

Product datasheet for **TL315676**

GCP4 (TUBGCP4) Human shRNA Plasmid Kit (Locus ID 27229)

Product data:

Product Type:	shRNA Plasmids
Product Name:	GCP4 (TUBGCP4) Human shRNA Plasmid Kit (Locus ID 27229)
Locus ID:	27229
Synonyms:	76P; GCP-4; GCP4; Grip76; MCCR3
Vector:	pGFP-C-shLenti (TR30023)
E. coli Selection:	Chloramphenicol (34 ug/ml)
Mammalian Cell Selection:	Puromycin
Format:	Lentiviral plasmids
Components:	TUBGCP4 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 27229). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.
RefSeq:	BC000966 , BC009870 , NM_001286414 , NM_014444 , NM_014444.1 , NM_014444.2 , NM_014444.3 , NM_014444.4 , NM_001286414.1 , NM_001286414.2 , BC000966.2 , BC009870.2 , BC012801 , BC012801.2 , NM_001286414.3 , NM_014444.5
UniProt ID:	Q9UGJ1
Summary:	This gene encodes a component of the gamma-tubulin ring complex, which is required for microtubule nucleation. In mammalian cells, the protein localizes to centrosomes in association with gamma-tubulin. Crystal structure analysis revealed a structure composed of five helical bundles arranged around conserved hydrophobic cores. An exposed surface area located in the C-terminal domain is essential and sufficient for direct binding to gamma-tubulin. Mutations in this gene that alter microtubule organization are associated with microcephaly and chorioretinopathy. Alternative splicing results in multiple transcript variants. [provided by RefSeq, May 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 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).