Factors from Human Embryonic Stem Cell-derived Fibroblast-like Cells Promote Topology-dependent Hepatic Differentiation in Primate Embryonic and Induced Pluripotent Stem Cells*\textsuperscript{S}

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Hsiang-Po Huang,\textsuperscript{a} Chun-Ying Yu,\textsuperscript{a} Hsin-Fu Chen,\textsuperscript{b,c} Pin-Hsun Chen,\textsuperscript{a} Ching-Yu Chuang,\textsuperscript{d} Sung-Jan Lin,\textsuperscript{e,f} Shih-Tsung Huang,\textsuperscript{g} Wei-Hung Chan,\textsuperscript{h} Tzuu-Huei Ueng,\textsuperscript{h} Hong-Nerng Ho,\textsuperscript{h,b,c} and Hung-Chih Kuo\textsuperscript{d,g} \textsuperscript{1}

From the Divisions of \textsuperscript{a}Medical Research and \textsuperscript{b}Reproductive Endocrinology and Infertility and the Departments of \textsuperscript{c}Dermatology and \textsuperscript{d}Anesthesiology, National Taiwan University Hospital, Taipei 10002, Taiwan, the \textsuperscript{e}Institute of Clinical Genomics, \textsuperscript{f}Biomedical Engineering, and \textsuperscript{g}Institute of Toxicology, National Taiwan University, Taipei 10617, Taiwan, the \textsuperscript{h}Institute of Clinical Medicine, Taipei Medical University, Taipei 11031, Taiwan, and the \textsuperscript{i}Genomics Research Center and the \textsuperscript{j}Institute of Cellular and Organismic Biology, Academia Sinica, Taipei 11574, Taiwan

The future clinical use of embryonic stem cell (ESC)-based hepatocyte replacement therapy depends on the development of an efficient procedure for differentiation of hepatocytes from ESCs. Here we report that a high density of human ESC-derived fibroblast-like cells (hESdFs) supported the efficient generation of hepatocyte-like cells with functional and mature hepatic phenotypes from primate ESCs and human induced pluripotent stem cells. Molecular and immunocytochemistry analyses revealed that hESdFs caused a rapid loss of pluripotency and induced a sequential endoderm-to-hepatocyte differentiation in the central area of ESC colonies. Knockdown experiments demonstrated that pluripotent stem cells were directed toward endodermal and hepatic lineages by FGF2 and activin A secreted from hESdFs. Furthermore, we found that the central region of ESC colonies was essential for the hepatic endoderm-specific differentiation, because its removal caused a complete disruption of endodermal differentiation. In conclusion, we describe a novel in vitro differentiation model and show that hESdF-secreted factors act in concert with regional features of ESC colonies to induce robust hepatic endoderm differentiation in primate pluripotent stem cells.

Patients with end stage and genetic liver diseases have been shown to benefit from liver cell replacement therapy (1). However, one of the major factors currently limiting the utilization of cellular replacement in the general population with severe liver disease is the lack of consistent sources of transplantable hepatocytes. Ideally, a proliferative stem cell population with robust capacity to generate hepatic cells in vitro will overcome the shortage of transplantable hepatic cells. Over the past several years, pluripotent embryonic stem cells (ESCs)\textsuperscript{3} have been successfully derived from nonhuman primates (2–4) and human embryos (5, 6). The ability of these cells to proliferate indefinitely and to develop into virtually any cell or tissue type, including hepatocytes, makes them likely candidates for cell-based treatment of human diseases.

Over the past few decades, studies in vertebrate models have described several signaling pathways critical for the embryonic development of hepatocytes (7, 8). For example, it has been shown that selective FGFs can substitute for the cardiac mesoderm to activate the expression of hepatic genes in the endoderm (9). In addition to the FGF pathway, bone morphogenetic factors (BMPs), which are highly expressed in the septum transversum mesenchyme, are capable of inducing early hepatic fate independently in mice (10). Subsequently, Wnt signaling plays an important role in enhancing the growth of the liver bud (7, 11). Other inductive signals, such as activin (12, 13) and Sonic hedgehog (14, 15), also contribute to endoderm or early liver development. Terminal maturation of hepatocytes requires additional signals, such as hepatocyte growth factor (HGF) (16, 17) and oncostatin M (17, 18).

