Sympathetic Neurons Regulate Cardiomyocyte Maturation in Culture

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Embryos devoid of autonomic innervation suffer sudden cardiac death. However, whether autonomic neurons have a role in heart development is poorly understood. To investigate if sympathetic neurons impact cardiomyocyte maturation, we co-cultured phenotypically immature cardiomyocytes derived from human induced pluripotent stem cells with mouse sympathetic ganglion neurons. We found that 1) multiple cardiac structure and ion channel genes related to cardiomyocyte maturation were up-regulated when co-cultured with sympathetic neurons; 2) sarcomere organization and connexin-43 gap junctions increased; 3) calcium imaging showed greater transient amplitudes. However, sarcomere spacing, relaxation time, and level of sarcoplasmic reticulum calcium did not show matured phenotypes. We further found that addition of endothelial and epicardial support cells did not enhance maturation to a greater extent beyond sympathetic neurons, while administration of isoproterenol alone was insufficient to induce changes in gene expression. These results demonstrate that sympathetic neurons have a significant and complex role in regulating cardiomyocyte development.

Keywords: cardiomyocyte maturation, sympathetic innervation, iPS cells, heart development, co-culture

INTRODUCTION

The heart is densely innervated by sympathetic nerves, which extend from cell bodies housed in the sympathetic ganglia derived from trunk neural crest cells (Kimura et al., 2012). Sympathetic activity provides critical regulation of cardiac function, increasing beat rate, relaxation, contractile force, and conduction velocity (Hasan, 2013; Vegh et al., 2016). In cardiac pathologies, altered innervation density and sympathetic dysfunction are associated with heart failure, hypertension, arrhythmogenesis, and sudden cardiac death (Chen et al., 2007; Kimura et al., 2007; Florea and Cohn, 2014; Shen and Zipes, 2014; Fukuda et al., 2015). Furthermore, in pediatric patients, sympathetic defects have been linked to sudden infant death syndrome, certain congenital heart defects, cardiac arrhythmic death, and hypertension (Kahn et al., 2003; Crump et al., 2011; Ohuchi et al., 2011). These innervation disorders may have foundations in the interactions between cardiomyocytes (CMs) and sympathetic neurons (SNs) during development. Understanding the function of the cardiac sympathetic nervous system in...
the embryo and its role in heart development would give valuable insight into fetal, pediatric, and adult heart disease.

Neuronal projections first appear in the dorsal mesocardium at embryonic day (E) 10.5 in mice (Hildreth et al., 2009). At this point, embryonic CMs are already responsive to adrenergic stimulation (Vegh et al., 2016). Projections reach the aorta and pulmonary trunk at E11.5 and begin to differentiate into parasympathetic and sympathetic branches. By E13.5, sympathetic axons have started to innervate the ventricles, guided by nerve growth factor-expressing vascular smooth muscle of the coronary veins and later arteries (Nam et al., 2013). They continue to expand well into postnatal life, eventually rivaling the density of capillaries within the myocardium (Freeman et al., 2014; Zaglia and Mongillo, 2017). Disrupting cardiac sympathetic patterning in the embryo leads to postnatal arrhythmia and sudden death (Ieda et al., 2007), while abolishing sympathetic function results in prenatal death due to cardiac failure (Kobayashi et al., 1995; Thomas et al., 1995; Zhou et al., 1995; Rohrer et al., 1996; Baker et al., 2012). We have studied the autonomic devoid Phox2b−/− mouse and observed embryo mortality at E13.5-14.5 (Pattyn et al., 1999; Mokshagundam et al., 2021). We performed echocardiography to measure cardiac function prior to death, but found no progressive phenotype in Phox2b−/− embryos, suggesting that sudden death due to arrhythmia was the most likely cause. These studies demonstrate that the sympathetic system is essential for cardiac function and embryo survival. Despite their extensive innervation and essential role, the long-term effects of SNs on developing CMs are largely unknown, and it remains unclear if SNs influence aspects of CM maturation.

To investigate the role of SNs in CM maturation, we performed in vitro co-culture of human induced pluripotent stem cell (hiPSC)-derived CMs with embryonic mouse SNs. CMs derived from hiPSC display an immature, fetal-like phenotype and are a well-used model for studying CM maturation (Uosaki et al., 2015; Guo and Pu, 2020). We found that co-culture with SNs up-regulated multiple genes for CM maturation, ranging from myofiber components to ion channels and calcium machinery. Furthermore, gap junction expression increased, as well as sarcomere organization and intracellular Ca2+ transient amplitudes. Sarcomere spacing, however, was shorter and sarcoplasmic reticulum (SR) Ca2+ stores were reduced in co-cultured CM, demonstrating that SNs can also have a limiting effect on some aspects of maturation. We then incorporated endothelial cells (ECs) and epicardial-derived vascular smooth muscle and fibroblasts (EPI) into our co-culture model, as demonstrated in the recent studies that these cell types enhance maturation of hiPSC-derived CMs (Lee et al., 2015; Tan et al., 2021). We found that the maturation benefits of SNs were equal to or exceeded those of ECs and EPI, while combining all cell types did not produce further enrichment. Our results therefore indicate that SNs have a significant impact on developing CMs and play a role similar to other support cells in CM maturation.

