

Product datasheet for **SC320272**

MSF (SEPT9) (NM_006640) Human Untagged Clone

Product data:

Product Type: Expression Plasmids
Product Name: MSF (SEPT9) (NM_006640) Human Untagged Clone
Tag: Tag Free
Symbol: MSF
Synonyms: AF17q25; MSF; MSF1; NAPB; PNUTL4; SEPT9; SeptD1; SINT1
Mammalian Cell Selection: Neomycin
Vector: pCMV6-AC (PS100020)
E. coli Selection: Ampicillin (100 ug/mL)

Fully Sequenced ORF: >OriGene sequence for NM_006640.3
CGGGACTCCCGCGCTGCTAAATATATCCGTAGGAATGGAGAGGGACCGGATCTCAGCCT
TGAAAAGATCTTTTGAGGTCGAGGAGGTCGAGACACCCAACCTCCACCCACCCCGGAGGG
TCCAGACTCCCTACTCCGAGCCACTGTGGCCAGCTCCACCCAGAAATCCAGGACTGG
GCGTGAAGAAGCTCAGAACCCTCGGCCGCCATGTGGACTCCCTAAGCCAACGCTCCCCCA
AGGCGTCCCTGCGGAGGGTGGAGCTCTCGGGCCCCAAGCGGCCGAGCCGGTGTCCCGGC
GCACTGAGCTGTCCATTGACATCTCGTCCAAGCAGGTGGAGAACGCCGGGCCATCGGCC
CGTCCCGGTTCCGGCTCAAGAGGGCCGAGGTGTTGGGCCACAAGACGCCAGAACCAGGCC
CTCGGAGGACGGAGATCACCATCGTCAAACCCAGGAGTCAGCCACCGGAGGATGGAGC
CCCCTGCCTCAAGGTCCCGGAGGTGCCACTGCCCTGCCACCGACGCACCCCCAAGA
GGGTGGAGATCCAGATGCCAAGCTGTGAGGGCCCCACGCCCCAGCCAGCCAGCCAGA
CCTTGGAGAATTCAGAGCCTGCCCTGTGTCTCAGCTGCAGAGCAGGCTGGAGCCCAAGC
CCCAGCCCCCTGTGGCTGAGGCTACACCCCGGAGCCAGGAGGCCACTGAGGCGGCTCCCA
GCTGCGTTGGCGACATGGCCGACACCCCCAGAGATGCCGGGCTCAAGCAGGCGCCTGCAT
CACGGAACGAGAAGGCCCGGTGGACTTCGGCTACGTGGGGATTGACTCCATCCTGGAGC
AGATGCGCCGGAAGGCCATGAAGCAGGGCTTCGAGTTCAACATCATGGTGGTCCGGCAGA
GCGGCTTGGGTAATCCACCTTAATCAACACCCTTTCAAATCCAAAATCAGCCGGAAGT
CGGTGCAGCCACCTCAGAGGAGCGCATCCCCAAGACCATCGAGATCAAGTCCATCACGC
ACGATATTGAGGAGAAAGCGTCCGGATGAAGCTGACAGTGATTGACACACCAGGGTTCCG
GGGACCACATCAACAACGAGAAGCTGCTGGCAGCCATCATGAAGTTCATCAATGACCAGT
ACGAGAAATACCTGCAGGAGGAGTCAACATCAACCGCAAGAAGCGCATCCCGGACACCC
GCGTCCACTGCTGCCTCTACTTCATCCCGCCACCGGCCACTCCCTCAGGCCCTGGACA
TCGAGTTTATGAAACGCCTGAGCAAGGTGGTCAACATCGTCCCTGTATCGCCAAGGCGG
ACACACTCACCTGGAGGAGAGGGTCCACTTCAAACAGCGGATCACCGCAGACCTGCTGT
CCAACGGCATCGACGTGTACCCCGAGAAGGAATTTGATGAGGACTCGGAGGACCGGCTGG
TGAACGAGAAGTCCGGGAGATGATCCCATTTGCTGTGGTGGCAGTGACCACGAGTACC
AGGTCAACGGCAAGAGGATCCTTGGGAGGAAGACCAAGTGGGGTACCATCGAAGTTGAAA