The future clinical use of ESC-based hepatocyte replacement therapy depends on the development of an efficient procedure for differentiation of hepatocytes from ESCs. To achieve this goal, researchers have successfully exploited the molecular signals for liver development outlined above to direct the development of mouse (19–21) and subsequently monkey (22, 23) and human (24–30) ESCs into hepatocyte-like cells (HLCs). Additionally, a few studies have demonstrated that mouse or monkey ESCs can be coaxed to generate HLCs in the presence of murine hepatocytes (31, 32) or human liver nonparenchymal cells (33).

\textsuperscript{3} The abbreviations used are: ESC, embryonic stem cell; hESdF, human ESC-derived fibroblast-like cell; HLC, hepatocyte-like cell; iPSC, induced pluripotent stem cell; BMP, bone morphogenetic factor; HGF, hepatocyte growth factor; MEF, mouse embryonic fibroblast; ICC, immunocytochemistry; QRT-PCR, quantitative RT-PCR; IVDS, in vitro differentiation stage; HBV, hepatitis B virus; hESdF-CM, hESdF-conditioned medium; EB, embryoid body.

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\textsuperscript{1} Both authors contributed equally to this work.

\textsuperscript{2} To whom correspondence should be addressed: No. 128, Sec. 2, Academia Rd., Nankang, Taipei, Taiwan. Tel.: 886-2-27899580, Ext. 201; E-mail: kuohuch@gate.sinica.edu.tw.
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Recent demonstrations of the generation of induced pluripotent stem cells (iPSCs) with defined transcription factors (34, 35) that allowed the derivation of patient- and disease-specific pluripotent stem cells (36, 37) without using human embryos have heightened interest in the in vitro hepatic differentiation potential of iPSCs (38–40). The demonstration of the hepatic differentiation capability of iPSCs suggests their possible application for in vitro personalized pharmacogenetics, toxicology, and metabolism studies and future in vivo transplantation trails.

In this study, we explored the potential of our previously established human ESC-derived fibroblast-like cells (hESdFs) (41) for derivation of hepatocytes from pluripotent stem cells. We found that exposure of monkey and human ESCs and human iPSCs to a high density of hESdFs induced robust hepatic endoderm differentiation in a regionally specific manner in ESC/iPSC colonies. Additionally, we elaborate the possible mechanisms responsible for such hepatic induction.

EXPERIMENTAL PROCEDURES

Cell Culture and Differentiation—Mitomycin C-inactivated mouse embryonic fibroblasts (MEFs) and hESdFs were generated as described previously (41). The monkey (ORMES-5 and ORMES-6) and human ESCs (NTU1, NTU2, and NTU3) (3) were grown on MEF feeders (2 × 10⁴ cells/cm²) in DMEM/F-12 medium plus 20% knock-out serum replacement (Invitrogen) and basic FGF (4 ng/ml; Sigma-Aldrich). To induce differentiation, ESCs and iPSCs were grown on a high density of hESdFs (1 × 10⁴ cells/cm²) in the ESC medium.

Immunocytochemistry (ICC)—ICC assay was performed as described previously (41). The antibodies used are listed in supplemental Table S1.

RNA Isolation, RT-PCR, and QRT-PCR—Assays were performed as described previously (41, 42). All of the primers used are listed in supplemental Table S2. All of the QRT-PCRs were performed in triplicate.

Quantitative Measurement of Albumin and Urea—Albumin and urea levels in the cell culture media were measured with ALB-G kits (Denka Seiken, Japan) or the ELISA kit for human albumin (Assaypro) and urea liquid kits (Sentinel Diagnostics), respectively. All of the steps were performed according to the manufacturers’ instructions. Colorimetric determinations were made using Toshiba 200FR (Toshiba).

