Platelets are key elements not only of hemostasis and thrombosis but also of tissue regeneration after injury and the pathophysiology of inflammation (Gawaz et al., 2005; Nesbitt et al., 2009). The production of platelets, thrombopoiesis, is regulated primarily by thrombopoietin (TPO)-mediated megakaryopoiesis within the BM (Patel et al., 2005; Schulze and Shivdasani, 2005). Notably, many patients with critical thrombocytopenia, caused by dysregulation of BM as a result of hematopoietic disease or aggressive chemotherapy, require platelet transfusions using platelet concentrates obtained through blood donation (Webb and Anderson, 1999). It is well known, however, that repeated transfusion induces antibodies in recipients against allo- genic human leukocyte antigen (HLA) on the transfused platelets (Schiffer, 2001). To establish a supply of identical platelet concentrates without loss of responsiveness as a result of immunorejection, particularly for patients with a rare HLA, human (h) induced pluripotent stem cells (iPSCs) represent a potentially abundant source.

Human (h) induced pluripotent stem cells (iPSCs) are a potentially abundant source of blood cells, but how best to select iPSC clones suitable for this purpose from among the many clones that can be simultaneously established from an identical source is not clear. Using an in vitro culture system yielding a hematopoietic niche that concentrates hematopoietic progenitors, we show that the pattern of c-MYC reactivation after reprogramming influences platelet generation from hiPSCs. During differentiation, reduction of c-MYC expression after initial reactivation of c-MYC expression in selected hiPSC clones was associated with more efficient in vitro generation of CD41a<sup>+</sup>CD42b<sup>+</sup> platelets. This effect was recapitulated in virus integration–free hiPSCs using a doxycycline–controlled c-MYC expression vector. In vivo imaging revealed that these CD42b<sup>+</sup> platelets were present in thrombi after laser–induced vessel wall injury. In contrast, sustained and excessive c-MYC expression in megakaryocytes was accompanied by increased p14 (ARF) and p16 (INK4A) expression, decreased GATA1 expression, and impaired production of functional platelets. These findings suggest that the pattern of c-MYC expression, particularly its later decline, is key to producing functional platelets from selected iPSC clones.

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RESULTS

Four-factor hiPSC-derived hematopoietic progenitors contribute to enhanced generation of MKs and platelets

Using VSV-G–pseudotyped retroviruses (Ory et al., 1996) harboring human reprogramming factors (OCT3/4, SOX2,
To determine the mechanism underlying the enhanced megakaryopoiesis exhibited by four-factor hiPSC-derived hematopoietic progenitors, we assessed the potential of progenitors

KLF4, and/or c-MYC), we sought to establish iPSCs from human dermal fibroblasts (HDFs). With our system, we consistently generated 200–300 hiPSC clones from 10^6 HDFs. For evaluation of pluripotency, established iPSC clones obtained through transduction with four or three factors (with or without c-MYC) and all clones showing a normal karyotype (not depicted) were examined for morphology, SSEA-4 expression (Fig. S1 A), other gene expression (Fig. S1 B), and the ability to form teratomas in vivo (Fig. S1 C). Our findings confirmed that exogenous OCT3/4, SOX2, KLF4, and c-MYC remained unexpressed in established iPSCs (Fig. S1 B).

To explore the hiPSC clones’ potential for differentiation into hematopoietic cells (Takayama et al., 2008), we evaluated several iPS-Sacs (Fig. S2 A) from individual clones (four-factor hiPSCs: TkDA3-1, -2, -4, -5, -9, -20, 201B6, and 201B7; three-factor hiPSCs: TkDN4-M, 253G1, and 253G4) and compared them to previously evaluated human (h) embryonic stem cells (ESCs; KhES clone, Kyoto University, Japan; Takayama et al., 2008). On day 15 of culture, iPS-Sacs that contained numerous hematopoietic-like round cells (Fig. S2 B) and showed expression of vascular endothelial growth factor type 2 receptor (VEGF-R2; Fig. S2 B) or platelet endothelial cell adhesion molecule 1 (CD31; not depicted) were deemed to be potentially suitable microenvironments from which to obtain hematopoietic progenitors, as was observed in hESC-derived structures (Takayama et al., 2008).

