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Electrospun acellular scaffolds for mimicking the natural anisotropy of the extracellular matrix†

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In tissue engineering, the use of scaffolds helps establish a synergistic relationship between the scaffolds and the tissues by improving cell–scaffold interaction. This interaction is enhanced when physiologically relevant biophysical cues are replicated in the artificial scaffolds. Here, we present a novel scaffold that mimics the natural anisotropy of the native extracellular matrix of tissues, fabricated by electrospinning a combination of three polymers: polycaprolactone (PCL), polyvinylidene fluoride (PVDF) and polyaniline (PANI). The scaffolds were characterized for their morphology, surface and mechanical properties. Rat cardiomyoblast (H9c2) cells, cultured on the PCL–PANI–PVDF scaffold, demonstrated cell alignment, penetration and proliferation across the entire surface area of the scaffold without any external chemical or physical stimuli. The PCL–PANI–PVDF scaffold, unlike other scaffolds, does not require post-processing or specific temperature conditions of storage, prior to use. These acellular scaffolds fabricated through polymer blending, open new avenues for research on functional acellular scaffolds for tissue engineering, based on synthetic materials.

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Introduction

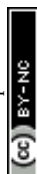
Scaffold-based tissue engineering is an important strategy in current regenerative research and reconstructive surgery.¹ Artificial matrices and scaffolds can emulate the natural microenvironment of cells and influence cell migration, differentiation, proliferation and survival for *ex vivo* cellular studies.^{2–4} However, in order to ensure appropriate cellular differentiation and proliferation, external stimulation or addition of growth factors is often required.⁵ For example, for connective tissues like the bone, the cells on the scaffolds are subjected to external stimuli such as constant compressive stress or axial stretching, to recreate the natural physical microenvironment.^{6,7} However, these external stimuli are ineffective in *in vivo* systems which must be minimalistic in design and reduce dependence on external factors. Modifying the material properties and surface morphology of a scaffold is one way of making the scaffold design simple, efficient and functional.^{8,9} In most synthetic scaffold-based processes, additional post-processing steps are required to make the scaffolds biocompatible and mechanically stable.¹⁰ These additional steps are time consuming and may further alter the physical structure and morphology of the scaffolds compromising their mechanical integrity.¹¹ Such properties are of vital importance since the cell–scaffold

interaction is dependent on several factors including the surface properties, morphology of the scaffold and its mechanical properties.¹² Functional scaffolds made from polyblend nanofibers fabricated by electrospinning are one of the strategies to address this challenge.¹³ The polyblend nanofibers are a combination of one or more polymers in a ratio that allows them to be electrospun into scaffolds using coaxial electrospinning.¹⁴ This technique allows the scaffold to take advantage of some of the native characteristics of the individual polymers, while gaining some new functionality.¹⁵

To achieve optimal cell–scaffold interaction, the surface and mechanical properties need to closely match with native tissues.¹⁶ The work presented here reports the fabrication of a 3D aligned nanofibrous scaffold, which closely replicates the mechanical anisotropy of the tissue and promotes cell proliferation without any external stimulation. Polycaprolactone (PCL), polyvinylidene fluoride (PVDF) and polyaniline (PANI) were used for the fabrication of the scaffold by coaxial electrospinning. The materials used were biocompatible, non-immunogenic and durable. PCL, a viscoelastic, aliphatic, biodegradable polyester is soluble in a number of organic solvents and compatible with many polymers which makes it an ideal candidate for blending with other polymers.¹⁷ PVDF is a piezoelectric and a mechanically strong polymer and PANI is a conductive polymer in the acidic state.^{18–20} A polymer scaffold composed of the three polymers provides the required flexibility in designing acellular scaffolds which mimic the surface and morphological properties of the natural extracellular matrix. Here, we developed a method to electrospin three distinct

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polymers by sequentially dissolving them in solvents with different boiling points and then coaxially electrospinning them to obtain a scaffold. The scaffold had an overall homogeneity composed of highly repetitive anisotropies analogous to native tissue. This unique morphology promotes cell adhesion, proliferation and migration in 3D without need for any growth factors, external stimuli and post processing.

