Rewirable gene regulatory networks in the preimplantation embryonic development of three mammalian species

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**Supplemental Material**

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Rewirable gene regulatory networks in the preimplantation embryonic development of three mammalian species

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Mammalian preimplantation embryonic development (PED) is thought to be governed by highly conserved processes. While it had been suggested that some plasticity of conserved signaling networks exists among different mammalian species, it was not known to what extent modulation of the genomes and the regulatory proteins could “rewire” the gene regulatory networks (GRN) that control PED. We therefore generated global transcriptional profiles from three mammalian species (human, mouse, and bovine) at representative stages of PED, including: zygote, two-cell, four-cell, eight-cell, 16-cell, morula and blastocyst. Coexpression network analysis suggested that 40.2% orthologous gene triplets exhibited different expression patterns among these species. Combining the expression data with genomic sequences and the ChIP-seq data of 16 transcription regulators, we observed two classes of genomic changes that contributed to interspecies expression difference, including single nucleotide mutations leading to turnover of transcription factor binding sites, and insertion of cis-regulatory modules (CRMs) by transposons. About 10% of transposons are estimated to carry CRMs, which may drive species-specific gene expression. The two classes of genomic changes act in concert to drive mouse-specific expression of MTF2, which links POU5F1/NANOG to NOTCH signaling. We reconstructed the transition of the GRN structures as a function of time during PED. A comparison of the GRN transition processes among the three species suggested that in the bovine system, POU5F1’s interacting partner SOX2 may be replaced by HMGB1 (a TF sharing the same DNA binding domain with SOX2), resulting in rewiring of GRN by a trans change.

[Supplemental material is available online at http://www.genome.org. The microarray data from this study have been submitted to the NCBI Gene Expression Omnibus (http://www.ncbi.nlm.nih.gov/geo) under accession nos. GSE18290 and GSE18319.]

Much of the information on evolution of gene regulatory networks (GRN) was generated from studies in invertebrates (Erwin and Davidson 2009). Comparison of fungi species revealed evolution of GRNs at both cis and trans levels (Scannell and Wolfe 2004). Invertebrate GRNs can be “rewired” at the “cis” level through changes in genomic regions that regulate gene transcription, or cis-regulatory modules (CRM). CRM can gain or lose transcription factor (TF) binding sites (TFBSs) by single nucleotide mutations, insertions and deletions of neutral sequence or TFBSs (Erwin and Davidson 2009). It has been argued that at the cis level, the transcription regulatory apparatus is a “programmable” computing machine whose function can be modulated through placements of TFBSs in the regulatory region of any gene (Buchler et al. 2003; Gertz et al. 2009). A notable advantage of encoding combinatorial control in the regulatory region, as opposed to in the regulatory proteins, is evolvability (Ptashne and Gann 2002): Unlike regulatory proteins, each cis-regulatory region controls the expression of a given gene and hence can be programmed with minimal pleiotropic effects. Contrary to this argument, changes to the TF that are produced by the cell (“trans” changes) appear to be widely tolerated by Escherichia coli and may even confer evolutionary advantages (Isalan et al. 2008). Moreover, several yeast species implement compensating cis and trans changes to achieve a conserved regulatory logic (Tsong et al. 2006). This study attempted to add vertebrate data to address these arguments by analyzing a relatively conserved developmental process in three mammalian species.

Mammalian preimplantation embryonic development (PED) encompasses the period from fertilization to implantation of the embryo in the endometrial lining of the uterus. Just after fertilization, the transcriptome of the embryo is comprised of maternally deposited transcripts. After several cell divisions, maternal transcripts are specifically degraded and are replaced by zygotic transcripts produced by the new diploid nucleus containing both maternal and paternal genes. This transition is termed zygote genome activation (ZGA). The timing of ZGA and maternal transcript degradation varies among species, but in humans and bovines it reportedly occurs between the four- and eight-cell stages of...
development, while in mouse, this transition occurs between the one- and two-cell stages of development, consistent with an overall accelerated pace of murine embryonic development (Braude et al. 1988; Kanka et al. 2003; Adjaye et al. 2007; Kues et al. 2008).

