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Article

Blockage of the Epithelial-to-Mesenchymal Transition Is Required for Embryonic Stem Cell Derivation

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SUMMARY

Pluripotent cells emanate from the inner cell mass (ICM) of the blastocyst and when cultivated under optimal conditions immortalize as embryonic stem cells (ESCs). The fundamental mechanism underlying ESC derivation has, however, remained elusive. Recently, we have devised a highly efficient approach for establishing ESCs, through inhibition of the MEK and TGF-β pathways. This regimen provides a platform for dissecting the molecular mechanism of ESC derivation. Via temporal gene expression analysis, we reveal key genes involved in the ICM to ESC transition. We found that DNA methyltransferases play a pivotal role in efficient ESC generation. We further observed a tight correlation between ESCs and preimplantation epiblast cell-related genes and noticed that fundamental events such as epithelial-to-mesenchymal transition blockage play a key role in launching the ESC self-renewal program. Our study provides a time course transcriptional resource highlighting the dynamics of the gene regulatory network during the ICM to ESC transition.

INTRODUCTION

Pluripotency is initiated in cells of the inner cell mass (ICM) and lapses shortly after implantation, coincident with the rise in lineage commitment. In vitro culture of ICM permits the generation of stable self-renewing pluripotent embryonic stem cells (ESCs) (Evans and Kaufman, 1981). However, ESC derivation is largely dependent upon the culture conditions. Under conventional medium, containing fetal calf serum and either feeder cells or leukemia inhibitory factor (LIF), only embryos from 129/Sv strain can efficiently give rise to ESCs and most strains of mice are refractory to ESC generation (Brook and Gardner, 1997). It is known that the 129/Sv strain has intrinsically more preimplantation epiblast (preEpi) cells than primitive endoderm (PE) cells when compared with refractory strains such as C57BL/6 or CBA (Batlle-Morera et al., 2008). So, preventing the formation of PE cells by induction of embryonic diapause (Brook and Gardner, 1997) or use of chemical substances that inhibit Fgf4 signaling (Ying et al., 2008) led to the formation of preEpi cells with efficient capability to generate ESCs even in serum- and feeder-free culture conditions. Also, when pre-blastocyst embryos have been used for ESC derivation, it can be assumed that these embryonic stages develop mainly into preEpi cells that subsequently develop into ESCs (Nichols and Smith, 2011). But, as pluripotent preEpi cells do not exhibit self-renewability, per se, the mechanism underlying this in vivo to in vitro conversion remains controversial (Loh et al., 2015).

To address the mechanisms underlying ICM to ESC conversion in the conventional culture condition, single-cell RNA sequencing (RNA-seq) analysis showed dramatic transcriptional and epigenetic gene expression changes during ICM to ESC transition (Tang et al., 2010). These changes include the simultaneous downregulation of Prame6/5/6/7, Gata6, and Cdx2 and upregulation of Dnmt3a and Hdac5, reflecting the higher expression of epigenetic modifiers during this cell transition process (Tang et al., 2010). This study suggested that reprogramming is a major event in the cell fate conversion of ICM to ESCs. However, this mechanism may be overshadowed by the low efficiency of generating ESCs from refractory mouse strains under serum-based culture conditions, as well as the tendency of the generated cells to undergo differentiation and their propensity to heterogeneously express key naive pluripotency transcription factors (TFs) (Hassani et al., 2012). Recent single-cell transcriptome analyses of mouse early embryos have demonstrated a high similarity between...
Figure 1. High-Resolution Transcriptome Trajectories from ICM to ESCs

(A) Isolated ICM and whole-blastocyst outgrowths (BOs) on days 3, 5, 7, and 9, and ESCs (in passage 20).

(B) Unsupervised hierarchical clustering of time course gene expression profiles for ESC derivation. Three independent biological replicates (nearly 20–30 ICMs and BOs for each replicate) were used except for BO-3 with two biological replicates. Transcript levels are mean-centered log2 scale values.

(C) Principal coordinate analysis (PCoA) of time course gene expression profiles. The red arrow shows the trajectory of cell expression profiles during ESC derivation.

(D) The number of differentially expressed genes (DEGs) between each of the consecutive time points. Red and blue bars indicate up- and downregulated genes, respectively.

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preEpi cells and ESCs grown in 2i (chemical inhibitors of MEK and GSK3) (Boroviak et al., 2015). The 2i condition supports the ground state of pluripotency, and triggers the efficient derivation and long-term maintenance of transcriptionally homogeneous ESCs. It has thus been suggested that 2i-grown ESCs are the actual self-renewing counterpart of naive preEpi cells, and that all preEpi cells are probably capable of becoming ESCs under 2i treatment (Boroviak et al., 2015).

We have recently introduced another efficient method for establishing ESCs from different refractory and non-permissive mouse strains by using PD0325901 and SB431542, chemical inhibitors of MEK, and transforming growth factor β (TGF-β) signaling pathways, respectively, which we named R2i (Hassani et al., 2014b). We showed that in the serum-free condition, R2i could support the ground state of pluripotency. Also, as well as from blastocysts, R2i could support the highly efficient ESC derivation from a wider developmental window, from single blastomeres of two to eight cell-stage mouse embryos with an efficiency of more than 2-fold higher than the 2i condition (Hassani et al., 2014a). Moreover, when cells underwent long-term passaging, they exhibited more genomic stability with R2i support compared with 2i (Hassani et al., 2014b). These salient features of R2i prompted us to evaluate the molecular mechanism(s) that underlie ICM to ESC transition. By conducting a gene expression profile at serial time points during the transition from ICM to ESCs, we discovered key regulatory genes and cellular events that are involved in this process. Besides the expression of preEpi-related genes, we found that the upregulation of DNA methyltransferases and an epithelial-to-mesenchymal transition (EMT) blockage probably have major impacts on the efficient generation of ESCs. This study reveals mechanistic insight into how transient pluripotent cells from the preimplantation embryo can perpetuate as self-renewing ESCs.

