

7930 Arjons Drive, Suite B San Diego, CA 92126, USA Phone: 1 (858) 265-6446 Fax: 1 (800) 380-4198 Email: orders@gentarget.com

Premade Anti-CD19 CAR-T Lentivirus

Note: All products are for Research Use Only and CANNOT be used in the treatment or diagnosis of disease.

Cat#	Product Name	Amounts
LVP1440	Anti-CD19-ScFv- <mark>4-1BB</mark> -CD3ζ (Puro)	
LVP1441	Anti-CD19-ScFv- <mark>4-1BB</mark> -CD3ζ (Bsd)	
LVP1442	Anti-CD19-ScFv- <mark>4-1BB</mark> -CD3ζ (GFP-Puro)	
LVP1443	Anti-CD19-ScFv- <mark>4-1BB</mark> -CD3ζ (RFP-Puro)	
LVP1444	Anti-CD19-ScFv- <mark>4-1BB</mark> -CD3ζ (no marker)	
LVP1445	Anti-CD19-ScFv- CD28-CD3ζ (Puro)	200 1 (4 408 1711 1)
LVP1446	Anti-CD19-ScFv- CD28-CD3ζ (Bsd)	200ul x (1x10 ⁸ IFU/ml) in PBS solution, pre-
LVP1447	Anti-CD19-ScFv- CD28-CD3ζ (GFP-Puro)	mixed with polybrene.
LVP1448	Anti-CD19-ScFv- CD28-CD3ζ (RFP-Puro)	
LVP1449	Anti-CD19-ScFv-CD28-CD3ζ (no marker)	
CAR-ctr1	CAR negative control: 4-1BB-CD3ζ	
CAR-ctr2	CAR negative control: Anti-CD19-ScFv-4-1BB	
CAR-ctr3	CAR negative control: CD28-CD3ζ	
CAR-ctr4	CAR negative control: Anti-CD19-ScFv-CD28	

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



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1. Product Description:

1) About Lentivirus,

GenTarget's lentivector system is Human Immunodeficiency Virus-1 (HIV) based plasmids for gene delivery. 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.

2) What is CAR-T,

CARs (Chimeric Antigen Receptors) is the method to modify T cells to produce a special structure on their surface. The structure consists of two parties, the antigen-binding and the T cell activation. The Antigen-Binding, like an antibody, targets a tumor surface antigen. The antigen-binding moieties usually the scFv's derived from antibodies (for example, targeting a tumor surface antigen), or Fab segment, or nature ligands binding their cognate receptor. The activation part consists of a chimeric molecule activation domain (1st generation CARs), or a fusion containing both activating and co-stimulatory properties (2^{nd} generation CARs). The 2nd-generation CARs encompasses the CD3- ζ chain and the cytoplasmic domain of a co-stimulatory receptor such as CD28, 4-1BB, CD80 CD40L. The so-called 3^{rd} - generation of CARs embed two co-stimulatory domains combined with an activation domain in their cytoplasmic domain.

CAR-modified T cells are tumor-targeted T cells, enhanced T cell expansion and persistence within tumor microenvironment, thus rapidly evolve a potential cancer immunotherapy. The encoded expression of cytokines in CAR-T cells can further alter the tumor microenvironment, showing great enhancement for this approach. For example, CD19-targeted, CAR-modified T cells expressing IL-12 showed greater efficacy than CAR-modified T cells alone.

3) About CD19,

CD19 is a marker of in most B cell leukemias and lymphomas but not in any normal tissue other than the B cell lineage, CD19 is used to diagnose cancers that arise from this type of cell, and used as the target for CD19-targeted therapies.



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4) About 4-1BB (CD137),

4-1BB (CD137) is a surface glycoprotein present on activated T Lymphocytes, CD4+ and CD8+ T cells. It binds to its ligand expressed on antigen-presenting cells (macrophages and activated B cells). It promotes T-cell antitumor activity. 4-1BB is one of the most widely studied co-stimulatory domain used in CAR design.

