

## Product datasheet for RC218271L3V

## OriGene Technologies, Inc.

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## NAGPA (NM\_016256) Human Tagged ORF Clone Lentiviral Particle

## **Product data:**

**Product Type:** Lentiviral Particles

Product Name: NAGPA (NM 016256) Human Tagged ORF Clone Lentiviral Particle

Symbol: NAGPA

Synonyms: APAA; UCE

Mammalian Cell Puromycin

Selection:

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

Tag: Myc-DDK

**ACCN:** NM\_016256

ORF Size: 1545 bp

**ORF Nucleotide** 

The ORF insert of this clone is exactly the same as(RC218271).

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 016256.2

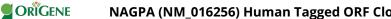
RefSeq Size: 2219 bp
RefSeq ORF: 1548 bp
Locus ID: 51172
UniProt ID: Q9UK23

Cytogenetics: 16p13.3

Domains: EGF

**Protein Families:** Transmembrane





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**Protein Pathways:** Lysosome

MW: 56.11 kDa

Hydrolases are transported to lysosomes after binding to mannose 6-phosphate receptors in **Gene Summary:** 

the trans-Golgi network. This gene encodes the enzyme that catalyzes the second step in the formation of the mannose 6-phosphate recognition marker on lysosomal hydrolases. Commonly known as 'uncovering enzyme' or UCE, this enzyme removes N-acetyl-Dglucosamine (GlcNAc) residues from GlcNAc-alpha-P-mannose moieties and thereby

produces the recognition marker. The encoded preproprotein is proteolytically processed by furin to generate the mature enzyme, a homotetramer of two disulfide-linked homodimers. Mutations in this gene are associated with developmental stuttering in human patients.

[provided by RefSeq, Oct 2015]