

# Product datasheet for MR206733L3V

### OriGene Technologies, Inc.

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## Acp2 (NM 007387) Mouse Tagged ORF Clone Lentiviral Particle

#### **Product data:**

**Product Type:** Lentiviral Particles

**Product Name:** Acp2 (NM\_007387) Mouse Tagged ORF Clone Lentiviral Particle

Symbol: Acp2

Acp; Acp-2; LAP Synonyms:

**Mammalian Cell** 

Selection:

Puromycin

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

Myc-DDK Tag: NM 007387 ACCN:

**ORF Size:** 1272 bp

**ORF Nucleotide** 

Sequence:

The ORF insert of this clone is exactly the same as(MR206733).

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.

RefSeq: NM 007387.1

RefSeq Size: 4669 bp RefSeq ORF: 1272 bp Locus ID: 11432 **UniProt ID:** P24638

Cytogenetics: 2 50.54 cM





#### **Gene Summary:**

The protein encoded by this gene belongs to the histidine acid phosphatase family, which hydrolyze orthophosphoric monoesters to alcohol and phosphate. This protein is localized to the lysosomal membrane, and is chemically and genetically distinct from the red cell acid phosphatase. Mice lacking this gene showed multiple defects, including bone structure alterations, lysosomal storage defects, and an increased tendency towards seizures. An enzymatically-inactive allele of this gene showed severe growth retardation, hair-follicle abnormalities, and an ataxia-like phenotype. Two isoforms are predicted to be produced from the same mRNA by the use of alternative in-frame translation termination codons via a stop codon readthrough mechanism. [provided by RefSeq, Oct 2017]