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Datasheet for ABIN5297959 Human ATP11C CRISPR gRNA + Cas9 in Lenti Particles

| Overview | |
|-----------------------|---|
| Quantity: | 300 µL |
| Gene: | ATP11C |
| Species: | Human |
| Insert: | gRNA + Cas9 |
| Vector: | Lentiviral Vector |
| Application: | Protein Expression (PExp), Genome Editing with Engineered Nucleases (GEEN) |
| Product Details | |
| Purpose: | Individual gRNA against ATP11C 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. |

U6 Forward Primer: 5'--TACGTCCAAGGTCGGGCAGGAAGA--3'

of cells, except ES cells or iPS cells.

Sequencing Primer:

Components:

Cas9 Nuclease is under the control of the SFFV promoter which should work for a vast majority

Lentiviral particles with an individual gRNA (300 μ L) for a specific sequence of ATP11C

| Target Details | | |
|---------------------|---|--|
| Gene: | ATP11C | |
| Alternative Name: | ATP11C (ATP11C Products) | |
| NCBI Accession: | NM_001010986 | |
| Application Details | | |
| 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, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (1991) | |