MATERIALS AND METHODS

Animals

C57BL/6J mice were purchased from the Jackson Laboratory. All experiments were performed under approval from the National Heart, Lung, and Blood Institute (NHLBI) Animal Care and Use Committee.

hiPSC-CM Co-Culture

CMs were differentiated from ND2.0 hiPS cells expressing a GFP reporter as previously described (Chen et al., 2011; Luo et al., 2014; Lin et al., 2017). CMs were cryopreserved on day 10 and thawed 24 h before co-culture. SNs were obtained from dissected sympathetic ganglia of E13.5 mouse embryos as previously described (Nam et al., 2013). Sympathetic ganglia were dissociated to single cell suspension with the papain dissociation system according to the manufacturer’s directions (Worthington). Epicardial cells were harvested from E13.5 mouse embryo hearts as previously described (Nam et al., 2013). HUVECs were purchased commercially (Lonza). CMs and support cells were co-cultured at a ratio of 10:1. Culture surfaces were coated in 0.11 mg/ml growth factor reduced Matrigel (Corning). For FACS and qPCR, cells were cultured on 12 well plates ( Falcon); for immunohistochemistry, cells were cultured on plastic coverslips (Nunc Thermostan); for Ca2+ imaging, cells were cultured on glass coverslips (Celltreat). Co-cultures were maintained in DMEM/F12 media ( Gibco 11330) supplemented with 64 μg/ml L-ascorbic acid (Sigma),13.6 ng/ml selenium (Sigma), 10 μg/ml transferrin (Sigma), 5% chemically defined lipid concentrate ( Gibco 11905), 5% fetal bovine serum (HyClone), 1% penicillin-streptomycin ( Gibco 15140), 20 μg/ml insulin (Sigma I 9278), 20 ng/ml nerve growth factor ( NGF, Millipore 01-125), 10 ng/ml human basic fibroblast growth factor (FGF-2, PromoCell C-60240), and 0.5 ng/ml human vascular endothelial growth factor-165 (VEGF165, PromoCell C-64420). Culture medium was changed every 2–3 days. All analyses were conducted on day 30 of cell culture.

GCaMP6s Knock-in

GCaMP6s-expressing CMs were developed as follows. We used CRISPR/Cas9-mediated targeting to knock-in GCaMP6s at the AAVS1 safe harbor locus (Chen et al., 2013; Lin et al., 2018). The AAVS1 locus is an ideal target for the introduction of cell markers or other transgenes as it is actively and stably expressed in pluripotent stem cells as well as in differentiated cells (Patterson et al., 2020). Donor plasmid containing GCaMP6s and AAVS1 homology arms was transfected along with a AAVS1-CRISPR/Cas9 expression plasmid into 2 million ND2.0 iPSCs by nucleofection (Supplementary Figure S1A). Transfected iPSCs were seeded in E8 medium for 2 days, 0.5 μg/μl of puromycin (Sigma-Aldrich) was added on day 3 and continued for 10 days thereafter. The medium was changed every day. The survived colonies were picked and screened by PCR. Genomic DNA was collected from single colonies, and the gene targeting was confirmed by S'-junction PCR (Supplementary Figure S1B). Primers AAVS1U-F2 (CTG
CCGTCTCTCTCTCTGTAGT) and PuroU-R (GTGGGCTTTGTACCGGTCAT) detected targeted allele. Primers AAVSEL-F (TTCCGGTACCTCTCAGTCC) and AAVSEL-R (GGCTCCATACGTAGCAAAACC) detected wild-type allele. A GaMP6s heterozygous knock-in iPSC clone, which contains both knock-in allele and wild-type AAVS1 allele, was then differentiated into cardiomyocytes for evaluation of GFP fluorescent transients that reflects calcium flux during contraction (Supplementary Figure S1C). Normal karyotype of the gene-edited ND2.0-GaMP6s iPSC clone was confirmed by G-band karyotyping (Supplementary Figure S1D).

Flow Cytometry

Single-cell suspensions were prepared by treating cultured cells with 124.8 units/ml collagenase type 1 (Worthington LS004214), 147.4 units/ml collagenase type 2 (Worthington LS004202), 400 units/ml DNaseI (Sigma DN25), and 5 μM CaCl2 in DMEM/F-12 (Gibco 11330) for 30 min at 37°C with gentle pipetting. Following dissociation with collagenase, cells were treated with 0.05% Trypsin-EDTA (Gibco 25300) for 3 min at 37°C. Cells were then resuspended in Leibovitz’s L-15 medium containing 1 mg/ml bovine serum albumin (Sigma A9148), 10 mM HEPES (Gibco 20278), and 5% FBS. Cells were incubated for an additional 30 min to allow cells to recover from trypsination and were then centrifuged (1,000 g) and resuspended in 1% penicillin-streptomycin (Gibco 15140), 250 nM 15630), 1% NaH2PO4, pH adjusted to 7.4 containing (in mM) 145 NaCl, 5.4 KCl, 2 CaCl2, 0.5 MgCl2, 5.5 glucose, 5 HEPES, 0.3 NaH2PO4, pH adjusted to 7.4.

