

Cite this: *Soft Matter*, 2014, 10, 7595

# Synthesis of colloidal microgels using oxygen-controlled flow lithography†

Harry Z. An,<sup>a</sup> H. Burak Eral,<sup>a</sup> Lynna Chen,<sup>b</sup> Michelle B. Chen<sup>c</sup> and Patrick S. Doyle<sup>\*a</sup>

We report a synthesis approach based on stop-flow lithography (SFL) for fabricating colloidal microparticles with any arbitrary 2D-extruded shape. By modulating the degree of oxygen inhibition during synthesis, we achieved previously unattainable particle sizes. Brownian diffusion of colloidal discs in bulk suggests the out-of-plane dimension can be as small as 0.8  $\mu\text{m}$ , which agrees with confocal microscopy measurements. We measured the hindered diffusion of microdiscs near a solid surface and compared our results to theoretical predictions. These colloidal particles can also flow through physiological microvascular networks formed by endothelial cells undergoing vasculogenesis under minimal hydrostatic pressure ( $\sim 5$  mm H<sub>2</sub>O). This versatile platform creates future opportunities for on-chip parametric studies of particle geometry effects on particle passage properties, distribution and cellular interactions.

Received 26th June 2014

Accepted 23rd July 2014

DOI: 10.1039/c4sm01400f

www.rsc.org/softmatter

## 1. Introduction

Controlled synthesis of non-spherical microparticles is an important problem in biomaterial design and drug delivery. Deviation from simple spherical symmetry in particle architecture dramatically improves the utility of these particles in a wide spectrum of biomedical applications, including multiplexed sensing and diagnostics of clinically relevant disease markers,<sup>1,2</sup> mimicry of naturally occurring biological entities (such as red blood cells<sup>3–7</sup> or platelets<sup>8,9</sup>), programmed encapsulation of drugs and imaging probes,<sup>10,11</sup> followed by their systemic or targeted delivery in due course.

In general, performance of designer particulate systems is the product of a complex interplay of material characteristics. This is especially true for microfabricated drug delivery vectors: modulating the size,<sup>12,13</sup> shape,<sup>14</sup> elasticity, and surface chemistry of the particles has been shown to delay internalization by immune cells,<sup>12</sup> influence intracellular trafficking,<sup>15,16</sup> and prolong *in vivo* circulation.<sup>5,17</sup>

In order to elucidate a complete set of design criteria for maximizing selectivity at the cell and tissue levels, synthesis methods need to afford independent control over the physicochemical properties of the resulting particles. A variety of versatile top-down particle fabrication techniques have emerged in the past few decades, chief among them

microfluidics,<sup>18</sup> lithography,<sup>19</sup> electrospraying,<sup>20</sup> template stretching,<sup>21,22</sup> or combinations thereof.<sup>23,24</sup> However, not all techniques can freely access the material composition, particle architecture, or length scale ( $\leq 10$   $\mu\text{m}$ ) relevant to biological assays, especially cellular uptake, and intravenous injections. For example, particles synthesized using droplet microfluidics are often restricted to spheres or deformations thereof due to surface energy minimization, with characteristic size  $\sim O$  (10  $\mu\text{m}$ ). Electrohydrodynamic jetting can be used to create many unique non-equilibrium particle structures. However, its ability to do so hinges heavily on prepolymer flow characteristics, which varies according to particle composition. Up to now, techniques for generating geometrically-complex, and structurally-anisotropic microparticles with size in the colloidal regime remain elusive.

