

## Product datasheet for **TL312581V**

### Glutathione S Transferase theta 2 (GSTT2) Human shRNA Lentiviral Particle (Locus ID 2953)

#### Product data:

Product Type:	shRNA Lentiviral Particles
Product Name:	Glutathione S Transferase theta 2 (GSTT2) Human shRNA Lentiviral Particle (Locus ID 2953)
Locus ID:	2953
Vector:	pGFP-C-shLenti (TR30023)
Format:	Lentiviral particles
Components:	GSTT2 - Human shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble control), 0.5 ml each, >10 <sup>7</sup> TU/ml.
RefSeq:	<a href="#">NM_000854</a> , <a href="#">NM_001302670</a> , <a href="#">NR_126442</a> , <a href="#">NR_126445</a> , <a href="#">NR_126446</a> , <a href="#">NM_000854.1</a> , <a href="#">NM_000854.2</a> , <a href="#">NM_000854.3</a> , <a href="#">NM_000854.4</a> , <a href="#">NM_001302670.1</a> , <a href="#">BC002415.1</a> , <a href="#">BC146929</a> , <a href="#">BC146940</a> , <a href="#">BC146943</a> , <a href="#">BM804805</a> , <a href="#">BM976631</a>
UniProt ID:	<a href="#">P0CG29</a>
Summary:	The protein encoded by this gene, glutathione S-transferase (GST) theta 2 (GSTT2), is a member of a superfamily of proteins that catalyze the conjugation of reduced glutathione to a variety of electrophilic and hydrophobic compounds. Human GSTs can be divided into five main classes: alpha, mu, pi, theta, and zeta. The theta class includes GSTT1, GSTT2, and GSTT2B. GSTT2 and GSTT2B are nearly identical to each other, and share 55% amino acid identity with GSTT1. All three genes may play a role in human carcinogenesis. The GSTT2 gene is a pseudogene in some populations. [provided by RefSeq, Sep 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 <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).