



Pre-made Lentiviral Particles for Target Overexpression (for human, mouse or rat genes / ORFs)

Amount: 200ul/vial (1×10^7 IFU/ml); or 200ul/vial (1×10^8 IFU/ml) for concentrated lentivirus.

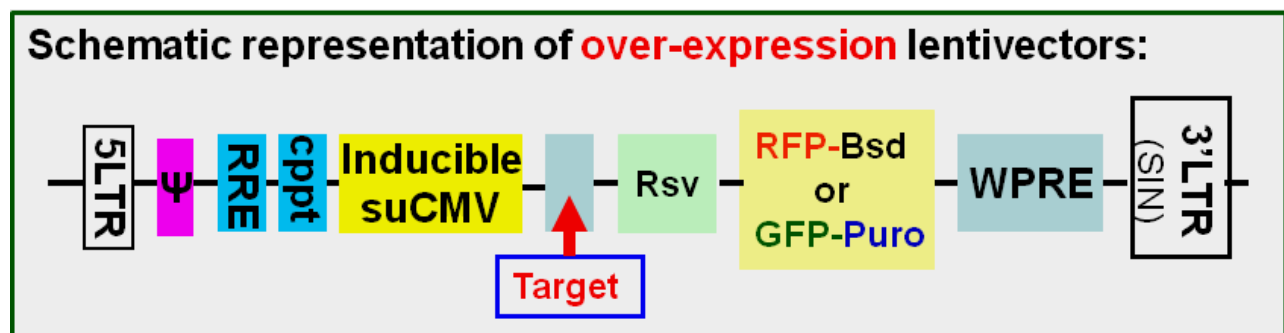
Storage: -70°C , avoid repeat freeze/thaw cycles. Stable for 6 months at -70°C .

Product Description:

GenTarget's Lentivector system is Human Immunodeficiency Virus-1 (HIV) based lentivector plasmids for gene expression and knockdown. The lentivectors are used to generate lentiviral particles (lentivirus) that can be transduced into most mammalian cell types, including stem cells, primary cells, and non-dividing cells both *in vivo* and *in vitro*. Lentiviral Particles stably integrate into the transduced cells' genome for long term expression, making lentivirus a great gene transfer agent.

Pre-made lentiviral particles for target over-expression are generated from GenTarget's [optional inducible lentiviral system](#) which means the lentivirus can be used for constitutive over-expression without need for any induction, or optionally, used for inducible expression when its repressor ([tetR](#)) is present (i.e., you purchase both tetR lentivirus and your desired target expression lentivirus, apply both lentivirus into your cells, then double antibiotic select the double transduced cells. The selected cells will express your target only after you add tetracycline inducer).

It adapted the most advanced biosafety features, including the self-inactivation feature in its 3' LTR, and only generates the replication-incompetent lentivirus.

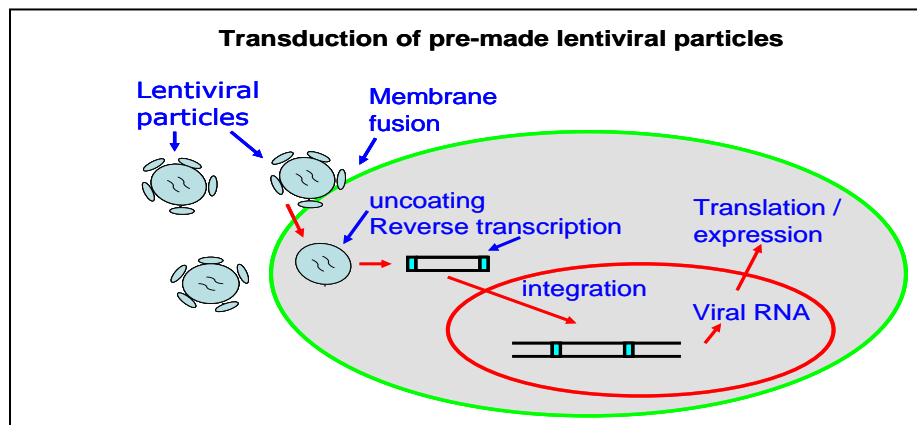


The lentivirus express a fully sequence-verified human, mouse, or rat gene under an optional inducible CMV promoter. (see lentivector's scheme above). Depend upon the product, a **RFP-Blasticidin** or **GFP-Puromycin** (Fluorescent-antibiotic



fusion) dual marker is included, which allows sorting or selection of the transduced cells by Fluorescent signal, and via antibiotic killing selection. The fluorescent signal provides a convenient, real-time means to monitor the particles' performance.

The [Negative Control Lentivirus](#) can be used to establish the controls for lentivirus treatment in a given cell line. The control lentivirus, CAT# [CMV-Null-RB](#) or [CMV-Null-GP](#), has the identical lentivector backbone as that of target expression, does not express any target but has the same dual selection. Please see also "[FAQs about premade lentiviral particles](#)".



