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Electrically Guiding Migration of Human Induced Pluripotent Stem Cells

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Abstract A major road-block in stem cell therapy is the poor homing and integration of transplanted stem cells with the targeted host tissue. Human induced pluripotent stem (hiPS) cells are considered an excellent alternative to embryonic stem (ES) cells and we tested the feasibility of using small, physiological electric fields (EFs) to guide hiPS cells to their target. Applied EFs stimulated and guided migration of cultured hiPS cells toward the anode, with a stimulation threshold of <math><30\text{ mV/mm}</math>; in three-dimensional (3D) culture hiPS cells remained stationary, whereas in an applied EF they migrated directionally. This is of significance as the therapeutic use of hiPS cells occurs in a 3D environment. EF exposure did not alter expression of the pluripotency markers SSEA-4 and Oct-4 in hiPS cells. We compared EF-directed migration (galvanotaxis) of hiPS cells and hES cells and found that hiPS

cells showed greater sensitivity and directedness than those of hES cells in an EF, while hES cells migrated toward cathode. Rho-kinase (ROCK) inhibition, a method to aid expansion and survival of stem cells, significantly increased the motility, but reduced directionality of iPS cells in an EF by 70–80%. Thus, our study has revealed that physiological EF is an effective guidance cue for the migration of hiPS cells in either 2D or 3D environments and that will occur in a ROCK-dependent manner. Our current finding may lead to techniques for applying EFs *in vivo* to guide migration of transplanted stem cells.

Keywords Human induced pluripotent stem (hiPS) cell · Electric field (EF) · Cell migration · Wound healing · Galvanotaxis

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Introduction

Human induced pluripotent stem (hiPS) cells show great potential for stem cell based therapies, as they are very similar to human embryonic stem (hES) cells in their ability to self-renew and to differentiate into many types of cell within the body, and are free from the ethical issues to immunological barriers associated with the use of hES cells [1]. While there has been a great deal of interest in the pluripotency and differentiation of hiPS cells, little is known about their migration and guidance cues.

Amongst the various challenging hurdles to successful stem cell therapy, is the poor homing to and integration of transplanted stem cells with target tissue. Poor healing responses and lack of regeneration may be partly due to the fact that very few therapeutic stem cells actually migrate to the damaged/diseased tissue to mount a reparative response [2, 3]. One example of such poor homing responses comes from a model of renal ischemia/reperfusion injury, in which intravenously injected mesenchymal stem cells (MSCs) sequestered mainly in the lung and liver, with less than 1% of injected MSCs reaching the injured kidney [4]. Also, homing of MSCs to infarcted areas in heart has been shown to be very poor, and this contributes to poor survival rate [5]. Effective techniques to stimulate and guide migration of stem cells may therefore offer new strategies to improve the efficiency of stem cell therapies.

Signals that guide and regulate the migration of stem cells include growth factors (GFs), chemokines, molecules of the extracellular matrix (ECM), and their respective receptors. The importance of stromal-derived factor-1 (SDF-1), a well studied chemokine expressed by a variety of cell types including stromal cells and stem cells, and its unique receptor CXCR4 in migration and homing of stem cells has been demonstrated. This SDF-1/CXCR4 signaling plays a critical role in the homing of hematopoietic stem cells [6], the directional migration of neural stem cells to sites of central nervous system (CNS) injury [7], and the directional migration of endothelial progenitor cells [8]. In addition to SDF-1/CXCR4, the fluocoylated CD44 has been shown to promote efficient adhesive interactions of MSCs with marrow vasculature and subsequent homing to endosteal surfaces [9]. Also, it has recently been suggested that insulin-like growth factor 2 (IGF-2)/IGF-2 receptor signaling is essential for homing of endothelial progenitor cells and vasculogenesis at ischemic or tumour sites [10]. In contrast, CD26 has been shown to be a negative regulator of hematopoietic stem cell homing [11].

Applied electric fields (EFs) may be another effective means to guide directional migration of hiPS cells. Damaged tissue and wounds produce weak, direct current (DC) EFs, which direct migration of many cell types. This phenomenon is known as galvanotaxis or electrotaxis [12].

With the aim of developing novel techniques to guide the migration of stem cells, we tested whether EFs direct the migration of hiPS cells. We found that: 1) applied EFs guide migration of isolated or clustered hiPS cells toward the anode, with a threshold voltage of <30 mV/mm; 2) EFs significantly increase migration speed of hiPS cells by 20% at 50 mV/mm, and 40% at 75 mV/mm; 3) EFs directed migration of hiPS cells in three-dimensional (3D) culture; 4) exposure to an EF did not alter expression of the stem cell markers SSEA-4 and Oct-4 in hiPS cells; 5) galvanotaxis of hiPS cells differed from that of hES cells in migratory speed, time-course of migration, to direction of migration; 6) Rho kinase (ROCK) inhibition significantly inhibited the directness of migration, but increased the speed of migration in hiPS cells.

