

Product datasheet for **SC126999**

ZNF195 (NM_007152) Human Untagged Clone

Product data:

Product Type:	Expression Plasmids
Product Name:	ZNF195 (NM_007152) Human Untagged Clone
Tag:	Tag Free
Symbol:	ZNF195
Synonyms:	HRF1; ZNFP104
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 SC126999 sequence for NM_007152 edited (data generated by NextGen Sequencing)

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ATGACTCTGTTGACGTTTCAGGGATGTGGCCATAGAATTCTCCCTGGAGGAGTGGAATGC
CTGGACCTCGCTCAGCAGAAATTTGTACAGGGATGTGATGTTGGAGAACTACAGAACTTG
TTCTCCGTTGGTCTCACTGTCTGTAAGCCAGGCCTGATCACCTGCCTGGAGCAACGAAAA
GAGCCCTGGAATGTGAAGAGACAGGAGGCAGACGGACATCCAGCTATGCTTCTCAT
TTTACCAAGACCTTCTGCCAGAGCAGGGCATAACAAGATGCATTCCCAAAAAGAATACTG
AGAGGATATGGAATTGTGGCCTTGATAATTTATATTTAAGGAAAGACTGGGAAAGTTTA
GATGAGTGAAGTTGCAAAAAGATTATAATGGACTTAACCAATGTTCACTCAACTACCCAT
AGCAAAATCTTTCAATATAATAAATATGTTAAAATCTTTGATAACTTTTCAAATTTACAT
AGACGTAATAAGTAATACTGGAGAGAAACCTTTCAAATGTCAAGAATGTGGCAAATCC
TTTCAAATGCTCTCATTCTAATGAACATCAGAAAATTCACACTGGAAAAAAATTTCAA
AAATGTGGAGAATGTGGCAAAACCTTTTCCAGTGCTCACACTTTACTGAACCTGAGAAC
ATTGACACTGGAGAGAAACCTTACAAGTGTCAAGAATGTAAACACGTCATTAACACTTGC
TCAGTCTTACTAAAAATAGAATTTATGCCGGAGGGGAACATTACAGATGTGAAGATTT
GGCAAAGTATTTAACCAGTGCTCCACCTTACTGAACATGAGCATGGTACTGAGGAAAAA
CCCTGCAAAATATGAAGAGTGCAGCAGTGTCTTTATCTCTTGCTCAAGCCTTTCTAATCAA
CAGATGATTTCTGCTGGAGAGAAGCTCTCCAAATGTGAAACATGGTACAAAGGTTTAAAC
CACAGCCCAATCCTTCCAAACACCAGAGAAAATGAGATTGGAGGGAAACCTTTCAAATGT
GAGGAATGTGACAGCATCTTCAAGTGGTCTCAGACCTTACTAAACATAAGAGAATTCAC
ACTGGTGAGAAACCATACAAATGTGACGAATGTGGGAAAGCCTATACACAGTCTCACAC
CTCAGTGAACACAGGAGGATTACACCCGAGAGAAAACCTACCAATGTGAAGAATGTGGG
AAGGCTTTCAGAACTTGTCAAGCCTTTTCAACCATAAGAGAATCATTCTGAAGAAAAA
CCCTACACGTGTGAAGAATGTGGCAACATCTTTAAGCAGTTATCAGACCTCACTAAGCAT
AAGAAAAACCCATACTGGAGAGAAGCCCTACAAATGTGACGAATGTGAAAAAACTTTACC
CAGTCTCCAACCTTATTGTACATAAGAGAATTCATACTGGAGAGAAACCTTACAAGTGT
GAAGAATGTGGCAGAGTCTTCAATGTGGTCTCAGACATTACCAACATAAGAAAAACCAT
ACTGGAGAGAAACCTTACAATGTGACGAATGTGGAAAAAACTTTACCCAGTCTCAAAC
CTTATTGTACATAAGAGAATTCATACTGGAGAGAAACCTTACAAGTGTGAAAAGTGTGGC
AAAGCCTTCAACAGTCTCACACCTGACTGTACATGAAAGCATTCACTACTGA
    
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Clone variation with respect to NM_007152.4

5' Read Nucleotide Sequence:

