Electrospun Carbon Nanotube-Based Scaffolds Exhibit High Conductivity and Cytocompatibility for Tissue Engineering Applications

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ABSTRACT: Carbon nanotubes (CNTs) are known for their excellent conductive properties. Here, we present two novel methods, “sandwich” (sCNT) and dual deposition (DD CNT), for incorporating CNTs into electrospun polycaprolactone (PCL) and gelatin scaffolds to increase their conductance. Based on CNT percentage, the DD CNT scaffolds contain significantly higher quantities of CNTs than the sCNT scaffolds. The inclusion of CNTs increased the electrical conductance of scaffolds from 0.0 ± 0.00 kS (non-CNT) to 0.54 ± 0.10 kS (sCNT) and 5.22 ± 0.49 kS (DD CNT) when measured parallel to CNT arrays and to 0.25 ± 0.003 kS (sCNT) and 2.85 ± 1.12 (DD CNT) when measured orthogonal to CNT arrays. The inclusion of CNTs increased fiber diameter and pore size, promoting cellular migration into the scaffolds. CNT inclusion also decreased the degradation rate and increased hydrophobicity of scaffolds. Additionally, CNT inclusion increased Young’s modulus and failure load of scaffolds, increasing their mechanical robustness. Murine fibroblasts were maintained on the scaffolds for 30 days, demonstrating high cytocompatibility. The increased conductivity and high cytocompatibility of the CNT-incorporated scaffolds make them appropriate candidates for future use in cardiac and neural tissue engineering.

1. INTRODUCTION

Heart disease was the leading cause of death in the U.S. in 2020, a statistic that has remained unchanged for the past 95 years. Cardiac tissue engineering offers promising treatments and investigative tools for these cardiac diseases and disorders, including cell-based pacemakers, cardiac patches, microfluidic “heart-on-a-chip” models, and regenerative therapies. Additionally, many are hopeful for the possibility of engineering cardiac constructs capable of being implanted in lieu of heart transplants, which are insufficient due to massive donor shortages and chronic immunogenic complications. Similar to cardiac tissue engineering, advances in neural tissue engineering show promise in spinal cord nerve regeneration, peripheral nerve regeneration, and microfluidic “brain-on-a-chip” models for high-throughput testing and disease modeling.

Because both cardiac and neural physiology depend on the electrical conduction of action potentials, conductive scaffolds are a common theme in tissue engineering research. In neural tissue engineering, chitosan–gelatin scaffolds doped with conductive hyaluronic acid-poly(3,4-ethylenedioxythiophene) (PEDOT) nanoparticles were shown to enhance neural stem cell proliferation and differentiation into neurons and astrocytes. Researchers have also explored polyaniline and polypyrrole to endow neural tissue engineering scaffolds with conductive properties. In cardiac tissue engineering, conductive scaffolds could be used to electrophysiologically mature human-induced pluripotent stem cell-derived cardiomyocytes (hiPSC-CMs). The immaturity of hiPSC-CMs compared to functional cardiomyocytes presents a roadblock to their use in cardiac tissue engineering due to the increased risk of introducing arrhythmias and the potential for poor electrical coupling with native tissues. To address this limitation, researchers have attempted to electrophysiologically mature hiPSC-CMs using long-term culture, electromechanical manipulations, electrical stimulation, protein regulation, and co-culture with sympathetic neurons. While successful, these approaches lack the microenvironmental control provided by a scaffold-based approach, particularly electrospinning, which allows selection of the material and
control over morphology. Additionally, these approaches would be impossible to utilize in vivo, unlike a scaffold, which could be implanted. Thus, there remains a knowledge gap regarding the ability of conductive scaffolds to mature hiPSC-CMs. Studies suggest that a similar roadblock exists for the use of iPSC-derived neurons for neural tissue engineering. Transplantation of iPSC-derived neurons into murine cortices failed to generate action potentials within 7 weeks. This indicates that iPSC-derived neurons similarly exhibit electrophysiological immaturity by default and could thus benefit from conductive scaffolds to mature them for neural tissue engineering applications.

We aim to endow scaffolds with conductivity by incorporating carbon nanotubes (CNTs) into electrospun polycaprolactone (PCL)—gelatin fibers. Electrospinning is ideal for the fabrication of scaffolds due to its production of fibers on the nano- to microscale, resulting in biomimicry of in vivo extracellular matrices (ECMs). Additionally, electrospinning offers a high degree of morphological tunability based on solution properties and electrospinning parameters. Finally, electrospun fibers are amenable to doping, coating, and post-processing, allowing for functionalization of scaffolds for specific tissue engineering needs. We selected PCL due to its high biocompatibility and excellent mechanical properties and gelatin due to its broad biocompatibility, promotion of cell adhesion, and cost-effectiveness. Gelatin lacks the mechanical strength and elasticity typically desired for tissue engineering scaffolds and degrades too rapidly in vivo for cardiac and neural tissue engineering. Conversely, PCL has excellent mechanical properties and takes longer to degrade, up to 2–3 years compared to gelatin, which can degrade within days. By using PCL–gelatin scaffolds, we tune the degradation profile and provide enhanced mechanical properties. However, PCL–gelatin scaffolds remain electrically nonconductive.

We thus endowed these PCL–gelatin scaffolds with CNTs to confer conductive properties, as CNTs are well known for their superior electrical conductivity. Due to this electrical conductivity, CNTs have been shown to promote differentiation and maturity of neural stem cells when formed into rope-like substrates and used to deliver electrical stimulation, enhance neurons’ electrical activity when used as direct substrates, and improve proliferation and neural differentiation of mesenchymal stem cells when loaded into electrospun scaffolds. Additionally, thin films of CNTs functionalized with hydroxyl acid have been shown to enhance both cardiac and neuronal differentiation of canine iPSC-CMs. Both mature cardiac and neuronal cells utilize electrophysiological signaling to propagate action potentials critical to their function. The ability of CNTs to enhance both cardiac and neuronal differentiation within canine iPSC-CMs suggests an underlying electrical mechanism. In addition to their conductive properties, the photoacoustic properties of CNTs have proven useful in neural tissue engineering. These photoacoustic properties have been shown to enable light-triggered depolarization of neurons without the need for genetic manipulation (such as the case in optogenetics), as well as to enhance neurite outgrowth.

CNTs enhance mechanical robustness in addition to conferring electrical conductivity. Experiments by Liu et al. revealed that CNTs increased Young’s modulus of electrospun poly(lactic-co-glycolic acid) scaffolds by 86% and tensile strength by 28% at a mere 0.5% concentration. Additionally, CNTs have been shown to improve cellular adhesion and confer antimicrobial properties. The cytocompatibility of CNTs with human cells is controversial due to seemingly conflicting data and differing metrics. In prior work, CNTs have been claimed to be cytocompatible based on the lack of cytotoxicity toward Schwann cells on CNT substrates.

