

Premade Lentiviral Particles for Human iPS Stem Factors

For generating induced pluripotent stem (iPS) cells or other applications.

FOR RESEARCH USE ONLY, not for diagnostic or therapeutic use.

Cat#	Product Name	amounts
<u>LVP003</u>	h OCT4 (RFP-Bsd) inducible particles	
<u>LVP004</u>	h SOX2 (RFP-Bsd) inducible particles	
LVP005	h NANOG (RFP-Bsd) inducible particles	
LVP006	h LIN28 (RFP-Bsd) inducible particles	
LVP007	h Myc (RFP-Bsd) inducible particles	
<u>LVP008</u>	h Klf4 (RFP-Bsd) inducible particles	
LVP311	h OCT4 (Neo) inducible particles	
LVP312	h SOX2 (Neo) inducible particles	200ul x
LVP313	h NANOG (Neo) inducible particles	(1 x10 ⁸ IFU/ml),
LVP314	h LIN28 (Neo) inducible particles	in DDC colution
LVP315	h cMyc (Neo) inducible particles	in PBS solution, premixed with 10x
LVP316	h KLF4 (Neo) inducible particles	Polybrene, 60ug/ml
LVP317	h OCT4 (EF1α) (puro) particles	
LVP318	h SOX2 (EF1α) (<mark>puro</mark>) particles	
LVP319	h NANOG (EF1α) (<mark>puro</mark>) particles	
LVP320	h LIN28 (EF1α) (<mark>puro</mark>) particles	
LVP321	h Myc (EF1α (<mark>puro</mark>) particles	
LVP322	h Klf4 (EF1α) (<mark>puro</mark>) particles	
LVP588	h OCT4 (EF1α) (<mark>RP</mark>) particles	
<u>LVP589</u>	h SOX2 (EF1α) (<mark>RP</mark>) particles	
<u>LVP590</u>	h NANOG (EF1α) (RP) particles	
<u>LVP591</u>	h LIN28 (EF1α) (RP) particles	
LVP592	h Myc (EF1α (RP) particles	
<u>LVP593</u>	h Klf4 (EF1α) (RP) particles	

Storage: -80 °C, avoid repeat freeze/thaw cycles. Products stable for 12 months.



1. Product Description:

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GenTarget's lentivector system is Human Immunodeficiency Virus-1 (HIV) based plasmids for gene expression and knockdown. The lentivectors are used to generate lentiviral particles (lentivirus) that can be transduced into almost all kinds of mammalian cells, 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 it a great gene transfer agent.

Conversion of fully differentiated mouse or human somatic cells into embryonic-like cells (so called induced Pluripotent Stem Cells or iPSCs) has attracted enormous attention. Multiple reports have demonstrated that iPS cells can be generated by using a set of transcription factors or stem cell factors delivered via expression virus or by expressed proteins. The main stem cell factors are: OCT3/4, SOX2, NANOG, LIN28, c-Myc, and KLF4, although the combination of reprogramming factors may be slightly different. iPSCs hold the promise of curing many human diseases and accelerating stem cell research.

GenTarget provides **two sets of premade lentiviral particles** for human or mouse iPS genes. Each stem factor was natively expressed (without any tags) under either an <u>optional inducible suCMV promoter</u> (set#1) or enhanced **EF1a** promoter (set#2).

Utilizing our <u>Inducible Lentiviral Vector</u> system (see vector scheme below), GenTarget has generated high-titer inducible lentiviral particles for all six **human and mouse** stem cell factors. Each factor is fully sequenceverified and matched to the CDs in the NCBI database (see table below). High titer lentiviral particles/ supernatant were produced in 293T packaging cells (Cat# <u>TLV-C</u>) with a packaging mix (Cat# <u>HT-pack</u>). They are pseudotyped with VSV-G glycoprotein. They are packaged in DMEM medium (containing 10% FBS and 10x polybrene as ready-to-use status), and supplied as 200 µl/per vial at ~ 1x 10⁷ IFU/ml.

All six stem factor were sequencing verified. Their sequences fully match to the CD region according to the NCBI's database (see table below).

