3D Bioprinting of Mature Bacterial Biofilms for Antimicrobial Resistance Drug Testing

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Abstract

The potential to bioprint and study 3D bacterial biofilm constructs could have great clinical significance at a time when antimicrobial resistance (AMR) is rising to dangerously high levels worldwide. In this study, clinically relevant bacterial species including *Escherichia coli*, *Staphylococcus aureus* (MSSA), Methicillin-resistant *Staphylococcus aureus* (MRSA) and *Pseudomonas aeruginosa* were 3D bioprinted using a double-crosslinked alginate bioink to form mature bacteria biofilms, characterized by confocal laser scanning microscopy (CLSM) and fluorescent staining. Solid and porous bacteria-laden constructs were reproducibly bioprinted with thicknesses ranging from 0.25 to 4 mm. We demonstrated 3D bioprinting of thicker biofilms (>4mm) than found in currently available *in vitro* models. Bacterial viability was excellent in the bioprinted constructs, with CLSM observation of bacterial biofilm production and maturation possible for at least 28 days in culture. Importantly, we observed the complete five-step biofilm life cycle *in vitro* following 3D bioprinting for the first time, suggesting the formation of mature 3D bioprinted biofilms. Bacterial growth was faster in thinner, more porous constructs whilst constructs crosslinked with BaCl$_2$ concentrations of above 10 mM had denser biofilm formation. 3D MRSA and MSSA biofilm constructs were found to show greater resistance to antimicrobials than corresponding two-dimensional (2D) cultures. Thicker 3D *E.coli* biofilms had greater resistance to tetracycline than thinner constructs over 7 days of treatment. Our methodology allowed for the precise 3D bioprinting of self-supporting 3D bacterial biofilm structures that developed biofilms during extended culture. 3D biofilm constructs containing bacterial biofilms produce a model with much greater clinical relevance compared to 2D culture models and we have demonstrated their use in antimicrobial testing.
Introduction

Biofilms can be defined as 3D structured communities of bacterial cells enclosed in a self-produced polymeric matrix, attached to a solid surface or substratum [1]. Bacterial biofilm formation is crucial to establishing chronic infections including respiratory infection[2], orthopaedic infection [3], heart valve infection (endocarditis) [4], and nosocomial infections [5]. In the case of acute infections, bacteria often exist in the planktonic (or free-swimming) state, allowing effective treatment with antimicrobials. However, once a biofilm develops infections are known to be 10-1000 times more resistant to antimicrobial agents, often rendering standard antimicrobial therapy ineffective without more invasive treatment such as surgery [6]. In the United States of America alone, there are 17 million new biofilm-associated bacterial infections that lead to estimated health care costs of $94 billion and 550,000 deaths each year [7]. According to the World Health Organization (WHO), urgent action is required to avoid a “post-antibiotic era”, in which common infections and minor injuries can once again kill; antimicrobial resistance is projected to result in 10 million deaths every year globally by 2050 [8]. Global concern about AMR is compounded by the fact that it has been 30 years since a new class of antibiotics was last introduced [9]. Therefore, increasing importance is being placed on drug screening, and in particular, antimicrobial susceptibility testing (AST), which requires suitable models that more closely resemble in vivo biofilm formation.

The minimum inhibitory concentration (MIC) of antimicrobial agents (defined as the lowest concentration of an antimicrobial agent at which visible bacterial growth is inhibited after overnight incubation) is frequently calculated during AST to assess antimicrobial efficacy and bacterial resistance [10]. Methods to determine the MIC based on 2D planktonic cultures of bacteria are well established [11]. However, determining the minimal biofilm eradicating concentration (MBEC) in biofilm infections is much more challenging. This is primarily because in vivo biofilm formation is three dimensional (3D) in architecture, which differs to most currently available laboratory models that tend to involve 2D biofilm culture [12-14]. AST of planktonic bacteria therefore tends to give misleading results that do not reflect the increased resistance of bacteria living in a 3D biofilm [15, 16]. This has significant clinical implications; for example, antimicrobial agents are usually chosen on the basis of their efficacy against 2D planktonic cultures which are more sensitive to treatment than 3D biofilms. Clinically this is well demonstrated by cystic fibrosis patients, where treatment of P.aeruginosa infection with antibiotics originally developed against planktonic cultures often becomes ineffective once biofilm formation occurs [15]. To develop novel antimicrobials capable of disrupting biofilm formation and resistance in future, 3D in vitro biofilm models more representative of clinical infection are required.

Most commonly used 2D biofilm culture methods attempt to simulate the nature of the in vivo environment by focussing on selected relevant factors such as materials, nutrients and, importantly, fluid flow including drip flow [16], rotating disk [17], microfluidics [18], and flow chamber architecture [19]. Unfortunately, none of these methods mimic the complexity of
the 3D microenvironment and host defence mechanisms [20] and unable to produce biofilm thicknesses beyond 100 µm [21, 22]. In contrast to the current in vitro models, in vivo biofilms can grow beyond 1000 µm in size and are often found embedded within a host’s extracellular matrix, leading to interactions with the host immune system which can further alter biofilm morphology and size [1, 23].

3D bioprinting has developed rapidly as a technique that can deposit living cells and biomaterials in user-defined patterns to build complex tissue constructs “from the bottom up” [24-27]. While there are elegant approaches on 3D bioprinting bacteria and their aggregates [28-32], there has been no report on demonstrating the formation of mature bacteria biofilms. However, the capacity to reliably and reproducibly 3D bioprint bacterial biofilms have several potential benefits. Embedded bacteria have been shown to have increased metabolic activity, AMR and plasmid stability compared to bacteria grown in [33, 34]. 3D bioprinted bacterial biofilms therefore could potentially mirror in vivo bacterial growth and behaviour more closely than traditional 2D models, increasing the potential to investigate critical bacterial quorum sensing (QS) and antimicrobial biofilm penetration [34, 35]. 3D bioprinting also increases the potential to produce biofilm constructs with predesigned dimensions, with a high degree of control possible over biofilm thickness and dimensions. Other benefits of 3D bioprinting biofilm include the potential creation of microbial fuel cells [36], biosensors [37] and biotechnological applications [37-39].