We detected considerable heterogeneity in the production of iPS-Sacs (a hallmark of the efficiency of hematopoietic progenitors) from iPSCs derived from the same source (i.e., TkDA3-1, -2, -4, -5, -9, or -20; Fig. 1 A), which was also consistent with previous observations in hESCs (Osafune et al., 2008). In particular, CD34+, but not CD34−, cells from iPS–Sacs showed successful colony formation in methycellulose colony assays (Fig. S2 C). The three-factor clone TkDN4-M, as well as KhES-3, appeared to have a greater potential for myeloid lineage hematopoiesis, as exemplified by the numbers of Sacs (Fig. 1 A, red bar) composed of CD34+ cells (Fig. 1 B, red bar), and the numbers of hematopoietic colonies formed from each Sac (Fig. 1 C, red bar). Nonetheless, the number of CD42b (GPIbα; von Willebrand factor receptor)+ MKs obtained with four-factor iPSC clones (e.g., TkDA3-2, TkDA3-4, and TkDA3-5) was higher than that obtained with TkDN4-M or KhES-3 when equal numbers of cells from iPS–Sacs were seeded onto fresh culture dishes in the presence of TPO, stem cell factor (SCF), and heparin (Fig. 1, D [day 22] and E; and Fig. S3 day 26; Takayama et al., 2008). For example, clone TkDA3-4 generated three times as many MKs as TkDN4-M or KhES-3 at the peak of production (Fig. 1 E, day 26; and Fig. S3). By days 22–38, phase-contrast imaging revealed the presence of proplatelets, a prerelease platelet form (Video 1), as well as mature MKs by May–Giemsa staining (Fig. S4). Moreover, flow cytometric analysis showed that 40–60% of floating cells expressed CD41a (integrin αIIbβ3 complex), a fibrinogen receptor, as well as CD42a (GPIX), GPIbα, and CD9, all of which are hallmarks of MKs (Fig. 1 F; Tomer, 2004; Takayama et al., 2008).

Figure 2. Time-dependent changes in qPCR induced by exogenous reprogramming genes in three-factor or four-factor iPSCs. mRNA encoding exogenous OCT3/4 (A), SOX2 (B), KLF4 (C), and c-MYC (D) in human ES cells (ESCs), TkDN4-M (three-factor iPSCs), TkDA3-2, TkDA3-4, and TkDA3-5 (four-factor iPSCs) on day 0 or their derivatives (on days 6, 10, 15, 22, and 26 after initiation of MK-lineage culture) were examined by qPCR as described in the Materials and methods section. TkDA3-4–derived mature MKs (day 26) was assigned a value of 1.0 (n = 4, means ± SEM from two independent experiments).

To determine the mechanism underlying the enhanced megakaryopoiesis exhibited by four-factor hiPSC-derived hematopoietic progenitors, we assessed the potential of progenitors
MK lineage culture) revealed expression of the exogenous (transgene [Tg]) reprogramming genes, which were not expressed before hematopoiesis (Fig. 2, A–D, day 0; and Fig. S1 B). Although qPCR after day 15 suggested that, in four-factor iPSCs, activation of \( \text{OCT3/4} \) Tg and/or \( \text{c-MYC} \) Tg might affect the enhanced megakaryopoiesis (Fig. 2, A and D), individual Tg activation was not dependent on the copy number in the genome (Fig. S1, D and E).

Thus, to confirm the functional effect of \( \text{OCT3/4} \) and/or \( \text{c-MYC} \) Tg on megakaryopoiesis from pluripotent stem cells, within iPSC-Sacs, based on colony-forming capacity (Fig. S2 D) and their surface markers (not depicted). We found no significant differences between TkDA3–4 (four-factor) and TkDN4–M (three-factor; Fig. S2 D), which means the potential and the capacity to drive most myeloid lineage commitment from iPSC-derived progenitors is independent of the clone type, or at least there was no detectable difference between the three- and four-factor iPSC clones we examined (not depicted).

In contrast, quantitative (q) PCR analysis of hematopoietic cells on days 22–26 (7–11 d after replating for selective MK lineage culture) revealed expression of the exogenous (transgene [Tg]) reprogramming genes, which were not expressed before hematopoiesis (Fig. 2, A–D, day 0; and Fig. S1 B). Although qPCR after day 15 suggested that, in four-factor iPSCs, activation of \( \text{OCT3/4} \) Tg and/or \( \text{c-MYC} \) Tg might affect the enhanced megakaryopoiesis (Fig. 2, A and D), individual Tg activation was not dependent on the copy number in the genome (Fig. S1, D and E).

Thus, to confirm the functional effect of \( \text{OCT3/4} \) and/or \( \text{c-MYC} \) Tg on megakaryopoiesis from pluripotent stem cells,
c-MYC levels in hiPSC-derived MKs determines the number of platelets generated per MK

We next tested whether iPSC-derived MKs actually yield platelets in vitro. We confirmed that four-factor iPSCs generate greater numbers of platelets than three-factor iPSCs or hESCs (Fig. 4, A and B; and Fig. S5). Moreover, we noticed that, at the peak of production (day 26), many more proplatelets and platelets were generated from TkDA3-4 iPSCs than from any other four-factor iPSCs (Fig. 4, B and C; and Fig. S5). We also noted that TkDA3-2 and TkDA3-5 iPSC-derived MKs showed an earlier peak, on day 22 (Fig. 1, D and E; and Fig. S3) and that there were fewer proplatelets in the dishes (Fig. 4 C), which suggests that most MKs promote apoptosis and/or senescence leading to inhibition of platelet release in those two clones.

