

SHFM3 (FBXW4) (NM_022039) Human Tagged ORF Clone

Product Type:	Expression Plasmids
Tag:	TurboGFP
Symbol:	SHFM3
Synonyms:	DAC; FBW4; FBWD4; SHFM3; SHSF3
Mammalian Cell	Neomycin
Selection:	
Vector:	pCMV6-AC-GFP (PS100010)
E. coli Selection:	Ampicillin (100 ug/mL)
ORF Nucleotide Sequence:	>RG209411 representing NM_022039 Red=Cloning site Blue=ORF Green=Tags(s)

ATGCGCGCGCGCGCGGGAGGAGGAGGAGGAGGAGGAGGCGGCTCGGAGTCGGCTGCCCGCCGCGGC
CGGGGCTGCGCTCTGGCGCTGCCGAGGAGCTGCTGCTGCTCATCTGCTCCTACCTGGACATGCGGGC
CCTCGCGCGCTGGCCAGGTGTGCCGTGGTCGGCGCTTACCAGTGCATCTGCTCTGGCGCGG
ATAGCCCGGGCCTCGCTCAACTCCGGCTTACGCGGCTCGGCACCGACCTGATGACCAGTGTCCAGTGA
AGGAACGAGTGAAGGTGTCTCAGAACTGGAGACTGGGCGCTGCCGAGAGGGGATTCTGCTGAAGTGAG
ATGCAGTCAGATGCCCTGGATGCAGCTAGAGGATGATTCTCTGTACATATCCAGGCTAATTTTCATCTG
GCCTACCACTTCCGTCCAGATGGTGCCAGCTTGAATCGTCGGCTCTGGGATCTTTGCTGGGCATGATG
AGGACGTTTGGCACTTTGTCTGGCCAACTCGCATATTTGTAAGTCAGAGGGGATGGGAAGATTGGCAT
TCATAAGATTACAGACACCTTGACCTGTCAAGTACTCGGCTCATGAACAGGAGGTGAATCTGTGGATTGC
AAAGGGGCGATCATTTGTGAGTGGCTCCAGGGACAGGACGGCCAAGTGTGGCCTTTGGCCTCAGGCCGGC
TGGGGCAGTGCTTACACACCATCCAGACTGAAGACCGAGTCTGGTCATTGCTATCAGCCCAATTACTCAG
CTCTTTTGTGACAGGGACGGCTTGTTCGGGGCACTTCTACCCCTGAGAATCTGGGACCTCAACAGTGG
CAGCTGATGACACACTTGGGCAGTGACTTCCCCAGGGGCTGGGGTGTGGATGTATGTATGAGTCCC
CTTTCACACTGCTGTCTGTGGCTATGACACCTATGTTTCGCTACTGGGACCTCCGCACACGCTGCCGAA
ATGTGTATGAGTGGGAGGAGCCCCACGACAGACCTGTACTGCCTGCAGACAGATGGCAACCACCTG
CTGGCCACAGGTTCTCTCTACTACGGTGTGTACGGCTGTGGGACCTCGCGCTCAAAGGGCTGCTGTCAGC
CTTCCCGCTGACCTGCACTCCCTCAGCAGCTCTGTGATGCTGCTCTCACCACCAAGCATCTCTA
TGCTGCCCTGTCTACAACCTCCAGCTCCTGGATTTTCAAAACCA



Protein Sequence: >RG209411 representing NM_022039
Red=Cloning site Green=Tags(s)

MAAAAGEEEEEAAARESAARPAAGPALWRLPEELLLLICSYLDMRALGRLAQVCRWLRRFTSCDLLWRR
IARASLNSGFTRLGTDLMTSVPVKERVKVSQNWRLGRCREGILLKWRCSQMPWMQLEDDSLYISQANFIL
AYQFRPDGASLNRRLPLGVFAGHDEDVCHFVLANSIHVSAGDGKIGIKIHSTFTVKYSAHEQEVNCVDC
KGGIIVSGSRDRTAKVWPLASGRLGQCLHTIQTEDRVWSIAISPLSSFTGTACCGHFSPLRIWDLNSG
QLMTHLGSDFPAGVLDVMEYSPFTLLSCGYDTYVRYWDLRTSVRKCVMEWEEPHDSTLYCLQTDGNHL
LATGSSYYGVVRLWDRRQRACLHAFPLTSTPLSSPVYCLRLTTKHLAALSYNLHVLDQNP

TRTRPLE - GFP Tag - V

Restriction Sites: SgfI-MluI

Cloning Scheme:



ACCN: NM_022039

ORF Size: 1236 bp

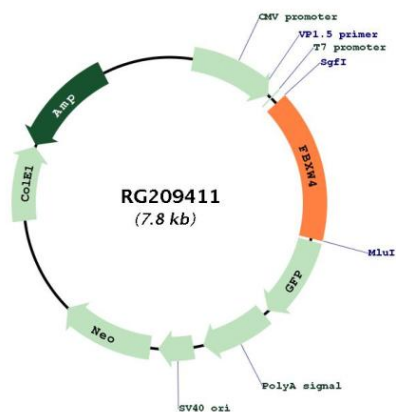
OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing variants is recommended prior to use. [More info](#)

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression varies depending on the nature of the gene.

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.
Note:	Plasmids are not sterile. For experiments where strict sterility is required, filtration with 0.22um filter is required.
RefSeq:	NM_022039.3 , NP_071322.1
RefSeq Size:	2303 bp
RefSeq ORF:	1704 bp
Locus ID:	6468
UniProt ID:	P57775
Cytogenetics:	10q24.32
Domains:	WD40, F-box
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]

Product images:



Circular map for RG209411