qPCR Analysis

Total RNA was extracted from the cultured cells as previously described (Li et al., 2013). First-strand cDNAs were synthesized with SuperScriptIII reverse transcriptase (Invitrogen) using random hexamer primers according to the manufacturer’s instruction. The same starting mass of RNA was used for all experiments. Quantitative mRNA expression analysis was performed with LightCycler 96 (Roche) using FastStart Universal SYBR Green master (Roche). A 10 μl reaction was performed, with cDNA diluted 1:4 and a final primer concentration of 250 nM. Reactions were run in triplicates and a Ct difference greater than 0.5 cycles was rejected. At least n = 2 biological samples were analyzed per gene for each experimental group. Data was analyzed by the 2−ΔΔCt method (Livak and Schmittgen, 2001). Glyceraldehyde 3-phosphate dehydrogenase (GAPDH) and β-actin (ACTB) were used as housekeeper genes, with the geometric mean of their Ct values acting as the first normalization step (Vandesompele et al., 2002). These housekeeping genes were selected following stability analysis (Vandesompele et al., 2002). Data was then normalized to the CM-only control cultures as the second step. A detailed description of the PCR primers is described in Supplementary Table S1.

Immunohistochemistry

Cell culture staining was performed as previously described (Nam et al., 2013). Cells were washed in cold PBS and fixed in 4% PFA/PBS for 15 min at room temperature. After washing in PBS, cells were incubated with primary antibodies overnight at 4°C in 0.1% Triton X-100/PBS with 10% heat inactivated goat serum. The following primary antibodies were used: bovine anti-cardiac troponin T antibody (mouse monoclonal, DSHB, 1:1,000) to detect CMs, rabbit anti-tyrosine hydroxylase antibody (Millipore, 1:200) to detect SNs, rabbit anti-Col1a1 antibody (Millipore, 1:500) to detect fibroblasts, Cy3-conjugated anti-α-SMA antibody (mouse monoclonal antibody, clone 1A4, Sigma, 1:1,000) to detect smooth muscle cells, rabbit anti-von Willebrand factor antibody (Sigma, 1:200) to detect endothelial cells, anti-α-actinin antibody (mouse monoclonal antibody, clone EA-53, Sigma, 1:500) to detect sarcomeres, and anti-Cx43 antibody (mouse monoclonal antibody, clone 2, BD Biosciences, 1:200) to detect gap junctions. For immunofluorescence detection, either Cy3-, Alexa-568-, or Alexa-647- conjugated secondary antibodies (Jackson ImmunoResearch or ThermoFisher Scientific, 1:250, 1 h at room temperature) were used. Coverslips were mounted with Prolong Gold Antifade Mounting solution (Thermo Fisher Scientific). All confocal microscopy was carried out on a Leica TCS SP5 confocal (Leica). To quantify sarcomere organization and spacing, sarcomeres were first segmented with ImageJ using a ridge detection algorithm (Steiger, 1998). The segmented images were then partitioned into 20 × 20 μm windows. For each window, the local sarcomere organization and spacing were calculated using automated analysis in Matlab (Suctiffle et al., 2018). Briefly, the co-occurrence matrix and Haralick correlation were computed for multiple angles, 0–180°, and offset distances, 0–38 μm, generating an m (number of angles) by n (number of offset distances) matrix of Haralick correlations (Haralick et al., 1973). The sarcomere score is the magnitude of the peak correlation, ranging between 0 and 2. The offset distance associated with the peak correlation is the sarcomere spacing. An idealized image of parallel stripes, all the same width, would give a maximum organization score of 2. After computing the organization and spacing for each window, the average values for the entire image was considered the overall sarcomere organization and spacing. Windows with no or insufficient sarcomeres were not included. Cx43 content was analyzed using custom Matlab scripts. Sample numbers are given in figure captions.

Live Imaging and Quantification Analysis of GaMP6s-hiPSC-CM

Intracellular Ca2+ transients were recorded from GaMP6s-expressing CM-only and CM + SN cultured on glass coverslips. Cells were incubated at 37°C for 1 hour in fresh medium and then coverslips were transferred to an imaging chamber perfused at 1 ml/min with 37°C Tyrode’s solution containing (in mM) 145 NaCl, 5.4 KCl, 2 CaCl2, 0.5 MgCl2, 5.5 glucose, 5 HEPES, 0.3 NaH2PO4, pH adjusted to 7.4 (Knollmann et al., 2006). Propranolol (1 μM) was applied during pre-incubation and imaging to isolate CM from adrenergic signaling. Propranolol was used during CM-only imaging as well in order to maintain experimental parameters. Ca2+ Imaging was performed with a Leica TCS SP5 confocal microscope (Leica). Time-lapse images were recorded at 22 frames/second with a 20X/0.7 NA objective. After several
transients were recorded, 10 mM caffeine was applied by injecting a bolus directly into the imaging chamber. Caffeine was washed out for 5 min between image acquisitions. Ca²⁺ images were analyzed using custom Matlab scripts (Mathworks, Natick, MA). For each time series, three ROIs (7.5 × 7.5 μm) were manually selected for analysis. We analyzed \( n = 45 \) ROIs from 6 CM-only coverslips and \( n = 54 \) ROIs from 7 CM + SN coverslips.