In this paper, we present a novel 3D bioprinting biofilm technology and report the first investigation of the formation of mature bioprinted 3D biofilms and measure their responses to antibiotic drug tests, and drug penetration. Mature biofilms with different thicknesses and structures were designed and bioprinted using a range of clinically relevant bacterial strains. In vitro AST was performed to compare the resistance of 2D cultures versus 3D printed biofilm constructs for the first time. Bioprinting of biofilm constructs with thicknesses greater than previously available in vitro models was also successfully performed.
Materials and Methods

Bacteria-laden bioink preparation

Brain Heart Infusion (BHI) broth (Sigma-Aldrich, UK) powder was dissolved in sterile deionized water to produce a 37 g/ L BHI Broth and then autoclaved. UV-sterilised sodium alginate powder (Protanal LF10/60FT, FMC Biopolymer, UK) was then dissolved in BHI Broth to produce a 4% (w/v) alginate solution. The alginate solution was subjected to magnetic stirring until reaching homogeneity and then sterilised through heating to boiling point (95°C) three times. Solutions consisting of 4% w/v sodium alginate and 0.4% w/v CaCl₂ were then mixed with a volume ratio of 1:1 to create a partially cross-linked 0.2% CaCl₂: 2% sodium alginate hydrogel in a 50 mL conical tube. The hydrogel solution was vortex mixed at room temperature at 1500 rpm for 5 min to produce a homogeneous, partially cross-linked alginate hydrogel. Alginate hydrogels were then stored at 4 °C prior to usage to prevent the growth of contaminants.

Bacterial strains and growth media

Bacterial strains were universally cultured in Brain Heart Infusion (BHI) broth at 37°C whilst shaking. Strains used included Escherichia coli (E.coli clinical isolate, ATCC 25922), Pseudomonas aeruginosa (P. aeruginosa, PAO1, wild type strain, ATCC 47085), Methicillin-sensitive staphylococcus aureus (MSSA, clinical isolate, ATCC 29213) and Methicillin-resistant staphylococcus aureus (MRSA, clinical isolate, ATCC 700788). Chosen strains were routinely maintained on BHI agar (Sigma-Aldrich, UK) plates and stocks kept frozen in glycerol (50% v/v) at -80°C.

Inoculum preparation

Bacterial strains taken from glycerol stocks were streaked on to a BHI agar plate and incubated at 37°C overnight. The following day a single colony was inoculated into 5 mL of BHI broth and incubated overnight at 37°C, with 200 rpm shaking (Mini shaker, Cleaver). The overnight cultures were harvested in the stationary phase after 18 h cultivation. The bacteria were collected by centrifugation (3,000 rpm, 4°C, 5 min) and washed three times with 9% sodium chloride (NaCl) to remove the residual BHI medium. In all experiments, the concentration of bacteria was determined by optical density spectrometry (Eppendorf BioPhotometer) and inoculated to 1.0 at wavelength 600 nm (OD₆₀₀nm=1.0). The inoculated suspension of each strain was prepared in 10 mL of 9% NaCl in a 50 mL centrifuge tube (Fisher Scientific, UK) and the cells harvested by centrifugation (3,000 rpm, 4°C, 5 min). Bacterial cell-pellets were then re-suspended in 500 µL of 0.2% CaCl₂: 2% sodium alginate hydrogel solution with a micropipette and dispensed into a 5 mL Luer-lock syringe (Fisher Scientific, UK). Connection to a further 5 mL Luer-lock syringe containing 4.5 mL 0.2% CaCl₂: 2% sodium alginate hydrogel warmed to 37°C allowed repeated, gentle mixing to be carried out back and forth between syringes containing bacteria and hydrogel (100 mixes back and forth), producing 5 mL bioink with homogeneously distributed bacteria.
Construct design

3D models consisting of a solid or lattice 10 mm x 10 mm square design with increasing vertical thicknesses (0.25 mm, 0.5 mm, 1 mm, 2 mm, 4 mm) were produced using Autodesk® Netfabb® software (Autodesk®, Inc, USA) and exported as an STL file. Open-source slicer software (Sli3er, Version 1.2.9) was used to load the STL files and generate G-code files using the following settings for bioprinting: layer thickness, 0.1 mm; infill pattern, rectilinear; infill density, 25%; speed, 10 mm/s; extrusion multiplier 1.2. G-code files corresponding to solid and lattice constructs with differing vertical thicknesses were then loaded onto the bioprinter.

Bioprinting

A three-axis (X-Y-Z), single nozzle 3D cell printer developed in our laboratory was used for bioprinting bioinks laden with different bacteria. This bioprinter represents an adapted, extrusion-based version of a previously developed microvalve-based bioprinter used in our lab to bioprint human cells including induced pluripotent stem cells [26, 39, 40]. Briefly, the bioprinter produces 3D constructs by coordinating the motion of a mechanically-driven syringe. The dispenser deposits extrudate consisting of hydrogel on a stationary Z-platform. As successive layers of extrudate are deposited, the z-platform moves downwards allowing structures to be bioprinted from the bottom up, layer-by-layer. Prior to use, the bioprinter was sterilized via UV exposure and wiped down with 70% ethanol. Sterility was maintained during bioprinting by placing the bioprinter in a laminar flow cabinet. Sterile 5 mL Luer-lock syringes containing bacterial bioink were attached to 25G printing nozzles and loaded into the bioprinter, allowing bioprinting into sterile 6-well culture plates to occur.

Secondary cross-linking of constructs

Ethylenediaminetetraacetic acid (EDTA), calcium chloride (CaCl₂) and barium chloride (BaCl₂) powders (Sigma-Aldrich, UK) were sterilised with ultraviolet (UV) light (three 30 min cycles). Solutions of 0.4% w/v CaCl₂, 10 mM BaCl₂, 20 mM BaCl₂, 40 mM BaCl₂ and 110 mM EDTA (Sigma-Aldrich, UK) were prepared in sterile deionised water. All solutions were then autoclaved at 121 °C for 30 minutes prior to experimental usage.

