

Product datasheet for **TL320522**

Retinoid X Receptor alpha (RXRA) Human shRNA Plasmid Kit (Locus ID 6256)

Product data:

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| Product Type: | shRNA Plasmids |
| Product Name: | Retinoid X Receptor alpha (RXRA) Human shRNA Plasmid Kit (Locus ID 6256) |
| Locus ID: | 6256 |
| Synonyms: | NR2B1 |
| Vector: | pGFP-C-shLenti (TR30023) |
| E. coli Selection: | Chloramphenicol (34 ug/ml) |
| Mammalian Cell Selection: | Puromycin |
| Format: | Lentiviral plasmids |
| Components: | RXRA - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 6256). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free. |
| RefSeq: | NM_001291920 , NM_001291921 , NM_002957 , NM_002957.1 , NM_002957.2 , NM_002957.3 , NM_002957.4 , NM_002957.5 , NM_001291921.1 , NM_001291920.1 , BC110998 , BC110998.1 , BC007925 , BC009882 , BC031356 , BC063827 , BC096246 , BM563384 , NM_001291921.2 , NM_002957.6 |
| UniProt ID: | P19793 |
| Summary: | Retinoid X receptors (RXRs) and retinoic acid receptors (RARs) are nuclear receptors that mediate the biological effects of retinoids by their involvement in retinoic acid-mediated gene activation. These receptors function as transcription factors by binding as homodimers or heterodimers to specific sequences in the promoters of target genes. The protein encoded by this gene is a member of the steroid and thyroid hormone receptor superfamily of transcriptional regulators. Alternative splicing of this gene results in multiple transcript variants. [provided by RefSeq, May 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).