Statistics
Data are presented as mean ± standard deviation. Data were compared using unpaired t-tests, Welch’s t-test, and Welch’s ANOVA with Dunnett’s T3 as indicated in the text and figure legends. \( p < 0.05 \) was considered significant. All statistical analysis was performed in GraphPad Prism (GraphPad Software, San Diego, CA).

RESULTS

Cardiomyocytes, Sympathetic Neurons, and Other Support Cells Intermix in Co-Culture
To understand how SNs support developing CMs, we established a long-term culture system (30 days) of GFP-expressing hiPSC-derived CMs (Lin et al., 2017) alone and with freshly isolated SNs dissected from E13.5 mouse embryos (Figure 1A). Cells were co-cultured on thin layers of Matrigel and grew to become 2–3 cell layers thick. CMs formed large clusters, which were surrounded by support cells (Figure 1C). CMs expressed the GFP reporter and CM marker cardiac troponin T (cTnT, Figure 1B). After 30 days, we observed SNs with tyrosine hydroxylase immunostaining (Figure 1C). Multiple sympathetic axons and axon bundles extended toward and crossed into CM clusters, confirming that the two cell types were interwoven and not sequestered in our co-culture (Figure 1C, Supplementary Figure S2). We performed further experiments with EPI and ECs as additional support cells (Figure 1A). EPI are epicardial-derived sources of cardiac fibroblasts and coronary vascular smooth muscle cells. These support cells grew adjacent to the CM clusters, but we observed few non-CM within the clusters themselves (Figures 1D,E). EPI-derived cells expressed the fibroblast marker Col1a1 and vascular smooth muscle marker α-smooth muscle actin (α-SMA, Figure 1D). ECs expressed von Willebrand factor (Figure 1E). Throughout the 30 days co-cultures, CMs beat spontaneously.

Sympathetic Neurons Up-Regulate Genes for Cardiomyocyte Maturation
We next examined changes in gene expression in co-cultured CM relative to CM-only controls. We performed quantitative PCR

FIGURE 1 | Co-culture protocol. (A) Diagram of cell sources and co-culture protocol. SNs were dissociated from E13.5 mouse sympathetic ganglia. GFP expressing hiPSC-CM were cultured alone (CM-only) or with SN (CM + SN) for 30 days. In other experiments, we included EPI and EC (CM + SN + EPI + EC). EPI were harvested from E13.5 mouse hearts and formed vascular smooth muscle and fibroblasts. ECs were HUVECS. (B–E) Co-cultures immunostained at day 30. (B) GFP-expressing hiPSC-derived CMs (green) were positive for cardiac troponin-T (red). (C) SNs expressed tyrosine hydroxylase (blue). (D) EPI cells showed α-smooth muscle actin (vascular smooth muscle, red) and Col1a1 (fibroblasts, cyan). (E) ECs are marked with von Willebrand factor (white). Scale bars 250 μm.
(qPCR) experiments on GFP+ CMs isolated from the co-cultures by FACS (Figure 2A). Relative expression is presented as a heatmap of log fold changes (Figure 2B). Genes tested were grouped into three categories: myofiber proteins, electrophysiology, and calcium cycling. When co-cultured with SNs, CMs showed increased expression of genes encoding structural components including TNNC1, TNNT2, MYL2, and ACTN2. Expression of ion channel genes such as KCNH2, KCNQ1, KCNJ2, SCN5A, and the Na+-Ca2+ exchanger (SLC8A1) were also increased, as well as genes encoding the calcium cycling machinery: ATP2A2, RYR2, PLN, S100A1. The gene encoding gap junction Cx43 (GJA1), however was not changed by co-culture with SNs. These data suggest that SNs can promote expression of genes for myofibril formation, electrophysiology functions, and calcium handling, all major facets of CM maturation.

We added additional support cells EPI and ECs to CM co-culture in order to better mimic the physiological environment. While CM + SN + EPI + EC showed up-regulation of multiple genes compared to CM-only controls, these additional support cells did not enhance expression beyond the changes observed in CM + SN (Figure 2B). In the case of some genes, such as RYR2 and PLN, expression increased to greater levels when SNs were the only support cell. We then performed further co-cultures with a single support cell: CM + EPI and CM + EC. Co-culture with EPI as the single support cell increased gene expression as well, although the change for most genes was slightly less than the multi-support cell or CM + SN conditions. EC had little or no effect on CM gene expression. In our co-culture, therefore, additional support cells did not enhance the effects of SNs.

