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PAPER

Surface acoustic waves induced micropatterning of cells in gelatin methacryloyl (GelMA) hydrogels

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Abstract

Acoustic force patterning is an emerging technology that provides a platform to control the spatial location of cells in a rapid, accurate, yet contactless manner. However, very few studies have been reported on the usage of acoustic force patterning for the rapid arrangement of biological objects, such as cells, within a three-dimensional (3D) environment. In this study, we report on a bio-acoustic force patterning technique, which uses surface acoustic waves (SAWs) for the rapid arrangement of cells within an extracellular matrix-based hydrogel such as gelatin methacryloyl (GelMA). A proof-of-principle was achieved through both simulations and experiments based on the in-house fabricated piezoelectric SAW transducers, which enabled us to explore the effects of various parameters on the performance of the built construct. The SAWs were applied in a fashion that generated standing SAWs (SSAWs) on the substrate, the energy of which subsequently was transferred into the gel, creating a rapid, and contactless alignment of the cells (< 10 s, based on the experimental conditions). Following ultraviolet radiation induced photo-crosslinking of the cell encapsulated GelMA pre-polymer solution, the patterned cardiac cells readily spread after alignment in the GelMA hydrogel and demonstrated beating activity in 5–7 days. The described acoustic force assembly method can be utilized not only to control the spatial distribution of the cells inside a 3D construct, but can also preserve the viability and functionality of the patterned cells (e.g. beating rates of cardiac cells). This platform can be potentially employed in a diverse range of applications, whether it is for tissue engineering, in vitro cell studies, or creating 3D biomimetic tissue structures.

1. Introduction

The control over the spatial location of cells within three-dimensional (3D) microenvironments is of particular interest when creating biomimetic tissue constructs that can recapitulate native tissues. To achieve precise patterning of cells within hydrogels, various approaches have been investigated [1, 2]. These approaches have either appeared in the form of indirect patterning (e.g. surface modifications for
differential cell adhesion, micro-molding [3] and photolithography [4]) or direct patterning (e.g. dielectrophoresis [5, 6], optical [7], magnetic patterning methods [8] and bioprinting [9, 10]). While indirect patterning methods have shown their ability to precisely position cells, they can be both expensive and time consuming. Additionally, one crucial shortcoming associated with indirect patterning techniques is the lack of proper control over cell density and proximity within the encapsulating biomaterial [1]. On the other hand, most direct patterning approaches rely on encapsulation of cells within hydrogels using electrical/magnetic/optical forces, or alternatively use inkjet or laser assisted printing approaches [11–20]. However, most of these approaches have major limitations in precisely positioning cells within hydrogel. For example, the main limitations of bioprinting technology is the relatively long amount of time necessary to correctly achieve the spatial location of cells within hydrogel and the lack of proper control over cell density. In addition, the magnetic approach needs to prove applicability in clinical settings due to the potential risk of the magnetic micro and nano particles on the metabolism and functionality of the cells as described in the literature [21, 22]. Further limitations are found when utilizing electrical forces, which largely depend on the conductive nature of the medium and can be cytotoxic to cells, altering their functionality. [23–26]. Finally, optical tweezers have also been employed to generate various patterns of cells in hydrogels. This technique, however, does not allow for the rapid organization of large populations of cells [27].

Aside from the approaches mentioned above, the application of surface acoustic waves (SAWs) has been recently reported as a non-invasive, simple and inexpensive approach towards the precise and rapid patterning of microparticles and cells [28–30]. SAWs can be easily stimulated under biocompatible conditions to create cell alignment without necessarily needing to tune the electrical conductivity or temperature of the assembly medium. This is because the SAWs technology depends neither on the conductivity of the assembly medium, nor does it generate significant heat during patterning. This technique has potential for various applications ranging from the focusing [30], separation [31], alignment [32, 33], and directionality of microparticles [34]. For instance, Shi et al have demonstrated the biocompatibility of this method using red blood cells and bacteria in two-dimensional (2D) manner [30]. To achieve manipulation of single microparticles, cells, and organisms using SAWs, the use of standing SAWs (SSAWs), has been investigated. SSAW usage was achieved by the creation and superposition of two SAWs while traveling in the opposite directions to one another [35,36]. Finally, complex 2D cell-patterning [37] and 3D cell manipulation in microfluidic devices have also been studied recently [38].

