

## **Product datasheet for TG305473**

## OriGene Technologies, Inc.

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## CDK5RAP2 Human shRNA Plasmid Kit (Locus ID 55755)

**Product data:** 

**Product Type:** shRNA Plasmids

Product Name: CDK5RAP2 Human shRNA Plasmid Kit (Locus ID 55755)

**Locus ID:** 55755

Synonyms: C48; Cep215; MCPH3

**Vector:** pGFP-V-RS (TR30007)

E. coli Selection: Kanamycin

Mammalian Cell Puromycin

Selection:

Format:

Retroviral plasmids

Components: CDK5RAP2 - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID =

55755). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free.

RefSeq: BC019577, NM 001011649, NM 001272039, NM 018249, NR 073554, NR 073555, NR 073556,

NR 073557, NR 073558, NM 018249.1, NM 018249.2, NM 018249.4, NM 018249.5,

NM 001011649.1, NM 001011649.2, NM 001272039.1, BC019577.1, BC004526, BC136275, BC143732, BC143734, BC143753, BC143755, BC143760, BC143762, BC143764, BC146782,

NM 018249.6, NM 001011649.3

UniProt ID: Q96SN8

Summary: This gene encodes a regulator of CDK5 (cyclin-dependent kinase 5) activity. The protein

encoded by this gene is localized to the centrosome and Golgi complex, interacts with CDK5R1 and pericentrin (PCNT), plays a role in centriole engagement and microtubule

nucleation, and has been linked to primary microcephaly and Alzheimer's disease. Alternative

splicing results in multiple transcript variants. [provided by RefSeq, Jan 2013]

**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).