RESULTS

Sampling Strategy for Gene Expression Profiling during the Transition from ICM to ESCs

To establish a gene expression signature throughout the course of ESC derivation, we initially collected samples from immunosurgically isolated ICMs from blastocysts at embryonic day (E) 3.5 and whole-blastocyst outgrowths (BOs) on days 3, 5, 7, and 9 post-plating, as well as from ESCs growing under R2i + LIF (R2i) conditions (Figure 1A). The mRNA extracted from these cells was pre-amplified and used for whole-genome analysis using the Illumina platform. Pairwise Pearson coefficient of the gene expression profiles of samples indicated a significant difference between the ICM cells and ESCs. Hierarchical clustering revealed three major clusters of ICMs, BOs, and ESCs (Figure 1B). Dimension reduction of gene expression data by principal coordinate analysis (PCoA) depicted a molecular trajectory that reflected the transitional cascades from ICM cells to ESCs (Figure 1C). Next, we examined the differentially expressed genes (DEGs) at the indicated time points. We found 2,849 DEGs that indicated alterations (1,647 up- and 1,202 downregulated) between ICM cells and BO-3, and 1,974 DEGs (799 up- and 1,175 downregulated) between BO-9 and ESCs (≥2-fold expression change, adjusted p ≤ 0.05) (Figure 1D). We performed an unsupervised time course of gene clustering followed by gene set analysis using the Enrichr web tool (http://amp.pharm.mssm.edu/Enrichr/; Figure S1A). We found evidence for pluripotency and the establishment of self-renewal in ESCs, as indicated by related functional annotation such as PluriNetWork and mitotic cell cycle in clusters IV and V, respectively. However, our data documented an alteration in gene expression between two major phases of the ICM to ESC transition. The first phase was detectable in the early stage of the process, between ICM cells and BO-3, whereas the second phase occurred in the late stage, between BO-9 and ESCs (Figures 1B and 1C).

We assumed that a 3-day gap between ICM cells and the first collected BOs almost hindered accurate interpretation of the extracted data. Furthermore, even though trophectoderm cells were not expanded (Figure 1A), the few such cells that existed probably resulted in gene alterations from BO-9 to ESCs. This result convinced us that a higher-resolution time course analysis from isolated ICM outgrowths (IOs) would give us better insight into the molecular cascades occurring during the cellular transition. Therefore, we collected immunosurgically isolated ICM cells and IOs on days 0.5, 1, 2, 3, and 5 post-plating.
A vs ICM

No. of DEGs

vs ICM

E2F5, Arhgap18, Ptsrd10, Akt3, Anapo11, Astb1, Wnt3a, Rab1, Rtn2, Mdm2, Stc8a1, Dnm1, Dnm1b, ...
Neu2, Neu3, Neu4, Arhgap18, Pdgfra, Rab31, Rtn2, ...

B

IO-0.5 vs ICM

| Rank | Functional annotation                  | p value |
|------|----------------------------------------|---------|
| 1    | GO terms: Mitotic cell cycle           | 5.76E-06|
| 2    | KEGG pathway: Small cell lung cancer  | 7.03E-03|
| 3    | GO terms: Membrane lipid catabolic process | 1.26E-04|
| 4    | KEGG pathway: Other glycan degradation | 2.98E-04|
| 5    | KEGG pathway: Metabolic pathways      | 8.20E-03|

| Rank (p value) |
|---------------|
| IO-1          |
| IO-2          |
| IO-3          |
| IO-5          |
| ESCs (P2)     |

C

ESC derivation efficiency %

D

No. of DEGs in IO vs different stages of IO

P2

P4

P15

E

P15 vs GO terms

| Rank (p value) |
|---------------|
| ICM: Mitotic cell cycle | 1(4.99E-17)|
| IO-0.5: regulation of cellular response to stress, DNA repair | 1(1.12E-06), 2(1.03E-06)|
| IO-1: DNA repair, mitotic cell cycle | 1(1.27E-07), 2(2.88E-11)|
| IO-2: organelle fission, nuclear division | 1(5.22E-17), 2(2.38E-11)|
| IO-3: positive regulation of macroautophagy | 1(4.74E-04)|
| IO-5: organelle fission, nuclear division | 1(5.76E-04), 2(2.09E-04)|

P15 vs GO terms

| Rank (p value) |
|---------------|
| ICM: Translation, gene expression | 1(2.77E-08), 2(4.55E-06)|
| IO-0.5: translation, gene expression | 1(4.37E-14), 2(4.64E-10)|
| IO-1: translation, gene expression | 1(2.25E-12), 2(8.34E-10)|
| IO-2: translation, gene expression | 1(1.27E-09), 2(3.69E-09)|
| IO-3: gene expression | 1(1.88E-05)|
| IO-5: regulation of epithelial cell migration | 1(3.63E-08)|

