

Product datasheet for **SC318898**

MSF (SEPT9) (NM_001113495) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	MSF (SEPT9) (NM_001113495) Human Untagged Clone
Tag:	Tag Free
Symbol:	SEPTIN9
Synonyms:	AF17q25; MSF; MSF1; NAPB; PNUTL4; SEPT9; SeptD1; SINT1
Mammalian Cell Selection:	Neomycin
Vector:	pCMV6-Entry (PS100001)
E. coli Selection:	Kanamycin (25 ug/mL)



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Fully Sequenced ORF: >SC318898 representing NM_001113495.
 Blue=Insert sequence Red=Cloning site Green=Tag(s)

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GCTCGTTTAGTGAACCGTCAGAATTTTGTAAACGACTCACTATAGGGCGGCCGGGAATTCGTGACTG
GATCCGGTACCGAGGAGATCTGCCGCCCGGATCGCC
ATGGGCTCAAGTTTCTGGGAAGGCCTGCAGGTGGCCGTAGGGCTGCCGACGGGTGCTGGCCCCAGGGT
CTGGATTCAGGGAGCCTGCAGAGGGAGGGCAGCTGGAGGCTGCTCCAGTGTGATTGTTACGAGGCAA
AGTAAGGAGACTCTGGGCCACGCTGGGCCGGGTGGATGGAGGCAAGGAAGTCTTCGCCGGGCAAG
GGCACCAGCTGTAGATGCCGGCAGCTTTCTCCTGGACACGGCCTGGAAGGCTGACAGGGTGTGGTGAG
TGCCACCGGCTCCCTGCCGTGGCTGGTCACTGGCTTACAGGCCTCCGTGGCAGGAGGAGGATGAC
CTTGCACTCTGCTGGCCACCAATTGGCAGTGACAGACAAGCCACTGAGGCGGCTCCAGCTGCGTTGGC
GACATGGCCGACACCCAGAGATGCCGGGCTCAAGCAGGCGCCTGCATCACGGAACGAGAAGGCCCCG
GTGGACTTCGGCTACGTGGGATTGACTCCATCTGGAGCAGATGCGCCGGAAGGCCATGAAGCAGGGC
TTCGAGTTCAACATCATGGTGGTCGGCCAGAGCGGCTTGGTAAATCCACCTTAATCAACACCCTCTTC
AAATCCAAAATCAGCCGGAAGTCGGTGCAGCCACCTCAGAGGAGCGCATCCCCAAGACCATCGAGATC
AAGTCCATCACGCACGATATTGAGGAGAAAGCGTCCGGATGAAGCTGACAGTGATTGACACACCAGGG
TTCGGGGACCACATCAACAACGAGAAGTCTGGCAGCCATCATGAAGTTCATCAATGACCAGTACGAG
AAATACCTGCAGGAGGAGGTCAACATCAACCGCAAGAAGCGCATCCCGGACACCCGCTCCACTGCTGC
CTCTACTTCATCCCCGCCACCGGCACTCCCTCAGGCCCTGGACATCGAGTTTATGAAACGCCTGAGC
AAGGTGGTCAACATCGTCCCTGTATCGCAAGGCGGACACACTCACCTGGAGGAGAGGGTCCACTTC
AAACAGCGGATCACCGCAGACCTGCTGTCAACGGCATCGACGTGATCCCCAGAAGGAATTTGATGAG
GACTCGGAGGACCGGCTGGTGAACGAGAAGTTCGGGAGATGATCCATTTGCTGTGGTGGCAGTGAC
CAGGATACCAGGTCAACGGCAAGAGGATCCTTGGGAGGAAGACCAAGTGGGTACCATCGAAGTTGAA
AACACCACACTGTGAGTTTGCCTACCTGCGGGACCTTCTCATCAGGACGCACATGCAGAACATCAAG
GACATCACAGCAGCATCCACTTCGAGGCGTACCGTGTGAAGCGCCTCAACGAGGCAGCAGCGCCATG
GCCAACGGCATGGAGGAGAAGGAGCCAGAAGCCCCGGAGATGTAG
ACGCGTACGCGGCCGCTCGAGCAGAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGAT
TACAAGGATGACGACGATAAGGTTTAAACGGCCGGC
  
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Restriction Sites: Sgfl-Mlul

Plasmid Map: □

ACCN: NM_001113495

Insert Size: 1425 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: [NM_001113495.1](#)

RefSeq Size: 3651 bp

RefSeq ORF: 1425 bp

Locus ID: 10801

UniProt ID: [Q9UHD8](#)

Cytogenetics: 17q25.3

Protein Families: Druggable Genome

MW: 52.9 kDa

Gene Summary: This gene is a member of the septin family involved in cytokinesis and cell cycle control. This gene is a candidate for the ovarian tumor suppressor gene. Mutations in this gene cause hereditary neuralgic amyotrophy, also known as neuritis with brachial predilection. A chromosomal translocation involving this gene on chromosome 17 and the MLL gene on chromosome 11 results in acute myelomonocytic leukemia. Multiple alternatively spliced transcript variants encoding different isoforms have been described.[provided by RefSeq, Mar 2009]

Transcript Variant: This variant (4) lacks three 5' exons but has an alternate 5' exon, as compared to variant 1. The resulting isoform (d) has a shorter and different N-terminus, as compared to isoform a.