Following bioprinting, constructs were cross-linked by submersion in ionic solutions of either 10, 20 or 40 mM BaCl₂ for 2 mins. Cross-linked constructs were then rinsed in phosphate-buffered saline (PBS) prior to incubation in BHI medium under standard culture conditions (37 °C, 5% CO₂, and 95% relative humidity). BHI media was replenished every second or third day and culture was performed atop a compact fixed-angle platform rocker (Grant Bio™ PMR-30 Compact Fixed-Angle Platform Rocker, Fisher Scientific, UK), to increase flow of media around the bioprinted constructs.

Fluorescence staining for biofilm viability

A commercial Film Tracer™ LIVE/DEAD™ biofilm viability kit (Thermo Fisher) was used for the assessment of biofilm viability based on staining with the membrane potential sensitive dye propidium iodide (PI) (490 nm excitation, red emission) and the nucleic acid stain SYTO-9 (488
nm excitation, green emission). In principle, bacteria with intact cell membranes stain fluorescent green, whereas bacteria with damaged membranes stain fluorescent red. Cell viability staining of bacteria was carried out by incubating biofilm constructs concomitantly with SYTO-9 (6.7 µM) and PI (40 µM) in 35 mm glass bottomed imaging dishes (Ibidi) at room temperature (RT) for 45 min to allow stain penetration.

**Biofilm morphotype analysis**

In this study, a Leica Microsystems TCS SP8 CARS microscope utilising a 25x objective (HC FLUOTAR L 25x/0.95 W) was used for all confocal fluorescence imaging measurements. To minimise or eliminate artefacts associated with simultaneous dual wavelength excitation, all dual labelled biofilms were sequentially scanned, frame-by-frame, first at 488 nm (Argon laser, 70 µW) then at 561 nm (DPSS laser, 80 µW). Line averaging (x2) was used to capture images with reduced noise. Fluorescence emission was then sequentially collected in the green and red regions of the spectrum respectively. Images were captured in a two-dimensional (2D) projection. For analysing spatial separation in the z-direction (thickness), step sizes between 40-140 µm were used and 3D reconstructions were performed using Leica imaging software (LAS X). Five image stacks were (typically 700 x 700 µm images over a depth of 40 - 140 µm) were acquired randomly from three independent constructs per BaCl₂ concentration per time point (15 stacks in total). The image stacks were then analysed using MATLAB 2016A software.

**Antibiotic susceptibility testing (AST)**

For all AST methods, inocula of the isolate tested were prepared according the inoculum preparation protocol described above.

The methicillin stock solution of 20 mg/mL was prepared in sterile dH₂O and diluted in BHI broth to obtain solutions with preliminary concentration in a range of 2.5 to 10 mg/mL. Investigation of the response of 3D biofilm constructs to methicillin was then made by initially culturing porous, 1 mm constructs containing MRSA or MSSA for 14 days to allow biofilm maturation to occur. The matured biofilm constructs were then transferred to sterile Corning™ 6-well microtiter plates (Sigma-Aldrich, UK). A 3 mL volume of each methicillin solution was dispensed into each well of the plate. Fresh BHI broth was then added without antibiotic into the positive control wells. The plates were sealed with an anaerobic film (Thermo Fisher Scientific, UK) and incubated under anaerobic conditions at 37°C for 24 h.

**2D Broth microdilution method**

Corning 96-well microtiter plates (Sigma-Aldrich, UK) were used for determining the MICs of the antimicrobial agents methicillin sodium salt (Sigma-Aldrich, UK). A methicillin concentration in a range of 0.02 to 5 mg/mL were used. The MRSA and MSSA inoculum plural (OD1.0) were prepared as described above. A 50 µL volume of each methicillin solution and a 50 µL of inoculated suspension were dispensed into each well of the microtiter plates respectively. The 96-well plates were then sealed with an anaerobic film (Thermo Fisher Scientific, UK) and incubated under anaerobic conditions at 37°C for 24 hours. The optical
density of inoculated culture wells was then measured using a plate reader (Multiskan Go, Thermo Scientific). Subsequently, MICs were read as the lowest concentration of an antimicrobial agent at which visible growth was inhibited.

3D Broth macrodilution method

Methicillin stock solution of 20 mg/mL was prepared in sterile dH₂O and diluted in BHI broth to obtain solutions with preliminary concentrations ranging from 2.5 to 10 mg/mL. Investigation of the response of 3D biofilm constructs to methicillin was then performed by exposing a series of porous, 1 mm MRSA or MSSA constructs to increasing concentrations of methicillin. MRSA and MSSA constructs were cultured for 14 days prior to methicillin exposure to allow biofilm maturation to occur. Mature MRSA and MSSA biofilm constructs were then transferred into sterile Corning® 6-well plates (Sigma-Aldrich, UK) and incubated in 3 mL volumes of either 2.5, 5 or 10 mg/mL methicillin solution. Positive-control wells containing fresh BHI broth, no methicillin and MRSA or MSSA constructs were also set up. The 6-well plates were sealed with an anaerobic film (Thermo Fisher Scientific, UK) and incubated under anaerobic conditions at 37°C for 24 hours. The optical density of inoculated culture wells was again measured using a plate reader (Multiskan Go, Thermo Scientific).

Biofilm antimicrobial penetration test

3D bioprinted E.coli biofilm constructs of 1mm and 2mm thickness and porous design were cultured for 5 days to allow significant biofilm formation to occur. Biofilm constructs were then washed x3 with phosphate buffered saline (PBS) solution to remove non-adherent bacteria. Antibiotic disks containing 30 µg tetracycline (Oxoid, UK) were then placed on top of E.coli biofilm constructs and incubated at 37°C for 7 days within BHI broth. The tetracycline disks located on top of the biofilm constructs were replaced daily to maintain consistent delivery of antibiotic.
Results and discussions

Developing long-term stability of bioprinted alginate hydrogels to allow observation of 3D biofilm formation

The schematic presented below (Scheme 1) elucidates our general methodology of bacterial biofilm bioprinting using a biocompatible bioink [40, 41], extrusion bioprinting and a step-wise ionic crosslinking process. Cultured bacteria were mixed into a partially-crosslinked hydrogel to produce a bioink with homogenous bacterial concentration. A home-built bioextrusion based bioprinter was then used to extrude the bioink to produce constructs with predesigned dimensions. Following bioprinting, secondary ionic cross-linking of the hydrogel was performed to increase construct stability, allowing prolonged culture and observation (up to 28 days).