Figure 1. Schematic portraying the fabrication of scaffolds. (A) Non-CNT scaffolds are electrospun from PCL and gelatin. (B) “Sandwich” CNT (sCNT) scaffolds are electrospun, CNT arrays are manually stretched over electrospun fibers, and another layer is electrospun. (C) Dual deposition CNT (DD CNT) scaffolds are fabricated by winding CNTs and then electrospinning fibers onto the same rotating collector. Cells are seeded onto all three scaffold types.
Others have argued that CNTs are not cytocompatible due to studies observing the loss of cell viability in immortalized human epidermal keratinocytes following exposure to unre-
refined CNTs. It has been more recently argued that the degree of CNT toxicity depends on many factors, including purity, dispersal, and fiber length. Recent research has shown that the structure, diameter, and length of CNTs affect their pulmonary toxicity and cytotoxicity. It has been proven that bent multiwalled CNTs (MWCNTs) exhibit less cytotoxicity than straight MWCNTs, and that cytotoxicity of submicron-
diameter carbon fibers increases with decreasing diameter and decreases with decreasing length.

In this work, we endeavored to incorporate CNTs into electrospun PCL–gelatin scaffolds to take advantage of their superior conductive and mechanical properties and to demonstrate their cytocompatibility, thus providing a basis for their use as conductive scaffolds in cardiac and neural tissue engineering applications. We incorporated CNTs into our electrospun PCL–gelatin scaffolds using two distinct methods: “sandwich” and dual deposition. Here, we compare the CNT content, morphology, mechanical properties, degradation profiles, hydrophobicity, electrical conductance, and cytocom-
patibility of the scaffolds.

2. MATERIALS AND METHODS

2.1. Scaffold Fabrication. Polycaprolactone (PCL) (M<sub>n</sub> = 90 g/mol, Sigma-Aldrich) was combined with type-A gelatin from porcine skin (gel strength ≈ 300 g Bloom, Sigma-Aldrich) in a 1:1 ratio and dissolved in 1,1,2,3,3,3-hexafluoro-1-propane (HFP) (Thermo Fisher) at 20% w/v concentration. During electrospinning, the solution was extruded at 4 mL/h and 14 kV was applied. The resultant electrospun PCL–gelatin fibers were collected onto copper shim with a 15 cm die–collector distance (Figure 1A).

Sprinnable, vertically aligned MWCNTs were grown in a tube furnace using a modified version of the chlorine-mediated chemical vapor deposition (CVD) route. The MWCNT arrays were grown on a quartz substrate at 760 °C with acetylene as the carbon precursor and FeCl<sub>2</sub> (anhydrous 99.5%, VWR) as the catalyst. At 760 °C, acetylene gas (99.5%, Machine and Welding Supply Company) was flowed at 600 sccm, chlorine gas (99.99%, Custom Gas Solutions) at 2 sccm, and carrier gas argon (99.999%, Machine and Welding Supply Company) at 398 sccm, while the system pressure was regulated at 5 Torr. The acetylene gas flow was stopped after 20 min from the start of the growth process, and the grown arrays were left in the argon and chlorine flow for 20 additional min. The system was then purged with argon during cooling. A detailed procedure of this CVD CNT growth method was previously published by the Bradford group. The resultant MWCNTs exhibited an average diameter of 39 ± 6 nm (measured by a field emission scanning electron microscope), an average length of 1 mm (measured by an optical microscope), an aspect ratio of ~25,650, and a purity of 99.67% (measured by TGA; only 0.33% iron oxide catalyst residue materials left after 900 °C air oxidation). Notably, 1 mm is longer than most MWCNTs.

CNT arrays were formed by drawing horizontally aligned CNT sheets from the vertically aligned MWCNT arrays, thus drawing a small bundle of nanotubes at the edge of a spinnable array to continuously transform vertically aligned MWCNTs into horizontally aligned MWCNT sheets. Continuous collection of the aligned MWCNT sheets around a rotating mandrel resulted in a flat sheet with the desired thickness.

CNTs were incorporated into electrospun PCL–gelatin scaffolds using two distinct methods. In the “sandwich” CNT (sCNT) method, PCL and gelatin fibers were electrospun onto static copper shim. CNT arrays were manually stretched over electrospun fibers and then another layer was electrospun on top (Figure 1B). In the dual deposition CNT (DD CNT) method, the same CNT arrays from the “sandwich” method were wound onto a copper shim affixed to a mandrel rotating at approximately 20 RPM. CNT arrays were wound around the entire circumference of the rotating mandrel 2.5 times. PCL and gelatin fibers were electrospun onto the same rotating mandrel, depositing on top of the CNT arrays (Figure 1C).

2.2. CNT Volume Percent Quantification. For sCNT and DD CNT scaffolds, 1 × 1 cm<sup>2</sup> samples (n = 10 per scaffold type) were cut. Scaffold samples were momentarily dipped in HFP (Thermo Fisher) and initial wet weights were acquired. Samples were then immerse in HFP for 1 h to dissolve the PCL and gelatin components of the scaffolds, leaving behind only the CNT component. Final wet weights were then measured. Assuming that the density of the PCL–gelatin electrospun fibers and the density of CNT arrays are constant, the mass percent of CNTs within scaffolds is equivalent to the volume percent of CNTs within scaffolds.

2.3. Electron SEM Assessment of Morphology. For each scaffold type, 0.75 × 0.75 cm<sup>2</sup> samples (n ≥ 5 per scaffold type) were cut. Samples were mounted with carbon adhesive onto SEM stubs and sputter-coated with 10 nm of gold and palladium and imaged with an electron microscope (Hitachi TM4000) at 10 kV. Three SEM images were taken of each sample, and 20 measurements of fiber diameter and pore size were taken for each image using ImageJ, an image analysis software freely available at imagej.nih.gov/ij. Mean fiber diameter and pore size were calculated, as well as fiber diameter and pore size distribution.

2.4. Quantification of Mechanical Properties Using the Modified Tensile Strip Test. For each scaffold type, 4 × 1 cm<sup>2</sup> samples (n = 5 for non-CNT and DD CNT, n = 4 for sCNT) were cut. Standard 3” × 5” index cards were cut into “C-cards” with a 2.54 cm gap for sample placement, into which samples were affixed between double-sided tape and lab tape (Supporting Figure 1A). The C-cards containing samples were placed into a tensile tester (MTS, Q-Test 5) using 2 N pinch clamp grips, and the C-card was partially cut to leave the samples between the grips (Supporting Figure 1B and 1C). Tensile strip tests were run using a 5 N load cell, 2.54 cm grip separation, 50% break sensitivity, and 0.01 mm/s strain rate. This procedure was adapted from the ASTM standard D5035, “Textile Strip Method.” From the resultant measurements of sample extension (mm) and applied load (kN), Young’s modulus for each sample was calculated.