Target	NCBI ID	Matched ORF position
h Myc	<u>NM_002467</u>	526-1890

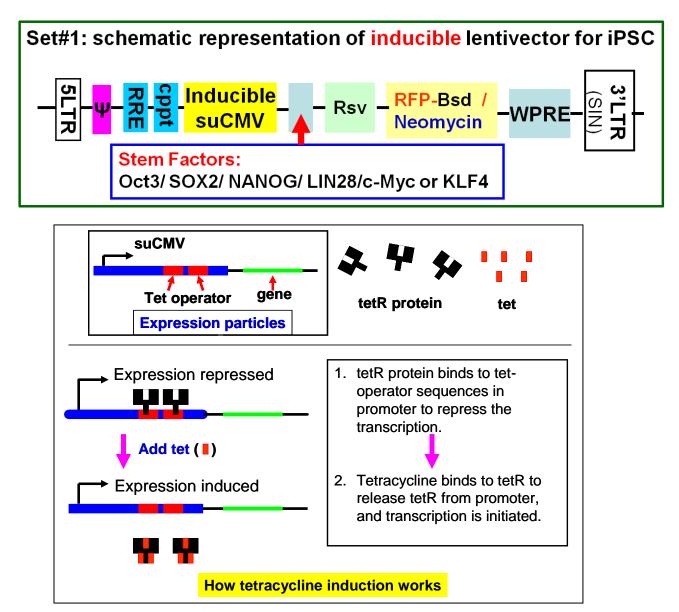




h Klf4	<u>NM_004235</u>	595-2034
h Oct3/4	<u>NM_002701</u>	55-1137
h SOX2	<u>NM_003106</u>	428-1381
h LIN28	<u>NM_024674</u>	115-744
h NANOG	<u>NM_024865</u>	217-1134

1) Set #1

Includes a tetracycline inducible **suCMV promoter** to drive iPS gene expression and contains a **RFP-Bsd** fusion dual or **Neomycin** selection marker under an **RSV promoter** (see vector map scheme below).



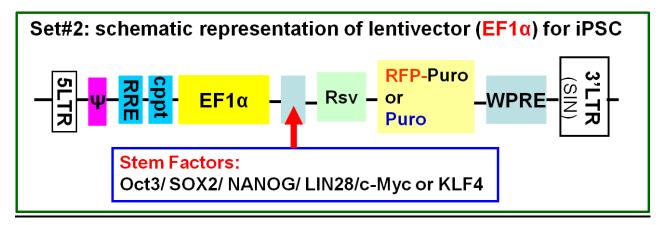




The particles in Set 1 can be used for regular constitutive high expression, or optionally for tetracycline-induced expression when the tetracycline regulator protein (tetR) is present in advance. For inducible expression, TetR must be expressed in advance to stop transcription; expression is then activated by adding tetracycline. This inducible expression is tetracycline dose dependent; in general, tetracycline is used at a final concentration of $1 \mu g/ml$. Please see the schematic above for the mechanism of inducible expression, and see our website for more details about our Inducible lentiviral system. GenTarget provides premade lentivirus expressing TetR with a variety of antibiotic markers. For general information about lentiviral particles, please refer to FAO about premade lentiviral particles.

2) Set #2

Uses the enhanced constitutive EF1a promoter to drive iPS gene expression (see vector map scheme below) with the option of using either a RFP**puromycin** fusion dual marker or the **puromycin** marker alone.



2. 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.

3. Attachment: iPS cell generation procedure for the Dox inducible system

(For reference only)



Day 0: Seed the parent cells:

- Seed human fibroblast cells at 1×10^5 cells/well in a 6-well plate, culture in 5ml of growth medium
- Incubate overnight at 37°C with 5% CO2

Day 1: Viral Transduction:

- Remove medium, add 2.5 ml of pre-warmed fibroblast growth medium, and then add 500µl of iPS lentivirus. Gentle mix for even distribution.
- Incubate overnight at 37°C with 5% CO2. [Note: set up inducible GFP positive control wells by adding 200ul/per well of GFP control particles]

Day 2: Change Medium

- At about 24 hours post-transduction, change to 5 ml growth medium.
- Incubated overnight at 37°C with 5% CO2.