Materials and Methods

Cell Culture

Human iPS cells (foreskin-derived iPS cell line, clone #3, a kind gift from Dr. James Thomson), were maintained on mouse embryonic fibroblasts (MEFs) (Globalstem Rockville, MD) in 6-well tissue culture plates (Thermo Fisher Scientific, Waltham, MA) with hESC medium: DMEM/F12 supplemented with glutamine, 20% knockout serum replacement, non-essential amino acids, 0.1 mM β -mercaptoethanol (all from Invitrogen, Inc., Carlsbad, CA), to fibroblast growth factor 2 (FGF2) (20 ng/ml, Chemicon). For experiments using feeder-free conditions, human iPS cells were transitioned from fibroblast feeders to hESC-qualified matrigel-coated galvanotaxis chamber with small colony clusters. Cells were allowed to grow in mTeSR™ maintenance medium (Stem Cell Technologies, Vancouver, BC, Canada) for approximately 12 h at 37°C with 5% CO₂ prior to experiments. For ROCK inhibition experiments, Y-27632 (Tocris Bioscience, Ellisville, MO) was added to the cells 30 min before exposure to an EF with the final concentration of 2 μ M. The foreskin-derived fibroblast (CRL-2522™, ATCC, Manassas, VA) was used as the control for hiPS cells.

Human embryonic stem cell line H7 (WiCell Research Institute, Madison, WI) was cultured in mTeSR™ maintenance medium at 37°C with 5% CO₂ with daily medium change. One day prior to the EF experiments, hESCs were treated with 1 mg/ml dispase (Invitrogen, Carlsbad, CA) and manually detached with a cell scraper. Small colony clusters were then plated in the hESC-qualified matrigel-coated electrotactic chamber in mTeSR™ culture medium, at a variety of cell densities up to $3\sim 5 \times 10^3$ cells/cm². Cells were allowed to attach and grow for approximately 12 h at 37°C with 5% CO₂ prior to experiments.

3D Matrigel Culture

For 3D culture of human iPS cells, a stepped dilution using a combination of hiPS cell-qualified matrigel and mTeSR™ medium was tested to optimize the culture condition. One day prior to the EFs experiments, hiPS cells were treated with 1 mg/ml dispase (Invitrogen, Carlsbad, CA) for 8 min and manually detached with a cell scraper. Cells at density of $0.5\text{--}1.0 \times 10^5/\text{ml}$ were transferred to a 0.5-ml tube and mixed well with matrigel at a volume ratio of 1:1 by gently pipetting up and down to distribute the colonies. 20 μl of matrigel and cells mixtures were then transfer to a custom designed 3D electrostatic chamber in 60-mm dishes, and placed in an incubator for 20 min to allow matrigel to polymerize. Once gel was formed, 4.5 to 5.0 ml mTeSR™ medium was added into the dishes. Cells were allowed to grow overnight at 37°C with 5% CO₂.

EFs Stimulation and Time Lapse Image Recording

Prior to EF exposure, mTeSR™ containing 25 mM HEPES buffer was added to one well of the chamber and the culture medium was observed to fill the opposite well assuring that the chamber was not clogged. EFs were applied to cells through two silver electrodes immersed in Steinberg's solution-filled reservoirs that were connected to wells at each end of the galvanotaxis chamber by two agar bridges (2% agar in Steinberg's solution). hiPS or hES cells were exposed to an EF with strength ranging from 30 to 200 mV/mm for 3–8 h at 37°C. Field strengths were measured directly at the beginning and the end of the experiment. Phase contrast (for 2D) and DIC (for 3D) time lapse imaging was performed using Nikon Eclipse imaging system equipped with a CO₂ and temperature control chamber. 3D scanning was performed at various Z focal planes across the whole thickness of matrigel containing hES or hiPS cells colonies. Data analysis was performed using the NIH Image J software.

Quantitative Analysis of Cell Behavior

Cell migration was analyzed as previously reported [36, 37]. We used the NIH Image J software to quantify migration directedness and trajectory speed by tracing the position of cell nuclei at frame intervals of 10 min. The directedness of migration was assessed as cosine θ , where θ is the angle between the EFs vector and a straight line connecting the start and end position of a cell. A cell moving directly along the field lines toward the anode would have a directedness of -1 ; a cell moving directly toward the cathode would have a directedness of 1 . A value close to 0

represents random migration. The cosine θ will range from -1 to $+1$, and an average of all the individual cells yields the directedness value for a population of cells, giving an objective quantification of the direction of cell migration. The trajectory speed (Tt/T) is the total migration distance of cells per minute; Dx/T is an x-axis displacement speed, which represents the migration along the EF vector.

Immunostaining

hiPS cells were seeded onto chambered glass-bottom dishes (Thermo Fisher Scientific) that were pre-coated with matrigel. Before or after EF stimulation, cells were washed with phosphate buffered saline (PBS), fixed with 2% paraformaldehyde for 15 min at room temperature. After 3 washes with PBS, cells were blocked with 3% bovine serum albumin (BSA) (Sigma-Aldrich). Cells were then incubated with Alexa Fluor 568 phalloidin (Invitrogen, USA) for 20 min at room temperature or primary antibodies against monoclonal mouse anti-SSEA-4 (Developmental Studies Hybridoma Bank, Iowa) at 1:50 dilution and rabbit anti-Oct-4 (Chemicon) at 1:500 dilution overnight at 4°C. After washes with PBS, cells were incubated with Alexa Fluor conjugate-secondary antibodies (5 $\mu\text{g}/\text{ml}$; Invitrogen) for 30 min at 4°C, stained with Hoechst 33342 (2 $\mu\text{g}/\text{ml}$; Invitrogen) for 5 min, mounted using Fluoro-Gel with Tris Buffer (Electron Microscopy Sciences, Hatfield, PA), and imaged using an Olympus BX51 fluorescence microscope (Olympus, Center Valley, PA).