2.5. Enzymatic Degradation by Weight. For each scaffold type, 0.5 × 0.5 cm<sup>2</sup> samples (n = 10 per scaffold type) were cut and placed into 12-well plates. The samples were immersed in a 1:1 solution of 0.25 mg/mL collagenase in 0.1 M Tris-HCl and 0.005 M CaCl<sub>2</sub> for the enzymatic degradation of gelatin and 0.0025 mg/mL Pseudomonas lipase (Type XIII, ≥15 units/mg solid, Millipore Sigma) in phosphate-buffered saline (PBS) for the enzymatic degradation of PCL. A 0.25 mg/mL collagenase concentration was used based on a modification of Alberti and Xu, who used a 1 mg/mL concentration of collagenase in 0.1 M Tris-HCl and 0.005

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M CaCl₂ to analyze degradation of tendon-derived collagen fibril-based tissue engineering scaffolds. The concentration was reduced from 1 to 0.25 mg/mL collagenase because the electrospun scaffolds' microfibers are significantly smaller than the tendon-derived collagen fibrils studied by Alberti and Xu. In 2018, Spearman et al. employed P. lipase to increase the degradation rate of electrospun PCL/PCL-polyglycolide nanocomposite fibers via cleavage of PCL’s ester bonds. Because the purpose of the present study was to analyze and not increase the degradation rate, the P. lipase concentration was reduced from 0.4 mg/mL used by Spearman et al. to 0.0025 mg/mL. The 12-well plates were kept in a heated chamber maintained at 37 °C for 4 weeks. Dry and wet weights were collected from six of the samples on days 0, 1, 7, 14, 21, and 28. Dry weights were acquired by weighing samples after they were desiccated overnight. Both dry and wet weights were collected to assess scaffolds under culture-like conditions (wet measurement) and to control for the effect of varying hydrophobicity on weight (dry measurement). Multiple SEM images were collected from one sample per each scaffold type on days 1, 7, 14, 21, and 28.

2.6. Contact Angle Measurement. For each scaffold type, 2.54 × 2.54 cm² samples (n = 3 per scaffold type) were cut. Contact angle with water droplets was measured using a goniometer (Dataphysics OCA System, FDS Corp.).

2.7. Electrical Analysis. For each scaffold type, 0.75 × 0.75 cm² samples (n = 10 per scaffold type) were cut. End-to-end resistance was quantified using a handheld multimeter (True RMS Multimeter, Keysight) with probes placed on the edges of the samples either parallel or orthogonal to CNT arrays. For all scaffold types, the probes were placed onto the sides of the scaffolds on which the cells were later seeded. Conductance was calculated as the inverse of resistance.

2.8. Cell Culture. NIH 3T3 murine fibroblasts were maintained at 37 °C and 5% relative humidity and cultured in high-glucose DMEM media (Corning) supplemented with 10% fetal bovine serum (Atlas Biologicals) and 1% Penicillin–Streptomycin (Fisher Scientific). Media changes occurred every other day.

NIH 3T3 cells were passaged and then seeded onto sterilized 1 × 1 cm² samples cut from each scaffold type at a density of 10⁴ cells/cm², with 3 × 10⁴ cells plated into a well without a scaffold as a control. Cellularized scaffolds were maintained in 1.5 mL each of DMEM media (Corning) supplemented with 10% fetal bovine serum (Atlas Biologicals) and 1% Penicillin–Streptomycin (Fisher Scientific).

2.9. Cytocompatibility Analyses. In preparation for the following cytocompatibility analyses, scaffolds were cut into 1 × 1 cm² samples and placed into 24-well plates. They were sterilized with 70% ethanol for 20 min, immersed in fresh PBS three times for 5 min each, and then stored in PBS at 4 °C overnight. The following day, a control well was coated with 0.1% gelatin in PBS. Cells were passaged and resuspended at 5 × 10⁵ cells/mL, 20 μL of the cell suspension was slowly added to each scaffold, and the samples were incubated at 37 °C and 5% CO₂ for 1 h to allow for cell adhesion, after which 1 mL of the appropriate media was added to each well.

2.9.1. Live/Dead Assays. For a preliminary cytocompatibility analysis of all scaffold types, LIVE/DEAD viability assays (Invitrogen) were performed using NIH 3T3 murine fibroblasts. LIVE/DEAD assays were performed with 3T3 cells on days 2, 7, 14, 21, and 27 after seeding (n ≥ 3 per scaffold type). Ten images were collected using a fluorescent microscope (EVOS FL Auto 2, Invitrogen), and cellularized scaffolds were returned to culture media. To determine viability, the quantity of live cells and dead cells for each image was determined using Celleste 5 software. Viability was then calculated as the fraction of live cells over total cells.

2.9.2. Proliferation Assays. Alamar Blue assesses cellular viability by quantifying fluorescence generated by resazurin as it reduces to resoruvin upon entering living cells due to their enzymatic activity. On days 1, 3, 5, and 7 post seeding, 150 μL of Alamar Blue reagent (Fisher Scientific) was added to each well of a plate containing 3T3 cells seeded onto 1 × 1 cm²
samples from each scaffold type (n = 3 per type) and one well with cells containing no scaffold as a control. The plate was incubated at 37 °C for 1 h. Following incubation, 100 μL of the media and reagent was added from each sample in triplicate to a 96-well plate. Fluorescence within the 96-well plate was quantified using a microplate reader (Synergy HT, BioTek) set to 540/25 λ excitation, 590/35 λ emission and maintained at 37 °C.

2.9.3. Immunocytochemistry. On day 14 post seeding, cellularized scaffold samples were fixed with 4% paraformaldehyde, permeabilized using 0.25% TritonX-100 in PBS and 0.1% Tween-20 in PBS, serum-blocked using 2% bovine serum albumin and 2% goat serum in 0.1% Tween-20 in PBS, stained with phalloidin, counterstained with Hoechst 33342, and imaged using a fluorescent microscope (EVOS FL Auto 2, Thermo Fisher).

To assess scaffold attachment and migration on CNT-based scaffolds (n = 3 per scaffold type), on day 7 post seeding, the cellularized scaffolds were fixed using 4% paraformaldehyde. Samples were paraaffin-sectioned and placed onto slides. Following sectioning, the slides were deparaffinized and permeabilized using a SafeClear II xylene substitute (Fisher Scientific), a progression of decreasing ethanol concentrations in deionized water, 0.1% TritonX-100 in PBS, and 0.1% Tween-20 in PBS. Samples were then serum-blocked using 2% bovine serum albumin and 2% goat serum in 0.1% Tween-20 in PBS, stained with phalloidin, counterstained with Hoechst 33342, and imaged using a fluorescent microscope (EVOS FL Auto 2, Invitrogen).