Materials and methods

Fabrication of scaffolds

All the chemicals were used as procured unless specified. Polycaprolactone ($M_w = 70\,000$ GPC; Scientific Polymer Products, USA), PVDF ($M_w = 534\,000$; Sigma Aldrich; France) and PANI (emeraldine base; avg $M_w \sim 50\,000$; Aldrich Chemistry, USA) were used for synthesis of the scaffolds including PCL and PCL–PVDF. The doped form of PANI–PCL solution was prepared by dissolving equal mass of emeraldine base and 10-camphorsulfonic acid (HCSA) (Aldrich Chemistry; France) in a solution of 10% PCL in chloroform. The PCL–PANI solution was filtered to remove suspensions larger than $0.22\ \mu\text{m}$ since the doped form of PANI is immiscible.²¹ DMF (Macron Fine Chemicals; USA) and acetone (Sigma Aldrich, USA) in the ratio 7 : 3 were used to prepare a 20% PVDF solution. PCL–PANI–PVDF scaffold was prepared by coaxially electrospinning from separate syringes which were controlled using syringe pumps (Harvard Pumps, EP-H11). The PCL–PANI (core solution) and 20% PVDF (sheath solution) were electrospun (EM-DIG; IME Technologies, Netherlands) at 18 kV using a rotating collector at a rotational speed of 750 rpm. The parameters used for fabrication of PCL and PCL–PVDF scaffolds are elaborated in the ESI.[†]

Characterization of scaffolds

The fiber samples were sputter coated with Au/Pd (5 nm thick) and visualized using a Field Emission Scanning Electron Microscope (Hitachi S-4700 FE-SEM) and a high-resolution transmission electron microscope (JEOL JEM-2010 TEM; HRTEM). The alignment of the fibers and average diameter were calculated from the images obtained using the directionality tool in the ImageJ software. Attenuated total reflectance (ATR) Fourier transform infrared (FTIR) spectroscopy (PerkinElmer FTIR-ATR 100 series) was collected in the range of $400\text{--}4000\ \text{cm}^{-1}$ to study the surface chemistry. X-Ray Diffraction (XRD) was used to analyse the scaffolds using XDS 2000 Diffractometer (Scintag XDS-2000 Powder) and a Cu-K α ($\lambda = 1.54\ \text{\AA}$) source of X-ray radiation. The X-ray spectra were analysed using the DMSNT software and the inbuilt Pearson VII profile fitting function. The mechanical properties of the scaffolds were analysed ($n = 3$) using a dynamic mechanical analyser (TA Instruments Q800 DMA).

Cell culture

Rat cardiomyoblast cells were purchased from ATCC (H9c2; CRL-1446TM), and cultured in Dulbecco's Modified Eagle's Medium, DMEM (Life Technologies, USA) supplemented with

10% fetal bovine serum, FBS (Life Technologies, USA) along with 1% penicillin–streptomycin (Life Technologies, USA). The scaffolds were cut into samples of approximately $1\ \text{cm}^2$ area, sterilized in 70% ethanol and dried in a sterile biosafety cabinet prior to cell seeding. Cells in exponential growth phase were trypsinized, counted using a haemocytometer, seeded (2×10^4 cells per cm^2) and incubated under standard cell culture conditions ($37\ ^\circ\text{C}$, 65% relative humidity, 5% CO_2).

Cell characterization

The cells were fixed using 4% paraformaldehyde (Electron Microscopy Sciences), permeabilized using 0.1% Triton-X 100, and immunostained with DAPI (Life Technologies, USA) and Alexa Fluor® 488 Phalloidin (Life Technologies, USA) for A-T regions of the nucleus and cytoskeleton (F-Actin) visualization respectively, according to the manufacturer's protocols. Cells were visualized *via* fluorescence microscopy (EVOS FL Auto) and electron microscopy after 24 h, 48 h, 72 h and 168 h of growth. The quantitative analysis of the cell orientation was carried out using ImageJ software (NIH). The cell orientation was calculated by randomly selecting image slices and analysing the deviation in orientation with respect to the orientation of the majority of the cells using the directionality plugin on ImageJ. The averages and standard deviations were obtained out of five replicates for each scaffold type. For FE-SEM the cells cultured on the scaffolds were sputter coated with a 5 nm thick Au–Pd layer, after drying with Hexamethyldisilazane (HMDS).