Statistical Analysis

Data are expressed as mean \pm SEM. Statistical differences between groups were assessed with unpaired, two-tailed Student's *t*-test, with statistical significance set at $p < 0.05$. Statistical analysis was performed using the software of SPSS 10.0.

Results

EF-Directed Migration of hiPS Cells

First of all we determined the migratory behavior of hiPS cells in small colonies of 20–150 cells. The colonies were relatively uniform and abundant. The following description, unless otherwise indicated, was from these colonies of hiPS cells. In the absence of an EF, no obvious directional migration was observed (Fig. 1a and b; supplemental movie 1). hiPS cells migrated very little and in random directions, as shown by the migration trajectories of individual cells (Fig. 1c). In an EF, hiPS cells migrated

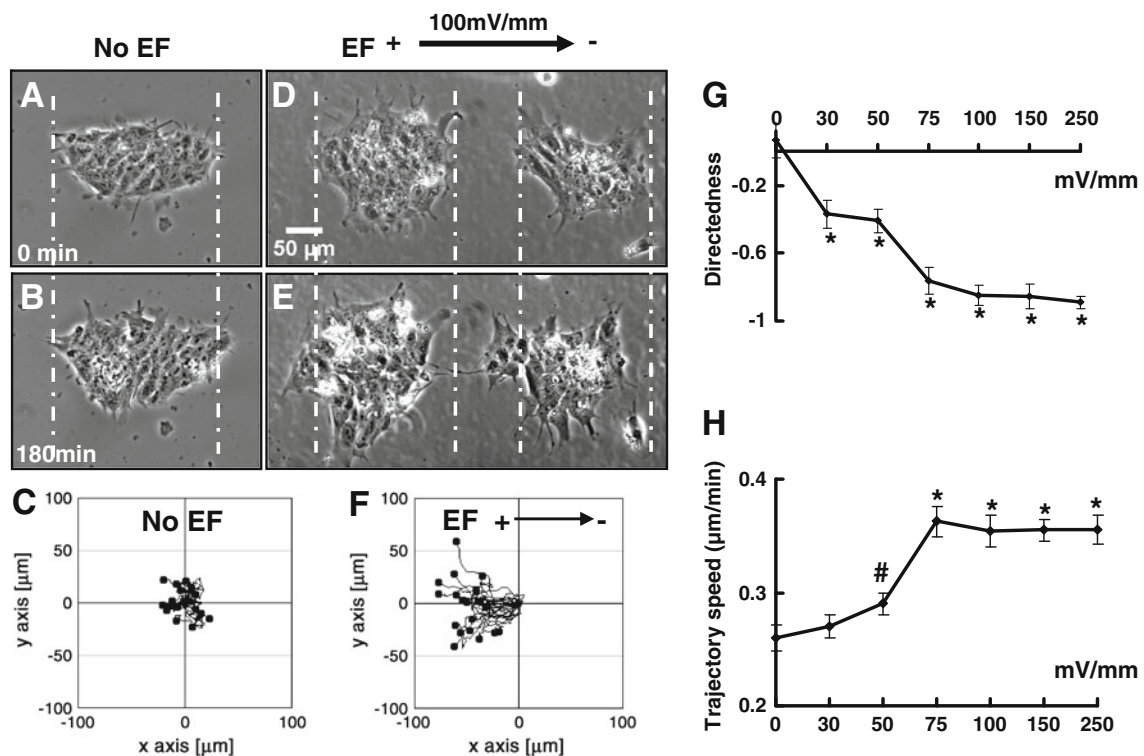


Fig. 1 Electric fields guided migration of hiPS cells in a voltage dependent manner. **a** and **b** In the absence of an EF, a hiPS cell colony showed no migration over a period of 3 h. **c** Tracking of individual cells in the colony showed very little migration and any migration was in random directions. **d** and **e** An EF instigated and directed migration of the two colonies of hiPS cells toward the anode (left). **f** Tracking of

individual cells showed the EF directed anodal migration, with increased motility. EF = 100 mV/mm; Scale bar, 50 μm. See Supplementary movies 1 and 2. **g** Migratory directedness of hiPS cells increased in a voltage dependent manner, with a threshold of 30 mV/mm or below. **h** EFs significantly increased the trajectory speed. #, $P < 0.05$; *, $P < 0.01$ compared to the no EF control

directionally toward the anode, at an increased speed (Fig. 1d, e and f; supplemental movie 2). Directional migration could be detected within ~10 min following the onset of the EF. Leading cells were seen to extend lamellipodia and filopodia toward the anode, while cells at the trailing edge were seen to retract extensions, and the whole colony migrated collectively. Cells largely maintained their relative position in the colony. We confirmed the EF-directed migration of hiPS cells by reversal of the EF polarity. Following reversal of the field polarity, hiPS cells changed their direction of migration and migrated toward the new anode (Fig. S1; supplemental movie 3).

