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Human C20RF72 shRNA in Retroviral Vector

Overview			
Quantity:	1 kit		
Gene:	C20RF72		
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 C2orf72.		
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 		

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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

Gene:	C20RF72	
Alternative Name:	C2orf72 (C2ORF72 Products)	
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, (