

Product datasheet for RC219532L4

ECM1 (NM_022664) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: ECM1 (NM_022664) Human Tagged Lenti ORF Clone

Tag: mGFP
Symbol: ECM1
Synonyms: URBWD

Mammalian Cell Puromycin

Selection:

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

E. coli Selection: Chloramphenicol (34 ug/mL)

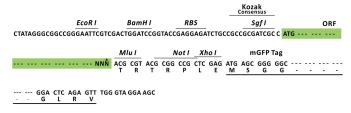
ORF Nucleotide The ORF insert of this clone is exactly the same as(RC219532).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





st The last codon before the Stop codon of the ORF.



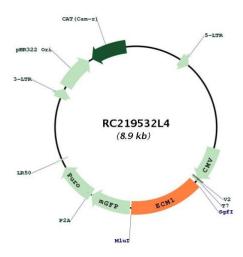
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Plasmid Map:



ACCN: NM_022664

ORF Size: 1245 bp

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.

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: 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: <u>NM 022664.2</u>

RefSeq Size: 1786 bp RefSeq ORF: 1248 bp



ECM1 (NM_022664) Human Tagged Lenti ORF Clone - RC219532L4

 Locus ID:
 1893

 UniProt ID:
 Q16610

 Cytogenetics:
 1q21.2

Protein Families: Secreted Protein, Transmembrane

MW: 46.1 kDa

Gene Summary: This gene encodes a soluble protein that is involved in endochondral bone formation,

angiogenesis, and tumor biology. It also interacts with a variety of extracellular and structural proteins, contributing to the maintenance of skin integrity and homeostasis. Mutations in this

gene are associated with lipoid proteinosis disorder (also known as hyalinosis cutis et mucosae or Urbach-Wiethe disease) that is characterized by generalized thickening of skin, mucosae and certain viscera. Alternatively spliced transcript variants encoding distinct

isoforms have been described for this gene. [provided by RefSeq, Feb 2011]