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ACACCACACTGTGAGTTTGCCTACCTGCGGGACCTTCTCATCAGGACGCACATGCAGA
 ACATCAAGGACATCACCAGCAGCATCCACTTCGAGGCGTACCGTGTGAAGCGCCTCAACG
 AGGGCAGCAGCGCCATGGCCAACGGCGTGGAGGAGAAGGAGCCAGAAGCCCCGGAGATGT
 AGACGCCACCCTGCCACCCCGGGATCCTGCCCCCAAGTCATTTCCGTCCCCCCCCCAG
 GCCCTCCCACCACCCATTTTATTTTATATGATTTTCTCCATTTGTCATCGTTCCCCACC
 CCTTCGACATGCTGCCAGAAACAAGGGAAGGGCCTCCCTCCGAGTGAGTCAGTGATGA
 GGCCGCGGCCTCCCCGAGTTGTGGGGAGGCTGCACTGGAGCCACAGGCAGGGGTGAGAG
 CACCCACTGAATTGACATGACCCTCTGTCCCCAGGCTGGCTCCCCGAGGGCTCAGAAGA
 GCAGTTCCGTTGTCAGATCATCCGTCTGTGTGGGTTCTCAGTGCCGGAGGCCTTGGGG
 TGGGGGCCAGGCCTCGCACTTGCAGAGGAGCCAGTGGGCTGCACGCTCCCCTCCATCCC
 CATCGGCCCTGTCCCCTGGAGTGTGTCAGAGCCAGGGGAGAATGCAGCCACCAGGAGC
 ACCTGGACCCCTGCCGCCACATGGTGTGGCCATCACTCAGCCCTACCCTGCCCTGC
 TCCTAAGGGTAGAAAACCCAGGGTCCCCTGCCACCGACTGCCAGCCACTCCAAGCCCC
 CTGGCAGCTGCCCTCCTGGAGCAGAAAGTGCCTTTATCTCAGCCATCCGCAGACTGCTT
 GGCCAGATGCGGGGACAGGCTGGAATGAGGGAGGCGTCTTCATCTCCCTGCCATCCCCT
 CTCACGCCACCCCGCCCCACCGGGCTCAGGTGCTGCTGATGCGCTGGGATCTGATTG
 AGGATAAAAAGGAAGGAGAGATGACCCCTACCCCTCATCCCCAGTTTTGAAAAGGTCT
 AAGCAAGTGAGTCTGGTGGAGGAGCTGAGGGAGGGAGCCATGGAAGTGCCAGAAGGAAG
 GTTGGCGGGGCACGTGTGGGCCGTGGCTTGGGCTGGTCAGAGTGGCGTGAGCTGCCCGG
 CGCCTGCCCTGCCAAGTGACCAGGGAAGTGTGTGTGTCCATGTGTATGCGTGTCCGT
 CTGTCTGTCTAGTGTCTGGGTTTGGCCCAAGACTGGGCTGTAGTTACATTAATGCCCAGC
 CAGCCACCCCTGCCACTCACCCCTCCTGGCCAGGCTTGTGACTCTCTGAGCTGGGGA
 GGTGGGAGGCCAGGCGAGCCTGACTCTGTTGATCTACCCGTGCCTGGGCCCTCCCCTCA
 GAGCCCATGGTAACGAACCCCTAGAAAAGGAGAGAACGGGCGTCAGGGGTGCACAGTCCAC
 AGCTGAAGAGCAAGGTTTCGTGGCAGCACGGCCCGGCCCTCACCTCTGTCCCCACGAG
 GGGACCCATGGGGCTGTCTTTCAGGGCACAGATGACCAAAGTCCCTTCTGCTTCTG
 TTACCTGTCTTGTCTTGGGGAGAAAGAGGGGCCTGATGAGACTCCACTCAGGTGCACAC
 ATCACCAGGTGCATCTGCAGGCACCGGGCTGGCTGCTTGCAGCCAGGAGAAGGTCAGCGA
 GAAGGAGTGTATGAGTGTGAGTGTGTGTCATGGAAGTTGGGGCACTGGGCGTCTGACTC
 CCTCCCCACCAAGAGAGGAAGGACCCCTCACCACCCCACTGGCGAGACAGTTTACTTT
 GCCGACTTGCCATGTTTTTGCAAAACCAAGATTTTGAAGGAAATGAGTGCCAGCGCCA
 GGGCCCAGGCCATGTGGCCTGCCAGCCTCAATGTCACCTTGGCGCGGGGTGGGGTGGGG
 GTGGGCAGCAGCATCCCAGCCTTGAGATGCTTCACTTTCCTTCTGTAAACCAGACTTTG
 AAAAATTGTTTCGTTTCATCAGGCTCTGTTCTCAATGGCCTTTTGTACGTGCCTCCCGA
 GAAATTTGCTTTTTGTATAAATGACAAAAGTGTGAAAATGTATTTCTGAAATAAATGT
 TTCAAATGCAGAAATAAAAAAAAAAAAAAAAAA

- Restriction Sites:** Please inquire
- ACCN:** NM_006640
- 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_006640.3](#), [NP_006631.2](#)

RefSeq Size: 4452 bp

RefSeq ORF: 1707 bp

Locus ID: 10801

UniProt ID: [Q9UHD8](#)

Cytogenetics: 17q25.3

Domains: GTP_CDC

Protein Families: Druggable Genome

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 (3, also known as alpha) lacks two 5' exons but has an alternate 5' exon, as compared to variant 1. The resulting isoform (c) has a shorter and different N-terminus, as compared to isoform a.