To determine whether the density of hiPS cells affects EF-directed migration, we analyzed migration of hiPS cells in dissociated culture and culture of large colonies (>300 cells). Both the dissociated cells and the large colonies migrated toward the anode in an EF (Fig. S2; supplemental movie 4, 5). Galvanotaxis of hiPS cells therefore remained very similar between dissociated cells and tightly connected cells in colonies, although their responses varied in terms of speed (dissociated cells: 0.50 ± 0.02 μm/min vs. large colonies: 0.04 ± 0.01 μm/min, $p < 0.05$) and directedness (dissociated

cells: -0.61 ± 0.06 vs. large colonies: -0.96 ± 0.01 , $p < 0.01$). Cells in colonies migrated more directionally, and the larger the colony, the more uniformly they migrated toward the anode (Fig. 1f, S2 D, S2 H).

Voltage-Dependent Galvanotaxis of hiPs Cells

In order to determine the threshold voltage required to induce galvanotaxis, we analyzed the voltage-dependent galvanotaxis of hiPS cells. Significant directional migration was observed at a field strength of 30 mV/mm with directedness of -0.37 ± 0.08 (Fig. 1g) ($p < 0.01$, when compared with the no EF control). The threshold voltage required to induce galvanotaxis of hiPS cells was therefore <30 mV/mm. EFs of higher voltage significantly increased directedness, which doubled in an EF of 75 mV/mm. Applied EFs also significantly increased trajectory speed. The increase in speed was dependent on field strength, with an EF of 75 mV/mm increasing the speed by 40% and this leveled out at higher voltages (Fig. 1h). Consistent with the directional migration, displacement along the χ axis (the field line toward the anode) in hiPS cells significantly increased following EF stimulation (Fig. S3). These results

indicate that a robust galvanotaxis of hiPS cells occurs *in vitro* in an EF of 75 mV/mm, under our experimental conditions, with a threshold of <30 mV/mm.

Galvanotaxis of hiPs Cells in 3D Matrigel

Next we tested whether hiPS cells would migrate in 3D culture and whether migration in 3D culture could be instigated and guided by applied EFs, as therapeutic use of EFs to guide cell migration will occur in a 3D environment. 3D electrostatic chambers were constructed, in which hiPS cells were cultured in 3D matrigel matrix. In the absence of an EF hiPS cells remained almost motionless over 5–8 h of observation in 3D culture (Fig. 2a and e). Directedness and trajectory speed were close to zero (Fig. 2c and d; supplemental movie 6, 8). Application of an EF instigated the formation of lamellipodia and filopodia at the anodal side of colonies of hiPS cells in 3D culture, and induced significant directional migration toward the anode in a voltage-dependent manner, with a threshold of <50 mV/mm (Fig. 2b, c, d and f). Both small and large colonies of hiPS cells extended polarized protrusions toward the anode and retracted protrusions facing the cathode (Fig. 2b and f; supplemental movie 7, 9). Cells migrated in the Z direction as well as toward the anode, therefore some cells gradually migrated out of focus. In large colonies, a group of leading cells formed, and the trailing cells migrated in lieu (Fig. 2f; supplemental movie 9). These results indicate that an EF can induce directional migration of otherwise stationary colonies of hiPS cells in a 3D environment. This may be of clinical significance as transplanted hiPS cells will reside in 3D tissues.

EF Exposure Did Not Alter Expression of Stem Cell Markers in hiPs Cells

In order to investigate whether EF exposure had any effect on the pluripotency of hiPS cells, we stained two stem cell markers, SSEA-4 and Oct-4, in hiPS cells following 5 h of EF exposure (100 mV/mm). Both SSEA-4 and Oct-4 were highly expressed in hiPS cells exposed to an EF (Fig. S4). The expression levels were similar to untreated control cells, indicating that EFs have no significant effect on the pluripotency of hiPS cells.

Galvanotaxis of hiPs Cells was Different from That of hES Cells

We compared the galvanotaxis of hiPS cells and hES cells because recent research has revealed differences in gene expression signatures between iPS cells and ES cells, although the pluripotency of hiPS cells and hES cells are thought to be similar [13]. Surprisingly, hES cells (H7)

migrated toward the cathode, in the opposite direction of hiPS cells (Fig. 3a; supplemental movie 11 vs. supplemental movie 10). The migratory directedness of both hiPS cells and hES cells increased in a time-dependent manner. However, hiPS cells were more sensitive than hES cells and responded better. hiPS cells showed earlier galvanotaxis with directedness reaching a significant level (-0.70 ± 0.08) following 1 h in an EF, while hES cells took over 3 h to show significant directional migration, with lower directedness values (0.14 ± 0.12). The directedness of hES cells never reached a level comparable to that of hiPS cells (Fig. 3b). hES cells showed a higher trajectory speed than hiPS cells in an EF, although no difference was observed between the two cell types in the absence of EFs (Fig. 3c).