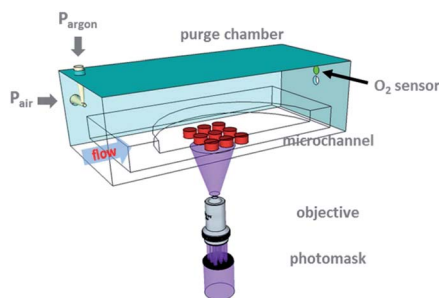
To ameliorate the experimental bottleneck on biocompatible colloidal microgel synthesis, we developed a modified flow lithography technique. This versatile method builds upon stop-flow lithography<sup>23</sup> (SFL), which allows microparticles of any 2D-extruded, mask-defined shape to be patterned from UV-cross-linkable oligomer formulations in a semi-continuous fashion. During normal operation, oxygen, a polymerization inhibitor, rapidly diffuses through the PDMS sidewalls, and gives rise to lubricating layers of thickness  $\delta$  above and below the particle. However, this fortuitous ability to easily create free-standing structures leads to other problems:  $\delta$  limits the feature resolution of the technique to  $\sim 2$   $\mu\text{m}$  in the out-of-plane direction.<sup>4</sup> Furthermore, for particle synthesis using conventional SFL, the minimum in-plane feature resolution depends on the channel height. In particular, 5  $\mu\text{m}$  diameter cylinders can only be synthesized using devices that are at least 10  $\mu\text{m}$  tall.<sup>23</sup> In this contribution, we control the inhibitory effects of ambient oxygen<sup>25</sup> *via* simple purge (Fig. 1). The improved experimental

<sup>a</sup>Department of Chemical Engineering, Massachusetts Institute of Technology, Cambridge, MA, 02139, USA. E-mail: pdoyle@mit.edu

<sup>b</sup>Department of Biological Engineering, Massachusetts Institute of Technology, Cambridge, MA, 02139, USA

<sup>c</sup>Department of Mechanical Engineering, Massachusetts Institute of Technology, Cambridge, MA, 02139, USA

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c4sm01400f



**Fig. 1** Experimental setup. Schematic drawing showing the fabrication of an array of colloidal discs ( $h \sim 1 \mu\text{m}$ ,  $2r < 10 \mu\text{m}$ ) under reduced oxygen atmosphere. A PEG-based prepolymer is flowed through a tiered PDMS synthesis device and polymerized via controlled pulses of mask-defined UV light through a microscope objective. The gel particles are then advected downstream for collection. The synthesis device is placed in a custom-made purge chamber, which allows the exchange of an air/argon mixture to control the overall ambient oxygen concentration. A RedEye™ oxygen sensor patch is mounted above the gas outlet to measure the oxygen concentration.

setup extends the particle synthesis capabilities of SFL to  $\sim 5 \mu\text{m}$  in diameter (in-plane resolution) and sub-micron in height, simultaneously.

In this article, we substantiate the colloidal nature of the microparticles fabricated using oxygen-controlled SFL by examining their diffusion in bulk and near an interface. In the latter case, we constructed a simple argument based on Boltzmann statistics for estimating the hindered diffusion coefficient near a planar wall.

In addition, we perform proof-of-concept experiments showing the flow of poly(ethylene glycol) (PEG)-based hydrogel colloids through multi-culture microfluidic vascular networks ( $\mu\text{VNs}$ ) formed by human umbilical vein endothelial cells undergoing vasculogenesis.<sup>26</sup> These self-organized, perfusable 3D  $\mu\text{VNs}$  were recently used to better understand extravasation events in tumor cell metastasis.<sup>27</sup> The *in vitro* microvascular network platform is a very useful tool for systematically studying transport of microparticles in real time, with increased physiological relevance compared to microfluidics-based flow assays (similar to other bio-inspired experiments<sup>4,28</sup>).

## 2. Experimental

### 2.1 Materials

Solutions of poly(ethylene glycol) diacrylate (PEGDA,  $M_n = 700 \text{ g mol}^{-1}$ , Sigma Aldrich) and 2-hydroxy-2-methyl-1-phenylpropan-1-one (Darocur® 1173, Sigma Aldrich) were used for particle synthesis. Methacryloxyethyl thiocarbamoyl rhodamine B ( $\lambda_{\text{ex}}/\lambda_{\text{em}} = 548/570 \text{ nm}$ , Polysciences) was used to label the gel particles for fluorescence imaging.