It has been reported that forced expression of c-Myc impairs maturation of MKs displaying polyploidization, leading to an increase in immature MKs (Thompson et al., 1996a). Indeed, we confirmed the appearance of immature MKs (Fig. 3, A–C). Flow cytometry revealed that on day 22, most of the EGFP+ or KO+ population was CD41a+CD42b+ in the c-MYC O/E specimens but not in the others (Fig. 3 D), although only mononuclear and lower ploidy cells were present (Fig. 3, E and F). These suggest that OCT3/4 O/E might not accelerate megakaryopoiesis (Fig. 3 D). Collectively then, these findings suggest that stronger expression of c-MYC in hESCs might promote lineage commitment into megakaryopoiesis without maturation.

Figure 4. Level of c-MYC reactivation in individual iPSC-derived MKs may determine the efficiency of platelet generation in vitro. (A and B) Numbers of CD41a+CD42b+ platelets generated from hESCs or hiPSCs on days 22 (A) and 26 (B; peak of platelet generation; n = 5, mean ± SEM). (C and D) Numbers of proplatelets (C) and platelets (D) derived from four-factor iPSCs and from ESC hematopoietic progenitors, with or without c-MYC transduction (ii; n = 5, mean ± SEM). Representative photomicrographs of proplatelets are derived from four-factor iPSCs. (D) Numbers of platelets per MK was calculated as the total number of platelets divided by the total number of MKs on day 26 (n = 5, mean ± SEM). (E) Representative flow cytometry dot plots show MKs derived from TkDA3-4 and KhES-3, with or without c-MYC transduction, on day 26.
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CD42b (GPIbα), as indicated by the recovery of CD42b expression in the presence of a metalloprotease inhibitor (Fig. S6 B; Nishikii et al., 2008). In contrast, platelet-like particles from hESC-derived MKs ectopically expressing c-MYC (c-MYC-O/E) showed significantly lower CD42b expression, a distinct pattern on dot plots (Fig. 4 E and Fig. S6 A), and no recovery of CD42b expression after administration of metalloprotease inhibitor (not depicted). Given the platelet generation per MK, it appears that forced expression of c-MYC in ESCs impairs platelet yield on day 26 (Fig. 4 D), which might recapitulate in TkDA3-2 or TkDA3-5 iPSC-MKS (Fig. 4 D).

How does the level of c-MYC expression control platelet generation from iPSCs?

The results so far suggest that excess c-MYC expression diminishes platelet yield. To confirm that hypothesis, we evaluated the time-dependent changes in the total expression of c-MYC and the genes involved in c-MYC activation and thrombopoiesis. qPCR analyses confirmed that total (endogenous plus exogenous) c-MYC expression in TkDN4-M (three-factor hiPSCs) or in four-factor hiPSCs (TkDA3-4 and TkDA3-5) on days 0, 15, 22, and 26. All levels were normalized to the level of GAPDH expression (n = 4 of two independent samples). The levels of c-MYC, p14ARF, and p16INK4A expression in an undifferentiated TkDA3-4 iPSC clone (day 0) or expression of the other genes (D–F) in TkDA3-4-derived mature MKs (day 26) was assigned a value of 1.0 (n = 4, means ± SEM).

Figure 5. Level of c-MYC expression affects INK4A/ARF locus genes and genes related to MK maturation during mega-karyopoiesis from pluripotent stem cells.

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Inducible c-MYC expression in iPSCs without reactivation exhibited behavior similar to that of iPSCs with reactivation, leading to efficient generation of functional platelets.