Scheme 1: Schematic of bacterial biofilm bioprinting process. Initial designs to be bioprinted were produced using computer-aided design (CAD) software. Following this, a partially cross-linked hydrogel was produced by mixing sodium alginate and calcium chloride (CaCl₂) together. Bacteria were then mixed into the hydrogel to produce a bioink with homogenously distributed bacteria. 3D bioprinting was then performed, using a custom-built bioprinter that uses mechanical force to extrude bioink from a syringe that is moved in the x-y-z plane. Bioprinted constructs of solid and porous design were then immersed in solutions of barium chloride (BaCl₂) for 2 mins to secondary cross-link the constructs. Following bioprinting and immersion cross-linking, the constructs were cultured in bacterial growth media, allowing analysis to be performed at selected time points.

The complex structure of 3D biofilms found in clinical infection take significantly longer to develop and mature than the simpler, 2D biofilm in vitro models which can be produced in overnight laboratory culture [3, 42]. Achieving sufficient stability in bioprinted bacterial construct was therefore essential to allow time for bacteria to associate, proliferate and deposit their own extracellular polymeric matrix to form a mature 3D biofilm structure. Alginate is a widely-adopted hydrogel for bioprinting and was chosen as the main component.
of our bacterial bioink due to its biocompatibility, low toxicity, low cost and ease of use [25, 43, 44].

In previous work we have developed the stability of alginate bioinks to allow the successful long-term 3D cell culture and differentiation of stem cells [25, 44]. This was achieved by double cross-linking alginate with calcium and then barium cations in a stepwise process [44]. We adapted this approach to produce double cross-linked bacterial bioink constructs with extended stability (>4 weeks) in culture. Other cations including strontium have been utilized elsewhere for this purpose; however, barium has been shown to give the strongest cross-linking effect, optimizing construct mechanical stability [45]. Initial cross-linking of sodium alginate hydrogel with calcium chloride created a hydrogel with sufficient viscosity to allow successful bioprinting of free-standing structures of both solid and porous design, ranging in thickness from 0.25 mm to 4 mm (figure 1a). By performing alginate hydrogel cross-linking prior to bioprinting, rather than extruding alginate onto a calcium-coated culture surface as performed in other literature, homogenous hydrogel cross-linking was achieved; this is essential to achieve good printability [28]. Further cross-linking occurred following bioprinting by exposure to solutions of barium chloride which further helped to maintain construct stability, extending the stability of constructs from within a week (with calcium-only cross-linking) to over 4 weeks in culture. (ESI, figure S1, figure S2). Bioprinting resolution with the hydrogel was sufficient to produce more intricate structures using a 32 g printing needle, corresponding to a 108 µm inner needle diameter (figure 1b).

Confocal laser scanning microscopy (CLSM) was used to observe 3D bioprinted biofilm formation. Standard light microscopes often struggle to image biofilm of more than 3-4 µm thickness as biofilm material above and below the focal plane tend to scatter light and interfere with direct measurement [46]. Contrastingly, CLSM allows optical sectioning of biofilms and, with image analysis, 3D reconstruction is possible [47].

The extended hydrogel stability after bioprinting allows observation of 3D biofilm formation for several weeks. Previous attempts reported elsewhere in the literature to 3D bioprint bacteria only demonstrated bacterial viability up to a maximum of 7-9 days, with no attempts made to perform antimicrobial testing on 3D bioprinted bacterial constructs [28, 30, 31]. The stability in culture of the bioprinted hydrogel-bacteria construct achieved in our study is therefore significant, as it allows for extended observation of bacterial growth as well as offering the potential to perform antimicrobial studies and further analysis of biofilm formation in 3D. Clinical biofilm infections are most often chronic in nature and develop over a period of weeks and even months; the stability of our bioprinted constructs may therefore facilitate greater potential to mirror clinical biofilms than currently available biofilm models [3, 7, 15, 48, 49].

**Investigating the influence of construct design and thickness on biofilm formation**
In order to mimic *in vivo* biofilms and to create an ideal *in vitro* 3D bioprinted biofilm model, solid and porous constructs were bioprinted in a range of thicknesses from 0.25 mm to 4 mm to investigate the ideal construct design and thickness for *E. coli* biofilm formation.

*E. coli* biofilm formation (or bacterial density) was greater in thinner (0.25 mm to 1 mm), constructs compared to thicker (4 mm) construct designs (*p* < 0.001, ANOVA) (figure 1c). However, thinner constructs of 0.25 mm and 0.5 mm thickness were not robust enough to allow physical manipulation and CLSM imaging to be performed after 14 days culture. This was presumed to be due to leaching of cations (Ca$^{2+}$ and Ba$^{2+}$) from the thin, relatively high-surface area constructs into surrounding culture media, resulting in decreased cross-linking; this is likely to have been exacerbated by regular media changes and culture atop a rocking device, increasing outwards diffusion of cations from the hydrogel-bacteria construct. In 4 mm thick constructs, reduced biofilm formation was observed in solid compared to porous constructs (*p* = 0.038, t-test) (figure 1c).

We believe the porous construct design facilitates convective fluid transport through the pore channels, enhancing nutrient and oxygen diffusion processes in comparison to non-porous, solid constructs. This would explain why the aerobic bacteria *E. coli* failed to proliferate and produce significant biofilm in the thick, solid constructs, with the optimal structure for *E. coli* being a 1 mm porous construct.
Figure 1. Thickness, structure and cross-linking of bioprinted constructs influences biofilm formation. (a) Solid and porous constructs with vertical thicknesses increasing from 0.25 mm to 4 mm were sequentially bioprinted and cross-linked by exposure to 20 mM BaCl2. Measured thickness correlated well with designed vertical thickness after measurement with digital callipers (ESI, Table 1). (b) Hydrogel printability was such that intricate structures could be printed with a 32G, 0.108 µm inner needle diameter needle. (c) 3D reconstructed CLSM z-stack images were acquired, allowing comparison of biofilm growth in solid and porous structures. Initial analysis at 5 days found that growth in solid constructs was slower than in corresponding porous constructs in all ranges of thicknesses. At day 14, 1 mm constructs appeared to have the greatest biofilm formation, whilst 0.5 mm and 0.25 mm constructs had insufficient mechanical stability to allow analysis. The sizes of the scale bars in the photograph and fluorescence images are 1 centimetre and 100 microns.

