

## **Product datasheet for TR518337**

## Atxn7l3 Mouse shRNA Plasmid (Locus ID 217218)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Atxn7l3 Mouse shRNA Plasmid (Locus ID 217218)

**Locus ID:** 217218

**Synonyms:** E030022H21Rik

Vector: pRS (TR20003)

**E. coli Selection:** Ampicillin

Mammalian Cell

Selection:

Puromycin

Format: Retroviral plasmids

Components: Atxn7l3 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

217218). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: NM 001098836, NM 001098837, NM 001098836.1, NM 001098837.1, BC059880, BC130269,

BM950535

UniProt ID: A2AWT3

Summary: Component of the transcription regulatory histone acetylation (HAT) complex SAGA, a

multiprotein complex that activates transcription by remodeling chromatin and mediating

histone acetylation and deubiquitination. Within the SAGA complex, participates in a

subcomplex that specifically deubiquitinates both histones H2A and H2B. The SAGA complex is recruited to specific gene promoters by activators such as MYC, where it is required for transcription. Required for nuclear receptor-mediated transactivation. Within the complex, it

is required to recruit USP22 and ENY2 into the SAGA complex. Regulates H2B

monoubiquitination (H2Bub1) levels. Affects subcellular distribution of ENY2, USP22 and

ATXN7L3B.[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 <u>techsupport@origene.com</u>.

If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.



**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



## 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).