Single-Cell Analyses of ESCs Reveal Alternative Pluripotent Cell States and Molecular Mechanisms that Control Self-Renewal

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SUMMARY

Analyses of gene expression in single mouse embryonic stem cells (mESCs) cultured in serum and LIF revealed the presence of two distinct cell subpopulations with individual gene expression signatures. Comparisons with published data revealed that cells in the first subpopulation are phenotypically similar to cells isolated from the inner cell mass (ICM). In contrast, cells in the second subpopulation appear to be more mature. Pluripotency Gene Regulatory Network (PGRN) reconstruction based on single-cell data and published data suggested antagonistic roles for Oct4 and Nanog in the maintenance of pluripotency states. Integrated analyses of published genomic binding (ChIP) data strongly supported this observation. Certain target genes alternatively regulated by OCT4 and NANOG, such as Sall4 and Zscan10, feed back into the top hierarchical regulator Oct4. Analyses of such incoherent feedforward loops with feedback (iFFL-FB) suggest a dynamic model for the maintenance of mESC pluripotency and self-renewal.

INTRODUCTION

Mechanisms that control self-renewal and the ways of manipulating them in predictive manner are among the key current problems in regenerative biology. Addressing these issues is necessary to realize the biomedical potential of both embryonic stem cells (ESCs) and induced pluripotent stem cells (iPSCs).

The stabilization of mouse ESCs (mESCs) is achieved by supplementing the culture medium with the cytokine leukemia inhibitory factor (LIF), which activates pluripotency genes via the Jak-Stat pathway and relieves their potential repression by Gsk3 (Niwa et al., 2009). LIF-mediated external signaling becomes dispensable in mESCs when the major mediators of signaling pathways ERK1/2 and GSK3 are inhibited by small molecules (“2i” conditions) (Martello et al., 2012). Under 2i conditions (Makows et al., 2012), the stability of the pluripotent state is not associated with extrinsic signals such as LIF, suggesting the presence of a self-sustained mechanism of pluripotency supported by an intrinsic pluripotency gene regulatory network (PGRN).

Molecular ESC analyses and the discovery of iPSC reprogramming attributed pluripotency and self-renewal functions to the transcriptional regulators OCT4, SOX2, NANOG, and others (Ivanova et al., 2006; Loh et al., 2006; Takahashi and Yamanaka, 2006). Since then, numerous PGRN versions have been proposed (Festuccia et al., 2013; Yang et al., 2010), and the list of factors continues to grow. However, the hierarchy of the PGRN, the order of regulatory links, and the principles of PGRN function remain largely elusive.

In our studies (see flowchart in Figure S1), hundreds of single mESCs grown either under serum + LIF or serum-free 2i + LIF conditions have been collected, and their expression signatures with respect to 46 pluripotency genes were retrieved using high-throughput microfluidic single-cell qRT-PCR (White et al., 2011). Clustering individual cells based on their gene expression profiles revealed the presence of two major cell subpopulations in cells grown under the serum + LIF condition. In contrast, under 2i conditions, the two populations collapsed into one, which is in agreement with recent data suggesting a reduction in gene expression heterogeneity in 2i versus LIF alone (Marks et al., 2012). Comparison of our single-cell data with published single-cell data (Kumar et al., 2014; Tang et al., 2010) established that one subpopulation detected under the LIF condition has a gene expression signature similar to that of the inner cell mass (ICM) (Boroviak et al., 2014), whereas the other subpopulation resembles
more mature epiblast cells from the embryo. Detection of subpopulations became possible here because of the large number of analyzed cells (96 cells on each chip, seven chips in total).

We integrated the single-cell data obtained in this study with the data available for knockdowns of major pluripotency transcription factors (Feng et al., 2009; Ivanova et al., 2006; Loh et al., 2006; Lu et al., 2009; Martello et al., 2012). PGRNs reconstructed based on the integrated data revealed network motifs such as incoherent feedforward loops (iFFL) (Goentoro et al., 2009; Milo et al., 2002; Papatsenko and Levine, 2011), linking OCT4 and NANOG with their target genes and suggesting an antagonistic interaction between OCT4 and NANOG. Certain genes alternatively regulated by OCT4 and NANOG (Sall4 and Zscan10) appear to feed back to Oct4 and Sox2. We discuss how these loops may stabilize OCT4 concentrations required for self-renewal.