Bioprinting of thick, anaerobic 3D biofilm constructs

Whilst the aerobic bacteria *E. coli* had limited growth in thicker bioprinted constructs (figure 1c), presumably due to limited diffusion of nutrients and oxygen, anaerobic bacteria have greater potential to thrive in oxygen-deplete conditions. As an opportunistic, nosocomial pathogen of immunocompromised individuals, the anaerobic strain *Pseudomonas aeruginosa* (*P. aeruginosa*) is well known for infecting the thick, oxygen-depleted mucus in the airways of cystic fibrosis (CF) patients, producing robust *in vivo* biofilms [2]. The culture conditions provided by the thick respiratory mucus in CF patients is somewhat analogous to those provided by our thick, non-porous hydrogel constructs. To investigate this, *in vitro* biofilm formation of *P. aeruginosa* (figure 2) was examined in non-porous, thick (2 mm and 4 mm) constructs (figure 2).

Figure 2. *Pseudomonas aeruginosa* (PAO1) formed anaerobic biofilms in thick constructs. (a) Photo images of 3D bioprinted PAO1 biofilm at day 0 (white colour) and matured biofilm at day 14 (blue-green colour). (b) 3D reconstructed CLSM Z-stack in 2D-projection and 3D reconstructed images (1:1 aspect ratio in x, y & z axes) of matured PAO1 biofilm formed at 2
mm and 4 mm thickness at day 14. The sizes of the scale bars in the photograph and fluorescence images are 1 centimetre and 100 microns.

*P. aeruginosa* was observed to undergo extensive colonisation and aggregation in 2 mm and 4 mm thick, non-porous structures, forming an extremely dense layer of biofilm (figure 2b). In contrast, much more limited bacterial growth and biofilm formation was observed via CLSM in 2 mm and 4 mm constructs inoculated with the aerobic bacteria *E. coli* (figure 1c). Strong blue-green pigmentation was also seen to form in 2 mm and 4 mm *P. aeruginosa* constructs over 14 days of culture (figure 2a); this is likely related to the expression of two metabolites, pyocyanin (blue) and pyoverdine (green), which is known to occur in *P. aeruginosa* to facilitate anaerobic respiration [50]. The prevalence of multidrug-resistant (MDR) anaerobes, including *P. aeruginosa*, is increasing worldwide with limited current therapeutic options [51, 52]. The extensive growth of *P. aeruginosa* and associated biofilm formation seen within our 3D bioprinted constructs therefore offers a novel and highly promising in vitro method of studying anaerobic bacterial biofilm infection.

**Capturing the in vitro life cycle of biofilm in 3D**

Biofilm formation is reported to occur in a five-step lifecycle (figure 3a), which begins with the attachment of planktonic cells to a biological or inert surface and culminates in mature biofilm formation[53]. However, due to factors including limited biofilm thickness, current in vitro models are unable to readily facilitate observation of the five-step process and complex microarchitecture development that occurs during biofilm formation [54].

As illustrated in figure 3a, Initially, ① free swimming planktonic bacteria were attached on the surface, ② Soon after, bacteria began to divide and aggregate together in small microcolonies and secrete quorum signals ③, which initiated up-regulation of various genes and virulence factors on a community-wide basis. Bacteria cells forming an extracellular biofilm matrix ④ by secrete copious polymers including polysaccharides, proteins and oligonucleotides. Biofilm continues to accumulate and consuming ambient nutrient and QC acceptors. As results of increased in shear stress and other cell signalling events, portions of biofilm started detaching or slough off ⑤ entirely. Dispersed cells can quickly revert to their planktonic form to colonise other sites, whilst retaining properties such as AMR [54].

The influence of BaCl₂ cross-linking concentration on bacterial growth was also analysed over 28 days by exposing porous, 1 mm constructs containing MRSA to a range of BaCl₂ concentrations (ESI, figure S3). Growth within all constructs was initially strong; however, it was perceptible that bacteria had a greater tendency to leach from constructs exposed to 10 mM BaCl₂, with greater biofilm formation seen in 20 mM and 40 mM constructs (figure 3c). A custom designed image processing algorithm, implemented in MATLAB2016a, was used to apply further statistical analysis to quantify biofilm formation (ESI, figure S4, figure S5). It was found that 10 mM of BaCl₂ provided less favourable conditions for biofilm formation compared to 20 mM and 40 mM constructs between days 4 and 23 (p<0.001, ANOVA). This
was presumed due to reduced cation (Ba\(^{2+}\)) cross-linking density allowing greater leaching of bacteria.

Figure 3. 3D reconstructed confocal laser scanning microscopy (CLSM) Z-stacks of 3D bioprinted biofilm images (a) The 5-step process of biofilm formation in 2D correlated with (b) cross-sectional and side-on CLSM images of 3D bioprinted biofilm formation. (c) Growth of MRSA in 1 mm, porous scaffolds exposed to increasing concentration of BaCl\(_2\) from 10 mM to 40 mM was examined over a 28 day period. Schematic (a) adopted from V. E. Wagner et al [2]. The sizes of the scale bars in the photograph and fluorescence images are 1 centimetre and 100 microns.

CLSM studies demonstrated superior biofilm formation in 10 mM, 20 mM and 40 mM constructs, with significant biofilm formation evident after 5 days. Initially, (1) individual planktonic bacteria were homogenously distributed in bioink at day 0 (figure 3c, Day 0). Although some bacteria may have left the construct, a high density remained and likely adhered to the bioink scaffold using cell surface displayed adhesin molecules. (2) soon after, bacteria began to divide and aggregate together in small microcolonies (figure 3c, Day 1-2)
with in the construct, which merged into larger communities (figure 3c, Day 3-5); ③ progressive deposition of an EPS matrix also occurred, ④ leading to mature biofilm formation (figure 3c, Day 14). Eventually, ⑤ regions of biofilm were seen to spontaneously disperse between days 23 and 28 as bacteria enzymatically dissolved the extracellular matrix [55], weaken the biofilm structure and release microbial cells spread and leak out of the construct (figure 3c, Day 23-28) into surrounding culture media (where new biofilms can be formed). It is important to observe that 3D bioprinted alginate constructs remain largely intact while the bacteria escaped from constructs (ESI, Figure S2) after day 23. This further confirms that the lower microbial cell density observed from Day 23-28 was consistent with the final stage of the biofilm lifecycle where bacteria leak out of the biofilm and spread rather than the degradation of the 3D alginate constructs.