Despite successful reports of the previously mentioned acoustic approaches toward generating 2D or 3D cell patterns and enabling new tissue engineering applications, the patterns must be sustained after removing the initial application of the SAWs. One alternative towards that end is to photo-crosslink the hydrogel matrix after patterning the cells. One example is Chen et al, where SSAWs were employed to pattern cell on a low-adhesion surface after sedimentation in fibrin hydrogel in a 2D manner, and crosslinking of the fibrin was achieved in about 20 min at room temperature [21].

More recently, the same group reported the use of bulk acoustic waves to pattern cells in horizontal planes, engineering multilayered, 3D brain-like constructs [39]. In their setup, the authors placed a piezoelectric transducer underneath a cell-contained pre-polymer, allowing for the emission of bulk acoustic waves within the fibrin hydrogel. In the study, the emission of bulk acoustic waves were used to form planar sheets of cells. While this approach was creative in its application, it may not be applicable for all settings of tissue engineering. This is mainly because bulk acoustic waves have volumetric characteristics, which makes this approach a challenge to integrate with small-scale, lab-on-a-chip techniques. As such, these issues favor the use of SAWs patterning for microfluidic assembly chambers and tweezer setups for biological and bioengineering applications.

Within our study, we make use of SSAWs and acoustic radiation force (ARF) assembly to spatially localize cells encapsulated within a photo-cross-linkable gelatin methacryloyl (GelMA) hydrogel matrix. These GelMA hydrogels are suitable platforms for encapsulated cells to easily attach, proliferate, and migrate with acceptable viability, which enables for 3D cell organization [40, 41]. By varying multiple parameters such as the frequencies corresponding to the SAWs, the concentration of GelMA molecules, as well as the duration of ultraviolet (UV) light exposure when curing GelMA hydrogels, we aim to further explore the usage of SSAWs as a contactless, direct patterning technique for localizing large numbers of cardiac cells within a hydrogel environment.

2. Experimental section

2.1. SAW setup fabrication

In this study, we used 128°Y-cut X-propagating 3 inch lithium niobate (LiNbO3). The photolithography process started with the cleaning of the piezoelectric substrates in an acetone bath (5 min sonication), subsequent rinsing under flowing water for 2 min, and blow-drying with N2. The substrate was treated in O2 Plasma for 3 min at 100 W. The sample was then spin-coated with photoresist (s1818, MicroChem, Newton, MA) for 30 s at 4000 rpm, and baked on a hot plate at 95 °C for 3 min. The transducer pattern on the mask
was transferred into the photoresist layer by UV light exposure for 5 s (MA-6, SUSS Micro Tec AG) and was then developed using a 1:1 (v:v) dilution of photoresist developer (MF CD-26, Microposit) for 1.5 min followed by a rinse under flowing RO water for 5 min. After blow-drying, the sample was treated with O₂ Plasma for 2 min at 60 W. A 20 nm titanium adhesion layer and a 100 nm gold layer were evaporated on their surfaces prior to the lift-off in acetone (at 45 °C) for 2 h. Finally, all unwanted parts of the metal layer were removed by lifting off the photoresist layer underneath, and the slanted-finger interdigital transducers (SFTTs) were obtained. The device was immobilized on the stage of a microscope (Zeiss Observer D1). A radio frequency signal generator (TTi TG5011 LXI) was connected to the SFTTs. Thereafter, coherent signals at a desired frequency with input power from −7 to −12dBm were provided to generate SAW on the lithium niobate substrate. The frequency of the generated acoustic wave was tuned by altering the input frequency of the transducers.

2.2. GelMA preparation

GelMA molecules was prepared according to our established protocol [40]. A 10% solution of gelatin from porcine skin (Sigma Aldrich) in Dulbecco’s phosphate buffer saline (DPBS, Gibco) was prepared by stirring for two hours at 60 °C. To initiate the methacrylation reaction, 0.025 ml g⁻¹ of methacrylic anhydride (Sigma Aldrich) were added dropwise to the mixture and subsequently incubated for 60 min at 60 °C under constant stirring at 500 rpm. Two volumes of DPBS were added to stop the reaction and dialysis was performed for 5 days. Subsequently, the GelMA solution was filtered and freeze-dried for 5 days.

