OriGene Technologies, Inc.
9620 Medical Center Drive, Ste 200
Rockville, MD 20850, US
Phone: +1-888-267-4436
https://www.origene.com
techsupport@origene.com
EU: info-de@origene.com
CN: techsupport@origene.cn

Product datasheet for SC205993

EVC2 (NM_001166136) Human 3' UTR Clone

Product data:

Product Type:	3' UTR Clones
Product Name:	EVC2 (NM_001166136) Human 3' UTR Clone
Vector:	pMirTarget (PS100062)
Symbol:	EVC2
Synonyms:	LBN; WAD
ACCN:	NM_001166136
Insert Size:	439 bp
Insert Sequence:	>SC205993 3'UTR clone of NM_001166136 The sequence shown below is from the reference sequence of NM_001166136. The complete sequence of this clone may contain minor differences, such as SNPs. Blue=Stop Codon Red=Cloning site GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC AAGGCCATGAGGGCCTTGGGCATGGACTAGCCCAAGGGAAAGACCTGCGGGAGCATCTGAAGAGAGAAG GGATGATTTTCTCCTGCCCGCCGGTGTTTGTGTGTTTATAATGCACAACTCGCAAATATAAATTGCACA TGCAGAAGGCACAGACCCCGTAGCGCATGCCAACTTGCAGGGACTCGGTTTAATCTTGTCTCATGAATT TCCAGATGGCCCACTCTCTTCCATATCACAAGGACATAAACACTCCTTCTTTCAGCCCCACCTCCCCAG GGCCCTGGAGGAGACCCCCACCCTGCAATCCACACCCCATCCTCTGCTGCAGAAGCTATGGTCTGTGTG GTGACAGCCAGATTCTCTACTCTTATGTTTTGTATTTGTTACATATTCTATTTTTATAAAGGGAATTTT AAAAAATAAATGTGTTTTGCACAAA ACGCGTAAGCGGCCGCGGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG
Restriction Sites:	SgfI-MluI
OTI Disclaimer:	Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences , e.g., single nucleotide polymorphisms (SNPs).
Components:	The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.
RefSeq:	NM_001166136.2

View online »

Summary: This gene encodes a protein that functions in bone formation and skeletal development. Mutations in this gene, as well as in a neighboring gene that lies in a head-to-head configuration, cause Ellis-van Creveld syndrome, an autosomal recessive skeletal dysplasia that is also known as chondroectodermal dysplasia. Mutations in this gene also cause acrofacial dysostosis Weyers type, also referred to as Curry-Hall syndrome, a disease that combines limb and facial abnormalities. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Oct 2009]

Locus ID: 132884

MW: 16.3