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(Figure 1E). In these analyses, we limited the period of study to day 5 due to a high similarity between the different time points of BO samples in the first run of the microarray. In addition, we used ESCs of different passage numbers (P2, P4, and P15) in our analyses. The negative control group for these experiments was the cultured IOs under SB + LIF (SB) conditions on days 1 and 3 post-plating (SBIO-1 and SBIO-3, respectively; Figure 1E). This control group was based on our previous observation that each component of R2i (PD or SB) in combination with LIF supports the long-term maintenance of ESCs. However, the efficiency of deriving ESCs under SB + LIF conditions was negligible or low, whereas it approximated 60% under PD + LIF (Hassani et al., 2014b). PCoA of the time course data depicted a parabolic trajectory of the ICM to ESC derivation process that bifurcated under SB conditions (Figure 1F). This analysis showed that the profile in SBIO-1 was close to the ICM to ESC trajectory, whereas SBIO-3 significantly differed from IO-3. We observed the highest number of DEGs in the sequential time points between IO-0.5 and ICM cells, at 2,704 altered genes (1,480 upregulated and 1,224 downregulated; adjusted p < 0.01; fold change >2; Figure 1G). This tangible gene alteration between ICM cells and IO-0.5 was reminiscent of the high number of DEGs between the ICM cells and BO-3 in the first run of our experiments. We observed that the alternations among consecutive samples are biologically relevant rather than a fluctuation among replicates (Figure S1B). Therefore, our study denoted a significant expression fluctuation from in vivo to in vitro, even in the short time period after this cultivation. However, the number of DEGs between late-stage outgrowths and ESCs of different passage numbers was dramatically dropped in the second set of experiments (Figure 1H). Overall, this new design for sampling provided a convenient platform for assessing the key events involved in the generation of ESCs from ICM cells.

A Gradual Molecular Transition from ICM to ESCs

Our first priority was to uncover the reason for the tremendous difference in the number of DEGs between the ICM cells and IO-0.5. Gene set analysis showed the reasonable enrichment score for genes involved in the mitotic cell cycle and membrane lipid catabolism for up- and downregulated genes, respectively (Figures 2A, 2B, and S2A). We sought to determine whether the gene expression alteration in the first interval could cause the establishment of ESCs, or only represented a stochastic fluctuation of gene expression during in vivo to in vitro transition. Therefore, we compared the expression profile of IOs from the other designated time points with the ICM cells (Figure 2A). This analysis revealed an upward trend in the number of DEGs for IOs with an increased time point interval and the ICM cells (Figure 2A). However, most of the altered genes revealed the same functional annotation identified in the comparison between ICM cells and IO-0.5, as well as pathways for small-cell lung cancer and relevant metabolism with regard to the up- and downregulated genes, respectively (Figures 2B and S2A). This representation of biological processes during the ICM to ESC transition appeared to be consistent with the acquisition of ESC self-renewal capability.

As the early evidence of self-renewal capability was found during ICM to IO-0.5 transition (Figures 2A and 2B), we sought to determine whether ESC identity was acquired quickly after ICM expansion in vitro. Therefore, we examined the temporal dependency on R2i culture conditions to determine the minimal time required for the efficient derivation of ESCs. We found that 1-day treatment with R2i led to the derivation of ESCs at approximately 60% efficiency, and that 2-day treatment induced ESCs at maximal derivation efficiency (Figure 2C). This finding revealed that the gene expression changes taking place during the first days of ICM culture supported the efficient establishment of ESCs. In the next step, we compared the expression profile of ESCs of different passages (P2, P4, and P15) with ICM cells and all the IOs (Figures 2D and 2E). The results showed that, with increasing passage number, ESCs are closer to late-stage IOs than to early-stage IOs (Figures 2D and 2E). Functional analysis of DEGs between IOs and ESCs at different passage numbers indicated the upregulation of nuclear division-related genes and the downregulation of genes involved in translation and gene expression processes. However, these differences declined as the interval between ESCs and IOs was reduced (Figures 2E and S2B–S2D). Nearly similar functional annotations were enriched by analyzing the differences in gene expression between ESCs of different passages (Figure S2E). Despite the high similarity between ESCs and late-stage

Figure 2. Transition from ICM to ESCs Is a Gradual Process

(A) The representation of DEGs between different stages of IOs and ESCs versus ICM.
(B) Functional annotation of up- and downregulated genes between IOs and ESCs of different stages versus ICM.
(C) Experimental schematic and the results of time course dependency of the derived ESCs under R2i culture conditions. For each experiment (rows), the red and green bars indicate the duration of time the cells were cultured in the negative control and R2i media, respectively. The efficiency of deriving ESCs is based on the number of Nanog-positive colonies derived from ten isolated ICM.
(D) The number of up- and downregulated genes between ESCs and R2i-treated IOs.
(E) Functional annotation of up- and downregulated genes between P15 and IOs of different stages.
IOs, functional analysis for downregulated genes highlighted the importance of pathways related to epithelial cell differentiation, organ formation, extracellular matrix organization, and regulation of epithelial cell migration (Figures S2B–S2D and 2E). This result suggested that preventing the differentiation or migration of epithelial cells should be considered for successfully establishing an immortal ESC line. Therefore, it appeared that, although key changes in gene expression occurred quite quickly during the conversion of ICMs to ESCs, cells had assumed an ESC identity rather gradually. These findings suggested the existence of a distinct core regulatory circuitry for the establishment of ESCs. This circuitry would form and be active promptly during the in vitro ICM culture on one hand, but would induce cells to acquire ESC identity gradually on the other hand.