### 2.2 Particle synthesis

The microfluidic devices were prepared by pouring polydimethylsiloxane (PDMS, Sylgard 184, Dow Corning) onto a 4 inch silicon wafer containing positive-relief SU-8 features. Procedures for device fabrication and assembly are discussed in

detail elsewhere.<sup>29</sup> The devices contain parallel rectangular channels  $100 \mu\text{m}$  in width and  $2 \mu\text{m}$  in height. Thickness over the synthesis section of each device was tuned between 1 to 3 mm by varying the amount of PDMS used. For synthesis experiments done in this work, a second layer of PDMS was poured near the inlet and outlet reservoirs to support modified pipette tips<sup>30</sup> (used to inject prepolymer mixtures) and blunt needles (used to collect particles).

All of the particles shown in this work were prepared using a modified stop-flow lithography (SFL) technique. The synthesis devices were placed inside a custom-made sealed chamber with multiple purge gas inlet and outlet apertures, and mounted on an inverted microscope (Axiovert 200, Zeiss). A prepolymer mixture consisting of 80% PEGDA, 15% photoinitiator and 5% fluorescent dye stock solution ( $1 \text{ mg mL}^{-1}$  PEG200), by volume, was injected into a synthesis device using modified pipette tips (Molecular BioProducts) under moderate forcing pressure ( $\sim 4 \text{ psi}$ ). Concurrently, air and argon purge streams (DPG4000-30, Omega) were introduced into the chamber according to a pre-determined calibration curve to achieve a targeted oxygen concentration, between 0% and 21%. The enclosed microfluidic device and prepolymer mixture were allowed to equilibrate with the exchanged gas mixture for up to 90 minutes, depending on the device thickness.

Colloidal microparticles were patterned by projecting mask-defined UV light from a Lumen 200 metal arc lamp (Prior Scientific, 100% setting,  $1250 \text{ mW cm}^{-2}$ ) through a wide excitation UV filter set (300–380 nm, 11000v2 UV, Chroma Technology) using a  $20\times$  objective. A shutter system (VS25, Uniblitz) interfaced with a custom-written Python automation script precisely controlled the duration of UV exposure (up to 250 ms). Microparticle formation was visualized using a cooled interline charge-coupled device camera (Clara, Andor).

Following polymerization, particles were transferred from the outlet reservoir of the microfluidic synthesis device to a clean 1.5 mL Eppendorf tube containing 750  $\mu\text{L}$  PBST ( $1\times$  phosphate buffered saline with 0.1% (v/v) Tween® 20). Particles were washed and re-suspended five times in PBST then stored at  $4^\circ\text{C}$  until further characterization.

### 2.3 Particle characterization

**Particle diffusivity measurements.** We used multiple particle tracking to measure the diffusion coefficient of colloidal discs suspended in aqueous buffer with 0% or 90% deuterium oxide, by volume. Particle samples ( $\sim 30 \mu\text{L}$ ) were loaded into observation chambers that consist of two parallel strips of parafilm sandwiched between a glass slide ( $24 \text{ mm} \times 60 \text{ mm}$ , no. 1.5, VWR) and a coverslip ( $18 \text{ mm} \times 18 \text{ mm}$ , VWR). Prior to assembly, the slide and coverslip were immersed in a base bath ( $1 \text{ M NaOH}$ ), rinsed with DI water and ethanol, and dried with compressed argon. The chamber was sealed on all sides by a thin layer of ultraviolet-cured optical glue (#65, Norland) to minimize drift and prevent evaporation.

The slides were transferred to an inverted microscope (Axiovert 200, Zeiss) equipped with an EC Plan Neofluar® objective ( $40\times$ ,  $\text{NA} = 0.75$ ) and observed under fluorescence mode.

Movies of up to 2500 frames were recorded using a cooled interline CCD camera (Clara, Andor) at a rate of 13.6 fps with an exposure time of 20 ms to minimize dynamic error. We used a public-domain MATLAB algorithm written by Kilfoil and coworkers<sup>31</sup> to extract two-dimensional particle trajectories from the movies. To extract particle mobility, trajectories were averaged to obtain the unbiased mean-squared displacement (MSD),  $\langle \Delta R(\tau) \rangle$  as a function of lag time,  $\tau$ . The static error ( $\xi^2$ ) due to camera noise was deduced from MSDs of immobilized discs at short lag time<sup>32</sup> to be  $10^{-4} \mu\text{m}^2$ .

