

Blocking autocrine VEGF signaling by sunitinib, an anti-cancer drug, promotes embryonic stem cell self-renewal and somatic cell reprogramming

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Maintaining the self-renewal of embryonic stem cells (ESCs) could be achieved by activating the extrinsic signaling, i.e., the use of leukemia inhibitory factor (LIF), or blocking the intrinsic differentiation pathways, i.e., the use of GSK3 and MEK inhibitors (2i). Here we found that even in medium supplemented with LIF, mESCs still tend to differentiate toward meso-endoderm lineages after long-term culture and the culture spontaneously secretes vascular endothelial growth factors (VEGFs). Blocking VEGF signaling with sunitinib, an anti-cancer drug and a receptor tyrosine kinase (RTK) inhibitor mainly targeting VEGF receptors (VEGFRs), is capable of maintaining the mESCs in the undifferentiated state without the need for feeder cells or LIF. Sunitinib facilitates the derivation of mESCs from blastocysts, and the mESCs maintained in sunitinib-containing medium remain pluripotent and are able to contribute to chimeric mice. Sunitinib also promotes iPSC generation from MEFs with only Oct4. Knocking down VEGFR2 or blocking it with neutralizing antibody mimicks the effect of sunitinib, indicating that blocking VEGF/VEGFR signaling is indeed beneficial to the self-renewal of mESCs. We also found that hypoxia-inducible factor alpha (HIF1 α) and endoplasmic reticulum (ER) stress are involved in the production of VEGF in mESCs. Blocking both pathways inhibits the expression of VEGF and prevents spontaneous differentiation of mESCs. Interestingly, LIF may also exert its effect by downregulating HIF1 α and ER stress pathways and subsequent VEGF expression. These results indicate the existence of an intrinsic differentiation pathway in mESCs by activating the autocrine VEGF signaling. Blocking VEGF signaling with sunitinib or other small molecules help to maintain the mESCs in the ground state of pluripotency.

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Introduction

The two essential properties of embryonic stem cells (ESCs), self-renewal and pluripotency, are regulated by a network of signaling from intrinsic factors and extrinsic stimulants such as cytokines and growth factors. Vari-

ous culture conditions, including the use of feeder cells, conditioned media, growth factors, and serum have been used to derive and maintain ESCs [1]. Leukemia inhibitory factor (LIF) is among the earliest cytokines discovered to be able to promote self-renewal of mESCs [2]. LIF binding leads to the dimerization of gp130 and LIF receptor and the activation of Jak/Stat3 signaling pathway. In combination with serum or bone morphogenetic proteins (BMPs), which induce the expression of proteins that suppress differentiation [3], LIF is able to maintain the self-renewal of mESCs. However, these cytokines fail to block the activation of ERK, a known pathway involved in the differentiation of mESCs [4]. This leads

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to the discovery that the combination of small-molecule inhibitors of ERK and GSK3 pathways is able to bypass the need of cytokine signaling and promote the self-replication of mESCs [5]. PD0325901 (MEK inhibitor) and CHIR99021 (GSK3 inhibitor), the so-called 2i, are now commonly used for culturing mESCs and induced pluripotent stem cells (iPSCs). In addition, other MEK or GSK3 inhibitors, such as pluripotin and 6-bromindirubin-3'-oxime (BIO) [6], and PKC inhibitors [7] have also been reported to promote the self-renewal of mESCs. However, the involvement of other signaling pathways in maintaining the balance between stemness and differentiation remains illusive and the demands for better and easier culture conditions remain high.

Vascular endothelial growth factors (VEGFs) are potent mediators of vasculogenesis in the fetus and angiogenesis in the adults. In mammals, five VEGF ligands have been identified so far. These ligands bind in an overlapping pattern to three receptor tyrosine kinases (RTKs), designated VEGF receptor-1, -2 and -3 (VEGFR1-3), and to co-receptors [8]. In mouse, at least three isoforms of VEGF (VEGFA, VEGFB and VEGFC) have been found and displayed different affinities for the known VEGF receptors and co-receptors, suggesting different roles of these VEGF isoforms [9].

Many early studies have shown that the VEGF and VEGFR family play critical roles in cardiovascular, haematopoietic and lymphatic development. Inactivation of a single *VEGFA* allele [10] or deletion of *VEGFR2* in mice [11] results in early lethality at embryonic stage due to deficient endothelial cell development and lack of vessels. In contrast, overexpression of *VEGFA* results in aberrant heart development, which also leads to lethality at E12.5-14 [12]. Haematopoietic and endothelial cells have been proposed to share a common early progenitor that expresses VEGF signaling system. Indeed, the survival [13] and migration [14] of haematopoietic stem cells are promoted by VEGF and VEGFRs. *In vitro*, VEGF has been reported to induce endothelium and cardiac myocyte differentiation from ESCs or iPSCs [15, 16]. VEGF-mediated angiogenesis is activated during tumor formation [17]. This leads to the development of anti-VEGF antibodies and small-molecule RTK inhibitors that target VEGFRs for tumor therapy. Anti-angiogenic drugs targeting the VEGF signaling pathways, including bevacizumab, sorafenib, sunitinib, and have been approved by FDA for the treatment of advanced carcinoma.

In this study, we sought to dissect the roles of autocrine VEGF/VEGFR signaling in the fate determination of mESCs and the generation of iPSCs and discuss the possible application of VEGFR inhibitor sunitinib in stem cell culture.

Results

mESCs tend to differentiate toward meso-endoderm lineages and produce VEGF during long-term culture

It is generally adopted that ESCs have to be cultured in special medium and passaged every 2-3 days to help maintain the pluripotency of the cells [3, 18]. Without passage, E14 cells cultured in mES medium for a long period tend to spontaneously differentiate even in the presence of LIF and with daily medium change (Figure 1A). On day 5, E14 cells started to lose the typical compact colony morphology of mESCs (Figure 1A). Similar phenomenon was also observed in E14 cells cultured in N2B27 medium supplemented with LIF or mES medium without LIF (Supplementary information, Figure S1A). The spontaneous differentiation was more obvious and occurred earlier (day 3) in mES medium without LIF (Supplementary information, Figure S1A). Quantitative PCR also confirmed the downregulation of pluripotency genes *Oct4*, *Sox2*, *Nanog* and *Rex1* (Figure 1B). Among the lineage-specific genes, the ectoderm markers *Nestin* and *Sox1* were not significantly altered during the spontaneous differentiation, while the mesoderm markers *Brachyury* and *SMA* and the endoderm markers *GATA4* and *Sox17* were all significantly increased (6.7-, 43.8-, 19- and 6.7-fold increase on day 8 relative to those on day 1, respectively; Figure 1C). The downregulation of epithelial cell markers *E-cadherin*, *EP-CAM*, *Ocln* and *Cldn3* and the upregulation of mesenchymal cell markers *N-cadherin*, *Snail*, *Slug* and *Fn* indicated that the epithelial-mesenchymal transition (EMT) occurs during long-term culture of mESCs (Figure 1D).

The above data suggested that mESCs tend to differentiate toward meso-endoderm lineages spontaneously during long-term culture. Therefore, we investigated additional genes and found that several genes involved in the vascular differentiation in mesoderm, including *VEGF*, *VE-cadherin* and *PECAM*, were all upregulated (Figure 1E). mRNA of *VEGFA* increased more than that of *VEGFB* (Figure 1F) after long-term culture and VEGFA protein secreted into the mES media (with LIF) also increased significantly (Figure 1G). Similar increase of *VEGF* expression was also observed in mESCs cultured in the serum-free N2B27 media (with LIF; Supplementary information, Figure S1B). The expression levels of a panel of growth factors were analyzed in E14 cells cultured in mES media (with LIF) for 1-8 days (Supplementary information, Figure S1C). Many growth factors were not detectable. A couple of growth factors, such as *FGF4* and *PDGFA*, did not show time-dependent change, even though their expression levels were relatively high. *VEGFA* expressed at a relatively high level and displayed

most striking upregulation (~6-fold) after 5 days of culture. Another growth factor, *PDGFB*, was also upregulated (~3-fold) in long-term culture, although its expression level was relatively low.

