Purification, Expansion and Characterization of Putative Murine Cardiac Progenitor Cells

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Abstract

Ischemic heart disease kills more people than any other condition. Medical treatment for ischemic heart disease is currently limited by the heart’s lack of regeneration after injury. Recent reports have suggested the existence of cardiac progenitor cells in the adult normal and diseased mammalian heart. The origin of these cells is unclear. We implemented novel culturing conditions to isolate putative cardiac progenitor cells (pCPCs) from the adult murine heart atrium with the properties of cardiac regeneration. Putative adult cardiac progenitor cells were purified and expanded from adult murine hearts by expansion medium supplemented with 2% fetal calf serum, epidermal growth factor, platelet-derived growth factor and leukemia inhibitory factor. Under the culture condition of 5% O2, these cells can be expanded beyond 42 days, expressed stem cell antigen (Sca-1) and cardiac-specific transcription factors. When treated with oxytocin in vitro, these cells express cardiac contractile proteins and when injected intramuscularly in the tibialis anterior muscle these cells give rise to cardiomyocyte-like cells. In contrast, transplantation of these pCPCs in uninjured heart did not result in cardiomyocyte differentiation suggesting that the heart environment is less permissive than the skeletal muscle of cardiosogenesis. These results suggest that cells from the adult murine heart selected with this culture conditions and transplanted in the skeletal muscle have cardiogenic potential. Thus, this approach warrants further investigation for the therapeutic development of strategies for cardiac tissue engineering and myocardial regeneration.

Keywords: Cardiac progenitor cells; Sca-1; Oxytocin; Cardiac regeneration

Introduction

Ischemic heart disease kills more people than any other condition [1]. Current medical treatment can partially improve the outcome of an infarct. Heart transplantation, the only current treatment to replace the lost muscle, is limited by the need for lifelong immunosuppression and limited supply of donor hearts [2,3]. Cell-based therapies are currently being investigated as alternative treatments to achieve cardiac repair [4].

Recent studies have shown that the adult mammalian heart cells can regenerate after injury [5,6]. Several different sources of origin for the cardiac progenitor populations have been examined and characterized. Putative cardiac progenitor populations have been reported including, c-Kit cell surface receptor expressing cells [7], Islet-1 marker expressing cells [8,9], side population cells (SP) [8,10,11], and cells expressing the stem cell antigen receptor (Sca-1+) [12,13]. It has been suggested that Sca-1+ cells play a regenerative role following myocardial infarction (MI) based on the observation that the number of endogenous Sca-1+ CD 31+ cells increased significantly 7 days after MI, as did the expression of Sca-1 protein [9]. Some studies suggest that c-Kit+ and cardiac SP cells in the heart may arise from the bone marrow [14,15].

Although differentiation capacity of the reported cells has been described in terms of protein expression characteristic of cardiogenesis, contractile and energetic compatibility between graft and host is more difficult to determine. A recent study showed poor survival and no long-term functional benefits of cardiac stem cell transplants [16]. We feel that until robust cardiac regeneration can be established with certainty at a functional level, the use of the term “putative” to describe examined cells is prudent. In this report, we therefore confirm the presence of putative cardiac progenitor cells (pCPCs), in the adult murine heart, using a modified culture condition protocol that results in high enrichment of Sca-1+ cells from the heart atrium [17]. These cells can differentiate in vitro and in vivo into putative cardiomyocytes. Notably, cardiomyocyte differentiation of transplanted cells was only observed during pCPC transplantation into the skeletal muscle, but not the uninjured heart, suggesting that the normal heart environment is less permissive than the skeletal muscle of cardiosogenesis.

Materials and methods

Animals

Cardiac cell suspensions were prepared from the enhanced green fluorescent protein (GFP) transgenic mice with C57BL/6 background (C57BL/6-Tg (CAG-EGFP) 10sb/J mice, 9-24 weeks of age, n=3). Animal studies were conducted according to NIH-approved guidelines, as well as approved guidelines of the institutional animal care and use committee of the University of Washington.

