

# Resource Summary Report

Generated by [PRECISE-TBI](#) on Sep 2, 2026

## pKanCMV-mRuby3-18aa-actin

RRID:Addgene\_74255

Type: Plasmid

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### Proper Citation

RRID:Addgene\_74255

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### Plasmid Information

**URL:** <http://www.addgene.org/74255>

**Proper Citation:** RRID:Addgene\_74255

**Insert Name:** actin

**Bacterial Resistance:** Kanamycin

**Defining Citation:** [PMID:26879144](#)

**Vector Backbone Description:** Backbone Marker:Michael Davidson Lab; Backbone Size:3900; Vector Backbone:pKanCMV; Vector Types:Mammalian Expression; Bacterial Resistance:Kanamycin

**Comments:** The specific substitutions used in the development of mRuby3 relative to its parent mRuby2 are N33R, M36E, T38V, K74A, G75D, M105T, C114E, H118N, Q120K, H159D, M160I, S171H, S173N, I192V, L202I, M209T, F210Y, H216V, F221Y, A222S, G223N.

**Plasmid Name:** pKanCMV-mRuby3-18aa-actin

**Record Creation Time:** 20220422T222453+0000

**Record Last Update:** 20260902T050124+0000

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### Ratings and Alerts

No rating or validation information has been found for pKanCMV-mRuby3-18aa-actin.

No alerts have been found for pKanCMV-mRuby3-18aa-actin.

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### Data and Source Information

Source: [Addgene](#)

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## Usage and Citation Metrics

We found 6 mentions in open access literature.

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

Dumoulin A, et al. (2024) A cell-autonomous role for primary cilium-mediated signaling in long-range commissural axon guidance. *Development (Cambridge, England)*, 151(17).

Cuhadar U, et al. (2024) Activity-driven synaptic translocation of LGI1 controls excitatory neurotransmission. *Cell reports*, 43(5), 114186.

Farrell RJ, et al. (2024) A ratiometric ER calcium sensor for quantitative comparisons across cell types and subcellular regions. *bioRxiv : the preprint server for biology*.

Cook DC, et al. (2023) GABABR silencing of nerve terminals. *eLife*, 12.

Lankford C, et al. (2022) Identification of HCN1 as a 14-3-3 client. *PloS one*, 17(6), e0268335.

Zhong H, et al. (2021) High-fidelity, efficient, and reversible labeling of endogenous proteins using CRISPR-based designer exon insertion. *eLife*, 10.