

# Resource Summary Report

Generated by [dkNET](#) on Sep 18, 2026

## p667-UBC-CDC42BPB-V5-miniTurbo\_IDG-K

RRID:Addgene\_198008

Type: Plasmid

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### Proper Citation

RRID:Addgene\_198008

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### Plasmid Information

**URL:** <http://www.addgene.org/198008>

**Proper Citation:** RRID:Addgene\_198008

**Insert Name:** CDC42BPB-V5-miniTurbo

**Organism:** Homo sapiens

**Bacterial Resistance:** Ampicillin

**Vector Backbone Description:** Backbone Marker:PEL; Backbone Size:9014; Vector Backbone:pDest667; Vector Types:Mammalian Expression, Lentiviral, Gateway Destination; Bacterial Resistance:Ampicillin

**Comments:** These plasmids were generated as part of the Illuminating the Druggable Genome (IDG) program sponsored by the NIH Common Fund. The goal of this program is to identify, gather, and distribute information and resources for proteins that currently are not well-studied yet belong to commonly drug-targeted protein families: protein kinases, non-olfactory G-protein coupled receptors (GPCRs), and ion channels. The IDG program is designed to develop fundamental research tools for understudied proteins, elucidate their function, and disseminate the IDG-related resources and data to the greater scientific community. These lentiviral gateway destination plasmids were generated, as part of the Kinase-Data and Resource Generating Center (DRGC), using the single fragment or multisite gateway cloning technology. They were used for generating proteomics-based protein-interaction and protein-proximity networks that can be accessed through the kinase-DRGC website <https://darkkinome.org/>. Each plasmid has either an N- or C-terminal fusion protein (Flag/ V5/V5-miniTurbo/V5-TurboID/miniTurbo-V5/TurboID-V5) tagged understudied kinase driven under either a CMV or Ubiquitin (UBC) promoter. All inserts in the pENTR plasmids used to generate the deposited destination plasmids were end-sequenced and insert size validated prior to multi-site gateway cloning. The combined insert size of the single fragment or three fragment destination plasmids were confirmed by restriction digestion (BsrGI, and EcoRV for pDest667; BsrGI, EcoRV, and BstZ17I for pDest663; and BsrGI for pHAGE) and all plasmids were partially sequenced to ensure in-

frame ORFs and fusion tags.

**Plasmid Name:** p667-UBC-CDC42BPB-V5-miniTurbo\_IDG-K

**Record Creation Time:** 20230414T080544+0000

**Record Last Update:** 20260912T093739+0000

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## Ratings and Alerts

No rating or validation information has been found for p667-UBC-CDC42BPB-V5-miniTurbo\_IDG-K.

No alerts have been found for p667-UBC-CDC42BPB-V5-miniTurbo\_IDG-K.

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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.