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## Human F11 CRISPR gRNA + Cas9 in Lenti Particles

| Overview              |                                                                                               |
|-----------------------|-----------------------------------------------------------------------------------------------|
| Quantity:             | 300 μL                                                                                        |
| Gene:                 | Factor XI (F11)                                                                               |
| Species:              | Human                                                                                         |
| Insert:               | gRNA + Cas9                                                                                   |
| Vector:               | Lentiviral Vector                                                                             |
| Application:          | Protein Expression (PExp), Genome Editing with Engineered Nucleases (GEEN)                    |
| Product Details       |                                                                                               |
| Purpose:              | Individual gRNA against F11 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'TACGTCCAAGGTCGGGCAGGAAGA-3'                                              |
| Components:           | Lentiviral particles with an individual gRNA (300 μL) for a specific sequence of F11          |

## **Target Details**

| Gene:             | Factor XI (F11)    |
|-------------------|--------------------|
| Alternative Name: | F11 (F11 Products) |
| NCBI Accession:   | NM_000128          |

| 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 ( |  |

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1991)