Key features:

- 1) High target expression levels driven by extremely strong suCMV promoter;
- 2) Easy transduction monitoring by the fluorescent signal;
- 3) Option for inducible expression if desired;
- 4) Dual selection markers;
- 5) Ready to use and easy to use: simply add it into your cell culture, No need any other reagents;

Transduction Protocols:

1. Transduction Protocol for Adhesive cells:

Note: Pre-made lentivirus is provided ready to use, so it can be simply added into your cell culture; the amount of virus to add depends on cell type. For quick transduction, add 50 μ l of virus into each well of 24-well-plate where cell density is 50% to 75%. After 72 hours (no need to change medium), visualize positive transduction rate by fluorescence microscopy.



For stable cell line generation, pass cells into medium containing antibiotic or perform fluorescence cell sorting followed by antibiotic selection.

Day 0:

Seed cells in complete medium at the appropriate density and incubate overnight.

Note: at the time of transduction, cells should be 50%-75% confluent. For example, seed HeLa cells at 0.5×10^5 /ml x 0.5ml in a well of a 24-well plate.

Day 1:

- Thaw the pre-made lentiviral stock at room temperature and add the appropriate amount of virus stock to obtain the desired MOI.
- Return cells to 37°C, CO₂ incubator. Do nothing.

Note: Try to avoid freezing and thawing. If you do not use all of the virus at one time, you may re-freeze the virus at -80 °C for future use; virus titer will decrease by ~10% for each freeze/thaw cycle.

Day 3:

At 48hr~72hr (Depend upon cell type) after transduction, check the transduction rate by fluorescence microscopy or calculate the exact transduction rate by flow cytometry (FACS or Guava).

Day 3 + (optional):

Sort transduced cells by FACS, or select by antibiotic killing. A pilot experiment should be done to determine the antibiotic's kill curve for your specific cell line (refer to the pertinent literature on generation of stable cell lines).

2. Transduction Protocol for Suspension Cells:

Grow cells in complete suspension culture medium; use a shaking flask in a CO₂ incubator if required.

Measure cell density (not grow over 3 million/ml), measured viability should be > 90%. Dilute cells into 1×10^6 cell/ml in complete medium.

Day 1:

- Thaw lentiviral particles at room temperature.
- Add premade lentiviral particles into the diluted cells at a ratio of: 50 to 100 µl virus per 0.5 ml of cells (Note: depending on cell type, you may need to use more or less virus).



- Grow cells in a shaking flask in a CO2 incubator.

Day 2:

At 24 hours after transduction, add an equal amount of fresh medium containing. Continue growing cells in CO2 incubator.

Day 3+:

At 48 hour to 72 hours (Depend upon cell type) after transduction, check fluorescence with a fluorescence microscope or calculate the transduction efficiency using a cell sorter such as FACS or Guava. Pass cells into 0.5 million/ml density in completed medium containing the corresponding antibiotic (**Note:** amount of antibiotic depends on cell type. A killing curve must pre-established). Sort for fluorescence positive cells and maintain antibiotic selection to generate a stable cell line.

Note: Filter wavelength settings:

GFP filter: ~Ex450-490 ~Em525;

RFP filter: ~Ex558 ~Em583;

Safety Precaution:

Gentarget lentiviral particles adapts must advanced lentiviral safety features (using the third generation vectors with self-inactivation SIN-3UTR), and the premade lentivirus is replication incompetent. However, please use extra caution when using lentiviral particles. Use the lentiviral particles in Bio-safety II cabinet. Wear glove all the time when handling Lentiviral particles! Please refer CDC and NIH's guidelines for more details regarding to safety issues.

References:

1. J Virol. 2000 November; 74(22): 10778–10784.
2. Hum Gene Ther (2003) 14: 1089-105.
3. Mol Ther (2002) 6: 162-8.
4. NIH Guidelines for [Biosafety Considerations for Research with Lentiviral Vectors](#). (Link).

Warranty:

This product is for research use only. It is warranted to meet its quality as described when used in accordance with its instructions. GenTarget disclaims any implied warranty of this product for particular application. In no event shall GenTarget be liable for any incidental or consequential damages in connection with the products. GenTarget's sole remedy for breach of this warranty should be, at GenTarget's option, to replace the products.