RESULTS

ESCs Grown in Serum + LIF Contain Two Cell Subpopulations

Potential medical applications involving ESCs or iPSCs require reproducible behavior of the cultured cells. From this perspective, the presence of cell subpopulations and phenotypic variations in stem cell cultures represent obstacles. Previously, subpopulations have been identified in mESCs grown under LIF conditions based on the expression of NANOG, REX1, and other pluripotency factors (Toyooka et al., 2008). Recent technologies allow the assessment of many genes in statistically sound numbers of cells (96 × 96 Fluidigm chips) (White et al., 2011), pushing the limits of detection of phenotypic variations and subpopulations.

Here single cell gene expression data were collected for 48 genes in 480 individual mESCs grown under serum + LIF conditions and in 192 individual cells grown under 2i + LIF conditions. The list of 48 genes included controls, core pluripotency transcription factors (Oct4, Sox2, Nanog, etc.), epigenetic regulators (Ctcf, Jarid2, etc.), and differentiation markers (Gata6, Pax6, etc.). The collected single-cell data were normalized using a relative normalization method (Experimental Procedures; Figures S2A–S2G; Table S1).

To establish the subpopulation structure, the single-cell gene expression data were clustered based on gene expression. Figure S2H shows hierarchical clustering, and Figure 1 shows co-expression clustering and principal component analysis (PCA) (Experimental Procedures) for serum + LIF cells. Hierarchical as well as co-expression clustering revealed two major cell clusters in serum + LIF cells, corresponding to two major mESC subpopulations (Figures 1A–1F; Figure S2H). Each cluster is characterized by a specific expression signature (Figure 1G). Cluster 1 cells occupy positions of ICM cells from published data (Tang et al., 2010; Figures 1H and 1I). Clustering data for 2i + LIF cells using the same set of methods showed that these cells occupy a position different from the two major LIF clusters (Figures 1B, 1D, 1F, 1J, and 1K). The detected variation of gene expression levels (Figures S3A–S3AB) was in agreement with previously reported reduced gene expression variations under 2i + LIF conditions (Wray et al., 2010). The fuzziness of the identified clusters may be attributed to the precision of the detection method and/or to intrinsic stochastic variation in expression levels not related to the deterministic expression signatures identified here (two signatures in serum + LIF and one in 2i + LIF).

Identities of Subpopulations

To characterize cell identities in the two mESC subpopulations, the corresponding cell clusters were compared with published single-cell data obtained from the ICM, epiblast cells, and ESCs (Boroviak et al., 2014; Tang et al., 2010). These analyses have shown (Figures 1H and 1I and 2) that cells in cluster 1 (~50% of cells grown in serum + LIF) have expression signatures similar to ICM from embryonic day 3.5 (E3.5) early blastocysts or even E2.5 morulae (Figures 2A–2C), whereas cluster 2 cells may be similar to more mature epiblast E5.5 cells. The previous view that mESCs have the identity of preimplantation epiblast cells (Plusa and Hadjantonakis, 2014) is not in contradiction with the current findings because the averaged expression signature for the two clusters (E2.5 and E5.5) may well resemble preimplantation epiblast cells (E4.0–E4.5).

To further characterize the cell identities in clusters 1 and 2, temporal expression profiles for genes explored in this study were retrieved from available ESC differentiation data (Hailesellasse Sene et al., 2007; Lu et al., 2009). Figure 2D shows that a majority of genes highly expressed in cluster 1 have peak expression in undifferentiated mESCs (day 0). In contrast, cluster 2 genes have peak expression at the onset of differentiation (12–24 hr, Figure 2E).

