

## Product datasheet for **RC206131L3V**

### **GPD1L (NM\_015141) Human Tagged ORF Clone Lentiviral Particle**

#### **Product data:**

Product Type:	Lentiviral Particles
Product Name:	GPD1L (NM_015141) Human Tagged ORF Clone Lentiviral Particle
Symbol:	GPD1L
Synonyms:	GPD1-L
Mammalian Cell Selection:	Puromycin
Vector:	pLenti-C-Myc-DDK-P2A-Puro (PS100092)
Tag:	Myc-DDK
ACCN:	NM_015141
ORF Size:	1053 bp
ORF Nucleotide Sequence:	The ORF insert of this clone is exactly the same as(RC206131).
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_015141.2</a> , <a href="#">NP_055956.1</a>
RefSeq Size:	4068 bp
RefSeq ORF:	1056 bp
Locus ID:	23171
UniProt ID:	<a href="#">Q8N335</a>
Cytogenetics:	3p22.3
Domains:	NAD_Gly3P_dh
Protein Pathways:	Glycerophospholipid metabolism



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**MW:** 38.4 kDa

**Gene Summary:** The protein encoded by this gene catalyzes the conversion of sn-glycerol 3-phosphate to glycerone phosphate. The encoded protein is found in the cytoplasm, associated with the plasma membrane, where it binds the sodium channel, voltage-gated, type V, alpha subunit (SCN5A). Defects in this gene are a cause of Brugada syndrome type 2 (BRS2) as well as sudden infant death syndrome (SIDS). [provided by RefSeq, Jul 2010]