**Confocal microscopy.** Colloidal microparticles were immobilized in a viscous solution (2% alginate,  $\eta/\eta_{\text{water}} = 18$ )<sup>33</sup> and loaded into a thin observation chamber. A confocal laser scanning microscope (LSM 700, Zeiss) equipped with a high numerical aperture, oil immersion objective (NA = 1.3) was used to best resolve the out-of-plane dimension. Successive x-y scans were acquired under fluorescence ( $\lambda = 568 \text{ nm}$ ) at intervals of 60 nm through representative particles. The images were processed using the 3D viewer feature in ImageJ (NIH).

## 2.4 Determination of oxygen concentration in purge chamber

The gas-phase oxygen concentration inside the purge chamber was monitored non-invasively using the RedEye<sup>TM</sup> oxygen sensing patches (FOXY, Ocean Optics) placed directly above the purge gas outlet (Fig. 1). The patches contain immobilized indicator dyes that quench in the presence of oxygen molecules. The degree of quenching correlates quantitatively with the partial pressure of oxygen ( $\text{pO}_2$ ) according to Stern–Volmer equation,  $t_0/t = 1 + K_{\text{SV}}\text{pO}_2$ , where  $t_0$  and  $t$  is the fluorescence lifetime in the absence and presence of oxygen, respectively,  $K_{\text{SV}}$  is the Stern–Volmer coefficient. Operational details of the oxygen sensing system were discussed previously.<sup>3</sup>

Ambient oxygen concentration was varied from 0% to 20.9% by adjusting the relative flow rates of argon and air into the chamber, and was measured from a linear calibration curve. A 2-point calibration was performed prior to each sensing experiment. Typically, fluorescence lifetime decreased from 2.848  $\mu\text{s}$  for 0% oxygen to 1.990  $\mu\text{s}$  for 20.9% oxygen.

## 2.5 Numerical solutions of mass transport equations

The simulation results shown in Fig. 3 were obtained by numerically integrating the conservation equations for oxygen and unconverted oligomers using a custom-written script in Fortran 77. Eqn (6) and (7) were discretized into 500 uniform mesh elements along the channel height ( $z$ ) direction between 0 (channel bottom) and 1 (channel top), and propagated in dimensionless time intervals of  $10^{-6}$  for  $\geq 10^4$  steps.

## 2.6 Particle flow experiments

The vasculogenesis flow devices (height = 100  $\mu\text{m}$ ) were molded in PDMS and permanently bonded to glass slides *via* oxygen plasma treatment. Device contains three parallel, spatially segregated gel regions. Each gel region is flanked by medium channels to provide adequate nutrients and gas exchange, and direct access to the openings of the vasculature. Human

umbilical vein endothelial cells (HUVECs) were encapsulated in fibrin gels and cultured alongside human lung fibroblasts (HLFs). Two columns of equally spaced trapezoidal PDMS posts, one on either side of the central HUVEC gel region, were introduced to (1) enable uniform gel-fluidic interfaces, and (2) minimize spillage into medium channels during gel filling. HLFs prevent nascent vascular networks from regressing after  $\sim 4$  days under non-contact co-culture conditions. For detailed procedures for device assembly, cell culture, and gel filling, see ref. 26.

To introduce colloidal particles into the microvascular networks, media from the reservoirs of the two channels flanking the endothelial cell channel was aspirated and 40  $\mu\text{L}$  of the particle suspension containing  $\sim 300\,000$  particles per mL was deposited into one of the reservoirs connected to the central media channel. This creates a pressure drop of 5.2  $\text{mmH}_2\text{O}$  across the vascular network, drawing the particles into the vasculature.

Videos were recorded, under both fluorescence and DIC conditions simultaneously on an Olympus confocal microscope (10 $\times$  objective, 2.4 $\times$  optical zoom) at a frame rate of 0.33 fps. Composite images were created using ImageJ showing the outlines of the microvascular network and the particle trajectory.

