

Resource Summary Report

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

pMSCV-dCasRx-ALKBH5-PURO

RRID:Addgene_175582

Type: Plasmid

Proper Citation

RRID:Addgene_175582

Plasmid Information

URL: <http://www.addgene.org/175582>

Proper Citation: RRID:Addgene_175582

Insert Name: dCasRx-ALKBH5

Organism: Other

Bacterial Resistance: Ampicillin

Defining Citation: [PMID:34181729](#)

Vector Backbone Description: Vector Backbone:pMSCV; Vector Types:Mammalian Expression, Lentiviral, CRISPR; Bacterial Resistance:Ampicillin

Plasmid Name: pMSCV-dCasRx-ALKBH5-PURO

Relevant Mutation: RfxCas13d R239A/H244A/R858A/H863A

Record Creation Time: 20220422T222015+0000

Record Last Update: 20260815T081052+0000

Ratings and Alerts

No rating or validation information has been found for pMSCV-dCasRx-ALKBH5-PURO.

No alerts have been found for pMSCV-dCasRx-ALKBH5-PURO.

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 6 mentions in open access literature.

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

Wang X, et al. (2026) m6A-Mediated Glycolysis by IL-37 Drives T Cell Metabolic Reprogramming to Regulate Colitis. *Advanced science* (Weinheim, Baden-Wurttemberg, Germany), e20472.

Lan Y, et al. (2025) Synonymous mutations promote tumorigenesis by disrupting m6A-dependent mRNA metabolism. *Cell*, 188(7), 1828.

Khan D, et al. (2024) Homozygous EPRS1 missense variant causing hypomyelinating leukodystrophy-15 alters variant-distal mRNA m6A site accessibility. *Nature communications*, 15(1), 4284.

Collignon E, et al. (2023) m 6 A RNA methylation orchestrates transcriptional dormancy during developmental pausing. *bioRxiv : the preprint server for biology*.

Collignon E, et al. (2023) m6A RNA methylation orchestrates transcriptional dormancy during paused pluripotency. *Nature cell biology*, 25(9), 1279.

Wang L, et al. (2022) m6A modification confers thermal vulnerability to HPV E7 oncotranscripts via reverse regulation of its reader protein IGF2BP1 upon heat stress. *Cell reports*, 41(4), 111546.