

Product datasheet for SC327400

NNT1 (CLCF1) (NM 001166212) Human Untagged Clone

Product data:

Product Type: Expression Plasmids

Product Name: NNT1 (CLCF1) (NM_001166212) Human Untagged Clone

Tag: Tag Free Symbol: CLCF1

Synonyms: BSF-3; BSF3; CISS2; CLC; NNT-1; NNT1; NR6

Mammalian Cell

Selection:

Neomycin

Vector:pCMV6-Entry (PS100001)E. coli Selection:Kanamycin (25 ug/mL)

Fully Sequenced ORF: >SC327400 representing NM_001166212.

Blue=Insert sequence Red=Cloning site Green=Tag(s)

GATCCGGTACCGAGGAGATCTGCCGCCGCGATCGCC

ATGTTAGCGTGCCTGTGCACGGTGCTCTGGCACCTCCCTGCAGTGCCAGCTCTCAATCGCACAGGGGAC
CCAGGGCCTGGCCCCTCCATCCAGAAAACCTATGACCTCACCCGCTACCTGGAGCACCAACTCCGCAGC
TTGGCTGGGACCTATCTGAACTACCTGGGCCCCCCTTTCAACGAGCCAGACTTCAACCCTCCCCGCCTG
GGGGCAGAGACTCTGCCCAGGGCCACTGTTGACTTGGAGTGTGGCGAAGCCTCAATGACAAACTGCGG
CTGACCCAGAACTACGAGGCCTACAGCCACCTTCTGTGTTACTTGCGTGGCCTCAACCGTCAGGCTGCC
ACTGCTGAGCTGCCCCAGCCTGCCCACTTCTGCACCAGGCCTGCTGGGCAGCATTGCG
GGCGTCATGGCAGCTCTGGGCTACCCACTGCCCCAGCCGCTGCCTGGGACTGAACCCACTTGGACTCCT
GGCCCTGCCCACAGTGACTTCCTCCAGAAGATGGACGACTTCTGGCTGCTGAAGGAGCTGCAGACCTGG
CTGTGGCGCTCGGCCAAGGACTTCAACCGGCTCAAGAAGAAGAAGATGCAGCCTCCAGCAGCTGCAGTCACC
CTGCACCTGGGGGCTCATGGCTTCTGA

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT

TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC

Restriction Sites: Sgfl-Mlul



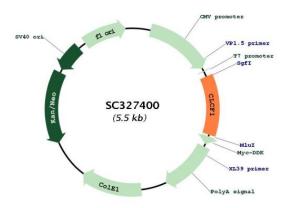
OriGene Technologies, Inc. 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



Plasmid Map:



ACCN: NM_001166212

Insert Size: 648 bp

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. 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.

RefSeq: <u>NM 001166212.1</u>



NNT1 (CLCF1) (NM_001166212) Human Untagged Clone - SC327400

RefSeq Size: 1779 bp
RefSeq ORF: 648 bp
Locus ID: 23529
UniProt ID: Q9UBD9
Cytogenetics: 11q13.2

Protein Families: Druggable Genome, Secreted Protein

Protein Pathways: Cytokine-cytokine receptor interaction, Jak-STAT signaling pathway

MW: 24.1 kDa

Gene Summary: This gene is a member of the glycoprotein (gp)130 cytokine family and encodes

cardiotrophin-like cytokine factor 1 (CLCF1). CLCF1 forms a heterodimer complex with cytokine receptor-like factor 1 (CRLF1). This dimer competes with ciliary neurotrophic factor (CNTF) for binding to the ciliary neurotrophic factor receptor (CNTFR) complex, and activates the Jak-STAT signaling cascade. CLCF1 can be actively secreted from cells by forming a complex with soluble type I CRLF1 or soluble CNTFR. CLCF1 is a potent neurotrophic factor, B-cell stimulatory agent and neuroendocrine modulator of pituitary corticotroph function. Defects in CLCF1 cause cold-induced sweating syndrome 2 (CISS2). This syndrome is characterized by a profuse sweating after exposure to cold as well as congenital physical abnormalities of the head and spine. Alternative splicing results in multiple transcript variants

encoding distinct isoforms.[provided by RefSeq, Oct 2009]

Transcript Variant: This variant (2) contains a distinct 5' UTR and lacks an in-frame portion of the 5' coding region, compared to variant 1. The resulting isoform (2) has a shorter N-

terminus when compared to isoform 1.