



pCXLE-hOCT3/4

(Plasmid #27076)

Purpose

Non-integrating (episomal) expression of human OCT3/4

Depositing Lab

Shinya Yamanaka

Publication

Okita et al Nat Methods. 2011 May;8(5):409-12 (How to cite ♥)

Sequence Information

Sequences (5)

Ordering

This material is available to academics and nonprofits only.

Item	Catalog #	Description	Quantity	Price (USD)		
P l asmid	27076	Standard format: Plasmid sent in bacteria as agar stab	1	\$85	Add to Cart	

Backbone

Vector backbone: pCXLE

Backbone size w/o insert (bp): 10180 Vector type: Mammalian Expression

Growth in Bacteria

Bacterial Resistance(s): Ampicillin, 100 µg/mL

Growth Temperature: 37°C **Growth Strain(s):** DH5alpha **Copy number:** High Copy

Gene/Insert

Gene/Insert name: OCT3/4
Species: H. sapiens (human)
Insert Size (bp): 1108

Entrez Gene: POU5F1 (a.k.a. (also known as) OCT3, OCT4, OTF-3,

OTF3, OTF4, Oct-3, Oct-4, Oct3/4)

Cloning Information

Cloning method: Restriction Enzyme
5' cloning site: EcoRI (not destroyed)
3' cloning site: EcoRI (not destroyed)
5' sequencing primer: pCAG-F
3' sequencing primer: WPRE-R
(Common Sequencing Primers)

Resource Information

A portion of this plasmid was derived from a plasmid made by: pCEP4 is from Invitrogen. CAG Promoter was from Dr. Jun-ichi Miyazaki of Osaka University Graduate School of Medicine. In publication using this plasmid, please cite: Efficient selection for high-expression

transfectants with a novel eukaryotic vector. Gene 108:193-200, 1991. Niwa, H., Yamamura, K. & Miyazaki, J.

Articles Citing this Plasmid:

• 43 References

Terms and Licenses

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• <u>UBMTA</u>

Industry Terms:

• Not Available to Industry

Trademarks:

• Zeocin® is an InvivoGen trademark.

How to cite this plasmid (Back to top ♠)

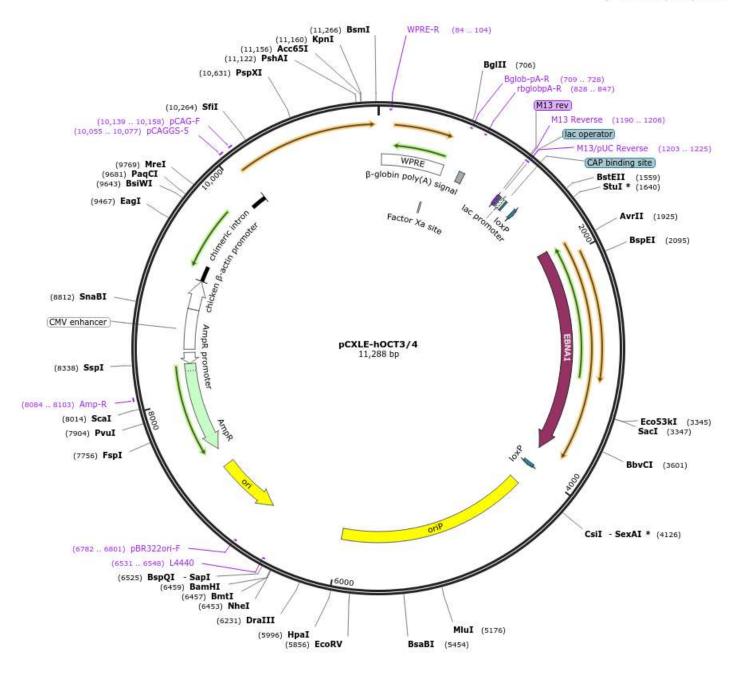
These plasmids were created by your colleagues. Please acknowledge the Principal Investigator, cite the article in which the plasmids were described, and include Addgene in the Materials and Methods of your future publications.

