

## OriGene Technologies, Inc.

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## Product datasheet for RC201904L4V

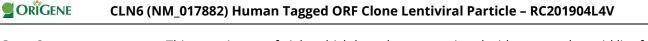
## CLN6 (NM\_017882) Human Tagged ORF Clone Lentiviral Particle

## Product data:

Product Type:	Lentiviral Particles
Product Name:	CLN6 (NM_017882) Human Tagged ORF Clone Lentiviral Particle
Symbol:	CLN6
Synonyms:	CLN4A; HsT18960; nclf
Mammalian Cell Selection:	Puromycin
Vector:	pLenti-C-mGFP-P2A-Puro (PS100093)
Tag:	mGFP
ACCN:	NM_017882
ORF Size:	933 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC201904).
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. <u>More info</u>
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 017882.1</u>
RefSeq Size:	2258 bp
RefSeq ORF:	936 bp
Locus ID:	54982
UniProt ID:	<u>Q9NWW5</u>
Cytogenetics:	15q23
Protein Families:	Transmembrane
MW:	35.9 kDa



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Gene Summary:This gene is one of eight which have been associated with neuronal ceroid lipofuscinoses<br/>(NCL). Also referred to as Batten disease, NCL comprises a class of autosomal recessive,<br/>neurodegenerative disorders affecting children. The genes responsible likely encode proteins<br/>involved in the degradation of post-translationally modified proteins in lysosomes. The<br/>primary defect in NCL disorders is thought to be associated with lysosomal storage function.<br/>[provided by RefSeq, Oct 2008]

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