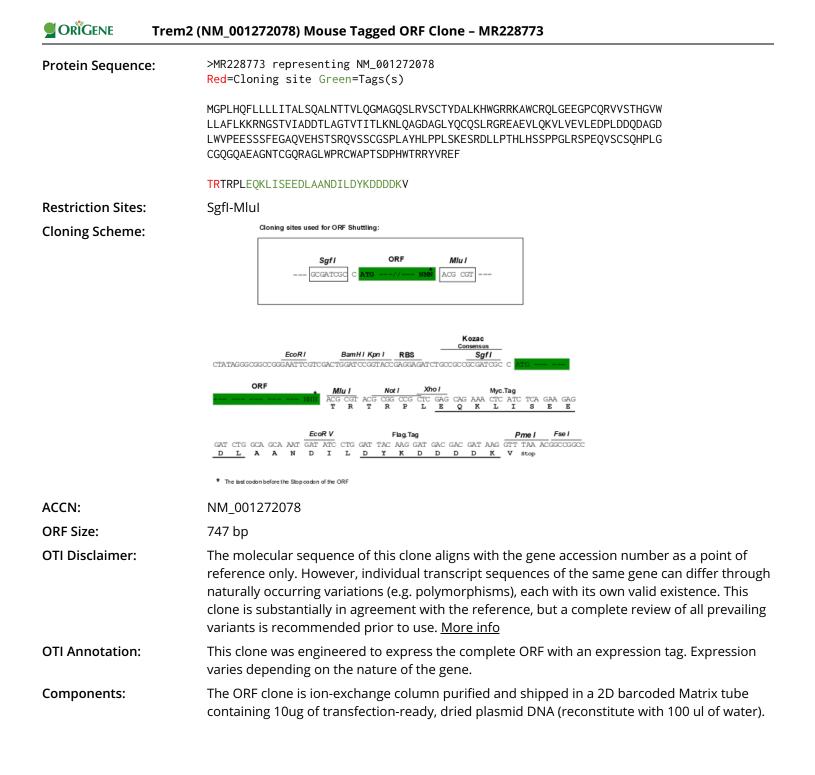
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Product Type:	Expression Plasmids
Product Name:	Trem2 (NM_001272078) Mouse Tagged ORF Clone
Tag:	Myc-DDK
Symbol:	Trem2
Synonyms:	Trem; TREM-2; Trem2a; Trem2b; Trem2c
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)
ORF Nucleotide Sequence:	>MR228773 representing NM_001272078 Red=Cloning site Blue=ORF Green=Tags(s)
	TTTTGTAATACGACTCACTATAGGGCGGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC GCC <mark>GCGATCGC</mark> C

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT ACAAGGATGACGACGATAAG**GTTTAA** 



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### **CRIGENE** Trem2 (NM\_001272078) Mouse Tagged ORF Clone – MR228773

Reconstitution Method:	<ol> <li>Centrifuge at 5,000xg for 5min.</li> <li>Carefully open the tube and add 100ul of sterile water to dissolve the DNA.</li> <li>Close the tube and incubate for 10 minutes at room temperature.</li> <li>Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid at the bottom.</li> <li>Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of shipping when stored at -20°C.</li> </ol>
RefSeq:	<u>NM 001272078.1, NP 001259007.1</u>
RefSeq Size:	1146 bp
RefSeq ORF:	750 bp
Locus ID:	83433
UniProt ID:	<u>Q99NH8</u>
Cytogenetics:	17 C
MW:	27.3 kDa
Gene Summary:	The protein encoded by this gene is part of the immunoglobulin and lectin-like superfamily and functions as part of the innate immune system. This gene forms part of a cluster of genes on mouse chromosome 17 thought to be involved in innate immunity. This protein associates with the adaptor protein Dap-12 and recruits several factors, such as kinases and phospholipase C-gamma, to form a receptor signaling complex that activates myeloid cells, including dendritic cells and microglia. In humans homozygous loss-of-function mutations in

this gene cause Nasu-Hakola disease and mutations in this gene may be risk factors to the

pathophysiological model for polycystic lipomembranous osteodysplasia with sclerosing leukoencephalopathy (Nasu-Hakola disease) and for inflammatory bowel disease. Alternative

splicing results in multiple transcript variants that encode different protein isoforms.

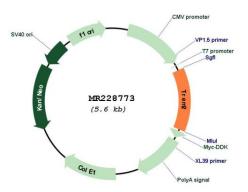
development of Alzheimer's disease. In mouse mutations of this gene serve as a

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[provided by RefSeq, Jan 2013]



### **Product images:**



Circular map for MR228773

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