Together, our analyses suggest that serum + LIF cells contain two subpopulations or major cell types: cluster 1, matching E2.5/E3.5 ICM cells, and cluster 2, matching E5.5 epiblast cells. Cluster 2 cells also express higher levels of genes associated with the onset of differentiation (Jarid2, Ctcf, Zscan10, and Lin28a). Cells in both clusters express similar levels of Oct4 and similar levels of differentiation markers (Figure S3D).

Pathways and Factors Stabilizing mESC States

Although the presence of an E2.5 morula/E3.5 ICM state (cluster 1) in ESCs is expected (ICM is the source of ESCs),
the presence and origin of the second subpopulation (cluster 2) is more obscure. To explore both states on a genome-wide level, differentiation time series datasets (Hailesellasse Sene et al., 2007; Lu et al., 2009) were searched for genes with a peak of expression in undifferentiated ESCs (0 hr), matching cluster 1 genes, and with expression peaks at the onset of differentiation (12–24 hr), matching genes in cluster 2. The results were validated computationally using published single-cell RNA sequencing (RNA-seq) data (Tang et al., 2010; Figure 3A). Five ICM-derived cells were selected, displaying the highest similarity to cells in cluster 1, and five ESC cells were selected, displaying the highest similarity with cells in cluster 2 (Figures 3B and 3C). Strikingly, 37 of 53 genes (69.8% of genes matching
cluster 1, 0 hr) and 135 of 189 genes (71.4% of genes matching cluster 2, 12–24 hr) were evaluated as true positives based on RNA-seq data (Figure 3D; Tables S2 and S3).

Among the genes predicted to be highly expressed in cluster 1 are self-renewal factors such as \textit{Klf5}, \textit{Prdm14}, and \textit{Tet2} (Table S3). Ectopic expression of KLF5 is sufficient to maintain ESCs in the absence of LIF (Parisi and Russo, 2011), and \textit{PRDM14} blocks differentiation toward extraembryonic endodermal fates (Ma et al., 2011). Genes predicted to be highly expressed in cluster 2 include \textit{p53}, \textit{Pml}, \textit{Tcea3}, \textit{Nodal}, and others. \textit{P53} may interfere the WNT signaling pathway and inhibit differentiation (Abdelalim and Tooyama, 2014). overexpression of \textit{Tcea3} in the presence of differentiation signals blocks in vitro differentiation. \textit{TCEA3} appears to work via the LEFTY1/NODAL pathway (Park et al., 2013), and all three genes (\textit{Tcea3}, \textit{Lefty1}, and \textit{Nodal}) were evaluated as true positives based on RNA-seq data (Figure 3D; Tables S2 and S3).
Nodal) were among the 135 genes predicted to be highly expressed in cluster 2. NODAL is essential for human ESCs (hESCs), which, phenotypically, are more mature than mESCs (James et al., 2005), suggesting a link between hESCs and the mature state (cluster 2) detected in mESCs in this study.

OCT4-NANOG Antagonism and Feedback Control of Pluripotency

Identification of cell clusters 1 and 2 under serum + LIF conditions suggested the presence of at least two groups of coordinately acting gene products. Co-expression clustering of the single-cell data with respect to genes supported this view (Figure 5A). Co-expression clustering and Bayesian network inference (Figures 4A–4C; Figures S4B–S4F) revealed that gene interactions within the gene clusters were mainly positive, whereas gene interactions between the gene clusters were mainly negative (Figure 4C).

To reconstruct a signed and hierarchical PGRN, additional expression data from published knockdown studies targeting Oct4, Nanog, Sox2, and Esrrb were taken into consideration (Feng et al., 2009; Ivanova et al., 2006; Loh et al., 2006). Data from single-cell co-expression analyses, Bayesian inference based on the same studies, and published RNAi knockdown studies were integrated (Table S4). The resulting PGRN was searched for network motifs connecting up to three nodes and comprising iFFLs (Milo et al., 2002). The information-processing potential of iFFLs is of great interest because these motifs combine both positive and negative regulatory links, creating opportunities for alternative output solutions. This search returned 41 network motifs that were integrated into the network shown in Figure 4A. Several iFFLs integrated OCT4 and
NANOG and their shared target genes. Strikingly, in many such motifs, OCT4 acts as an activator and NANOG as a repressor (Figure 4B), suggesting a high significance for OCT4-NANOG antagonism. A number of shared OCT4-NANOG target genes were found in gene cluster 2 (Figure 4C). A few of these (Sall4 and Zscan10) have been suggested as Oct4 regulators (Tan et al., 2013; Yang et al., 2010; Yu et al., 2009), providing a feedback-based control system to the PGRN (Figure 4D).

