

## **Product datasheet for TR318810**

## **SYT8 Human shRNA Plasmid Kit (Locus ID 90019)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** SYT8 Human shRNA Plasmid Kit (Locus ID 90019)

**Locus ID:** 90019

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: SYT8 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

90019). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: NM 001290332, NM 001290333, NM 001290334, NM 138567, NM 138567.1, NM 138567.2,

NM 138567.3, NM 138567.4, NM 001290334.1, NM 001290333.1, NM 001290332.1, BC038591, BC144657, BC144659, BC144660, BC144661, BC160127, NM 138567.5

UniProt ID: Q8NBV8

**Summary:** This gene encodes a member of the synaptotagmin protein family. Synaptotagmins are

membrane proteins that are important in neurotransmission and hormone secretion, both of which involve regulated exocytosis. Expression of the encoded protein in human pancreatic islets has been connected to activity of the promoter for the insulin gene, on the same chromosome several hundred kilobases away (PMID: 21336277 and 22928559). This

association would link response to gluclose to insulin secretion. Alternative splicing results in multiple transcript variants encoding different isoforms. [provided by RefSeq, Mar 2014]

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 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. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).