Our co-culture results indicated that SNs increased CM gene expression. We then examined whether SN-derived β-adrenergic
signaling was the underlying factor, rather than the actual SN cells. To address this, we cultured CMs under chronic exposure to 1 μM isoproterenol, a non-selective β-agonist. Isoproterenol was added at each media change for the entire 30-days culture. Gene expression analysis revealed that the β-agonist reduced expression of several genes and had little to no effect on others. These results suggest that a different mechanism is responsible for the expression changes in CM + SN co-culture and that β-adrenergic signaling has no, or possibly a detrimental, effect on CM maturation.
Sarcomere Maturation Is Slightly Delayed in Cardiomyocytes Co-Cultured With Sympathetic Neurons

We next examined structural changes in the co-cultured CMs. We first analyzed sarcomere morphology in CMs with α-actinin staining (Figures 3A vs. 3B vs. 3C). CMs in all culture conditions showed parallel Z-disks distributed along the length of the cell. Sarcomere organization, which represents the alignment and regularity of spacing, was significantly increased in CM + SN (0.32 ± 0.05) compared to both CM-only controls (0.24 ± 0.04, p < 0.001) and CM + SN + EPI + EC (0.25 ± 0.05, p < 0.01, Welch’s ANOVA with Dunnett’s T3, Figure 3D). However, sarcomere spacing was significantly reduced by co-culture with SNs (CM + SN: 1.9 ± 0.07 μm, CM-only: 2.1 ± 0.08 μm, p < 0.0001, Figure 3E). When adding EPI and EC, sarcomere spacing recovered somewhat (2.0 ± 0.08 μm, p < 0.01, Figure 3E), but was still less than CM-only levels (p < 0.01). These additional cells may inhibit or counteract the effects of the SNs. In the developing heart, then, SNs may have a complex, dual role in sarcomere maturation, improving the alignment of Z-disks, but slightly delaying their spacing.

Sympathetic Neurons Increase Cx43 in Cardiomyocytes

Gap junction formation is a critical step in CM maturation (Scuderi and Butcher, 2017). We performed the connexin 43 (Cx43) gap junction protein immunostaining and analyzed the amount of Cx43 relative to CM area (Figure 3I). Cx43 expression was significantly increased in CM + SN compared to CM-only controls (0.03 ± 0.03 CM + SN vs. 0.008 ± 0.006 CM-only, p < 0.05, Figure 3F vs. 3G,I). Cx43 levels did not increase further when EPI and EC were added along with SN (0.05 ± 0.04, p = 0.61, Figure 3F vs. 3H,I). Therefore, SNs may have an important role in promoting Cx43 formation in CMs. Interestingly, in the above-mentioned qPCR analysis, mRNA for the Cx43 gene (GJA1) did not show a large fold change relative to CM-only controls (Figure 2B). These results indicate that SNs might enhance translation or other post-transcriptional events, leading to greater levels of the Cx43 gap junction protein.

Ca2+ Handling Is Changed Significantly by Co-Culture With Sympathetic Neurons

To investigate functional changes in response to co-culture with SN, we analyzed intracellular Ca2+ transients during CM contraction (Figures 4A,B). GCaMP6s-expressing CMs and SNs were co-cultured on Matrigel-coated glass coverslips for 30 days and then imaged under perfusion with a warmed Tyrode’s buffer. Ca2+ imaging was conducted at the intrinsic, spontaneous beat rate of CMs (Figures 4C,D). We found that the amplitude of the Ca2+ transient was significantly greater in CMs co-cultured with SNs compared to CM-only controls (2.87 ± 1.52 CM + SN vs. 2.05 ± 1.38 CM-only, p < 0.01, unpaired t-test, Figure 4E). Time to 90% relaxation (RT90), however, was longer in CM + SN (2.20 ± 0.76 s vs. 1.54 ± 0.50 s, p < 0.0001, Figure 4F). The larger Ca2+ amplitude is indicative of a more mature phenotype, while longer RT90 suggests less mature Ca2+ clearance. Taken together, these data demonstrate that SNs have a significant, yet bi-directional, affect the development of CM Ca2+ transients.

We further examined development of the SR Ca2+ stores by administration of caffeine. In response to caffeine stimulus, both CM-only and CM + SN showed an increase in the Ca2+ peak followed by temporary cessation of Ca2+ flux (Figure 4G vs. 4I). The relative change in the caffeine peak compared to baseline was significantly reduced in CM + SN (0.50 ± 0.32 CM + SN vs. 1.32 ± 0.88 CM-only, p < 0.0001, Welch’s t-test, Figure 4G vs. 4H,I). This smaller caffeine response indicates that CMs co-cultured with SNs have significantly less mature SR Ca2+. The variability of the caffeine response, however, was significantly greater in CM-only controls (0.23 ± 0.09 CM + SN vs. 0.67 ± 0.40 CM-only, p < 0.05, unpaired t-test, Figure 4G vs. 4H,I). We measured this variability as the standard deviation across all Ca2+ ROIs in a single coverslip. These results show a significant role for SNs in development of the SR and its function in Ca2+ dynamics.

