

## Product datasheet for **TL308494**

### UGT2B7 Human shRNA Plasmid Kit (Locus ID 7364)

#### Product data:

Product Type:	shRNA Plasmids
Product Name:	UGT2B7 Human shRNA Plasmid Kit (Locus ID 7364)
Locus ID:	7364
Synonyms:	UDPGT 2B7; UDPGT2B7; UDPGT 2B9; UDPGTh-2; UDPGTH2; UGT2B9
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	UGT2B7 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 7364). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	<a href="#">NM_001074</a> , <a href="#">NM_001330719</a> , <a href="#">NM_001349568</a> , <a href="#">NM_001074.1</a> , <a href="#">NM_001074.2</a> , <a href="#">NM_001074.3</a> , <a href="#">BC030974</a> , <a href="#">BC030974.1</a> , <a href="#">NM_001074.4</a>
UniProt ID:	<a href="#">P16662</a>
Summary:	The protein encoded by this gene belongs to the UDP-glycosyltransferase (UGT) family. UGTs serve a major role in the conjugation and subsequent elimination of potentially toxic xenobiotics and endogenous compounds. This protein is localized in the microsomal membrane, and has unique specificity for 3,4-catechol estrogens and estriol, suggesting that it may play an important role in regulating the level and activity of these potent estrogen metabolites. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2017]
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).