Rho/ROCK Signaling was Required for Galvanotaxis of hiPs Cells

To examine the underlying signaling mechanisms involved in the galvanotaxis of hiPS cells, we focused on the Rho/ROCK signaling pathway, as Rho kinase (ROCK) inhibition has been used to promote the survival of iPS cells [14], and we have previously demonstrated its involvement in EF-directed migration [15]. hiPS cells were treated with Y-27632 (2 μ M), a specific inhibitor of ROCK, for 30 min prior to EFs application. EF-induced anodal migration of hiPS cells was significantly suppressed (Fig. 4b; supplemental movie 15 vs. supplemental movie 14). Quantitative analysis showed a sharp decline in migratory directedness from -0.89 ± 0.03 in the no drug controls, to -0.22 ± 0.08 in the Y-27632-treated hiPS cells, suggesting a critical role for Rho/ROCK signaling in the galvanotaxis of hiPS cells (Fig. 4e). Y-27632-treated cells showed increased motility both in the absence and presence of an EF (Fig. 4c and d; supplemental movie 13 vs. supplemental movie 12; supplemental movie 15 vs. supplemental movie 14). The trajectory speed of Y-27632-treated hiPS cells increased by 36% and 34% in the absence and presence of an EF respectively (Fig. 4f). Treatment with Y-27632 induced two distinct morphological changes: 1) loss of close cell-cell contacts; 2) multiple protrusions (Fig. 4a and b; Fig. S5; supplemental movies 13 and 15).

Discussion

With the aim of developing novel techniques to guide human stem cells, to facilitate stem cell therapy, we investigated the application of EFs to guide hiPS cell migration in both 2D and 3D cultures. We reported that: 1) a DC EF guided migration of hiPS cells in 2D culture; colonies showed significantly greater directionality than dissociated cells; 2) hiPS cells remained immotile in a 3D

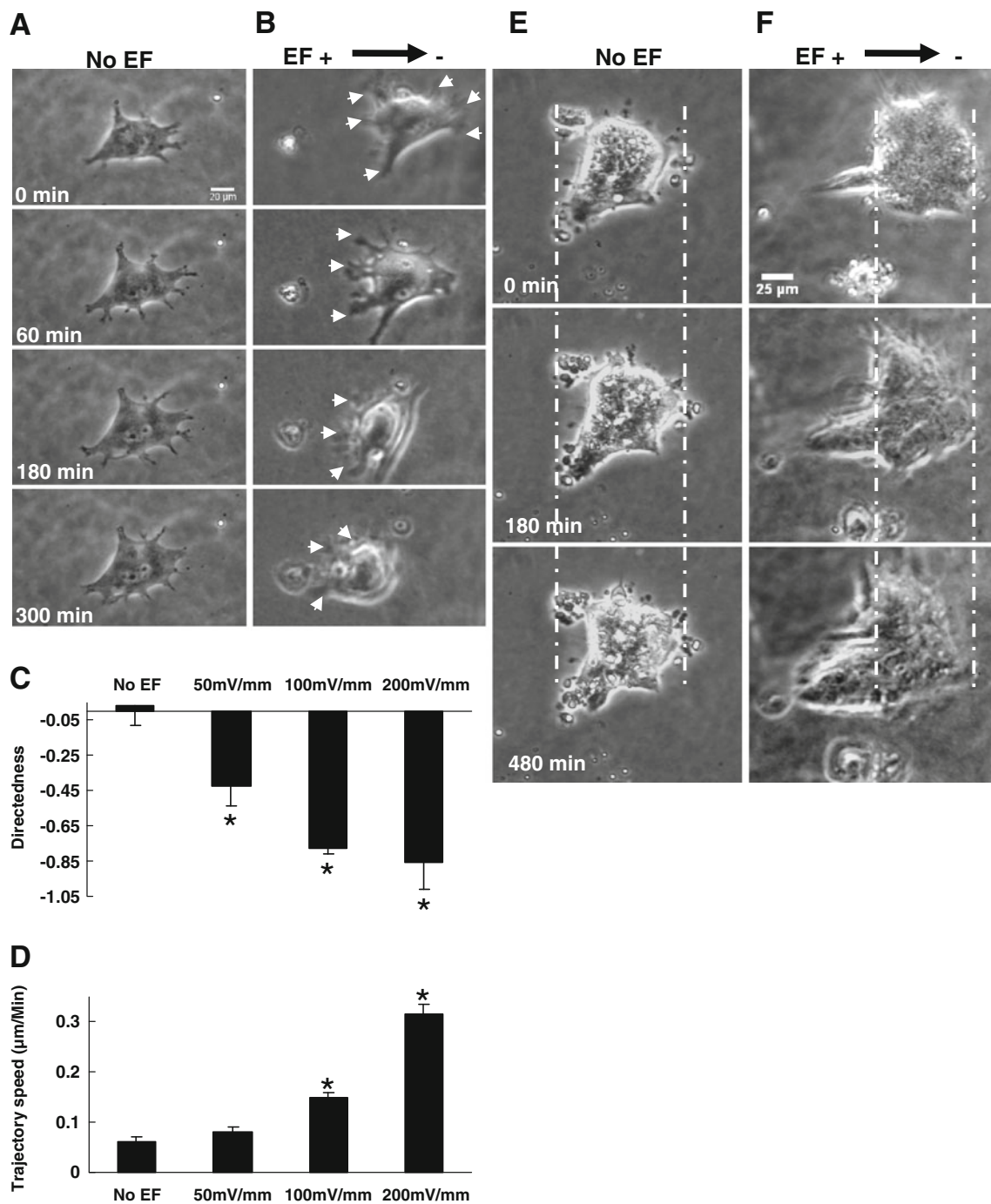


Fig. 2 EFs stimulate and guide migration of hiPS cells in 3D matrigel. **a** and **e** Small and large colonies of hiPS cells remained immotile in control culture without an EF over 5 and 8 h, respectively. **b** A small colony extended multiple protrusions to the anode and migrated in the same direction. **f** A large colony enlarged a leading process toward the anode and the whole colony migrated to the anode