After the flow experiment, vessels were fixed with 4% paraformaldehyde (Electron Microscopy Sciences) for 15 min, permeabilized with 0.01% Triton X (Sigma) and stained for actin using Alexa Fluor<sup>®</sup> 488 Phalloidin (Invitrogen) at 1 : 200 dilution. Fluorescent confocal images (Olympus IX81) were then taken and processed using the IMARIS imaging software (Bitplane).

# 3. Results and discussion

## 3.1 Modified stop-flow lithography

To synthesize sub-micron tall particles, we build upon the stop-flow lithography (SFL) technique introduced by Doyle and co-workers.<sup>23</sup> In conventional SFL, pulses of mask-defined UV light are projected through a microscope objective to generate free-standing gel structures within a microfluidic device. The particles of height  $h$  are flanked above and below by lubricating layers of thickness,  $\delta$ , which is normally  $\sim 0$  (1  $\mu\text{m}$ ). The boundary of the inhibition layer is determined by the delicate balance between oxygen diffusion and consumption *via* the inhibition reaction with propagating oligomers. Therefore, in order to fabricate microgels with  $h \sim \delta$ , ways of controlling ambient oxygen concentration during synthesis needed to be explored. We note that the use of inert sheath flows is insufficient as the minimum gap height achievable is  $\sim 2 \mu\text{m}$  (ref. 34) for any reasonable polymerization time due to diffusive mixing at the fluid interfaces.

One facile method of exercising oxygen control is through the use of a purge chamber. A schematic of the modified flow lithography experimental setup is shown in Fig. 1. The chamber is made in-house and latches onto the microscope stage. During particle synthesis, oxygen concentration inside the chamber can be altered by passing air and argon (which contains 20.9% and

0% O<sub>2</sub> by volume, respectively) in the desired proportions through the two purge gas inlet apertures. Placing the two nozzles at a right angle facilitates gas mixing. We maintain the flow rates of the purge gases for the duration of the experiment in order to prevent gas leakage through any microscopic cracks on the chamber walls (Fig. S1†).

Use of the purge chamber introduces an additional conditioning step in the normal experimental workflow. At the beginning of each synthesis experiment, a brief waiting period is required for (1) the individual purge gas streams to mix and (2) the microfluidic device and its contents to equilibrate with the air/argon mixture within the chamber. In practice, the former occurs more rapidly than the latter. We placed a RedEye™ oxygen sensor patch above the purge gas outlet to monitor the oxygen concentration in the gas phase (Fig. 1). The patch contains an immobilized dye molecule that quenches in the presence of O<sub>2</sub>. To make chamber oxygen readings, we point a bifurcated optical fiber (RE-BIFBORO-2) at the RedEye™ patch. The fiber directs a blue LED excitation light toward the patch, and excites the sensor dye. The red emitted signal is then collected by the optical probe and transmitted to a fluorometer, which measures the fluorescence lifetime,  $t$ .  $t$  is related to the partial pressure of oxygen by the well-known Stern–Volmer equation.<sup>35</sup> Fig. 2(b) shows the chamber oxygen concentration as a function of time for a range of air inlet pressures. In all cases, the oxygen concentration reached an equilibrium value approximately 1 minute after the onset of purge, indicating efficient gas mixing.

A second time delay is associated with the synthesis device and the prepolymer solution flowing within the microchannel coming to equilibrium with the surrounding chamber gas mixture. Duration of the delay or response time is determined by the thickness of the PDMS layer above the synthesis microchannel. To demonstrate this, we assembled devices of varying

