## Product datasheet for TG312813

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200
Rockville, MD 20850, US
Phone: +1-888-267-4436
https://www.origene.com techsupport@origene.com
EU: info-de@origene.com
CN: techsupport@origene.cn

## GCLC Human shRNA Plasmid Kit (Locus ID 2729)

Product data:

Product Type:
Product Name:
Locus ID:
Synonyms:
Vector:
E. coli Selection:

Mammalian Cell
Selection:
Format:
Components:

RefSeq:

UniProt ID:
Summary:
shRNA Design:
shRNA Plasmids
GCLC Human shRNA Plasmid Kit (Locus ID 2729)
2729
GCL; GCS; GLCL; GLCLC
pGFP-V-RS (TR30007)
Kanamycin
Puromycin

Retroviral plasmids
GCLC - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 2729). $5 \mu \mathrm{~g}$ purified plasmid DNA per construct 29-mer scrambled shRNA cassette in PGFP-V-RS Vector, TR30013, included for free.

NM 001197115 NM 001498, NM 001498.1 NM 001498.2, NM 001498.3 NM 001197115.1, BC039894, BC022487, BC047788, NM 001498.4 NM 001197115.2
P48506
Glutamate-cysteine ligase, also known as gamma-glutamylcysteine synthetase is the first ratelimiting enzyme of glutathione synthesis. The enzyme consists of two subunits, a heavy catalytic subunit and a light regulatory subunit. This locus encodes the catalytic subunit, while the regulatory subunit is derived from a different gene located on chromosome 1p22-p21. Mutations at this locus have been associated with hemolytic anemia due to deficiency of gamma-glutamylcysteine synthetase and susceptibility to myocardial infarction.[provided by RefSeq, Oct 2010]

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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.

## Performance <br> 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).

