

## Product datasheet for MR208769L3V

## OriGene Technologies, Inc.

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## **Atp5a1 (NM\_007505) Mouse Tagged ORF Clone Lentiviral Particle**

**Product data:** 

**Product Type:** Lentiviral Particles

**Product Name:** Atp5a1 (NM\_007505) Mouse Tagged ORF Clone Lentiviral Particle

Symbol: Atp5a1

**Synonyms:** Al035633; AL022851; AL023067; Atpm; D18Ertd206e; Mom2

Mammalian Cell

Selection:

Puromycin

**Vector:** pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK
ACCN: NM 007505

ORF Size: 1662 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(MR208769).

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 007505.2, NP 031531.1

 RefSeq Size:
 2443 bp

 RefSeq ORF:
 1662 bp

 Locus ID:
 11946

 UniProt ID:
 Q03265

Cytogenetics: 18 52.38 cM





## **Gene Summary:**

Mitochondrial membrane ATP synthase (F(1)F(0) ATP synthase or Complex V) produces ATP from ADP in the presence of a proton gradient across the membrane which is generated by electron transport complexes of the respiratory chain. F-type ATPases consist of two structural domains, F(1) - containing the extramembraneous catalytic core, and F(0) - containing the membrane proton channel, linked together by a central stalk and a peripheral stalk. During catalysis, ATP synthesis in the catalytic domain of F(1) is coupled via a rotary mechanism of the central stalk subunits to proton translocation. Subunits alpha and beta form the catalytic core in F(1). Rotation of the central stalk against the surrounding alpha(3)beta(3) subunits leads to hydrolysis of ATP in three separate catalytic sites on the beta subunits. Subunit alpha does not bear the catalytic high-affinity ATP-binding sites (By similarity). Binds the bacterial siderophore enterobactin and can promote mitochondrial accumulation of enterobactin-derived iron ions (By similarity). [UniProtKB/Swiss-Prot Function]