

## Product datasheet for **RC222295L2V**

### APXL (SHROOM2) (NM\_001649) Human Tagged ORF Clone Lentiviral Particle

#### Product data:

Product Type:	Lentiviral Particles
Product Name:	APXL (SHROOM2) (NM_001649) Human Tagged ORF Clone Lentiviral Particle
Symbol:	APXL
Synonyms:	APXL; HSAPXL
Mammalian Cell Selection:	None
Vector:	pLenti-C-mGFP (PS100071)
Tag:	mGFP
ACCN:	NM_001649
ORF Size:	4848 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC222295).
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. <a href="#">More info</a>
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:	<a href="#">NM_001649.2</a>
RefSeq Size:	7445 bp
RefSeq ORF:	4851 bp
Locus ID:	357
UniProt ID:	<a href="#">Q13796</a>
Cytogenetics:	Xp22.2
Protein Families:	Druggable Genome
MW:	176.2 kDa


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**Gene Summary:**

This gene represents the human homolog of *Xenopus laevis* apical protein (APX) gene, which is implicated in amiloride-sensitive sodium channel activity. It is expressed in endothelial cells and facilitates the formation of a contractile network within endothelial cells. Depletion of this gene results in an increase in endothelial sprouting, migration, and angiogenesis. This gene is highly expressed in the retina, and is a strong candidate for ocular albinism type 1 syndrome. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Mar 2016]