Addition of the recombinant protein mVEGF₁₆₄ or

PDGFBB induced loss of mESC morphology on day 3, 2 days earlier than the spontaneous differentiation (Figure 1H and Supplementary information, Figure S2A), indicating that activation of both pathways leads to differentiation. However, knockdown of *VEGFA* but not *PDGFB*

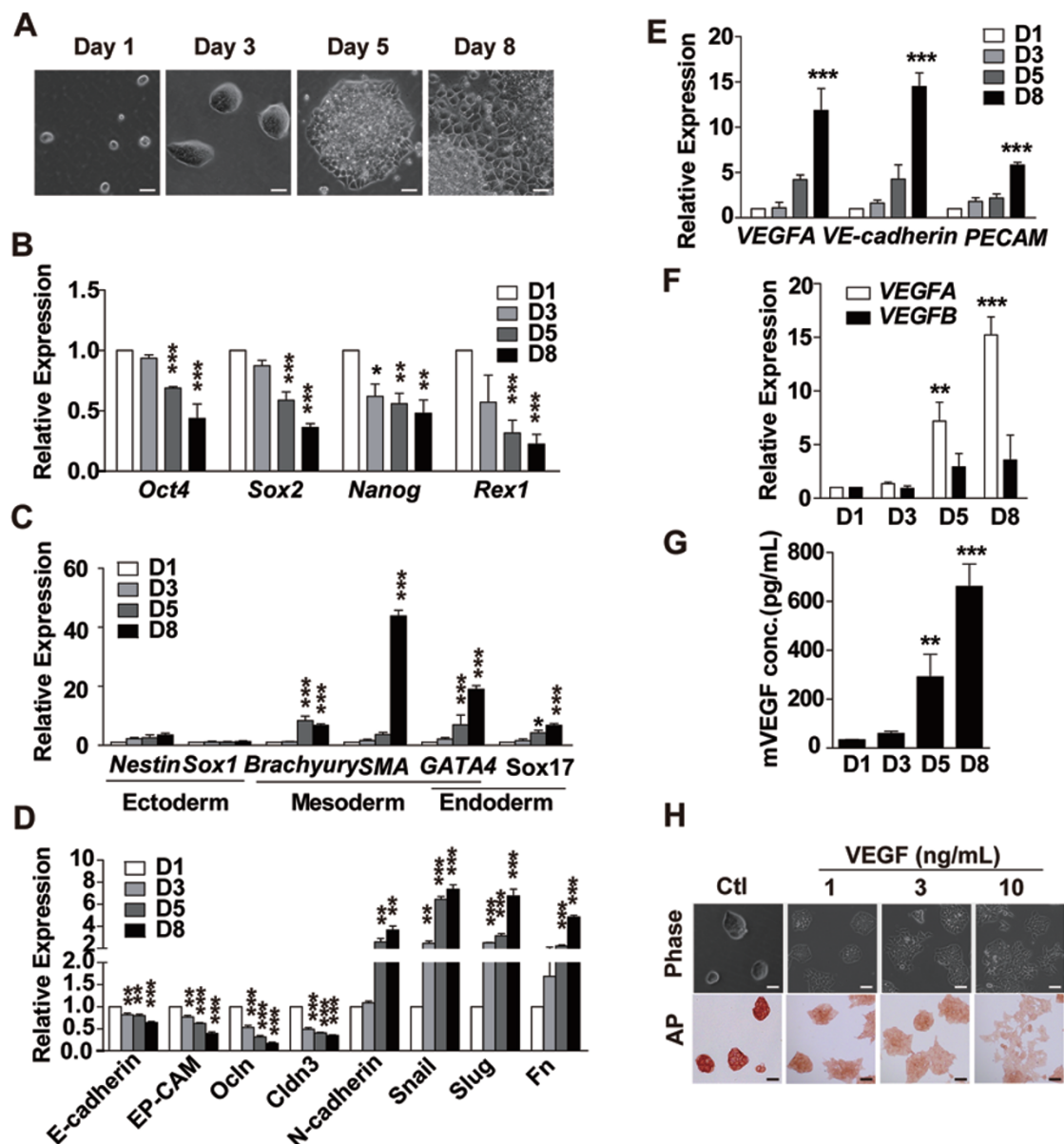


Figure 1 mESCs differentiate toward meso-endoderm lineage spontaneously during long-term culture. **(A)** Morphology of E14 cells cultured in mES medium containing 1 000 U/ml LIF without passage for 1-8 days. Medium was changed every day. Scale bar, 50 μ m. **(B-F)** qRT-PCR analysis of pluripotency genes **(B)**, lineage-specific genes (mesoderm: *Brachyury* and *SMA*, endoderm: *GATA4*, *Sox17*, ectoderm: *Nestin* and *Sox1*) **(C)**, genes involved in EMT **(D)**, vascular marker genes **(E)** and VEGF genes **(F)** in mESCs described in **A** on days 1, 3, 5 and 8. **(G)** ELISA analysis of VEGF secretion from mESCs described in **A**. **(H)** VEGF-induced morphology and alkaline phosphatase (AP) change in E14 cells. Scale bar, 50 μ m. E14 cells were cultured in mES medium containing 1 000 U/ml LIF for 48 h, then vehicle or 1, 3, 10 ng/ml mVEGF₁₆₄ was added into the medium for another 48 h prior to analysis. Data are mean \pm SEM ($n = 3$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ vs day 1. Independent experiments were repeated at least three times.

with shRNA in E14 cells significantly blocked the differentiation induced by LIF withdrawal (Supplementary information, Figure S2B-S2E), suggesting that PDGFB might induce differentiation in a paracrine fashion, while VEGFA in an autocrine way. Taken together, these data suggest that during long-term culture, mESCs secrete growth factors such as VEGFA, which promote cells to differentiate toward meso-endoderm lineages.

Blocking VEGF receptors with sunitinib promotes mESC self-renewal in the absence of LIF and de novo derivation of pluripotent ESCs

VEGF might promote spontaneous differentiation of mESCs in an autocrine way, thus we investigated whether blocking VEGF signaling with small-molecule inhibitors would help to maintain the self-renewal of mESCs. In mES media deprived of LIF, E14 cells showed spontaneous differentiation on day 3 (Figure 2A). 2i (PD0325901 and CHIR99021) effectively maintained the mESC morphology up to day 5 without passage, while sunitinib, a multi-targeted RTK inhibitor mainly inhibiting VEGF and PDGF receptors, showed a similar effect in maintaining the self-renewal of mESCs (Figure 2A). Dose-response study indicated that 1 μ M sunitinib was best in maintaining the undifferentiated state of mESCs in LIF-free condition (Supplementary information, Figure S3A). In the presence of LIF, both sunitinib and 2i were able to maintain the mESC morphology up to day 8 without passage (Supplementary information, Figure S3B). In the absence of LIF, mESCs could be maintained for 3-4 passages, even though the alkaline phosphatase (AP) staining results revealed the loss of pluripotency at passage 1 (Figure 2B). Both sunitinib and 2i helped to maintain the mESC morphology and positive AP staining beyond passage 10 (Figure 2B). Immunocytochemistry data showed that mESCs maintained in sunitinib⁺ LIF⁻ medium expressed high levels of pluripotency markers, including Oct4, Nanog and SSEA1, in passages up to 10, while cells in the sunitinib⁻ LIF⁻ medium completely lost the expression of these genes at passage 4 (Supplementary information, Figure S4A). Quantitative PCR results revealed that both sunitinib and 2i almost completely reversed the upregulation of lineage-specific genes, including *SMA*, *Desmin*, *Nestin* and *VE-cadherin*, and the loss of pluripotency genes, such as *Oct4*, *Sox2*, *Nanog* and *Rex1*, caused by the removal of LIF (Supplementary information, Figure S4B and S4C). And LIF deprivation-induced changes in the EMT-related genes, including *E-cadherin*, *EP-CAM*, *Ocln*, *Cldn3*, *N-cadherin*, *Snail*, *Slug* and *Fn*, were also blocked by both sunitinib and 2i (Supplementary information, Figure S4D). Sunitinib (1 μ M) was also very effective in maintaining the

self-renewal of iPSCs generated from OG2 mice carrying a transgenic GFP gene driven by *Oct4* promoter in the absence of LIF (Supplementary information, Figure S3C).

To confirm the intact pluripotency, E14 cells cultured in medium containing sunitinib, 2i or LIF (passage 11) were induced to form embryoid body (EB) after removal of sunitinib, 2i or LIF. No difference was observed in EB formation between cells previously maintained in different media. qRT-PCR analysis revealed the upregulation of lineage-specific genes and downregulation of pluripotent genes in EBs formed by E14 cells previously maintained in all three media (Supplementary information, Figure S5A). For immunofluorescent staining, EBs were reseeded into gelatin-coated plates and cells spontaneously differentiating into three germ layers could be identified (Supplementary information, Figure S5B). Stimulation by mVEGF₁₆₄ (50 ng/ml) promoted EB differentiation toward vascular cells with increased expression of vascular lineage markers, including SMA, PECAM and VE-cadherin (Supplementary information, Figure S5C and S5D). For *in vivo* differentiation, mESCs from passage 11 were injected into the thigh of SCID mice. Teratoma formation was observed after 3-4 weeks. H&E staining revealed that these tumors contain tissues derived from all three germ layers, including gastrointestinal-like epidermis structure (ectoderm), cartilage-like structure (mesoderm) and neural tube-like structure (endoderm; Figure 2C and Supplementary information, Table S1). Sunitinib seems to exert its anti-differentiation effect by blocking VEGF-induced ERK and GSK3 β activation (Supplementary information, Figure S6A) rather than by direct activation of the Stat3 pathway like LIF (Supplementary information, Figure S6B).

