

Product datasheet for **TG320572**

WNT5A Human shRNA Plasmid Kit (Locus ID 7474)

Product data:

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| Product Type: | shRNA Plasmids |
| Product Name: | WNT5A Human shRNA Plasmid Kit (Locus ID 7474) |
| Locus ID: | 7474 |
| Synonyms: | hWNT5A |
| Vector: | pGFP-V-RS (TR30007) |
| E. coli Selection: | Kanamycin |
| Mammalian Cell Selection: | Puromycin |
| Format: | Retroviral plasmids |
| Components: | WNT5A - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 7474). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free. |
| RefSeq: | NM_001256105 , NM_003392 , NM_003392.1 , NM_003392.2 , NM_003392.3 , NM_003392.4 , NM_001256105.1 , BC064694 , BC064694.1 , BC074783 , BM728706 , NM_003392.7 |
| UniProt ID: | P41221 |
| Summary: | The WNT gene family consists of structurally related genes which encode secreted signaling proteins. These proteins have been implicated in oncogenesis and in several developmental processes, including regulation of cell fate and patterning during embryogenesis. This gene encodes a member of the WNT family that signals through both the canonical and non-canonical WNT pathways. This protein is a ligand for the seven transmembrane receptor frizzled-5 and the tyrosine kinase orphan receptor 2. This protein plays an essential role in regulating developmental pathways during embryogenesis. This protein may also play a role in oncogenesis. Mutations in this gene are the cause of autosomal dominant Robinow syndrome. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Jan 2012] |
| 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 techsupport@origene.com . If you need a special design or shRNA sequence, please utilize our custom shRNA service . |



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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. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).