Day 3: Re-plate the transduced cells to feeder cells

- At three days post-transduction, trypsinize cells and centrifuge at 200 x g for 5 minutes
- Resuspend in Fibroblast Cell Growth Medium
- Re-plate in a 150mm MEF Feeder Dish
- Incubate overnight at 37 °C with 5% CO2

Day 4: Induce Reprogramming using Dox

- At 24 hours after re-seeding, replace Fibroblast Cell Growth Medium with 2.0 ml Dox-Induction Medium containing 2µg/ml Dox. [Note: set up a negative control well without Dox.]
- Incubated Cells overnight at 37°C with 5% CO2.

Day 5+: Change Induction Medium

- Change Dox-Induction Medium every 48 hours
- Continue to pass the cells until they show typical human ES cell morphology

Day 14++: Select iPS cell colonies

- Pick the iPS cell colonies that conform to proper cell morphology using a sterile glass picking tool.
- Trypsinize each individually isolated iPS cell colony and pass into each well of a 24-well feeder plate.

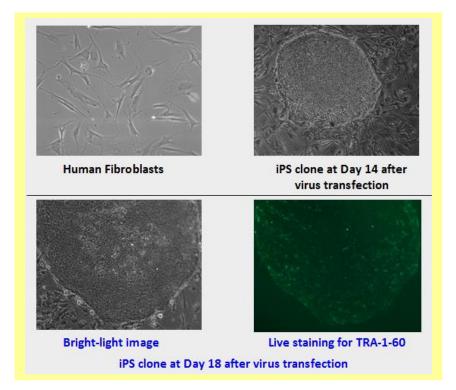
Passage, and Expansion of iPS cell colonies

- Incubate a 24-well plate at 37°C and 5% CO2,
- Replace culture medium with fresh medium without Dox every 48 hours.
- Passage into an appropriate size plate for iPS cell expansion (the process takes about 6-10 days).
- Monitor iPS cell colony growth and morphology, and validate the iPS colonies. Save iPS cells in cryogenic vials.



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iPS cell sample images:



4. References:

- 1) <u>NIH stem cell training program (Link)</u>.
- Masaki Ieda, Ji-Dong Fu, et al. (2010). Direct Reprogramming of Fibroblasts into Functional Cardiomyocytes by Defined Factors. Cell 142, 375-386.
- 3) Takahashi, K. and Yamanaka, S. (2006). Induction of pluripotent stem cells from mouse embryonic and adult fibroblast cultures by defined factors. Cell 126, 663-676.
- Yu, J., Vodyanik, M.A., Smuga-Otto, K., Antosiewicz-Bourget, J., Frane, J.L., Tian, S., Nie, J., Jonsdottir, G.A., Ruotti, V., Stewart, R., Slukvin, I.I., and Thomson, J.A. (2007). Induced pluripotent stem cell lines derived from human somatic cells. Science 318, 1917-1920.
- 5) Park, I.H., et al., Reprogramming of human somatic cells to pluripotency with defined factors. Nature, 2008. 451(7175): p. 141-6.
- 6) Shao, L., et al., Generation of iPS cells using defined factors linked via the self-cleaving 2A sequences in a single open reading frame. Cell Res., 2009. 19(3): p. 296-306.
- 7) NIH Guidelines for Biosafety Considerations for Research with Lentiviral Vectors. (Link).
- 8) <u>CDC guidelines for Lab Biosafety levels (Link).</u>



