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Human NBPF14 shRNA in Retroviral Vector

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Characteristics:

Overview		
Quantity:	1 kit	
Gene:	NBPF14	
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 NBPF14.	
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. 	

• 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. 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** Gene: NBPF14 Alternative Name: NBPF14 **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, (

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