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Mouse CTNND1 CRISPR gRNA + Cas9 in Lenti Particles

| Overview | |
|-----------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Quantity: | 300 μL |
| Gene: | CTNND1 |
| Species: | Mouse |
| Insert: | gRNA + Cas9 |
| Vector: | Lentiviral Vector |
| Application: | Protein Expression (PExp), Genome Editing with Engineered Nucleases (GEEN) |
| Product Details | |
| Purpose: | Individual gRNA against Ctnnd1 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 Ctnnd1 |

Target Details

| Gene: | CTNND1 |
|-------------------|--------------------------|
| Alternative Name: | Ctnnd1 (CTNND1 Products) |
| NCBI Accession: | NM_007615 |

| Application Notes: | Recommended for quality control: Restriction Enzyme Digest and Sequencing |
|--------------------|-----------------------------------------------------------------------------------------|
| Restrictions: | For Research Use only |
| Handling | |
| Format: | Viral Particles |
| Storage: | -80 °C |
| Expiry Date: | 12 months |
| Publications | |
| Product cited in: | Johnson Drugan Millar Evans: "29" in: Val. 1363 Jesua Nuclaia saide research, pp. 29-20 |

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