

Resource Summary Report

Generated by [nidm-terms](#) on Sep 19, 2026

pCXLE-hOCT3/4

RRID:Addgene_27076

Type: Plasmid

Proper Citation

RRID:Addgene_27076

Plasmid Information

URL: <http://www.addgene.org/27076>

Proper Citation: RRID:Addgene_27076

Insert Name: OCT3/4

Organism: Homo sapiens

Bacterial Resistance: Ampicillin

Defining Citation: [PMID:21460823](#)

Vector Backbone Description: Backbone Size:10180; Vector Backbone:pCXLE; Vector Types:Mammalian Expression; Bacterial Resistance:Ampicillin

Plasmid Name: pCXLE-hOCT3/4

Record Creation Time: 20220422T222121+0000

Record Last Update: 20260919T091631+0000

Ratings and Alerts

No rating or validation information has been found for pCXLE-hOCT3/4.

No alerts have been found for pCXLE-hOCT3/4.

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 52 mentions in open access literature.

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

Loureiro JR, et al. (2026) The insertion of an ATTTC repeat in an Alu element hyperactivates a neurodevelopmental enhancer in spinocerebellar ataxia type 37. *Cell reports*, 45(4), 117146.

Marteau S, et al. (2025) Generation of a FAM189A2/ENTREP1 knockout human induced pluripotent stem cell line using CRISPR/Cas9 technology. *Stem cell research*, 89, 103857.

Almaghrabi R, et al. (2025) A heterozygous CEBPA mutation disrupting the bZIP domain in a RUNX1 and SRSF2 mutational background causes MDS disease progression. *Nature communications*, 16(1), 5489.

Urdaneta KE, et al. (2025) Generation of patient-derived and gene-corrected hiPSC lines from Hereditary Hemorrhagic Telangiectasia type 2 patients with ACVRL1 c.1042delG mutation. *Stem cell research*, 88, 103812.

Pierre B, et al. (2024) Generation of CRISPR/Cas9 edited human induced pluripotent stem cell line carrying the heterozygous p.H695VfsX5 frameshift mutation in the exon 10 of the PKP2 gene. *Stem cell research*, 76, 103341.

Duboscq-Bidot L, et al. (2024) Generation of CRISPR-Cas9 edited human induced pluripotent stem cell line carrying BAG3 V468M mutation in its BAG domain. *Stem cell research*, 74, 103294.

MacCarthy CM, et al. (2024) Highly cooperative chimeric super-SOX induces naive pluripotency across species. *Cell stem cell*, 31(1), 127.

Kim M, et al. (2023) Generation of Induced Pluripotent Stem Cells from Lymphoblastoid Cell Lines by Electroporation of Episomal Vectors. *International journal of stem cells*, 16(1), 36.

Chemla A, et al. (2023) Generation of two induced pluripotent stem cell lines and the corresponding isogenic controls from Parkinson's disease patients carrying the heterozygous mutations c.815G > A (p.R272Q) or c.1348C > T (p.R450C) in the RHOT1 gene encoding Miro1. *Stem cell research*, 71, 103145.

Baden P, et al. (2023) Glucocerebrosidase is imported into mitochondria and preserves complex I integrity and energy metabolism. *Nature communications*, 14(1), 1930.

Sukparangsi W, et al. (2022) Establishment of fishing cat cell biobanking for sustainable conservation. *Frontiers in veterinary science*, 9, 989670.

Mencke P, et al. (2022) Generation and characterization of a genetic Parkinson's disease-patient derived iPSC line DJ-1-delP (LCSBi008-A). *Stem cell research*, 62, 102792.

Ader F, et al. (2022) Generation of CRISPR-Cas9 edited human induced pluripotent stem cell line carrying FLNC exon skipping variant. *Stem cell research*, 58, 102616.

Mencke P, et al. (2022) Generation of isogenic control DJ-1-delP GC13 for the genetic Parkinson's disease-patient derived iPSC line DJ-1-delP (LCSBi008-A-1). Stem cell research, 62, 102815.

Mengel D, et al. (2021) A de novo STUB1 variant associated with an early adult-onset multisystemic ataxia phenotype. Journal of neurology, 268(10), 3845.

de Rus Jacquet A, et al. (2021) The LRRK2 G2019S mutation alters astrocyte-to-neuron communication via extracellular vesicles and induces neuron atrophy in a human iPSC-derived model of Parkinson's disease. eLife, 10.

Inzunza J, et al. (2021) Generation of nonviral integration-free human iPS cell line KISCOi001-A from normal human fibroblasts, under defined xeno-free and feeder-free conditions. Stem cell research, 51, 102193.

Liu R, et al. (2021) Generation of two induced pluripotent stem cell lines from blood cells of a prostate cancer patient carrying germline mutation in CHEK2. Stem cell research, 53, 102299.

Bliley JM, et al. (2021) Dynamic loading of human engineered heart tissue enhances contractile function and drives a desmosome-linked disease phenotype. Science translational medicine, 13(603).

Vermeer MC, et al. (2021) Gain-of-function mutation in ubiquitin-ligase KLHL24 causes desmin degradation and dilatation in hiPSC-derived engineered heart tissues. The Journal of clinical investigation, 131(17).