For your Materials & Methods section:

pCXLE-hOCT3/4 was a gift from Shinya Yamanaka (Addgene plasmid # 27076; http://n2t.net/addgene:27076; RRID:Addgene_27076)

For your References section:

A more efficient method to generate integration-free human iPS cells. Okita K, Matsumura Y, Sato Y, Okada A, Morizane A, Okamoto S, Hong H, Nakagawa M, Tanabe K, Tezuka KI, Shibata T, Kunisada T, Takahashi M, Takahashi J, Saji H, Yamanaka S. *Nat Methods*. 2011 May;8(5):409-12 10.1038/nmeth.1591 PubMed 21460823



Yamanaka Lab's Plasmid Stock # 5/-7/

Name: pCXLE - hOct 3/4

Size: //, 3 k bp

Constructed by: K. Okita

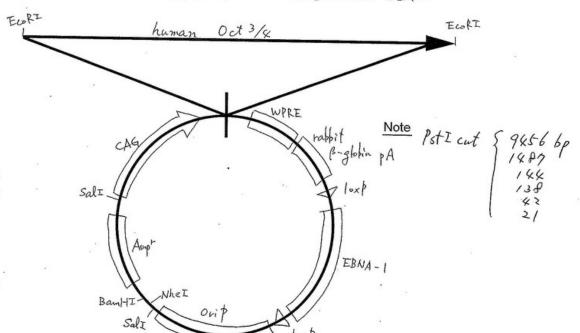
Date: May 2009

Insert

Description: $p To po - h O ct \frac{3}{4}$ (# 43 - 8)

Size: /, /k bp

Enzyme used: Ecolo



Vector

Description:

PCXLE-GFP. (#51-69) Size: 10.2k bp

Enzyme used:

EcoRI





pCXLE-hSK (Plasmid #27078)

Purpose

Integration-free (episomal) expression of human SOX2 and KLF4

Depositing Lab

Shinya Yamanaka

Publication

Okita et al Nat Methods. 2011 May;8(5):409-12 (How to cite ♥)

Sequence Information

Sequences (8)

Ordering

This material is available to academics and nonprofits only.

Item	Catalog #	Description	Quantity	Price (USD)		
P l asmid	27078	Standard format: Plasmid sent in bacteria as agar stab	1	\$85	Add to Cart	

Backbone

Vector backbone: pCXLE

Backbone size w/o insert (bp): 10180 Vector type: Mammalian Expression

Growth in Bacteria

Bacterial Resistance(s): Ampicillin, 100 µg/mL

Growth Temperature: 37°C **Growth Strain(s):** DH5alpha **Copy number:** High Copy

Gene/Insert

Gene/Insert name: SOX2, KLF4 **Species:** H. sapiens (human) **Insert Size (bp):** 2513

Entrez Gene: KLF4 (a.k.a. (also known as) EZF, GKLF)

Entrez Gene: SOX2 (a.k.a. (also known as) ANOP3, MCOPS3)

Cloning Information

Cloning method: Restriction Enzyme
5' cloning site: EcoRl (not destroyed)
3' cloning site: EcoRl (not destroyed)
5' sequencing primer: pCAG-F
3' sequencing primer: WPRE-R
(Common Sequencing Primers)

Resource Information

A portion of this plasmid was derived from a plasmid made by: pCEP4 is from Invitrogen. CAG Promoter was from Dr. Jun-ichi Miyazaki of Osaka University Graduate School of Medicine. In publication using this plasmid, please cite: Efficient selection for high-expression transfectants with a novel eukaryotic vector. Gene 108:193-200, 1991.

Niwa, H., Yamamura, K. & Miyazaki, J. Mutation(s) made by Provider lab: Two genes are connected with the foot-and-mousht desease virus 2A self-cleaving sequence.