Identification of Transcriptional Signature in IOs

To identify the genes that play a key role in the ICM to ESC transition process, we categorized DEGs in both the R2i and control (SB) groups. We generated the gene expression profile of the SB group at two serial time points (SBIO-1 and SBIO-3) and subsequently compared their DEGs with the R2i group at the same days (IO-1 and IO-3). The outcome consisted of eight heatmaps (Figure 3A, Table S1). Heatmaps I–IV showed the list of genes that were downregulated in at least one of the aforementioned time points in the SB versus R2i groups, while heatmaps V–VIII showed the upregulated genes (Figure 3A). Pathway enrichment analysis indicated that most genes in heatmaps I–IV were associated with signaling pathways that regulated pluripotency, chromatin modification, and DNA metabolism. Specifically, heatmap III contained the largest number of DEGs, which revealed several pluripotency markers such as Esrrb, Sox2, Tbx3, and Rarg, and epigenetic modifiers including Wdr3, Suz12, Ezh2, Eed, Nrsu2, Dmnt3b, and Dmnt3l (Figures 3A, 3B, and S3A). Thereafter, we denoted the list of genes in heatmaps I–IV as R2i-specific and downregulated in SB groups, suggesting that these genes play an important role on the road to pluripotency. In contrast, heatmaps V–VIII (SB-specific gene list) highlighted mainly developmental pathways that included cell morphogenesis, cell-cell adhesion, and protein processing. In these gene lists, the greatest numbers of DEGs represented in heatmap VII contained some key upregulated factors such as Pitx2, Sna1, Itgb4, Sox17, Igf2, Fgf10, and Dab2, suggesting that the SB-treated IOs exited from the path of pluripotency (Figures 3A, 3B, and S3A).

Interestingly, our result depicted that the upregulation of some relevant pluripotency-related markers was observed in the SB-specific gene list, which included Stat3, Dusp9, and Id1, downstream of LIF and BMP4, respectively. This finding raised the question of whether TGF-β inhibition could sustain the expression of some pluripotency-related genes during the ICM to ESC transition. To address this question, we classified the expression profile of a comprehensive list of pluripotency-related genes (366 genes extracted from the PluriNetWork and the literature; Table S2) into common pluripotency, R2i-specific pluripotency, and SB-specific pluripotency groups (Figure 3C). A total of 233 well-known TFs that regulated pluripotency was put in the common pluripotency list, which included Pou5f1 (Oct4), Zfp42 (Rex1), Kfl4, Ctnmb1 (β-Catenin), and Sall4. In addition, 13 of 366 genes were in the SB-specific pluripotency list, which included Stat3, Id1, Dkk1, and Podxl. This analysis demonstrated that TGF-β inhibition, like fibroblast growth factor (FGF) signaling inhibition, plays a pivotal role in supporting the dynamic expression of pluripotency-related markers, while SB alone was insufficient for ESC derivation. It has been proposed that some of the pluripotency factors induce differentiation and play a central role in the determination of different cell states by orchestrating different gene expression profiles (Loh et al., 2015). Consistent with this notion, co-occupation of various sites with Smad2/3 and different master TFs has been found to direct different responses in ESCs, myoblasts, and B cells (Mullen et al., 2011). In contrast, R2i-specific pluripotency markers, such as Nanog, Klf2, Dppa3, Sox2, Esrrb, Utf1, and Nrsu2, have unveiled key regulators in the path to efficiently deriving ESCs from ICM. We showed that 5 (Esrrb, Gbx2, Sox2, Klf2, and Nanog) of 12 essential TFs for naive pluripotency (Dunn et al., 2014).
(legend on next page)
were in the R2i-specific pluripotency list, suggesting that they play key roles during ESC derivation (Figure S3B). To test this finding, we overexpressed Nanog in the ICM cells of E3.5 blastocysts obtained from Nanog-inducible mice in order to generate ESCs in N2B27 medium in the absence of SB or PD, resulting in 40% ESC derivation efficiency (Figure 3D). These data indicated that our strategy to classify the expression of pluripotency markers during the expansion of ICM could point to the key players involved in ESC derivation.