Since the E14 cells were derived from 129/Ola mice with cream coat color, it would be difficult to assess the chimerism between E14 cells and ICR mice (with white coat color). Therefore, we generated mESCs from the blastocysts of C57BL/6 mice, which have black coat color, and cultured them in mES medium containing sunitinib (1 μ M) and 2 000 U/ml LIF. mESCs were later expanded in mES medium containing only sunitinib (1 μ M) (Figure 2D). Cells from passages 6 and 11 were injected into the blastocysts of ICR mice, and chimeras were generated successfully (Figure 2E). Taken together, these data indicate that blocking VEGF signaling by sunitinib help to maintain the pluripotency of mESCs, and the effect is comparable to 2i.

Knocking down VEGFR promotes long-term self-renewal of mESCs in the absence of LIF

Although mainly targeting VEGFRs and PDGFRs,

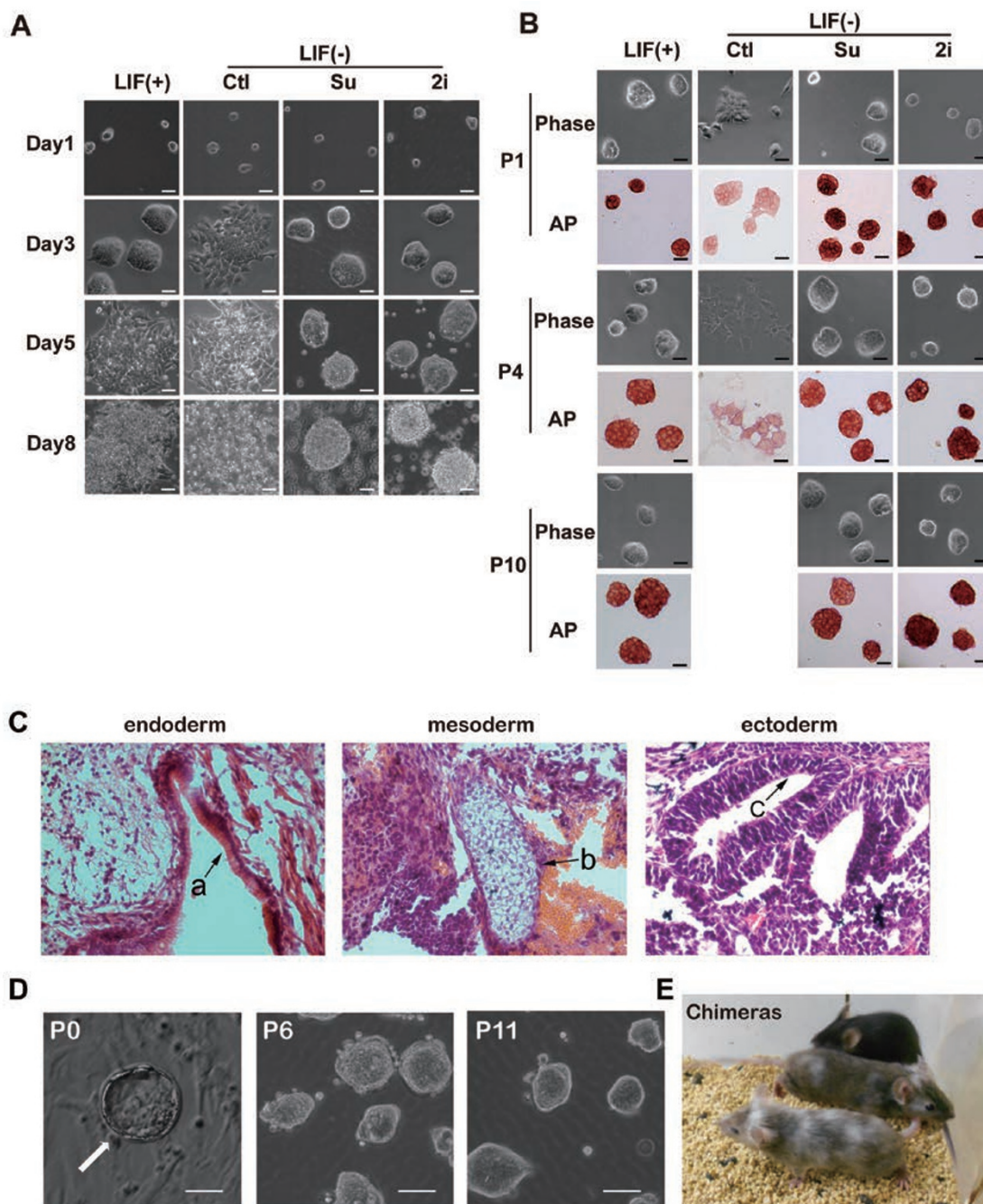
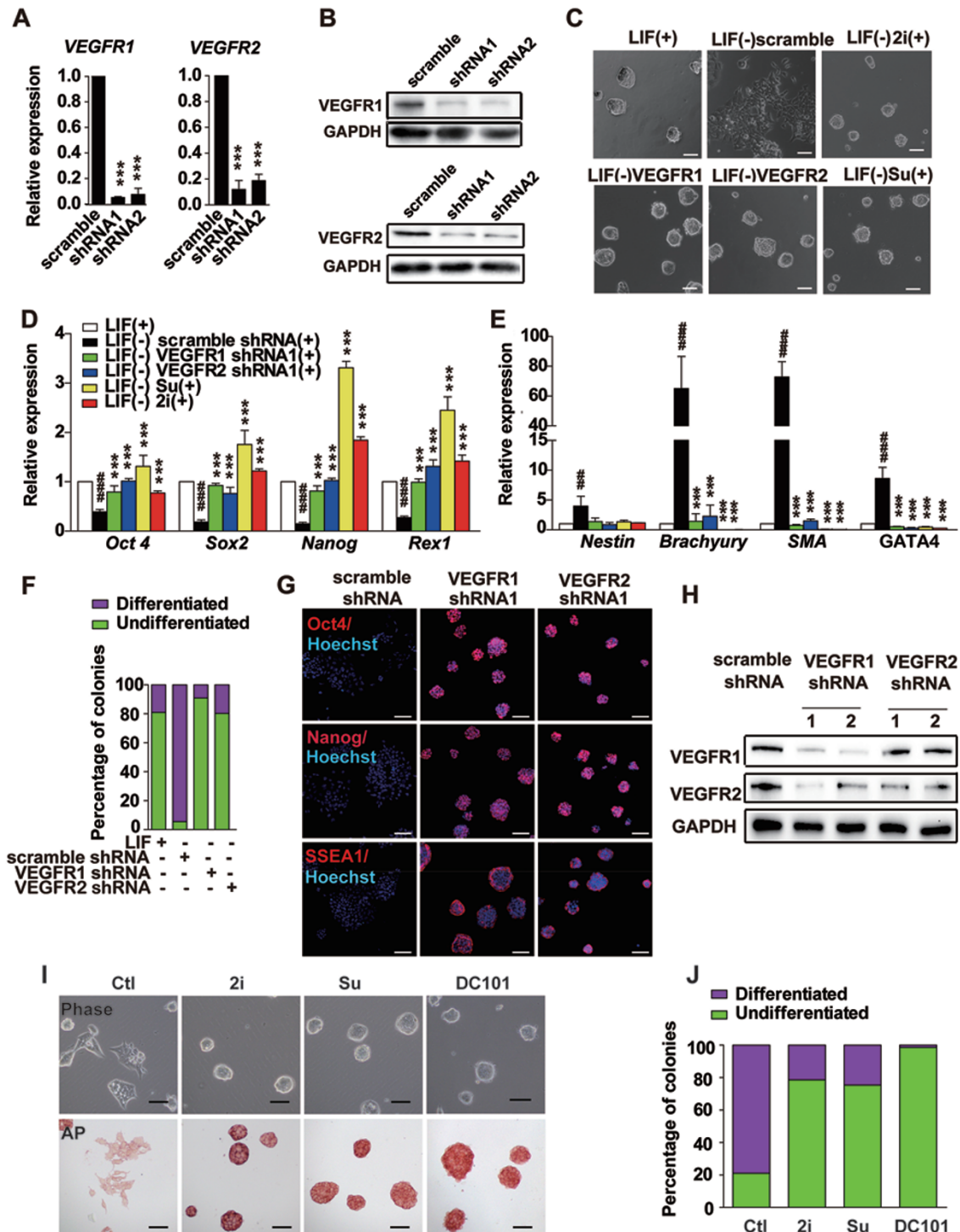


