

Product datasheet for RR212919L4

Aprt (NM_001013061) Rat Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: Aprt (NM_001013061) Rat Tagged Lenti ORF Clone

Tag: mGFP
Symbol: Aprt

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(RR212919).

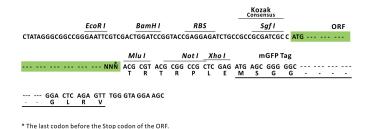
Sequence:

ence.

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





ACCN: NM 001013061

ORF Size: 540 bp



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Aprt (NM_001013061) Rat Tagged Lenti ORF Clone - RR212919L4

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 001013061.1</u>, <u>NP 001013079.1</u>

19q12

 RefSeq Size:
 543 bp

 RefSeq ORF:
 543 bp

 Locus ID:
 292072

 UniProt ID:
 P36972

Cytogenetics:

Gene Summary: Catalyzes a salvage reaction resulting in the formation of AMP, that is energically less costly

than de novo synthesis.[UniProtKB/Swiss-Prot Function]