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Preparation and expansion of pCPCs from the adult murine heart

Atrium and ventricle heart portions were separated and cardiomyocytes depleted cardiac cell suspensions were prepared as follows. Minced cardiac muscle tissue was digested with 2mg/ml collagenase type IV, 1.2 U/ml dispase II (Worthington, Lakewood, NJ), and 2mM CaCl₂ in PBS for 45 minutes at 37°C. Samples were pipetted up and down every 15 minutes to fracture clumps and dissociate mononuclear cells. The digestion was neutralized by adding twice the original volume HAM’S/F10 supplemented with 15% horse serum (HyClone, Logan, UT) followed by filtration with a 40 μm nylon cell striainers (BD Falcon, Franklin Lakes, NJ) then centrifuged at 300g for 5 minutes. To eliminate erythrocytes, the cell pellet was treated with 5 ml hemolytic buffer (155 mM NH₄Cl, 10 mM KHCO₃, and 0.1 mM EDTA in H₂O) for 4 minutes at room temperature. Cell suspensions were counted and centrifuged again at 300g for 5 minutes. 2 x10⁶ cells were plated in 5 ml expansion medium on standard polystyrene tissue culture dishes (BD Falcon, Franklin Lakes, NJ). Expansion medium consisted of 60% low glucose DMEM (HyClone, Logan, UT), 40% MCDB 201, 2% fetal calf serum (FCS), and 0.1 mM dexamethasone, and 10⁻⁴ M ascorbic acid 2-phosphate (all from Sigma, St. Louis, MO). 100 U penicillin, 1000 U streptomycin (HyClone, Logan, UT), supplemented with 1x10⁻⁴ units/ml leukemia-inhibitory factor (LIF-ESGRO, Millipore), 10 ng/mL EGF (Sigma) and 10 ng/mL PDGF-BB (R&D). Cultures were incubated at 37°C, 5%CO₂ and 5%O₂. Media was initially changed at day three and subsequently replaced every 15 minutes. To eliminate erythrocytes, the cell pellet was treated with 5 ml of original volume HAM’S/F10 supplemented with 15% horse serum (HyClone, Logan, UT) per manufacturer’s instructions. Primers for cardiomyocyte gene amplification were as follows. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH): sense, 5'-CTC GTC CGC TAG ACA AAA TGG-3', antisense, 5'-CGC TTC TGG AAC ATG GTG-3'; GATA-4: sense, 5'-CCC GGG CTG TCA TCT CAC TA-3', antisense, 5'-GTC CCC CAG CTT ATT TTG-3'; GATA-6: sense, 5'-AGC AGG ACC CTT CGA AAC G-3', antisense, 5'-GGC GCT CTG TGG CTT GAT G-3'; Nkx2.5: sense, 5'-CCA AGT CTC CTT CTC TTC-3', antisense, 5'-GAG ACC CTT CGA AAC G-3', antisense, 5'-GGC CCT CTG TGG CTT GAT G-3'; Myf-5: sense, 5'-AGA AGG ACA AAC C-3', antisense, 5'-AGC ACC GCC TCT CTT CTC-3', antisense, 5'-AGT GGT GCC TCT GTA AAG-3'; C-kit: sense, 5'-AGG CTG CGC TTT CCT CAG T-3', antisense, 5'-GCC GCA AAT CGC AAA TCT TT-3'; Oct4: sense, 5'-CTG GGC GTT CTC TTT GGA AA-3', antisense, 5'-TCG GCC GCC TAC GAA ACA AG-3'; Nanog: sense, 5'-TCT CAA GTA CTC AGG AGC ACA AG-3'; antisense, 5'-GTG CGT AGC ACG GAA ACC TCA CTA-3'; Ki67: sense, 5'-AGT TGG GGT TGG TCG TGG TCT CTA-3', antisense, 5'-AGC CCA GTT CCG AAA TCT TT-3'; biotinylated anti-mouse or goat antibody (Sigma, St. Louis, MO) at 1:500 dilution, monoclonal rat anti-Oct4 antibody (BD Pharmingen, San Jose, CA), at a 1:400 dilution, followed by anti-rat Alexa 594 conjugated antibody (Invitrogen, Carlsbad, CA). Antibodies were diluted and combined in PBS + 0.3% bovine serum albumin (BSA) at a ratio of 600 ng of each antibody per 10⁵ cells, except for Sca-1 antibody, which was added at 300 ng per 10⁵ cells due to its strong avidity. 10⁵ cells per single color controls were stained with 200 ng of each antibody/fluorophore along with one containing only cells for unstained control. Samples and controls with antibodies in PBS + 0.3% BSA were incubated on ice in the dark for 45 minutes. Cells were washed and centrifuged for 5 minutes at 300 g for unpermeabilized cells or 800 g for permeabilized cells then resuspended in PBS+0.3% BSA. FACS- analysis was performed using a BD FACSAria II (BD Biosciences, San Jose, CA). Software compensation of single color controls were used to correct the potential spectral overlap between color filters and fluorophores. The cells positive for Sca-1, c-kit, Nanog, and Nkx2.5 were analyzed by the FlowJo v7.54 (TreeStar, Ashland, OR).