(left). Scale bar, 20 μm in A and B; 25 μm in E and F. See Supplementary movies 6, 7, 8 and 9. **c** and **d** EFs significantly increased trajectory speed (C) and migratory directedness (D) of hiPS cells in 3D matrigel in a voltage-dependent manner, with a threshold of <math><50\text{ mV/mm}</math>. *, $P<0.01$ compared to the no EF control

environment in the absence of an EF, while a small EF induced significant directional migration; 3) electrically guided migration of hiPS cells showed voltage dependence with a threshold voltage of <math><30\text{ mV/mm}</math>; 4) EF exposure

did not affect the expression of stem cell markers; 5) hiPS cells migrated toward the anode with greater sensitivity and directedness compared to hES cells which migrated toward the cathode; 6) ROCK inhibition by Y-27632 significantly

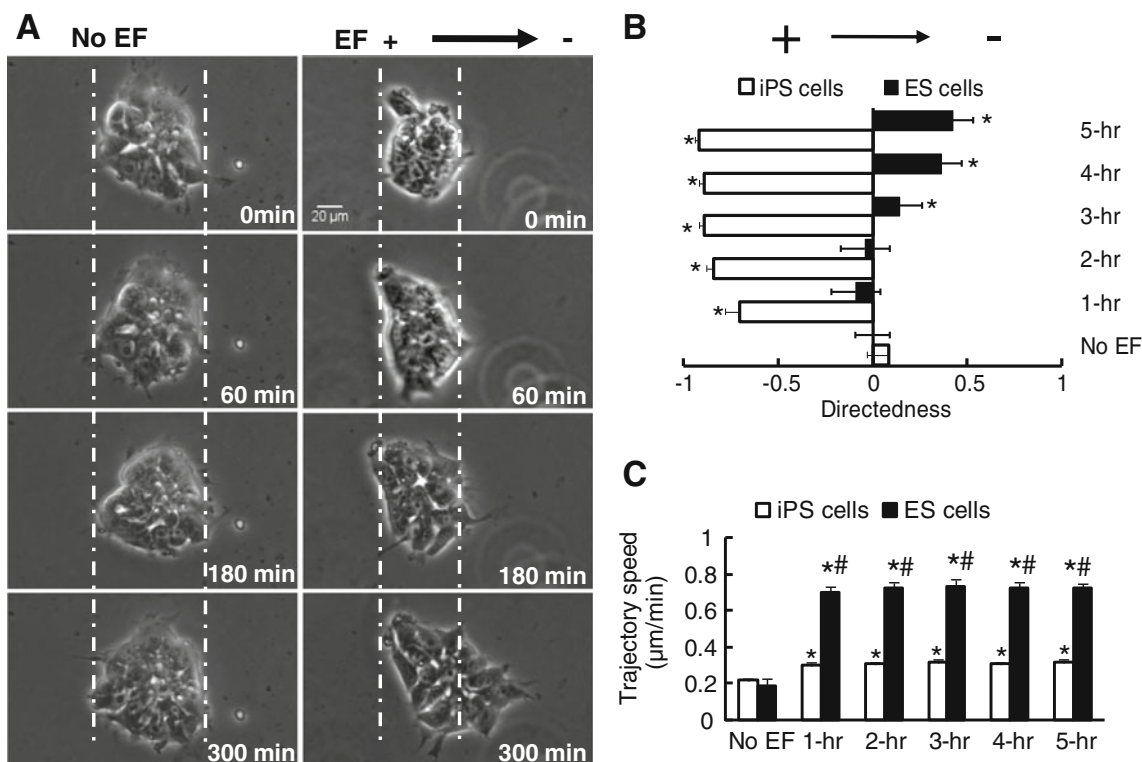


Fig. 3 Galvanotaxis of hiPS cells was different from that of hES cells. **a** No migration (left panel) of a colony of hES (H7) cells occurred in the absence of an EF. Right panel showed directional migration toward the cathode (right) of a hES cell colony in the presence of an EF. Scale bar, 20 μm . **b** hiPS cells and hES cells migrated in opposite directions in an EF. hiPS cells migrated toward the anode, whereas hES cells migrated toward the cathode. hiPS cells manifested robust

galvanotaxis with directedness of -0.70 ± 0.08 within 1 h in an EF, whereas in the same field, hES cells presented a directedness value of only 0.14 ± 0.12 after 3 h. **c** EFs significantly increased the trajectory speed in both hiPS and hES cells. *, $P < 0.01$ when compared to the no EF control; #, $P < 0.01$ compared to hiPS cells. EF = 100 mV/mm. See Supplementary movies 10 and 11

reduced the directedness of galvanotaxis but increased motility.

