

## **Product datasheet for TL308219**

## OriGene Technologies, Inc.

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## **ZNF215 Human shRNA Plasmid Kit (Locus ID 7762)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** ZNF215 Human shRNA Plasmid Kit (Locus ID 7762)

**Locus ID:** 7762

Synonyms: BAZ-2; BAZ2; ZKSCAN11; ZSCAN43

**Vector:** pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** ZNF215 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 7762).

5µg purified plasmid DNA per construct

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

RefSeq: NM 013250, NM 001354853, NM 001354854, NM 001354855, NM 001354856,

NM 001354857, NM 001354858, NM 001354859, NM 001354860, NR 149005, NM 013250.1,

NM 013250.2, BC014538, NM 013250.4

UniProt ID: Q9UL58

**Summary:** This gene is imprinted in a tissue-specific manner with preferential expression in the testis,

and encodes a zinc finger protein that belongs to a family of zinc finger transcription factors. The encoded protein contains an N-terminal SRE-ZBP, Ctfin51, AW-1, and Number 18 (SCAN)

domain, a kruppel-associated box A (KRABA) domain, and four C-terminal zinc finger

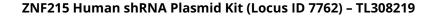
domains. This gene is located within one of three regions on chromosome 11p15 associated with Beckwith-Wiedemann syndrome, called Beckwith-Wiedemann syndrome chromosome region-2 (BWSCR2), and is thought to play a role in the etiology of this disease. [provided by

RefSeq, Aug 2017]

**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 custom shRNA service.







## 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).