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Human MKRN4P shRNA in Retroviral Vector

Overview		
Quantity:	1 kit	
Gene:	MKRN4P	
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 MKRN4.	
Brand:	HuSH-29™	
Vector Backbone:	pRS	
Promoter:	U6 Promoter	
Selectable Marker:	Puromycin	
Bacterial Resistance:	Ampicillin	
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 oligonucleotides. These oligonucleotide sequences were computer designed for optimal suppression of gene 	

expression and	minimal	off-target	effects.
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- All shRNA sequences are verified through DNA sequencing analysis.
- Components:
- Gene-specific shRNA expression pRS vectors, 5 ug plasmid DNA per vial.
- · Four unique constructs per gene.
- HuSH 29-mer NonEffective Scrambled pRS 5 ug plasmid DNA.

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Target Details

- Target Details	
Gene:	MKRN4P
Alternative Name:	MKRN4
Application Details	
Application Notes:	 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.
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.
Publications	
Product cited in:	Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (