

7930 Arjons Drive, Suite B San Diego, CA 92126 Phone: (858) 6788683 Fax: (800) 3804198

Email: orders@gentarget.com

Pre-made Reporter Lentivirus for EGR1-Promoter Signal Pathway

Cat#	Product	Amounts
	Name	
<u>LVP1119-P</u>	EGR1- GFP (Puro)	
Or	Lentivirus	
LVP1119-P-PBS LVP1120-P		_
or	EGR1-RFP (Puro) Lentivirus	
LVP1120-P-PBS		
LVP1121-P	FCD4 Luc (Dura)	
or	EGR1- Luc (Puro) Lentivirus	
LVP1121-P-PBS	Lentivirus	
<u>LVP1122-P</u>	EGR1-Rluc (Puro)	
Or 1)/D4400 D DD0	Lentivirus	
LVP1122-P-PBS		_
<u>LVP1119-B</u> or	EGR1- GFP (Bsd)	200ul, ~1 x 10 ⁷ IFU/mL in
LVP1119-B-PBS	Lentivirus	
LVP1120-B	FOR4 PER (5-1)	DMEM containing 10% FBS
or	EGR1- RFP (Bsd) Lentivirus	
LVP1120-B-PBS		Or
<u>LVP1121-B</u>	EGR1- Luc (Bsd) Lentivirus	200ul, ~1 x 10 ⁸ IFU/mL in
or		
LVP1121-B-PBS	20111111100	PBS solution
<u>LVP1122-B</u>	EGR1- Rluc (Bsd) Lentivirus	
or LVP1122-B-PBS		
LVP1119-N		
or	EGR1- GFP (Neo)	
LVP1119-N-PBS	Lentivirus	_
LVP1120-N	EGR1- RFP (Neo)	
or	Lentivirus	
LVP1120-N-PBS		
LVP1121-N or	EGR1- Luc (Neo)	
LVP1121-N-PBS	Lentivirus	
LVP1122-N	F004 B: 4	
or	EGR1- Rluc (Neo)	
LVP1122-N-PBS	Lentivirus	
<u>LVP1119-R</u>	EGR1- GFP (RFP)	
or	Lentivirus	
LVP1119-R-PBS		
<u>LVP1121-R</u>	EGR1- Luc (RFP)	
or LVP1121-R-PBS	Lentivirus	
LVP1122-R	 	
or	EGR1- Rluc (RFP) Lentivirus	
LVP1122-R-PBS		



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LVP1120-G or LVP1120-G-PBS	EGR1- RFP (GFP) Lentivirus	
LVP1121-G or LVP1121-G-PBS	EGR1- Luc (GFP) Lentivirus	
LVP1122-G or LVP1122-G-PBS	EGR1- Rluc (GFP) Lentivirus	

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

Product Description:

Lentiviral system is a gene delivery tool using lentivectors for gene expression or knockdown. 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.

EGR1 pathways:

EGR-1 (Early growth response protein 1) is a mammalian zinc finger transcription factor. The full human EGR1 promoter sequence contains regulatory elements including two Sp1 sites, an AP1 site, two cAMP response elements and an Egr-1 binding site, five serum response elements (SREs). This promoter is activated by ionizing radiation, demonstrating radiation-induced upregulation. It is also up-regulated in response to growth factors or stress that activated by the MAP kinase kinase MEK1. Therefore, dependent on specific cell type, the activation of EGR1 promoter involves multiple signal transduction pathway.

The feature that the EGR1 promoter is induced by ionizing radiation, various mitogenic stimuli, malignant transformation signals, makes it a useful *in vivo* tool for cancer research.

Product Principle:

GenTarget developed a set of EGR1 reporting lentivirus to monitor EGR1-promoter signaling pathway. Those reporting lentivirus has a **luminescent report** or a **fluorescent report** under the <u>full sequence of human EGR1 promoter</u>. Those EGR1 reporter are used to measure all pathway signals that activate EGR1 transcription factor. Those reporter monitor both increases and



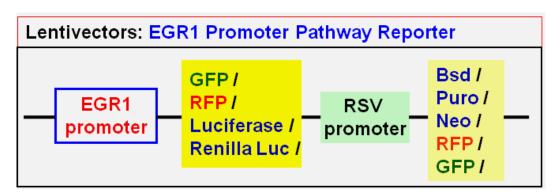
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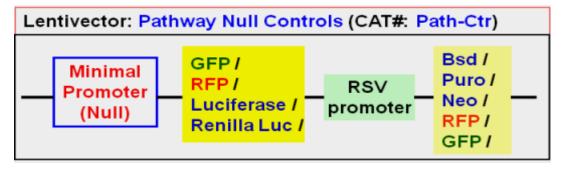
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decreases in the activity of EGR1 pathway, and determine the effect of various treatments, such as gene knockdown, over-expression, and chemical compounds on EGR1 activity.

The reporter's signal can be easily and rapidly readout via plate assays (luciferase assay or by fluorescent microscope or FACS sorter).

Those reporting lentivirus also contains a constitutively expressed fluorescent selection marker or an antibiotic selection marker under the RSV promoter, which makes it easier to select the stably infected signal reporting cells (to generate pathway specific sensor cell lines), or provides internal reference for virus transduction efficiency when a fluorescent marker is under the RSV promoter. A set of control lentivirus use the same lentivector backbones except the minimal promoter does not contain any signal pathway's TRE sequences. The control lentivirus are used to set the reference for specificity of pathway signal response upon treatment. See the scheme below for lentivector's core expression cassette.





The premade, ready-to-use reporter lentivirus provides a much easier tool to monitor the activity of EGR1-Promoter signaling pathways in virtually any mammalian cell type. It also allows to generate your own reporting cell line in your desired cell type for study or screen of pathway specific gene-



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knockdown, over-expression, or chemical / drug/protein treatment in the cell based assay.

Lentivirus are HIV-based, pseudotyped with VSVG envelope protein, produced in 293T cells. All particles were tested to be free bacterial and mycoplasma contamination. Virus titers were tested lot by lot.

Key Application for Pathway Signaling Lentivirus:

- Create signal pathway specific cell lines which can provide a Highthroughput, live cell based assays for signal transduction tests;
- 2. Identify or validate the signaling pathway specific drugs (drug discovery and validation);
- 3. Analyze the pathway-specific responses to proteins, peptides, or hormones;
- 4. Analyze the pathway-specific responses to gene activation, over-expression, knockdown, knockout, or mutagenesis;
- 5. Screen for pathway-specific stimulus or for the transcriptional activators that response to specific pathway's TRE elements;
- 6. makes it easy to measure the transcriptional and post-transcription regulation in response to signal pathway stimulus.

Product Formats:

The pre-made lentivirus provided in two formats:

- Packaged in 10% of FBS in DMEM containing 10% FBS and 60ug/ml of polybrene (10x);
- 2. Particles were concentrated and buffer exchanged in PBS without any human or animal origin components. The virus in PBS are used for any cell types that requires non-serum in the culture medium, or best for the hard-to-infect cell types.

The lentivirus are ready and easy to use, simply add 50ul into one well of your cell culture in 24-well plate, and select or sort the positive transduced cells at 2-3 days post virus transduction (for sensor cell line assay). Or simply go for Estrogen receptor signal induction without the selection (for transient assay). The readout can be easily monitored by luciferase assay or via the Fluorescent microscope or readers depending on product report type.

For more details about premade particles, please see <u>FAQ for pre-made</u> <u>lentiviral particles</u> (.pdf).



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Transduction Protocols:

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 (when applicable). For stable cell line generation, pass cells into medium containing antibiotic for selection, or perform fluorescence cell sorting.

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 $0.5 \times 10^5/\text{ml} \times 0.5\text{ml}$ 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.

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

Day 3:

At ~72hr after transduction, check the transduction rate by fluorescence microscopy or calculate the exact transduction rate by flow cytometry (FACS or Guava). You can now treat the cell for signal pathway assay.

Day 3 + (optional):

Sort transduced cells by FACS, and select for antibiotic resistance. 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). The selected stable cells will be used for signal pathway assay with desired treatment.

Next: Treat the cell with signal pathway inducer, and analyze the pathway reporter expression (Fluorescent readout or luciferase assay).

Signal pathway assay recommendations:



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1. **Treatment:** the reporter's inducible expression is dose and time dependent upon induction or treatment. You may need to optimize the best treatment amount and the time point.

2. Controls:

- Virus transduction controls: Gentarget's signal report lentivirus contains a non-inducible fluorescent marker as the internal normalization control (only applicable to the lentivirus containing a fluorescent marker under Rsv promoter). The embed internal control fluorescent signal also monitors the lentivirus transduction efficiency in assay cell types. When the internal control is not available, you can mix a regular luciferase or fluorescent marker lentivirus under a constitutive promoter (see GenTarget's <u>luciferase lentivirus</u> and <u>fluorescent lentivirus</u> products).
- No-response controls: if desired, you also can use the Pathway-control lentivirus that made in the same lentivector backbone but without any TRE in its minimal promoter. (Note: the minimal CMV promoter has no or little activity in most cell types). This control virus serves for the specificity of any treatment or as for the establishment of the basal signal profile.
- Positive controls: If applicable, apply the characterized pathway stimulus as the pathway positive induction controls, such as treated with known inducer, proteins, peptide or compounds.
- 3. **Make triplicates** for each condition for assay reproducibility.
- 4. **Assay cell number**: you may need to carry out a cell titration to determine the optimal cell number for the signal reporter assay.

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 Biosafety II cabinet. Ware glove all the time at handling Lentiviral particles! Please refer CDC and NIH's guidelines for more details regarding to safety issues.

References:

- DNA Seq. 2000;10(6):429-32.
- Eur J Biochem. 2002 Jan;269(1):337-46.



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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.

Note: Filter wavelength settings:

GFP filter: ~Ex450-490 ~Em525; RFP filter: ~Ex545 ~Em620;

Related Products: GenTarget's pre-made lentivirus product category.

Product	Product Description
Category	(please click category name to see product's pages)
<u>.Human,</u>	Premade lentivirus expressin a human, mouse or rat gene with
mouse or rat	RFP-Blastididin fusion dual markers.
ORFs.	
<u>Pathway</u>	Premade lentivirus reporter express a luminescence or fluorescent
Reporter	report (firefly Luciferase, Renilla luciferase, RFP or GFP
~ 11 ~ 1.0	fluorescent marker) under a pathway specific promoter.
Cell-Specific	Premade lentivirus reporter for targeting expression of a
Reporter	luminescence or fluorescent report (firefly Luciferase, Renilla
	luciferase, RFP or GFP fluorescent marker) under a cell type
	specific promoter.
Fluorescent	Preamde lentivirus express human codon optimized fluorescent
<u>markers</u>	protein, GFP / RFP/ CFP/ BFP / YFP.
<u>Luciferase</u>	Premade lentivirus for all kinds of luciferase protein expression:
expression.	firefly and Renilla with different antibiotic selection markers.
. <u>CRE</u>	Premade lentivirus for expressing nuclear permeant CRE
<u>recombinase</u>	recombinase with different flurescent and antibiotic markers.
<u>LoxP</u>	Premade lentivirus expressing "LoxP-GFP-Stop-LoxP-RFP"
<u>ColorSwitch</u>	cassette, used to monitor the CRE recombination event in vivo.
CRISPR /hu	Preamde lentivirus express humanzied wild-type Cas9
CAS9	endonuclease for genomic editing with CRISPR
. <u>TetR</u>	Premade lentivirus expressin TetR (tetracycline regulator) protein,
<u>inducible</u>	the repressor protein for the inducible expression system.
expression	
<u>repressor</u>	



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<u>iPS factors</u>	Premde lentivirus for human and mouse iPS (Myc, NANOG, OCT4, SOX2, FLF4) factors with different fluorescent and antibitoic markers
<u>T-antigen</u>	Express SV40 large T antigen with different selection markers
Expression	
<u>.Cell</u>	Premade lentivirus for cell organelle imaging. The fluorescent
<u>Organelle</u>	marker GFP/RFP/CFP was sub-cellular localized in different
<u>imaging</u>	cell organelle for living cell imaging.
LacZ	Express different full length β- galactosidase (lacZ) with different
expression	selection markers
Anti-miNA	Pre-made lentivirus expression a specific anti-miRNA cassette.
lentivirus	
Fluorescent-	Pre-made lentivirus expression a "GFP/RFP/CFP-ORF" fusion
ORF fusion.	target.
Pre-made	Premade shRNA lentivirus for knockdown a specific genes (P53 ,
shRNA	LacZ, Luciferase and more).
<u>lentivirus</u>	
_microRNA	Premade lentivirus expression human or mouse precursor
and anti-	miRNA. And anti-miRNA lentivector and virus for human and
microRNA	mouse miRNA.
lentivirus	
Negative	Premade negative control lentivirus with different markers:
control	serves as the negative control of lentivurs treatment, for
lentiviruses	validation of the specificity of any lentivirus target expression
.3.16.711 0000	effects.
Other	Ready-to-use lentivirus, expressing specific enzymes with
Enzyme	different selection markers.
expression	
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