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>OriGene 5' read for NM_007152 unedited
ATTTTGAATACGAACTCTATAGGGCGCCGCGAATTCGGCACGAGGGTCCCCTGCTAG
CCGTAGGCCGTGTGACCCGACGGCACCAGGAGATCCAGAAGTGAACGCCAGGCTCTCTG
GAGGCCAGGAGATGACTCTGTTGACGTTTCAGGGATGTGGCCATAGAATTTCCCTGGAGG
AGTGGAAATGCCTGGACCTCGCTCAGCAGAATTTGTACAGGGATGTGATGTTGGAGAACT
ACAGAAAATTGTTCTCCGTTGGTCTCACTGTCTGTAAGCCAGGCCTGATCACCTGCCTGG
AGCAACGAAAAGAGCCCTGGAATGTGAAGAGACAGGAGGCAGCAGACGGACATCCAGCTA
TGTCTTCTCATTTTACCAAGACCTTCTGCCAGAGCAGGGCATAACAAGATGCATTCCCAA
AAAGAATACTGAGAGGATATGGAATTTGTGGCCTTGATAATTTATATTTAAGGAAAGACT
GGGAAAGTTTAGATGAGTGAAGTTGCAAAAAGATTATAATGGACTTAACCAATGTTTCA
CAACTACCCATAGCAAAATCTTTCAATATAATAAATATGTTAAAATCTTTGATAACTTTT
CAAATTTACATAGACCGTATATAAGTAATACTGGAGAGAAACCTTTCAAATGTCAAGAAT
GTGGCANATCCTTTCAAATGCTCTCATTCTAATGAACATCAGAAAATTCACACTGGAA
AAAAATTTCAAAAATGTGGAGAATGTGGCANAACCTTTATCCAGTGCTCACACTTACCT
GACCTGAGAACATTGACACTGGNAGAGAACCTTACCAGTGTCAAGAATGGTACAACGTCAT
TTAAAACCTGCTCAGTCTTACTAAAATAGAANTTATGCCCGAG
    
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3' Read Nucleotide Sequence:	>OriGene 3' read for NM_007152 unedited GACCGCGCCGCAATCTAGAGTCGAGTTTTTTTTTTTTTTTTTCTCTCTACAAAATTT TACTCACGCATCTTACATTTTCAGTGTCTTTTACCTTTTACGGACAAGTATATCAGTGATG CACACCTAAGAGAAAAGAATTTCTCATATCTCTGACGCAGCAACAAGTACCATGTGTTT TCACATCAGCACTAGAAATGAAGACACAGCATCTACTGATTTGAGAATAGAATGACATCA ATATTCGCTTTTCAAAAATTTAACATTTTCAATGCAAAAAGTATATTTGAATGTAA TTATAACTCTACAAAACACTCAGTCTCATCCTCTTTTAACTTAAATATAAATGATCTAT TTTAAGTCTAGATGTCTCTGTAGACTCAACACAGGGATTTCAGTGCATTTCTGAACGTGT CCGTGCTCTATTCTTTCTCTTCAAATCCTCCATTGTTTACTTAAATTGGATTTTGCTT ATTTTTCCATTTAGTGTATTTGCAAAATATCTTTTAGTATGAACTCTAGTGTCTTCTAAG CTATCATTTTTGAACAAATGTTATTCCACATTTATGAGATTTGTTGGGTTTCTCTACAAA TTTTCTGAAAGTTTAAGCAACTGATGCAAGTCTTCCATCTAGTATAAATTGAACATTTTT TTCAGAAGAAATACTCTTGTGTGCTCTGGAGACTATATTTTCATGAAAGGCTTTCAATA GTAATTACATTTATGATAACTTTATTAAGTCTGAACTCTGATGTAAGTGGGGATGCGAG CAGATACTAACGGCCTTGCCCTATTTTATTTGTTATTTCTTTCTCAGTATGAATGCTT TCATTACAGTCAGGTGTGAGACTGGGTGAGGCTTTGCCCACTTTTACACTGTAGGTTCT CTCCATATGAATCTCTTAGACATAAGGTTGG
Restriction Sites:	NotI-NotI
ACCN:	NM_007152
Insert Size:	2560 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_007152.1 , NP_009083.1
RefSeq Size:	2394 bp
RefSeq ORF:	1890 bp
Locus ID:	7748
UniProt ID:	O14628
Cytogenetics:	11p15.4
Domains:	KRAB, zf-C2H2
Protein Families:	Transcription Factors

Gene Summary:

This gene encodes a protein belonging to the Krueppel C2H2-type zinc-finger protein family. These family members are transcription factors that are implicated in a variety of cellular processes. This gene is located near the centromeric border of chromosome 11p15.5, next to an imprinted domain that is associated with maternal-specific loss of heterozygosity in Wilms' tumors. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Feb 2012]

Transcript Variant: This variant (3) lacks two alternate in-frame exons in the 3' coding region, compared to variant 1, resulting in an isoform (3) that is shorter than 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.