

# Product datasheet for TR701809

## Akr1c2 Rat shRNA Plasmid (Locus ID 291283)

## **Product data:**

#### OriGene Technologies, Inc.

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| Product Type:                | shRNA Plasmids  |
|------------------------------|---|
| Product Name:                | Akr1c2 Rat shRNA Plasmid (Locus ID 291283)  |
| Locus ID:                    | 291283  |
| Synonyms:                    | Akr1c21   |
| Vector:                      | pRS (TR20003)   |
| E. coli Selection:           | Ampicillin  |
| Mammalian Cell<br>Selection: | Puromycin   |
| Format:                      | Retroviral plasmids   |
| Components:                  | Akr1c21 - Rat, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =<br>291283). 5μg purified plasmid DNA per construct<br>29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.   |
| RefSeq:                      | <u>NM 001013057, NM 001013057.1, BC078957</u>   |
| UniProt ID:                  | Q6AYQ2  |
| Summary:                     | NADP-dependent 17-alpha-hydroxysteroid dehydrogenase that converts 5-alpha-androstane-<br>3,17-dione into androsterone. Has lower 3-alpha-hydroxysteroid dehydrogenase activity. Has<br>broad substrate specificity and acts on various 17-alpha-hydroxysteroids, 17-ketosteroids, 3-<br>alpha hydroxysteroids and 3-ketosteroids. Reduction of keto groups is strictly stereoselective.<br>Reduction of 17-ketosteroids yields only 17-alpha-hydroxysteroids. Likewise, reduction of 3-<br>ketosteroids yields only 3-alpha-hydroxysteroids (By similarity).[UniProtKB/Swiss-Prot<br>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 <u>techsupport@origene.com</u> . If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .   |



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### Serigene Akr1c2 Rat shRNA Plasmid (Locus ID 291283) – TR701809

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).

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