

## OriGene Technologies, Inc.

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## Product datasheet for TL312749

## alpha 1 Glycine Receptor (GLRA1) Human shRNA Plasmid Kit (Locus ID 2741)

## **Product data:**

Product Type:	shRNA Plasmids
Product Name:	alpha 1 Glycine Receptor (GLRA1) Human shRNA Plasmid Kit (Locus ID 2741)
Locus ID:	2741
Synonyms:	HKPX1; STHE
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	GLRA1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 2741). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	NM 000171, NM 001146040, NM 001292000, NM 000171.1, NM 000171.2, NM 000171.3, NM 001292000.1, BC074980, BC074980.2, BC114947, BC114967, NM 001292000.2, NM 000171.4, NM 001146040.2
UniProt ID:	<u>P23415</u>
Summary:	The protein encoded by this gene is a subunit of a pentameric inhibitory glycine receptor, which mediates postsynaptic inhibition in the central nervous system. Defects in this gene are a cause of startle disease (STHE), also known as hereditary hyperekplexia or congenital stiff- person syndrome. Multiple transcript variants encoding different isoforms have been found. [provided by RefSeq, Dec 2015]
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 <u>techsupport@origene.com</u> . If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .



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Performance	OriGene guarantees that the sequences in the shRNA expression cassettes are verified to
Guaranteed:	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 aPCP to

guaranteed to produce 70% or more gene expression knock-down provided a minimum are 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. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

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