2.3. Cell isolation

Neonatal rat ventricular cardiomyocytes were obtained by isolation based on a well-known procedure, approved by the Institute’s Committee on Animal Care [42]. For this purpose, 2 day old Sprague-Dawley rats were used as model animals. To guarantee physiologically reliable and reproducible cell behavior, cardiomyocytes were applied in experimentation directly after the aforementioned isolation procedure. In addition to that, to separate cardiac fibroblasts, the cells were enriched by pre-plating for one hour. In this process, cardiomyocytes were harvested from the supernatant media. The remaining cardiac fibroblasts were continually kept in culture using DMEM (Gibco, USA) holding 10% FBS (Gibco, USA), 1% L-Glutamine (Gibco, USA) and 100 units ml⁻¹ penicillin–streptomycin (Gibco, USA). This was continued until 90% confluence was reached. Consecutively, the cells were detached (by trypsin) and harvested. Prior to the application of electrical stimulation, the multilayer samples were kept in culture for up to 5 days.

2.4. Preparation of the experimental setup

Following the described isolation, cardiomyocytes were mixed with cardiac fibroblasts in a 1:1 ratio to achieve a final concentration of 2 million cells ml⁻¹. The cell mixture was suspended in a 5% GelMA solution. For this study, a glass slide was placed on the LiNbO₃ wafer coupled with water, and a PMMA chamber was placed on top of the glass slide. The dimensions of the charmer were 0.6 cm (length) × 0.4 cm (width) × 0.8 mm (height). A 45 μl droplet of cell encapsulated GelMA solution was released in the chamber and was covered with a TMSPMA glass slide. The SAWs were generated using the signal generator (TTi TG5011 LXI). After the application of SAW, samples were exposed to UV immediately and were placed in 12 well plates, each filled with 1 ml of Dulbecco’s modified eagle medium (DMEM). In total, for each of the applied frequencies, 12 samples were prepared. The values of the frequencies applied in this study are 3.4, 4.6 and 6.4 MHz. The multi-well plate was then placed in an environmental incubator for 2 h to allow cardiomyocytes and cardiac fibroblasts to adhere before replacing with medium. To reduce the risk of contamination, the medium was changed 24 and 48 h after seeding.

2.5. Viability assay (live/dead) and actin DAPI staining

Viability test was performed by incubating cell-laden samples for 15 min at 37 °C with a mixture of 2 μl ml⁻¹ ethidium homodimer-1 (EthD-1, red, dead cells, Invitrogen) and 0.5 μl ml⁻¹ calcein AM (live cells, green, Invitrogen). Fluorescence Microscopy was performed on a Zeiss Observer.D1 microscope using an X-Cite Series 120Q fluorescence source. We used three samples to calculate cellular viability of patterned cardiac fibroblasts at the 6.4 MHz resonant frequency with short UV exposure time (<10 s). Samples were fixed in 4% paraformaldehyde (Electron Microscopy Sciences), washed in DPBS and subsequently stained with phalloidin (life Technologies, labeled with Alexa F 594) and DAPI as described in the manufacturer’s manual. Microscopy was performed as described above.

2.6. Finite element modeling

Finite element simulations were conducted to evaluate the principles of the device and to determine the mathematical characteristics of the cells manipulation and patterning in the proposed setup. In acoustic force patterning, the ARF $F_{\text{rad}}$, created by the scattering of acoustic waves on the cells, will manipulate the position of the cells. The time-average ARF as a gradient of a potential function $F_{\text{rad}}$ on a single small spherical cell of radius $a$, density $\rho_c$, and bulk modulus $K_c$ placed in a standing wave field is given by [43]:

$$F_{\text{rad}} = -\nabla U_{\text{rad}},$$

(1)
Table 1. Simulation model parameters at 37 °C.

