

Resource Summary Report

Generated by [NIF](#) on Aug 24, 2026

pET His6 High pl protein 68 TEV LIC cloning vector (2-Hpl-68)

RRID:Addgene_55211

Type: Plasmid

Proper Citation

RRID:Addgene_55211

Plasmid Information

URL: <http://www.addgene.org/55211>

Proper Citation: RRID:Addgene_55211

Bacterial Resistance: Ampicillin

Vector Backbone Description: Vector Backbone:pET; Vector Types:Bacterial Expression; Bacterial Resistance:Ampicillin

Comments: This plasmid is an empty vector. Your gene can be inserted with a LIC cloning protocol. All 2-series vectors work as single-expression vectors, as well as transfer vectors for our polycistronic system. The LIC cloning site is flanked by 5 pairs of restriction sites, so that your gene can easily be subcloned into our polycistronic destination vectors (2D, 2E, or 2Z). 2-Hpl-68 has a TEV-cleavable N-terminal His6-High pl protein fusion tag. The high pl can improve solubility of your target protein. To clone into this vector, add LIC v1 tags to the 5' end of your PCR primers. Forward - 5'-TACTTCCAATCCAATGCA-3' Reverse - 5'-TTATCCACTTCCAATGTTATTA-3' Linearize the plasmid with SspI and gel purify. When digesting the DNA with T4 polymerase, use dCTP for insert and dGTP for vector.

Plasmid Name: pET His6 High pl protein 68 TEV LIC cloning vector (2-Hpl-68)

Record Creation Time: 20220422T222323+0000

Record Last Update: 20260815T081648+0000

Ratings and Alerts

No rating or validation information has been found for pET His6 High pl protein 68 TEV LIC cloning vector (2-Hpl-68).

No alerts have been found for pET His6 High pl protein 68 TEV LIC cloning vector (2-Hpl-68).

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 1 mentions in open access literature.

Listed below are recent publications. The full list is available at [NIF](#) .

Clerici M, et al. (2018) Structural basis of AAUAAA polyadenylation signal recognition by the human CPSF complex. Nature structural & molecular biology, 25(2), 135.