2.10. Statistical Analyses. Results are presented as mean ± standard error of the mean. Statistical significance was tested using analysis of variance (ANOVA). Probability values of p < 0.05 were considered statistically significant.

3. RESULTS

3.1. CNTs can be Incorporated into Electrospun Scaffolds Using the “Sandwich” or the Dual Deposition Method. CNTs were successfully incorporated (Figure 2). The resultant scaffolds are shown with schematics, SEM images, and day 2 LIVE/DEAD assays using NIH 3T3 murine fibroblasts (Figure 2). From the visual analysis of the SEM images, it is evident that the CNTs are more evenly distributed in the DD CNT scaffolds (Figure 2H, Supporting Figures 2–4, Table 1) than in the sCNT scaffolds (Figure 2E, Supporting Figures 2–4, Table 1).

Table 1. Mean Values ± Standard Error of the Mean for Volume Percent of CNTs Within Scaffolds; n = 10 per Scaffolding Type

| Scaffold Type | volume percent of CNTs (%) | non-CNT | sCNT | DD CNT |
|---------------|----------------------------|---------|------|--------|
|               | 0.00 ± 0.00                | 26.39 ± 2.05 | 51.75 ± 5.66 |

“Denotes p < 0.05 significant difference between non-CNT and sCNT. “Denotes p < 0.005 significant difference between non-CNT and DD CNT.

Figures 2–4, Table 1). It is also evident from the greater presence of CNTs in SEM images that the CNT concentration is higher for DD CNT (Figure 2H) scaffolds than for sCNT (Figure 2E) scaffolds. These conclusions are both also apparent from gross visual analysis of the scaffolds (Figure 2A,D,G).

On day 2 post seeding, LIVE/DEAD assays performed using NIH 3T3 murine fibroblasts showed good cellular proliferation on all three scaffold types. For representative images of each scaffold type, there were 100% live and 0% dead cells for non-CNT scaffolds (Figures 2C), 95.48% live and 4.52% dead for sCNT scaffolds (Figure 2F), and 98.55% live and 1.45% dead for DD CNT scaffolds (Figure 2I). For all scaffold types, dead cells were far outnumbered by live cells, indicating no significant cytotoxicity. Notably, almost all cells formed clusters on sCNT (Figure 2F) scaffolds, with some cells forming clusters on non-CNT (Figure 2C) scaffolds, and no cluster formation on DD CNT scaffolds (Figure 2I). This indicates that the sCNT scaffolds present a more heterogeneous microenvironment, wherein cells are more attracted to certain areas of the scaffold than others, migrating to and forming clusters within these areas.

3.2. Volume Percent of CNTs is Significantly Higher Within DD CNT Scaffolds than Within sCNT Scaffolds. As visible in high-magnification SEM images (Figure 3) and as quantified based on the volume percent of CNTs within scaffolds (Table 1), DD CNT scaffolds have the highest quantity of CNTs within them (51.75 ± 5.66%), followed by sCNT scaffolds (26.39 ± 2.05%). Non-CNT scaffolds, obviously, contain no CNTs (0.00 ± 0.00%). The differences in volume percent of CNTs are significant between all three groups.

3.3. Inclusion of CNTs Significantly Increases Fiber Diameter and Pore Size. The inclusion of CNTs via both sCNT and DD CNT methods significantly increased the mean fiber diameter of the resultant scaffolds from 1.05 ± 0.02 to 20.21 ± 0.92 μm (sCNT) and 39.53 ± 3.92 μm (DD CNT) (Table 1). The DD CNT method increased the fiber diameter significantly more than the “sandwich” method (Table 1). The inclusion of CNTs via both fabrication methods also significantly increased the mean pore size of the scaffolds (Tables 1 and 2).

3.4. Inclusion of CNTs Using “Sandwich” and Dual Deposition Methods Significantly Increased Both Mean Young’s Modulus and Mean Failure Load of Scaffolds. Stress-strain curves show that sCNT and DD CNT scaffolds are stronger and more ductile with higher failure loads and Young’s moduli, while non-CNT scaffolds are more brittle with lower failure loads and Young’s moduli (Figure 4). Inclusion of CNTs using the “sandwich” and dual deposition methods significantly increased their Young’s moduli from 6.47 ± 0.16 MPa (non-CNT) to 19.76 ± 1.95 MPa (sCNT) and 59.32 ± 6.40 MPa (DD CNT) (Figure 4B). This is equivalent to a 205.4 and 816.9% increase in Young’s moduli at an 0.25% CNT load.

This is consistent with our results, as we used a significantly higher ratio of CNTs to electrospun fibers and thus achieved significantly higher Young’s modulus. Additionally, inclusion of CNTs significantly increased the mean failure load of scaffolds from 0.13 ± 0.012 kN for non-CNT scaffolds to 0.33 ± 0.002 kN for sCNT scaffolds and 0.38 ± 0.016 kN for DD CNT scaffolds (Figure 4C).
Figure 3. High-magnification SEM images of non-CNT (A), sCNT (B), and DD CNT (C) scaffolds exhibiting higher CNT content in DD CNT scaffolds than sCNT scaffolds and no CNTs in non-CNT scaffolds. Scale bars = 10 μm.

Table 2. Mean Values ± Standard Error of the Mean for Fiber Diameter and Pore Size, With Summaries of Their Distributions; n = 10 Per Scaffold Type for Mean Fiber Diameters and Pore Sizes and n = 4 With 160 Measurements Each Per Scaffold Type for Distribution Analyses

|                          | non-CNT                      | sCNT                        | DD CNT                      |
|--------------------------|------------------------------|-----------------------------|-----------------------------|
| mean fiber diameter (μm) | 1.05 ± 0.02<sup>a</sup>      | 20.01 ± 0.92<sup>b,c</sup> | 39.53 ± 3.92<sup>b,c</sup> |
| fiber diameter distribution | unimodal, skewed right (Supporting Figure 3A) | unimodal, skewed right (Supporting Figure 4A) | unimodal, skewed right (Supporting Figure 5A) |
| mean pore size (μm<sup>2</sup>) | 9.70 ± 1.30<sup>c</sup> | 618.27 ± 70.77<sup>a,b</sup> | 9778.36 ± 1041.81<sup>b,c</sup> |
| pore size distribution | unimodal, skewed right (Supporting Figure 3B) | unimodal, skewed right (Supporting Figure 4B) | bimodal, skewed right (Supporting Figure 5B) |

<sup>a</sup>Denotes <i>p < 0.05</i> significant difference between non-CNT and sCNT. <sup>b</sup>Denotes <i>p < 0.05</i> significant difference between sCNT and DD CNT. <sup>c</sup>Denotes <i>p < 0.05</i> significant difference between non-CNT and DD CNT.

