

Product datasheet for RC220584L3V

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

MEF2C (NM_002397) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: MEF2C (NM_002397) Human Tagged ORF Clone Lentiviral Particle

Symbol: MEF2C

Synonyms: C5DELq14.3; DEL5q14.3

Mammalian Cell

Selection:

Puromycin

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

 Tag:
 Myc-DDK

 ACCN:
 NM_002397

ORF Size: 1419 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC220584).

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.

RefSeg: NM 002397.2

 RefSeq Size:
 4077 bp

 RefSeq ORF:
 1422 bp

 Locus ID:
 4208

 UniProt ID:
 Q06413

 Cytogenetics:
 5q14.3

Domains: MADS

Protein Families: Transcription Factors





MEF2C (NM_002397) Human Tagged ORF Clone Lentiviral Particle - RC220584L3V

Protein Pathways: MAPK signaling pathway

MW: 51 kDa

Gene Summary: This locus encodes a member of the MADS box transcription enhancer factor 2 (MEF2) family

of proteins, which play a role in myogenesis. The encoded protein, MEF2 polypeptide C, has both trans-activating and DNA binding activities. This protein may play a role in maintaining the differentiated state of muscle cells. Mutations and deletions at this locus have been associated with severe cognitive disability, stereotypic movements, epilepsy, and cerebral malformation. Alternatively spliced transcript variants have been described. [provided by

RefSeq, Jul 2010]