To further confirm whether an increase and subsequent decline in c-MYC is critical for megakaryopoiesis, leading to an efficient platelet yield, we prepared a Sendai viral vector (SeV) harboring the four reprogramming factors expression in SeV-iPSC clones (passage number 4) derived from CB, HDF-A (adult), and HDF-N (neonate). A sample of HDFs transduced with SeV is used as a positive control for the SeV Tg. (C) Scheme of c-MYC induction in SeV-iPSC-derived hematopoietic cells. Hematopoietic progenitors derived from SeV-iPSCs were transfected with DOX-inducible c-MYC O/E vector on day 15 and analyzed on day 26. In Protocol a, DOX was added only from days 15 to 22. In protocol b, DOX was added from days 15 through 26. (D) Representative Western blots of cell lysates with c-MYC O/E (DOX-on; protocol b) or without c-MYC O/E (DOX-off; Protocol a) on day 26. The α-tubulin levels indicate same protein value. (E–G) Numbers of CD42b (GPIb) MKs (E), proplatelets (F), and platelets (G) on day 26 derived from 10^5 hematopoietic progenitors transfected with vehicle or DOX-inducible c-MYC O/E vector in protocol a or protocol b (n = 4, means ± SEM). (H) Numbers of platelets per MK generated on day 26 of culture (peak of platelet generation; n = 4, means ± SEM). Numbers of platelets per MK were calculated as the total number of platelets divided by the total number of MKs on day 26.
integrin activation in human platelets (from peripheral blood [PB]), PB-based aged platelets (48-h incubation; Bergmeier et al., 2003; Nishikii et al., 2008), iPSC platelets, and ESC platelets. The aged platelets were tested because iPSC-derived platelets were heterogeneously produced from MKs at various stages in culture, so that many of the platelets produced could have become aged (Nishikii et al., 2008). Conformational changes in integrins are required for platelet aggregation and stable thrombosis in vivo (Shattil et al., 1985). Indeed, although PB-based aged platelets were nonresponders, the integrin activity of TkDA3-4 (four-factor iPSCs) platelets was comparable to that of TkDN4-M (three-factor iPSCs) platelets, which showed a weaker response than human PB platelets (Fig. 7, A and B; and Fig. S8 A). Notably, c-MYC O/E-dependent iPSC-derived platelets showed little binding (Fig. 7, A and B). In contrast, platelets produced from SeV-iPSCs-MKs in the absence of c-MYC O/E after its activation responded well to ADP stimulation (Fig. 7 C). We therefore conclude that c-MYC activation and decline during MK differentiation may lead to the generation of functional platelets from iPSCs. We also examined expression of P-selectin (CD62P) on platelets in the presence of 50 µM ADP and observed weak but positive P-selectin expression in iPSC-derived platelets (Fig. S8 B).

**Human iPSC-derived platelets function normally in vitro and in vivo**

To assess the effect of c-MYC reactivation on the functionality of platelets from TkDA3-4, we compared agonist-induced c-MYC modulates platelet generation from iPSCs | Takayama et al. we confirmed that day 22 was the most suitable point to turn off O/E. The results also showed that continuous c-MYC O/E from days 22 to 26 still increased the number of MKs (Fig. 6 E), whereas the lack of c-MYC O/E from days 22 to 26 increased the total numbers of proplatelets (Fig. 6 F) or CD41a+CD42b+ platelets (Fig. 6 G). An increase in platelet yield per MK was also evident with the absence of c-MYC O/E after day 22 (Fig. 6 H), confirming the effect of c-MYC expression on megakaryopoiesis.

**Figure 7. Integrin activation and the structure of human iPSC platelets are comparable to those in human PB-derived platelets.**

(A–C) Integrin activation in fresh human platelets (Fresh-P), aged human platelets (48-h incubation at 37°C; Aged-P), TkDN4-M (three-factor iPSC) platelets (3f-iPSC-P), TkDA3-4 (four-factor iPSC) platelets (4f-iPSC-P), and ESC platelets (ESC-P), with or without c-MYC O/E. The binding of PAC-1 (indicative of platelet activation) to individual platelets was quantified in the absence and presence of 50 µM ADP using flow cytometry (A) Representative flow cytometry dot plots. Square indicates CD42b+ platelets. (B) Mean fluorescence intensity (MFI) of bound PAC-1, obtained from square gate in A. Error bars depict means ± SEM for four independent experiments (duplicate). (C) Representative flow cytometry analysis of PAC-1-bound platelets generated from integration-free SeV-iPSCs subjected to biphasic activation and, thereafter, decline of c-MYC expression as protocol b shown in Fig. 6 C. Square indicates CD42b+ platelets. (D) Spreading of iPSC platelets on fibrinogen. Human CD41a (red) and phalloidin (green) were used to identify F-actin fibers. Arrowheads indicate lamellipodia. Arrows indicate actin stress fibers. Bars, 5 µm. (E) Transmission electron micrographs of hiPSC (TkDA3-4) platelets on day 26. Bar, 3 µm.
In contrast, as in the previous paragraph, we recently showed that culture at 37°C influences the degradation of platelets by causing them to shed the extracellular domain of GPIbβ, which is required for initial adhesion to the injured vessel wall (Nishikii et al., 2008; Fig. S6 B). In vivo, platelets lacking the GPIbβ extracellular domain are quickly cleared from the circulation (Bergmeier et al., 2003), leading to insufficient levels of circulating platelets after transfusion (Nishikii et al., 2008). To avoid this without affecting platelet yield, we applied GM6001, a nonspecific inhibitor to metalloproteases, for 2 d before collection of cultured platelets and confirmed that the shedding of GPIbβ was dose-dependently inhibited by GM6001 (Fig. S6 B; Nishikii et al., 2008).}

We next sought to develop a transfusion model to assess platelet circulation in vivo. Using a NOG (nod-scid/IL-2 γc-null) mouse thrombocytopenia model induced by irradiation (2.0 Gy, 9 d beforehand), flow cytometric analyses performed...
2 and 24 h after transfusion (~1.0–1.2 × 10^7 platelets/mouse) showed that four-factor iPSC platelets were consistently present and that the percentage circulating was similar to that obtained with fresh human platelets (platelet chimerism of human CD41a+/mouse CD41a−; 3–10%, 2 h after transfusion; Fig. 8 A).