Immunostaining of cultured pCPCs

Immunofluorescence staining was done using a monoclonal Cy3 conjugated anti-

Animal surgery and cell transplantation

Two types of transplant studies were conducted; cultured cells directly injected intramuscularly in the tibialis anterior (TA) muscle
and the heart. Mice were anesthetized with isofluorane. 10^6 cultured pCPCs resuspended in 30 µl PBS were directly injected into the TA muscle of 2-month-old C57BL6 mice (n= 8) and 2x10^6 cultured cells in 80 µl PBS were directly injected into the heart of 12-month-old C57BL6 mice (n= 3). All mice were sacrificed 2 weeks after the injection. Euthanasia and animal injections were in compliance with local guidelines and protocols approved by the Institutional Animal Care and Use Committee.

Immunofluorescent staining of transplanted tissues

10-µm cryosections of transplanted tissue embedded in optimal cutting temperature compound (OCT) on glass slides were stained sequentially for GFP and troponin-T and were also stained with 49,6-Diamidino-2-phenylindole dihydrochloride (DAPI), purchased from Vector Labs (Burlingame, CA). Polyclonal goat anti-GFP Ab (ABCAM, Cambridge, MA) was used at 1: 100 dilution followed with anti-goat Alexa 488 antibody (Invitrogen, Carlsbad, CA) diluted at 1:1000. Polyclonal rabbit anti-troponin-T Ab (ABCAM, Cambridge, MA) was used at 1: 1000 dilution, followed with anti-rabbit Alexa 594 antibody (Invitrogen, Carlsbad, CA) diluted at 1:1000. Polyclonal rabbit anti-connexin 43 Ab (ABCAM, Cambridge, MA) was used at 1: 500 followed with anti-rabbit Alexa 594 antibody (Invitrogen, Carlsbad, CA) diluted at 1:1000. All immunofluorescence described in this manuscript was detected using a Zeiss Axiovert 200 fluorescence microscope (Thornwood, NY). Photographs were taken with an onboard monochrome AxioCam MRm camera and colored using Adobe Photoshop (San Jose, CA).

Masson’s trichrome staining of transplanted tissues

10-µm cryosections of transplanted tissue embedded in optimal cutting temperature compound (OCT) on glass slides were stained for 3 min in 50% hematoxylin solution (Protocol® Fisher Scientific, Kalamazoo, MI), rinsed in tap water, stained for 4 min in Biobech scarlet-acid fuchsin (Sigma-Aldrich, St. Louis, MO), rinsed in 0.5% acidified water (5ml glacial acetic acid per 1L H<sub>2</sub>O), immersed for 10 min in PMA-PTA solution (5ml Phosphomolybdic acid 10%/w/v, 5ml Phosphotungstic acid per 10ml H<sub>2</sub>O - reagents from Sigma-Aldrich, St. Louis, MO), rinsed in acidified water, stained for 4 min in 2% fast green solution (in 195ml H<sub>2</sub>O and 2ml glacial acetic acid, 4g fast green FCF from Sigma-Aldrich, St. Louis, MO), rinsed in acidified water, dehydrated in 3 changes of 100% ethanol and cleared in xylene. Prior to light-microscopy analysis, stained slides were covered with cover slips using a resinsious mounting medium.