Figure 4. Mechanical characterization of scaffolds. (A) Representative stress−strain curves from one sample per scaffold type. (B) Mean Young’s modulus. n ≥ 4 for each scaffold type. (C) Mean failure load. n ≥ 4 for each scaffold type. *Denotes <i>p < 0.05</i> significant difference between groups. Black = non-CNT, light gray = sCNT, and dark gray = DD CNT.

3.5. Inclusion of CNTs Reduced the Enzymatic Degradation Rate As Quantified By Change in Wet and Dry Weights. Based on both mean dry (Figure 5A) and wet (Figure 5B) weights, non-CNT scaffolds exhibited the steepest degradation profile, followed by sCNT scaffolds, and then DD CNT scaffolds. The CNTs remained visible in SEM images up to day 7 in sCNT scaffolds (Figure 5C,G) and up to day 28 in DD CNT scaffolds (Figure 5C,O).

3.6. Inclusion of CNTs Significantly Increased Water Contact Angle of Scaffolds. The inclusion of CNTs by both “sandwich” and dual deposition methods significantly increased the mean water contact angle of scaffolds from 73.2 ± 0.9° for non-CNT scaffolds to 143.9 ± 1.2° for sCNT scaffolds and 103.9 ± 1.0° for DD CNT scaffolds. Hence, the non-CNT scaffolds were the most hydrophilic, followed by DD CNT scaffolds, and then sCNT scaffolds (Figure 6). Additionally, the sCNT scaffolds exhibited a significantly larger mean water contact angle than the DD CNT scaffolds.

3.7. Inclusion of CNTs Significantly Increased End-to-End Conductance of Scaffolds. CNT inclusion using the “sandwich” and dual deposition methods increased the end-to-end conductance of the scaffolds from 0.00 ± 0.00 kS (both parallel and orthogonal), with a greater increase in conductance for the DD CNT scaffolds (5.22 ± 0.49 kS parallel, 2.85 ± 1.12 kS orthogonal) than the sCNT scaffolds (0.54 ± 0.10 kS parallel, 0.25 ± 0.003 kS orthogonal) (Figure 7B). This increased conductance was significant for DD CNT scaffolds but not for sCNT scaffolds. The increased conductance in DD CNT scaffolds correlates to the excellent electrical conductivity of CNTs. The conductance increased more for DD CNT than for sCNT scaffolds because the dual deposition method yielded a higher volume percent of CNTs (Table 1) than the “sandwich” method.

The increased conductance in sCNT and DD CNT scaffolds exhibited anisotropy, with conductance becoming significantly higher when measured parallel to CNT arrays (0.54 kS sCNT, 5.22 kS DD CNT) compared to when measured orthogonal to CNT arrays (0.25 kS sCNT, 2.86 kS DD CNT) (Figure 7B). This anisotropy of conductance is directly correlated to the inherent geometric anisotropy of the embedded CNT networks, in which the fibers are aligned toward a particular axis.
3.8. All Scaffold Types Demonstrate Cytocompatibility With Murine Fibroblasts for Up To 1 Month Post Seeding. Based on fluorescent microscopy of the month-long progression of LIVE/DEAD assays using NIH 3T3 cells, all scaffold types demonstrated high cytocompatibility with minimal to no cytotoxicity (Figure 8). Cells attached and proliferated well to all scaffolds, both without CNTs (Figure 8A–E) and with CNTs incorporated via the “sandwich” (Figure 8F–J) and dual deposition (Figure 8K–O) methods. Visual observation revealed that the cells also successfully migrated through the layers of the scaffolds for all scaffold types, indicating a high degree of biointegration (Figure 8N) and suggesting that the scaffolds’ structural microenvironment is physiologically favorable toward intravasation of the murine fibroblasts. By day 28, based on quantities of cells observed in the images, the cells proliferated the most on the DD CNT scaffolds (Figure 8O), followed by the non-CNT scaffolds (Figure 8E), and then the sCNT scaffolds (Figure 8J).

Metabolic activity is a good indicator of cytocompatibility because it quantifies cellular proliferation. NIH 3T3 cells demonstrate excellent metabolic activity when seeded onto the scaffolds, with no significant difference in metabolic activity between cells seeded onto scaffolds vs without scaffolds nor the significant difference between cells seeded onto scaffolds with vs without CNTs (Figure 9). On days 1, 3, and 7 post seeding, there was no significant difference overall between Alamar Blue fluorescence (RFU) in all groups (non-CNT scaffolds, sCNT scaffolds, DD CNT scaffolds, and non-scaffold cell-only scaffolds exposed to collagenase and P. lipase. (A) Mean dry weight of scaffolds over 28 days. n = 6 per scaffold type. (B) Mean wet weights of scaffolds over 28 days. n = 6 per scaffold type. (C) SEM images of scaffolds over 28 days. CNTs indicated by arrows. n = 1 per scaffold type. Scale bars = 200 μm.

Figure 6. Water contact angle. Representative images of one sample per (A) non-CNT, (B) sCNT, and (C) DD CNT scaffold type. n = 3 per scaffold type.
control). On day 5, there was a significant difference between groups, with the DD CNT scaffolds exhibiting the highest cell viability (2909.11 RFU), followed by non-CNT scaffolds (2709.78 RFU), and the cell-only control (2571.00 RFU), with sCNT scaffolds (2146.78 RFU) exhibiting the lowest cell viability. This is consistent with the visual assessment of LIVE/DEAD assays (Figure 9), which show the lowest rate of cellular proliferation on sCNT scaffolds, compared to non-CNT and DD CNT scaffolds, on both day 2 and day 7.

On day 14 post seeding, phalloidin and Hoechst 33342 staining (Figure 8) revealed that NIH 3T3 cells continued to attach and proliferate on scaffolds both without CNTs (Figure 8A) and with CNTs (Figure 9B,C). Furthermore, the morphology of the cells remained intact and unchanged, as indicated by the staining of actin filaments green and nuclei blue (Figure 9C). It is evident from the images that the cells attached to the sCNT (Figure 9B) and DD CNT (Figure 9C) scaffolds because cells are only located where the scaffolds are present, forming an "edge" within the images.