Another major event during PED is the differentiation and specialization of cells. Immediately after ZGA, embryos undergo compaction, a process during which dividing embryonic cells become a tightly organized cell mass with an indistinguishable membrane. In most mammalian embryos, compaction is usually observed between the eight- and 16-cell stages. After compaction, the embryo is named the morula, which becomes the blastocyst through formation of an internal cavity and further cellular differentiation. At least two cell types can be distinguished in the blastocyst: an inner cell mass (ICM) and an outer layer of cells termed the trophectoderm. The ICM contributes to all three germ layers present in the mature adult (known as pluripotency). Embryonic stem (ES) cells were derived from ICM, and ES cells have been used as surrogates to analyze the molecular mechanisms in ICM and preimplantation embryos (Ogawa et al. 2008). The progression of PED is highly conserved among mammals: All progress through the same morphologic stages. Perhaps the most marked difference is the amount of time spent at each stage. The other notable interspecies differences appear after the blastocyst stage, including that bovine blastocysts initiate the gastrulation process before implantation, and the formation and functions of the placenta and yolk sac (Gossler 1992). In this study, we chose the highly conserved PED stages, i.e., from zygote to blastocyst stages for cross-species GRN comparison.

Results

Dynamic gene expression landscapes of preimplantation development in three species

Human, mouse, and bovine embryos were collected at representative stages of PED, including oocyte (bovine only), one-, two-, four-, eight-, 16-cell (bovine only), morula, and blastocyst. Developmental stage was determined by direct microscopic visualization of each collected embryo (Fig. 1A). RNA from each stage was collected and analyzed with species-specific Affymetrix GeneChips. Biological replicates of the same developmental stage exhibited strong correlations, which were comparable to the sample correlations in previous bovine and mouse studies (Fig. 1B; Hamatani et al. 2004; Wang et al. 2004; Kues et al. 2008). Hierarchical clustering analyses revealed global transcriptional similarities among the biological replicates of the same developmental stages together, with a few exceptions of the samples at neighboring stages (Fig. 1C). The hierarchical order of all the samples almost perfectly reflected the relative time differences of all the developmental stages. The largest changes of expression were observed between four- and eight-cell stages in human and mouse and between eight- and 16-cell stages in the bovine system, in keeping with the previously reported timing of ZGA in these species.

In order to explore the conservations and changes of gene expression patterns across species, we developed a Bayesian clustering method to simultaneously cluster the timecourse expression data and automatically determine the cluster number (Methods). Nine clusters were generated for 5941 orthologous gene triplets (including 1036 TF triplets) (Supplemental Fig. S1). Of these orthologous triplets, 2122 (35.7%, 383 TF triplets) fell into clusters with consistent expression patterns in all the three species, and the remaining 3290 orthologous triplets (55.4%, 444 TF triplets) showed at least two different expression patterns among these species. The large differences in the TF expression patterns suggested...
that there might be a substantial interspecies difference of GRN structures. To test if this result is sensitive to the input genes, we repeated this analysis with the 927 orthologous gene triplets (including 371 TF triplets) that exhibited significantly variable expression (ANOVA P-value < 0.05) during PED in at least two of the three species. Among them, 383 (40.2%) orthologous triplets (149 TF triplets) showed different expression patterns among these species, consistent to the estimated large-scale GRN changes.

The three mammals share more maternally deposited than ZGA-activated RNAs

The transition from predominance of maternal transcripts to zygotic transcripts is a critical aspect of PED. We identified maternally deposited (MT) and ZGA-activated transcripts (ZGAT) in each species based on trends in transcript levels (Fig. 2B). Comparing the conservation levels of the human MTs vs. human ZGAT,

Figure 2. Cross-species comparison of gene expression. (A) Pictorial description of trends in transcription of a subset of genes with distinct roles during PED. Human, mouse, and bovine probe sets are depicted by red, green, and blue curves, respectively. When a gene is targeted by multiple probe sets, all probe sets are shown. The maternally deposited transcript for TCF7 is degraded at two-, four-, and eight-cell stages in mouse, human, and bovine embryos, respectively. SIN3A is highly abundant in zygotes, and exhibits consistent decreasing expression patterns in all three species. POU5F1 exhibits a consistent up-regulation in all three species, however its expression peaked at the eight-cell stage in mouse and at the morula stage in human and bovine. MTF2 transcript exhibits a reversed expression pattern in mouse and in the other two species, as reproduced by two probe sets in each species. (B) Conserved and species-specific MT and ZGAT. A percentage of MT, 33.7%, is shared in all three species, whereas only 3.2% of the ZGAT is shared in all three species. MT has a 10.7-fold larger chance of being shared across species than ZGAT (P-value < 10^-30).
Rewiring of ZGA genes by TFBS usage
We hypothesized that the large-scale changes of ZGA across species is associated with changes of TFBSs of TFs that activate gene expression in morulas and ICM. To test this hypothesis, we intersected human and mouse ZGA genes with the genes that are bound by POU5F1, SOX2, and NANOG in human ES cells (Lister et al. 2009) and mouse ES cells (Chen et al. 2008) (Methods). POU5F1, SOX2, and NANOG are hereafter referred to as ZGA TFs. Among the ZGA genes that are conserved, human-specific and mouse-specific, 58, 64, and 335 of them are bound by ZGA TFs in a conserved, human-specific and mouse-specific manner (Supplemental Fig. S2B). Surprisingly, the TFBSs of ZGA TFs are not enriched in the conserved ZGA genes ($\chi^2$ test with continuity correction $P$-value > 0.31). Instead, mouse-specific ZGA genes are enriched with mouse-specific binding of ZGA TFs ($P$-value $< 2 \times 10^{-6}$). The large changes of ZGATs and the TFBSs of ZGA TFs across species may suggest a large scale of GRN rewiring events during mammalian PED. We note that there can be other ZGA TFs besides POU5F1, SOX2, and NANOG, and therefore this analysis is simply a primitive attempt to estimate the extent of rewiring events associated with ZGA.