To further assess and confirm the functionality of TkDA3-4 hiPSC platelets in vivo, we used the same NOG mouse model (Fig. 8 A) with high-spatiotemporal resolution confocal laser microscopy to visualize the behavior of individual platelets upon initiation of adhesion to an injured vessel wall and during the subsequent steps in thrombus formation under flow within the vessel (Takizawa et al., 2010). iPSC platelets stained with tetramethylrhodamine ethyl ester, which is incorporated into the external lipid layer of the cell membrane, were transfused into NOG mice (2 Gy, 14 d beforehand), after which we confirmed they circulated as individual platelets (Fig. 8 B, red; and Video 2). Then using our novel laser injury thrombus model (Takizawa et al., 2010), we clearly observed that iPSC platelets initially adhered to the injured vessel wall, coordinating with host platelets and ultimately leading to thrombus formation and vessel occlusion (Fig. 8 C; and Videos 3–5). Thus, four-factor iPSC platelets appear capable of mediating hemostasis and thrombosis in vivo.

**DISCUSSION**

Megakaryocytic lineage-restricted c-Myc expression in mouse models showed this gene to be a positive regulator of MK progenitor proliferation (Thompson et al., 1996a,b) and to be required for TPO/c-mpl signaling in megakaryopoiesis (Chanprasert et al., 2006). Notably, c-Myc deficiency also accelerates megakaryopoiesis, but with lower ploidy (up to 8n), and augments immature platelet release accompanied by an increase in the platelet count. In our hiPSC culture system, however, total cellular c-Myc expression (endogenous plus reactivation of exogenous) appeared to be at an appropriate, and probably restricted, level suitable for promoting platelet generation (Fig. 4 B, Fig. 5 A, and Fig. 6 G, day 26). Increasing MYC levels enables cells to move from quiescence to S phase, even in the absence of mitogens (Eilers et al., 1991), possibly through activation of target genes (cyclin D1(D2,D3)/Cdk4/6) and cyclin E/Cdk2). In that regard, although MYC activity, per se, is required for normal cell proliferation (Murphy et al., 2008), it is recognized that above a certain threshold c-Myc expression may induce the onset of oncogenesis. Moreover, excessively high MYC levels induce senescence via activation of the Arf–Mdm2–p53 pathway (Eischen et al., 1999; Murphy et al., 2008) and suppression of key regulators of MK maturation, such as GATA1, B1-tubulin, and NF-E2, which likely inhibits maturation (Fig. 5, B–F) and may also induce apoptosis in some cells (Askew et al., 1991; Evan et al., 1992).

Four-factor hiPSC-derived hematopoietic progenitors also generate much larger numbers of platelets than three-factor iPSCs or hESCs (Fig. 4, A and B; and Fig. S5), possibly as a result of MK proliferation mediated through c-Myc reactivation (Fig. 3 A; Fig. 4, A and B; and Fig. S3). However, the number of platelets per MK derived from individual four-factor hiPSC clones (TkDA3-2, -4, and -5; with c-Myc) and ESCs (with or without c-Myc) on day 26 differs (Fig. 4 D). Nonetheless, TkDA3-4, which is the most efficient platelet-producing clone (Fig. 4, B and D; and Fig. S5), shows weaker c-Myc expression than other four-factor iPSCs or c-Myc-O/E MKs (Fig. 5 A). Indeed, DOX-inducible expression of c-Myc O/E in genome integration-free iPSCs (SeV-iPSCs) confirmed our hypothesis (Fig. 6, A–H). In addition, when we considered the possibility that deregulated MK maturation leads to release of nonfunctional platelets, we found differences in platelet functionality between hESCs or SeV-iPSCs, with or without c-Myc, and iPSC clones (Fig. 7, A–C). Immature megakaryocytic cell lines, such as Meg01, are reportedly capable of releasing CD41a+ platelet-like particles, but they show poor functionality in vitro (Takeuchi et al., 1998). We also found that hESC-derived CD41+ particles showed significantly less CD42b expression when c-Myc was ectopically expressed in MKs (Fig. 4 E and Fig. S6 A) and showed poor functionality in vitro (Fig. 7, A and B). In contrast, the function of CD42b+ platelets generated from TkDA3-4 was intact and able to mediate hemostasis in vivo, although some of the yield was likely made up of nonfunctional aged platelets (Fig. 7 and Fig. 8 C).