PDMS thickness (the channel height stayed the same at 2 μm). The practical lower threshold (~2 mm) is set by the minimum thickness necessary to reliably secure fluidic connections into the device inlet and outlet reservoirs. Using a canonical composition (80% v/v PEGDA700, and 15% v/v Darocur®1173, and 5% v/v rhodamine B), we synthesized cuboids of decreasing size in 2 μm tall channels under constant influx of pure argon (1.0 psi). Fig. 2(a) shows the time necessary for the smallest feature (5 μm square) to stick to the confining PDMS surfaces due to lack of oxygen inhibition near the channel periphery. The observed linear trend in log-log format, with a power-law slope of 0.5, is consistent with a simple diffusive mechanism, in which the PDMS layer above the synthesis device acts as the dominant barrier for mass transfer. We estimate the characteristic diffusion coefficient to be ~O(10<sup>−9</sup> m<sup>2</sup> s<sup>−1</sup>), on par with the diffusivity of oxygen<sup>36</sup> and argon<sup>37</sup> through PDMS. In order to limit the overall equilibration time prior to synthesis to ~5 min, we developed a tiered PDMS synthesis device (Fig. S2†) using a two-step curing process. First, we poured a thin layer (~1 mm) of PDMS over the device wafer and partially cured in the oven at 65 °C for 25 min. Then, shortened pipette tips (200 μL) were used to shield the central synthesis portion of each device and allow a second layer of PDMS to coat only the inlet and outlet regions, where rapid gas transport is unessential. The sacrificial pipette tips (Fig. S2B†) were removed from the wafer once the PDMS fully cured after baking overnight at 65 °C.

Fabrication of free-standing gel particles using SFL requires non-zero oxygen concentrations in the purge chamber in order to avoid particle sticking. As such, we systematically calibrated the equilibrium chamber oxygen concentration for a wide range of air pressures using the RedEye™ sensor. The concentration measurements were collapsed using a simple model derived from the Blasius correlation for high Re flow in smooth pipes<sup>43</sup> (Fig. 2(c)). This approach is justified since the Re for purge gas

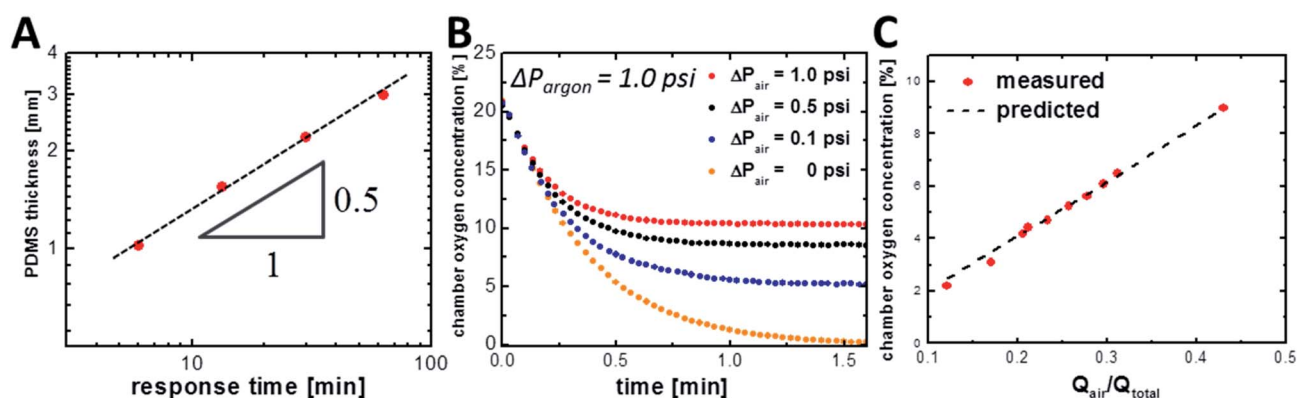


Fig. 2 Timescale comparison. (A) Thickness of the PDMS synthesis device determines the threshold response time (defined as the duration of continuous gas purge needed to reach equilibrium). The measured response time obeys a power-law trend, with exponent of 0.5 (a dashed line is drawn to guide the eye), which suggests a simple diffusion-based mechanism, with a characteristic coefficient ~O(10<sup>−9</sup>) m<sup>2</sup> s<sup>−1</sup>, typical of oxygen<sup>35</sup> and argon<sup>36</sup> diffusion through porous PDMS. (B) The amount of oxygen in the chamber for a range of air feed pressures: 0 psi (orange circles), 0.1 psi (blue circles), 0.5 psi (black circles), 1.0 psi (red circles). Pressure of the argon stream is fixed at 1 psi. Efficient mixing at the gas stream inlets result in short time lags (<1.5 min) until a final chamber oxygen concentration plateau is reached. (C) Equilibrium oxygen concentration in the purge chamber as a function of volume fraction of air introduced. The empirical measurements (red circles) made using the RedEye™ oxygen sensors are in excellent agreement with a simple model of high Re gas mixing derived from the Blasius correlation (dotted line, eqn (2) in text).



