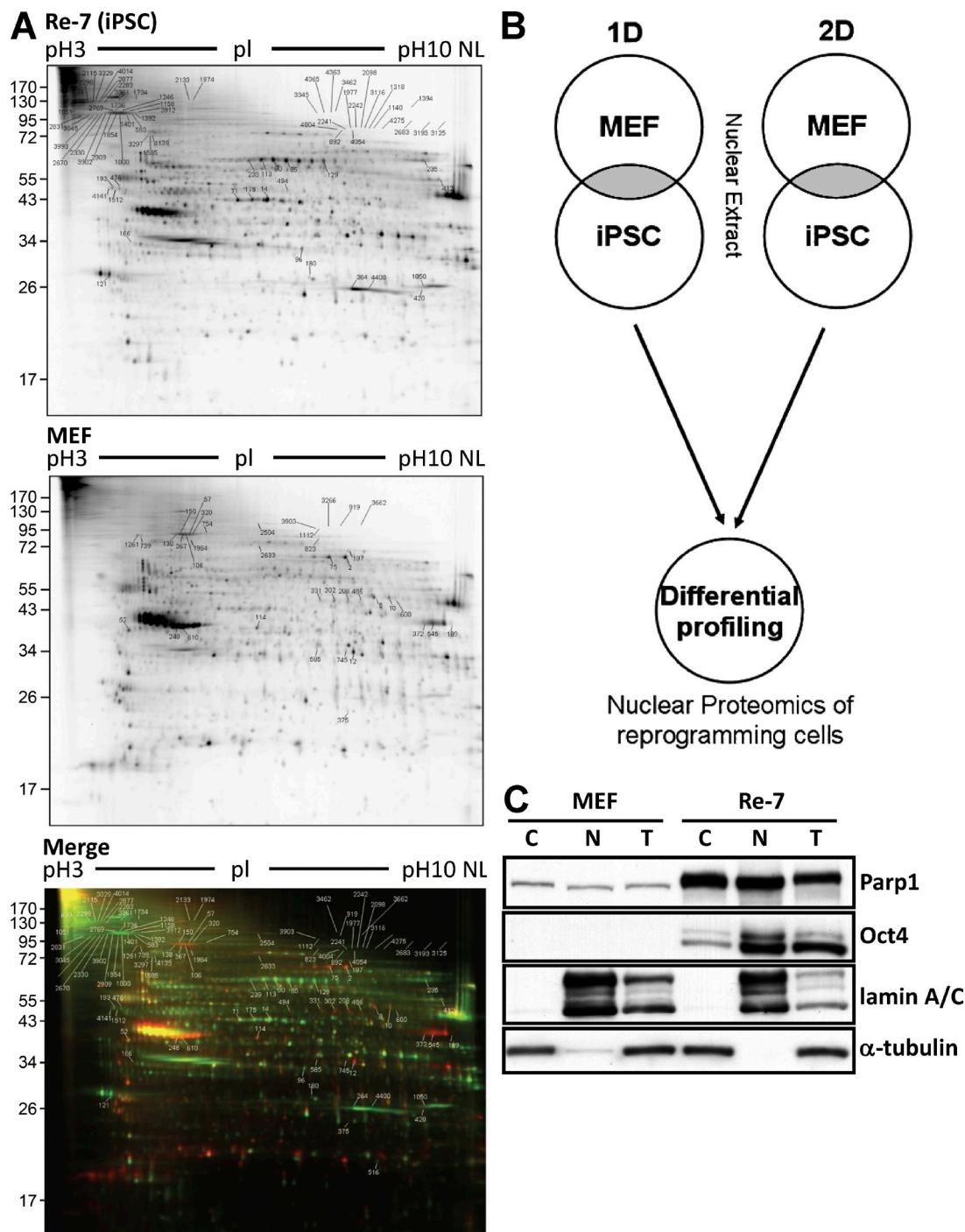


## SUPPLEMENTAL MATERIAL

Chiou et al., <http://www.jem.org/cgi/content/full/jem.20121044/DC1>

**Figure S1. Establishment of the differential profiling of nuclear proteins from pluripotent stem cells and MEFs by MS-based proteomics.** (A) 2D-differential gel electrophoresis (2D-DIGE) and silver staining. Then the identified candidate proteins confirmed by LC-MS/MS. (B) Based on the statistical comparative analysis between the databases of 1D LC-MS/MS and 2D-differential gel electrophoresis, we sorted the hubs of nuclear proteins between the nuclear profiles from MEFs and iPSCs. (C) Detection of Parp1 and Oct4 expression in nuclear fractions and whole-cell lysates of MEFs and iPSCs. C: cytoplasmal fraction; N: nuclear fraction; T: total cell lysate.





