In Vivo Binding Data Suggest the Existence of an OCT4-SOX2/NANOG Composite Element

To identify a potential link between OCT4 and NANOG, available in vivo binding data (chromatin immunoprecipitation [ChIP]) for OCT4, SOX2, and NANOG and their binding motifs were retrieved and analyzed (Figure S5). Comparison of all published binding motifs revealed a secondary motif component present adjacent to the core of the published OCT4, NANOG, and SOX2 motifs (Figures 5A–5F). In the case of the OCT4 and SOX2 motifs, the secondary motif indicates the presence of a well-known OCT4-SOX2 element (Yuan et al., 1995). Most of the published NANOG motifs performed very poorly in the computational validation test (Figure 5D), with the exception of two motifs (shown in the figure as CHEN2008 and ChIPMunk) that contained a secondary motif component resembling the consensus OCT4 binding site.

De novo motif reconstruction based on ChIP data from multiple sources (ChIPMunk; Table S5; Kulakovskiy et al., 2013a) produced a similar bipartite consensus sequence for all three binding motifs, suggesting the presence of a composite OCT4-SOX2/NANOG element. In this element, SOX2 and Nanog binding sites overlap (Figure 5D), suggesting competition between Nanog and the OCT4-SOX2 complex. The competition may prevent activation of the target genes by OCT4 and appear as their repression by Nanog (Figure 5G). Distribution of the composite element in the Oct4, Nanog, Sall4, and Zscan10 loci is shown in Figures S6A–S6D. Distributions of the OCT4-SOX2/NANOG element and ChIP-seq peaks were analyzed in the loci of genes highly

Figure 4. Regulatory Gene Network and Feedback Control of Pluripotency

(A) Part of the hierarchical gene network, including all identified iFFLs.

(B) Connectivity pattern emerging from iFFL analyses. Oct4 activates genes repressed by Nanog.

(C) Antagonistic relationships between gene clusters 1 and 2. Edge directions were predicted using Bayesian inference (Figure S4). Signs (activation/repression) were assigned based on coexpression clustering.

(D) Generalized representation of the network shown in (C) with feedback control of Oct4 by cluster 2 factors such as SALL4.

(E) Gene expression signature (as in Figure 1G) for the genes shown in (C) and (D).

(F) Differentiation expression profiles of the genes shown in (C) and (D) (integrated data).
A significantly larger number (p < 0.05, Fisher’s exact test) of OCT4-SOX2/NANOG elements were found in cluster 2 genes (Table S6), supporting their alternative regulation by OCT4 and NANOG as established based on expression studies (Figures 4A and 4B).

Identified Network Motifs Suggest a Mechanism for Stabilization of OCT4 Concentration

The identified iFFL feedback (iFFL-FB) motifs were explored using quantitative models based on transcriptional interactions (Papatsenko and Levine, 2011). iFFLs without feedback are essential for threshold responses of target genes (Goentoro et al., 2009 Figure 6A). The addition of a positive feedback from the target gene to the upstream regulator creates a dynamic system in which the prevalent non-zero solution is monostability (Figure 6B). Specific properties of the iFFL-FB motif include self-compensation, when proportional scaling of regulatory parameters (Figure 6C; Supplemental Experimental Procedures) has a small effect on the steady-state concentration of the upstream regulator (Figure 6C). Indeed, proportional increases in both the activation and the repression strengths in iFFL-FB (orders of magnitude) compensate each other, and the steady-state concentration of OCT4 remains nearly the same (Figure 6C).