**Transition in DNA Methylation Level during the ICM to ESC Conversion**

As R2i-specific pluripotency markers significantly presented in day 3 (Figures 4A and 4B), we sought to classify the gene expression patterns by performing an unsupervised time course clustering on DEGs between SBIO-3 and IO-3 (Figure 4C). Clusters I–III represented the upregulated genes in SBIO-3 compared with IO-3, which exhibited different expression patterns in the R2i-treated group. Gene ontology analysis highlighted the increased activities of several biological processes that were associated mainly with tissue morphogenesis, metabolic pathways, and focal adhesion (Figure 4C). In contrast, clusters IV–VII represented downregulated genes in the SBIO-3. Gene enrichment analysis revealed that the genes in clusters IV–VII were associated with PluriNetWork, the DNA metabolic process, and chromatin modification. Clusters IV (80 genes) and V (86 genes) contained some key TFs for pluripotency (Tbx3, Esrrb, and Nanog), which were most likely expressed in a steady-state pattern from ICM cells to ESCs. Clusters VI (165 genes) and VII (435 genes) consisted of parabolic and gradually upregulated patterns of genes from ICM cells to ESCs, which included some epigenetic modifiers such as Dnmt3b, Dnmt3l, Chd8, Mts1, Suz12, Eed, Wibr3, and Mat2b (Figure 4C). To better understand the expression dynamics of epigenetic modifier genes during the derivation of ESCs, we conducted qRT-PCR analysis to quantify the expression of Dnmt1, Dnmt3b, Dnmt3l, Sirt1, Ezh2, Suz12, and Mat2b (Figure 4A). This experiment confirmed that the expression of Dnmt1, Dnmt3b, and their partners, was upregulated in this cell transition process. To examine the DNA methylation pattern, we measured the methylation level of the individual CpG sites for three classes of repetitive elements by deep hairpin-bisulfite sequencing. As approximately 40% of the genome consists of repetitive elements, we selected major Satellites (mSat), the 5′ UTR of L1Md_Tf (L1), and a class of LTR-retrotransposons (IAP-LTR1) (Arand et al., 2012). Our data showed that the DNA hyper-methylation during ESC derivation was due mainly to increased amounts of hemi-methylation at CpG positions (Figure 5B). We observed the highest amount of non-CpG methylation at mSat in R2i-treated IOs (Figure S4; Table S3). Non-CpG methylation depended on the presence of Dnmt1 mediated by Dnmt3a/3b in ESCs under serum conditions (Arand et al., 2012). This finding showed that early changes in DNA methylation probably occurred in tandem with an increased activity in de novo methyltransferases.

Next, we sought to determine the impact of increased global DNA methylation levels on the transition between the in vivo to in vitro pluripotent cell states. Inhibition of DNA methyltransferases by RG108 led to reduced efficiency of deriving ESCs under R2i conditions, but had no adverse effect under 2i conditions (Figure 5C). It had previously been reported that 2i could induce global hypomethylation (Ficz et al., 2013; Leitch et al., 2013); however, our results indicated the upregulation of de novo methyltransferases for the efficient establishment of ESCs in R2i. Unexpectedly, this finding implied that blockage of the ERK pathway, in parallel with TGF-β inhibition, led to a high expression of epigenetic modifiers and DNA methylation-related genes in the transition from ICM to ESCs. This analysis revealed a major difference between 2i- and R2i-grown ESCs, suggesting the existence of a different pluripotency circuitry of these two naive representative conditions.

**Blocking EMT Is Required for ESC Derivation**

Our gene expression profile analysis highlighted a difference between ESCs and IOs in epithelial differentiation and migration (Figures 2E and S2B–S2D). Interestingly, a gene set enrichment analysis between DEGs from R2i- and SB-treated IOs displayed an enrichment of EMT-related genes (Figure S5A). Thus, we sought to assess whether EMT might play a critical role in ESC derivation. To this end, the expression levels of genes involved in EMT and mesenchymal-to-epithelial transition (MET), such as Snail, Eomes, Dab2, Cdh1, and Klf4, as well as key pluripotency genes, were analyzed in R2i-treated IOs using qRT-PCR, and serum + LIF-treated IOs were used as a negative control (Figure 6A). Our results showed an upregulation of epithelial motif changes and downregulation of mesenchymal genes.
markers and a downregulation of mesenchymal markers in R2i-treated IOs compared with the negative control. Indeed, Snail, Eomes, and Dab2 exhibited higher expression levels in the presence of serum, correlating with a decrease in ESC derivation efficiency and an increase in differentiating IOs (Figure S5B). Next, we decided to ascertain whether blocking EMT is required for ESC derivation. To this end, we overexpressed Snail-2A-Tdtomato (EMT inducer) and Klf4-2A-Tdtomato (MET inducer) in R2i- and SB-treated IOs. In the presence of Snail, IOs started to differentiate and ESCs could not be derived under R2i conditions (Figures 6B and 6C). In contrast, 14 of 14 ESC lines were established from R2i-treated IOs overexpressing Tdtomato alone as a control. Conversely, Klf4 overexpression led to a 30% efficiency in ESC derivation in SB-treated IOs (Figures 6D, 6E, and S5C), whereas no ESC lines were derived in the Tdtomato control under the same conditions. Finally, we noticed that supplementation of 2i with TGF-β1 impeded the transition from ICM cells to ESCs (Figure 6F). Taken together, these findings demonstrate the crucial role of EMT blockage for the ICM to ESC transition. Contrary to what was expected, we showed that TGF-β1 inhibition induced EMT, while with the synergistic effect of MEK inhibition blocked EMT. Overall, our data

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**Figure 5. DNA Methylation Changes during ESC Derivation**

(A) Time course expression of epigenetic modifier genes during ESC derivation using qRT-PCR. Three biological replicates were used for this experiment. Error bars represent the SD. Here, * represents significant DEGs between ICM and ESCs, p ≤ 0.01. Statistical analysis was performed by one-way ANOVA and post hoc Tukey test.

(B) Hairpin-bisulfite amplicon sequencing of CpG dyads at retrotransposable elements (L1MdTf, IAP-LTR1, and mSat). The bars sum up the DNA methylation status of all CpG dyads. The map next to the bar represents the distribution of methylated sites. Each column shows neighboring CpG dyads and each line represents one sequence read. The reads in the map are sorted first by fully methylated sites, then by hemi-mCpG dyads. Red, fully methylated; light green and dark green, hemi-mCpG; blue, unmethylated CpG. Two biological replicates for IAP-LTR1 and mSat and one biological replicate for L1MdTf were used and * represents significant difference of DNA methylation between ICM, IOs, and ESCs, p ≤ 0.01. The numbers of reads and methylation state are shown in Table S3.