flow estimated using a typical pressure drop of 1 psi was >4000. Applying the Blasius equation, gas flow rate,  $Q_i$ , where  $i$  stands for either air or argon, can be expressed as:

$$Q_i = 2.26 \left( \frac{\Delta P_i}{L_i} \right)^{4/7} (\rho_i^3 \eta_i)^{-1/7} d^{19/7} \quad (1)$$

$\Delta P/L$  is the pressure drop per unit length of tubing,  $d$  is the nozzle diameter ( $\sim 1$  mm),  $\rho$  and  $\eta$  are the density and kinematic viscosity, respectively. Therefore, setting  $L_{\text{argon}} = L_{\text{air}}$ , the volume fraction of air introduced  $Q_{\text{air}}/Q_{\text{total}}$  is:

$$\frac{Q_{\text{air}}}{Q_{\text{total}}} = \frac{\Delta P_{\text{air}}^{4/7}}{\Delta P_{\text{air}}^{4/7} + c \Delta P_{\text{argon}}^{4/7}} \quad (2)$$

here,  $c = (\rho_{\text{air}}^3 \eta_{\text{air}} / \rho_{\text{argon}}^3 \eta_{\text{argon}})^{1/7}$  and is numerically equal to 0.845 once we substitute in values for the physical properties (Table S1†). To achieve symmetric mixing, length of the tubing for the argon needs to be decreased by  $\sim 18\%$ . The experimental data and model predictions (Fig. 2(c)) are in excellent agreement. The raw oxygen concentration data as a function of air pressure appears in Fig. S3.†

### 3.2 Modeling of photopolymerization

The importance of controlling ambient oxygen concentration in enabling thin particle synthesis can be understood using a simple reaction-diffusion model. This one-dimensional description of particle formation in a microfluidic channel was first formulated by Dendukuri *et al.*<sup>25</sup> It has since been adapted<sup>44</sup> and further validated *via* experiments.<sup>3</sup>

This model considers the spatial and temporal variation of dissolved oxygen concentration  $[O_2]$  and un-crosslinked oligomers  $[M]$  within a prepolymer mixture between confining PDMS channel boundaries ( $z$ -direction). Using mass action kinetics, we can write the governing equations as:

$$\frac{\partial [O_2]}{\partial t} = D_{\text{ox}} \frac{\partial^2 [O_2]}{\partial z^2} - k_o [O_2] [\dot{X}] \quad (3)$$

$$\frac{\partial [M]}{\partial t} = -k_p [M] [\dot{X}] \quad (4)$$

The RHS terms of eqn (3) correspond to oxygen diffusion through PDMS (with  $D_{\text{ox}}$  as the diffusion coefficient) and reaction with propagating radical species,  $\dot{X}$  (with rate constant  $k_o$ ), respectively. This model neglects the diffusion of oligomers due to their bulky size relative to that of oxygen. As a result, only the propagation reaction (with rate constant  $k_p$ ) contributes to the decrease in  $[M]$  over time. The two expressions are coupled *via* the inhibition reaction where oxygen quenches further polymerization by reacting with propagating radical species.  $[\dot{X}]$  can be estimated by invoking the quasi-steady state approximation, and setting the rate of radical generation ( $r_a$ ) in a thin film of prepolymer at height  $z$  equal to the rate of consumption by the inhibition and termination reactions:

$$[\dot{X}] = \frac{-k_o [O_2] + \sqrt{(k_o [O_2])^2 + 4k_t r_a}}{2k_t} \quad (5)$$

where  $r_a = \psi \varepsilon [\text{PI}] I_o \exp(-\varepsilon [\text{PI}] z)$ .  $\psi$  is the quantum yield of the initiating radical species.  $\varepsilon$  and  $[\text{PI}]$  is the molar extinction coefficient (at 365 nm) and concentration of the photoinitiator, respectively.  $I_o$  is the incident light intensity.  $k_t$  is the rate constant for the bimolecular termination reaction. For detailed derivation of the model equations and full reaction mechanism, see ref. 25.