DISCUSSION

Multiple aspects of CM maturation were modified by co-culture with SNs, including gene expression, sarcomeres, gap junctions, and Ca2+ handling. SNs up-regulated genes important to the contractile apparatus, electrophysiology, and calcium machinery, improved sarcomere organization, generated greater Cx43 levels in cells, and increased Ca2+ transient amplitudes. These results suggest a positive effect of SNs on CM maturation. However, SNs were also responsible for shorter sarcomere lengths, longer Ca2+ relaxation, and decreased release of Ca2+ from the SR, indicating delayed maturation. This bimodal role may be important to regulate CM development in vivo. Incorporating EPI and ECs as additional support cells produced only modest changes, lending further support to a significant role for SNs in guiding CM maturation.

Relative gene expression by qPCR revealed that multiple genes related to CM maturation increased expression when co-cultured with SNs. Genes encoding both thin (TNNC1, TNNT2) and thick (MYL2) myofibril filaments were up-regulated in the presence of SNs. Additionally, ACTN2, which encodes the major Z-disc α-actinin protein, was increased. However, TNNJ3, the mature isoform of troponin-I (Bedada et al., 2014), and the MYH7/ MYH6 ratio (Guo and Pu, 2020) showed only small effects, suggesting that these isoform switches may not be influenced by SNs. We furthermore found up-regulation of membrane bound ion channels, which are required to propagate the CM action potential. Genes for the major voltage-dependent Na+ channel (SCN5A) and inward rectifying K+ channel (KCNJ2) were both increased in CM + SN co-cultures. During the action potential, membrane depolarization triggers Ca2+-induced Ca2+ release, which facilitates contraction, followed by removal of Ca2+ from the cytoplasm as the CM relaxes (Bers, 2008). The main components for Ca2+ cycling all showed increased expression in CMs co-cultured with SNs, including L-type Ca2+ channel...
(CACNA1C), ryanodine receptor (RYR2), Na+-Ca2+ exchanger (SLC8A1), sarco/endoplasmic reticulum Ca2+-ATPase (SERCA, ATP2A2), and phospholamban (PLN). Interestingly, some of the largest expression increases occurred in RYR2 and PLN, which may relate to the changes in Ca2+ dynamics in CM + SN discussed below. Taken together, these data show that SNs modify CM gene expression and promote a more mature profile.

We found that SNs increased sarcomere organization in CMs. Randomly oriented myofibrils with condensed Z-bodies is a hallmark of immature CMs (Snir et al., 2003; Kamakura et al., 2013; Lundy et al., 2013). In both CM-only and CM + SN, Z-disks are visible and mostly aligned to the CM long axis, similar to previous 30 days culture (Kamakura et al., 2013). In the presence of SNs, however, alignment has progressed further and

![FIGURE 4](image-url)
sarcomeres are more uniformly spaced. These features can enhance the contractile force of CMs. Despite this increased organization, CMs co-cultured with SNs exhibited slightly shorter sarcomere lengths, indicating that SNs may inhibit sarcomere elongation. As CMs mature, sarcomere length—the distance between Z-disks—increases, reaching 2.2 μm in adult human CMs (Yang et al., 2014). We suggest, therefore, that SNs enhance organization but delay elongation of sarcomeres. Mechanisms of sarcomere assembly and maturation are slowly being uncovered (Guo et al., 2021), and further research is required to understand how these processes are regulated and interpret the effects of this dual role on CM function.

Cx43 increased in hiPSC-derived CMs co-cultured with SNs. Interestingly, the gene, GJA1, did not show increased expression, suggesting that SNs may regulate translation or another post-transcription mechanism. Cx43 forms the gap junctions in CMs, allowing for propagation of the action potential. In adult CMs, Cx43 is localized to the intercalated disc between individual cells. We did not observe any localization pattern in control or CM + SN. The effect of SN, therefore, seems limited to expression of the protein, but not trafficking to cell termini. In human CMs, Cx43 localization is a largely postnatal process and may take up to 7 years to completely organize at the intercalated disk (Vreeker et al., 2014). SN-promoted Cx43 expression, therefore, may be important to support later development of electrical conduction. A recent study using 3D hiPSC-derived CM/EC/fibroblast microtissues proposed that greater cAMP in CMs leads to enhanced organization but delay elongation of sarcomeres. As CMs mature, sarcomere length increases in CM + SN co-cultured hiPSC-CMs. While SN + CM indeed have less SR Ca2+, this increase would likely be due to greater influx of Ca2+ through the L-type Ca2+ channel. An alternative scenario, though, is that CM + SN release a greater fraction of their SR Ca2+ during normal beating. More sensitive ryanodine channels is one possible mechanism. Upon introducing caffeine, there would be less reserve Ca2+, therefore resulting, in part, to the reduced caffeine amplitude. Additional assays testing the activation of ryanodine receptors and activity of the L-type Ca2+ channel could provide insight to this issue.