Results and discussion

Coaxial electrospinning helps in formation of functional scaffolds from otherwise immiscible polymer blends. In the case of PCL–PANI–PVDF, a heterogenous functional scaffold composed of PCL, PVDF and PANI was formed by preparing PCL–PANI core solution and coaxially electrospinning it with PVDF sheath solution. The differential rate of solubilities of the polymers in solvents and their polarity helped in obtaining fibers with an aligned morphology. The PCL–PANI–PVDF fibers when observed by the naked eye have a repetitive pattern composed of thick diameter fibers aligned along the rotation of the cylinder during electrospinning (Fig. 1A).

Visualizing the fibers under the scanning electron microscope revealed a well-aligned and dense formation of a 3D fibrillar network of PCL–PANI–PVDF (low magnification: Fig. 1B and high magnification: Fig. 1C) with periodic structural anisotropies. The morphological characterization of the PCL and PCL–PVDF samples were done using FESEM (Fig. S1[†]). The PCL and PCL–PVDF fibers had an aligned morphology and had no beading in the fibers. The diameter of the fibers in the PCL–PANI–PVDF scaffold was measured to be between 30 nm to 2000 nm with a deviation of 8.68° in alignment. The fibers overlapped forming multiple layers leading to a very dense three-dimensional structure for the scaffold. The morphology was obtained by controlling the jet elongation and polymer flow rate which in turn was influenced by the sol–gel and the electrospinning parameters (electric field and environmental



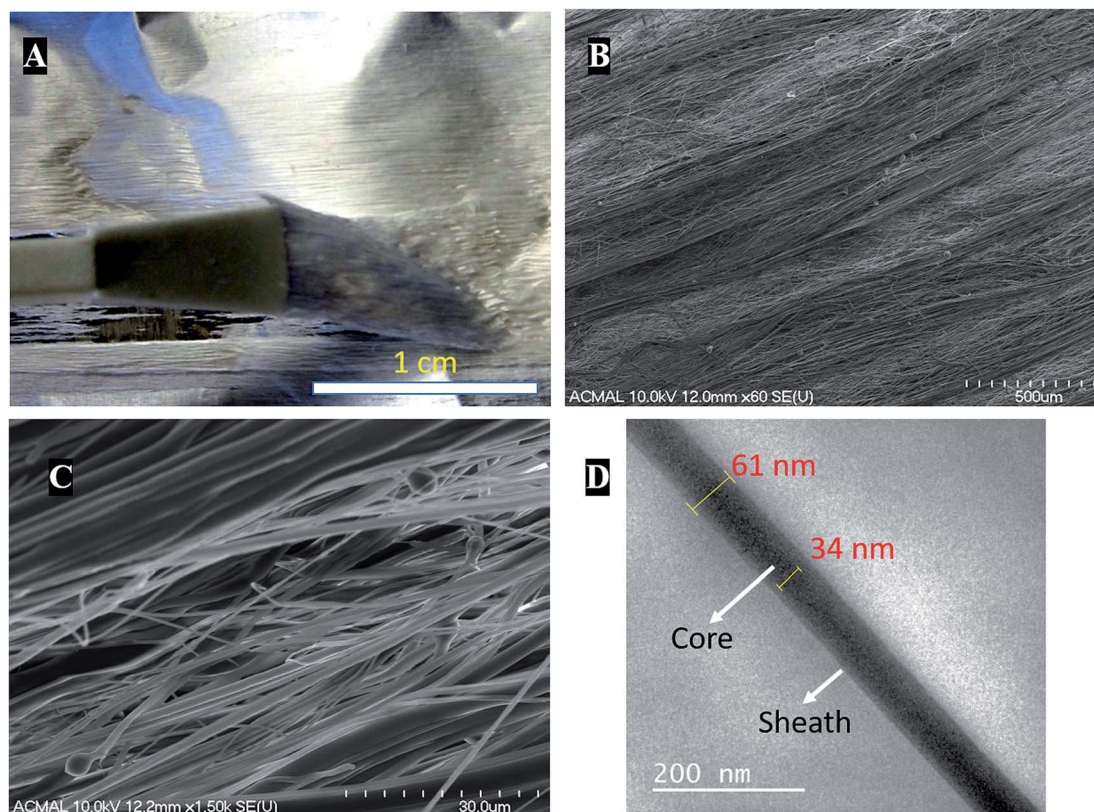
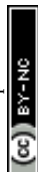