Transposons contribute to species-specific gene expression
There is a significant overlap between transposon-associated elements and TFBS-rich regions in mammalian genomes. One notable example includes association between the TPS3 binding sites and transposons in the human genome (Wang et al. 2007; Bourque et al. 2008), and one of the TPS3 binding transposons was stabilized in humans (Wang et al. 2007); however, it is yet to be determined to what extent the transcriptome is modulated by transposons. Therefore, we attempted to assess the overall impact of murine-specific transposons on the PED transcriptome.

Previously reported chromatin immunoprecipitation with massively parallel sequencing (ChIP-seq) mapping of mouse ES cells identified binding sites of 16 transcriptional regulators to the mouse genome (Chen et al. 2008). Based on this information, we identified the binding regions that resided in murine-specific transposable regions (Fig. 3A). CTCF-binding regions exhibited the largest number of overlap with known transposons (10,313 overlapping regions, representing 26.0% of total CTCF-binding regions), among which 9074 (22.9%) overlapped with the B2 family of transposons. In addition to the CTCF-binding regions, the binding regions for the following factors exhibited nontrivial overlaps ($\geq 10\%$ of the binding regions of a factor) with transposons: ESRRB, SOX2, NANOG, SMAD1, TCFCP21, STAT3, EP300, and POU5F1. Transposon families B2, ERVK, and B4 were most strongly associated with the binding regions of these transcription regulators. This suggests that these families of transposable elements contain the DNA binding motifs of the listed TFs. On the contrary, 10 (0.2%) and 1 (0%) of Suz12 and RNA polymerase II (Pol II) binding regions overlapped with transposons, far below the expected amount of overlaps (16.3%), suggesting that the insertion of transposable elements to the binding regions may negatively affect the function of these factors.

Gain of functional TFBSs and CRM by species–specific transposons
To confirm the functional importance of the transposons to gene regulation, we further characterized two murine-specific transposons that carry functional TFBSs and CRM and, in turn, regulate mouse-specific gene expression in early development. One such transposon is Lx8, a member of non-LTR retrotransposons, which we estimated to be inserted into the murine genome after the divergence of murine and primate species, ~60 million years ago (Supplemental Fig. S3). A copy of Lx8 is present upstream of the we found 69.4% human MT and 40.0% human ZGAT were shared in mouse (ratio = 69.4/40.0 = 1.74, $P$-value $< 10^{-10}$), 63.8% hMT and 18.5% human ZGAT were shared in bovine (ratio = 3.45, $P$-value $< 10^{-20}$), 49.0% human MT and 8.5% human ZGAT were shared in both mouse and bovine (ratio = 5.76, $P$-value $< 10^{-20}$). These data showed that human MT is more conserved than human ZGAT in mammals (Fig. 2A,B). Comparisons benchmarked against mouse or bovine MT and ZGAT generated similar results. Changing the algorithmic thresholds for detection of MT and ZGAT did not affect this conclusion (Supplemental Fig. S2A).
cold shock domain–containing protein A (Csda) gene in the mouse genome. Csda is a transcription repressor, highly conserved among eukaryotes from yeast to human. In the mouse, Csda is known to be critical for several processes in reproduction and development, including spermatogenesis and in utero fetal development. The function of Csda in other mammals has not been tested. Transcription of Csda in mouse PED starts at the time of ZGA (two-cell stage), with an apparent pause during the four-cell stage, followed by continued increase in transcription from the eight-cell stage to the morula stage (Fig. 4). The increase in Csda transcription during mouse PED was correlated with Pou5f1 expression (R = 0.41, permutation P-value = 0.046). Like Pou5f1, Csda transcripts were restricted to the ICM at the blastocyst stage (Yoshikawa et al. 2006); however, ZGA of CSDA was not conserved in human or bovine, suggesting that the mouse Csda is associated with species-specific regulatory sequences. The insertion of Lx8 upstream of the mouse Csda gene produces a consensus POU5F1 binding site ([T/A]ATG[C/T]AAAT). This POU5F1 TFBS was functional in mouse ES cells: Analysis of published ChIP-seq data (Chen et al. 2008) revealed that the POU5F1–SOX2-bound regions together with the Pol II-bound Csda promoter. ChIP-chip analyses of human ES cells did not identify any POU5F1, SOX2, or NANOG binding sites upstream of the human CSDA gene. Furthermore, motif scans based on the POU5F1 and NANOG motifs did not detect any high-affinity TFBS within 20 kilobases (kb) upstream or downstream of the human CSDA transcription start site (TSS). The data produced by gene expression, DNA motif, and TF binding analyses collectively suggest that the unique, transposon-mediated change in the regulatory sequence of the mouse Csda gene is associated with a gain of function.

