

Product datasheet for **TG304883**

DUSP10 Human shRNA Plasmid Kit (Locus ID 11221)

Product data:

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|---------------------------|---|
| Product Type: | shRNA Plasmids |
| Product Name: | DUSP10 Human shRNA Plasmid Kit (Locus ID 11221) |
| Locus ID: | 11221 |
| Synonyms: | MKP-5; MKP5 |
| Vector: | pGFP-V-RS (TR30007) |
| E. coli Selection: | Kanamycin |
| Mammalian Cell Selection: | Puromycin |
| Format: | Retroviral plasmids |
| Components: | DUSP10 - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 11221). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free. |
| RefSeq: | NM_007207 , NM_144728 , NM_144729 , NR_111939 , NR_111940 , NM_007207.2 , NM_007207.3 , NM_007207.4 , NM_007207.5 , NM_144729.1 , NM_144729.2 , NM_144728.1 , BC031405 , BC031405.1 , BC020608 , BC063826 , NM_007207.6 |
| UniProt ID: | Q9Y6W6 |
| Summary: | Dual specificity protein phosphatases inactivate their target kinases by dephosphorylating both the phosphoserine/threonine and phosphotyrosine residues. They negatively regulate members of the MAP kinase superfamily, which is associated with cellular proliferation and differentiation. Different members of this family of dual specificity phosphatases show distinct substrate specificities for MAP kinases, different tissue distribution and subcellular localization, and different modes of expression induction by extracellular stimuli. This gene product binds to and inactivates p38 and SAPK/JNK. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Apr 2014] |
| 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).