

Resource Summary Report

Generated by [RRID](#) on Sep 2, 2026

[pTubb3-MC](#)

RRID:Addgene_87112

Type: Plasmid

Proper Citation

RRID:Addgene_87112

Plasmid Information

URL: <http://www.addgene.org/87112>

Proper Citation: RRID:Addgene_87112

Insert Name: GFP

Organism: Synthetic

Bacterial Resistance: Kanamycin

Defining Citation: [PMID:27851729](#)

Vector Backbone Description: Vector Backbone:pMC.BESPX; Vector Types:CRISPR; Bacterial Resistance:Kanamycin

Comments: It is necessary to remove plasmid backbone with minicircle technology (<https://www.systembio.com/minicircle-dna-vectors/overview>).

Plasmid Name: pTubb3-MC

Record Creation Time: 20220422T222559+0000

Record Last Update: 20260902T050420+0000

Ratings and Alerts

No rating or validation information has been found for pTubb3-MC.

No alerts have been found for pTubb3-MC.

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](#) .

Li JY, et al. (2024) The ergodicity question when imaging DNA conformation using liquid cell electron microscopy. *Proceedings of the National Academy of Sciences of the United States of America*, 121(3), e2314797121.

Pham AM, et al. (2023) Merkel Cell Polyomavirus Large T Antigen Induces Cellular Senescence for Host Growth Arrest and Viral Genome Persistence through Its Unique Domain. *Cells*, 12(3).

Droogers WJ, et al. (2022) Duplex Labeling and Manipulation of Neuronal Proteins Using Sequential CRISPR/Cas9 Gene Editing. *eNeuro*, 9(4).

Kozisek T, et al. (2021) Comparison of promoter, DNA vector, and cationic carrier for efficient transfection of hMSCs from multiple donors and tissue sources. *Molecular therapy*. *Nucleic acids*, 26, 81.

Han B, et al. (2021) Bi-FoRe: an efficient bidirectional knockin strategy to generate pairwise conditional alleles with fluorescent indicators. *Protein & cell*, 12(1), 39.

Suzuki K, et al. (2019) Precise in vivo genome editing via single homology arm donor mediated intron-targeting gene integration for genetic disease correction. *Cell research*, 29(10), 804.