A second transposon in the mouse genome is associated with species-specific gene expression during PED. This transposon, Orr1b1-int, a member of the ERV3 family of retroviruses, is inserted upstream of the murine Mtf2 gene (Fig. 5A,D). Orr1b1-int is a “younger” transposon than Lx8 (Supplemental Fig. S3), but may
have more potent gene regulatory effects than Lx8. Orr1b1-int contains a CRM comprised of two TFBSs (POU5F1 and a NANOG), in contrast with the single POU5F1 site in Lx8. ChIP-seq data revealed strong binding of POU5F1 and NANOG to their corresponding TFBSs within this Orr1b1-int, as reflected by large sequence counts: (the NANOG site ranked 434 among 10,345 detected NANOG sites, and the POU5F1 site ranked 66 among 3736 detected POU5F1 sites) (Chen et al. 2008). The POU5F1 binding on this Orr1b1-int region was five times stronger than on the Lx8 Csda enhancer. For both NANOG and POU5F1, the minimal overlaps of the ChIP-seq reads pinpointed to sites that matched their known DNA binding motifs (Fig. 5D), further confirming that the TF–DNA binding was a result of the presence of TFBSs within the Orr1b1-int sequence. These data, together with the gain of another NANOG TFBS by mutation (see below), offer an explanation for the dramatic difference in MTF2 expression among the three mammals analyzed in this study (Fig. 2A). Although it is not known why MTF2 transcripts are present in only human and bovine oocytes, the dramatic, mouse-specific induction of the Mtf2 gene during PED is consistent with the gain of murine-specific regulatory sequences.

**Turnover of TFBSs caused by single nucleotide mutations**

Besides transposons, we postulated that single nucleotide changes may cause TFBS turnover and thus be responsible for interspecies differences in mammalian PED. Associating TFBSs with single nucleotide mutation with the genes with species-specific expression, we found three single nucleotide mutations that appear to influence gene expression by modulating the binding affinities of the TFBSs. Two of the three mutations locate within the same TFBS upstream to the Frat2 gene, and the third mutation locates upstream to the Mtf2 gene.

FRAT2 belongs to the GSK-3-binding protein family and is a positive regulator of the WNT signaling pathway. Expression of the human and bovine Fra2 genes is strongly induced in late PED, whereas the mouse Frat2 gene appears to be transcriptionally

![Figure 4](https://example.com/figure4.png)
silent throughout PED. (Supplemental Fig. S4A). The absence of expression in mouse is attributable to an A-to-C mutation at the center position of the POU5F1 binding site (Supplemental Fig. S4C,D), which eliminated the POU5F1 binding, as indicated by a chromatin immunoprecipitation with a microarray hybridization (ChIP-chip) peak on the human POU5F1 site (Boyer et al. 2005), but no ChIP-seq signal in the genomic neighborhood of the mouse Frat2 gene (Chen et al. 2008) (Supplemental Fig. S4B). This mutation at the center of the DNA binding motif caused a loss of the TFBS. Furthermore, the human Frat2 exhibits larger induction of expression in response to the increased concentration of POU5F1 during PED, as compared with the bovine system. This delicate difference is attributable to a primate-specific T-to-A mutation at the second position (a peripheral position) of the same POU5F1 TFBS (Supplemental Fig. S4C,D). This mutation made the human TFBS identical to the consensus sequence of the POU5F1 motif, thus generating a stronger TFBS than the bovine version (Matys et al. 2003). In summary, in the case of an orthologous group of POU5F1 TFBSs, two mutations are present in DNA motif. The central mutation was consistent with a dramatic change on gene expression, and the peripheral mutation was reflected by a delicate change of expression.

