

## Product datasheet for **RC216239L3V**

### SHANK2 (NM\_133266) Human Tagged ORF Clone Lentiviral Particle

#### Product data:

Product Type:	Lentiviral Particles
Product Name:	SHANK2 (NM_133266) Human Tagged ORF Clone Lentiviral Particle
Symbol:	SHANK2
Synonyms:	AUTS17; CORTBP1; CTTNBP1; ProSAP1; SHANK; SPANK-3
Mammalian Cell Selection:	Puromycin
Vector:	pLenti-C-Myc-DDK-P2A-Puro (PS100092)
Tag:	Myc-DDK
ACCN:	NM_133266
ORF Size:	3783 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC216239).
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_133266.3</a>
RefSeq Size:	8872 bp
RefSeq ORF:	3786 bp
Locus ID:	22941
UniProt ID:	<a href="#">Q9UPX8</a>
Cytogenetics:	11q13.3-q13.4
MW:	135.7 kDa


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**Gene Summary:**

This gene encodes a protein that is a member of the Shank family of synaptic proteins that may function as molecular scaffolds in the postsynaptic density of excitatory synapses. Shank proteins contain multiple domains for protein-protein interaction, including ankyrin repeats, and an SH3 domain. This particular family member contains a PDZ domain, a consensus sequence for cortactin SH3 domain-binding peptides and a sterile alpha motif. The alternative splicing demonstrated in Shank genes has been suggested as a mechanism for regulating the molecular structure of Shank and the spectrum of Shank-interacting proteins in the postsynaptic densities of the adult and developing brain. Alterations in the encoded protein may be associated with susceptibility to autism spectrum disorder. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Feb 2014]