Fig. 1 (A) Digital photograph of the aligned morphology of the PCL–PANI–PVDF scaffold as visible to the naked eye. (B) Field emission scanning electron microscopic (FESEM) images of the PCL–PANI–PVDF scaffolds exhibiting aligned morphology and overlapping fibers. Large and small diameter fibers are interspersed having the same directionality. (C) A higher magnification image reveals the interlocked pattern of large and small diameter nanofibers and their close assembly. (D) High resolution Transmission Electron Microscopy (HR-TEM) image of PCL–PANI–PVDF after staining with osmium(VIII) oxide.

factors).²² The solvents used for preparing the solutions have different physical and chemical properties, which alters the surface tension, subsequently affecting the jet elongation.²³ Bands of larger diameter fibers were interspersed by smaller diameter fibers forming a pattern. Large diameter fibers have a lower critical uptake velocity than small diameter nanofibers.²⁴ The uptake velocity was controlled to introduce differential alignment with the larger diameter fibers having greater alignment compared to the smaller diameter fibers, producing a repetitive pattern. Transmission electron microscopy images of the fibers revealed a coaxial fiber composed of a thick core (PCL–PANI) and a relatively thin sheath (PVDF) (Fig. 1D). The thickness of the fiber and core was measured to be 61 ± 7 nm and 35 ± 3 nm, respectively.

Infrared spectroscopy was used for analyzing the surface functional groups of the PCL–PANI–PVDF scaffold as seen in Fig. 2A, PCL–PVDF and PCL scaffold as seen in Fig. S2.† The spectrum from PCL–PANI–PVDF showed distinctive peaks pertaining to PCL at 1722 cm^{-1} (C=O stretching), 1167 cm^{-1} (C–O stretching), 2944 cm^{-1} (asymmetric CH_2 stretching), 2866 cm^{-1} (symmetric CH_2 stretching) and a strong intensity peak at 1293 cm^{-1} (C–O and C–C stretching in the crystalline phase of PCL). Although, the peaks of PVDF overlapped with PCL and PANI, there was a low intensity peak at 840 cm^{-1} corresponding

to the CH_2 rocking in PVDF. The N–H stretching at 3442 cm^{-1} was specific to PANI. The PCL–PANI–PVDF scaffold has signature peaks corresponding to the three polymers. The XRD spectra from PCL–PANI–PVDF showed major peaks at 21.6° and 24.0° (Fig. 2B). The degree of crystallinity was calculated to be 0.653 confirming that the fibers were molecularly oriented. The significant peaks corresponding to PCL and PVDF are present. The PCL–PANI–PVDF scaffold has a high draw ratio of the polymer chains while electrospinning, thereby aligning the fibers along the axis of the rotating collector. This arrangement also provides more room for alignment of the microfibrils. From the FESEM images, it can be seen that there was branching in the fibers of larger diameters which confirmed the formation of microfibrillar regions. The presence of these microfibrillation increases the crystallinity of PCL–PANI–PVDF fibers, which agrees well with the presence of peaks at 1293 cm^{-1} corresponding to the crystalline phase. The surface characterization of PCL and PCL–PVDF was done using XRD (Fig. S3†). The mechanical properties of the PCL–PANI–PVDF scaffold were characterized from the stress–strain–stiffness graph (Fig. 2C and D). The microfibrillation and alignment of the fibers helps provide a higher stiffness in an uniaxial direction.²⁵ The stiffness is a very important parameter that affects the cellular responses through mechanoreceptors. The PCL–