Figure 2 Sunitinib maintains the self-renewal and pluripotency of mESCs. **(A)** Morphology of E14 cells cultured in mES media containing 1 000 U/ml LIF, or in mES media without LIF but containing 2i (3 μ M PD0325901 and 3 μ M CHIR99021) or sunitinib (Su, 1 μ M), for 1-8 days. Media were changed every day. Scale bar, 50 μ m. **(B)** Morphology and alkaline phosphatase (AP) staining of E14 cells at passages 1, 4 and 10 maintained in mES media containing 1 000 U/ml LIF, or in mES media without LIF but containing 2i (3 μ M PD0325901 and 3 μ M CHIR99021) or sunitinib (1 μ M). Scale bar, 50 μ m. **(C)** H&E staining of teratomas generated with E14 cells maintained with sunitinib (passage 11). Typical structures of the three embryonic germ layers are shown as follows: a, gastrointestinal-like epithelium (endoderm); b, cartilage-like structure (mesoderm); c, neural tube-like structure (ectoderm). Scale bars, 20 μ m. **(D)** Blastocyst outgrowth (P0, the arrow) on feeder cells and mES medium containing sunitinib and 2 000 U/ml LIF. mESCs were subsequently maintained in mES media containing sunitinib for multiple passages. Scale bar, 20 μ m. **(E)** Chimeric mice generated with mESCs derived and maintained with sunitinib (passage 6).

sunitinib also blocks other RTKs including KIT, FLT3, and RET [19]. To avoid the possible off-target effect of sunitinib, we designed shRNAs specifically targeting *VEGFR1* and *VEGFR2*, two main receptors conveying VEGFA signaling [20]. Both receptors were expressed in mESCs and the effectiveness of shRNAs was confirmed

with quantitative RT-PCR and western blot analysis (Figure 3A and 3B). When LIF was deprived for 2 days, E14 cells expressing scrambled shRNA almost completely lost the compact colonial morphology, while knocking down either *VEGFR1* or *2* helped maintain the mESC-like morphology (Figure 3C). The reduced expression of



pluripotency genes and the elevated expression of lineage-specific genes caused by LIF removal could also be reversed by knockdown of either *VEGFR1* or *2* (Figure 3D and 3E). About 80% of E14 colonies maintained in LIF-containing medium were AP positive, while < 10% of E14 cells expressing scramble shRNA remained to be AP positive after LIF removal (Figure 3F). Knocking down either *VEGFR1* or *2* restored the percentage of AP-positive colonies to more than 80% (Figure 3F). Immunocytochemistry data also showed that VEGFR-knockdown E14 cells (passage 4) maintain high levels of pluripotency markers, including Oct4, Nanog and SSEA1, in LIF-free mES medium, while the scramble shRNA did not prevent the loss of all these markers (Figure 3G). It was intriguing to find that knocking down either *VEGFR1* or *2* almost completely resembled the effect of sunitinib and blocked the spontaneous differentiation induced by LIF removal. It has been reported that VEGFR1 is a critical determinant of VEGFR2 abundance, while VEGFR2 is the key receptor directly responsible for endothelial cell signaling stimulated by VEGF [21]. Therefore, we examined whether knockdown of one VEGFR affect the level of the other VEGFR. In agreement with the previous report, knocking down *VEGFR1* also reduced the expression of *VEGFR2*, but knocking down *VEGFR2* did not affect the expression of *VEGFR1* (Figure 3H).

To further confirm the role of VEGFR2 in mESC differentiation, DC101, a neutralizing antibody of VEGFR2 [22], was tested along with 2i and sunitinib in mESCs cultured without LIF. As demonstrated in Figure 3I and 3J, DC101 blocked mESC differentiation induced by LIF withdrawal as effectively as sunitinib and 2i. Taken together, these data indicate that autocrine activation of VEGFRs, especially VEGFR2, in mESCs is one of the major forces that promote the exit from self-renewal and entry into lineage commitment.

Hypoxia-inducible factor alpha and endoplasmic reticulum stress are involved in the VEGF production in mESCs during long-term culture

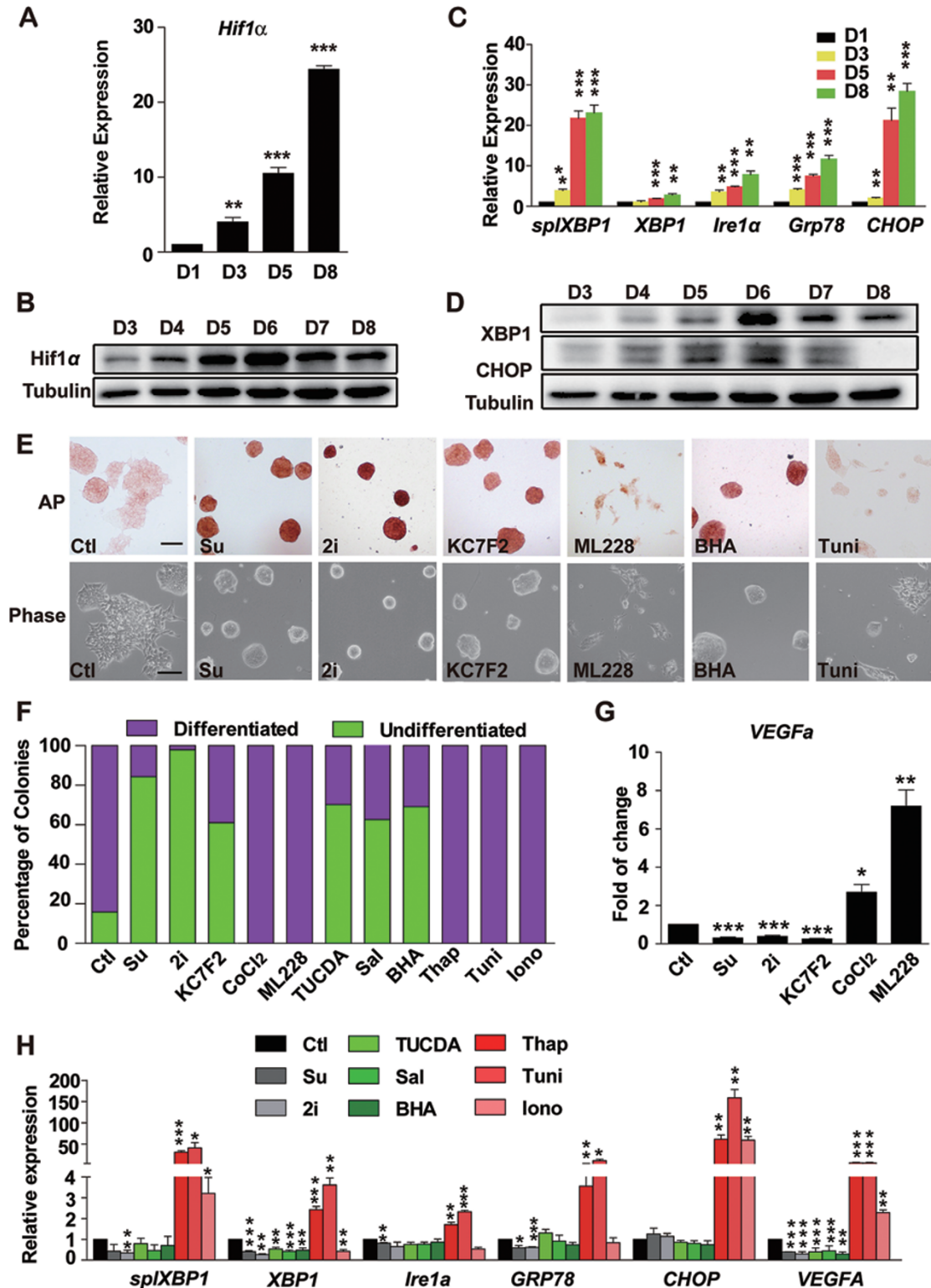
Hypoxia-inducible factor alpha (HIF1 α) is one of the major factors that regulate VEGF expression and angiogenesis in cancers [23]. There is still controversy in the roles of hypoxia in maintaining the self-renewal of ESCs. Some have reported that hypoxia promotes ESC propagation [24] and facilitates the generation of iPSCs [25], while others found that hypoxic priming of mESCs accelerates vascular-lineage differentiation through HIF1-dependent mechanisms [26, 27]. Nevertheless, we found that both the mRNA and protein level of HIF1 α increased significantly on day 3 during the long-term culture without passage but with daily medium change (Figure 4A and 4B). In LIF-free mES media, HIF1 α inducers CoCl₂ [28] and ML228 [29] led to elevated expression of *VEGFA* and promoted cell differentiation further (Figure 4E-4G). In contrast, KC7F2 [30], a selective HIF1 α inhibitor, reduced the expression of *VEGFA* (Figure 4G) and blocked the spontaneous differentiation caused by the removal of LIF for 2 days (Figure 4E and 4F). Sunitinib and 2i also reduced the expression of *VEGFA* (Figure 4G) and differentiation induced by LIF deprivation (Figure 4E and 4F).

