

Product datasheet for **TG306568**

beta Arrestin 1 (ARRB1) Human shRNA Plasmid Kit (Locus ID 408)

Product data:

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| Product Type: | shRNA Plasmids |
| Product Name: | beta Arrestin 1 (ARRB1) Human shRNA Plasmid Kit (Locus ID 408) |
| Locus ID: | 408 |
| Synonyms: | ARB1; ARR1 |
| Vector: | pGFP-V-RS (TR30007) |
| E. coli Selection: | Kanamycin |
| Mammalian Cell Selection: | Puromycin |
| Format: | Retroviral plasmids |
| Components: | ARRB1 - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 408). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free. |
| RefSeq: | BC003636 , NM_004041 , NM_020251 , NM_004041.1 , NM_004041.2 , NM_004041.3 , NM_004041.4 , NM_020251.1 , NM_020251.2 , BC003636.2 , BM681392 , NM_004041.5 , NM_020251.4 |
| UniProt ID: | P49407 |
| Summary: | Members of arrestin/beta-arrestin protein family are thought to participate in agonist-mediated desensitization of G-protein-coupled receptors and cause specific dampening of cellular responses to stimuli such as hormones, neurotransmitters, or sensory signals. Arrestin beta 1 is a cytosolic protein and acts as a cofactor in the beta-adrenergic receptor kinase (BARK) mediated desensitization of beta-adrenergic receptors. Besides the central nervous system, it is expressed at high levels in peripheral blood leukocytes, and thus the BARK/beta-arrestin system is believed to play a major role in regulating receptor-mediated immune functions. Alternatively spliced transcripts encoding different isoforms of arrestin beta 1 have been described. [provided by RefSeq, Jan 2011] |
| 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).