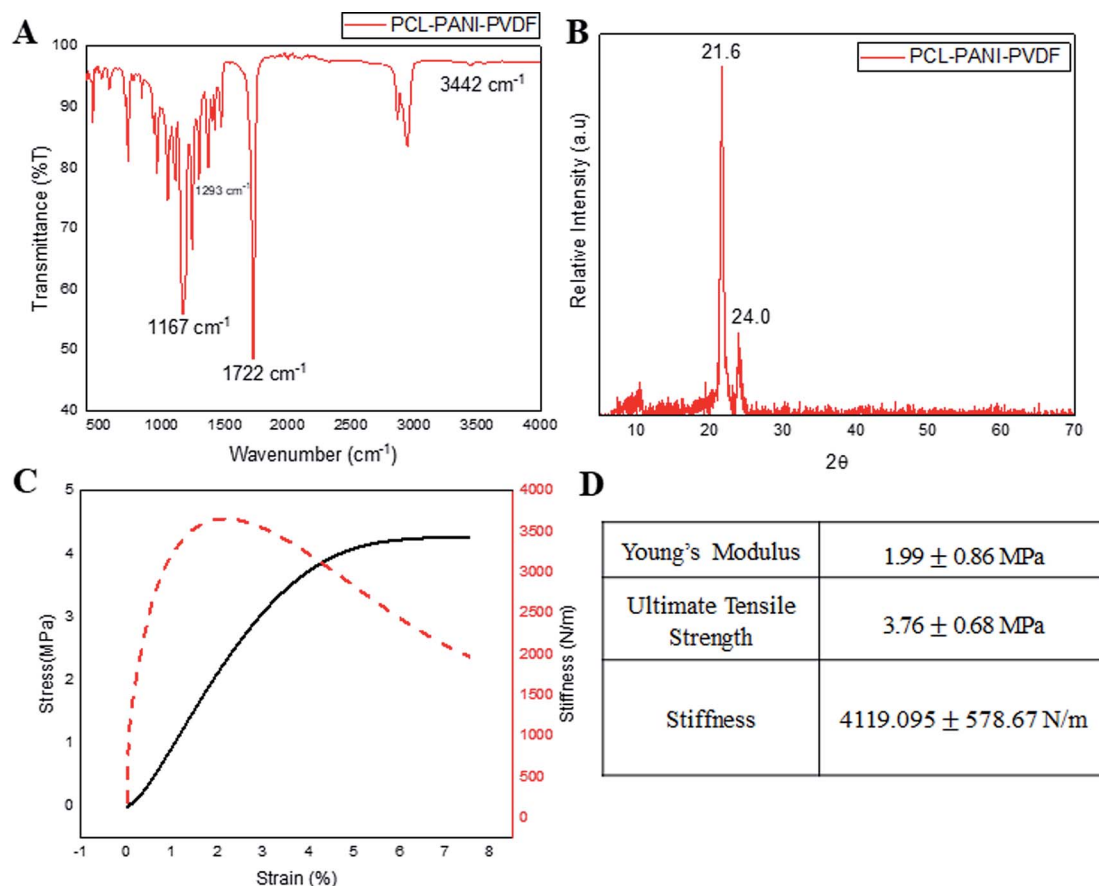


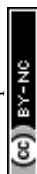
Fig. 2 The surface and mechanical characterization of the PCL–PANI–PVDF scaffolds was done. (A) Surface characterization was done using FTIR spectroscopy. The peaks unique to the functional groups present in PCL, PVDF and PANI were identified. (B) Further surface analysis was done using XRD analysis which showed two crystalline peaks characteristic of crystalline PCL nanofibers. (C) The mechanical characterization of the scaffolds was done by measuring the stress–strain response of the scaffold under isothermal conditions. (D) The table summarizes the mechanical properties of the scaffold.