ELISA—Haptoglobin, activin A, and FGF2 levels in hESdF-conditioned media were determined by ELISA kits for human haptoglobin (Alpcro Diagnostics, Salem, NH), activin A (R & D, Minneapolis, MN), and FGF2 (Sigma) according to the manufacturers’ instructions.

Periodic Acid-Schiff Staining—The cells were fixed in 4% paraformaldehyde, treated with 0.5% periodic acid for 10 min, washed three times with distilled water, and treated with Schiff’s buffer for 20 min. Afterward, the cells were washed with distilled water and stained with hematoxylin.

Ethoxyresorufin-O-deethylase—Ethoxyresorufin-O-deethylase assays were performed as described previously (43). The detailed procedure is included in the supplemental “Materials and Methods”.

Microarray Analysis and Data Processing—Ten μg of total RNA purified by TRIzol (Invitrogen) was used for cDNA synthesis and to generate biotin-labeled cRNA probe, which was hybridized to an Affymetrix Rhesus Macaque genome array (Affymetrix, Santa Clara, CA) by the Affymetrix Gene Expression Service Laboratory at Academia Sinica in Taiwan. The chips were scanned with an Affymetrix GeneChip Scanner 7G, and GeneSpring X software (Agilent, Santa Clara, CA) was used for data mining. Raw data were normalized by robust multichip average, and the weakly expressed signals (<20% of total samples) were excluded. The filtered genes were clustered by Pearson centered complete clustering. The average log intensities of the biological replicates were normalized to the average log intensity of ESCs to obtain the changes in expression between different conditions. Genes highly expressed in the liver, pancreas, or lung were identified with reference to UniGene transcriptome cluster annotation (National Center for Biotechnology Information). The raw microarray data are available through the Gene Expression Omnibus (GEO, GSE19964).

Statistical Analysis—All of the in vitro results were derived from triplicate experiments if not otherwise indicated. The results are presented as the means ± S.D. Grubb’s test was used to detect possible outliers. Student’s t test was used to examine the significance of differences between group means, and differences with p values less than 0.05 were considered significant.

RESULTS

hESdFs Induced and Promoted Regional Differentiation in Monkey and Human ESC Colonies—To see whether hESdFs could support the undifferentiated status of multiple primate ESC lines, we evaluated the growth pattern of monkey ORMES-5 and ORMES-6 (3) and human NTU1, NTU2, and NTU3 ESCs (44) on mitotically inactivated hESdFs. Surprisingly, we found that hESdFs induced the formation of a small flattened area with differentiated cells in the central area of ESC colonies of all examined ESC lines after prolonged co-culture without splitting. Subsequently, the early flattened area expanded, resulting in the formation of differentiated foci containing HLCs (Fig. 1). Moreover, we found that a higher density of hESdFs (5 × 10⁴ cells/cm²) further enhanced the differentiation and eventually led to the formation of differentiated foci containing HLCs in the majority of human and monkey ESC colonies. In contrast, only a few ESC colonies were able to develop into HLC foci in the MEF co-culture group. We divided the differentiation process into three stages according to the morphological changes seen in the differentiated cells along the course of focus formation and expansion. The first 5-day period (days 1–5) after the seeding of ESCs on hESdFs was defined as in vitro differentiation stage (IVDS) 1. The colonies growing at this stage normally contained cells exhibiting characteristic ESC morphology, occasionally with a small differentiated area (Fig. 1). The following 5-day period (days 6–10) was defined as IVDS2. At this stage an extended flattened area with spiky differentiated cells started to emerge in the central region (Fig. 1). At approximately day 15 (IVDS3), differentiated cells exhibiting typical hepatocyte morphological features, such as compact cell-cell contacts, polygonal shapes, and distinctive round
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To further dissect lineage information along the course of differentiation, we performed ICC analysis on monkey and human ESC colonies at different differentiation stages. We found that cells in the central area of ESC colonies expressed a high level of SOX17 at IVDS1 (Fig. 2B). Subsequently at IVDS2, expression of SOX17 and down-regulation of OCT4 were mainly detected in the peripheral ring region of the foci area of ESC colonies (Fig. 2B), whereas cells in the foci expressed SOX17 sporadically and weakly (Fig. 2B). These results suggest that most endoderm-like cells in the foci might have already differentiated into early hepatic cells at IVDS2. Double ICC for SOX17 and the hepatocyte markers AFP or CK18 demonstrated that CK18 and AFP were expressed in the majority of cells in the IVD2 foci, with sporadic cells co-expressing both SOX17 and CK18 or AFP (Fig. 2C). However, the expression of CK18 or AFP was completely absent from the SOX17-expressing cells surrounding the foci (Fig. 2C). Consistent with the RT-PCR results, the differentiation of both mesodermal and ectodermal lineages was mostly suppressed in the presence of hESFs because Brachyury expression was absent in all stages, and only a few occasional SOX1-expressing cells were found outside the foci (Fig. 2B).

