

Product datasheet for MC209666

Zfp57 (NM_001168502) Mouse Untagged Clone

Product data:

Product Type: Expression Plasmids
 Product Name: Zfp57 (NM_001168502) Mouse Untagged Clone
 Tag: Tag Free
 Symbol: Zfp57
 Synonyms: G19; Zfp-57
 Vector: pCMV6-Entry (PS100001)
 E. coli Selection: Kanamycin (25 ug/mL)
 Cell Selection: Neomycin
 Fully Sequenced ORF: >MC209666 representing NM_001168502
 Red=Cloning site Blue=ORF Orange=Stop codon

TTTTGTAATACGACTCACTATAGGGCGCCGGGAATTCGTCGACTGGATCCGGTACCGAGGAGATCTGCC
 GCC**GCGATCGC**C

ATGGCAGCTAGGAAACAGCCATCCAGGACACCAGTCAGTTATGAGGACGTGGCAGTGTCTTTCACCCAGG
 AAGAATGGGAATATCTTACTTCTACACAGAAGACCCTTTACCAGAAAGTGATGTCAGAAACCTTCAAGAA
 CCTGACATTTGTGCGAAGCAAGAAGAAACCTCAAGAACCTAGCTCAGATCTGCAAGATAAGAACGAGGAG
 CAGGAGAAGTCTCCAGTTGCACAGGGGTATTCAAAGGTGGACCACTCTTTTTCTGTCTGACCTGTGGCA
 AATGTTTTCAAAAAGAACACCTTCTCTTAAATCACCAGTTTCTGTGAGGTCCCGGAGGCTGGCAGTCAC
 AAATCCACAAAGCCGCAAGGCAAGGGCTACAAGGCTCAGCATCGTGGAGAGAGGCCCTTTCTTTGTAAT
 TTCTGTGGCAAGACTTACCGTGATGCTTCTGGACTGAGCCGTCACCGACGTGCTCATTTAGGTTATAGGC
 CCCGTTTATGCCCTGAGTGTGAAAGTGTTCGGGATCAGTCTGAGGTCAACCGTCACCTGAAGGTACA
 CCAAAACAAGCCAGCAGCTAGCAACCAGGCTGGCAACCAGGCTAGCAACCAGAGGCTGAAGAGTAGGGTT
 CCACCTACAACACCTAGATCCCAAGCGCCCGCCCTCAAGTATGTGAAAGTGATCCAGGGACCAGTGGCCA
 GGGCTAAGGCACGGAACAGCGGAGCCTCGACCCTGAATGTCAGATCCAACCTATTACAGTGGTCCGTTT
 AAGAGAAAAGATCTCTTGTCCCTATTGTACATAACGTTTACCATGAGAACCTGTCTCTTAACCCACCTC
 AAGATCCACTTCAGACGTCAACCAACAGCACTTCTGCTGCAAAGAGTCGGCCCACTCATCCAACACAC
 TCAGAATGCAAAGATCTACACTTGCCCGTCTGTGACAGCTCCTTTAGGGGAAAGGAGAGCCTGCTGGA
 TCACTTGTGCTGCCAAAGACCAATCAGATTAGTAAATGCTGGGAAATCCTGGGTCATTTGCTCGGCTAT
 CTTTATGAACCCGTGGTGTGGGAAATATTTTTAAAGTAAAGGACTCCTCGGAAAGAGGATGGAATCCA
 GGAGGAGAAGACGGAACGTGCCTGCACTGAGAATCCTGAAACAGAAGGCCTGTCTGGGAAAGGTAGGGT
 GGCTCCGTGGGAAATGGAGGGTCCACCAGCCCTGAGAGTCTGTGACAGAAGAAGATTCGGACTGA

ACGCGTACGCGGCCGCTCGAGCAGAAACTCATCTCAGAAGAGGATCTGGCAGCAAATGATATCCTGGATT
 ACAAGGATGACGACGATAAGGTTTAA



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Restriction Sites:	Sgfl-Mlul
ACCN:	NM_001168502
Insert Size:	1257 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).
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:	<ol style="list-style-type: none">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_001168502.1 , NP_001161974.1
RefSeq Size:	2088 bp
RefSeq ORF:	1257 bp
Locus ID:	22715
UniProt ID:	Q8C6P8
Cytogenetics:	17
Gene Summary:	<p>Transcription regulator required to maintain maternal and paternal gene imprinting, a process by which gene expression is restricted in a parent of origin-specific manner by epigenetic modification of genomic DNA and chromatin, including DNA methylation. Acts by controlling DNA methylation during the earliest multicellular stages of development at multiple imprinting control regions. Required for the establishment of maternal methylation imprints at SNRPN locus. Acts as a transcriptional repressor in Schwann cells. Binds to a 5'-TGCCGC-3' consensus sequence and recognizes the methylated CpG within this element. [UniProtKB/Swiss-Prot Function]</p> <p>Transcript Variant: This variant (3) differs at the 5' end and uses an alternate, in-frame acceptor splice site at one of the coding exons compared to variant 1. This results in a shorter isoform (2) missing three internal aa compared to isoform 1. Sequence Note: This RefSeq record was created from transcript and genomic sequence data to make the sequence consistent with the reference genome assembly. The genomic coordinates used for the transcript record were based on transcript alignments.</p>