

# Resource Summary Report

Generated by [NIF](#) on Jul 29, 2026

## pSAR-MT

RRID:Addgene\_16485

Type: Plasmid

### Proper Citation

RRID:Addgene\_16485

### Plasmid Information

**URL:** <http://www.addgene.org/16485>

**Proper Citation:** RRID:Addgene\_16485

**Bacterial Resistance:** Ampicillin

**Defining Citation:** [PMID:8755583](#)

**Vector Backbone Description:** Backbone Size:0; Vector Backbone:pSAR-MT; Vector Types:Mammalian Expression; Bacterial Resistance:Ampicillin

**Comments:** The pSAR-MT inducible expression plasmid was constructed as follows. Two oligonucleotides (5'-GATCTCGAGCTCCCTGCA-3' and 5'-GGGAGCTCGA-3') were annealed, and the resulting fragment was cloned into the BamHI and PstI sites of the scaffold attachment region (SAR)-containing plasmid p8. The resulting plasmid, pJM7, consisted of two SAR sequences flanking a short polylinker containing restriction sites for XhoI and PstI. The mutant metallothionein (MT) promoter was amplified by PCR and cloned into pCEP4, linearized with BglII and NotI, yielding pCEP-MT. A Sall-BamHI fragment containing an intron from the globin gene was then inserted into the XhoI and BamHI sites of pCEP-MT, yielding pCEP-MTi. Finally, the MT promoter/intron/poly(A) region was removed from pCEP-MTi by digestion with Sall and cloned into the XhoI site of pJM7, yielding pSAR-MT.

**Plasmid Name:** pSAR-MT

**Record Creation Time:** 20220422T221931+0000

**Record Last Update:** 20260725T074159+0000

### Ratings and Alerts

No rating or validation information has been found for pSAR-MT.

No alerts have been found for pSAR-MT.

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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.