

Product datasheet for **SC111098**

AIF (AIFM1) (NM_004208) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	AIF (AIFM1) (NM_004208) Human Untagged Clone
Tag:	Tag Free
Symbol:	AIF
Synonyms:	AIF; AUNX1; CMT2D; CMTX4; COWCK; COXPD6; DFNX5; NADMR; NAMSD; PDCD8; SEMDHL
Mammalian Cell Selection:	None
Vector:	<u>pCMV6-XL5</u>
E. coli Selection:	Ampicillin (100 ug/mL)



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Fully Sequenced ORF: >OriGene ORF within SC111098 sequence for NM_004208 edited (data generated by NextGen Sequencing)

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ATGTTCCGGTGTGGAGGCCTGGCGGCGGGTGCTTTGAAGCAGAAGCTGGTGCCCTTGGTG
CGGACCGTGTGCGTCCGAAGCCCAGGCAGAGGAACCGGCTCCCAGGCAACTTGTTCAG
CGATGGCATGTTCTCTAGAACTCCAGATGACAAGACAAATGGCTAGCTCTGGTGATCA
GGGGGCAAAATCGATAATTCTGTGTTAGTCCTTATTGTGGGCTTATCAACAGTAGGAGCT
GGTGCCATGGCTACAAGACTATGAAAGAGGATGAAAAAGATACAATGAAAGAATTTCA
GGGTTAGGGCTGACACCAGAACAGAAACAGAAAAAGGCCGCTTATCTGCTTCAGAAGGA
GAGGAAGTTCCTCAAGACAAGGCCTCAAGTCAATGTTCTTTCTGCTAATTGGTGGAGGC
ACAGCTGCTTTTCTGTCAGCCAGATCCATCCGGGCTCGGGATCCTGGGGCCAGGGTACTG
ATTGTATCTGAAGATCCTGAGCTGCCGTACATGCGACCTCCTCTTTCAAAGAAGTGTGG
TTTTCAGATGACCCAAATGTCACAAAGACACTGCGATTCAAACAGTGAATGGAAGAG
AGAAGCATATATTTCCAGCCACCTTCTTTCTATGTCTCTGCTCAGGACCTGCCTCATATT
GAGAATGGTGGTGTGGCTGTCTCACTGGGAAGAAGGTAGTACAGCTGGATGTGAGAGAC
AACATGGTGAACTTAATGATGGCTCTCAAATAACCTATGAAAAGTGCTTGATTGCAACA
GGAGGTAATCCAAGAAGTCTGTCTGCCATTGATAGGGCTGGAGCAGAGGTGAAGAGTAGA
ACAACGCTTTTTCAGAAAGATTGGAGACTTTAGAAGCTTGAGAAGATTTACGGGAAGTC
AAATCAATTACGATTATCGGTGGGGGCTTCTTGGTAGCGAACTGGCCTGTGCTCTTGGC
AGAAAGGCTCGAGCCTTGGGCACAGAAGTGATTCAACTCTTCCCCGAGAAAGGAAATATG
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ATCAAGCTGAAAGACGGCAGGAAGGTAGAACTGACCACATAGTGGCAGCTGTGGGCCTG
GAGCCCAATGTTGAGTTGGCCAAGACTGGTGGCCTGGAATAGACTCAGATTTTGGTGGC
TTCCGGGTAAATGCAGAGCTACAAGCAGCTCTAACATCTGGGTGGCAGGAGATGCTGCA
TGCTTCTACGATATAAAGTTGGGAAGGAGCGGGTAGAGCACCATGATCACGCTGTTGTG
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ATGTTCTGGAGTGATTTGGGCCCGATGTTGGCTATGAAGCTATTGGTCTTGTGGACAGT
AGTTTGCCACAGTTGGTGTGTTTTGCAAAGCAACTGCACAAGACAACCCCAAATCTGCC
ACAGAGCAGTCAGGAACTGGTATCCGATCAGAGAGTGAGACAGAGTCCGAGGCCTCAGAA
ATTACTATTCTCCAGCACCCCGCAGTTCACAGGCTCCCGTCCAGGGGGAGGACTAC
GGCAAAGGTGTCATCTTCTACCTCAGGGACAAAGTGGTCGTGGGGATTGTGCTATGGAAC
ATCTTTAACCGAATGCCAATAGCAAGGAAGATCATTAAAGGACGGTGAGCAGCATGAAGAT
CTCAATGAAGTAGCCAACTATTCAACATTCATGAAGACTGA

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Clone variation with respect to NM_004208.3

