

## **Product datasheet for TL306779**

## OriGene Technologies, Inc.

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## **AKAP11 Human shRNA Plasmid Kit (Locus ID 11215)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** AKAP11 Human shRNA Plasmid Kit (Locus ID 11215)

**Locus ID:** 11215

Synonyms: AKAP-11; AKAP220; PPP1R44; PRKA11

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

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Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** AKAP11 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 11215).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 016248, NM 144490, NM 016248.1, NM 016248.2, NM 016248.3, NM 144490.1,

BC156135, BC172498, NM 016248.4

UniProt ID: 09UKA4

Summary: The A-kinase anchor proteins (AKAPs) are a group of structurally diverse proteins, which have

the common function of binding to the regulatory subunit of protein kinase A (PKA) and confining the holoenzyme to discrete locations within the cell. This gene encodes a member

of the AKAP family. The encoded protein is expressed at high levels throughout

spermatogenesis and in mature sperm. It binds the RI and RII subunits of PKA in testis. It may serve a function in cell cycle control of both somatic cells and germ cells in addition to its putative role in spermatogenesis and sperm function. [provided by RefSeq, Jul 2008]

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>.







## 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).