A NANOG TFBS resides upstream of the Mtf2 gene in the mouse, but not in human or cow. ChIP-seq data accumulated over 62 sequence reads, overlapping on a single nucleotide in the mouse genome. (C) Electrophoretic mobility shift assay results confirm that mutation of the mouse binding site back to the version in other mammals (A-to-T) eliminates NANOG binding in vitro. (A>T) A-to-T mutation; (A>C) A-to-C mutation. (D) A mouse-specific transposable element inserts a CRM for POU5F1 and NANOG into the region upstream of Mtf2: This CRM is bound by POU5F1 and NANOG in mouse ES cells. (E) Expression levels of Mtf2, Pou5f1, and Nanog in mouse ES and trophoblastic stem (TS) cells. (F) Mtf2 knockdown in mouse ES cells affects the expression of other transcription factors and signaling proteins, including the down-regulation of another ICM-specific gene, Notch3. pSUPER and Luci are control vectors that carry an empty vector (pSUPER) and shRNA against a luciferase gene (Luci). RNAi-1 and RNAi-2 are knockdowns mediated by two different shRNAs.
at a location corresponding to a NANOG binding site (Fig. 5A). The position in question is occupied by an A in the mouse and a T in all other mammals. This suggests that the murine lineage contains a distinct T-to-A mutation (Fig. 5B). We sought to assess whether the putative "ancestral" sequence (not containing the T-to-A mutation) might harbor a NANOG TFBS. To this end, we mutated the functional binding site from A (mouse) back to T (ancestral). This mutation abrogated NANOG binding to the site in vitro (Fig. 5C).

We therefore reconstructed the dynamic transitions of GRNs during PED in the human, murine, and bovine systems. For each stage of PED, we identified the interacting modules of genes (in each species) by jointly modeling the expression data and a reference protein interaction network with the MATISSE program as described in Methods (Fig. 6A; Ulitsky and Shamir 2007). The expression data from a given developmental stage were used in conjunction with expression data from adjoining stages: e.g., one-, two-, and four-cell expression data were used for reconstructing the interacting modules at the two-cell stage. The reference protein interaction network was generated by integrating the data from the Human Protein Reference Database (Mishra et al. 2006) with high throughput yeast two-hybrid screens (Ulitsky and Shamir 2007) and mass spectrometry analyses (Wang et al. 2006; Ewing et al. 2007). After constructing the state transition of the GRNs (http://biocomp.bioen.uiuc.edu/ped), we focused our analyses on the modules that contained the key regulatory genes that were induced in ES cells, as opposed to differentiated cells (Fig. 6B). Although expression data from the blastocyst-stage embryos were generated from a mixture of both ICM and trophectodermal cells, the reconstructed GRN in human blastocysts resembles the GRN of human ES cells (Muller et al. 2008). This similarity is highlighted by the interaction of POU5F1, SOX2, NANOG, SALL4, and C1orf103. The resemblance of the two GRNs lends credibility to the GRN reconstruction procedure and also suggests the genes involved in the POU5F1 interaction module exhibited a strong coexpression pattern during the morula-to-blastocyst transition.

Dynamic changes in GRNs during PED

The static GRNs active in human ES cells were recently described (Muller et al. 2008), although the dynamic processes that are capable of generating pluripotent GRNs remain elusive (Mikkelsen et al. 2008). Since many varieties of ES cells are procured from the inner cell mass of blastocyst-stage embryos, a more comprehensive understanding of the molecular events that drive PED could illuminate the mechanisms by which pluripotency-associated GRNs active in ES cells are constructed. Species-specific differences in the dynamic development of these GRNs may also help us better understand the different behaviors of ES cells from different species.
Discussion

An evolutionary model for TFBS insertion by transposons

It has been recently observed that species-specific transposable elements carried a substantial fraction of TFBSs that are associated with TFs in vivo (Wang et al. 2007; Bourque et al. 2008). The influence of these transposable elements on the transcriptome has not been extensively explored. Our analyses suggest that although tens of thousands of transposons bind with TFs in vivo, the majority of such TF-bound transposons do not affect the expression of the surrounding genes. We estimate that 10% of murine-specific transposons, generally those that carry CRMs, may affect gene expression. These data suggest a model for the evolution of cis-regulatory sequences by insertion of transposons. The TFBSs associated with transposons are inserted into the host genome and bind with corresponding TFs with similar stoichiometric coefficients as the parent TFBSs. The majority of these inserted TFBSs do not affect the expression of the surrounding genes and are evolutionarily neutral. A small fraction of the transposons, especially those that carry CRMs and attract a module of TFs, can change the expression of surrounding genes, and may be subject to positive or negative selections.

