

Product datasheet for RC205682L2V

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

MSX1 (NM 002448) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: MSX1 (NM 002448) Human Tagged ORF Clone Lentiviral Particle

Symbol: MSX^{*}

Synonyms: ECTD3; HOX7; HYD1; STHAG1

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_002448

ORF Size: 909 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC205682).

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 002448.3, NP 002439.2

 RefSeq Size:
 1940 bp

 RefSeq ORF:
 912 bp

 Locus ID:
 4487

 UniProt ID:
 P28360

Cytogenetics: 4p16.2

Domains: homeobox

Protein Families: Druggable Genome, Transcription Factors





ORIGENE

MW: 31.5 kDa

Gene Summary:

This gene encodes a member of the muscle segment homeobox gene family. The encoded protein functions as a transcriptional repressor during embryogenesis through interactions with components of the core transcription complex and other homeoproteins. It may also have roles in limb-pattern formation, craniofacial development, particularly odontogenesis, and tumor growth inhibition. Mutations in this gene, which was once known as homeobox 7, have been associated with nonsyndromic cleft lip with or without cleft palate 5, Witkop syndrome, Wolf-Hirschom syndrome, and autosomoal dominant hypodontia. [provided by RefSeq, Jul 2008]