To elucidate the spatial-temporal gene expression pattern in the differentiating ESC colonies, we looked at RNA expression from ORMES-6 ESC, cells at IVDS2 and IVDS3, and cells in the central foci (IVDS2-C) and peripheral area (IVDS2-P) of ESC colonies at IVDS2, by cDNA microarray analysis. The global gene expression analysis revealed a dramatic increase in the expression of genes highly expressed in the hepatic lineage at IVDS2 and IVDS3 (Fig. 3, A and B). Hierarchical clustering analysis showed that the IVDS2-C expression pattern was similar to that at IVDS3, whereas the IVDS2-P pattern was closer to ESCs, indicating differences in the fate of cells in the central and peripheral regions of IVDS2 colonies (Fig. 3, A and C). To ascertain whether hESF co-culture can induce nonhepatic endoderm, we compared the expression patterns of subsets of genes highly expressed in the pancreatic or pulmonary endoderm in cells from IVDS2 and IVDS3; we found that the majority of these genes were down-regulated at IVDS2 and IVDS3 (Fig. 3D). Taken together, these results demonstrate that hESFs induced a stepwise endodermal-to-hepatic differentiation of ESCs along with morphological change in the ESC colonies emanating from the central region.

Primate ESCs Gave Rise to Differentiated Cells Exhibiting Mature Hepatic Phenotypes in hESF Co-culture—To determine whether HLCs derived from our hESF culture system displayed mature characteristics of hepatic lineages, we analyzed the expression patterns of early hepatocyte marker genes AFP and HNF4A, mature hepatocyte marker genes, such as Albumin, CK8, CK18, α-1AT, TOT2, and CYP3A4, and cholangiocyte markers CK19 and AQP1 in the HLCs collected from IVDS3 by RT-PCR. All of these genes were highly expressed in the HLCs from both monkey and human ESCs (NTU1 and 3) co-cultured with hESF derived either from H9 (see Fig. 5A and supplemental Fig. S3A) or NTU1 (supplemental Fig. S3B). QRT-PCR assay further demonstrated that albumin expression was dramatically increased at IVDS3 in both human and monkey ESCs (supplemental Fig. S4A). Moreover, hESCs had...
FIGURE 2. hESdFs promote a sequential endodermal-to-hepatic differentiation in human and monkey ESCs. A, QRT-PCR analysis of IVDS 1, 2, and 3 for pluripotency-related markers (Oct4 and Nanog), mesodermal markers (Brachyury-MIXL1), endodermal markers (SOX17, GATA4, and CXCR4), ectodermal markers (SOX1 and PAX6), and hepatocyte markers (AFP, CK8, CK18, albumin, and CYP7A1). B, the expression of Oct4 (pluripotency-related), early ectodermal marker SOX1, early mesodermal Brachyury, and early endodermal SOX17 in differentiating human and monkey ESC colonies of IVDS 1 and 2 were analyzed by ICC. C, co-expression of endodermal marker SOX17 with hepatocyte marker CK18 or AFP is as indicated. IVDS2 foci derived from monkey ESC were stained for SOX17 (red) and CK18 (green) or AFP (green). Scale bar, 25 μm.
increased expression of cholangiocyte markers CK19 and AQP1 from IVDS2 to IVDS3, and monkey ESCs had a higher expression of CK19 at IVDS2 (supplemental Fig. S4A). ICC analysis also showed that the majority of the primate ESC-derived HLCs expressed key hepatocyte proteins, such as AFP, CK7, CK18 (Fig. 4, A and B, and supplemental Fig. S3), and albumin (Fig. 4, A and B, and supplemental Fig. S3). However, only a small subpopulation of the HLCs expressed cholangiocyte marker, CK19 (supplemental Fig. S4A). These results demonstrate that hESdFs derived from different parental ESC lines were able to promote hepatic differentiation in most populations in multiple primate ESC lines.