ES cells better represent ICM than EpiSCs

Pluripotent stem cells have been derived from several embryonic source tissues in mouse: ES cells are derived from the inner cell mass of blastocyst stage embryos, and epiblast stem cells are derived from the late epiblast layer of post-implantation mouse embryos (mEpiSCs) (Brons et al. 2007). mEpiSCs and mES cells exhibit nonidentical TF binding patterns and gene expression patterns. In terms of both enhancer usage and gene expression patterns, mES cells resemble ICM, whereas mEpiSCs are different. Based on these data, it was suggested that mES cells and mEpiSCs are two distinct pluripotent states representing cells of the preimplantation embryo and later epiblast cells (Brons et al. 2007), which is in line with our approach of using TF-binding data in ES cells to study TF-binding in PED.

Pluripotency is a conserved phenotype that can be sustained by evolvable GRNs

Human and mouse ES (hES and mES) cells are similar in the sense that they are both pluripotent cell populations that are capable of self-renewal that are derived from the ICM of blastocyst embryos. Even so, the growth factor requirements for hES and mES maintenance are different: mES cells require the presence of LIF, whereas hES cells do not. The differences in media conditions are thought to reflect differences in critical signaling networks responsible for maintenance of pluripotency and self-renewal capability between species. For example, TF Foxd3 is required for mES cell self-renewal, but its expression appears to be nonessential for hES (Ginis et al. 2004). Similarly, STAT3, a TF downstream to LIF signaling, is also required for self-renewal and the maintenance of pluripotency of mES cells, but it seems to be dispensable in hES cells (Schuringa et al. 2002). Compared to mES cells, mEpiSCs are functionally more similar to hES cells in several ways, including their responses to BMB4 and Activin/Nodal signaling (Brons et al. 2007).
observations led to the hypothesis that pluripotent cell identity can be established and maintained through more than one GRN. These GRNs may share core components universally indispensable for pluripotency. Peripheral components, though critical for cell fate, can be implemented using alternative designs. Because ES cells are derived from the ICM of blastocyst embryos, the natural processes of building pluripotent GRN are reflected by the transitions of the maternal cellular environment to ZGA-activated GRN. The comparison of GRN across multiple stages of PED suggested novel factors, such as MTF2 and HMGB1 for building a pluripotent GRN. These factors serve as candidates of dispensable reprogramming factors, which may be tested for potential enhancement of reprogramming efficiency. Since these factors may already be expressed in some tissues, they may contribute to customizing the combination of reprogramming factors for each tissue.

Methods

IVF embryos

The IVF conditions were optimized for each species. To minimize the influence of IVF conditions to cross-species comparison, we subsequently based the comparisons on either relative expression changes or coexpression patterns (Stuart et al. 2003; Gilad et al. 2006; Tirosh et al. 2007).

Human embryos

Supernumerary human embryos were obtained from patients at the Boston IVF Clinic through informed consent with the approval of the Internal Review Board of Harvard University. Embryos thus obtained were cultured in a two-step culture system (Sage/Biopharma) in microdrops at 37°C and 5% CO2 under oil (Sage) as described previously (Kaeberlein and Powers 2007). Embryos from each stage were segregated into three groups. Each group was considered to be a biologic replicate: Embryos within each replicate were pooled for further analysis. The blastocyst embryos were graded according to the Gardner scale (Brook and Gardner 1997; Van Royen et al. 1999).

Mouse embryos

Mouse embryos were obtained as described previously (Wang et al. 2004). Briefly, female mice were impregnated, and embryos were flushed from the oviducts. Individual embryos were morphologically staged by light microscopy.

