

# Resource Summary Report

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

## Double UP mNeon to mScarlet Rac1-N17

RRID:Addgene\_125137

Type: Plasmid

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### Proper Citation

RRID:Addgene\_125137

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### Plasmid Information

**URL:** <http://www.addgene.org/125137>

**Proper Citation:** RRID:Addgene\_125137

**Insert Name:** mNeon/mScarletP2A Rac1-N17

**Organism:** Other

**Bacterial Resistance:** Ampicillin

**Defining Citation:** [PMID:32508591](#)

**Vector Backbone Description:** Backbone Size:6734; Vector Backbone:pCax; Vector Types:Mammalian Expression; Bacterial Resistance:Ampicillin

**Comments:** Intent is to introduce low doses of pCAG Cre to recombine a subset of cells, such that mNeon can act as an internal control to mScarlet for use in in utero electroporation or other related techniques. Rac1-N17 is expressed following a P2A site, so that Cre+ cells will be both mScarlet positive and also contain an untagged Rac1-N17. Note: Addgene NGS is unable to fully resolve the CAG promoter sequence. Please refer to the depositor's sequence for this section of the plasmid.

**Plasmid Name:** Double UP mNeon to mScarlet Rac1-N17

**Relevant Mutation:** Removed Not1 site in mScarlet. 17th AA in Rac1 changed to N

**Record Creation Time:** 20220422T221656+0000

**Record Last Update:** 20260902T041603+0000

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### Ratings and Alerts

No rating or validation information has been found for Double UP mNeon to mScarlet Rac1-N17.

No alerts have been found for Double UP mNeon to mScarlet Rac1-N17.

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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.