indicated hepatic markers. The nuclei were stained with DAPI (Fig. 4A). ICC analysis of hepatocyte marker expression in human (NTU1) and monkey (ORMES-6) ESC-derived HLCs showed that the majority of the HLCs were positively stained by the indicated hepatic markers. The nuclei were stained with DAPI (blue). Scale bar, 25 μm. C, temporal production of haptoglobin, albumin, and urea hepatocyte proteins during differentiation and maturation of monkey ESC-derived HLCs under hESdF co-culture conditions. The production of haptoglobin, albumin, and urea in the medium of differentiated cells increased in parallel with differentiation duration (Med, medium; 2W, 2 weeks; 4W, 4 weeks; 8W, 8 weeks; 12W, 12 weeks). The data correspond to the averages and S.D. of triplicate experiments, except the albumin data, which are the averages of four independent experiments. Significant differences between each time point and media are labeled (*, p < 0.05). D, ethoxyresorufin-O-deethylase assay showed that differentiated cells had a higher activity than control cells. E, periodic acid-Schiff staining showed glycogen storage in HLCs (lower panel), whereas the hESdF feeder cells were negative. F, immunoperoxidase staining showed positive nuclear staining of HBcAg (arrows, upper panels) in monkey ESC-derived HLCs infected with serum from patients with HBV infection for 3 days. The feeder cells (left lower panel) were negative. An HBV-infected liver section (right lower panel), stained for both cytoplasmic and nuclear (arrows); HBcAg was the positive control. Scale bar, 25 μm.

Next, we investigated the functional properties of the ESC-derived HLCs. Conditioned media previously incubated with monkey HLCs for 2, 3, 4, 8, and 12 weeks were collected for hepatocyte functional tests, including ureagenesis and haptoglobin and albumin synthesis. These monkey HLCs exhibited urea, haptoglobin, and albumin synthesis that significantly increased in parallel with the duration of hepatic differentiation (Fig. 4C), indicating enhanced hepatic maturation over the course of prolonged culture. In addition to monkey HLCs, we also tested the albumin synthesis ability of HLCs differentiated from human ESCs (NTU1) using ELISA and found that albumin secretion was significantly increased after co-culture with hESdFs for 4 or 8 weeks compared with that without differentiation (supplemental Fig. S5). The increase of albumin in both human and monkey HLCs was comparable with the increase using the three-step differentiation protocol (supplemental Fig. S6). These HLCs also exhibited other characteristics of hepatocytes, including activation of cytochrome P-450 activity (Fig. 4D) and glycogen storage (Fig. 4E), as demonstrated by an ethoxyresorufin-O-deethylase assay and by periodic acid-Schiff staining, respectively. Remarkably, we found that a small number of monkey ESC-derived HLCs could be infected by human hepatitis B virus (HBV), as shown by the positive nuclear staining for HBV core antigen (Fig. 4F). These results demonstrate that this hESdF co-culture system did support the production of mature and functional hepatocytes from both human and monkey ESCs.