Endoplasmic reticulum (ER) stress is another major pathway that regulates the expression of VEGF under adverse environmental conditions [31]. It has been reported that hESCs must resolve ER stress related to culture conditions to survive and maintain pluripotency [32]. ER stress has also been reported to participate in ESC differentiation induced by retinoic acid [33]. Therefore, we further investigated the expression of genes related to ER stress during long-term culture of mESCs. Quantitative PCR indicated that the expression of several genes involved in ER stress, including *XBPI* and *splXBPI* (an alternative splicing form of *XBPI* in re-

Figure 3 VEGFR knockdown recapitulates the effect of sunitinib. **(A, B)** Validation of the shRNAs targeting VEGFRs by qRT-PCR and western blot. **(C)** Morphology of E14 cells expressing the indicated shRNA cultured in mES media without LIF. Cells treated with sunitinib or 2i were used as positive control. Scale bar, 50 μ m. **(D, E)** qRT-PCR analysis of pluripotency genes (*Oct4*, *Sox2*, *Nanog*, *Rex1*) **(D)** and lineage-specific genes (mesoderm: *Brachyury* and *SMA*, endoderm: *GATA4*, ectoderm: *Nestin*) **(E)** in cells expressing the indicated shRNA cultured in mES media without LIF (passage 4). Cells treated with sunitinib or LIF were used as positive control. Data are mean \pm SEM ($n = 3$). $^{###}P < 0.01$, $^{####}P < 0.001$ vs LIF (+) condition; $^{*}P < 0.05$, $^{**}P < 0.01$, $^{***}P < 0.001$ vs scramble shRNA. **(F)** E14 cells expressing the indicated shRNA were cultured in LIF-free mES media for 3 days at an initial density of 10 000 cells/well in 24-well plate. The percent of undifferentiated and differentiated colonies were calculated according to AP staining and morphology. Independent experiments were repeated at least three times. **(G)** Immunofluorescence staining of pluripotency markers (Oct4, Nanog, SSEA1) in cells described in **E** (passage 4). Cell nuclei were stained with Hoechst. Scale bar, 80 μ m. **(H)** Western blot analysis of VEGFR1 and 2 in cells transfected with scramble shRNA, or shRNAs targeting *VEGFR1* or *2* for 48 h. **(I)** Morphology and AP staining of E14 cells cultured in mES medium without LIF but containing 2i (3 μ M PD0325901 and 3 μ M CHIR99021), sunitinib (1 μ M) or DC101 (20 μ g/ml) for 3 days. **(J)** The percent of undifferentiated and differentiated colonies corresponding to **I**.

sponse to ER stress), *Ire1α*, *Grp78* and *CHOP*, increased with varying degrees from day 1 to day 8 (Figure 4C). Among them, *splXBP1* and *CHOP* increased dramatically on day 5 and the protein levels of XBP1 and CHOP

increased in a time-dependent manner and peaked on day 6 (Figure 4D). In the absence of LIF, ER stress inducers thapsigargin, tunicamycin and ionomycin [34] lead to more severe differentiation and higher expression of ER



stress-related genes and *VEGFA* (Figure 4E, 4F and 4H), while tauroursodeoxycholic acid, salubrinal and BHA [34], the ER stress inhibitors, blocked the expression of ER stress-related genes and *VEGFA* caused by the removal of LIF (Figure 4H), and thus prevented cell differentiation (Figure 4E and 4F).

It was reported that the retina of LIF-knockout mice displayed increased microvessel density due to increased VEGF expression [35]. We also found that LIF suppressed the expression of *VEGFs* in E14 cells during long-term culture (Supplementary information, Figure S7A). And such effect was likely due to the suppressed expression of HIF1 α and ER stress-related genes by LIF (Supplementary information, Figure S7B).

Taken together, these data suggest that both HIF1 α and ER stress are involved in the expression of *VEGFA* in mESCs, blocking these pathways is beneficial for the maintenance of pluripotency and self-renewal of mESCs. Interestingly, blocking the VEGF pathways with sunitinib decreased the expression of ER stress-related genes and *VEGFA* (Figure 4G and 4H), indicating the possible existence of a feed-forward mechanism between VEGF and HIF1 α /ER stress pathways. LIF may also promote self-renewal of mESCs by reducing *VEGF* expression via downregulation of HIF1 α and ER stress pathways.

Sunitinib promotes the generation of mouse iPSCs with four or reduced numbers of transcription factors

A number of small molecules, including 2i, pluripotin, BIO and PKC inhibitors, which support ESC self-renewal by inhibiting signal pathways involved in differentiation, have been reported to enhance the generation of iPSCs. Therefore, we investigated whether sunitinib can also promote reprogramming. Using a 96-well-plate-based screening system with OG2 MEFs carrying a transgenic *Oct4* promoter-driven GFP reporter [36], we found in 4 factor (4F: Oct4, Sox2, Klf4 and c-Myc)-mediated reprogramming, 1 μ M sunitinib (added from day

3 to 14) was able to induce significantly more GFP⁺ colonies (~8-fold increase) than vehicle control (Figure 5A). We also treated the 4F-transduced MEF cells with 1 μ M sunitinib for various durations starting from day 3. Sunitinib started to show beneficial effect after 5-day treatment (day 3-8). In fact, the longer the treatment duration, the better the results (Figure 5B), supporting the notion that sunitinib can promote the self-renewal of pluripotent stem cells. Indeed, sunitinib also supported the propagation of iPSCs in the absence of LIF (Supplementary information, Figure S3C).

Combinations of small molecules have been reported to facilitate the generation of iPSCs with reduced numbers of transcription factors. In 2F (Oct4 and Klf4)-transduced MEFs, sunitinib (1 μ M) facilitated iPSC generation alone or in combination with RepSox (1 μ M), VPA (0.5 mM), CHIR99021 (3 M) and parnate (2 μ M; Figure 5C). More remarkably, sunitinib (1 μ M) alone was able to enhance the efficiency of 1F (Oct4)-mediated reprogramming (Figure 5D). Sunitinib also doubled the number of GFP⁺ colonies induced by Oct4 in combination with RepSox (1 μ M), VPA (0.5 mM), CHIR99021 (3 μ M) and parnate (2 μ M; Figure 5D). Knockdown of *VEGFR1* or *VEGFR2* also facilitated 4F-mediated reprogramming (Supplementary information, Figure S8G).

A series of iPSC lines were established by using 4F, 3F, 2F or 1F plus sunitinib. PCR analysis of genomic DNA confirmed the integration of correct viral factors in these iPSC lines (Supplementary information, Figure S8A). Quantitative RT-PCR revealed the reactivation and expression of the endogenous *Oct4*, *Sox2*, *Nanog* and *Rex1* (Supplementary information, Figure S8B) and the silencing of viral genes (Supplementary information, Figure S8C). 1F- and 2F-iPSCs were further characterized. These iPSCs maintain GFP⁺- and ES-like morphology, and express typical pluripotency markers, such as AP, SSEA1 and Nanog (Figure 5E and Supplementary information, Figure S8D). Bisulfite genomic sequenc-

Figure 4 Role of HIF1 α and ER stress in VEGF expression and pluripotency of mESCs. **(A, B)** qRT-PCR **(A)** and western blot analysis **(B)** of HIF1 α in E14 cells cultured in mES media containing 1 000 U/ml LIF for 1-8 days. Tubulin served as a loading control for western blot. **(C, D)** qRT-PCR **(C)** and western blot analysis **(D)** of genes involved in ER stress in E14 cells cultured in mES medium containing 1 000 U/ml LIF for 1-8 days. **(E)** Morphology (bottom) and AP staining (top) of E14 cells cultured in mES medium without LIF but containing sunitinib (1 μ M), 2i, ML228 (1 μ M), KC7F2 (1 μ M), BHA (10 μ M) or tunicamycin (Tuni, 2.5 μ g/ml) for 48 h. Scale bar, 50 μ m. **(F)** Statistical data of the percent of undifferentiated and differentiated colonies in E14 cells cultured in LIF-free mES medium containing the indicated compounds for 3 days, including sunitinib, 2i, KC7F2 (1 μ M), CoCl₂ (30 μ M), ML228 (1 μ M), TUCDA (0.1 mM), salubrinal (Sal, 10 μ M), BHA (10 μ M), thapsigargin (Thap, 0.4 μ M), tunicamycin (Tuni, 2.5 μ g/ml) and ionomycin (Iono, 3.6 μ M). **(G)** qRT-PCR analysis of *VEGFA* in E14 cells cultured in LIF-free mES medium containing sunitinib, 2i or the indicated HIF1 α regulators (same concentrations as in **F**) for 3 days. Data are mean \pm SEM ($n = 3$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ vs control. **(H)** qRT-PCR analysis of the expression of ER stress-related genes and *VEGFA* in E14 cells cultured in LIF-free mES medium containing sunitinib, 2i or the indicated ER stress regulators (same concentrations as in **F**) for 3 days. Data are mean \pm SEM ($n = 3$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ vs control. Independent experiments were repeated at least three times.