<table>
<thead>
<tr>
<th></th>
<th>5% GelMA pre-polymer solution</th>
<th></th>
<th>Cell</th>
</tr>
</thead>
<tbody>
<tr>
<td>Speed of sound⁴</td>
<td>1534 m s⁻¹</td>
<td>Speed of sound⁴</td>
<td>1525 m s⁻¹</td>
</tr>
<tr>
<td>Viscosity³</td>
<td>0.0025 Pa s</td>
<td>Poisson ratio⁴</td>
<td>0.5</td>
</tr>
<tr>
<td>Density⁵</td>
<td>1020 Kg m⁻³</td>
<td>Density⁵</td>
<td>1060 Kg m⁻³</td>
</tr>
<tr>
<td>Viscous boundary layer thickness⁶</td>
<td>0.35 μm</td>
<td>Bulk modulus⁶</td>
<td>2.4 GPa</td>
</tr>
<tr>
<td>Bulk modulus⁶</td>
<td>2.4 GPa</td>
<td>Cell diameter⁸</td>
<td>10 μm</td>
</tr>
</tbody>
</table>

* Measured;  
* Calculated as δ = Sqrt(t(y/(ρvf)) from [47];  
* Calculated as k = ρc² from [47];  
* Assume to resemble water;  
* Calculated as (μc²)(1 + v)/(3(1 – v)) from [47].

\[
U_{rad} = (4\pi/3)\alpha^3 \left[ \left( f_1/(2\rho_g c_g^2) \right) \langle p^2 \rangle - (3f_2\rho_g/4) \langle v^2 \rangle \right],
\]

\[
f_1 = 1 - k_g/k_c \quad \text{and} \quad f_2 = 2(\rho_c - \rho_g)/(2\rho_c + \rho_g)
\]

in which, \( \rho_g \) and \( \rho_c \) are the density of the fluid (GelMA pre-polymer solution) and cell, \( c_g \) is the speed of sound in fluid, \( k_g \) and \( k_c \) are bulk modulus of the fluid and cell, and finally \( p \) and \( v \) represent pressure and velocity fields, respectively. All of the simulations were performed in COMSOL Multiphysics 5.0. As mentioned, for SFITs, each sub-channel was considered a uniform IDT electrode; so the simulations were conducted for cross sectional planes (x-z plane or side view) of the channel containing GelMA pre-polymer solution. The same pattern will repeat in other cross sectional planes of the mentioned sub-channel area. In the above-described cross-sectional model, we took into account the whole setup including piezoelectric substrate, glass slide and GelMA solution.

Material properties of LiNbO₃ used in our simulations are characterized in previous studies [48]. Also, material parameters corresponding to GelMA pre-polymer solution and cells are tabulated on table 1. Making use of the CFD eigenfrequency module, the resonance frequency of the device was obtained. Subsequently, the generated acoustic pressure field was modeled using a frequency domain approach. Finally, a particle-tracing module was utilized to simulate the effects of the pressure field on the movements of cells within the GelMA pre-polymer solution. The simulations associated with the top view of the GelMA solution were performed using the values obtained from the previously described cross sectional simulations. So, for the simulations in the top view, we modeled GelMA solution exiting by defining acceleration on the walls within the frequency domain and then used particle tracing for evaluating cell movements.

2.7. Characterization of beating behavior

Spontaneous beating behavior was assessed every day by light microscopy (Nikon, Eclipse, Ti-S, Japan/Zeiss Observer D1) at 10× magnification using a CCD camera connected to the microscope alongside a video capture program. During the imaging process, the temperature of the sample environment was kept constant at 37 °C through the use of a heating chamber. To calculate the beating frequency, an in-house, custom-developed MATLAB script was employed to analyze the video sequences, frame by frame. Three independent experiments with a minimum of three replicates each were used for the calculation.

2.8. Statistical analysis

The statistical significance was determined using two-way ANOVA statistical analysis provided by the GraphPad package (GraphPad Prism 5.02, GraphPad Software). The same software package was used to arrive at the mean and standard deviation of the presented plots.