**hESdFs Promote Hepatic Differentiation in Human iPSC Cells**—To determine whether human iPSCs are equally as amenable to induction by hESdFs as ESCs, we generated iPSCs by reprogramming human foreskin fibroblasts with methods described previously (34). After various in vitro and in vivo characterizations, three iPSC clones (iPSC-CFB 10, 37, and 50) (supplemental Fig. S7) were selected for experiments. The iPSC colonies treated with hESdFs formed hepatic foci after 10 days of differentiation, although clone 37 was less efficient than in the other two clones (Fig. 5A). RT-PCR showed that the iPSC derived HLCs from IVD3 expressed hepatic genes, such as α-1-AT, CYP3A4, TAT, and TDO2 (Fig. 5B). QRT-PCR analysis showed that the expression levels of hepatic genes varied among the HLCs of different iPSC clones (Fig. 5C), indicating that different iPSC clones responded differently to the stimulation from hESdFs. Finally, ICC analysis further indicated that the majority of the HLCs from all three lines expressed CK18 and AFP, but only HLCs from iPSC-CFB-10 expressed abundant albumin (>70% cells; Fig. 5D).

**hESdFs Produced Factors Supporting Hepatic Differentiation of Primate ESCs**—Next, we reasoned that such hepatic differentiation of ESCs/iPSCs might result from factors produced by hESdFs. Thus, we investigated whether hESdF-conditioned
FIGURE 5. Derivation of human iPSCs and characterization of iPSC-derived HLCs under hESdF co-culture conditions. A, phase contrast images of three human iPSC lines (cell lines 10, 37, and 50), which showed characteristic hepatocyte-like morphology after 3 weeks co-culture with feeder hESdFs. B, RT-PCR analysis of hepatic markers (CK18, CK19, AFP, Albumin, HNF4α, α-1-AT, CYP3A4, CYP7A1, TAT, TDO2, and HGF) expression in the HLCs derived from human ESCs (NTU1) and monkey ESCs (ORMES6) and three different iPSC clones, and in control ESCs (NTU1) without differentiation. C, QRT-PCR analysis of total RNA isolated from HLCs derived from iPSC-CFB-10, -37, and -50 clones for CK8, CK18, HNF4α, α-1-AT, CYP3A4, CYP7A1, TAT, TDO2, and HGF expression. For each sample, relative expression levels were normalized to corresponding levels in human ESCs (NTU1). The data correspond to the averages and standard deviations of triplicate experiments. D, ICC analysis of hepatocyte markers in human iPSC-derived HLCs. HLCs derived from iPSC-CFB-10 clone at day 15 of differentiation were double-stained with anti-human CK18 (green) and AFP (red) or albumin (red) as indicated. The nuclei were stained with DAPI (blue). Scale bar, 25 μm.

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We found that medium (hESdF-CM) alone could induce ESC differentiation. Furthermore, QRT-PCR analysis (Fig. 6C) showed that SOX1 was up-regulated in the regenerating cells, but the expression levels of endodermal and mesodermal marker genes were dramatically reduced compared with their counterparts in the colonies without excision (Fig. 6C). These results indicated that removal of the central area of ESC colonies in hESdF co-culture conditions likely disrupted the progress of hepatic endoderm differentiation and converted the differentiation direction from endoderm to ectoderm and therefore highlight the critical role of central region cells for hepatic endoderm differentiation.

DISCUSSION

Several protocols (22–30) have been published that claim to effectively coax human or monkey ESCs to differentiate into hepatocyte foci (Fig. 7A) and instead resulted in emergence of SOX1-expressing neural rosettes (Fig. 7B). Furthermore, QRT-PCR analysis (Fig. 7C) showed that SOX1 was up-regulated in the regenerating cells, but the expression levels of endodermal and mesodermal marker genes were dramatically reduced compared with their counterparts in the colonies without excision (Fig. 7C). These results indicated that removal of the central area of ESC colonies in hESdF co-culture conditions likely disrupted the progress of hepatic endoderm differentiation and converted the differentiation direction from endoderm to ectoderm.

FGF2 and/or activin A in hESdFs by lentiviruses stably expressing short hairpin RNAs against human FGF2 and/or activin A transcripts and grew ORMES-6 ESCs on the modified hESdFs (Fig. 6, C and D). Under these conditions, the typical morphological change and hepatocyte focus formation induced by hESdFs were significantly suppressed at IVDS3 (Fig. 6F). Furthermore, FGF2 and/or activin A knockdown in hESdFs also reduced the expression level of endoderm and hepatocyte marker genes in the differentiating ESCs at IVDS3 (Fig. 6F). Taken together, these experiments indicate that factors produced by hESdFs were directly responsible for hepatic development in primate ESCs.