Our co-culture experiments showed that SNs significantly affect Ca2+ dynamics in hiPSC-derived CMs. We observed larger Ca2+ amplitudes in CM + SN vs. CM-only controls, but slower relaxation times. Previous studies have demonstrated that stem cell-derived CMs express Ca2+ handling proteins, although at a lower level than adult CMs (Satin et al., 2008). Our qPCR data showed increased expression of Ca2+ handling machinery such as CACNA1C, SLCA8A1, RYR2, ATP2A2, and PLN in SN co-cultured hiPSC-CMs. While up-regulation of these genes is consistent with greater Ca2+ amplitudes, the reason for slower relaxation is less clear. A possible explanation is that the less mature SR Ca2+ stores in SN co-cultured CM may make these CMs more reliant on membrane bound Ca2+ channels for Ca2+ removal, prolonging their RT90. It is important to note, however, that we analyzed CMs at intrinsic beat rates, and the slower beat rates of CM + SN may contribute to the greater Ca2+ amplitudes (Figure 4). Field stimulation at different pacing rates would offer further insight into the effects of SN on Ca2+ cycling. Stem cell-derived CMs have been shown to exhibit functional features such as Ca2+-induced Ca2+ release and SERCA activity (Satin et al., 2008; Itzhaki et al., 2011; Hwang et al., 2015). In particular, it has been demonstrated in vitro that the caffeine triggered Ca2+ release increases with culture time as the CMs develop SR Ca2+ stores (Satin et al., 2008). However, in our study, we found that the caffeine Ca2+ release was significantly reduced in CM + SN. This caffeine response suggests that CM + SN have smaller SR Ca2+ stores compared to CM-only, indicating that SNs lead to a less mature SR Ca2+ phenotype. It is intriguing, however, that at baseline, CM + SN show a larger Ca2+ amplitude. If the CM + SN indeed have less SR Ca2+, this increase would likely be due to greater influx of Ca2+ through the L-type Ca2+ channel. An alternative scenario, though, is that CM + SN release a greater fraction of their SR Ca2+ during normal beating. More sensitive ryanodine channels is one possible mechanism. Upon introducing caffeine, there would be less reserve Ca2+, therefore resulting, in part, to the reduced caffeine amplitude. Additional assays testing the activation of ryanodine receptors and activity of the L-type Ca2+ channel could provide insight to this issue.

Although the magnitude of the caffeine triggered Ca2+ release was reduced, CM + SN showed a more consistent response to the caffeine stimulus. In CM-only cultures, the caffeine response varied greatly, with some CMs exhibiting only a modest caffeine peak while others showed a more than 3-fold change compared to their baseline amplitude. CM + SN, however, had a much narrower band of responses. This difference may reflect a role for SNs in regulating development of Ca2+ handling apparatuses. Without SNs, maturation of Ca2+ handling may progress more haphazardly, generating a heterogenous mix of CMs. With SNs, Ca2+ handling development may be more tightly controlled. An intriguing question is if more uniform Ca2+ handling and, possibly, lower SR Ca2+ could be protective against arrhythmia early in development. Without uniform Ca2+ handling, a variegated response to environmental stimuli, for example, could lead to discordant ventricular contraction or rhythm defects. Additionally, delayed afterdepolarizations (DADs) occur when high luminal SR Ca2+ stimulates spontaneous Ca2+ release (Bers, 2008). These afterdepolarizations can trigger arrhythmias in some cases. With lower SR Ca2+, a DAD is less likely to occur. Later in development, these effects would need to dissipate, especially to grow the SR Ca2+. But there may be a window where SNs provide some protection to the developing heart. Further research can explore these potential roles of SNs.

The significance of non-CM cells in the heart has long been recognized and various cell types have been used to improve stem cell-derived CMs (Burrudge et al., 2014; Lee et al., 2015; Masumoto et al., 2016; Scuderii and Butcher, 2017; Zhang et al., 2017; Giacomelli et al., 2020; Tan et al., 2021). To generate a more physiologic environment, we added EPI and EC to perform CM + SN + EPI + EC co-culture. CM + SN + EPI + EC showed improvements in gene expression and Cx43 level over CM-only controls. Compared to CM + SN, however, the addition of EPI and ECs did not have a strong effect. Changes in gene expression were similar, and in some cases genes were more highly expressed in CM + SN, such as RYR2 and PLN. CM + EPI
also achieved similar changes in gene expression compared to CM + SN + EPI + EC. In the case of Cx43, there was no difference when EPI and EC were added. We did note that the EPI and ECs may have countered the effects of the SNs on sarcomere organization and spacing, recovering both to at or near the CM-only levels. Together, these data suggest that multiple support cells do not add up to generate further maturation. Our CM + SN + EPI + EC results demonstrate that SNs can influence CM maturation similar to other, more widely studied, support cells.

SNs have received relatively little attention as a potential regulator of CM maturation. Most studies thus far have focused on their role in modulating CM proliferation, which remains unclear (Tampakakis and Mahmoud, 2021). Neonatal rat SN and CM co-culture improved contractility and increased L-type Ca²⁺ channel expression, which is in line with our findings (Lloyd and Marvin, 1990; Ogawa et al., 1992). Co-culture of stem cell-derived CMs with SNs have demonstrated that SNs can control CM beat rate, but effects on CM maturation were not investigated (Takeuchi et al., 2012; Sakai et al., 2017; Takayama et al., 2020). In summary, our study demonstrates that SNs have a significant impact on multiple facets of CM maturation, giving insight into their critical role during development.