3. Results and discussion

In an exploration of potential approaches towards directing cell patterning in tissue engineering, we have investigated the use of ARF to localize cells without contact. This is done by using SAWs to relocate cells inside the GelMA pre-polymer solution. Figure 1(A) illustrates the concept of employing SAWs to localize cells as a way to direct cell patterning. Pressure fluctuation generated by SAWs lead to periodic distribution of nodes (where the pressure values drop to zero) and anti-nodes (where the pressure values are at a maximum). In acoustic force patterning, the ARF, which is a time-averaged net force created by pressure fluctuations, will manipulate the position of the cells. Based on the properties associated with the type of cells and the fluid surrounding the cells, this force can push the cells toward the pressure nodes. Depending on the frequency of the SAWs, the above-described patterning of cells can be tunable to a desirable spatial distance. To do this, our device was designed to generate multiple frequencies on a single platform. Supplementary figure 1 shows the peak of resonance frequencies exhibited by the transducers on the wafer. To manipulate the position of the cells within the GelMA pre-polymer solution, various peak frequencies were applied. Here, the maximum and minimum distance of the slanted fingers can limit the range of working frequencies of the setup. In the other words, the generated wavelength of the piezo-electric substrate in the excited subchannel area is equal to the period distance of the slanted fingers [49]. To generate these SAWs, piezoelectric transducer elements are used, as shown in figures 1(B) and (C). The piezoelectric transducers used for this setup are known as...
SFITs. A SFIT can be considered as a cascade of sub-channels, which are called interdigital transducers (IDTs). Each sub-channel possesses its own resonance frequency, which means the SFIT is capable of operating over a range of frequencies \[49\]. Therefore, utilizing a frequency that is equal to the resonance frequency of a sub-channel, creates resonance, which in turn, produces waves that correspond to the width of the sub-channel. Figures 1(B) and (C) show the schematic and depiction of the piezoelectric transducers utilized in the experimental setup, which produced one-directional waves and subsequently patterned cells in parallel lines. Furthermore, to create more complex structures within GelMA hydrogel, we extended the fabrication of the experimental setup in a manner that allowed for the creation of SSAWs (in two directions perpendicular to one another), which would reorganize the cells inside the pre-polymer solution. A schematic of the two-directional setups and a depiction of the apparatus are illustrated in figures 1(D) and (E).

Due to the promising potential of GelMA hydrogel’s use in cell encapsulation, we aimed to examine its effectiveness for achieving 3D localization of cells while maintaining long-term cell viability. One way to achieve such patterning in 3D is to generate SSAWs on the surface of the piezoelectric substrate, which can then be transferred into the bulk volume of the GelMA pre-polymer solution. As schematically shown in figure 2(A), a glass treated with 3-(trimethoxysilyl) propyl methacrylate (TMSPMA) was used as an intermediate layer to integrate the waves into a poly(methyl methacrylate) (PMMA) channel. The chamber was then filled with 5% GelMA pre-polymer solution, encapsulating the cells, and was covered with a coverslip (thickness of 0.18 mm). Next, UV light was irradiated from the topside of the device for 10–30 s to crosslink the GelMA pre-polymer solution and maintain the generated pattern of the cells.

Figures 2(B)–(G) displays numerical modeling in a cross section of GelMA pre-polymer solution, demonstrating ARF’s propensity for cell manipulation in the described setup. Pressure fields that resulted from the

Figure 1. (A) The concept of employing SSAWs to localize and pattern cells. ARF will push cells toward pressure nodes and pattern cells in parallel lines; (B) schematic of the experimental setup employed to generate SSAWs, leading to the localization of the cells encapsulated within the GelMA pre-polymer solution; (C) real image, top-view, of the piezoelectric transducers utilized in the experimental setup. The non-uniform patterns of the electrodes allow variations of the frequency of the generated waves; (D) top-view schematic of the two-directional piezoelectric setup and the GelMA pre-polymer solution; (E) real image of the experimental setup including the transducers and piezoelectric substrate.
transfer of the SAWs into the GelMA segment are illustrated in figure 2 (B). The transferred waves reflect off the hard wall boundaries such as PMMA and glass, creating rectangular standing waves. This pressure field causes the creation of nodes and anti-nodes, which are shown in the figure 2 (B) as elliptical and circular dotted lines, respectively. The trajectories associated with the cellular movements are presented in color. This is done with the minimum velocities shown in blue and the maximum shown in red. The colored disks in the center of the circles are denoting the final destinations of the cells. (C) after 0; (D) 0.5; and (E) 1 s; (F) acoustic radiation force will reduce as the cells approaching the nodes and therefore; (G) the velocity of cells will decrease when they are going toward them.

