-online.com **denomics**





Human RIMBP3B shRNA in Retroviral Vector

| Overview | |
|-----------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Quantity: | 1 kit |
| Gene: | RIMBP3B |
| 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 RIMBP3B. |
| 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: RIMBP3B Alternative Name: RIMBP3B Application Details Application Notes: Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA

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