5' Read Nucleotide Sequence:	<p>>OriGene 5' read for NM_004208 unedited</p> <pre>TTTTAGGATTTTGTAAATACGATCTCACTATAGGGCGGCCGCAATCGGCACGAGGGAGAG GAAAGGGAAGGAGGAGGTCCCGAATAGCGGTCCGCCGAAAGTTCGGTGTGGAGGCCTGGC GGCGGGTGCTTTGAAGCAGAAGCTGGTGCCTTGGTGCGGACCGTGTGCGTCCGAAGCCC GAGGCAGAGGAACCGGCTCCCAGGCAACTGTTCCAGCGATGGCATGTTCTCTAGAAGT CCAGATGACAAGACAAATGGCTAGCTCTGGTGCATCAGGGGGCAAATCGATAATTCTGT GTTAGTCCTTATTGTGGCTTATCAACAGTAGGAGCTGGTGCCTATGCCTACAAGACTAT GAAAGAGGATGAAAAAGATACAATGAAAGAATTCAGGGTTAGGGCTGACACCAGAACA GAAACAGAAAAAGGCCGGTTATCTGCTTCAGAAAGGAGAGGAAGTTCCTCAAGACAAGGC GCCAAGTCATGTTCTTTCTGCTAATTGGTGGAGGCACAGCTGCTTTTGCTGCAGCCAG ATCCATCCGGGCTCGGGATCCTGGGGCCAGGGTACTGATTGTATCTGAAGATCCTGAGCT GCCGTACATGCGACCTCCTTTTCAAAGAAGTGTGGTTTTTCCAGATGACCCACATGTCAC AAAGACTGCGATTCAAACAGTGAATGAAAAAGAGAGAAGCATATATTTCCAGCCACC TTCTTTCTATGTCTCTGCTCANGACCTGCCTCATATTGAGAATGGCGCTGTGGCTGTCT CACTGTGAAGAAGTAGTACACCTGGATGTGACAGACAACAGGGTAAAACCTAATGATGGC TCTCCAATAACCTATGAAAAGTGCCTTGATTGCAACAAGTAGGTAATCCAAACAAGTCTG TCTTGCCATTGATACGGCTGGAC</pre>
3' Read Nucleotide Sequence:	<p>>OriGene 3' read for NM_004208 unedited</p> <pre>ACCGCGGCCGCAATCTANAGTCGAGTTTTTTTTTTTTTTTTTTTGAAGAACAAGATTTAT TTCATATGTGAACATTAAGAATTTACCTACATAGTTGAAAATATTCACAAAGGACTTGA TCATTCACACTCATAACAGAGAAAGTCTGCTGAATAAAAAAATGCTCCTTTACCCATTC GACCTCCTCTCAGGGGCTGCAGTGGGTTTGCCAATCCACTGTGGGGCTTCAGTCTTCAT GAATGTTGAATAGTTTTGCTACCTCCATTGAAAACCTATGCTGGCCCCCGGGCCTTAAA AGAACCCCTGCTTTTGGGGCTTGGGGAAAAAAAAGGTTCAAAAAAAAACCCCCCAA CCCTTTTTTCTCTGGGGGAAAAAAAACCTTTTTCGGGGAAACCCCCCCCCCC GAGGAGGGATCCTTTTAAACCCCGGGGGGGGGGGGGGAGGAAAAAAAATTTTTG TGGCGGGGGGCTTTTTTTTTTTTTTTTTTTTGGAGGGCAACACCCCTCTTTCT TTTTTTGAAAAAAAAGGGGTGTGTTTTTACCCTCATTTCTTTTCAAAAAAACAC CCCGCGGGGGTGTCTCCTCCCAAAAAACAAAAATAATTTTTTATTCGGTACGCG CGGGGGCGCTCCACCACACAACGAATTAGTGGGGCGAGGGGGGGCCGGCGAATACC AACACACTATTTTTTTTTTTTTTCTCTCTCTCCCTCCCCCAACCCACCCGAGACG GGGGGGGTGGCGGGGCCCCCCCTCTCTTATNTTACACAAAAACCCAC CCTCCTCTCTCTCCCACTCAGATACTATATGCATTGCGCGTCTCTCCCACT TAATAACCAAAAAC</pre>
Restriction Sites:	NotI-NotI
ACCN:	NM_004208
Insert Size:	2000 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_004208.2 , NP_004199.1
RefSeq Size:	2215 bp
RefSeq ORF:	1842 bp
Locus ID:	9131
UniProt ID:	O95831
Cytogenetics:	Xq26.1
Domains:	pyr_redox
Protein Families:	Druggable Genome, Transmembrane
Protein Pathways:	Apoptosis
Gene Summary:	<p>This gene encodes a flavoprotein essential for nuclear disassembly in apoptotic cells, and it is found in the mitochondrial intermembrane space in healthy cells. Induction of apoptosis results in the translocation of this protein to the nucleus where it affects chromosome condensation and fragmentation. In addition, this gene product induces mitochondria to release the apoptogenic proteins cytochrome c and caspase-9. Mutations in this gene cause combined oxidative phosphorylation deficiency 6 (COXPD6), a severe mitochondrial encephalomyopathy, as well as Cowchock syndrome, also known as X-linked recessive Charcot-Marie-Tooth disease-4 (CMTX-4), a disorder resulting in neuropathy, and axonal and motor-sensory defects with deafness and cognitive disability. Alternative splicing results in multiple transcript variants. A related pseudogene has been identified on chromosome 10. [provided by RefSeq, Aug 2015]</p> <p>Transcript Variant: This variant (1) encodes the longest isoform (AIF).</p>