

Product datasheet for TL312551

OriGene Technologies, Inc.

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Glycogenin 1 (GYG1) Human shRNA Plasmid Kit (Locus ID 2992)

Product data:

Product Type: shRNA Plasmids

Product Name: Glycogenin 1 (GYG1) Human shRNA Plasmid Kit (Locus ID 2992)

Locus ID: 2992

Synonyms: GSD15; GYG

Vector:pGFP-C-shLenti (TR30023)E. coli Selection:Chloramphenicol (34 ug/ml)

Mammalian Cell

Puromycin

Selection:

Format: Lentiviral plasmids

Components: GYG1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 2992).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 001184720, NM 001184721, NM 004130, NM 004130.1, NM 004130.2, NM 004130.3,

NM 001184721.1, NM 001184720.1, BC000033, BC031096, NM 004130.4

UniProt ID: P46976

Summary: This gene encodes a member of the glycogenin family. Glycogenin is a glycosyltransferase

that catalyzes the formation of a short glucose polymer from uridine diphosphate glucose in an autoglucosylation reaction. This reaction is followed by elongation and branching of the polymer, catalyzed by glycogen synthase and branching enzyme, to form glycogen. This gene is expressed in muscle and other tissues. Mutations in this gene result in glycogen storage

disease XV. This gene has pseudogenes on chromosomes 1, 8 and 13 respectively.

Alternatively spliced transcript variants encoding different isoforms have been identified.

[provided by RefSeq, Sep 2010]

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>.





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).