5) About CD28,

CD28 is transmembrane protein expressed on T cells. It provides costimulatory signals for inducing T cell activation and proliferation. CD28 costimulatory domains in CARs led to enhanced anti-malignancy efficacy. CD28 is widely used as the costimulatory domain in CARs.

6) About CD3ζ,

CD3 ζ (T-cell receptor zeta) is expressed by T cells and NK cells. It together with T-cell receptor and CD3 γ , δ , ϵ chain, forms the TCR-CD3 complex. CD3-zeta is the most commonly used activation component of CARs. It transmits an activation signal to the T cell after the antigen is bound. It can be coupled with additional co-stimulatory signaling for the complete activation.

2. Gentarget's CAR-T Lentivirus:

To generate the genetically modified T cells that kill certain types of cancer cells, Gentarget Inc constructs the tumor-specific CAR lentivectors, expressing the 2nd or 3rd generation of chimeric antigen receptor (CAR) or a T cell receptor (TCR).

CAR-T targeted on CD19: Those lentivirus express the CAR construct of "anti-CD19-ScFV (VL-VH)-CD8 hinge-4-1BB-CD3 ζ ", or "anti-CD19 ScFV (VL-VH)-CD8 hinge-CD28-CD3 ζ ", under the enhanced EF1a promoter that tested strong in T cells. A signal leader with 3x FLAG tag, is constructed in front of Anti-CD19 ScFV. The Anti-CD19 is the verified clone FMC63. The stimulatory domain of either human 4-1BB (CD137) or human CD28 is constructed at downstream of Anti-CD19, and followed by the activation signal of CD3 ζ .

For the validation of CAR-T specificity, two types of CAR-T controls are constructed by either remove Anti-CD19 ScFV recognition domain or remove CD3 ζ activation domain.

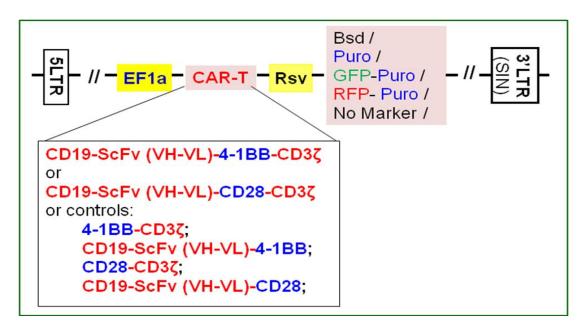
To avoid any unwanted immnuo-antigen, the lentivirus does not contain any antibiotic marker, or as desired, contains an antibiotic selection marker, or



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Fluorescent-Antibiotic fusion dual selection marker under Rsv Promoter. See the core structure of the lentivirus in the following scheme.



The VSV-G pseudotyped CAR-T Lentivirus are generated from 293T cells, and the concentrated lentivirus is provided in PBS solution at the titer of 1×108 IFU/ml, as 200ul aliquots.

For more details, please see *FAQs for pre-made lentiviral particles* (.pdf).

3. Key features of Gentarget's CAR-T Lentivirus:

- Idea as the validation assay for T cell CAR killing in CD19 positive cancer cells;
- High titer lentivirus for the high transduction rate in T cells;
- High promoter strength in T cells by enhanced EF1a promoter;
- Easy transduction verification by the GFP or RFP signal (when desired);
- Dual selection markers: transduced cells can be sorted via a fluorescent signal or selected for antibiotic resistance (when desired);
- Ready to use: simply add 50 µl into your cell culture in a 24-well plate, and leave the virus on for 48hr to 72 hours. (Note: depending upon your specific needs, you may transduce with different MOIs for different levels of expression.)



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4. Transduction Protocol in T Cells (lymphocyte):

Note: Pre-made lentivirus is provided ready to use, 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% or at the cell number as 1 million/ml for suspension cells. 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 or perform fluorescence cell sorting.