(C) Efficiency of ESC derivation upon inhibition of DNA methyltransferases by using RG108 in R2i- and 2i-treated IOs.
Figure 6. Blocking EMT Is Required for ESC Derivation

(A) Time course expression of EMT and MET gene markers was measured by qRT-PCR in R2i- and serum + LIF-treated IOs. All data were calibrated to ICM, which is considered 1. Error bars reflect SD of results derived from three biological replicates. Here, * represents significant differentially expressed genes in R2i- and serum + LIF-treated IOs as well as ESCs. *p ≤ 0.05, **p ≤ 0.01, error bar, ± SD.

(B) Overexpression of Snail-2A-tdTomato in OG2 ICM, which contain an Oct4-GFP reporter transgene, with R2i treatment. A construct with tdTomato alone was used as a negative control. Induction of EMT through the forced expression of Snail impaired ICM to ESC transition. tdTomato expression correlates with the transduction rate and expression of the gene of interest.

(C) Efficiency of ESC derivation upon Snail overexpression.

(D) Overexpression of Klf4-2A-tdTomato on E3.5 ICM (B6 × C3H) F1 in N2B27 supplemented with SB + LIF prevents EMT and supports the derivation of ESC lines.

(E) Efficiency of ESC derivation upon Klf4 overexpression on SB-treated IOs.

(F) The impact of TGF-β1 on the generation of ESCs was assessed in 2i-treated IOs after NANOG immunostaining. Blue, DAPI.
demonstrate that blocking EMT is an essential step in the ESC derivation process. When EMT blocking occurs, a TGF-β inhibitor (SB) combined with LIF could support the long-term maintenance of ESC self-renewal (Hassani et al., 2014b).

R2i Could Maintain preEpi Cell Identity in IOs

We also conducted a parallel transcriptional study on the development of ESCs in both in vitro and in vivo conditions. Recently, Boroviak et al. (2015) reported single-cell RNA-seq data for the lineage-specific profile of naïve pluripotency in early embryogenesis. However, those authors had adopted their protocol to small group of cells, instead of single cells, to be able to detect scarce transcripts, which provided a good substrate for comparison with our bulk transcriptome data. We observed a range of different early embryo lineage-feature gene expressions, from the ICM to post-implantation cells in R2i-treated IOs (Figure 7A; Table S4). Our study revealed a maximum similarity of different ICMs and early-passage ESCs (P2 and P4) to E4.5 preEpi cells. Late-passage ESCs (P15) showed slightly less similarity to the E4.5 preEpi cells (Figure 7B), while functional analysis of up- and downregulated genes in this comparison highlighted the similar pathways (Figure S6A) that were enriched between P15 ESCs and different ICMs (Figures 2E and S2D).

A comparison of R2i and SB treatment revealed that the number of ICM-related genes decreased in R2i- versus SB-treated IOs, while the number of gene markers related to preEpi and postEpi cells increased (Figure S6B). This analysis indicated the exclusive expression of diapaused-specific markers, such as Tcfap2c, Gbx2, Utf1, Sox2, Esrb, Thx3, Klf2, Nanog, Dppa3, and Msc, in R2i-specific clusters (Figures S6B and S6C). We also found that the expression of nearly one-third of the genes in heatmap II in Figure 3A were comparable with preEpi- and diapaused-related indicators, suggesting that the expression of these genes is important throughout the entire process of ESC derivation (Figure 7C).

We then compared our finding with a previously reported dynamic single-cell transcriptional dataset for identifying markers during the preEpi and PE bifurcation of the ICM (Gerovska and Arauzo-Bravo, 2016). We found that R2i-treated IOs noticeably represented the expression profile of E4.5 preEpi cells, while SB-treated IOs showed the different expressed genes such as Sox17, Sox7, and Gata6, which indicated an expression pattern of E4.5 PE (Figures S6D and S6E). In particular, one-third of genes in cluster VI, which were downregulated in R2i versus SB on both days 1 and 3, represented the PE transcriptome profile. A comparison of different staged ICMs and early-stage embryos revealed significantly expressed PGC markers, which included Ifitm3, Fragliis, Dppa3, Stella, Zp3, Ifitm1, and Ifitm2 (Boroviak et al., 2015; Gerovska and Arauzo-Bravo, 2016), as well as Tex11, Tex14, Piwil2, and Dppa1 (Figure 7D). This observation not only elucidated the preEpi cell origin of ESCs but also magnified the role of PGC-related genes during the ICM to ESC transition. Overall, this dataset depicted that ICM-derived ESCs proceeded through the preEpi stage, and that the successful derivation of ESCs relied on FGF inhibition. However, epigenetic modifications and EMT blockage were required for the perpetual self-renewal capability of preEpi cells.

DISCUSSION

An optimized regimen is required for dissecting the molecular mechanism of ESC derivation. Dual inhibition of FGF and TGF-β creates high genomic stability and an efficient ICM to ESC transition (Hassani et al., 2014b). We have employed this system to produce a temporal transcriptome profile. In addition, a negative control group (SB), which not only maintains self-renewal but also has similar components with R2i, helped us to develop an appropriate strategy for detecting regulatory genes that play a key role in ESC derivation.

