

Resource Summary Report

Generated by [RRID](#) on Aug 26, 2026

pLenti-DsRed_IRES_EGFP

RRID:Addgene_92194

Type: Plasmid

Proper Citation

RRID:Addgene_92194

Plasmid Information

URL: <http://www.addgene.org/92194>

Proper Citation: RRID:Addgene_92194

Insert Name: DsRed-IRES-EGFP

Organism: Synthetic

Bacterial Resistance: Ampicillin

Defining Citation: [PMID:27779468](#)

Vector Backbone Description: Backbone Size:8000; Vector Backbone:pLenti-GPS-GAW-IRES; Vector Types:Mammalian Expression, Lentiviral; Bacterial Resistance:Ampicillin

Plasmid Name: pLenti-DsRed_IRES_EGFP

Record Creation Time: 20220422T222623+0000

Record Last Update: 20260815T082228+0000

Ratings and Alerts

No rating or validation information has been found for pLenti-DsRed_IRES_EGFP.

No alerts have been found for pLenti-DsRed_IRES_EGFP.

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 7 mentions in open access literature.

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

Taglini F, et al. (2024) DNMT3B PWWP mutations cause hypermethylation of heterochromatin. EMBO reports, 25(3), 1130.

Slein MD, et al. (2024) Effector functions are required for broad and potent protection of neonatal mice with antibodies targeting HSV glycoprotein D. Cell reports. Medicine, 5(2), 101417.

Yan J, et al. (2024) Improving prime editing with an endogenous small RNA-binding protein. Nature, 628(8008), 639.

Shi P, et al. (2023) Collateral activity of the CRISPR/RfxCas13d system in human cells. Communications biology, 6(1), 334.

Slein MD, et al. (2023) Antibody effector functions are required for broad and potent protection of neonates from herpes simplex virus infection. bioRxiv : the preprint server for biology.

Huang YH, et al. (2022) Systematic Profiling of DNMT3A Variants Reveals Protein Instability Mediated by the DCAF8 E3 Ubiquitin Ligase Adaptor. Cancer discovery, 12(1), 220.

Wang R, et al. (2021) Genetic Screens Identify Host Factors for SARS-CoV-2 and Common Cold Coronaviruses. Cell, 184(1), 106.