**Table S2.** The sequences of the primers for quantitative RT-PCR and RT-PCR

Gene, accession no.	Primer sequence (5' to 3')	Product size bp
<b>Quantitative RT-PCR</b>		
Parp1, NM_007415.2	F: CGGAGAGGCTTATCGAGTGGAGTA; R: AAGCAGGAGAAGTGGTACCGAGTGTG	157
Nanog, NM_028016.2	F: TGGTCCCCACAGTTGCCTAGTTCT; R: GGTAGAAGAACAGGGCTGCCTGA	192
Gapdh, NM_008084.2	F: CTCATGACCACAGTCCATGC; R: TTCAGCTCTGGATGACCTT	155
<b>RT-PCR</b>		
Nanog, NM_028016.2	F: CCTTGAGCCGTTGGCCTTCAGAT; R: GCTGCCACATGGAAAGGCT	808
Dppa5aL (Esg1), NM_025274.3	F: CGCCCACACAGGTACTAAAACCTCCTG; R: ACAGTGGCCACAGCTAACCTGC	515
ErasL, NM_181548.2	F: CCAACTGTCCGGTCAGATCCGC; R: GCCTCCTGGGCCCTCTGAATCT	713
mRex1 (Zfp42), NM_009556.3	F: GCCAGCAGCTCCTGCACACA; R: TGAGCTGCCCAACCCCTCA	631
Nat (Eif4g2), NM_001040131.2	F: ATTCTTCGTTGCAAGCCGCAAAGTGGAG; R: AGTTGTTGCTGCGGAGTTGTCATCTCGTC	223

F, forward; R, reverse.

**Table S3.** The sequences of shRNA and the primers for Parp1 promoter constructs

shRNA/construct	Sequence/primer (5' to 3')
shParp1	
TRCN0000071209	GCCCTTGGAAACATGTATGAA
shParp2	
Oligo Sequence	CCGGCGGTTACCAGTCTCTCAAGAACTCGAGTTCTGAGAGACTGTAACCGTTTT
Parp1 promoter constructs	
-2,000 fragment	F: CCATGATAACAATGACATGGCGAAAG; R: CTCTCGCGTACTCCACTCGATAAA
-1,650 fragment	F: AGACAGGGTTCTCTGTAGCCCT; R: CTCTCGCGTACTCCACTCGATAAA
-1,100 fragments	F: GAAAAAGAGCAGCAACAGAGGAAGC; R: CTCTCGCGTACTCCACTCGATAAA
-600 fragments	F: GATGAGAACACACGACATCTGTTGT; R: CTCTCGCGTACTCCACTCGATAAA
-125 fragments	F: CTAGGCATCAGTAATCTATCCTGAG; R: CTCTCGCGTACTCCACTCGATAAA
C1 c-Myc site	F: TTCCAGACAGGGTTCTGTGAG; R: CAAGTCCTCTGAAAGAGTAACCACT
C2 c-Myc site	F: CTTTATAGTGAGACCCCTGCATCACG; R: TAGATCA G GCTGGCCAAGAACCTCAC
C3 c-Myc site	F: AAGCCTTCTGGGACAAACCGACAA; R: TCAGGATAGATTACTGATGCCTAGC

F, forward; R, reverse.

**Table S4.** List of proteins tested by antibodies

Protein	Assay	Antibody	Origin	Dilution	Incubation period
Oct4	WB	mmAb	Cell Signaling Technology	1:1,000	Overnight
Oct4	IF	rmAb	Cell Signaling Technology	1:200	Overnight
Sox2	WB	rmAb	Cell Signaling Technology	1:1,000	Overnight
Klf4	WB	rpAb	Cell Signaling Technology	1:1,000	Overnight
c-Myc	WB	rmAb	Cell Signaling Technology	1:1,000	Overnight
Parp1	WB, IF, IP	rmAb	Cell Signaling Technology	1:1,000, 1:200, 1:100	Overnight
Nanog	WB	rpAb	Millipore	1:1,000	Overnight
Lamin A/C	WB	rmAb	Cell Signaling Technology	1:1,000	Overnight
Pcna	WB	mpAb	Cell Signaling Technology	1:2,000	Overnight
Topoisomerase II $\alpha$	WB	rpAb	Cell Signaling Technology	1:500	Overnight
Parp2	WB	mmAb	Millipore	1:500	Overnight
Chd1L	WB	mmAb	Abcam	1:50	Overnight
Ssrp1	WB	mmAb	Santa Cruz Biotechnology, Inc.	1:300	Overnight
DNA ligase III	WB	rpAb	GeneTex	1:500	Overnight
Ku-70(Xrcc-6)	WB	gpAb	Santa Cruz Biotechnology, Inc.	1:500	Overnight
Poly(ADP-ribose)	WB, IF	mmAb	Enzo Life Sciences	1:1,000, 1:100	Overnight
$\beta$ -Actin	WB	mmAb	Millipore	1:10,000	Overnight
$\alpha$ -Tubulin	WB	mmAb	Sigma-Aldrich	1:10,000	Overnight

Abbreviations: WB, Western blot; mmAb, mouse mAb; rmAb, rabbit mAb; rpAb, rabbit polyclonal Ab; gpAb, goat pAb; FC, flow cytometry; IF, immunofluorescence; IP, immunoprecipitation.