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### Human TAP1 CRISPR gRNA + Cas9 in Lenti Particles

| Overview              |  |
|-----------------------|--|
| Quantity:             | 300 μL   |
| Gene:                 | TAP1   |
| Species:              | Human  |
| Insert:               | gRNA + Cas9  |
| Vector:               | Lentiviral Vector  |
| Application:          | Protein Expression (PExp), Genome Editing with Engineered Nucleases (GEEN)                     |
| Product Details       |  |
| Purpose:              | Individual gRNA against TAP1 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 TAP1          |

# Target Details Gene: TAP1

Alternative Name: TAP1 (TAP1 Products)

NCBI Accession: NM\_000593

#### **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)