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## **Human C160RF47 shRNA in Retroviral Vector**

Overview	
Quantity:	1 kit
Gene:	C16orf47 (C16ORF47)
Species:	Human
Insert:	shRNA
Vector:	Retroviral Vector
Application:	RNA Interference (RNAi)
Product Details	
Purpose:	Pre-designed Hush-29 shRNAs in viral vectors with proven effectiveness for knock-down of Human C16orf47.
Brand:	HuSH-29™
Vector Backbone:	pRS
Promoter:	U6 Promoter
Selectable Marker:	Puromycin
Bacterial Resistance:	Ampicillin
Expression Type:	Transient, Stable
Specificity:	<ul> <li>The HuSH shRNA gene-specific expression cassettes were optimized to include both the termination signal for RNA Pol III and GC content targeted at 50 % to further improve the quality of the gene-specific shRNA expression vectors.</li> <li>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.</li> </ul>
Characteristics:	<ul> <li>The shRNA gene-specific expression cassettes are prepared using synthetic oligonucleotides.</li> <li>These oligonucleotide sequences were computer designed for optimal suppression of gene</li> </ul>

## **Product Details** expression and minimal off-target effects. · All shRNA sequences are verified through DNA sequencing analysis. • Gene-specific shRNA expression pRS vectors, 5 ug plasmid DNA per vial. Components: · Four unique constructs per gene. · HuSH 29-mer NonEffective Scrambled pRS 5 ug plasmid DNA. **Target Details** C16orf47 (C16ORF47) Gene: Alternative Name: C16orf47 **Application Details** · Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA Application Notes: constructs 72 hrs post transfection.

Restrictions: For Research Use only

### Handling

Format:

Lyophilized

Storage:

4 °C/-20 °C

Storage Comment:

The dried plasmids can be stored at 4°C. However, once reconstituted with dH2O, the plasmids must be stored at -20°C.

• 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..

#### **Publications**

Product cited in: Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (1991)