- 1) Seed 0.5 ml of T lymphocyte cells (CD4+ and CD8+) in complete medium, into one wall in 24 well/plate, at the cell concentration of 1 million cells/ml, at the time of lentivirus transduction (i.e. 0.5 million cells/well);
- 2) Thaw the Lentivirus at room temperature, and add 50 ul of the CAR-T lentivirus into each well in 24 w/p. You can scale up the transduction when needed for larger culture volume; No need to add polybrene as the provided lentivirus are pre-mixed with 10x polybrene.
- 3) Return cells to 37°C, CO₂ incubator;

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

- 4) At 48hr to 72hr post transduction, transfer the cells to flask in completed medium (when applicable, with puromycin selection) for the transduced cell enrichment and expansion; or analysis via FAC.
- 5) When cell grow to the desired cell number (about one more week culture), check the transduction rate by fluorescence microscopy or by flow cytometry, or simply go for the validation of the anti-CD19 CAR-T assays (For example, use our <u>CD19 expression cell lines</u> as targeted killing assay which can be measured via GFP signal decay via plate readout format or via FAC.

Note: Filter wavelength settings:

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

5. 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. Ware glove all the time at



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handling Lentiviral particles! Please refer CDC and NIH's guidelines for more details regarding to safety issues.

6. References:

- 1. David M; The journal of immunology,2015
- 2. Porter DL et al; N Engl J Med. Aug 25 2011;
- 3. Kochenderfer JN, et al; J Clin Oncol. Feb 20 2015.
- 4. NIH Guidelines for Bio-safety Considerations for Research with Lentiviral Vectors. (Link).
- 5. CDC guidelines for Lab Bio-safety levels (Link).

7. 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.

8. 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 humanzied wild-type Cas9 endonuclease, the dCas9 , gRNAs, CRISPR gene editing research	
Epigenomic: CRISPRi and	"dCas9-Protein" fusion Lentivirus for epigenomic modification, resulted in CRISPR interference (CRISPRi)	



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Category	(please click into each category's page)
CRISPRa	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
<u>Infectious</u> <u>Antigens</u>	Llentivirus 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 verifeid and customized shRNA lentivirus for target knockdown,
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.
<u>Luciferase</u> <u>expression</u>	Premade lentivirus for all kinds of luciferase protein expression: firefly and Renilla, Red-Luc and more, with different antibiotic selection markers.
<u>Fluorescent</u> <u>Markers</u>	Lentivirus express all commonly used fluorescent proteins: GFP, RFP, CFP, BFP YFP, niRFP, unstable GFP and others.
<u>Luminescent</u> <u>Imaging</u>	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 fusioned to a fluorescent protein, great tools for live-cell imaging and for dynamic investigation of subcellular signal pathways.



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Category	(please click into each category's page)
Cytoskeleton	A fluorescent marker (GFP, RFP or CFP) fusion with a
Imaging	cellular structure protein, provides a convenient tool for
	visualization of cytoskeletal structure
<u>Unstable GFP</u>	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"
<u>fusion</u>	fusion target.
	Premade lentivirus for expressing nuclear permeant
CRE recombinase	CRE recombinase with different flurescent and antibiotic
	markers.
CRE, Flp	Lentivirus expressing "LoxP-GFP-Stop-LoxP-RFP" or
<u>ColorSwtich</u>	"FRT-GFP-Stop-FRT-RFP" cassette, used to monitor the
	CRE or Flp recombination event in vivo.
05455	lentivirus expressing SEAP under different promoters
SEAP Reporter	(TetCMV, EF1a, CAG, Ubc, mPGK, Actin-beta or a signal
	pathway responsive promoter),
Total Donwooder	Premade lentivirus expressin TetR (tetracycline
TetR Repressor	regulator) protein, the repressor protein for the
	inducible expression system.
rtTA Expression	rtTA binds to the tetracycline operator element (TetO) in
ITTA EXPLESSION	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
II 3 Tactors	fluorescent and antibitoic markers
LacZ expression	Express different full length β- galactosidase
<u>Euce expression</u>	(lacZ) with different selection markers
	Premade negative control lentivirus with different
Negative control	markers: serves as the negative control of lentivurs
lentiviruses	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



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Product Category	Product Description (please click into each category's page)	
lentivirus	cells and for in vivo manipulation of sperm cells, or stem cells.	