Several potential methods to direct migration of stem cells have been explored, in order to overcome the barrier of poor homing and integration of stem cells with the targeted tissues. These methods include: 1) enhancing the chemotaxis of stem cells by genetic manipulation of chemokines and their receptors such as SDF-1/CXCR-4 [8]; 2) increasing the migratory potential of stem cells by cytokine pretreatment or its receptor overexpression [16, 17]; 3) removing the extracellular matrix (ECM) barrier by matrix metalloprotease stimulation to accelerate the migration of stem cells [18]. In addition to those biochemical methods, a biophysical approach using a magnetic field has been shown to offer a guidance cue for the migration of neural stem cells [19]. However, effective and practical techniques to guide stem cell migration have not yet been established.

We have shown here that small applied EFs can be an effective signal to stimulate and direct migration of hiPS cells. Endogenous EFs occur in the body at sites of wounds and damaged tissue [20–22]. Applied EFs guide migration

of many types of cells. We have recently provided evidence that EFs could be a predominant guidance cue for corneal epithelial wound healing, over-riding contact inhibition release, free edge, injury stimulation and other well-accepted guidance cues [12].

There are several important points to consider when developing methods of electrical stimulation and guidance of hiPS cells for possible practical use. First is the threshold of applied EFs. 3D tissues are volume conductors in which EFs dissipate exponentially. To maintain a constant high-voltage gradient in electrically conductive tissue inevitably involves issues such as electrode products, heat generation, effects on other types of cells, and permeability of blood vessels. Low voltage offers many advantages. In our experiments, an EF of 30 mV/mm induced significant directional migration. This is a voltage below the thresholds for galvanotaxis of fibroblasts and endothelial cells [23, 24]. Application of an EF of this strength may therefore selectively guide migration of hiPS cells without significant effects on fibroblasts and endothelial cells. The second point is the migration of clusters of cells. Large