To clarify the underlying mechanism, we examined the association between platelet generation and gene expression. In c-Myc-O/E MKs, expression of GATA1 (Vyas et al., 1999; Rylski et al., 2003), a key regulator for MK maturation and polyploidization, was suppressed, whereas INK4A and ARF expression was greatly up-regulated (Fig. 5, B–D; Eischen et al., 1999; Murphy et al., 2008). Consequently, the maturation phase was inhibited, as exemplified by the presence of hypoploid cells without proplatelets (Fig. 3, E and F; and Fig. 4 C) and the weak expression of mature MK markers B1-tubulin and NF-E2 (Fig. 5, E and F), as well as platelet factor 4 (not depicted; Patel et al., 2005; Schulze and Shvidosani, 2005). These MKs only generated a few nonfunctional CD41a+CD42b+ platelets (Fig. 4, D and E; Fig. 5 A; and Fig. 7 B). In contrast, MKs derived from TkDA3-4 transiently showed appropriately high levels of c-Myc expression, with no effect on expression of INK4A/ARF locus genes (Fig. 5, A [day 15], B, and C), after which c-Myc expression declined (Fig. 5 A, days 22 and 26). From days 22 to 26, the pattern of GATA1, B1-tubulin, and NF-E2 expression was opposite that of c-Myc expression (Fig. 5, D–F), indicating that reduction of c-Myc after day 22 (immature MKs) may be required for MK maturation and generation of functional platelets (e.g., day 26). We therefore suppose that because most MKs derived from TkDA3-5, which produced fewer platelets than TkDA3-4 (Fig. 4 D), showed sustained c-Myc activity, leading to up-regulation of INK4A/ARF locus genes (Fig. 5), those MKs were unable to complete the maturation phase, even though GATA1, B1-tubulin, and NF-E2 expression was enhanced (Fig. 5, D–F). We further confirmed this phenomenon using integration-free SeV-iPSCs in a DOX-inducible gene expression system. Removal of c-Myc O/E from day 22 until day 26 increased the total numbers of proplatelets (Fig. 6 F) and...
CD42b+ functional platelets (Fig. 6 G and Fig. 7 C), as compared with the numbers seen with continuous c-MYC O/E until day 26. We therefore conclude that increased expression of c-MYC in hematopoietic progenitors may promote megakaryopoiesis, leading to increased MK generation (Fig. 6 E), but that a sustained increase in c-MYC after the MK progenitor stage may impair MK maturation, thereby diminishing platelet release (Fig. 6, F and G).

Both hESCs and hiPSCs were previously shown to differentiate into hematopoietic cells (Wang et al., 2004; Vodyanik et al., 2005; Ma et al., 2008; Choi et al., 2009; Yokoyama et al., 2009), although with the exception of hESC-derived natural killer cells (Woll et al., 2009), only in vitro functionality has been reported. In this paper, we demonstrated that platelets derived from hiPSCs via a mechanism involving limited c-MYC reactivation (Fig. 4 B and Fig. 5 A) are capable of thrombus formation in vivo.

Recent studies suggest that the source of the somatic cells and introduction of reprogramming factors without c-Myc (retroviral vectors, plasmids, proteins, Sendai viral vectors, and so on), also known as L-Myc/L-MYC, are essential elements for selection of efficient and safe iPSC clones (Okita et al., 2007; Nakagawa et al., 2010). We propose that selection of the iPSC clones most suitable for their purpose should also be considered. Our analysis of multiple hiPSC clones accompanied by Tg genome integration or a DOX-inducible expression system shows that time-dependent changes in c-MYC expression, specifically up-regulation and then down-regulation within an appropriate time span, facilitates generation of a novel platelet transfection system derived from hiPSCs. It is noteworthy that in vitro generation of platelet concentrates custom made from HLA-identical donors or the patients themselves are not subject to immune rejection and do not require donor blood. We propose that iPSC platelets could be an invaluable resource for patients requiring repeated platelet transfusion and that this system should enable us to investigate as yet unresolved aspects of the mechanisms underlying thrombocytopenia.

MATERIALS AND METHODS

Cells, reagents, viral vectors, and mice. All reagents were obtained from Sigma-Aldrich unless indicated otherwise. The human ESC clone Kyoto hiESCs (Khes) 3 was obtained from the Institute for Frontier Medical Science, Kyoto University (Kyoto, Japan) after approval for hESC use was granted by the Minister of Education, Culture, Sports, Science, and Technology of Japan. PB was provided from healthy volunteers approved by ethic committee of the Institute of Medical Science at University of Tokyo for experiments and use of viral vectors were approved by the committees of the JEM of Japan. Original hiPSC clones, 201B6, 201B7, 253G1, and 253G4 (Kyoto, Japan). Semi-qRT-PCR. hiPSCs were lysed with Trizol (Invitrogen), after which aliquots of suspended cells (1–3 × 10(6) cells) were in-digested with BglII, and NdeI, and NcoI for c-MYC or Oct3/4, respectively, Southern blotting analysis. 7 µg of genomic DNA was digested with BglII, HindIII, and EcoRI, and was electrophoresed on a 0.8% agarose gel and transferred to a nylon membrane (GE Healthcare). The membranes were hybridized with a 32P-labeled probe and exposed to X-ray film.
Hybridized with radioactively labeled DNA probes (c-MYC-exon3 or Oct3/4-exon1) in PerfectHyb Plus Hybridization buffer (Sigma-Aldrich) at 55°C overnight with constant agitation. After washing, signals were visualized using the FAL-5100 imaging system (FujiFilm).

