

## Product datasheet for **TL302484**

### PIGQ Human shRNA Plasmid Kit (Locus ID 9091)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	PIGQ Human shRNA Plasmid Kit (Locus ID 9091)
Locus ID:	9091
Synonyms:	c407A10.1; DEE77; EIEE77; GPI1
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	PIGQ - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 9091). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">BC010094</a> , <a href="#">NM_004204</a> , <a href="#">NM_148920</a> , <a href="#">NM_148920.1</a> , <a href="#">NM_148920.2</a> , <a href="#">NM_004204.1</a> , <a href="#">NM_004204.2</a> , <a href="#">NM_004204.3</a> , <a href="#">BC006377</a> , <a href="#">BC006377.2</a> , <a href="#">NM_004204.5</a> , <a href="#">NM_148920.3</a>
UniProt ID:	<a href="#">Q9BRB3</a>
Summary:	This gene is involved in the first step in glycosylphosphatidylinositol (GPI)-anchor biosynthesis. The GPI-anchor is a glycolipid found on many blood cells and serves to anchor proteins to the cell surface. This gene encodes a N-acetylglucosaminyl transferase component that is part of the complex that catalyzes transfer of N-acetylglucosamine (GlcNAc) from UDP-GlcNAc to phosphatidylinositol (PI). Alternatively spliced transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Jun 2012]
shRNA Design:	These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .



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**Performance  
Guaranteed:**

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at [techsupport@origene.com](mailto:techsupport@origene.com). Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).