

## Product datasheet for **TL300782**

### **PHEMX (TSPAN32) Human shRNA Plasmid Kit (Locus ID 10077)**

#### **Product data:**

|                                  |   |
|----------------------------------|---|
| <b>Product Type:</b>             | shRNA Plasmids  |
| <b>Product Name:</b>             | PHEMX (TSPAN32) Human shRNA Plasmid Kit (Locus ID 10077)  |
| <b>Locus ID:</b>                 | 10077   |
| <b>Synonyms:</b>                 | PHMX, PHEMX, TSSC6, MGC22455  |
| <b>Vector:</b>                   | pGFP-C-shLenti (TR30023)  |
| <b>E. coli Selection:</b>        | Chloramphenicol (34 ug/ml)  |
| <b>Mammalian Cell Selection:</b> | Puromycin   |
| <b>Format:</b>                   | Lentiviral plasmids   |
| <b>Components:</b>               | TSPAN32 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 10077). 5µg purified plasmid DNA per construct<br>29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.   |
| <b>RefSeq:</b>                   | <a href="#">NM_005705</a> , <a href="#">NM_139022</a> , <a href="#">NM_139023</a> , <a href="#">NM_139024</a> , <a href="#">NM_139022.1</a> , <a href="#">NM_139022.2</a> ,<br><a href="#">NM_005705.4</a> , <a href="#">NM_139024.3</a> , <a href="#">BC016693</a> , <a href="#">NM_139022.3</a>   |
| <b>UniProt ID:</b>               | <a href="#">Q96QS1</a>  |
| <b>Summary:</b>                  | This gene, which is a member of the tetraspanin superfamily, is one of several tumor-suppressing subtransferable fragments located in the imprinted gene domain of chromosome 11p15.5, an important tumor-suppressor gene region. Alterations in this region have been associated with Beckwith-Wiedemann syndrome, Wilms tumor, rhabdomyosarcoma, adrenocortical carcinoma, and lung, ovarian and breast cancers. This gene is located among several imprinted genes; however, this gene, as well as the tumor-suppressing subchromosomal transferable fragment 4, escapes imprinting. This gene may play a role in malignancies and diseases that involve this region, and it is also involved in hematopoietic cell function. Alternatively spliced transcript variants have been described, but their biological validity has not been determined. [provided by RefSeq, Jul 2008] |
| <b>shRNA Design:</b>             | 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).