Hematopoietic differentiation of hiPSCs. To differentiate hiPSCs into hematopoietic cells, we used the same protocol we established with hESCs (Takayama et al., 2008). In brief, small clumps of hiPSCs (~100 cells treated with PBS containing 0.25% trypsin, 1 mM CaCl₂, and 20% KSR) were transferred onto irradiated C3H10T1/2 cells and co-cultured in hematopoietic cell differentiation medium, which was refreshed every 3 d. On days 14–15 of culture, the iPSC-Sacs were collected into a 50-ml tube, gently crushed with a pipette and passed through a 40-µm cell strainer to obtain hematopoietic progenitors, which were transferred onto freshly irradiated feeder cells and cultured in differentiation medium established as previously in human ES cells (Takayama et al., 2008). The medium was refreshed every 3 d, and nonadherent cells were collected and analyzed from days 22 to 38.

DOX-inducible c-MYC lentiviral vector. The c-MYC gene-inducible lentiviral vector was based upon EV-TRE-mOXS-Ubc-cTA-12G (Kobayashi et al., 2010) and modified by replacing the mOXS cassette with c-MYC gene. Viral supernatant was generated as previously described (Eto et al., 2007).

Western blotting experiments. Experiments were performed as previously described (Eto et al., 2007; Nishikii et al., 2008). In brief, 45 µg of cell lysates treated with TNE buffer (10 mM Tris-HCl, pH 7.8, 150 mM NaCl, 1% NP-40, and 1 mM EDTA), supplemented with protease inhibitor cocktail (Roche), were separated by electrophoresis on 10–20% SDS-polyacrylamide gradient gel (Bio-Rad Laboratories) and transferred to a polyvinylidene difluoride membrane (Millipore), followed by visualization with SuperSignal West Pico Chemiluminescent Substrate (Thermo Fisher Scientific).

Immunohistochemical and flow cytometric analyses of ES- and iPSC-Sacs. Immunohistochemical staining of iPSC-Sacs was performed on day 14 or 15. Intact iPSC-Sacs were fixed with 10% methanol in PBS, after which they were stained first with an antibody against human VEGF-R2 and then with a secondary antibody and observed using an epifluorescence microscope (DM IRBE; Leica). Round cells within the ES- and iPSC-Sacs were stained with anti-human CD31-PE, CD34-FITC, CD38-APC, CD41a-APC, CD45-Alexa Fluor 405, or VEGF-R2–APC antibodies and analyzed by flow cytometry.

Hematopoietic colony-forming cell assay. Hematopoietic colony-forming cell assays were performed in MethoCult H4434 semisolid medium (STEMCELL Technologies Inc.) supplemented with 50 ng/ml human TPO. 10,000 hematopoietic progenitors from within an iPSC-Sac were plated in 1.5 ml of medium and cultivated for 14 d. The colonies were then collected, stained with Hemacolor (Merck), and observed under a microscope.

Flow cytometric analysis of MKs. Nonadherent cells on days 22–38 were prepared in PBS containing 2% FBS and stained with combinations of antibodies for 30 min on ice. Samples were then washed with PBS and analyzed by flow cytometry (FACSARia, BD).

Viral transduction of hematopoietic progenitors within ES-sacs. A total of 107 hematopoietic progenitors harvested from within an ES-sac or iPSC-sac on day 15 of culture were suspended in hematopoietic differentiation medium containing 50 ng/ml of human SCF, 100 ng/ml of human TPO, 25 μM heparin, and 10 mg/ml protamine sulfate and then transduced with viral supernatant for vehicle, OCT3/4-KO, SOX2-EGFP, KLF4-EGFP, c-MYC-EGFP, or DOX-inducible c-MYC. The cells were then replated into a 6-well plate precoated with C3H10T1/2 cells and centrifuged at 900 rpm for 60 min at 32°C. The viral transduction was performed three times with 12-h intervals in between. To induce c-MYC O/E, 1 µg/ml DOX was added to the culture medium from days 15 to 22 or 26 (Fig. 5 C).

qRT-PCR. cDNA samples were prepared as described in the previous section. Real-time PCR was performed using a kit (TaqMan Gene Expression Master Mix [Applied Biosystems] or SYBR Premix Dimer Eraser [Takara Bio, Inc.]) according to the manufacturer’s instructions. Signals were detected using an ABI7900HT Real-Time PCR System (Applied Biosystem). Primer sets for GAPDH, c-MYC, p14ARF, p16INK4A, GATA-1, β1-tubulin, and NF-E2 p45 were determined using the Universal Probe Library Set for humans (https://www.roche-applied-science.com/sis/rtpcr/upl/index.jsp?id=UP030000). Primer sets for exogenous OCT3/4, SOX2, KLF4, c-MYC, and endogenous c-MYC are listed in Table S1.