Changing environmental and intrinsic conditions (temperature, stress, starvation, and epigenetic changes) may affect global system parameters such as the rate of protein synthesis. However, in the context of the iFFL-FB model, the concentration of Oct4 will tolerate changing global conditions. PCA of OCT4 and SALL4 expression levels in single cells revealed a bimodal dependence between the factors, in which OCT4 and SALL4 levels are correlated positively over a broad range of OCT4 concentrations but correlated negatively at high OCT4 concentrations (Figures 6D and 6E). This observed bimodal dependence between OCT4 and SALL4 is consistent with the model predictions (compare Figures 6F and 6G).
Modifications to the original iFFL-FB model have been considered, including high cooperativity models (Figure S7A), NANOG self-repression models (Figure S7B), and OCT4 dual regulation models (Figure S7C). The OCT4 dual regulation models (Marikawa et al., 2011; Papatserko and Levine, 2008) produce very robust bistable solutions, sustaining 4-fold proportional scaling of regulatory parameters (Figures S7C–S7L, see also Movie S1).

**DISCUSSION**

Major Cell States Defining ESCs and Self-Renewal

Analyses of single-cell gene expression data in mESC populations have demonstrated the presence of two major subpopulations or states in cells grown in the presence of serum + LIF (LIF) and a single subpopulation in cells grown under serum-free 2i + LIF conditions (2i) (Figure 1). Cell
identities in the two LIF subpopulations were established using published data (Halleselassie Sene et al., 2007; Lu et al., 2009; Tang et al., 2010). Cells in one subpopulation matched, by their expression profiles, cells from the E2.5 morula/E3.5 ICM, and cells in the other subpopulation matched epiblast cells (E5.5; Figure 2).

Earlier studies of NANOG-GFP-expressing mESC lines have demonstrated the restoration of a subpopulation structure from isolated GFP high and low fractions, suggesting a dynamic equilibrium between the two cell states (Chambers et al., 2007). A possible reverse exchange between alternative ESC states has been also suggested based on the expression of differentiation markers (Canham et al., 2010; Morgani et al., 2013). Taken together, these observations suggest a view of LIF-supported self-renewal as “stalled” differentiation (Figure 7). If the progress of normal differentiation (as observed in the embryo) is blocked or stalled at the onset of differentiation, then the cell identities “revert” back to an ICM-like state.

Several genes were found outside of the identified clusters (states). For example, OCT4 and SOX2 have similar expression levels in both states, potentially suggesting their general maintenance functions in self-renewal. The expression of differentiation markers was also similar (Figure S3D), suggesting that both LIF states are pluripotent. Conversely, STAT3 and TCF3 were highly expressed in cells occupying an intermediate or transitional position between the two major LIF states (Figure 2). Both of these transcription factors mediate signaling pathways, emphasizing the role of signaling in cell fate transitions between the two LIF subpopulations.

**Molecular Signatures of ICM-like and Transient Pluripotent States**

The presence of an ICM-like state in ESCs grown under LIF conditions is expected given the origin of ESCs. Our current studies show that the major determinants of the ICM state are TBX3, KLF4, NANOG, ESRRB, and other factors associated with pluripotency (Figure 1; Figure S3). TBX3 shows the greatest differences, with much higher levels in the ICM-like state. Both Tbx3 and KLF4 are activated by LIF, suggesting that LIF may be the reason for the formation of the two ES subpopulations. Interestingly, pulsed expression of KLF2 and NANOG promotes conversion of epiblast-like hESCs into a more naive ground state, which may be analogous to the ICM-like mESC state (Takashima et al., 2014). Epigenetic modifiers promoting pluripotency, such as WDR5 (Ang et al., 2011) and others, were also highly expressed in the ICM-like state (Table S3).