As previously mentioned, one of the advantages of using SAWs, as opposed to bulk acoustic waves, is for their ease of use in microfluidic setups, as well as their capability in generating cell patterns in 3D space (in spite of being surface waves). To elaborate on this further, simulations revealed that if the height of the photo-crosslinked hydrogel in the z direction is thicker than ~80 μm, then the position of the pressure nodes and subsequently patterned cells will vary from one longitudinal plane to another. This phenomenon is evident from the simulations presented in figure 2(B); demonstrating a shift in the position of the pressure nodes (shown with elliptical dotted lines) from one z-plane to another. In addition, this phenomenon can also be seen on the parallel cell lines pattern (supplementary figure 1); lying beside the bright cells on the surface of the hydrogel construct, other cells can also be observed in the middle of the parallel cell lines, with dimmer brightness. ARF is negligible in nodes or anti-nodes, and is at its maximum amplitude between them. In figure 2(B), referring to the midline between node and anti-node shown as the ‘X–Y’ line, the ARF (and therefore the cells’ velocities) decrease when moving from the midline toward acoustic node. This is further illustrated and quantified in figures 2(F) and (G): as expected, when the cells which are initially on the X–Y line approach the acoustic node, the ARF decreases, and as a result, the velocity of the cells decrease as well.

To investigate the capability of the acoustic force assembly technique in enabling more complex structures within GelMA pre-polymer solution, as mentioned earlier, we extended the fabrication of the
experimental setup in a manner that allowed for the creation of SSAWs (in two-directions perpendicular to one another). A schematic of the bi-directional setup and a depiction of the apparatus are illustrated in figures 1(D) and (E). The top-view results for the complex orthogonal patterns formed by this two-directional setup are also shown in figure 3. The superposition of the two perpendicular SSAWs results in the formation of a square-like pressure field, with double the amplitude of each one-directional SSAWs (figure 3(A)). This pressure field forms net-like, patterns of the cells from top view (figure 3(B) and supplementary figure 3). The pressure derived from two-directional application of the SSAWs is plotted in figure 3(C). In this plot, the pressure distribution along the line X–Y is depicted by a solid red line. This pressure distribution is related to the time corresponding to the peak of the first half cycle of the wave. This peak is associated with a timing of \( t = 1/(4f) \) where \( t \) refers to the time and \( f \) is the frequency. Similarly, the pressure distribution along the line X–Y at the negative peak of the second half-cycle of the wave is illustrated with the dotted blue line. This timing is associated with \( t = 3/(4f) \). Such plots not only help us gain an understanding of the cells’ patterning, but also provides further information regarding the cells’ localization (e.g. higher amplitude pressure will results in faster organization of the cells). This data, in addition to the data associated with the one-directional setup, serve as proof-of-principle evidence, suggesting the promise of the described acoustic force assembly technique for its use in cell localization.

For the cellular alignments associated with the one-directional setup shown in figure 1(C), simulations of patterning from the top view (C–C view of figure 2(E)) are illustrated in figures 4(A)–(C). As shown in these figures, the pressure field causes the cells to pattern in parallel lines. Following the simulation investigations, attempts were made to experimentally assess the prospect of cell-alignment within GelMA pre-polymer solution using ARF. To do so, 2 million cardiac fibroblasts and cardiomyocyte cells were mixed with a GelMA pre-polymer solution, and the mixture was loaded into a rectangular-shaped assembly chamber with 0.8 mm in depth. The random distribution of cells prior to the application of acoustic forces is demonstrated in figure 4(D). Subsequent to turning on the piezoelectric transducers, SAWs are generated on the surface of the LiNbO_3 substrate, a piezoelectric material, with a reasonable level of optical transparency and high electro-mechanical coupling. Following generation of the SSAWs on the surface of the piezoelectric substrate, the energy created is transferred into the bulk volume of the cell-containing GelMA pre-polymer solution, resulting in a pressure fluctuation that can affect positioning of the cells within the solution (supplementary movie 1). The entire assembly time for the samples varied from one to another (based on the input power, frequency, and physical properties of GelMA pre-polymer solution), but generally remained under 10 s. Different

![Figure 3](image-url)

*Figure 3.* (A) Top-view pressure fields resulted from the orthogonal acoustic wave interaction created using numerical simulation, assuming a working frequency of 4.6 MHz; (B) corresponding simulated and experimental patterns, when zoomed in. The scale bar shows 100 \( \mu \)m; (C) pressure fluctuation in GelMA pre-polymer resulted from transferred two-directional SSAWs. Pressure along the dotted X-Y line in the positive peak \((t = 1/(4f))\) and negative one \((t = 3/(4f))\) are depicted with solid red and blue dotted lines, respectively. \((t \) refers to the time and \( f \) refers to the frequency).
As depicted in UV light crosslinking of the hydrogel pre-polymer and cardiomyocytes after the use of ARF assembly and tuning, the proposed setup does possess the control frequencies were used to prove that through frequency microscope images of patterned cardiac Figures 4 the assembled pattern remains stable and intact. After UV irradiation and stopping the standing waves, frequencies of 3.4 MHz and 6.4 MHz, respectively. acoustic standing waves corresponding to resonant acoustic force assembly approach. As indicated earlier, the spatial distribution can be varied by altering the input resonant frequency. However, it is of equal importance to consider overall cell viability and proliferation when utilizing techniques to align cells.