In vitro differentiation of pCPCs with oxytocin

To induce cardiomyocyte differentiation, we used oxytocin treatment as previously described by Matsuura et al. [18]. In brief, pCPCs in the density of 2 x 10^4 cells per cm<sup>2</sup> were cultured on 1% gelatin-coated dishes with Dulbecco’s Modified Eagle Medium (DMEM) supplemented with 10% FBS, 100 U penicillin, and 1000 U streptomycin at 37°C in humidified air containing 5% CO<sub>2</sub>. Twenty-four hours after seeding, the cells were treated with 100 nM oxytocin (RayBiotech, Norcross, GA). After treatment, the medium was changed every three to four days. Total time in culture was 21 days. RT-PCR from RNA extractions and immunohistochemistry staining were used to determine the extent of cardiomyocytes differentiation. pCPCs cultured in the expansion medium without oxytocin served as control.

In vitro differentiation of pCPCs with vascular endothelial growth factor (VEGF) and fibroblast growth factor 4 / hepatocyte growth factor (HGF)

Three pCPCs lines were exposed to VEGF or FGF4 and HGF to determine their capacity to differentiate into endothelial and hepatocyte-like cells respectively as previously described [19,20]. pCPCs were seeded at a density of 2 x 10^4 cells per cm<sup>2</sup> in 24-well plates in expansion medium overnight. After 12 hours, pCPCs formed an adherent monolayer, expansion medium was removed, cell monolayers were washed with PBS and medium was replaced by serum free medium containing 60% low glucose DMEM (Hyclone, Logan, UT), 40% MCDB 201, 1X insulin transferrin selenium (ITS), 1X linoleic bovine serum albumin (LA-BSA), 10° M dexamethasone, and 10° M ascorbic acid 2-phosphate (all from Sigma, St. Louis, MO), 100 U penicillin, 1000 U streptomyacin (Hyclone, Logan, UT)) supplemented with 10 ng/mL VEGF (R&D) for endothelial differentiation or 10ng/mL FGF4 and 20ng/mL HGF (R&D) for hepatocyte differentiation. Cultures were incubated at 37°C, 5%CO<sub>2</sub> and 5%O<sub>2</sub> and respective media were changed every 4 days for 14 days.

results

pCPCs proliferated and expanded beyond 45 days in culture

Cardiomyocytes depleted cardiac cell suspensions in the expansion medium did not survive after 2 weeks in atmospheric oxygen conditions (20% O<sub>2</sub>). Under physiological hypoxia (5% O<sub>2</sub>), many cells died early in culture, however, after the first week of culture, spindle like cells that form colonies were observed. These cells were later termed putative cardiac progenitor cells (pCPCs). The colonies from the atrium proliferated at a rate of 13-15 cell divisions within 42 days and were viable beyond 42 days in culture. However, the cells from the ventricle proliferated slower and reached senescence early in culture (Figure 1). Therefore all subsequent studies described herein were done with pCPCs derived from atrium.

Cultured cells express Sca-1, smooth muscle actin and cardiac transcription factors

pCPCs grown in expansion medium stained positive for Sca-1.
and smooth muscle actin (SMA) at day 40 (supplementary figure 1). RT-PCR of cultured cell at the beginning (passage 0) slightly express Klf4, GATA-4 and GATA-6 while the cultured cells (after 42 days) had increased Klf4, GATA-4 and GATA-6 expression and also expressed Nanog, c-kit and Nkx2.5. Early and late cultures were always negative for the pluripotent transcription factor Octamer-4 (Oct4) and cardiac myosin heavy chain, Myh6 (Figure 2).