Bovine embryos

Ovaries were washed, and 2–6 mm follicles were aspirated. Only oocytes surrounded by several cell layers of dense cumulus were selected for culture. Oocytes were washed three times in TL-HEPES (0.22 mM sodium pyruvate, 3 mg/mL BSA at pH 7.4), and we placed maturation media that had been equilibrated in 5% carbon dioxide and air (39°C, high humidity). The maturation medium consisted of TC199 supplemented with 3 mL of bovine LH and FSH (Siuoh Biochemical), 0.25 mM sodium pyruvate, and 10% fetal calf serum. After 20 h of incubation in maturation media, oocytes were removed, washed in TL-HEPES and placed in groups of 10 oocytes each into 44-μL microdrops of IVF-Talp (Biowhittaker) supplemented with 0.22 mM sodium pyruvate, and 6 mg/mL essentially fatty acid-free BSA. Live sperm were separated by centrifugation in a percoll gradient (Sigma, P1985). Eggs were fertilized by adding one million/mL of sperm, 2.0 μg of heparin (Sigma), pencillamine, hypotaurine, and epinephrine (Sigma) to each drop of fertilization media. Oocytes were removed from the fertilization media after 18–22 h, washed in TL-HEPES, placed in a 1.5-mL tube, and vortexed for 2 min to remove cumulus cells. Twenty-five cumulus-free oocytes were placed in each 50-μL drop of SOF culture medium, supplemented with 8 mg/mL of fatty acid–free BSA, nonessential and essential amino acids, and pyruvate.

Transcription profiling

RNA was isolated from embryos using the PicoPure Isolation system (Arcturus) according to the manufacturer’s instructions. Two rounds of linear amplification were performed for each sample to generate cDNA probes from total RNA as previously described (Baugh et al. 2001) using the PicoAmp system (Arcturus). The cDNA produced by this system was converted to aRNA and labeled using a T7-based Invitro-Transcription Labeling system (Enzo Lifesciences). Labeled aRNA was hybridized to the Affymetrix U133 2.0 plus, MOE430A 2.0 and Bovine Genome Genechips array (Affymetrix) according to the manufacturer’s instructions. These expression data were submitted to the NCBI Gene Expression Omnibus (GEO) (http://www.ncbi.nlm.nih.gov/geo) under accession no. GSE18290. The raw data for each species were normalized and modeled with dChip software.

Clustering analysis

We developed a Bayesian clustering method to cluster the timecourse expression data (W. Huang, X. Cao, G. Lin, D. Xie, and S. Zhong, in prep.; see Supplemental material). The cluster generating process is modeled with a Dirichlet Process (Rasmussen 2000) and the time course profile of each cluster is modeled with a mixed-effects model with splines (Luan and Li 2003). The advantages of this method are: (1) it explicitly utilizes the timecourse data structure to minimize the adverse effects of noise in the data; (2) it automatically identifies the proper cluster number; (3) it is based on a fully described probabilistic model and therefore it minimizes the use of heuristic algorithmic steps. The difference in developmental time had little contribution to the detected difference of expression patterns. For example, maternal deposited RNA for TC7, the canonical TF for the WNT signaling pathway, started degradation at four-, two-, and eight-cell stages in human, mouse, and bovine, respectively (Fig. 2A), but TC7 was grouped into clusters of consistent expression patterns.

Identifying genes with mouse-specific expression changes

For the mouse vs. human comparison, we defined mouse-specific expression change as a gene exhibiting a non-flat expression trajectory in mouse PED, but its orthologous human gene exhibiting a flat expression trajectory in human PED, ignoring the bovine expression data. For the mouse vs. human and bovine comparison, we defined mouse-specific expression change as a gene exhibiting a non-flat expression trajectory in mouse PED, but its orthologous genes exhibiting flat expression trajectories in both human PED and bovine PED. To detect if a gene has a flat expression trajectory, we compared the expression levels between the first time point and every later time point, using t-tests. If any of the t-tests showed a significant difference (P-value < 0.05), then this gene was considered to have a non-flat trajectory. If none of the t-tests showed a significant difference, this gene was considered to not have any expression changes.

Detection of maternal and ZGA transcripts

The MT genes were required to have presence call in all the replicates at one-cell stage (oocyte and one-cell stages for bovine). The ZGA genes were identified with the following criteria: There is a detectable increase in expression levels during PED. To detect an
increase of expression, we compared every time point to its previous time point. A detectable increase was defined as: fold change > 2, difference of dChip expression index > 100 and two-sample comparison P-value < 0.05.

ChIP-seq data

ChIP-seq data in human and mouse ES cells were obtained from GSE17917, GSE18292 (Lister et al., 2009), and GSE11431 (Chen et al., 2008). ChIP-seq identified TFBSs were associated with a gene if it is within a 50,000-bp flanking region (25,000 upstream and downstream) of the transcription start site of the gene. A ZGA gene is called bound by a ZGA TF, if it is bound by any of the three Tfs: Pou5f1, Sox2, and Nanog.