Sagittal sectioning was performed on cellularized scaffolds, thus allowing imaging of the cellularized scaffolds in cross section. From the staining of these sections, we observed that NIH 3T3 cells migrated into the microstructure of sCNT and DD CNT scaffolds with full-thickness penetration. Fibroblasts (indicated by their nuclei, stained blue with Hoechst 33342) on both sectioned sCNT (Figure 10A–B) and DD CNT (Figure 10C–D) scaffolds on day 7 post seeding migrated throughout the entire thickness of the scaffolds (Figure 10). Cells migrated more thoroughly on the sCNT scaffolds than on the DD CNT scaffolds, with 66.8% of cells migrating to the inner 50% of sCNT scaffolds, in contrast to only 42.4% of cells migrating to the inner 50% of DD CNT scaffolds. However, this is likely affected by the varying scaffold thicknesses; the sCNT scaffolds (Figure 10B) are thinner than the DD CNT scaffolds (Figure 10D).

4. DISCUSSION

The volume percent of CNTs within DD CNT scaffolds is significantly higher than within sCNT scaffolds. The volume percent of CNTs within both sCNT and DD CNT scaffolds is, obviously, significantly higher than within non-CNT scaffolds. The inclusion of CNTs into the scaffolds significantly increased both their fiber diameter (Figure 1A, Supporting Figures S2–S4) and pore size (Figure 1B, Supporting Figures S2–S4). The significant increase in scaffolds’ mean fiber diameter when CNTs were incorporated is likely due to the

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Figure 7. End-to-end conductance of scaffolds. (A) Measurement of end-to-end resistance parallel and orthogonal to CNT arrays. (B) Mean end-to-end conductance for electrospun scaffolds. n = 10 per scaffold type. *Denotes p < 0.05 significant difference between scaffold types and † denotes p < 0.05 significant difference between measurement directions. Black = non-CNT, light gray = sCNT, and dark gray = DD CNT.

Figure 8. Cytocompatibility of scaffolds with NIH 3T3 cells. LIVE/DEAD assays of non-CNT, sCNT, and DD CNT scaffolds on day 2 (A, F, K), 7 (B, G, L), 14 (C, H, M), 21 (D, I, N), and 28 (E, J, O). Live stained green and dead stained red with cells-only live (P) and dead (Q) controls. Scale bars = 275 μm.
highly electrically conductive properties of CNTs. Electrospinning operates through the application of a high voltage to a polymer droplet as it is extruded through a syringe needle. When the electrostatic forces overcome the surface tension forces of the droplet, a charged jet erupts. The jet deposits onto a negatively charged or grounded collector as the solvent simultaneously evaporates, generating a dried, fibrous mat. When the electrostatic forces overcome the surface tension forces of the droplet, a charged jet erupts. The jet deposits onto a negatively charged or grounded collector as the solvent simultaneously evaporates, generating a dried, fibrous mat. Therefore, the deposition of electrospun fibers depends on the electrostatic attraction between the charged jet and the collection plate. In the “sandwich” and dual deposition fabrication methods, CNTs were present in the scaffolds upon which fibers were electrospun. Thus, we hypothesize that the high electrical conductance of the scaffolds increased electrostatic attraction between the charged jet and the surface of collection, resulting in increased fiber diameter. The charged surface area of the fibers wound around the mandrel contributed to this effect.

We hypothesize that the significant increase in mean pore size when CNTs were incorporated into the scaffolds is also due to electrostatic processes during electrospinning. The increase in pore size was due to increased electrostatic repulsion between formerly and newly deposited electrospun fibers. Therefore, the deposition of electrospun fibers depends on the electrostatic attraction between the charged jet and the collection plate. In the “sandwich” and dual deposition fabrication methods, CNTs were present in the scaffolds upon which fibers were electrospun. Thus, we hypothesize that the high electrical conductance of the scaffolds increased electrostatic attraction between the charged jet and the surface of collection, resulting in increased fiber diameter. The charged surface area of the fibers wound around the mandrel contributed to this effect.

Figure 9. (A) Metabolic activity over 1 week as quantified by Alamar blue assays and (B) phallolidin-stained F-actin filaments green and Hoechst 33342-stained nuclei blue on (A) non-CNT, (B) sCNT, and (C) DD CNT scaffolds. Scale bars = 100 μm. *Denotes p < 0.05 significant difference between timepoints. Black = non-CNT, light gray = sCNT, and dark gray = DD CNT.

Figure 10. NIH 3T3 migration into scaffolds. Hoechst 33342-stained nuclei of 3T3 cells on day 7 post seeding on cross sectioned sCNT (A, B) and DD CNT (C, D) scaffolds, merged with trans-fluorescent microscope images of scaffolds (B, D). Dotted lines separate the outer 25% of scaffolds from the inner 50%. Scale bars = 275 μm.
fibers containing CNTs. Additionally, the increased pore size could be due to the less efficient packing of fibers because of their increased volume and decreased quantity.

Inclusion of CNTs also increased the mechanical robustness of the electrospun scaffolds, as shown by their significant increase in Young’s modulus (Figure 4D) and failure load (Figure 4D, Table 3). We previously found that decellularized ECM from cardiac tissues of porcine hearts exhibits average Young’s moduli of 5.36 ± 0.14 kPa from the left ventricle and 16.69 ± 0.32 kPa from the sinoatrial node.60 Jacot et al. discovered that Young’s modulus of healthy adult cardiac tissues remains in the range of 10–15 kPa.61 While the former data lacks cellular influence on elastic properties, focusing only on the contribution of the ECM, and the latter reference refers to nonhuman, nonadult cardiac tissues, it is likely that Young’s modulus of healthy adult cardiac tissues remains in the range of 5–20 kPa due to the extreme physiological similarity between porcine and human hearts. Our scaffolds, with Young’s moduli of 8.33 MPa (non-CNT), 19.76 ± 1.95 MPa (sCNT), and 74.67 MPa (DD CNT) greatly exceed this 10–15 kPa range by several orders of magnitude. Upon initial observation, this could appear problematic; however, high Young’s moduli are attractive for their durability, and many materials commonly implanted into cardiovascular tissues have Young’s moduli far exceeding that of native cardiac tissues and are mechanically tolerated without issue. For example, vascular grafts comprising commercial expanded polytetrafluoroethylene (ePTFE) exhibit Young’s modulus of 17.4 MPa.62 Commercial sutures commonly used in cardiac bioprostheses exhibit even higher Young’s moduli, ranging from 2,315.30 MPa in Gore-Tex to 13.06 GPa in Vicryl. A review by Li et al. reported that Young’s moduli of polymers used in implantable prosthetic heart valves are mostly in the range of 10–100 MPa, with one outlier (ePTFE) exhibiting a modulus of 413 MPa.63 Because these vascular grafts, sutures, and synthetic materials are commonly used and FDA-approved and have Young’s moduli similar to or exceeding those of our scaffolds, it is unlikely that the discrepancy in Young’s moduli, though large, would have any negative effect within our intended uses both as in vitro platforms for the electrophysiological maturation of hiPSC-CMs or as in vivo engineered cardiac tissue scaffolds for implantation.