To the best of our knowledge, we have demonstrated for the first time the processes involved in mature 3D biofilm formation in vitro over a 28-day period using bioprinting (figure 3c). This allows direct correlation to the 5-step process governing biofilm formation in 2D to be made (figure 3a).

**Comparison of 2D vs 3D in vitro antimicrobial susceptibility testing (AST)**

To compare the susceptibility of 2D and 3D bacterial cultures to treatment, we utilised 3D bioprinted biofilms as an in vitro model with comparison made to 2D bacterial cultures. *Staphylococcus aureus* (*S. aureus*) was chosen for investigation as a major human pathogen[56]. Although most commonly associated with skin and soft tissue infections, *S. aureus* is also responsible for a range of serious invasive infections, including osteomyelitis, necrotising pneumonia, endocarditis and bacteraemia [56]. Infections caused by *S. aureus* are increasing worldwide, with over 52% of intensive care unit (ICU) infections reported to be caused by *S. aureus* [57]. Most strains of *S. aureus*, including methicillin-susceptible *S. aureus* (MSSA), are sensitive to β-lactam antibiotics and are responsive to treatment. However, there is a growing worldwide prevalence of methicillin resistant *S. aureus* (MRSA) infections, which have repeatedly been associated with a worse patient outcome compared to infections caused by methicillin sensitive *S. aureus* (MSSA) [58]. Furthermore, the efficacy of first-line treatments for MRSA such as vancomycin is dwindling [59]. Antibiotic resistance studies are therefore essential to allow the development of novel anti-biofilm therapies against MRSA and MSSA biofilms.

The broth microdilution method was used to determine the lowest concentration (MIC) of methicillin antibiotic that prevented visible growth of MRSA and MSSA in 2D culture (figure 4a). The broth macrodilution method was then used to determine the minimal biofilm eradicating concentration (MBEC) in 3D bioprinted MRSA and MSSA biofilm culture models (figure 4c). The MIC and MBEC were determined by a visual inspection of culture wells and correlated with measurements of absorbance of light through treated culture wells in both cases (figure 4b & 4d). Due to resistance to methicillin, MRSA had a higher MIC than MSSA in 2D (figure 4a) and a higher MBEC than MSSA in 3D culture as expected (figure 4c). However, for both MRSA and MSSA, the MBEC calculated in 3D culture was significantly higher than the
MIC for 2D culture. Whilst 0.16 μg/mL methicillin prevented visible growth of 2D MSSA culture, the MBEC for MSSA in 3D culture appeared to be at least 15 times higher at 2.5 mg/ml. Similarly, although 1.25 μg/mL methicillin appeared to prevent 2D growth of MRSA, growth of MRSA in 3D culture still occurred with greater than 10 mg/ml methicillin. Therefore, for both MRSA and MSSA, a far higher dose of methicillin was required to treat biofilm growth than was required to treat 2D infection. This result is in keeping with previous reports suggesting that biofilm formation can cause a 10 to 1,000-fold increase in bacterial tolerance to antimicrobial treatment compared to 2D, planktonic cultures [33, 55].

![Graph showing MIC and MBEC for MSSA and MRSA in 2D and 3D cultures.](image)

**Figure 4. In vitro antimicrobial susceptibility testing (AST).** (a) The MICs were determined by broth microdilution methods. An MIC of methicillin of 0.16 μg/mL was required to prevent visible growth of MSSA, whilst for MRSA the MIC of methicillin was 1.25 μg/mL (figure 4a). (b) Optical density measurement of the methicillin-containing culture. No significant change in absorbance was observed when methicillin concentrations were increased beyond the MIC calculated for MRSA or MSSA in 2D. (c) The MBECs were determined by broth macrodilution method. MBECs appeared to be at least 2.5 mg/mL for MSSA, and greater than 10 mg/mL for MRSA on inspection. (d) Measurement of the light absorbance of the culture broth surrounding the MRSA and MSSA constructs supported these findings, with far higher doses of methicillin required to reduce bacteria growth and therefore the measured broth light absorbance than in 2D cultures.

**Biofilm thickness influences response to treatment**
AST methods such as MIC calculation do not distinguish between bactericidal and bacteriostatic effects of antibiotics, and crucially do not provide information on the degree of antimicrobial biofilm penetration or eradication [1, 4, 50, 60-64]. Utilising 3D bioprinted biofilms as an in vitro model, we sought to investigate the relationship between bacterial biofilm thickness and susceptibility to antimicrobial treatment. Sensitivity of E. coli to tetracycline was first confirmed in 2D culture (ESI, figure S6). Bioprinted E. coli constructs of 1 and 2 mm thickness were then grown for 5 days to allow biofilm maturation, before exposure to 30 µg tetracycline discs which were changed every 24 h for seven days, mimicking a course of clinical antimicrobial treatment (figure 5a). It was apparent that 2 mm constructs remained opaque whilst 1 mm constructs became increasingly transparent in response to tetracycline exposure (figure 5a). CLSM imaging of the constructs after 7 days of tetracycline exposure demonstrated that E. coli biofilms had greater viability in 2 mm constructs, whilst bacteria located below the tetracycline disc in 1 mm constructs had largely been destroyed (figure 5b).

