

## **Product datasheet for TL313902**

## 9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com

OriGene Technologies, Inc.

EU: info-de@origene.com CN: techsupport@origene.cn

## **CIB2 Human shRNA Plasmid Kit (Locus ID 10518)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** CIB2 Human shRNA Plasmid Kit (Locus ID 10518)

**Locus ID:** 10518

**Synonyms:** DFNB48; KIP2; USH1J

Vector: pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

**Components:** CIB2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 10518).

5µg purified plasmid DNA per construct

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

**RefSeq:** <u>NM 001271888</u>, <u>NM 001271889</u>, <u>NM 001301224</u>, <u>NM 006383</u>, <u>NR 125435</u>, <u>NM 006383.1</u>,

NM 006383.2, NM 006383.3, NM 001271889.1, NM 001271888.1, NM 001301224.1,

BC047381, BC047381.1, BC033108, BC057292, BM908035, NM 001301224.2,

NM 001271888.2, NM 001271889.2

UniProt ID: 075838

**Summary:** The protein encoded by this gene is similar to that of KIP/CIB, calcineurin B, and calmodulin.

The encoded protein is a calcium-binding regulatory protein that interacts with DNA-dependent protein kinase catalytic subunits (DNA-PKcs), and it is involved in photoreceptor

cell maintenance. Mutations in this gene cause deafness, autosomal recessive, 48 (DFNB48), and also Usher syndrome 1J (USH1). Alternative splicing results in multiple transcript

variants. [provided by RefSeq, Jul 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 <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).