Increased Young’s moduli could also appear problematic for neural tissue engineering, particularly considering that softer substrates with lower Young’s moduli are typically preferred in these applications.64 However, previous research has shown that CNT-based electrospun scaffolds not only show no cytotoxicity toward rat mesenchymal stem cells but actively enhance their proliferation and neural differentiation. These electrospun thermoplastic urethane scaffolds loaded with MWCNTs at 1.5, 2.5, and 3.5% concentrations exhibited Young’s moduli of 3.94, 10.01, and 9.80 MPa, respectively, lower but in the same order of magnitude as our CNT-based scaffolds’ Young’s moduli.44 Even neuronal stem cells cultured directly onto CNTs exhibited neurite elongation and physiological maturation when electrically stimulated.65 These results indicate that the robust mechanical properties of CNTs do not impede cell attachment, proliferation, and maturation in neural tissue engineering constructs. Hence, it is unlikely that the increase in Young’s modulus due to CNT incorporation will preclude our scaffolds from being used in neural tissue engineering applications.

Inclusion of CNTs by the “sandwich” and dual deposition methods significantly increased the failure load of the scaffolds. The respective mean failure loads of 0.13 ± 0.012, 0.33 ± 0.0002, and 0.38 ± 0.016 kN for the non-CNT, sCNT, and DD CNT scaffolds (Figure 4C) become 0.325 ± 0.03, 0.825 ± 0.0005, and 0.95 ± 0.04 N/mm², respectively, when the scaffold area is accounted for, which far exceed the 2–4 mN/mm² contractile force exerted by healthy, adult CMs in vitro,66 indicating that there should be no issue with material failure and the scaffolds would tolerate contraction of attaching CMs in vivo. The scaffolds are predicted to also tolerate iPSC-CMs with similar (although immature) contractile physiology seeded onto them in vitro for electrophysiological maturation. As the neural tissue is noncontractile, withstand contractile force is a nonissue for most neural tissue engineering applications.

Both mean dry (Figure 5A) and wet (Figure 5B) weights indicate that non-CNT scaffolds degrade the fastest, followed by sCNT, and then DD CNT scaffolds (Table 3). Thus, we conclude that the inclusion of CNTs slows the overall degradation profile of the electrospun PCL–gelatin scaffolds. This is confirmed by SEM images showing CNT arrays remaining intact as PCL and gelatin fibers degrade (Figure 5C). The CNTs remained present longer in DD CNT than sCNT scaffolds due to the higher volume percent of CNTs (Table 1). This difference in the degradation profile between the PCL–gelatin fibers and the CNTs is attractive for in vivo applications because it offers the possibility of CNTs remaining to facilitate electrophysiological conduction while PCL–gelatin fibers degrade and are replaced by de novo native tissues.

### Table 3. Mean Values ± Standard Error of the Mean For Morphological, Electrical, and Mechanical Data for Each Scaffold Type

|                          | non-CNT          | sCNT             | DD CNT            |
|--------------------------|------------------|------------------|-------------------|
| volume percent of CNTs (%) | 0.00 ± 0.00⁶⁻⁶²  | 26.39 ± 2.05⁶⁻⁶² | 51.75 ± 5.66⁶⁻⁶²  |
| mean fiber diameter (µm)  | 1.05 ± 0.02⁶⁻⁶²  | 20.01 ± 0.92⁶⁻⁶² | 39.53 ± 3.93⁶⁻⁶²  |
| mean pore size (µm²)      | 9.70 ± 1.30⁶⁻⁶²  | 618.27 ± 70.77⁶⁻⁶² | 9778.36 ± 1041.83⁶⁻⁶² |
| parallel conductance (kS) | 0.00 ± 0.00⁶⁻⁶²  | 0.54 ± 0.10⁶⁻⁶²  | 5.22 ± 0.49⁶⁻⁶²  |
| orthogonal conductance (kS)| 0.00 ± 0.00⁶⁻⁶²  | 0.25 ± 0.003⁶⁻⁶² | 2.85 ± 1.12⁶⁻⁶²  |
| Young’s modulus (MPa)     | 6.47 ± 0.16⁶⁻⁶²  | 19.76 ± 1.95⁶⁻⁶² | 59.32 ± 6.40⁶⁻⁶²  |
| failure load (kN)         | 0.13 ± 0.012⁶⁻⁶² | 0.33 ± 0.0002⁶⁻⁶² | 0.36 ± 0.016⁶⁻⁶²  |
| degradation rate (1- fastest) | 1                | 2                | 3                |
| water contact angle (°)   | 73.2 ± 0.9⁶⁻⁶²   | 143.9 ± 1.2⁶⁻⁶²  | 103.9 ± 1.0⁶⁻⁶²  |

*Denotes p < 0.05 significant difference between non-CNT and sCNT. ⁵Denotes p < 0.05 significant difference between sCNT and DD CNT. ⁶Denotes p < 0.05 significant difference between non-CNT and DD CNT.
degradation profile of the fibers can be tailored by modifying the ratio of PCL to gelatin to match the rate of de novo tissue formation for the tissue type of interest.

Inclusion of CNTs into electrospun PCL–gelatin scaffolds significantly increased their water contact angle (Figure 6, Table 3). This indicates that the CNTs are hydrophobic and endow the scaffolds with surface hydrophobicity. The effect is greater for sCNT scaffolds than for DD CNT scaffolds, resulting in significantly higher mean water contact angles. We hypothesize that this is because of the difference in thickness of the outermost layer of electrospun fibers between sCNT and DD CNT scaffolds. Although the volume percent of CNTs is higher in DD CNT scaffolds than in sCNT scaffolds, the electrospun fibers within the sCNT scaffolds are divided into two sections within the “sandwich” structure: above the CNT array and below the CNT array. Thus, though the overall volume percent of electrospun fibers is lower for DD CNT scaffolds, the outermost layer of electrospun fibers that the water droplet interacted with was thicker in the DD CNT scaffolds due to the previously discussed electrostatic effect of the CNTs. This thicker layer of electrospun PCL–gelatin fibers provided a more hydrophilic surface to attenuate the hydrophobicity of the CNTs. Thus, DD CNT scaffolds are more suitable as substrates for mammalian cells, which typically prefer hydrophilic surfaces. This factor likely contributed to the higher proliferation rate on DD CNT than on sCNT scaffolds observed in LIVE/DEAD assays (Figure 8) and higher metabolic activity (Figure 9).

