

## Product datasheet for RC228654L3V

## OriGene Technologies, Inc.

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## EVC2 (NM\_001166136) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

Product Type: Lentiviral Particles

**Product Name:** EVC2 (NM\_001166136) Human Tagged ORF Clone Lentiviral Particle

Symbol: EVC2

Synonyms: LBN; WAD

Mammalian Cell Puromycin

Selection:

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK

**ACCN:** NM\_001166136

ORF Size: 3684 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC228654).

Sequence:

**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.

**RefSeq:** NM 001166136.1, NP 001159608.1

 RefSeq Size:
 4864 bp

 RefSeq ORF:
 3687 bp

 Locus ID:
 132884

 UniProt ID:
 Q86UK5

**Cytogenetics:** 4p16.2

**Protein Families:** Transmembrane

MW: 139.9 kDa







## **Gene 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]