ing of the *Oct4* and *Nanog* promoters revealed that both were demethylated in 1F-iPSCs as in mESCs, while in MEFs they were both hypermethylated (Supplementary information, Figure S8E). Intramuscular injection of the 1F-iPSCs into SCID mice led to teratoma formation in 3-4 weeks, containing tissues derived from all three germ

layers, including epidermis structure (ectoderm), muscle and cartilage structure (mesoderm) and epithelium tube structure (endoderm; Figure 5F). The 1F- and 2F-iPSCs were also able to generate chimera mice when injected into the blastocysts of ICR mice (Figure 5G and Supplementary information, Figure S8F), indicating that the

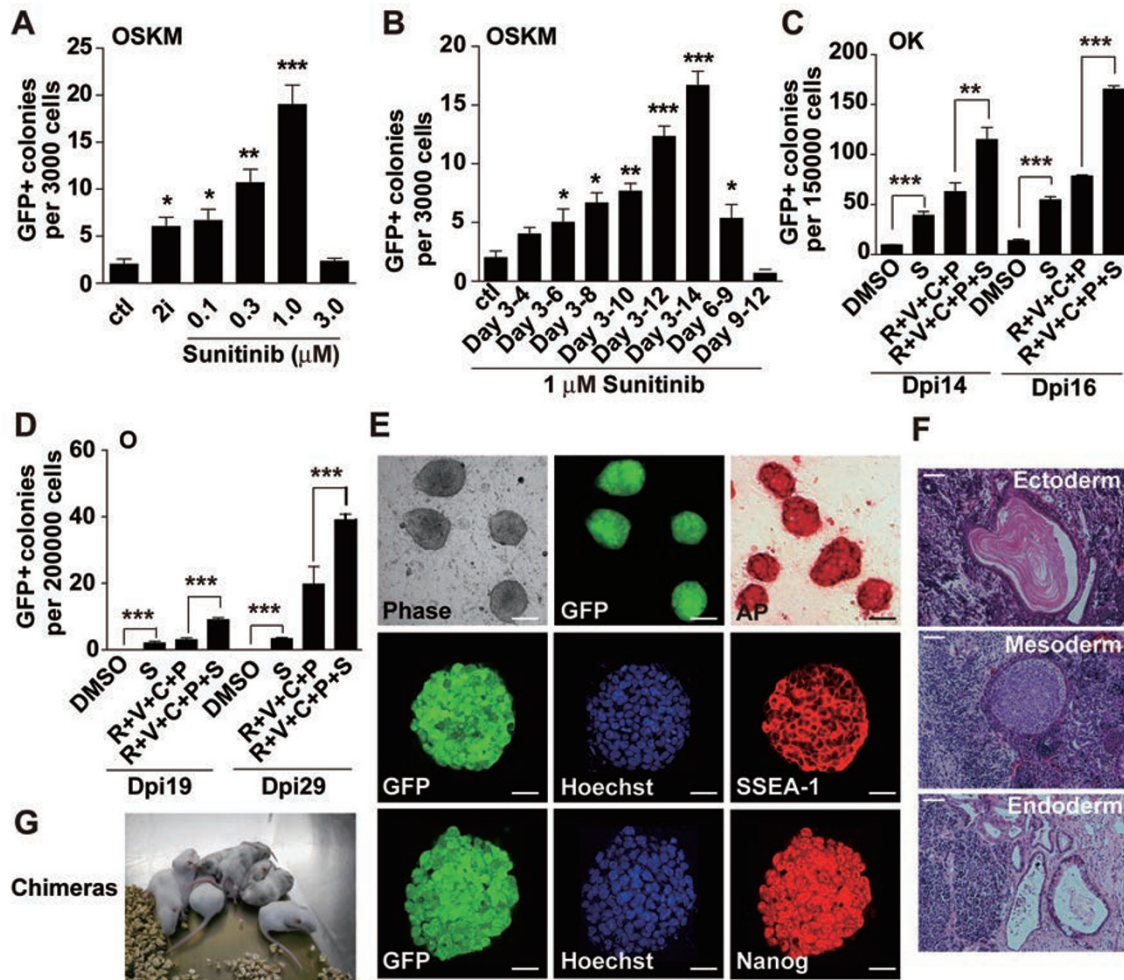


Figure 5 Sunitinib promotes reprogramming of somatic cells. **(A)** Dose response of sunitinib in 4F-mediated iPSC induction. Sunitinib at the indicated concentration was added into 3 000 4F-transduced MEFs from day 3 to day 14. **(B)** Time response of sunitinib in iPSC induction. Sunitinib (1 μM) was added at the indicated time periods, and GFP⁺ colonies were counted on day 14. Data are mean SEM (n = 3). *P < 0.05, **P < 0.01, ***P < 0.001 vs control. Independent experiments were repeated at least three times. **(C)** Sunitinib facilitated iPSC generation from 2F (OK)-transduced MEFs alone or in combination with several known chemicals. For OK-mediated reprogramming, 150 000 OK-infected cells in 6-well plates were treated with 1 μM sunitinib (S) alone or various combinations of compounds (including 1 μM R, RepSox; 0.5 mM V, VPA; 3 μM C, CHIR and 2 μM P, Parnate) from day 3 to day 14. **(D)** Sunitinib facilitated iPSC generation from Oct4-transduced MEFs alone or in combination with several known chemicals. For Oct4-mediated reprogramming, 200 000 Oct4-infected cells in 6-well plates were treated with 1 μM sunitinib (S) alone or the indicated combinations of compounds from day 3 to day 14. GFP⁺ colonies were counted on day 19 and day 29. **(E)** Top: morphology, GFP expression and AP staining in iPSC clones induced with 1F (Oct4) plus sunitinib (1 μM). Scale bar, 50 μm. Middle and bottom: immunofluorescent staining of SSEA-1 and Nanog in the same clone. Scale bar, 10 μm. **(F)** HE-stained sections of teratomas formed with iPSC clone described in **E**. Typical structures of the three embryonic germ layers are shown: epidermis (ectoderm), muscle and cartilage (mesoderm) and epithelium (endoderm). Scale bar, 20 μm. **(G)** chimeric mice produced with iPSC clone described in **E**.

iPSCs generated with sunitinib were pluripotent.

Other RTK inhibitors also facilitate the self-renewal of mESC and the generation of iPSCs

Apart from sunitinib, a number of RTK inhibitors whose targets include VEGFR and PDGFR have been developed and some have been used as anti-neoplastic drugs [19]. We tested whether these drugs or compounds could also be used to maintain the self-renewal of mESCs and facilitate factor-mediated reprogramming. E14 cells were seeded into 96-well plates at a density of 1 000 cells per well in mES medium and allowed to grow with LIF for 1 day, and then media were changed into mES media without LIF but containing various compounds for 2 days. Then, cells were stained with AP staining kit. Cell colonies could be characterized into differentiated or

undifferentiated categories according to the morphology and AP activity. All of the 10 RTK inhibitors, including axitinib, sorafenib, dovitinib, lenvatinib, motesanib, nintedanib, vandetanib, pazopanib, tivozanib and semaxinib, showed beneficial effect in maintaining the self-renewal of mESCs, with sunitinib and dovitinib being the most effective (Figure 6A). We also tested these compounds in our 96-well-plate-based iPSC induction systems with 4 factors, and all compounds significantly enhanced 4F-mediated reprogramming at their optimal concentrations, with sunitinib, dovitinib, axitinib and vandetanib showing the best results (Figure 6B).

