

## Product datasheet for RC222818L4V

## 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

## NT5C3 (NT5C3A) (NM 001002010) Human Tagged ORF Clone Lentiviral Particle

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** NT5C3 (NT5C3A) (NM\_001002010) Human Tagged ORF Clone Lentiviral Particle

Symbol: NT5C3

Synonyms: cN-III; hUMP1; NT5C3; P5'N-1; P5N-1; p36; PN-I; POMP; PSN1; UMPH; UMPH1

Mammalian Cell

Selection:

Puromycin

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

Tag: mGFP

**ACCN:** NM\_001002010

ORF Size: 1008 bp

**ORF Nucleotide** 

OTI Disclaimer:

The ORF insert of this clone is exactly the same as(RC222818).

Sequence:

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: <u>NM 001002010.1</u>

 RefSeq Size:
 1688 bp

 RefSeq ORF:
 996 bp

 Locus ID:
 51251

 UniProt ID:
 Q9H0P0

Cytogenetics: 7p14.3

**Protein Families:** Transmembrane





## NT5C3 (NT5C3A) (NM\_001002010) Human Tagged ORF Clone Lentiviral Particle - RC222818L4V

**Protein Pathways:** Metabolic pathways, Nicotinate and nicotinamide metabolism, Purine metabolism, Pyrimidine

metabolism

MW: 37.8 kDa

**Gene Summary:** This gene encodes a member of the 5'-nucleotidase family of enzymes that catalyze the

dephosphorylation of nucleoside 5'-monophosphates. The encoded protein is the type 1 isozyme of pyrimidine 5' nucleotidase and catalyzes the dephosphorylation of pyrimidine 5' monophosphates. Mutations in this gene are a cause of hemolytic anemia due to uridine 5-

prime monophosphate hydrolase deficiency. Alternatively spliced transcript variants encoding multiple isoforms have been observed for this gene, and pseudogenes of this gene

are located on the long arm of chromosomes 3 and 4. [provided by RefSeq, Mar 2012]