Articles Citing this Plasmid:

• 312 References

Terms and Licenses

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Industry Terms:

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Trademarks:

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How to cite this plasmid (Back to top ↑)

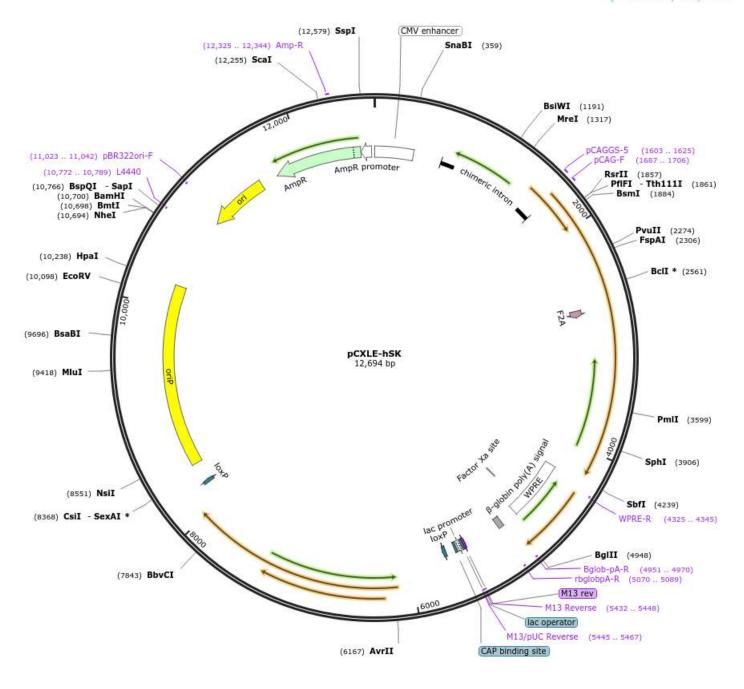
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For your Materials & Methods section:

pCXLE-hSK was a gift from Shinya Yamanaka (Addgene plasmid # 27078; http://n2t.net/addgene:27078; RRID:Addgene_27078)

For your References section:

A more efficient method to generate integration-free human iPS cells. Okita K, Matsumura Y, Sato Y, Okada A, Morizane A, Okamoto S, Hong H, Nakagawa M, Tanabe K, Tezuka KI, Shibata T, Kunisada T, Takahashi M, Takahashi J, Saji H, Yamanaka S. *Nat Methods*. 2011 May;8(5):409-12 10.1038/nmeth.1591 PubMed 21460823



Yamanaka Lab's Plasmid Stock # 51-74

Name: pCXLE-hSK

Size: /2.7k bp

Constructed by: K. Okita

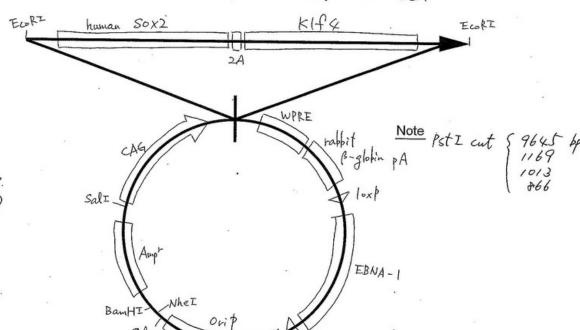
Date: May 2009

Insert

Description: pBS-hSox2-2A-KIF4 (# 46-38)

Size: 2,5k bp

Enzyme used: Ecokt



Vector

Description:

PCXLE-GFP. (#51-69)

Size: 10, 2 k bp

Enzyme used:

EcoRI





Purpose

Integration-free (episomal) expression of human L-MYC and LIN28

Depositing Lab

Shinya Yamanaka

Publication

Okita et al Nat Methods. 2011 May;8(5):409-12 (How to cite ♥)

Sequence Information

Sequences (5)

Ordering

This material is available to academics and nonprofits only.

Item	Catalog #	Description	Quantity	Price (USD)		
Plasmid	27080	Standard format: Plasmid sent in bacteria as agar stab	1	\$85	Add to Cart	

Backbone

Vector backbone: pCXLE

Backbone size w/o insert (bp): 10180 Vector type: Mammalian Expression

Growth in Bacteria

Bacterial Resistance(s): Ampicillin, 100 µg/mL

Growth Temperature: 37°C Growth Strain(s): DH5alpha Copy number: High Copy

Gene/Insert

Gene/Insert name: L-MYC

Alt name: LIN28

Species: H. sapiens (human) Insert Size (bp): 1871

Entrez Gene: LIN28A (a.k.a. (also known as) CSDD1, LIN-28, LIN28,

ZCCHC1, lin-28A)

