

# Resource Summary Report

Generated by [dkNET](#) on Sep 4, 2026

## pCoofy49

RRID:Addgene\_55187

Type: Plasmid

### Proper Citation

RRID:Addgene\_55187

### Plasmid Information

**URL:** <http://www.addgene.org/55187>

**Proper Citation:** RRID:Addgene\_55187

**Bacterial Resistance:** Kanamycin

**Defining Citation:** [PMID:23410102](#)

**Vector Backbone Description:** Backbone Marker:Novagen; Backbone Size:7701; Vector Backbone:pET, pCoofy4; Vector Types:Bacterial Expression; Bacterial Resistance:Kanamycin

**Comments:** C-terminal tags are separated by a stop codon. Depending on the LP2 primer used in SLIC, the desired C-terminal tag(s) can be fused to the target gene. Use the one of the following linearization primers pairs for SLIC cloning. Choose one LP1 forward vector primer and one LP2 reverse vector primer: LP1 forward vector primer: SLIC Primer (3C site) 5' GGGCCCCTGGAACAGAACTTCCAG 3' LP2 - select an LP2 primer below depending on which C-terminal tags you want to include fused to your gene of interest: none SLIC Primer 5' CGCCATTAACCTGATGTTCTGGGG 3' 10His- SLIC Primer 5`GAGCATCATCATCATCACAC 3' StrepOne - SLIC Primer 5`AGCGCTTGGAGCCACCCGCAG 3' S-Tag - SLIC Primer 5`AAAGAAACCGCTGCTGCTAAATTCG 3' CBP - SLIC Primer 5`ATGAAGCGGCGGTGGAAGAAAAAC 3' HPC4 - SLIC Primer 5`GAGGACCAGGTGGACCCCGG 3' ??54CPD - SLIC Primer 5`GGCAGCGGCAAGATCCTGCAC 3' Test each preparation of the plasmid by transforming it into non-resistant cells (ex. DH5a) to ensure negative selection by ccdB kills all the transformed cells as expected.

**Plasmid Name:** pCoofy49

**Record Creation Time:** 20220422T222323+0000

**Record Last Update:** 20260829T080717+0000

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## Ratings and Alerts

No rating or validation information has been found for pCoofy49.

No alerts have been found for pCoofy49.

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## Data and Source Information

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

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## Usage and Citation Metrics

We have not found any literature mentions for this resource.