The properties of the transient LIF state (E5.5) may provide a clue regarding the nature of self-renewal. Addition of LIF to ESCs promotes JAK-STAT3 and other pathways, which, in turn, activate transcription factors such as TBX3, KLF4, TCFCP2II, and others (Chen et al., 2008).
This process may slow down the natural progression of the ICM-like state toward differentiation and promote the formation of the transient or stalled pluripotent state. The transient state is characterized by increased expression of many epigenetic factors, such as MBD3, JARID2, or CTCF. Increased levels of these molecules suggest that the transient state is a turning point toward differentiation in which the cells are inherently ready for silencing their genomes and losing pluripotency. However, LIF or 2i conditions force the expression of the core pluripotency factors. Therefore, overriding the natural progression and “rolling back” the cell fates to the ICM-like state. Genome-wide predictions (Figure 3) produced a longer list of genes highly expressed in the transient pluripotent state. Some of these, such as the TCEA3/LEFTY1/NODAL group, are involved in blocking differentiation rather than promoting pluripotency.

The above cases suggest the existence of molecular mechanisms or checkpoints demarcating a transition toward differentiation. The transient state, with its high expression levels of differentiation-blocking genes, may be one such checkpoint. What controls “crossing the border” events, and how do the switches work? Reconstruction of gene networks has pointed to feedback control mechanisms involving downstream OCT4 targets such as Sall4 or Zscan10, which are also highly expressed in the transient pluripotent state.

**SALL4 May Be among the Key Self-Renewal Factors**

Our network reconstructions (Figure 4) revealed an interesting connectivity pattern in which OCT4 and NANOG alternatively regulate genes highly expressed in the transient state (cluster 2). The literature suggests that at least two genes from this group, Sall4 and Zscan10, are not only regulated by the core factors but also provide feedback control to the system by regulating Oct4 (Yang et al., 2010; Yu et al., 2009; Figure S6). Feedback control in dynamic systems is required to achieve equilibria or steady states under changing (initial) conditions.

Several recent studies suggest that the OCT4-NANOG-SALL4 network regulates pluripotency via a feedback control mechanism (Tan et al., 2013; Yang et al., 2010). Systematic knockdown studies of 100 transcription factors (KD-100) in mESCs place SALL4 in fourth position (after ESRRB, OCT4, and SOX2) based on the number of downstream targets (Nishiyama et al., 2013). Known SALL4 functions include blocking trophodermal commitment (Oron and Ivanova, 2012), regulating extraembryonic endodermal genes (Lim et al., 2008), stimulating human hematopoietic stem cells (Yang et al., 2011), and many others.

Our current analyses of in vivo binding data for core pluripotency factors suggest the presence of a composite DNA element combining OCT4/SOX2 and NANOG binding sites (Figure 5). Both Sall4 and Zscan10 loci contain this element, suggesting the direct alternative regulation of these two genes by Oct4 and Nanog, which is also supported by expression data. The composite element includes the known OCT4/SOX2 element, in which the part of the SOX2 site overlaps with the Nanog site. Binding of NANOG to the composite element may interfere with binding of SOX2, therefore disrupting OCT4-SOX2 synergy, which may appear as direct repression by Nanog. Quantitative modeling of the network motif (iFFL-FB) loop combining Oct4, Sall4, and Nanog demonstrates how the interplay of these factors may stabilize OCT4 concentration and, therefore, safeguard pluripotency.

**iFFL-FB: Self-Compensatory Network Motifs that Stabilize Pluripotency**

The identified iFFL-FB motifs and their variants were explored using quantitative models (Figure 6). One of the most frequent solutions for such motifs is monostability. For certain modifications of iFFL-FB motifs, bistable solutions are frequent but not robust to parameter variations. Instead, monostable solutions display a remarkable robustness: the equilibrium concentration of OCT4 in these solutions sustained proportional scaling of regulatory parameters over orders of magnitude. Indeed, proportionally increasing both activation and repression rates in the context of the iFFL-FB compensates for the outcome and maintains a stable equilibrium concentration of OCT4.

