

# Resource Summary Report

Generated by [RRID](#) on Aug 20, 2026

## [pSBtet-Hyg](#)

RRID:Addgene\_60508

Type: Plasmid

### Proper Citation

RRID:Addgene\_60508

### Plasmid Information

**URL:** <http://www.addgene.org/60508>

**Proper Citation:** RRID:Addgene\_60508

**Bacterial Resistance:** Ampicillin

**Defining Citation:** [PMID:25650551](#)

**Vector Backbone Description:** Backbone Size:2021; Vector Backbone:pUC19; Vector Types:Mammalian Expression, Transposon; Bacterial Resistance:Ampicillin

**Plasmid Name:** pSBtet-Hyg

**Record Creation Time:** 20220422T222348+0000

**Record Last Update:** 20260815T081733+0000

### Ratings and Alerts

No rating or validation information has been found for pSBtet-Hyg.

No alerts have been found for pSBtet-Hyg.

### Data and Source Information

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

### Usage and Citation Metrics

We found 9 mentions in open access literature.

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

Edupuganti RR, et al. (2025) Integrator promotes the association of TFIID and RNA polymerase II to maintain pluripotency during development. *Molecular cell*, 85(15), 2937.

Kumari S, et al. (2025) FLAG Immunoprecipitation-Based Mapping of the In Vivo Assembled Spliceosomal C\* Complex. *International journal of molecular sciences*, 26(20).

Laarne M, et al. (2025) A homozygous single-nucleotide variant in TNNT1 causes abnormal troponin T isoform expression in a patient with severe nemaline myopathy: A case report. *Journal of neuromuscular diseases*, 12(5), 689.

Aleksashin NA, et al. (2023) A highly efficient human cell-free translation system. *bioRxiv* : the preprint server for biology.

Sarparanta J, et al. (2023) Extension of the DNAJB2a isoform in a dominant neuromyopathy family. *Human molecular genetics*.

Bram Y, et al. (2022) Dual-Reporter System for Real-Time Monitoring of SARS-CoV-2 Main Protease Activity in Live Cells Enables Identification of an Allosteric Inhibition Path. *ACS bio & med chem Au*, 2(6), 627.

Johari M, et al. (2021) Missense mutations in small muscle protein X-linked (SMPX) cause distal myopathy with protein inclusions. *Acta neuropathologica*, 142(2), 375.

Yeung TK, et al. (2021) One-step multiplex toolkit for efficient generation of conditional gene silencing human cell lines. *Molecular biology of the cell*, 32(14), 1320.

Gerbracht JV, et al. (2020) CASC3 promotes transcriptome-wide activation of nonsense-mediated decay by the exon junction complex. *Nucleic acids research*, 48(15), 8626.