

Product datasheet for **TG500619**

Ephb4 Mouse shRNA Plasmid (Locus ID 13846)

Product data:

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| Product Type: | shRNA Plasmids |
| Product Name: | Ephb4 Mouse shRNA Plasmid (Locus ID 13846) |
| Locus ID: | 13846 |
| Synonyms: | AI042935; Htk; MDK2; Myk1; Tyro11 |
| Vector: | pGFP-V-RS (TR30007) |
| E. coli Selection: | Kanamycin |
| Mammalian Cell Selection: | Puromycin |
| Format: | Retroviral plasmids |
| Components: | Ephb4 - Mouse, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 13846). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free. |
| RefSeq: | BC016505 , NM_001159571 , NM_010144 , NM_010144.1 , NM_010144.2 , NM_010144.3 , NM_010144.4 , NM_010144.5 , NM_010144.6 , NM_001159571.1 , BC013342 , BC090839 |
| UniProt ID: | P54761 |
| Summary: | Receptor tyrosine kinase which binds promiscuously transmembrane ephrin-B family ligands residing on adjacent cells, leading to contact-dependent bidirectional signaling into neighboring cells. The signaling pathway downstream of the receptor is referred to as forward signaling while the signaling pathway downstream of the ephrin ligand is referred to as reverse signaling. Together with its cognate ligand/functional ligand EFNB2 it is involved in the regulation of cell adhesion and migration, and plays a central role in heart morphogenesis, angiogenesis and blood vessel remodeling and permeability. EPHB4-mediated forward signaling controls cellular repulsion and segregation from EFNB2-expressing cells.[UniProtKB/Swiss-Prot Function] |
| 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).