PANI–PVDF had a much higher matrix stiffness in comparison to PCL and PCL–PVDF (Fig. S4†). The PCL–PANI–PVDF scaffold had a maximum stiffness of 4119.095 ± 578.67 N m⁻¹. Higher matrix stiffness (>4 kN m⁻¹) is generally favored since it promotes cell proliferation, alignment and elongation.^{26,27}

The H9c2 cells were fixed and stained with DAPI and Alexa Fluor® 488 Phalloidin to visualize the nucleus and the actin filaments respectively as seen in Fig. 3. The cell–scaffold interaction is different on each type of scaffold. The cells on PCL scaffold clumped together and lacked projections typical of cardiomyoblasts while the cells on PCL–PVDF scaffold exhibited relatively better cell alignment and cell spreading, the cells were circular, and nuclei were not elongated as inferred from Fig. S5†. Neither PCL nor PCL–PVDF scaffolds supported cell penetration into the scaffold till day 7. In the PCL–PANI–PVDF scaffold the cells appeared aligned with actin filaments stretching across the length of the fibers. The cells also appeared to have penetrated through the different layers of the scaffold with similar actin distribution. The distinct elliptical nuclei indicate nearly identical cell separation and spread across the scaffold as seen in Fig. S5†. The morphology is beneficial for cellular alignment as inferred from the cell

deviation graph in Fig. S6†. The FESEM images of PCL–PANI–PVDF scaffold (Fig. 4a and b) clearly demonstrate cells attached to the aligned fibers of the scaffold. The cells were not only attached but also integrated into the scaffold confirming cellular infiltration. In addition, there was no clumping or crowding of the cells and the overall alignment and integrity of the scaffold was maintained. The FESEM images of cells on PCL and PCL–PVDF scaffolds (Fig. S7†) showed that cells had little or no cell spreading. The cells were clumped in regions and didn't infiltrate through the scaffold.

In summary, we have fabricated acellular scaffolds with mechanical anisotropy and inherent electroactive properties due to its component polymers ensuring cell penetration, migration and alignment. External stimuli or growth factors were not used to promote cell growth. The polymer blend used can be tuned to alter chemical composition and mechanical structure paving the way for research into acellular scaffolds for various applications like cardiac patches, skin grafts and organ regeneration. The scaffold also has the benefit of being readily available and can be synthesized with limited resources and stored at room temperature.