Most notably, CNT incorporation into scaffolds by both the “sandwich” and dual deposition methods significantly increased their end-to-end conductance (Figure 7, Table 3). The CNT-attributed increase in conductance measured greater parallel to the arrayed CNT fibers compared to measurements made in the orthogonal direction. This was observed due to the anisotropy of the conductive network formed by the CNTs. Electrical stimulation has been shown effective in promoting iPSC-CM differentiation and synchronizing the spontaneous beating exhibited by iPSC-CMs, and electrical stimulation of CNTs specifically has been shown to provide cardiomyogenic cues to mesenchymal stem cells, directing their differentiation toward CM lineages. Furthermore, electrical stimulation has been shown to facilitate in vitro maturation of iPSC-CMs, resulting in increased action potential conduction velocity and calcium ion flux shifting toward physiological values for one study and shifting calcium transients toward physiological values while enhancing membrane N-cadherin signaling, stress-fiber formation, and sarcomeric length shortening when combined with mechanical stimulation in another. The increased electrical conductance of the scaffolds presented in this work potentially offers an attractive substrate for in vitro electrophysiological maturation of hiPSC-CMs. Based on our electrical conductance data and results observed in previously mentioned studies, we hypothesize that under the application of electrical stimulation, hiPSC-CMs would exhibit enhanced gap-junctional coupling facilitated by the high electrical conductance of our CNT-based scaffolds, enhancing propagation of action potentials between neighboring hiPSC-CMs and enabling the development of physiologically realistic calcium transients.

Electrical stimulation is also often used in neural tissue engineering. As previously mentioned, researchers cultured neural stem cells onto a rope comprised of single-walled CNTs. The constructs were electrically stimulated 2 days post-seeding. It was concluded based on ENO2 and MECP2 expression that this electrical stimulation promoted the early differentiation of neural stem cells into neurons and the maturation of the neurons post differentiation. A conductive scaffold composed of composite polypyrrole and silk fibroin was found to enhance viability, proliferation, migration, and expression of neurotrophic factors within Schwann cells when electrically stimulated. Due to their high conductance, our CNT-based scaffolds are similarly amenable to electrical stimulation for neural tissue engineering applications.

As previously mentioned, debate exists surrounding whether CNTs are cytotocompatible and thus suitable as biomaterials for use in implants or culture systems. Our data indicates that the inclusion of CNTs into our scaffolds results in high cytotocompatibility, resulting in negligible cytotoxicity toward NIH 3T3 murine fibroblasts for up to 1 month (Figure 8A) based on fluorescent microscopy of LIVE/DEAD assays and a week-long Alamar blue study done on 3T3 cells (Figure 9). Of the cellular proliferation assays performed on days 1, 3, 5, and 7, the only significant difference in cellular metabolic activity was observed on day 5 (Figure 9). The difference was likely significant on day 5 because this was the timepoint with the highest overall cellular proliferation before contact inhibition caused a decline in metabolic activity. On day 5, DD CNT scaffolds exhibited the highest proliferation, followed by non-CNT scaffolds, the cell-only control, and finally the sCNT scaffolds. We hypothesize that the cells preferred the homogenous microstructure of the DD CNT and non-CNT scaffolds to the heterogeneous microstructure of the sCNT scaffolds. The sCNT scaffolds exhibit more heterogeneity in structure than the DD CNT and non-CNT scaffolds because the layers of CNT arrays and electrospun PCL–gelatin fibers are arranged in a “sandwich” geometry (Figure 2).

Based on both the viability assays (Figure 8) and proliferation assays (Figure 9), cells appeared to prefer DD CNT scaffolds to sCNT scaffolds. In addition to the previously discussed heterogeneity of sCNT scaffolds, scaffold morphology also contributed to this difference. DD CNT scaffolds exhibit significantly greater fiber diameter and pore size than sCNT scaffolds (Table 2), resulting in increased surface area, which allows more attachment and proliferation sites for cells. Fibroblasts maintained their cellular morphology when seeded onto CNT-based scaffolds, visualized with phalloidin and Hoechst 33342 staining performed on day 14 post seeding (Figure 9C). Finally, fibroblasts migrated into both sCNT and DD CNT scaffolds, traversing their thickness by day 7 post seeding, as shown by phalloidin staining on sectioned cellularized scaffolds (Figure 10). This indicates that the CNT-based scaffolds provide an attractive and biomimetic microenvironment for cellular attachment, proliferation, and migration, the hallmark of a robust biomaterial for tissue engineering applications. Furthermore, these scaffolds can be used as conductive substrates for the application of electrical stimulation to cells, which has been explored as a method for in vitro maturation of hiPSC-CMs and neural stem cells.

5. CONCLUSIONS
We here present two types of electrospun, CNT-based scaffolds, which are highly electrically conductive and cytotocompatible. The use of electrospinning as a fabrication method allows for easy tunability of the resulting scaffolds, whose morphology can be easily controlled by modifying solution properties and electrospinning parameters. Both the
“sandwich” and dual deposition methods of CNT incorporation are amenable to modifications in the electrospinning portion of the fabrication process. Due to their increased electrical conductance and high cytocompatibility, these scaffolds hold potential for the development of novel cardiac and neural tissue-engineered constructs. Neural applications include spinal cord and peripheral nerve regeneration, neuronal growth substrates, and microfluidic models of the brain. Cardiac applications include cellular pacemakers, three-dimensional (3D) printed cardiac tissues, and cardiac patches for tissue repair following myocardial infarction. Additionally, these scaffolds show promise for incorporation into in vitro platforms to electrophysiologically mature hiPSC-CMs, making them relevant for use in a myriad of cardiac tissue engineering applications.

ASSOCIATED CONTENT

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/10.1021/acsomega.2c01807.

Supporting information uploaded separately contains the experimental setup for tensile testing and mean distributions of fiber diameter and pore size for each scaffold type (non-CNT, sCNT, and DD CNT) (PDF).

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AUTHOR CONTRIBUTIONS

T.C.S.: Conceptualization, methodology, investigation, validation, analysis, data curation, and writing original draft; J.T.: Investigation, methodology, data curation, visualization, writing review, and editing; N.M.: Investigation; K.M.A.: Investigation; M.L.: Investigation, methodology, and writing original Draft; P.D.B.: Supervision, resources; M.A.D.: Supervision, resources, writing review and editing, and funding acquisition; J.M.G.: Supervision, writing review and editing, funding acquisition, and resources

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NOTES

The authors declare no competing financial interest.

ABBREVIATIONS

CM, cardiomyocyte; CNT, carbon nanotube; CVD, chemical vapor deposition; DD CNT, dual deposition carbon nanotube; ECM, extracellular matrix; ePTFE, expanded polytetrafluoroethylene; hiPSC, human-induced pluripotent stem cell; hiPSC-CM, human-induced pluripotent stem cell-derived cardiomyocytes; MWCNT, multiwalled carbon nanotube; PCL, polycaprolactone; PEDOT, poly(3,4-ethylenedioxythiophene); sCNT, “sandwich” carbon nanotube

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