Electron microscopic observation of hiPSC-derived platelets. Platelet pellets were fixed with a mixture of 0.5% glutaraldehyde and 2% paraformaldehyde in 0.1 M phosphate buffer, pH 7.4, for 60 min at 4°C. After washing with phosphate buffer, the samples were post-fixed with 1% osmium tetroxide in phosphate buffer for 60 min on ice. After dehydration, samples on coverslips were infiltrated with and embedded in Epoxy resin. Ultrathin sections (60–80 nm thick) were cut and stained with 2% uranyl acetate in 70% methanol and Reynolds’ lead citrate and observed in a transmission electron microscope (1200EX, JEOL) operating at 80 kV.

Flow cytometric analysis of platelets. Washed platelets were prepared as described previously (Takayama et al., 2008). The resultant platelet pellets were resuspended with PBS and stained with anti–human CD41a (integrin αIIbβ3 complex)-APC, GPIb-IX-FITC, GPIa, or CD9-PF for 30 min at room temperature. The platelets were then diluted in 200 µl PBS and analyzed by flow cytometry. Platelet numbers were estimated using true count beads (BD).

In vitro functional analysis of platelets derived from human iPSCs. Collected platelets were resuspended in an appropriate volume of modified Tyrode-Hepes buffer (10 mM Hepes, pH 7.4, 12 mM NaHCO₃, 138 mM NaCl, 5.5 mM glucose, 2.9 mM KCl, and 1 mM MgCl₂) and eventually used after addition of 1 mM CaCl₂. To investigate integrin αIIbβ3 activation, 50-µl aliquots of fresh PB- or PB-based aged, hESC-, and hiPSC-derived platelets (ES platelets and iPSC platelets) were incubated with PE-conjugated anti-GPIbα and FITC-conjugated PAC-1 (Shattil et al., 1985) or FITC-conjugated CD62P (P-selectin) in the absence or presence of human thrombin or ADP for 20 min at room temperature. The binding of PAC-1 to platelets was quantified using an Aria flow cytometer. Non-specific binding was determined in the presence of 10 µM tirofiban, a specific antagonist of human integrin αIIbβ3 (Pfeilnick et al., 1993). Specific binding was defined as total minus non-specific binding.

In vivo imaging by iPSC-platelets. Details of this method are provided elsewhere (Nishimura et al., 2008; Takizawa et al., 2010). In brief, to visualize analyze iPSC platelet function, including circulation and thrombus formation in the mesentery of living animals, mice were anesthetized and a small incision was made in the abdominal wall. Intravital imaging was then performed through this small (~3 mm) window. FITC-dextran (20 mg/kg body weight) was injected into the tail vein for visualization of host blood cell dynamics. iPSC-derived platelets were stained with 5 µM TMRE for 15 min, washed, and injected into the mice. To induce thrombus formation, hematoporphyrin (1.8 mg/kg body weight) was also administrated. Serial two-color images were obtained for 20 s at 30 frames/s using a high-speed spinning-disk confocal laser scanning microscope (CSU-X1) and a pair of EM charge-coupled device cameras (eXon). All experiments were approved by the University of Tokyo Ethics Committee for Animal Experiments and strictly adhered to the guidelines for animal experiments of the University of Tokyo.

Statistical analysis. All data are presented as the mean ± SEM. We used two-tailed Student’s t tests for statistical analysis; values of P < 0.05 were considered significant.

Online supplemental material. Fig. S1 depicts the character of hiPSCs derived from HDFs. Fig. S2 shows the generation of hematopoietic progenitors.
from hiPSCs. Figs. S3 and S4 show the time courses of MK generation. Fig. S5 shows the time course of platelet generation. Fig. S6 depicts the representative dot plots of PB-derived and hiPSC-derived platelets. Fig. S7 describes the time-dependent change of exogenous and endogenous c-MYC expression in hiESCs and hiPSCs. Fig. S8 shows the in vitro functionality of iPS-PC platelets. Video 1 shows a typical proplatelet formation. Video 2 demonstrates in vivo imaging of circulating iPS-PC platelets. Videos 3–5 demonstrate in vivo imaging of iPS-PC platelets or platelets in thrombi by laser-induced vessel wall injury.

Table S1 is a list of primers used in this study. Online supplemental material is available at http://www.jem.org/cgi/content/full/jem.20100844/DC1.

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