The Central Region of Colonies Is Constitutionally Essential for Hepatocyte Differentiation—In our system, hepatocyte differentiation always followed focus formation in the central region of ESC colonies. To explore how the central foci formation affects the subsequent hepatic differentiation, we manually removed the central foci area (composed of ~20–40% of the monkey ESC colonies) at IVDS2 and investigated the growth pattern of the excised colonies (Fig. 7A). The empty central region recovered after 3 days by replacement of cells from the surrounding region (Fig. 7A). Interestingly, further co-culturing the recovered colonies with hESdF failed to regenerate the hepatocyte foci (Fig. 7A) and instead resulted in emergence of SOX1-expressing neural rosettes (Fig. 7B). Furthermore, QRT-PCR analysis (Fig. 7C) showed that SOX1 was up-regulated in the regenerating cells, but the expression levels of endodermal and mesodermal marker genes were dramatically reduced compared with their counterparts in the colonies without excision (Fig. 7C). These results indicated that removal of the central area of ESC colonies in hESdF co-culture conditions likely disrupted the progress of hepatic endoderm differentiation and converted the differentiation direction from endoderm to ectoderm and therefore highlight the critical role of central region cells for hepatic endoderm differentiation.
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hESdF secreted factors promoting hepatic differentiation in pluripotent stem cells. A, QRT-PCR analysis of hepatic markers TAT, AFP, albumin, TDO2, and CYP3A4 in ESCs cultured under EB-based differentiation conditions (29), three-step culture conditions, and hESdF co-culture conditions and in hESdF-CM (27). The cells were harvested for analysis 10 days post-differentiation. Gene expression levels are presented as the relative change of the 2^{-ΔΔCT} value relative to that of day 10 EB-based differentiation conditions. The data correspond to the averages and standard deviations of triplicate experiments. B, RT-PCR analysis of BMP4, EGF, FGF2, WNT3A, FGF4, activin A, Nodal, HGF, oncostatin M (OSM), and VEGF in hESdFs. Actin was used as a loading control, and a reaction without reverse transcriptase (-RT) was applied to rule out genomic DNA contamination. C and D, ELISAs for activin A (C) and FGF2 protein (D) expression in conditioned media derived from MEFs, hESdFs, and hESdFs transduced with short hairpin RNA specific to INHBA (shInA) and/or FGF2 (shFGF2). E, QRT-PCR analysis for hepatic markers in monkey ESCs co-cultured with activin A and/or FGF2 knockdown hESdFs for 15 days. Gene expression levels relative to those in hESdF co-cultured cells are shown. F, morphology of monkey ESC colonies co-cultured with hESdFs and shInA and/or shFGF2 hESdFs for 15 days. The number and size of hepatic foci were dramatically reduced in shFGF2 and/or shInA hESdF group(s). Small hepatic foci (arrows) were occasionally found in the ESC colonies cultured in the modified hESdFs.

hepatocytes. The majority of the protocols described involve an initial differentiation step of embryoid body (EB) formation (22, 23, 25, 28, 29), and others include direct differentiation of ESCs on MEF (24, 30) or Matrigel-coated plates (26, 27). Subsequent strategies mostly involve multiple culture steps with sequential supplements of small molecules and recombinant growth factors to the culture medium. In this study, we demonstrate that the hESdF co-culture system is at least as efficient for hepatocyte differentiation as the previously published protocols (22–30). EB formation has been the most popular method exploited to differentiate ESCs into many cell types including hepatocytes; however, the uncontrolled nature of EB differentiation has made this approach less likely to generate scalable specific cell types with satisfactory purity. Monolayer-adherent culture conditions, on the other hand, have been used successfully to direct primate ESC differentiation into various cell types including hepatocytes (26, 27).

