

Resource Summary Report

Generated by [RRID](#) on Sep 3, 2026

LV-Cre pLKO.1

RRID:Addgene_25997

Type: Plasmid

Proper Citation

RRID:Addgene_25997

Plasmid Information

URL: <http://www.addgene.org/25997>

Proper Citation: RRID:Addgene_25997

Insert Name: Cre recombinase

Organism: bacteriophage

Bacterial Resistance: Ampicillin

Defining Citation: [PMID:20526348](#)

Vector Backbone Description: Backbone Size:6543; Vector Backbone:pLKO.1; Vector Types:Mammalian Expression, Lentiviral, Cre/Lox; Bacterial Resistance:Ampicillin

Comments: Second AgeI site is introduced preventing AgeI-EcoRI shRNA subcloning. TRC library shRNAs can be subcloned as an Accl fragment.

Plasmid Name: LV-Cre pLKO.1

Record Creation Time: 20220422T222116+0000

Record Last Update: 20260902T045130+0000

Ratings and Alerts

No rating or validation information has been found for LV-Cre pLKO.1.

No alerts have been found for LV-Cre pLKO.1.

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 18 mentions in open access literature.

Listed below are recent publications. The full list is available at [RRID](#) .

Sunami T, et al. (2026) IGF2BP3 is essential for the growth heterogeneity of colorectal adenoma cells by regulating MYC. *iScience*, 29(7), 116535.

Cho MG, et al. (2024) MRE11 liberates cGAS from nucleosome sequestration during tumorigenesis. *Nature*, 625(7995), 585.

Zhang X, et al. (2024) Chandipura viral glycoprotein (CNV-G) promotes Gectosome generation and enables delivery of intracellular therapeutics. *Molecular therapy : the journal of the American Society of Gene Therapy*, 32(7), 2264.

Lorenzo-Martín LF, et al. (2024) Spatiotemporally resolved colorectal oncogenesis in mini-colons ex vivo. *Nature*, 629(8011), 450.

Wang L, et al. (2023) A single-cell atlas of bovine skeletal muscle reveals mechanisms regulating intramuscular adipogenesis and fibrogenesis. *Journal of cachexia, sarcopenia and muscle*, 14(5), 2152.

Jana S, et al. (2023) Transcriptional-translational conflict is a barrier to cellular transformation and cancer progression. *Cancer cell*, 41(5), 853.

Kantzer CG, et al. (2022) ID1 and CEBPA coordinate epidermal progenitor cell differentiation. *Development (Cambridge, England)*, 149(22).

Karagyaour M, et al. (2022) A Novel Cre/lox71-Based System for Inducible Expression of Recombinant Proteins and Genome Editing. *Cells*, 11(14).

Kato S, et al. (2021) Precision modeling of gall bladder cancer patients in mice based on orthotopic implantation of organoid-derived tumor buds. *Oncogenesis*, 10(4), 33.

Maru Y, et al. (2021) Probing the tumorigenic potential of genetic interactions reconstituted in murine fallopian tube organoids. *The Journal of pathology*, 255(2), 177.

Maru Y, et al. (2021) Kras activation in endometrial organoids drives cellular transformation and epithelial-mesenchymal transition. *Oncogenesis*, 10(6), 46.

Isaev K, et al. (2021) Pan-cancer analysis of non-coding transcripts reveals the prognostic onco-lncRNA HOXA10-AS in gliomas. *Cell reports*, 37(3), 109873.

Cai EY, et al. (2020) Selective Translation of Cell Fate Regulators Mediates Tolerance to Broad Oncogenic Stress. *Cell stem cell*, 27(2), 270.

Aono S, et al. (2019) β -Catenin/TCF4 Complex-Mediated Induction of the NRF3 (NFE2L3) Gene in Cancer Cells. *International journal of molecular sciences*, 20(13).

Guo Y, et al. (2019) Chemical suppression of specific C-C chemokine signaling pathways enhances cardiac reprogramming. *The Journal of biological chemistry*, 294(23), 9134.

Gantner ML, et al. (2016) Complementary Roles of Estrogen-Related Receptors in Brown Adipocyte Thermogenic Function. *Endocrinology*, 157(12), 4770.

Lin J, et al. (2016) MicroRNA-181b inhibits thrombin-mediated endothelial activation and arterial thrombosis by targeting caspase recruitment domain family member 10. *FASEB journal : official publication of the Federation of American Societies for Experimental Biology*, 30(9), 3216.

Moriwaki K, et al. (2015) A RIPK3-caspase 8 complex mediates atypical pro-IL-1 β processing. *Journal of immunology (Baltimore, Md. : 1950)*, 194(4), 1938.