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

Product	Product Description
Category	(please click into each category's page)
<u>Pathway</u>	Repoter Lentivirus for all kinds of pathway screening
<u>Reporter</u>	assays
<u>Cell</u>	Lentivirus for cell immortalization: Large T-antigen,
Immortalization	hTERT, EBNA1/EBNA2, HpV16-E6/E7, Adenovial E1A, Kras_G12V, HOXA9, et al.
	Lentivirus products for immuno therapy research: CAR
	and TCR; Assay Cell Lines for T-cell targeted killing
<u>ImmunoOncology</u>	assay and other cell-based assays; over-expression
<u>Research</u>	lentivirus products for the immune response targets;
	Cell surface antigens (CDs); immune checkpoint /
	Receptors; CRISPR gene Repair and knock-IN lentivirus;
	CRISPR knockout lentivirus;
<u>CAR-T, TCR</u>	CARs Lentivirus: Anti-CD19 /CD20 /CD22 /BCMA
<u>Lentivirus</u>	/hHER2 /HLA-A2 /TGFβ; TCRs : MART-1/ NY-ESO1/
	CD1d-a-GalCer/ TRaV3-F2A-TRβV5-6;
CRISPR Gene	Preamde lentivirus express humanzied wild-type Cas9
<u>Editing</u>	endonuclease, the dCas9 , gRNAs, CRISPR gene editing
	research
Epigenomic:	"dCas9-Protein" fusion Lentivirus for epigenomic
CRISPRi and	modification, resulted in CRISPR interference (CRISPRi)
<u>CRISPRa</u>	or activation (CRISPRa).
	a set of reporter lentiviruses to express a luminescence
Cell-Specific	or fluorescent reporter (firefly Luciferase, Renilla
<u>Reporter</u>	luciferase, RFP or GFP fluorescent marker) under a
	tissue specific promoter
Infectious	Llentivirus that express all kinds of infectious antigens
Antigens	with C-term 6His-tag.
<u>Virus Like</u>	Lentiviral Like Particles, pseudo-typed with a different
Particles (VLP)	envelope proteins.
Non-integrating	Integration Defective Lentivirus, express different
LV	targets for transient expression without the unwanted
	insertional mutagenesis.
<u>shRNA</u>	Knockdown verifeid and customized shRNA lentivirus for
<u>Knockdown</u>	target knockdown,
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Product	Product Description
Category	(please click into each category's page)
microRNA	Premade lentivirus expression human or mouse
lentivirus	precursor miRNA. And anti-miRNA lentivector and
	virus for human and mouse miRNA.
<u>Anti-miNA</u>	Pre-made lentivirus expression a specific anti-miRNA
<u>lentivirus</u>	cassette.
Human and	Premade lentivirus expressin a human, mouse or rat
mouse ORFs	gene with RFP-Blastididin fusion dual markers.
Luciferase	Premade lentivirus for all kinds of luciferase protein
expression	expression: firefly and Renilla, Red-Luc and more,
	with different antibiotic selection markers.
Fluorescent	Lentivirus express all commonly used fluorescent
<u>Markers</u>	proteins: GFP, RFP, CFP, BFP YFP, niRFP, unstable GFP
Lunche e e such	and others.
Luminescent Imaging	Lentivirus express Nano-Latern as Bio-probes for in vivo
Imaging	imaging of sub-cellular structural organization and
Sub-cellular	dynamic processes in living cells and organisms Lentivirus contain a well-defined organelle targeting
Imaging	signal fusioned to a fluorescent protein, great tools for
Indging	live-cell imaging and for dynamic investigation of sub-
	cellular signal pathways.
Cytoskeleton	A fluorescent marker (GFP, RFP or CFP) fusion with a
Imaging	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	Pre-made lentivirus expression a "GFP/RFP/CFP-ORF"
fusion	fusion target.
	Premade lentivirus for expressing nuclear permeant
CRE recombinase	CRE recombinase with different flurescent and antibiotic markers.
CDE Elp	
<u>CRE, Flp</u> <u>ColorSwtich</u>	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.



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Product	Dreduct Description	
	Product Description	
Category	(please click into each category's page)	
	lentivirus expressing SEAP under different promoters	
SEAP Reporter	(TetCMV, EF1a, CAG, Ubc, mPGK, Actin-beta or a signal	
	pathway responsive promoter),	
	Premade lentivirus expressin TetR (tetracycline	
TetR Repressor	regulator) protein, the repressor protein for the	
	inducible expression system.	
	rtTA binds to the tetracycline operator element (TetO) in	
rtTA Expression	the presence of doxycycline (Dox). Used for Tet-On /OFF	
	inducible system.	
	Premde lentivirus for human and mouse iPS (Myc,	
iPS factors	NANOG, OCT4, SOX2, FLF4) factors with different	
	fluorescent and antibitoic markers	
LacZ expression	Express different full length β- galactosidase	
	(lacZ) with different selection markers	
	Premade negative control lentivirus with different	
Negative control	markers: serves as the negative control of lentivurs	
<u>lentiviruses</u>	treatment, for validation of the specificity of any	
	lentivirus target expression effects.	
Other Enzyme	Ready-to-use lentivirus, expressing a specific enzymes	
expression	with different selection markers.	
<u>Ultra titer</u>	Ultra-titer lentivirus used for the hard-to-transduced	
<u>lentivirus</u>	cells and for in vivo manipulation of sperm cells, or stem	
	cells.	