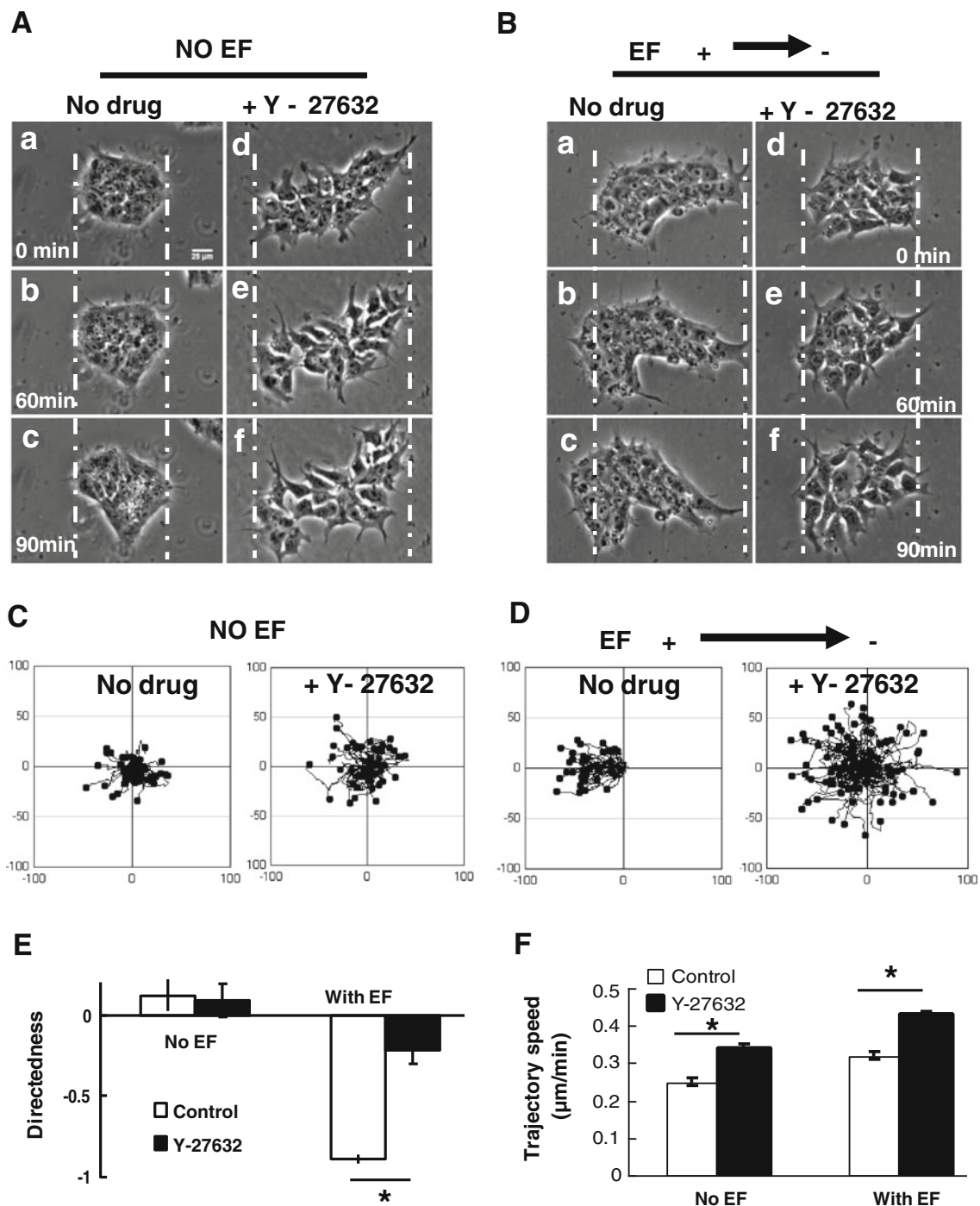


Fig. 4 Rho A/ROCK signalling required for galvanotaxis of hiPS cells. **a** and **b** Y-27632, a ROCK inhibitor, induced a tendency for dissociation of cells from the colony to inhibited galvanotaxis. **c** and **d** Inhibition of Rock by Y-27632 increased the motility of hiPS cells and inhibited the directional migration. **e** and **f** Quantitative analysis

showed a sharp decline in migratory directedness and a significant increase in the trajectory speed in Y-27632-treated hiPS cells. Scale bar, 25 µm. EF = 100 mV/mm. See Supplementary movies 12, 13, 14 and 15

numbers of hiPS cells will need to be used for transplantation. Clusters of cells are normally immotile, although some cells may move within the clusters (Figs. 1a, 2a and e). This poses a problem for the transplanted cells to integrate

with the host tissue. Very encouragingly, we have shown that an applied EF was able to stimulate and induce directional migration of clusters of hiPS cells. Thirdly, transplanted cells have to migrate in a 3D environment. We have shown that

hiPS cells did not migrate in 3D in matrigel in the absence of an EF (Fig. 2a and e). Excitingly, applied EFs stimulated and guided migration of hiPS cells. Exogenously applied EFs have been safely used in the treatment of chronic wounds, neurorehabilitation, and spinal cord injury [25–27]. Our results thus provide a basis for developing stimulation techniques for guiding migration of transplanted hiPS cells *in vivo* where they need to migrate in 3D.

There are a number of important differences in galvanotaxis between hiPS cells and hES cells, although the biological potency and epigenetic state of iPS cells are indistinguishable from those of ES cells [28]. Surprisingly, hES (H7) cells migrated in a direction opposite to that of hiPS cells in an EF; hES cells migrated toward the cathode (Fig. 3). Since cathodal migration is the most popular migratory response in many different types of cells, we employed the foreskin dermal fibroblasts, from which the hiPS cells were generated in this study, as the cell control, and found that the non-programmed fibroblast also migrate toward the anode in EFs, same to that observed in hiPS cells (supplemental figures 6 and 7; supplemental movies 16, 17, 18 and 19). In addition, hES cells were less sensitive to an EF. The same voltage (100 mV/mm) took 3–4 times longer time to induce a directional response in hES cells and the response was much weaker (Fig. 3b). Thus, these results revealed new differences in cellular behavior between iPS and ES cells. Whether these differences are due to the different gene expression signatures will need to be studied [13]. Nevertheless, electrical stimulation might be particularly suitable in therapies using iPS cells compared to ES cells.

ROCK is known to be important to maintain iPS and ES cell cultures and we found that it critically influenced galvanotaxis of hiPS cells. ROCK is the downstream effector kinase of Rho A, a family member of the Rho GTPases. Rho A/ROCK signaling has been implicated in cytoskeleton remodeling, cell adhesion, membrane protrusion, and cell migration [29–31]. Recent studies demonstrated that inhibition of ROCK with Y-27632 was an effective means of increasing the survival rate of stem cells including hES and hiPS cells [14, 32]. The ROCK inhibitor therefore is considered as a promising molecule to aid in the expansion of pluripotent stem cells for regenerative medicine [33]. We and others have previously found an important role for ROCK in galvanotaxis [15, 34]. Y-27632 at 2 μ M, a concentration much lower than that normally used for enhancing the survival rate of stem cells, dramatically decreased the directedness of hiPS cells in an EF by 75%. ROCK activation thus is involved in galvanotaxis of hiPS cells. A significant decline in directional migration following ROCK inhibition was associated with an increase in the migratory speed of hiPS cells (Fig. 4e and f). Reduced cell-cell contact and increased filopodium formation in hiPS cells also occurred following ROCK inhibition in hiPS cells (Fig. 4a and b, Fig. S5). Our results are consistent with a

previous report suggesting ROCK inhibition promoted cell motility in mouse ES cells [35]. When considering guiding stem cell migration electrically, the effect of ROCK inhibition will be an important factor.

In summary, a very small EF (30 mV/mm) is an effective cue to guide and stimulate migration of hiPS cells. Most importantly, applied EFs can direct migration of large clusters of hiPS cells which are normally stationary in a 3D environment. This response depends on Rho/ROCK signaling. The size, duration and direction of applied EFs can be relatively easily controlled. Electrical stimulation may offer a practical approach to facilitate stem cell therapy, where guided cell migration and integration with host tissue are needed. These results may lead to techniques for applying EFs *in vivo* to guide migration of transplanted stem cells.

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Author contribution MZ and YH designed the research; JZ, MC, QZ and XZ performed the research; WD and RL contributed reagents and cell lines; JZ and MZ analyzed the data; JZ and MZ wrote the paper with help from WD to XZ.

Conflicts of interest None

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