**pCPCs are Sca-1-** cells by flow cytometric analysis

Flow cytometric analysis of pCPCs after 6 weeks in expansion medium demonstrated majority of the cells with Sca-1, Nanog and Nkx2.5 expression, while c-Kit was only slightly expressed (supplementary figure 2). pCPCs were also positive for Nkx2.5 by staining (Figure 3A).

**Oxytocin-treated pCPCs expressed cardiomyocyte properties in vitro**

pCPCs supplemented with oxytocin in the expansion media showed induction of morphological changes including cellular arrangement and elongation. The control cells, pCPCs without oxytocin in the expansion medium, did not show these morphological changes (supplementary figure 3). Immunostaining confirmed the induction of cardiac sacromeric myosin (MF-20) in oxytocin-treated cells but not in the control cells (Figure 3B), although the staining pattern was perinuclear which suggest an immature phenotype [21]. We examined the gene expression of cardiac transcription factors and cardiac structural proteins by RT-PCR. Before treatment with oxytocin, only Nkx2.5 and GATA4 cardiac transcription factors were slightly expressed. Cardiac structural protein, troponin-I, was expressed after three weeks of oxytocin treatment (Figure 4). However, cardiac myosin heavy chains were not expressed (not shown) also consistent with an immature phenotype.

**Failure of pCPCs to differentiate into non-cardiomyocyte lineages**

pCPCs after 6 weeks in expansion medium were induced to differentiate into endothelial-like and hepatocyte-like cells. 14 days after exposure to differentiation media cells were analyzed by RT-PCR for the endothelial marker, von Willebrand Factor (vWF), and the hepatocyte marker, albumin, respectively. All three induced pCPC lines were negative for vWF and albumin (supplementary figure 4).

**pCPCs differentiated into cardiomyocyte-like cells when transplanted in skeletal muscle**

The atrium-derived pCPCs from GFP transgenic mice, which were transplanted into the skeletal muscle of wild type mice, formed new muscular bundles 14 days after transplant. These GFP positive muscular bundles were not positive for skeletal specific myosin (data not shown), however, they stained positive for cardiac Troponin-T and their sizes were smaller than the endogenous mature muscular bundles from the skeletal muscle (Figures 5A-B). Untransplanted skeletal muscles were negative for cardiac Troponin-T staining, indicating the cardiac specificity of this antibody. In contrast to the cytoplasmic GFP pattern in transplanted cells at injection tract (Figures 5A-B, arrow) which were not Troponin-T positive, the GFP pattern in Troponin-T positive bundles was cross-striated, which has been previously described in transgenic cardiomyocytes expressing GFP driven by cardiac-actin
promoter [22]. Some Troponin+/GFP+ muscular bundles were bi-nucleated which is commonly seen in cardiomyocytes (Figure 5B). In addition, these new muscular bundles were positive for smooth muscle actin, which was expressed by pCPCs in culture. Furthermore, transition of smooth muscle actin positive muscular bundles that co-expressed cardiac Troponin-T is seen in the transplanted cells (Figure 5C), indicating progression from early cardiogenic markers to more mature markers [23,24]. The cross-striation of the Troponin+/GFP+ muscular bundles differ from skeletal muscle fibers and unlike the skeletal muscle fibers which show a continuum syncytium of myonuclei along the muscle fibers, the Troponin+/GFP+ muscular bundles seem to have gap junctions connecting them (Figure 5A). Thus we stained for connexin 43, a gap junction protein expressed in cardiac intercalated disks, which was positive in the cell-cell membrane boundaries exclusively in the transplanted tissue that co-stained with GFP and Troponin-T (Figure 6).