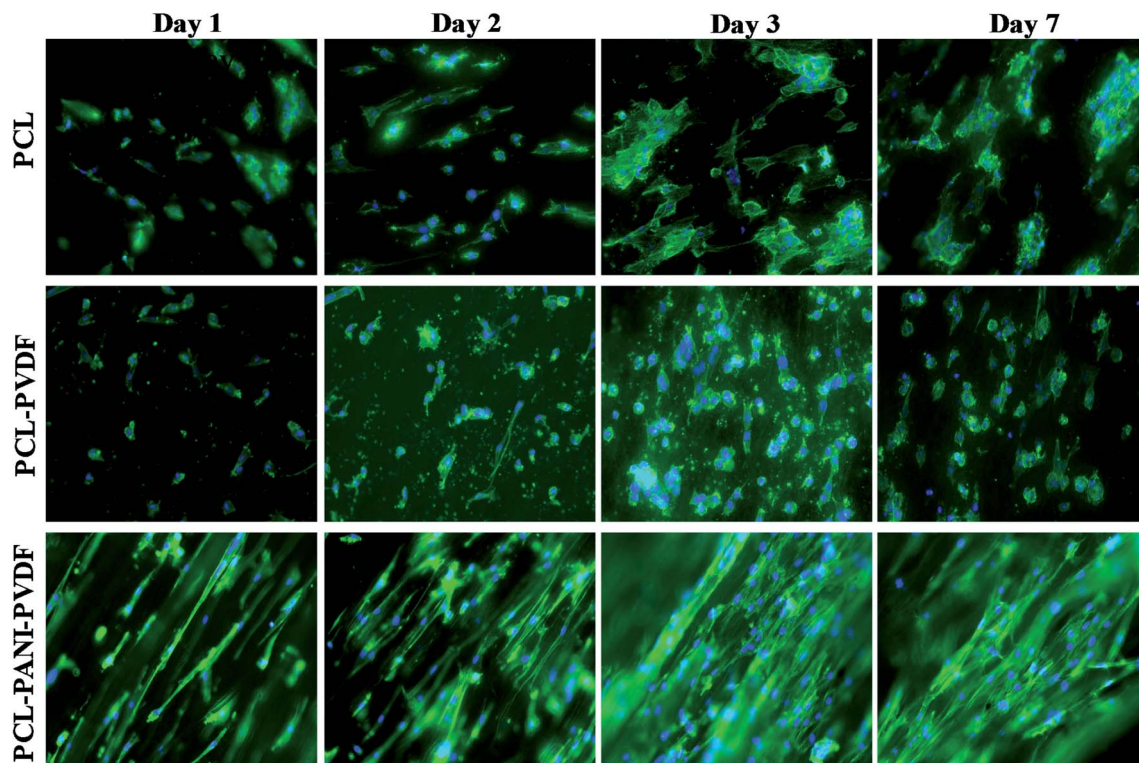


Fig. 3 Fluorescent microscope images of rat cardiomyoblasts (H9c2) cells on PCL, PCL–PVDF and PCL–PANI–PVDF on days 1, 2, 3 and 7. The nuclei were stained with DAPI (blue) and F-actin filaments were stained with Alexa Fluor® 488 Phalloidin (green). Images captured at 20 $\times$ magnification. The cells are aligned on PCL–PANI–PVDF and have a defined orientation as compared to PCL and PCL–PVDF scaffolds. The cells also infiltrated the scaffolds in the PCL–PANI–PVDF as indicated by the blurred regions present at different depths of the scaffold.

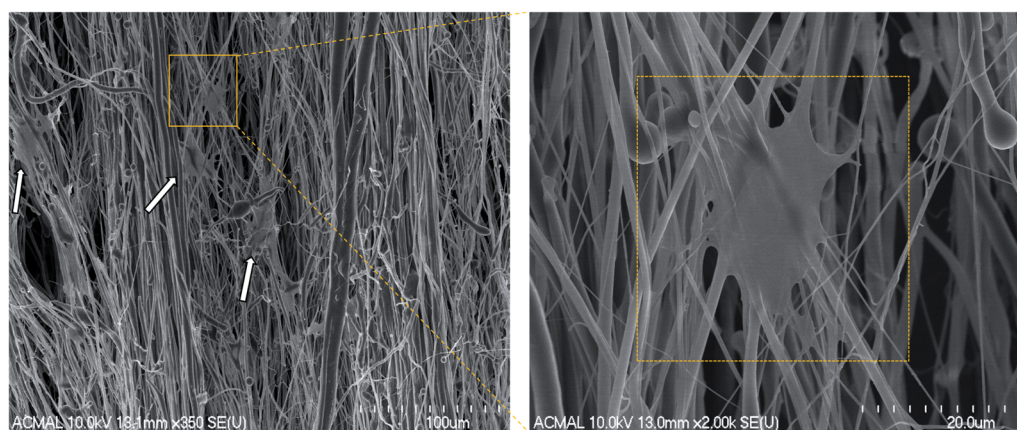


Fig. 4 Field Emission Scanning Electron Microscopy (FESEM) images of rat cardiomyoblasts (H9c2) cells on PCL–PANI–PVDF on day 1. The white arrows point to the location of the cells on the scaffold. The interaction of the cells is clearly visible on the high magnification image (right).

Conclusion

Here we have successfully demonstrated the alignment of cells on an acellular scaffold without the use of an external environment or additional growth factors. The scaffold is composed of a blend of three polymers: PCL, PANI and PVDF. Coaxial electrospinning was used to obtain the scaffolds with controlled surface morphology. The fibers were characterized for their

surface morphology, mechanical strength, surface properties, and the cell–scaffold interaction was demonstrated using rat cardiomyoblasts (H9c2). The cell–scaffold interaction was compared to electrospun aligned PCL and PCL–PVDF scaffolds. The cells on the PCL–PANI–PVDF scaffold exhibited cellular alignment and proliferated along the direction of fibers without any external stimuli in comparison to PCL and PCL–PVDF scaffolds.



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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