Discussion

ESCs are isolated from the inner cell mass (ICM) of

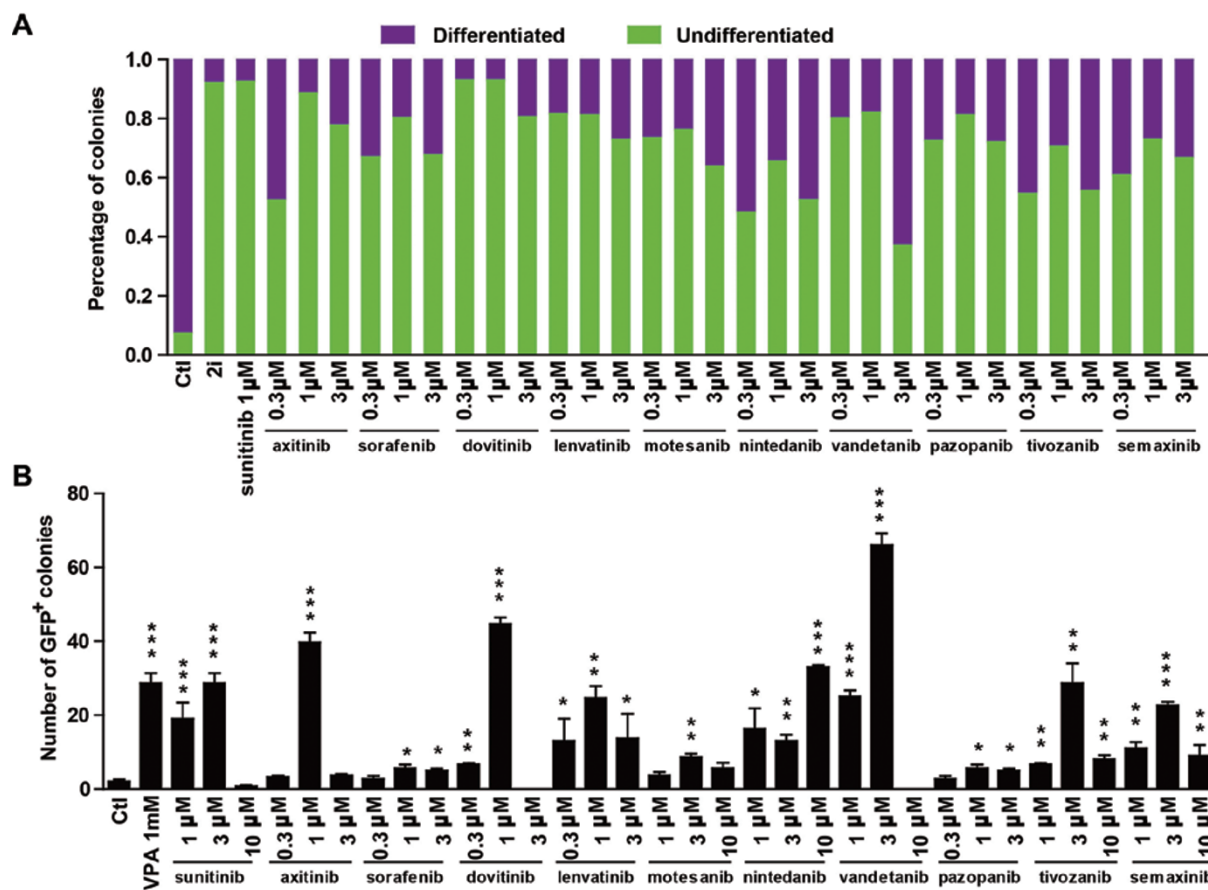


Figure 6 Other RTK inhibitors targeting VEGFRs facilitate the self-renewal of mESCs and the generation of iPSCs. **(A)** E14 cells were cultured in LIF-free mES media containing sunitinib or other inhibitors at the indicated concentrations for 3 days. The initial cell density was 1 000 cells/well in 96-well plate. The percent of AP⁺ (undifferentiated) and AP⁻ (differentiated) colonies were calculated. 2i culture was used as a positive control. **(B)** MEFs transduced with Yamanaka 4 factors were treated with sunitinib or the indicated inhibitors (added from day 3 to day 14) and GFP⁺ colonies were counted on day 14. VPA (1 mM, added from day 3 to day 8) treatment was used as a positive control. Data are mean \pm SEM ($n = 3$). * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ vs vehicle control.

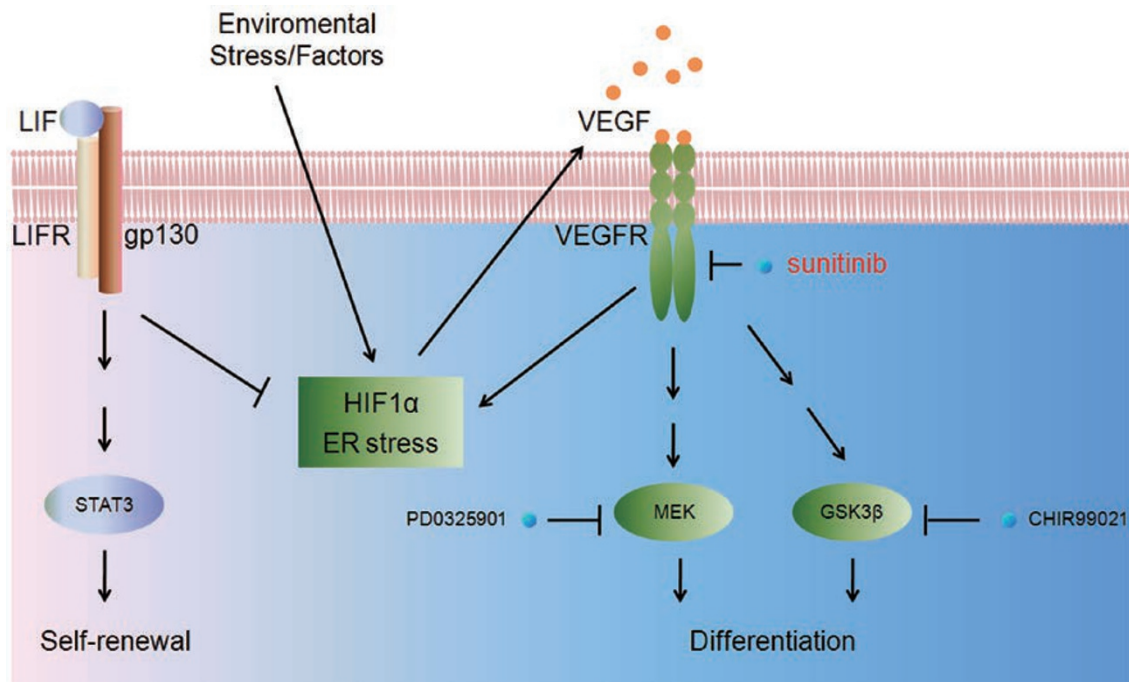


Figure 7 Schematic representation of the autocrine VEGF signaling in mESCs. Production of VEGF is activated via HIF1 α and/or ER stress, presumably due to environmental stress in the *in vitro* culture. VEGF activates VEGFRs expressed on the surface of the mESCs, inducing the activation of MEK, GSK3 β and possibly other pathways that promote differentiation and lineage commitment of mESCs. Blocking VEGFRs with sunitinib inhibits both the MEK and GSK3 pathways and the feed-for-ward loop between HIF1 α /ER stress and VEGF/VEGFR, and thus prevents the spontaneous differentiation of mESCs.

early embryos. It is quite remarkable that the pluripotent cells in the ICM only exist transiently as they undergo programmed differentiation, while their pluripotency could be captured *in vitro*. The initial isolation and maintenance of mESCs relies on ‘feeder cells’ and serum supplementation, but later studies identified LIF and BMP4 as the key extracellular stimulants required for the sustained proliferation and self-renewal of mESCs [2, 37]. Similarly, hESCs can also be cultured *in vitro*, although they require the activation of fibroblast growth factor (FGF2) and TGF β /activin/Nodal pathways for maintenance [38, 39]. In addition, modulation of the WNT pathway in mouse and human ESCs by inhibiting GSK3 β activity, or adding WNT ligands, also enables short-term maintenance of the undifferentiated state [40].

Many of these growth factors exert their actions in ESCs in an autocrine manner. For example, FGF2 is believed to be secreted by hESCs to induce the expression of TGF β family members such as Nodal, TGF β and activin that activate the SMAD2/3 pathway which help maintain pluripotency [38]. In contrast, FGF4 is produced in mESCs as an autocrine factor that activates the ERK pathway and promotes exit from self-renewal into lineage commitment [4]. The latter leads to the dis-

covery that the combination of inhibitors of ERK and GSK3 pathways (2i) are able to sustain the self-renewal of mESCs in the absence of extrinsic signals [5]. It has been suggested that mESCs cultured under these conditions are in a ground or ‘naive’ state and equivalent to cells from the early epiblasts. In this study, we identified VEGF as another autocrine factor that promotes lineage commitment of mESCs even in the presence of LIF. Blocking VEGF signaling with sunitinib, or by knocking down VEGFRs, not only maintained the mESCs in the undifferentiated state without the need for feeder cells or LIF, but also facilitated transcription factor-mediated pluripotency reprogramming in MEFs.

It is well accepted that VEGF produced by tumor cells acts on neighboring VEGFR-expressing endothelial cells to promote angiogenesis for continued tumor growth [41]. This concept has been challenged by reports demonstrating expression of the VEGFRs on tumor cells. An intriguing finding related to the autocrine function of VEGF signaling in tumors was that VEGF secreted by tumor cells triggers tumor cell proliferation by binding VEGFR1 expressed on the same cell [42]. Moreover, in patients with lung cancer, an increase in gene copy number of *VEGFR2* is associated with chemoresistance

and shorter survival [43]. Autocrine VEGF signaling has also been reported to play important roles in all aspects of breast tumor biology, including adhesion, survival, migration and invasion of the cancer cells [44]. In physiological condition, cell-autonomous VEGF signaling pathway has been demonstrated to be required for the homeostasis of blood vessels in the adult even though it is dispensable for the angiogenic cascade [45]. In ESC or iPSC cultures, the addition of exogenous VEGF has been demonstrated to induce endothelium and cardiac myocyte differentiation [15, 16], but the autocrine function of VEGFs has rarely been discussed. Our data demonstrated that the mESCs express both VEGF and VEGFRs, and the activation of this pathway leads to spontaneous differentiation. Interestingly, blocking the VEGF pathways with sunitinib decreased the expression of *VEGFA* (Figure 4G and 4H) in mESCs, indicating the possible existence of an autocrine feed-forward loop which contributed to the continuous increase of VEGF secretion in long-term ESC culture. Such autocrine feed-forward loop of VEGF/VEGFRs has also been demonstrated in lung cancer [46].