Transplantation of pCPCs in heart resulted in fibrotic tissue at the injection site

Due to autofluorescence of the heart tissue specially in necrotic and infarcted areas [25], prior to transplantation we labeled the GFP+ pCPCs with PKH26, a red fluorescent dye that irreversibly binds the cell membrane and has successfully been used for in vivo transplantation of the cells [26]. Using this double tracking system, we found GFP+/PKH26+ pCPCs in the needle track surrounded by fibrotic tissue 14 days after transplantation into the heart (Figure 7). These GFP+PKH-26+ cells did not acquire the morphology of mature cardiomyocytes and did not stain positive for cardiac troponin (Figures 7 G-N, inset in M and N), indicating that in contrast to the skeletal muscle transplant, pCPCs transplanted in the heart did not differentiate into cardiomyocytes but the injection elicited a fibrotic response which was also observed in the mock injected hearts. It is likely this fibrotic response impaired the capacity of pCPCs to undergo cardiogenesis in the heart.

Discussion

We describe a new culture method for selection of the putative cardiac progenitor cells from the adult murine heart. This method is a modification of previous methods used to isolate and culture-expand stem cells from the bone marrow, including multipotent adult progenitor cells (MAPC), multipotent adult stem cells (MASC) and marrow-isolated adult multilineage inducible (MIAMI) cells [17,27-29]. Using the expansion medium containing 2% FCS, EGF, PDGF-BB, Beltrami et al. [29] successfully isolated human multipotent cells from the heart, which expressed undifferentiated cell marker Octamer-4 (Oct4) with the capacity to differentiate into 3 germ layers. Unlike Beltrami et al, we isolated cells which did not express Oct4 but were positive for other pluripotent stem cell genes such as Nanog and KLF4. Mononuclear cardiac cells did not express Nkx2.5 and c-kit immediately after isolation but rather up-regulated the expression of Nkx2.5 and c-kit over time in culture. This indicates first that we did not have contamination of cardiomyocytes or committed cardiac progenitor cells early in culture, and second, that up-regulation of...
Figure 6: pCPCs transplanted in skeletal muscle formed muscle bundles that expressed connexin 43, the gap junction protein expressed in cardiac intercalated disks. (A and B) Masson's trichrome staining of the transplanted TA muscle showed pCPC transplanted cells in the injection track (pointed by arrow) forming muscle like tissues (B). Notice that pCPCs did not form fibrotic tissues (no green or blue collagen tissue was seen in the injection site). (C-F) Transplanted TA muscle was stained with anti-GFP (Alexa 488) (D), connexin 43 (Alexa 594) (E) and DAPI (F, merged with DAPI). (C) Montage of the pCPCs transplanted tissue along the TA muscle identified as GFP+ and connexin 43 positive. (D-F) Higher magnification of transplanted tissue showed connexin 43 expression at cell-cell membrane boundaries. (G) Masson's trichrome staining of transplanted TA muscle of over 100 µm distant to (A) also showed transplanted pCPCs in the injection track. (I) An adjacent slide of area depicted by arrowhead was positive for cardiac Troponin-T Alexa 647. In this area transplanted pCPCs identified as GFP+ were also connexin 43 positive. Scale bars = 100 µm.

Figure 7: pCPCs transplanted in the heart did not express cardiac troponin and were found in fibrotic tissue. (A) Transplanted pCPCs identified as GFP+/PKH26+ cells (green and red fluorescent, respectively) were located in the left ventricular wall. The bright red florescence emitted from PKH26 greatly facilitated the screening of the heart tissue to locate the transplanted cells. (B) H&E staining showed fibrotic-like tissue in the transplanted area. (C) Higher magnification of area pointed by an arrow in (B). (D) Masson's trichrome staining showed collagen deposition (in blue) in the transplanted area. (E) Higher magnification of area pointed by an arrowhead in (D). (F) Higher magnification of area pointed by arrow in (D). (G-J) Transplanted pCPCs identified as GFP+/PKH26+ did not exhibit cardiomyocyte morphology and did not stain positive for cardiac Troponin T. (K-N) Another transplanted heart clearly showed GFP+/PKH26+ cells in the injection track were negative for cardiac Troponin T (M-N insets). Scale bars = 100 µm.
Nkx2.5 and c-kit in culture indicates either commitment of endogenous cardiac stem cells into cardiac progenitor cells or an in vitro artifact resulting in expression of these cardiac progenitor cell markers in a cell population that does not have endogenous cardiogenic function. The pCPCs isolated with our protocol did not differentiate into endothelial cells or hepatocytes using the same protocols previously reported for differentiation of multipotent adult progenitor cells (MAPC) from the bone marrow [19,20], suggesting that the pCPCs isolated with our method from the adult atrium are committed to the cardiac cell fate. This unipotentiality or restriction to the cardiac lineage was also observed when pCPCs were transplanted in the skeletal muscle resulting only in cardiac-like cells. Additionally, although the pCPCs expressed some of the pluripotency transcription factors, such as Nanog and KLF4, they did not express the Oct4 pluripotent marker which may explain their limited differentiation capacity.