Further research is required to understand how SNs act to influence CM maturation. Our qPCR results suggest that β-adrenergic signaling is not sufficient to achieve the changes we observe in CM + SN co-culture. A limited number of previous studies on β-adrenergic stimulation and immature CMs have left the issue mostly unsettled. Neonatal rat CMs showed decreased cell size but no effect on proliferation in response to isoproterenol treatment (Kreipke and Birren, 2015). In E11.5 mouse embryonic CMs, β-adrenergic stimulation inhibited proliferation via decreased phosphorylation of Erk and Akt (Peridooni et al., 2017). Studies with hiPSC-CMs have shown conflicting results with regards to β-adrenergic stimulation: it either results in CM hypertrophy (Zhi et al., 2013) or has no effect on cell size (Foldes et al., 2014). In the case of α-adrenergic signaling, the agonist phenylephrine promoted hypertrophy in hESC-CMs, but had no impact on hiPSC-CMs (Foldes et al., 2011; Foldes et al., 2014). In the clinical setting, chronic exposure to catecholamines contributes to progressive heart failure and be cardiotoxic (Lymperopoulos et al., 2013). A previous study exposing iPSC-CM engineered tissues to norepinephrine for 7 days showed features of heart failure, but these were significantly blunted in serum-containing media (Tiburcy et al., 2017). Our qPCR results showed reduced expression of several genes after 30-days exposure to isoproterenol. It is possible that these changes are due to heart failure development in hiPSC-CMs. However, few studies have examined the effects of chronic catecholamine exposure on immature CMs, and it will require additional experiments to determine if there are cardiotoxic effects.

The physical presence of SNs in our CM + SN co-cultures may be the significant factor modulating CM maturation, rather than adrenergic signaling. Multiple sympathetic axons extended toward and invaded CM clusters, potentially enabling cell-cell contacts. A previous study with neonatal rat CM and SN showed changes in CM morphology, synapsin puncta, and cadherin accumulation at sites of cell-cell overlap (Shcherbakova et al., 2007). If similar specialized zones occur in our co-culture model, it may allow for greater cell-cell interaction between SN and CM. Further experiments such as RNA-sequencing will be needed to understand what signaling pathways are changed in CM + SN vs. CM-only conditions. While a future such experiment may reveal new interactions between CM and SN, there remains much to be learned about adrenergic signaling. Although prior work has not shown a definitive role, it is clear from Th⁻/⁻, Dbh⁻/⁻, and other mouse models that catecholamines are essential for embryonic heart function and embryo survival (Thomas et al., 1995; Zhou et al., 1995; Baker et al., 2012; Osuala et al., 2012). In the Phox2b⁻/⁻ model, embryos die mostly between E13.5-E14.5. At this stage, SNs have just reached the ventricle, innervating only a small portion of the dorsal subepicardium and are not yet in contact with CMs (Nam et al., 2013). Before SNs reach the heart, intrinsic cardiac adrenergic cells are required to sustain catecholamine levels (Ebert et al., 2008). In early embryonic CMs, therefore, adrenergic signaling is necessary, while the actual presence of SNs may be dispensable. Later CM development may require contact with SNs. SNs therefore likely act through a combination of adrenergic stimulation and local cell-cell signaling to influence CMs. As seen in the results of this study, SNs can have an enhancing or delaying effect on CM maturation.

In conclusion, we found that SNs had a significant influence on hiPSC-derived CM maturation. Gene expression, Cx43 quantity, sarcomere organization and some aspects of Ca²⁺ handling were improved in CM + SN co-culture, while maturation of sarcomere spacing and SR Ca²⁺ were delayed. Adding EPI and ECs as other support cells did not enhance maturation beyond CM + SN, highlighting the significance of SNs. We assessed that SNs act through more than catecholamines, as culture of CM with isoproterenol did not replicate the changes in gene expression. These data suggest a complex role for SN in CM development. Future research will examine the influence of SNs other facets of CM maturation. Understanding this relationship could provide insight into fetal and pediatric arrhythmias, in particular sudden cardiac death as well as other disorders.

**DATA AVAILABILITY STATEMENT**

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

**ETHICS STATEMENT**

The animal study was reviewed and approved by National Heart, Lung, and Blood Institute (NHLBI) Animal Care and Use Committee.
AUTHOR CONTRIBUTIONS

WK designed and performed experiments and wrote the manuscript. IG-P designed and performed experiments. WL assisted with some experiments. HU, ET, and CK provided experimental materials and procedures. JZ, YL, and KP provided GFP-expressing hiPSC-CMs and GCaMP6s-expressing hiPSC-CMs. Y-SM directed this study and wrote the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fcell.2022.850645/full#supplementary-material
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