In terms of biology, the identified self-compensatory feature of the iFFL-FB motif suggests that the equilibrium concentration of OCT4 may be resistant to global changes in cell metabolic and biochemical reaction rates caused by changing temperature, stress, starvation, as well as global chromatin configuration. The models described in this study (Figures 6B and 6C) predict that, under such changing conditions, the concentration of OCT4 would remain constant, therefore safeguarding or buffering pluripotency. Interestingly, the major impact of global parametric changes appears to be “absorbed” by SALL4. Therefore, SALL4 acts as a buffering component of the system, whereas OCT4 is the buffered component. NANOG, in this configuration, mediates the negative feedback in the context of OCT4-NANOG-SALL4 iFFL-FB. The measured distribution of OCT4 and SALL4 expression levels in single cells is consistent with the proposed model, assuming changing metabolic rates during mESC transitions from one state to another (Figures 6D–6G).

Although iFFL-FB models predict monostable solutions, cells grown under LIF conditions form at least two subpopulations. Is there a contradiction between the data and the model? Apparently, LIF media may be considered an “enforced” condition where the stability of the pluripotent...
states and self-renewal are maintained artificially. Instead, the 2i condition, producing a single subpopulation of cells, may represent a more natural, self-sustainable PGRN steady state in which most of the signaling is bypassed by the chemical inhibitors. In this case, a single predicted state is in agreement with the single detected cell subpopulation. Under enforced LIF conditions (two subpopulations), two iFFL-FB loops may be involved in stabilizing either the ICM-like or transient state or, possibly, both. Moreover, switching between states may be explained not only by bistable solutions (Figure S7C), but may be the consequence of more sophisticated processes in which the regulatory parameters undergo changes and generate new dynamic solutions (bifurcation).

In summary, we identified and characterized two cell subpopulations present in mESC, proposed a molecular mechanism explaining the antagonistic actions of Oct4 and Nanog in the pluripotency gene regulatory network, developed quantitative models explaining the stabilization of Oct4 concentration as well as pluripotency states, and proposed a “state exchange model” to explain self-renewal.

EXPERIMENTAL PROCEDURES

mESC Culture and Data Collection
CCE mESCs were cultured on gelatin-coated plates in high-glucose DMEM supplemented with 15% fetal bovine serum, 100 mM non-essential amino acids, 1 mM L-glutamine, 0.1 mM β-mercaptoethanol, 100 U/ml penicillin/streptomycin, 1 mM sodium pyruvate, and 1000 U/ml LIF. When transferring to 2i serum-free conditions, cells were seeded on gelatin-coated plates in 2i medium composed of a 50/50 mixture of Neurobasal and DMEM/F12 media, N2 and B27 supplements, 0.05% BSA, 100 U/ml penicillin/streptomycin, 1 mM L-glutamine, 1000 U/ml LIF, 3 μM CHIR99021, and 1 μM PD0325901. For single-cell gene expression analyses, 4’,6-diamidino-2-phenylindole (DAPI)-stained live cells were sorted into 96-well PCR plates containing pre-amplification mix, Taqman probes, Superscript III RT/Platinum Taq mix, and CellsDirect reaction mix (Invitrogen). Reverse transcription, pre-amplification, and amplification were performed as follows: 50°C for 15 min and 95°C for 2 min, followed by 21 cycles of 95°C for 15 s and 60°C for 4 min. A well containing no sorted cells served as a negative control, and a well with 100 cells served as a positive control for the population state. RT and pre-amplification products were diluted and run on a Biomark Fluidigm machine following the manufacturer’s instructions using 96.96 dynamic arrays. Each probe set was run in duplicate.

Single-Cell Data Filtering and Normalization
Single-cell gene expression data were first filtered to remove cells with multiple failed probes. The threshold (<60% of failed measurements) was established based on the dependence of marker concentration on the fraction of failed cells (Figure S2E). This procedure returned a total of 552 cells for seven Fluidigm chips, including 375 cells for LIF (5 × 96 cells initially) and 177 cells for 2i (2 × 96 cells initially) conditions. The single-cell data were processed using relative normalization. First, data from each individual cell were normalized using the linear zero-mean method, and technical replicates were averaged. Next the data for each gene from each chip were normalized in the same way, therefore eliminating chip-to-chip variations (Figure S2). Finally the normalized data were integrated for multiple chips (either for five LIF chips or for two 2i chips) and ranked. The ranks were converted to a 0–100 range, with failed readings assigned a value of 0. All published single-cell data used in the comparisons were rank-normalized using the same procedure to match the data format. Comparisons between the raw cycle threshold (CT) values and the normalized data are given in Figures S2F and S2G. Raw and normalized single-cell data produced in this study are available in Table S1.

