

## Product datasheet for **TG309983**

### **RAB7 (RAB7A) Human shRNA Plasmid Kit (Locus ID 7879)**

#### **Product data:**

Product Type:	shRNA Plasmids
Product Name:	RAB7 (RAB7A) Human shRNA Plasmid Kit (Locus ID 7879)
Locus ID:	7879
Synonyms:	CMT2B; PRO2706; RAB7
Vector:	pGFP-V-RS (TR30007)
E. coli Selection:	Kanamycin
Mammalian Cell Selection:	Puromycin
Format:	Retroviral plasmids
Components:	RAB7A - Human, 4 unique 29mer shRNA constructs in retroviral GFP vector(Gene ID = 7879). 5µg purified plasmid DNA per construct 29-mer scrambled shRNA cassette in pGFP-V-RS Vector, TR30013, included for free.
RefSeq:	<a href="#">NM_004637</a> , <a href="#">NM_004637.1</a> , <a href="#">NM_004637.2</a> , <a href="#">NM_004637.3</a> , <a href="#">NM_004637.4</a> , <a href="#">NM_004637.5</a> , <a href="#">BC008721</a> , <a href="#">BC008721.2</a> , <a href="#">BC013728</a> , <a href="#">BC014200</a> , <a href="#">NM_004637.6</a>
UniProt ID:	<a href="#">P51149</a>
Summary:	RAB family members are small, RAS-related GTP-binding proteins that are important regulators of vesicular transport. Each RAB protein targets multiple proteins that act in exocytic / endocytic pathways. This gene encodes a RAB family member that regulates vesicle traffic in the late endosomes and also from late endosomes to lysosomes. This encoded protein is also involved in the cellular vacuolation of the VacA cytotoxin of Helicobacter pylori. Mutations at highly conserved amino acid residues in this gene have caused some forms of Charcot-Marie-Tooth (CMT) type 2 neuropathies. [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 <a href="mailto:techsupport@origene.com">techsupport@origene.com</a> . If you need a special design or shRNA sequence, please utilize our <a href="#">custom shRNA service</a> .



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**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](mailto: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).