

# Resource Summary Report

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

## AAV\_NLS-dSaCas9-NLS-VPR

RRID:Addgene\_68495

Type: Plasmid

### Proper Citation

RRID:Addgene\_68495

### Plasmid Information

**URL:** <http://www.addgene.org/68495>

**Proper Citation:** RRID:Addgene\_68495

**Insert Name:** dSaCas9

**Organism:** Synthetic

**Bacterial Resistance:** Ampicillin

**Defining Citation:** [PMID:26344044](#)

**Vector Backbone Description:** Vector Backbone:pX600-AAV-CMV::NLS-SaCas9-NLS-3xHA-bGHpA Plasmid #61592; Vector Types:Mammalian Expression, AAV; Bacterial Resistance:Ampicillin

**Plasmid Name:** AAV\_NLS-dSaCas9-NLS-VPR

**Record Creation Time:** 20220810T080342+0000

**Record Last Update:** 20260815T082330+0000

### Ratings and Alerts

No rating or validation information has been found for AAV\_NLS-dSaCas9-NLS-VPR.

No alerts have been found for AAV\_NLS-dSaCas9-NLS-VPR.

### Data and Source Information

**Source:** [Addgene](#)

### Usage and Citation Metrics

We found 8 mentions in open access literature.

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

Mesaki K, et al. (2023) Immunomodulation of the donor lung with CRISPR-mediated activation of IL-10 expression. The Journal of heart and lung transplantation : the official publication of the International Society for Heart Transplantation, 42(10), 1363.

Mikaeili H, et al. (2023) Molecular basis of FAAH-OUT-associated human pain insensitivity. Brain : a journal of neurology, 146(9), 3851.

Jensen TI, et al. (2021) Targeted regulation of transcription in primary cells using CRISPRa and CRISPRi. Genome research, 31(11), 2120.

Olson A, et al. (2020) Targeted Chromatinization and Repression of HIV-1 Provirus Transcription with Repurposed CRISPR/Cas9. Viruses, 12(10).

Marina RJ, et al. (2020) Evaluation of Engineered CRISPR-Cas-Mediated Systems for Site-Specific RNA Editing. Cell reports, 33(5), 108350.

Matharu N, et al. (2019) CRISPR-mediated activation of a promoter or enhancer rescues obesity caused by haploinsufficiency. Science (New York, N.Y.), 363(6424).

Fang L, et al. (2019) A Simple Cloning-free Method to Efficiently Induce Gene Expression Using CRISPR/Cas9. Molecular therapy. Nucleic acids, 14, 184.

Haldeman JM, et al. (2019) Creation of versatile cloning platforms for transgene expression and dCas9-based epigenome editing. Nucleic acids research, 47(4), e23.