Estimation of transposon ages

All the Lx8 and Orr1b1-int transposons were downloaded from the RepeatMasker track on the UCSC Genome Browser (http://genome.ucsc.edu). The ages of the repeats were estimated by age = divergence/substitution rate (Bourque et al., 2008). The substitution rate of the mouse genome used in this analysis was 4.5 × 10^{-9} (Mouse Genome Sequencing Consortium 2002).

Detection of TFBS-carrying transposons

Coordinates of ChIP-seq detected binding regions of 16 transcription regulators in mouse ES cells were downloaded from GEO (Chen et al., 2008). Coordinates of transposable elements were downloaded from the UCSC Genome Browser, and the murine specific transposons were identified by the annotation of RepBase (Jurka et al., 2005). We took the intersection of these two sets of sequences and obtained a list of TFBS-carrying transposons.

Detection of species-specific expression

The map of orthologous human, mouse, and bovine probe sets was downloaded from the Affymetrix website (http://www.affymetrix.com/Auth/analysis/downloads/na28/ivt/Bovine.na28.ortholog.csv.zip). To resolve ambiguous mapping, for each gene we chose the probe set with the largest mean signal as the representative probe set for that gene. A gene was identified as exhibiting species-specific expression when either of the following two criteria was satisfied: (1) It was called “absent” in all time points in one species, but it had “presence” calls in other species. (2) When it was present in all species, its fold change of early (one- to eight-cell) and late (morula and blastocyst) embryos were computed. The fold change was more than twice as large in one species as compared to another.

shRNA-mediated RNA knockdown

Feeder-free E14 mouse ES cells were cultured at 37°C with 5% CO2. All cells were maintained on gelatin-coated dishes in DMEM (GIBCO), supplemented with 15% heat-inactivated FBS (GIBCO), 0.055 mM β-mercaptoethanol (GIBCO), 2 mM l-glutamine, 0.1 mM MEM nonessential amino acid, 5000 units/mL penicillin-streptomycin, and 1000 units/mL LIF (Chemicon), as described previously. Transfection of shRNA constructs was performed using Lipofectamine 2000 (Invitrogen) according to manufacturer’s instructions. Briefly, 1.5 μg of plasmid DNA was transferred into ES cells on 60-mm plates for RNA extraction. Puromycin (Sigma) selection was introduced 1 d after transfection at 1.0 μg/mL and maintained for 2 and 4 d before harvesting.

shRNA targeting specific genes was designed as previously described (Reynolds et al., 2004; Ui-Tei et al., 2004). The 19-nucleotide (nt) hairpin-type shRNAs with a 9-nt loop were cloned into pSUPER.puro (BglIII and HindIII sites, Oligoengine). Two shRNA, targeting different regions of respective transcripts, were designed for MTF2 to ensure specificity. These short nucleotides are: GGA GACTGTTATTTAAAGA, GCCACCATGCGTTTACA, pSUPER.puro constructs expressing shRNA against luciferase (Firefly) were used as controls.

EMSA

Electrophoretic mobility shift assay (EMSA) was used to analyze the binding affinities of putative NANOG TFBSs. The mouse wild-type “AGTGCTCA” was compared to two site-directed mutagenesis sites “AGTGCTCT” and “AGTGCTTC”, denoted as “A > T” and “A > C,” respectively. A recombinant NANOG protein (GST-tagged) was used for EMSA. The full-length NANOG protein was cloned into the pET42b (Novagen) vector, expressed and purified with GSH-sepharose beads (Amersham). DNA oligonucleotides (Proligo) labeled with biotin at the 5’ end of the sense strands were annealed with the antisense strands and were purified with agarose gel DNA extraction kit (Qiagen). A LightShift Chemiluminescent EMSA Kit (Pierce Biotechnologies) was used. One-hundred nanograms of protein were added to a 5-μL reaction mixture and incubated for 20 min at room temperature. The binding mixture was resolved on 6% native polyacrylamide gels. The gels were transferred to Bio-dyne B nylon membranes (Pierce Biotechnologies) using Western blot techniques and detected using chemiluminescence.

Microarray data for Mtf2 knockdown ES cells and trophoblast stem (TS) cells

Two Mtf2 knockdown ES cell samples and two samples with control vectors were analyzed with Agilent Whole Mouse Genome Oligo Microarray Kit, representing over 41,000 mouse genes and transcripts. These expression data were submitted to GEO under accession no. GSE18319. Affymetrix MU74Av2 microarray data for ES and trophoblast stem (TS) cells were obtained from the GEO database (http://www.ncbi.nlm.nih.gov/geo/, accession no. GSE3766).

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References


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