Clustering Cell Gene Expression Profiles
Clustering of data with respect to cell expression profiles was performed using three different methods: hierarchical clustering (de Hoon et al., 2004), co-expression clustering (Stuart et al., 2003), and PCA (de Hoon et al., 2004). In the case of co-expression clustering, failed measurements were ignored, and the corresponding probabilistic similarity matrix was constructed by calculating correlation p values given the number of non-failed readings in all pairwise comparisons. The p values were corrected using the Benjamini-Hochberg technique to establish an appropriate similarity cutoff (p < 0.01 was adopted). Visualization of the resulting co-expression clusters was performed with force-directed layout using the Cytoscape program (Figures 1A–1D; Smoot et al., 2011).

Gene Network Reconstruction
Gene networks were reconstructed using the following methods: co-expression clustering (Stuart et al., 2003) of single-cell data using an approach described for cells but applied to genes (clustering columns versus rows), Bayesian network inference using Banjo software (Hartemink, 2005), mutual information analysis to detect non-linear interactions between genes, and reconstruction based on genome-wide knockdown studies compiled from multiple sources (Martello et al., 2012) (Supplemental Experimental Procedures; Table S4).

Analysis of Published ChIP Datasets
Available ChIP data and existing binding motif models (positional weight matrices [PWMs]) for OCT4, SOX2, and NANOG were collected from available sources (Chen et al., 2008; Heinz et al., 2010; Kulakovskiy et al., 2013b; Loh et al., 2006; Portales-Casamar et al., 2010; Figure SS; Table SS). The ChIPMunk for classic PWMs (Weirauch et al., 2013) and diChIPMunk (Kulakovskiy et al., 2013a; Levitsky et al., 2014) programs were used for de novo motif discovery. Raw published ChIP data were processed, and receiver operating characteristic (ROC) curves were constructed as described earlier (Kulakovskiy et al., 2013a). Sequences for motif discovery were extracted with an offset of ±150 bp with respect to the peak summits. Transcription factor binding site (TFBS) predictions were produced using diChIPMunk dinucleotide PWMs for the OCT4/SOX2/NANOG subtypes of the SOX2/OCT4 composite element.
Quantitative Modeling

Quantitative models for network motifs were constructed using ordinary and stochastic differential equations describing interactions between genes in transcriptional networks (Supplemental Experimental Procedures). Model solutions were analyzed using the NetExplore web server (http://line.bioinfolab.net/nex/netexplore.htm) (Papatsenko and Lemischka, 2015). At least 10^4 solutions were analyzed. The explored additive and cooperative models have four open edge-specific parameters (each specific to one of the four regulatory links) and three global parameters (common to all nodes). For all node/edge absolute concentrations, cooperativity levels and synthesis rates were identical. The NANOG self-repression model has five open parameters, and the OCT4 dual regulation model has six edge-specific open parameters plus three node-specific parameters for cooperative interactions.

SUPPLEMENTAL INFORMATION

Supplemental Information includes Supplemental Experimental Procedures, seven figures, six tables and one movie and can be found with this article online at http://dx.doi.org/10.1016/j.stemcr.2015.07.004.

AUTHOR CONTRIBUTIONS

I.R.L., D.P., and H.D. conceived the study. H.D. collected and analyzed single-cell data; D.P. implemented the quantitative models; I.V.K. and V.J.M. carried out the integrative analysis of ChIP data; A.W. and B.D.M. consulted on data analysis; and D.P., H.D., and I.R.L. wrote the manuscript.

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