Sca-1 positive pCPCs did not express c-kit at the beginning of culture indicating absence of c-kit positive cells immediately in culture. This result is the same as previously reported by Oh et al. [8]. Because c-kit+ cardiac progenitor cells decline rapidly after birth [30], we hypothesize that our culture conditions either select for a very small c-kit+ population that outgrows in culture, or that these culture conditions induce dedifferentiation. The latter is unlikely, since the pCPCs generated under these culture conditions only differentiated into cardiac-like cells and not into cells of any other lineage.

Most of the protocols reported to induce beating cardiomyocytes have used co-culture methods with mature cardiomyocytes [10] or induction with 5-aza-deoxycytidine (5-azadC), an inhibitor of DNA methylation previously reported to promote the differentiation of pluripotent P19 embryonal carcinoma cells [8,9]. We induced cardiomyocytes differentiation with oxytocin treatment in DMEM plus 10% FBS for 21 days, which showed expression of cardiac proteins by immunohistochemistry staining and cardiac-specific genes by RT-PCR. However, the phenotype of oxytocin-induced pCPCs was immature and no beating cells were observed in our cultures. In contrast to the studies reported by Matsuura et al. [18], which resulted in beating cardiomyocytes when induced with oxytocin, we cultured the cells for a shorter period of time and we did not replate the cells at confluence.

Nonetheless, pCPCs transplanted into the skeletal muscle formed muscle bundles that expressed mature cardiac proteins, suggesting these cells turned into mature cardiomyocytes. Furthermore, pCPCs that generated cardiac muscle bundles in the skeletal muscle stained positive for connexin 43 at cell-cell boundaries across the transplanted tissue, indicating that pCPCs differentiated into cardiac muscle bundles with gap junctions. In contrast, pCPCs transplanted into the heart in vivo did not differentiate into cardiomyocytes, but instead, resulted in fibrosis at the site of injection. This finding lends further evidence to the idea that the adult heart may not be a very permissive environment for cardiomyogenesis, and in turn, the heart’s response to injury is scar formation [16,31]. In light of the fact that the heart-derived pCPCs did differentiate into cardiomyocyte-like cells in skeletal muscle, further studies may be warranted investigating the endogenous factors in the skeletal muscle that permit cardiogenesis and may be missing in the heart environment. Identification of such cardiogenesis-permitting growth factors could open up a new venue for strategies to coax the injured heart into regeneration with exogenous growth factor therapy. Although, transplantation of the GFP+ cells into the wild type host cannot rule out cell fusion, the lack of expression of skeletal muscle specific myosin in the transplanted pCPCs excludes fusion in the skeletal muscle transplants.

One of the limitations of the present study is the absence of determination of the identity of the cell type capable of becoming pCPC in culture. Future studies should aim at prospectively isolating pCPCs using stem cell markers such as c-kit and Sca-1. Unfortunately, humans do not have the Sca-1 gene [32,33]. Although we have not yet determined if similar cardiac progenitor cells can be derived from the human heart, our method, which does not depend on any cell surface markers, may contribute to a protocol for human CPCs culture for future cell-based therapies.

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