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## Human NPY4R CRISPR gRNA + Cas9 in Lenti Particles

| Overview              |   |
|-----------------------|---|
| Quantity:             | 300 μL  |
| Gene:                 | NPY4R   |
| Species:              | Human   |
| Insert:               | gRNA + Cas9   |
| Vector:               | Lentiviral Vector   |
| Application:          | Protein Expression (PExp), Genome Editing with Engineered Nucleases (GEEN)                    |
| Product Details       |   |
| Purpose:              | Individual gRNA against PPYR1 in Lentiviral Particles with a Titer of >1x10e7 IU/mL. (sgRNA   |
|                       | and Cas9 in a single vector)  |
| Vector Backbone:      | pLenti-U6-sgRNA-SFFV-Cas9-2A-Puro   |
| Promoter:             | U6 Promoter, SFFV Promoter  |
| Selectable Marker:    | Puromycin   |
| Bacterial Resistance: | Ampicillin  |
| Expression Type:      | Stable, Transient   |
| Sequence:             | Sequence available upon placing order   |
| Specificity:          | GRNAs are designed for use with Cas9 Nuclease only.   |
|                       | Cas9 Nuclease is under the control of the SFFV promoter which should work for a vast majority |
|                       | of cells, except ES cells or iPS cells.   |
| Sequencing Primer:    | U6 Forward Primer: 5'TACGTCCAAGGTCGGGCAGGAAGA3'   |
| Components:           | Lentiviral particles with an individual gRNA (300 μL) for a specific sequence of PPYR1        |

## **Target Details**

| Gene:             | NPY4R                  |
|-------------------|------------------------|
| Alternative Name: | PPYR1 (NPY4R Products) |
| NCBI Accession:   | NM_005972              |

| pplication Notes: | Recommended for quality control: Restriction Enzyme Digest and Sequencing |
|-------------------|---|
| estrictions:      | For Research Use only   |
| Handling          |   |
| rmat:             | Viral Particles   |
| orage:            | -80 °C  |
| piry Date:        | 12 months   |

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