

Product datasheet for **SC101935**

SHFM3 (FBXW4) (AK056917) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	SHFM3 (FBXW4) (AK056917) Human Untagged Clone
Tag:	Tag Free
Symbol:	SHFM3
Synonyms:	DAC; FBW4; FBWD4; SHFM3; SHSF3
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)



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Fully Sequenced ORF: >NCBI ORF sequence for AK056917, the custom clone sequence may differ by one or more nucleotides

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AGTGGTTAGGTGAGAGAAAAGGCGAGCTCTGATCTCCAGCAGGATTAAGTTGCTCCCCTCTCTATTGAT
TTACCTTGATGCTAATTTAGGCCTGTGCCAGTCAGAGGCTGCCATAGCCTTAAGAGATGCCTGGCCG
GGCGCGTTGGCTCAGCCTGTAATTCCAGCACTTTCGGAGGCCAAGGTGGGCGGATCACAGGTCAAGAA
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GAGCTCGTGGGCTTCTGGCCAGGCACCACTGGGGCTACACCCCTCACAGTGTCCAAGATGTCTTCCC
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CAGCGTCCGAAATGTGTATGGAGTGGGAGGAGCCCCACGACAGCACCCTGTACTGCCTGCAGACAGAT
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CCAGATGGTTCCTTTCTGCCCTTCTGGAAGGAAAGGTGAGGCTGCCAATAGCCTCCTGGCACCAGCCA
GACCTCACCCCTGACCAACCTCTCGGGGCTGGGGTTCATTCTGGGGCACTGTGGCCTGTTTTGCTTT
GAAACCAAGAAAGAGCAAAGGGAACCCAGCAGTCTGAGTGAGTTCTGAGCCAGCCCTACCTCAGGCTGG
CTGTTGAGACATGCTACAATTTTCATTTTTGTAATAAAGCTTATTGTTTAC
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5' Read Nucleotide Sequence:	>OriGene 5' read for AK056917 unedited GTAACGTCAGAATTGTAACGACTCCTAAAGGCGGCCGGAATTCGGCAGGAGGGCTGGG GTGCTGGATGTCATGTATGAGTCCCCTTTCACACTGCTGTCTGTGGCTATGACACCTAT GTTCGCTACTGGGACCTCCGACCAGCGTCCGGAAATGTGTCATGGAGTGGGAGGAGCCC CACGACAGCACCTGTACTGCCTGCAGACAGATGGCAACCACCTGCTGGCCACAGTTCC TCCTACTACGGTGTGTACGGCTGTGGGACCGCGTCAAAGGGCCTGCCTGCACGCCTTC CCGCTGACGTCGACTCCCCTCAGCAGCCCTGTGTACTGCCTGCGTCTACCACCAAGCAT CTCTATGCTGCCCTGTCTTACAACCTCCACGTCCTGGATTTTCAAACCCATGACCGTCA GGGCCACCCTGCCTCTGGGCCAGGAAACCAGTACTCAGGGACTTCTCTTGCCTGGAG GGTGCACTGATAGCTCCTCCTACTGCCCACTGTGCTCCTGGCCTGTGACCCAGTGC TCAGGCACCTTGCACTAGAGGCTTCTGACTCCTGGGACTTTGGAGCTTACCAGAGATGCA GTCCCTCCAGGAACCTGTTGGAGAGGCAGGACCTGCTGCTTTAGAGTGGGCTGAACCC GGGCCTTGGCTCCCTGTTGGCCAGAGCAAGGATCTGGCCTGGAGAGGCCATCTATAC CCCTTATTAGAGCCATGACAGCCTACAGAGTGAGGTGAGGTGCTCCACCTTCCCAGATG GTTCCTTCTGCCCTTCTGGAAGAAAAGTGAGGCTGCCAATAGCCTCCTGGCACCAGC CAGACCTCACCTTGACCAACCTCTCGGGCTGGGGGTTCAATCCTGGGGCACTA
Restriction Sites:	NotI-NotI
ACCN:	AK056917
Insert Size:	1200 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:	AK056917.1
RefSeq Size:	3205 bp
RefSeq ORF:	3205 bp
Locus ID:	6468
Cytogenetics:	10q24.32
Protein Families:	Druggable Genome

Gene Summary:

This gene is a member of the F-box/WD-40 gene family, which recruit specific target proteins through their WD-40 protein-protein binding domains for ubiquitin mediated degradation. In mouse, a highly similar protein is thought to be responsible for maintaining the apical ectodermal ridge of developing limb buds; disruption of the mouse gene results in the absence of central digits, underdeveloped or absent metacarpal/metatarsal bones and syndactyly. This phenotype is remarkably similar to split hand-split foot malformation in humans, a clinically heterogeneous condition with a variety of modes of transmission. An autosomal recessive form has been mapped to the chromosomal region where this gene is located, and complex rearrangements involving duplications of this gene and others have been associated with the condition. A pseudogene of this locus has been mapped to one of the introns of the BCR gene on chromosome 22. [provided by RefSeq, Jul 2008]