

Product datasheet for RC202748L2V

OriGene Technologies, Inc.

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Metallothionein (MT2A) (NM_005953) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Metallothionein (MT2A) (NM 005953) Human Tagged ORF Clone Lentiviral Particle

Symbol: Metallothionein
Synonyms: MT-2; MT-II; MT2

Mammalian Cell

Selection:

None

Vector: pLenti-C-mGFP (PS100071)

Tag: mGFP

ACCN: NM_005953

ORF Size: 183 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC202748).

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 005953.2

 RefSeq Size:
 466 bp

 RefSeq ORF:
 186 bp

 Locus ID:
 4502

 UniProt ID:
 P02795

 Cytogenetics:
 16q13

 MW:
 6 kDa





Metallothionein (MT2A) (NM_005953) Human Tagged ORF Clone Lentiviral Particle – RC202748L2V

Gene Summary:

This gene is a member of the metallothionein family of genes. Proteins encoded by this gene family are low in molecular weight, are cysteine-rich, lack aromatic residues, and bind divalent heavy metal ions, altering the intracellular concentration of heavy metals in the cell. These proteins act as anti-oxidants, protect against hydroxyl free radicals, are important in homeostatic control of metal in the cell, and play a role in detoxification of heavy metals. The encoded protein interacts with the protein encoded by the homeobox containing 1 gene in some cell types, controlling intracellular zinc levels, affecting apoptotic and autophagy pathways. Some polymorphisms in this gene are associated with an increased risk of cancer. [provided by RefSeq, Sep 2017]