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Human OR4F16 shRNA in Retroviral Vector

| Overview | |
|-----------------------|--|
| Quantity: | 1 kit |
| Gene: | OR4F16 |
| 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 OR4F16. |
| 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 |

Product Details expression and minimal off-target effects. • 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. Target Details Gene: OR4F16 Alternative Name: OR4F16 (OR4F16 Products) Application Details • Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA

constructs 72 hrs post transfection.

For Research Use only

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Handling

Restrictions:

| 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, (|

• 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..