VEGFs are dramatically upregulated under hypoxic conditions. Hypoxia allows the stabilization of HIFs that bind to the promoter region of *VEGFA* [47]. Under different cultivation conditions, the role of hypoxia and HIFs in the fate determination of ESCs remains controversial. It has been reported that hypoxia prevents ESC differentiation and promotes iPSC induction [24]. Conversely, other report suggested that hypoxia or HIFs enhance the differentiation of hESC and mESC toward cardiomyocytes [48]. Exogenous expression of HIF-1 α also promotes cardiac differentiation of mESCs [49], while HIF-1 α -knockout mESCs do not produce spontaneously beating cells [50]. Recently, VEGF, the important downstream protein of HIF1 α , was proved to be indispensable in effective cardiac myocyte differentiation of human iPSCs [16]. In addition, hypoxic priming of mESCs accelerates vascular-lineage differentiation through HIF1-mediated VEGF expression and Oct4 suppression [27]. Our findings support the notion that the activation of HIF1 α and subsequent VEGF secretion from mESCs facilitated the meso-endoderm lineage commitment of these cells. Blocking HIF1 α reduced the expression of VEGF and promoted self-renewal of mESCs. Our data also demonstrated that ER stress plays a role in VEGF secretion in mESCs. It has been reported that hESCs must resolve ER stress related to culture conditions to survive and maintain pluripotency [32]. Blocking ER stress also reduced the expression of VEGF and promoted self-renewal of mESCs. Interestingly, we also demonstrated that LIF might also exert its self-renewal-promoting effect by inhibiting HIF1 α and ER stress pathways, thus blocking

the activation of the autocrine VEGF signaling in mESCs (Figure 7).

Sunitinib (marketed as Sutent) is an orally available multi-targeted RTK inhibitor that was approved by the FDA for the treatment of renal cell carcinoma and imatinib-resistant gastrointestinal stromal tumor. Although mainly targeting VEGFRs and PDGFRs, it also inhibits other RTKs including KIT, FLT3, and RET [19]. In our study, we have demonstrated the expression of VEGF/VEGFRs and PDGF/PDGFR in mESCs, and the activation of PDGF pathway might induce differentiation in a paracrine fashion, while the activation of VEGF pathway might induce differentiation in an autocrine way. Although we have demonstrated the critical role of VEGFR2 in spontaneous differentiation of mESCs by the knockdown and neutralizing antibody studies, sunitinib might exert its effect by blocking additional RTKs. For example, SCF and its receptor c-Kit play important roles in hematopoiesis, spermatogenesis and melanogenesis [51]. Mice that do not express SCF or c-Kit both die from severe anemia [51]. SCF may serve as guidance cues that direct hematopoietic stem cells to their stem cell niche. Flt3 is expressed on many hematopoietic progenitor cells, and play an important role in the development of haematopoietic stem cells and progenitor cells. Mutation in *Flt3* gene is associated with acute myeloid leukemia [52]. CSF-1R, also known as macrophage CSF-1R, controls the differentiation and function of macrophages [53]. Mutations in CSF-1R are associated with chronic myelomonocytic leukemia and type M4 acute myeloblastic leukemia [54]. Although the expression and function of these RTKs have rarely been studied in mESCs, their activation all seem to lead to differentiation toward meso-endoderm lineages. Therefore, blocking any of these RTKs may also contribute to the self-renewal-promoting effect of sunitinib.

In conclusion, we have discovered the critical roles of autocrine VEGF signaling in the spontaneous differentiation of mESCs in *in vitro* culture. Blocking VEGF signaling with sunitinib, other RTK inhibitors targeting VEGFR, VEGFR neutralizing antibody, or by knocking down VEGFRs, was capable of maintaining the mESCs in the undifferentiated state without the need for feeder cells or LIF. Sunitinib could also be used to facilitate the derivation of mESCs from the blastocysts, and the mESCs maintained in sunitinib-containing medium remained pluripotent and were able to contribute to chimeric mouse. Sunitinib was also able to promote iPSC generation from MEFs. These results indicate the existence of an intrinsic differentiation pathway of mESCs by activating the VEGF signaling in an autocrine way. Blocking VEGF signaling with sunitinib or other small molecules

help to maintain the mESCs in the ground state of pluripotency.

Materials and Methods

ES cell culture

E14 and OG2-ES cells were maintained feeder free on gelatin-coated plastics in mES media (DMEM with 15% FBS (Hyclone 30070.03), 2 mM GlutaMAX, 0.1 mM non-essential amino acids (NEAA), 0.1 mM β -mercaptoethanol, 100 U/ml penicillin and 100 μ g/ml streptomycin) supplemented with 1 000 U/ml LIF and 2i (3 μ M CHIR99021 and 3 μ M PD0325901) and passaged every 3 days. In testing conditions, ES cells were cultured in mES media supplemented with LIF, 2i, growth factors (mVEGF₁₆₄ and hPDGF-BB) or other chemicals as indicated. In serum-free conditions, ES cells were cultured in N2B27 media (DMEM/F12 with 1/200 B27 supplement, 1/100 N2 supplement, 2 mM GlutaMAX, 0.1 mM NEAA and 0.1 mM β -mercaptoethanol) supplemented with LIF, 2i, growth factors or other chemicals as indicated.

Derivation of ES cells with sunitinib

Blastocysts from female ICR mice (Shanghai SLAC Lab Animal) were flushed from uterine horns with M2 medium (Sigma-Aldrich) 3.5 days *post coitum*. Isolated blastocysts were plated onto irradiated MEF feeder layer in mES medium containing sunitinib (1 μ M) and 2 000 U/ml LIF. To obtain ES cell colonies, blastocyst outgrowths were disaggregated with trypsin and re-plated on gelatin-coated plastic wells with sunitinib and LIF. ESCs were expanded by passaging in mES medium supplemented with sunitinib (1 μ M). After six passages in sunitinib-containing mES medium, cells were injected into the C57BL/6 blastocysts to test for chimeric generation.

Mouse iPSC generation

OG2 mice, which carry a transgenic GFP gene driven by Oct4 promoter, were mated with C57BL/6J. MEFs were isolated from E13.5 embryos heterozygous for the Oct4-GFP transgenic allele. Gonads and internal organs were removed before MEF isolation. MEFs were grown in DMEM supplemented with 10% FBS, 2 mM GlutaMAX, 0.1 mM NEAA, 100 U/ml penicillin and 100 μ g/ml streptomycin. MEFs in early passages (up to passage 3) were used for further experiments.

Retroviruses were produced by transfection of plat-E cells with pMXs retroviral vectors containing the coding sequences of mouse Oct4, Sox2, Klf4 and c-Myc (Addgene). MEFs were seeded at a density of 150 000 cells/well into six-well plate 18 h before infection. Virus-containing supernatants, supplemented with 8 μ g/ml polybrene, were added into the plates of MEF culture and centrifuged at 2 500 rpm for 90 min. Medium was changed immediately after virus transduction and this day is counted as “day 0”. Two days post virus infection, MEFs were reseeded at a density of 3 000 cells/well into 96-well plates pre-seeded with irradiated MEF feeder cells, supplemented with mES medium containing 1 000 U/ml LIF. On day 6, culture medium was replaced with KSR medium (knockout DMEM supplemented with 15% KSR, 2 mM L-GlutaMAX, 0.1 mM NEAA, 0.1 mM β -mercaptoethanol, 1 000 U/ml LIF, 100 U/ml penicillin and 100 μ g/ml streptomycin). Chemicals were added from day 3 with various durations. GFP⁺ colonies were counted using a high-content screening instrument Acumen (TTP

Labtech) on day 14 post infection. Colonies were also trypsinized and analyzed using guava easyCyte Flow Cytometer (Millipore).

For reduced factor-induced iPSC, 150 000 MEFs cells (2F, Oct4 and Klf4) or 200 000 cells (1F, Oct4) per well were seeded in six-well plate 18 h before infection. Supernatants containing 2F or 1F were used for infection as above. Medium was changed immediately after virus transduction and this day is counted as “day 0”. MEFs were infected twice, 2 days post virus infection, and MEFs were supplemented with KSR medium containing 10 ng/ml vitamin C, 1/100 N2 supplement and 10 ng/ml bFGF. Chemicals were added from day 3 to day 14.

Statistic analysis

Values are reported as the mean \pm SEM. *P*-values were calculated by Student's *t*-test. All graphs are plotted with GraphPad Prism 5 software.

Detailed methods are described in the Supplementary information, Data S1.

Acknowledgments

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