As discussed previously, current methods of studying antimicrobial biofilm penetration and eradication suffer significant limitations. However, 3D bioprinted biofilms could offer hope for a novel and reproducible method of studying antimicrobial biofilm penetration and eradication in 3D. In the clinical environment 3D bioprinted biofilms could feasibly be generated from bacterial samples taken from patients in a similar manner to our experiment; this would allow antimicrobials to be selected on the basis of their ability to achieve biofilm penetration and eradication in patient-specific infections. Furthermore, it is recognised that 3D cultures (such as our 3D bioprinted biofilm) more closely resemble the in vivo biofilm, when compared to traditionally used 2D in vitro cultures [15, 64-66].
Figure 5. Biofilm thickness determines response to treatment (a) 1 mm and 2 mm thick constructs containing *E.coli* were bioprinted and allowed to mature for 14 days before 30 µg tetracycline discs were placed directly on top of them. Discs were changed every 24 h to maintain a high dose of tetracycline delivery to the constructs. Over a 7 days period, visible clearing of biofilm occurred within the 1 mm construct below the area of tetracycline exposure. (b) CLSM Z-stack images of the 1 mm and 2 mm constructs was performed after exposure to tetracycline discs. Whilst the majority of bacteria were found to be dead below the area of tetracycline disc exposure in the 1 mm construct, greater evidence of biofilm survival in the 2 mm construct was observed. The sizes of the scale bars in the photograph are 1 centimetre.

Conclusions

In conclusion, mature bacterial biofilm constructs were reproducibly 3D bioprinted for the first time using clinically relevant bacteria. By deploying a methodology originally developed to enable 3D culture and differentiation of bioprinted stem cells [25], we have been able to demonstrate for the first time 3D bioprinted mature biofilm formation, dispersal and morphology over 28 days, as well as the antibiotic tolerance of clinically relevant bacterial biofilms in 3D. Our methodology also significantly prolongs the viability of bacteria cultured in 3D bioprinted constructs compared to previous studies. Future ability to investigate clinically relevant bacterial biofilms in a biocompatible, cost-effective 3D model that more closely resembles *in vivo* conditions than traditional 2D culture methods is therefore increased.
A high degree of control was achieved over biofilm construct thickness and design, with the production of biofilms thicker (>4 mm) than currently available in vitro models also achieved. We observed that anaerobic bacteria continued to thrive in constructs of greater than 4 mm thickness, demonstrating the potency of these infections. To our best knowledge, the 4 mm thick aerobic bacteria biofilm formation is the thickest 3D bioprinted in-vitro biofilm construct ever reported, allowing easy observation of antimicrobial biofilm penetration.

We observed that 3D biofilm constructs had greater resistance to antimicrobial treatment than 2D cultures, underlining the significance of biofilm formation in clinical infection. Thicker biofilms were also seen to have greater resistance to antimicrobial therapy than thinner biofilms, even over a prolonged period of treatment.

With rising worldwide antimicrobial resistance, 3D bioprinted biofilm technology could become a key weapon to aid the discovery of novel therapeutic targets and increase the understanding of biofilm formation.

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**References**


Electronic Supplementary Information

3D Bioprinting of Mature Bacterial Biofilms for Antimicrobial Resistance Drug Testing

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Determination of bacterial viability in 3D bioprinted constructs

Three-dimensional (3D) bioprinted biofilms were dissolved in EDTA to be able to enumerate and characterise them. Briefly, non-adherent bacteria were removed before by washing the construct with PBS then transferring to a new well plate. The constructs were dissolved in 1 ml EDTA (110 mM) for 1 h then thoroughly mixed before appropriate dilutions in 9% sodium chloride (NaCl) and plated on Brain Heart Infusion (BHI) agar plates and incubated at 37°C overnight to determine viable colony forming unites per millilitre (CFU/mL).

![Figure S1. Enumerate of 3D bioprinted biofilm growth in increasing barium chloride (BaCl2) concentration over the period of 5 days.](image)

(a) Photo images of construct after dissolved in EDTA solution. (b) CFU count in the construct dissolved in EDTA solution.
Long-term stability of printed alginate hydrogels for 3D biofilms

Cross-linking a sodium alginate hydrogel with calcium chloride created a hydrogel with viscosity sufficient to allow bioprinting of free-standing structures ranging in size and thickness. Further cross-linking following bioprinting by exposure to solutions of barium chloride helped to maintain individual construct stability further, which extended the stability of the hydrogel from within a week to over 4 weeks.

Figure S2. Photo images of bacterial constructs over a period of 4 weeks. Growth of MRSA in 1 mm, porous scaffolds exposed to increasing concentration of BaCl₂ from 10 mM, 20 mM and 40 mM. The sizes of the scale bars in the photograph are 1 centimetre.
Reproducible bioprinting of 3D bacterial biofilms with controlled dimensions

Successful bioprinting relies in part on combining a suitable bioprinting technique with an appropriate bioink. To achieve this, a bacteria-friendly hydrogel was developed with reliable bioprinting characteristics. Cross-linking a sodium alginate hydrogel with calcium chloride created a hydrogel with viscosity sufficient to allow bioprinting of free-standing structures ranging in size and thickness. Further cross-linking following bioprinting by exposure to solutions of barium chloride (BaCl₂) helped to maintain individual construct stability further. This approach allowed a range of construct thicknesses from 0.25 mm to 4 mm to be successfully bioprinted.

Table S1. Printing parameters before and after BaCl₂ crosslinking. Solid and porous constructs with vertical thicknesses increasing from 0.25 mm to 4 mm were sequentially bioprinted and cross-linked by exposure to 20 mM BaCl₂. Measured scaffold thickness following bioprinting correlated well with designed thickness prior to and after barium chloride crosslinking.