This analysis has identified a group of significantly expressed TFs and epigenetic modifiers in the ICM, different ICMs, and ESCs. Our result was consistent with previously reported ICM-specific genes such as Pramel4, Pramel5, and Pramel7 (Tang et al., 2010). In a systemic data analysis, we observed approximately 3,000 DEGs between the ICM and day 0.5 (Figure 2A). Instead of a fluctuated gene expression stemming from the in vivo to in vitro transition, we observed that these large numbers of DEGs could establish the path toward the generation of ESCs. We demonstrated that only 2 days were needed for deriving ESCs at maximal efficiency under R2i conditions (Figure 2C). However, we have suggested that ICM cells do not suddenly assume an ESC identity; the longer the cultivation period, the more likely it is that the cells will acquire ESC characteristics. We observed that, while the number of DEGs decreased between the late-stage ICMs and established ESCs, ESCs showed downregulation of the EMT biological process compared with these ICMs (Figure 2E). Despite the expression of common pluripotency factors, SB-treated ICMs exited from pluripotency likely because they followed the EMT program. Our results are consistent with this notion, and have suggested that the ectopic expression of Snail, an EMT inducer, could impede the establishment of ESCs (Figure 6B). Normally, the corresponding protein presents quickly after the onset of gastrulation (Hemavathy et al., 1997). However, Snail surprisingly presents in SB-treated ICMs as a downstream protein of the TGF-β signaling pathway. This data has illustrated that inhibition of TGF-β without
Figure 7. R2i-Treated IOs Exhibited preEpi Cell Characteristics
(A) DEGs between in vivo embryonic samples (Boroviak et al., 2015) and ICM to ESC time course samples for this study (≥ 2-fold expression change, adjusted p ≤ 0.01).
(B) Volcano plots show fold change in gene expression (x axis, log2 fold change, adjusted p ≤ 0.01) between E4.5 preEpi cells (Boroviak et al., 2015), IO-5, and ESCs of different passages (this study). The y axis is the negative log10 of Benjamini-Hochberg adjusted p value. Blue, green, and red dots represent downregulated, non-significant, and upregulated genes, respectively.
(C) The comparability of the gene expression for some heatmap II-specific genes (from Figure 3) between in vitro IOs (this study) and single-preEpi cells (EPI1-3) or single-diapaused cells (DIA1-2; Boroviak et al., 2015).
(D) The expression of PGC-related genes in R2i- and SB-treated IOs.
suppression of FGF signaling could not block EMT and allow embryonic cells to exit from pluripotency. It has been reported that MEK suppression activates Klf4 through a phosphorylation pathway (Kim et al., 2012) and consequently induces MET (Chen et al., 2011). Accordingly, we observed that forced expression of Klf4 in SB-treated IOs led to the generation of ESCs (Figure 6D). We also detected the downregulation of epithelial markers (Cdh1 and Klf4) and the upregulation of mesenchymal markers (Snail, Eomes, and Dab2) under conventional conditions, i.e., serum + LIF (Figure 6A). Therefore, we concluded that the lack of EMT blockage was the main reason for the inefficient generation of ESCs from a refractory cell strain under serum or SB culture conditions. When EMT blockage occurs, SB or serum could support the long-term maintenance of ESC self-renewal.

We believe that the strategies developed in this study can help identify the most important gene regulators for deriving pluripotent ESCs. Based on significant changes in gene expression patterns and morphological changes between treatment (R2i) and the control (SB) groups on day 3 (Figure 4), we postulated that DEGs between these two groups was crucial for ICM to ESC transition. The data showed that pluripotency factors such as Pou5f1, Zfp42, Klf4, Sall4, Ctnmb1, and Klf6 were not significantly altered between the treatment and control groups up to the third day of ESC derivation. We classified these 233 pluripotency markers as common pluripotency factors. However, the pluripotency markers that were significantly upregulated in R2i compared with SB, including Nanog, Sox2, Klf4, Esrrb, Gbx2, Jam2, Dppa3, Msc, and Wif1c2, could be the genes driving the establishment of ESCs (Figure 3C).

Of all the TFs that play an essential role in maintaining pluripotency (Dunn et al., 2014), half were included in the R2i-specific gene list (Figure S3B). Therefore, we proposed that the overexpression of these genes, such as Nanog, would undermine the necessity of a special regimen, such as R2i and 2i (Figure 3D).