Attachment: GenTarget's pre-made lentivirus product categories.

| Product Category | Product Description (please click into each category's page) |
|---|--|
| Pathway Reporter | Repoter Lentivirus for all kinds of pathway screening assays |
| Cell Immortalization | Lentivirus for cell immortalization: Large T-antigen, hTERT, EBNA1/EBNA2, HpV16-E6/E7, Adenovial E1A, Kras_G12V, HOXA9, et al. |
| ImmunoOncology Research | Lentivirus products for immuno therapy research: CAR and TCR; Assay Cell Lines for T-cell targeted killing assay and other cell-based assays; over-expression lentivirus products for the immune response targets; Cell surface antigens (CDs); immune checkpoint / Receptors; CRISPR gene Repair and knock-IN lentivirus; CRISPR knockout lentivirus; |
| CAR-T, TCR Lentivirus | CARs Lentivirus: Anti-CD19 /CD20 /CD22 /BCMA /hHER2 /HLA-A2 /TGFβ; TCRs : MART-1/ NY-ESO1/ CD1d-α-GalCer/ TRαV3-F2A-TRβV5-6; |
| CRISPR Gene Editing | Preamde lentivirus express humanized wild-type Cas9 endonuclease, the dCas9 , gRNAs, CRISPR gene editing research |
| Epigenomic: CRISPRi and CRISPRa | " dCas9-Protein " fusion Lentivirus for epigenomic modification, resulted in CRISPR interference (CRISPRi) or activation (CRISPRa). |
| Cell-Specific Reporter | a set of reporter lentiviruses to express a luminescence or fluorescent reporter (firefly Luciferase, Renilla luciferase, RFP or GFP fluorescent marker) under a tissue specific promoter |
| Infectious Antigens | Lentivirus that express all kinds of infectious antigens with C-term 6His-tag. |
| Virus Like Particles (VLP) | Lentiviral Like Particles, pseudo-typed with a different envelope proteins. |
| Non-integrating LV | Integration Defective Lentivirus, express different targets for transient expression without the unwanted insertional mutagenesis. |
| shRNA Knockdown | Knockdown verified and customized shRNA lentivirus for target knockdown, |



| Product Category | Product Description (please click into each category's page) |
|--|---|
| microRNA lentivirus | Premade lentivirus expression human or mouse precursor miRNA . And anti-miRNA lentivector and virus for human and mouse miRNA. |
| Anti-miNA lentivirus | Pre-made lentivirus expression a specific anti-miRNA cassette. |
| Human and mouse ORFs | Premade lentivirus expressin a human, mouse or rat gene with RFP-Blastididin fusion dual markers. |
| Luciferase expression | Premade lentivirus for all kinds of luciferase protein expression: firefly and Renilla, Red-Luc and more , with different antibiotic selection markers. |
| Fluorescent Markers | Lentivirus express all commonly used fluorescent proteins: GFP, RFP, CFP, BFP YFP, niRFP, unstable GFP and others. |
| Luminescent Imaging | Lentivirus express Nano-Latern as Bio-probes for in vivo imaging of sub-cellular structural organization and dynamic processes in living cells and organisms |
| Sub-cellular Imaging | Lentivirus contain a well-defined organelle targeting signal fused to a fluorescent protein, great tools for live-cell imaging and for dynamic investigation of sub-cellular signal pathways. |
| Cytoskeleton Imaging | A fluorescent marker (GFP, RFP or CFP) fusion with a cellular structure protein, provides a convenient tool for visualization of cytoskeletal structure |
| Unstable GFP | Lentivirus express the the destabilized GFP (uGFP) which provides fast turnover responses in signal pathway assay and in knockdown / knockout detection |
| near-infrared RFP | The near-infrared Red fluorescent (niRFP) expression Lentiviurs provides the whole-body images with better contrast and brighter images |
| Fluorescent-ORF fusion | Pre-made lentivirus expression a " GFP/RFP/CFP-ORF " fusion target. |
| CRE recombinase | Premade lentivirus for expressing nuclear permeant CRE recombinase with different flurescent and antibiotic markers. |
| CRE, Flp ColorSwitch | Lentivirus expressing "LoxP-GFP-Stop-LoxP-RFP" or "FRT-GFP-Stop-FRT-RFP" cassette, used to monitor the CRE or Flp recombination event in vivo. |



| Product Category | Product Description (please click into each category's page) |
|---|--|
| SEAP Reporter | lentivirus expressing SEAP under different promoters (TetCMV, EF1a, CAG, Ubc, mPGK, Actin-beta or a signal pathway responsive promoter), |
| TetR Repressor | Premade lentivirus expressing TetR (tetracycline regulator) protein, the repressor protein for the inducible expression system. |
| rtTA Expression | rtTA binds to the tetracycline operator element (TetO) in the presence of doxycycline (Dox). Used for Tet-On /OFF inducible system. |
| iPS factors | Premade lentivirus for human and mouse iPS (Myc, NANOG, OCT4, SOX2, FGF4) factors with different fluorescent and antibiotic markers |
| LacZ expression | Express different full length β-galactosidase (lacZ) with different selection markers |
| Negative control lentiviruses | Premade negative control lentivirus with different markers : serves as the negative control of lentivirus treatment, for validation of the specificity of any lentivirus target expression effects. |
| Other Enzyme expression | Ready-to-use lentivirus, expressing a specific enzymes with different selection markers. |
| Ultra titer lentivirus | Ultra-titer lentivirus used for the hard-to-transduced cells and for in vivo manipulation of sperm cells, or stem cells. |