<table>
<thead>
<tr>
<th>Solid Construct</th>
<th>CAD thickness</th>
<th>0.25 mm</th>
<th>0.5 mm</th>
<th>1 mm</th>
<th>2 mm</th>
<th>4 mm</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before BaCl₂ Crosslinking thickness (mm)</td>
<td>0.33 ± 0.06</td>
<td>0.63 ± 0.09</td>
<td>1.1 ± 0.06</td>
<td>2.2 ± 0.19</td>
<td>4.2 ± 0.29</td>
<td></td>
</tr>
<tr>
<td>After BaCl₂ Crosslinking thickness (mm)</td>
<td>0.27 ± 0.11</td>
<td>0.57 ± 0.08</td>
<td>1.2 ± 0.24</td>
<td>2.2 ± 0.29</td>
<td>4.1 ± 0.27</td>
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</tr>
</tbody>
</table>

<table>
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<tr>
<th>Porous Construct</th>
<th>CAD thickness</th>
<th>0.25 mm</th>
<th>0.5 mm</th>
<th>1 mm</th>
<th>2 mm</th>
<th>4 mm</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before BaCl₂ Crosslinking thickness (mm)</td>
<td>0.29 ± 0.06</td>
<td>0.51 ± 0.08</td>
<td>1.0 ± 0.13</td>
<td>2.2 ± 0.24</td>
<td>4.1 ± 0.31</td>
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</tr>
<tr>
<td>After BaCl₂ Crosslinking thickness (mm)</td>
<td>0.28 ± 0.09</td>
<td>0.56 ± 0.08</td>
<td>1.1 ± 0.12</td>
<td>2.0 ± 0.23</td>
<td>4.2 ± 0.19</td>
<td></td>
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</tbody>
</table>
Fluorescence staining for Biofilm viability

Biofilm viability staining unlisted a commercial Film Tracer™ LIVE/DEAD™ biofilm viability kit (Thermo Fisher) for the assessment of viability is based on the staining with the membrane potential sensitive dye propidium iodide (PI) (490 nm excitation, red emission) and the nucleic acid stain SYTO-9 (488 nm excitation, green emission). Confocal laser Scanning Microscopy (CLSM) was used for enumeration and morphological observation of 3D bioprinted biofilm formation. A magnification 25x water immersion objective was used in all imaging experiments. Images were captured in two-dimensional (2D) projection.

Figure S3. Confocal laser scanning microscopy (CLSM) images of 3D bioprinted biofilms over a period of 28 days. Growth of MRSA in 1 mm, porous scaffolds exposed to increasing concentration of BaCl₂ from 10 mM to 40 mM.
**Image processing algorithm and statistical analysis**

The data flow for the algorithm to analyse the 3D reconstruction of the undisturbed biofilm samples was carefully considered to make sure that the analysis was robust with respect to the cellular physiology and the uncertainty of the signal to noise ratio due to environmental and microscope conditions. The algorithm has a main function using unsupervised learning to cluster the data by their statistical properties, and it was chosen to apply $k$-means clustering to deploy automated segmentation. There were four benefits: single parameter, no pre-processing, fully automated and high accuracy when comparing to conventional thresholding method, which was not robust on non-uniform or non-flattened images.

The algorithm was developed using MATLAB 2016a software with its built-in statistics and machine learning, and image processing toolboxes. A fixed threshold value and connected volume filtration was used for all image stacks. Five image stacks were taken randomly from three independent constructs per BaCl$_2$ concentration per time point (15 stacks in total). Firstly, the 2D-projection fluorescent images were imported to the software to train the basic algorithm by extracting the green channel. An unsupervised $k$-means segmentation method was applied to the channel, where $k$ was set to be 3, which was a suitable value to separate the cellular and background signals. This was gained for the technical experience due to uncertainty of signal to noise levels. The calculations were on average completed with 3 – 5 seconds from a i7 CPU computer. Once the segments were identified, they were then correlated to the fluorescent image to identify the corresponding cellular structures. Thirdly, the cells were counted and analysed using the labelling connected component method to identify individual microcolony groups. With this method, it was possible to isolate groups and when to have no pixelated connection. Finally, the mean and standard deviation of the numbers of groups, group areas, and area variations were calculated for statistical purposes.
### K-Mean Segmentation

<table>
<thead>
<tr>
<th>Day</th>
<th>Total number of microcolonies</th>
<th>Total area of microcolonies</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 0 - 40nM</td>
<td>2114</td>
<td>80450</td>
</tr>
<tr>
<td>Day 1 - 40nM</td>
<td>548</td>
<td>246480</td>
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<tr>
<td>Day 3 - 40nM</td>
<td>344</td>
<td>280219</td>
</tr>
<tr>
<td>Day 5 - 40nM</td>
<td>130</td>
<td>451139</td>
</tr>
<tr>
<td>Day 14 - 40nM</td>
<td>0</td>
<td>490000</td>
</tr>
<tr>
<td>Day 28 - 40nM</td>
<td>1111</td>
<td>98855</td>
</tr>
</tbody>
</table>

Figure S4. Analysis of CLSM z-stacks and adaptive segmented images with calculated total number and total area of microcolonies.

Figure S5. Quantification of biofilm formation over time was performed using unsupervised k-means segmentation and a labelling connected component function on MATLAB to threshold areas of biofilm formation out from background noise on images. The resultant graph of biofilm area produce over time demonstrates that 20 mM and 40 mM crosslinking provided superior conditions for biofilm formation and with 10 mM found to provide less favourable conditions for biofilm formation as compared to 20 mM and 40 mM constructs.
**Antibiotic selection - Disk diffusion test**

The BHI agar plates were inoculated with *E.coli* suspension as described in the material and methods session (inocula preparation). Briefly, 100 µL of OD 1.0 inoculated suspension was placed on the centre of the BHI agar plate and spread evenly over the surface by a L-loop. Antibiotic disks containing 30 µg tetracycline (Oxoid), 1U and 10U penicillin (Oxoid) were placed on the plates. The agar plates containing the bacteria inoculum and antibiotics disks are further incubated at 37°C for 24 h. During the incubation, the antibiotics diffuse into the BHI agar with the antibiotic concentration decreasing with increasing distance from the disk. The microbiological determination of the inhibition zone sizes is shown in (figure S6), evident inhibitory effect resulting from the application of 30 µg tetracycline disk (figure S6A).

![Disk diffusion test](image)

**Figure S6. Disk diffusion test for determine antibiotic susceptibility from the discovery of antibiotics.** Disk diffusion assays involve placing multiple antibiotic-impregnated disks onto the BHI agar surface inoculated with *E.coli* and measuring the diameter of zones of inhibition to qualitatively determine antibiotic susceptibility. (A) Photograph showing lack of *E.coli* colonies in the vicinity of 30 µg tetracycline, which is considered to be susceptible to the antibiotic. The bacteria grow within the predetermined zone width (B) 1U penicillin and (C) 10U penicillin, which is considered resistant to the antibiotic.