Entrez Gene: MYCL (a.k.a. (also known as) L-Myc, LMYC, MYCL1,

bHLHe38)

Cloning Information

Cloning method: Restriction Enzyme
5' cloning site: EcoRI (not destroyed)
3' cloning site: EcoRI (not destroyed)
5' sequencing primer: pCAG-F
3' sequencing primer: WPRE-R
(Common Sequencing Primers)

Resource Information

A portion of this plasmid was derived from a plasmid made by: pCEP4

is from Invitrogen. CAG Promoter was from Dr. Jun-ichi Miyazaki of Osaka University Graduate School of Medicine. In publication using this plasmid, please cite: Efficient selection for high-expression transfectants with a novel eukaryotic vector. Gene 108:193-200, 1991. Niwa, H., Yamamura, K. & Miyazaki, J. Mutation(s) made by Provider lab: Two genes are connected with the foot-and-mouth disease virus 2A self-cleaving sequence.

Articles Citing this Plasmid:

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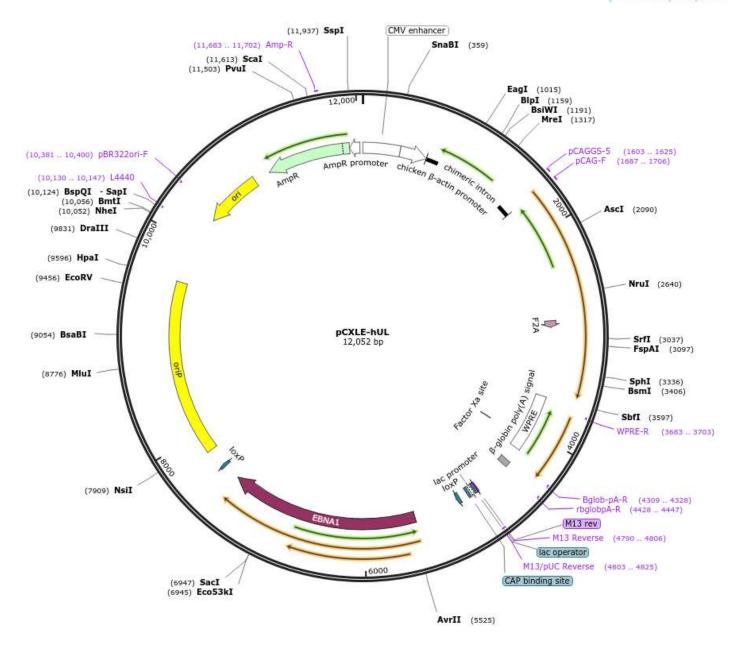
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For your Materials & Methods section:

pCXLE-hUL was a gift from Shinya Yamanaka (Addgene plasmid # 27080 ; http://n2t.net/addgene:27080 ; RRID:Addgene_27080)

For your References section:

A more efficient method to generate integration-free human iPS cells. Okita K, Matsumura Y, Sato Y, Okada A, Morizane A, Okamoto S, Hong H, Nakagawa M, Tanabe K, Tezuka KI, Shibata T, Kunisada T, Takahashi M, Takahashi J, Saji H, Yamanaka S. *Nat Methods*. 2011 May;8(5):409-12 10.1038/nmeth.1591 PubMed 21460823



Yamanaka Lab's Plasmid Stock # 52-28

Name: pCXLE-hUL.

Size: /2 / k bp

Constructed by: X. Okita

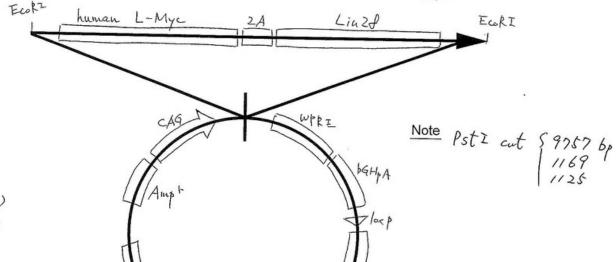
Date: 2009.5.19

Insert

Description: pcx - hUL

Size: 1,9 k bp

Enzyme used: EcokI



Vector

Description:

(#\$1-69)

Size: 10,2k bp

Enzyme used:

EcokI