In this study, the efficient hepatocyte differentiation induced by the hESdF co-culture system required ESC/iPSCs to be cultured as a monolayer adhering to the culture surface. The monolayer-adherent culture helped to maintain the structural integrity of the two-dimensional ESC colonies and therefore possibly provided a foundation for establishing an unknown feature in the central region of ESC/iPSC cell colonies that is susceptible to differentiation signal(s) secreted by hESdFs.

We showed that FGF2 and activin A secreted by hESdFs played critical roles in mediating the hepatic endoderm differentiation in primate pluripotent stem cells. This finding is well supported by results of previous reports, indicating that activin A and FGF2 have positive effects in promoting ESCs to differentiate into endodermers (28, 33). Unlike most protocols involving supplementation of extraneous factors (28, 33), it is not possible to manipulate the expression level of the factors secreted by hESdFs along the course of differentiation.

However, we still observed efficient hepatocyte differentiation with the presence of mature hepatocytes. Therefore, the persistent presence of factors such as activin A and FGF2, which are important for early stages of differentiation, may not affect the subsequent differentiation and maturation of hepatocytes. Additionally, other growth factors secreted by hESdFs (Fig. 5B), such as BMP4, EGF, and VEGF, which have been shown to be required for hepatic specification of mouse ESC-derived endoderm (21, 45) and to enhance hepatocyte specification of ESCs, respectively, were very likely responsible for the hepatic specification, differentiation, and maturation following the endodermal differentiation mediated by hESdF secreted FGF2 and activin A (21). Finally, the nonparenchymal cells generated along the hepatic lineages differentiation may also contribute to the differentiation and maturation process by secreting growth factors, because HGF expression was normally detected in hepatic foci after 3 weeks of differentiation (Fig.
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5B). Taken together, these results strongly suggest that a combination of hESdF and differentiated ESC secreted factors might be responsible for the hepatocyte differentiation and maturation in our system.

In the hESdF co-culture system, the central regions of ESC colonies always transformed into endoderm-like foci, a step that preceded hepatocyte differentiation. Removal of central foci disrupted the progress of hepatic endoderm differentiation and resulted in the acquisition of ectodermal fate by the regenerated cells. We can only speculate about why the central region of the ESC colonies appears so important for hepatic differentiation. It seems unlikely that the hESdF feeders created a growth factor/cytokine signal gradient along the central to peripheral axis of the ESC colonies, because the differentiation foci could still be readily induced in hESdF-conditioned medium alone.

Alternatively, there might be heterogeneous populations of pluripotent cells in ESC colonies that are assembled in specific topological compartments in the two-dimensional ESC/iPSC colonies; perhaps only those allocated to the central region of ESC/iPSC colonies could properly respond to hESdF-secreted factors to form hepatic foci. This hypothesis is consistent with a recent observation showing that subpopulations of ESCs with distinct tissue-specific fates can be selected from pluripotent cultures (46) and was supported by our microarray analysis comparing the gene expression patterns of the central and rim regions of the ESC colonies (Fig. 3). Future identification of the molecular mechanism by which the distinct hepatocyte-promoting capability of the central region of ESC colonies is established may lead to more efficient control of hepatic differentiation.

Consistent with a previous study (40), we found that different iPSC lines have different capabilities to generate hepatic lineages in response to differentiation stimuli (e.g. hESdF treatment), even though they are all regarded as fully reprogrammed iPSC lines according to currently used standards (supplemental Fig. S7) for iPSC characterization. These discrepancies may be attributed to currently unknown factors that modulate hepatocyte differentiation, which were somehow defective in some of the iPSC lines. It will be of interest to further explore the mechanisms that constitute the discrepancies among different iPSC clones.

In summary, we have developed a simple method that allows scalable hepatocyte generation from primate pluripotent stem cells. Furthermore, we provide compelling evidence indicating that hESdF-secreted factors are responsible for the topology-dependent differentiation of endodermal cells to hepatocytes in ESC/iPSC colonies. These findings provide opportunities to further explore the mechanisms underpinning hepatic lineage differentiation and may be useful for future pharmacotoxicology and transplantation applications.

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