In this study, we also found a dramatic difference in the DNA methylation patterns between ICM cells and ESCs. The passive and active demethylation of maternal and paternal genomes resulted in a hypomethylated epigenetic landscape for ICM cells (Liao et al., 2015). It has been shown that ESCs have higher DNA methylation levels than ICM cells under conventional conditions, as opposed to 2i conditions (Leitch et al., 2013). DNA methylation and DNA methyltransferases expression have been broadly implicated in early embryo development and long-term maintenance of pluripotent cells (Arand et al., 2012; Ficz et al., 2013; Habibi et al., 2013; Marks et al., 2012; Smith et al., 2012), but the exact epigenetic events that occur during the derivation of ESCs have remained elusive (Tang et al., 2010). Here, we have observed that the expression of epigenetic modifier genes, such as Dnmt3b, Dnmt3l, Chd8, Mts1, Suz12, Eed, Wdr3, and Mat2b, was significantly increased in the intermediate stages of ESC derivation, and followed an upward trend during the course of the culture (Figure 4C). Next, we measured the level of DNA methylation in the CpG dyads of IAP-LTR1 and L1Md_tf retrotransposable elements and mSat. Consistent with the results of previous studies (Ficz et al., 2013; Habibi et al., 2013; Smith et al., 2012), this entire dataset has highlighted the role of epigenetic modifiers and DNA methylation, pointing toward a reconstruction process that shapes a different epigenetic landscape for ESCs. In contrast with previous observations (Ficz et al., 2013; Okashita et al., 2014), blockage of the ERK pathway was probably required for a high expression of epigenetic modifiers and DNA methylation-related genes during ICM to ESC transition. Interestingly, we found that RG108-mediated inhibition of DNA methylation impeded the efficient derivation of ESCs in the presence of R2i, yet had no adverse effects under 2i conditions. This analysis highlighted a major difference between the R2i and 2i conditions in the regulation of pluripotency. It has previously been demonstrated that the augmented BMP4 signaling pathway plays a key role in R2i pluripotency, whereas it does not make a difference in the 2i condition (Gomes Fernandes et al., 2016; Hassan et al., 2014b). Despite the high degree of similarity between the R2i and 2i conditions, as well as the strength of R2i in supporting pluripotency, i.e., in establishing and maintaining ESCs from single blastomeres (Hassani et al., 2014a) and embryonic germ cells from the PGCs of mice and rats (Attari et al., 2014; Mohammadi et al., 2015), these two conditions work in different ways.

To determine the state of R2i pluripotent cells in comparison with in vivo pluripotent cells, we compared our transcriptome data with that of early embryonic cells (Figure 7). Although a considerable number of preEpi- and PE-specific markers demonstrated concurrent expression patterns in both the R2i- and SB-related clusters, the enrichment of substantial preEpi- and PE-specific genes in the R2i and SB groups, would suggest a specified path for the generation of ESCs from ICM cells. We observed that R2i treatment could support the expression of diapause- and PGC-related genes, which were associated with the preEpi cell state. A comparison of 2i-grown cells with early embryonic cells indicated that there was greater similarity with them and E4.5 preEpi cells (Boroviak et al., 2015). In this study, we also demonstrated that R2i could sustain the preEpi-specific gene expression during the process of establishing ESCs from the ICM. IOs and early-passage ESCs were more similar to preEpi cells than were P15 ESCs. This was consistent with data by Boroviak et al. Therefore, despite certain differences between 2i and R2i for maintaining pluripotency, both conditions produced ESCs that
exhibited high similarity to E4.5 preEpi cells. However, our results demonstrated that the establishment of ESCs requires a reprogramming phenomenon such as DNA methylation regulation and EMT blockage.

To the best of our knowledge, this is the first study reporting that EMT prevention is required for the derivation of ESCs from the ICM. In addition to the maintenance of preEpi-specific marker gene expression, prevention of TGF-β and MAPK under R2i conditions tended to inhibit the EMT process in normal development of the early embryo and to maintain pluripotency in vitro. From the perspective of human ESCs, it would be of tremendous interest to investigate the role of EMT or cell adhesion in the derivation of primed and naive human ESCs.

**EXPERIMENTAL PROCEDURES**

**ESC Derivation and Sample Collection**

For ESC derivation, blastocysts/isolated ICMs were prepared as described in the Supplemental Experimental Procedures. The isolated ICMs were washed twice in PBS, and then selected for microarray analysis. ESCs were derived by transferring blastocysts/isolated ICMs on gelatin-coated plates (0.1%, Sigma-Aldrich) that contained N2B27 defined medium supplemented with R2i (which included 1 μM PD0325901 [Stemgent] and 10 μM SB431542 [Sigma-Aldrich]) and 1,000 U/mL LIF (ESGRO, Millipore). This time was designated as day 0. For the first microarray analysis, the samples included isolated ICMs (day 0), BOs on days 3, 5, 7, and 9 (BO3–9) after plating, and ESCs on passage 20, in three biological replicates. We chose and pooled approximately 20–30 isolated ICMs and BOs for each biological replicate.

For the second microarray analysis, we collected new samples in a time resolution experiment that included immunosurgically isolated ICM, IOs, on days 0.5, 1, 2, 3, and 5 (IO0.5–5), and passages 2, 4, and 15 of ESCs (P2–15). In addition, IOs which were cultivated isolated ICMs were washed twice in PBS, and then selected for microarray analysis. The isolated ICMs were cultured in N2B27 supplemented with SB431542 + LIF on days 1 (SBIO-1) and 3 (SBIO-3) considered as the negative controls. Approximately 30–40 ICM or IOs were picked and pooled in two biological replicates.

**ACCESSION NUMBERS**

Gene profiling data was submitted to the GEO repository with accession number GEO: GSE87793.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, six figures, and four tables and can be found with this article online at http://dx.doi.org/10.1016/j.stemcr.2017.08.006.

**AUTHOR CONTRIBUTIONS**

M.T., S.N.-H., and A.S. developed the study. N.T. performed some of the EMT experiments and discussed the EMT results. K.A. discussed the EMT results. J.A. and J.W. performed the DNA methylation experiments. B.G. discussed the gene expression results. D.S. assisted with animal and embryo manipulations. M.A. analyzed the microarray data. M.P. performed the time course derivation and supplemental experiments. H.G. supervised the project, provided financial support, and discussed the results. H.S. designed the experiments, discussed the results, and approved the manuscript. H.B. provided financial support, designed and analyzed experiments, discussed the results, and approved the manuscript.

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