To optimize the UV crosslinking times, the cellular viability of patterned cardiac fibroblasts in GelMA hydrogel after different UV exposure times is quantified. (supplementary figure 4). A significant loss of cell viability was observed for longer UV exposure times. Therefore, a short UV exposure time of around 10 s was selected to make GelMA hydrogel. To evaluate the cytocompatibility of the acoustic force assembly method, the viability of the aligned cardiac cells within the GelMA hydrogel was evaluated up to 7 days (figure 5 and supplementary figure 5). A live/dead assay was performed and quantified for cardiac fibroblasts immediately following the patterning at the 6.4 MHz resonant frequency. As shown in figure 5(A), close to 90% cellular viability was revealed at short UV exposure time (<10 s), demonstrating the minimal effect of the acoustic force assembly method on cell viability. Furthermore, to assess morphological behavior of the cardiac cells during the culture period,
cardiac fibroblast and cardiomyocytes within the 5% GelMA hydrogel were cultured up to 7 days post-assembly, and were stained with F-actin and DAPI. Figures 5(B) and (C) represent the images of F-actin/DAPI staining of cells after 5 and 7 days, respectively. On day 5, the encapsulated patterned cardiac cells demonstrated spreading, and by day 7 they started forming interconnected cellular networks. Finally, to validate the functionality of such patterned constructs, figure 5(D) is shown to demonstrate the development of beating over time (movie 2). As evident from the plot, the beating rates increased with the number of cell culture days that have elapsed and was at its peak on day 9; after which the rate of beating started to decline rapidly. The decrease in beating rate with increasing culture time was also shown in previous studies, in which rat neonatal cardiomyocytes were cultured on hydrogel-based tissue-engineered models [50, 51]. This decrease might be attributable to the phenotypical transition of fetal cardiomyocytes towards the neonatal stage [51]. These results confirm that the SSAWs assembly approach reported in this study could be employed to align cells within a pre-defined assembly setup. Most importantly, this technique allowed cells to remain viable and grow into functional constructs that demonstrated cardiac beating.

4. Conclusion

In this study, we have demonstrated that the acoustic force assembly technique, in combination with GelMA hydrogels (possessing adjustable physical, chemical, and mechanical characteristics) can provide a setting to facilitate the generation of various 3D cellular patterns and constructs in a rapid and contactless manner. Towards this purpose, SSAWs were applied on the surface of the piezoelectric substrate, resulting in energy transfer into the cell-laden GelMA pre-polymer solution, leading to rapid alignment of the cardiac cells. Simulations were conducted to gain an understanding of cell behaviour and to find the optimal design parameters for our acoustic setup. Experimentation was also conducted in one and two-directional setups based on in-house fabricated piezoelectric transducers. This allowed us to study and refine the effect of various parameters, including the optimal operating frequency of the setup. Moreover, it was demonstrated that the acoustic force assembly method can be utilized to control spatial distribution of the cells while maintaining cell viability and function (e.g. demonstrating cardiac beating). Therefore, the acoustic force assembly approach presented here offers a novel technique for the engineering of complex 3D tissues. Patterning techniques are fundamental for investigation of cellular behaviours such as cell–cell interaction and drug development studies in various settings [35]. One limitation of the one-directional patterning method presented in this study is that the created structures is limited to line patterns. However, in the context of cardiac tissue engineering and in order to capture the proper functionality, it is vital to recapitulate the 3D aspects of the microenvironment of the cardiac fibres [52]. Another limitation of our approach is that the 3D patterned constructs are
produced in micro dimensions only. Despite the limitations, it makes it uniquely suitable for micro-fluidic devices, with unique advantages for a number of applications such as drug screening.

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