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Datasheet for ABIN3760409 Human C3P1 shRNA in Lentiviral Vector (GFP tag)

Overview

Quantity:	1 kit
Gene:	C3P1
Species:	Human
Fusion tag:	GFP tag
Insert:	shRNA
Vector:	Lentiviral Vector
Application:	RNA Interference (RNAi)

Product Details

Purpose:	Pre-designed Hush-29 shRNAs in viral vectors with proven effectiveness for knock-down of Human LOC388503.
Brand:	HuSH-29™
Vector Backbone:	pGFP-C-shLenti
Promoter:	U6 Promoter
Selectable Marker:	Puromycin
Bacterial Resistance:	Chloramphenicol
Expression Type:	Transient, Stable
Specificity:	 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. 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.
Characteristics:	The shRNA gene-specific expression cassettes are prepared using synthetic

Product Details	
	 oligonucleotides. These oligonucleotide sequences were computer designed for optimal suppression of gene expression and minimal off-target effects. All shRNA sequences are verified through DNA sequencing analysis.
Components:	 Gene-specific shRNA in pGFPC-shLenti vector, 4 unique constructs per gene, 5 ug per vial. HuSH 29-mer Scrambled in pGFP-C-shLenti 5 ug plasmid DNA.

Target Details

C3P1
LOC388503 (C3P1 Products)
• 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 Research Use only
Lyophilized
4 °C/-20 °C
The dried plasmids can be stored at 4°C. However, once reconstituted with dH2O, the plasmids must be stored at -20°C.

Publications

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