The governing equations, eqn (3) and (4), can be non-dimensionalized using

$$\tau = t D_{\text{ox}} / H^2, \quad \eta = z / H$$

$$\theta = [O_2] / [O_2]_{\text{eq}}, \quad \xi = [M] / [M]_0$$

to obtain:

$$\frac{\partial \theta}{\partial \tau} = \frac{\partial^2 \theta}{\partial \eta^2} - \text{Da}_1 \theta \left( -\theta + \sqrt{\theta^2 + \alpha \exp(-\beta \eta)} \right) \quad (6)$$

$$\frac{\partial \xi}{\partial \tau} = -\text{Da}_2 \xi \left( -\theta + \sqrt{\theta^2 + \alpha \exp(-\beta \eta)} \right) \quad (7)$$

where  $H$  is the channel height,  $[O_2]_{\text{eq}}$  is the equilibrium concentration of dissolved oxygen in the oligomer, and  $[M]_0$  is the initial concentration of oligomer species. Resulting expressions for the dimensionless parameters are as follows:

$$\text{Da}_1 = \frac{k_o^2 H^2 [O_2]_{\text{eq}}}{2 D_{\text{ox}} k_t}, \quad \text{Da}_2 = \frac{k_p k_o H^2 [O_2]_{\text{eq}}}{2 D_{\text{ox}} k_t}$$

$$\alpha = \frac{4\psi \varepsilon [\text{PI}] I_o k_t}{E k_o^2 [O_2]_{\text{eq}}^2}, \quad \beta = \varepsilon [\text{PI}] H$$

$\text{Da}_1$  and  $\text{Da}_2$  are Damkohler numbers. The former quantifies the relative rates of oxygen inhibition and oxygen diffusion, while the latter quantifies the relative rates of chain propagation and oxygen diffusion. We note that the expression for  $\alpha$  differs from its counterpart in the original contribution by a factor of  $E$ , which denotes the energy content of a mole of photons at 365 nm ( $328 \text{ kJ mol}^{-1}$ ).<sup>25</sup> This conversion factor was not explicitly stated previously. However, we do so here to maintain dimensional consistency.

To computationally show the feasibility of creating thin particles by controlling the chamber oxygen concentration, we numerically integrated eqn (6) and (7) to obtain  $\theta$  and  $\xi$  given the parameters listed in Table 1 and the following boundary conditions:

$$\theta(0, \tau) = 1, \quad \theta(1, \tau) = 1, \quad \theta(\eta, 0) = 1$$

which assume that the oligomer contains its equilibrium concentration of  $O_2$  both initially and at the PDMS walls.

In order to vary  $[O_2]$  in the chamber, we assumed 1) the equilibrium dissolved oxygen concentration  $[O_2]_{\text{eq}}$  in the

**Table 1** Summary of model parameters used in the simulation

Parameter	Value	Units	Source
$k_p$	25	$\text{m}^3 \text{mol}^{-1} \text{s}^{-1}$	Ref. 38
$k_t$	$2.52 \times 10^3$	$\text{m}^3 \text{mol}^{-1} \text{s}^{-1}$	Ref. 38
$k_o$	$5 \times 10^5$	$\text{m}^3 \text{mol}^{-1} \text{s}^{-1}$	Ref. 39
$D_{\text{ox}}$	$2.84 \times 10^{-11}$	$\text{m}^2 \text{s}^{-1}$	Ref. 40
$H$	2	$\mu\text{m}$	Measured
$I_o$	$1.25 \times 10^4$	$\text{W m}^{-2}$	Measured
$[\text{PI}]$	990	$\text{mol m}^{-3}$	Calculated
$\varepsilon$	1.6	$\text{m}^3 \text{mol}^{-1} \text{m}^{-1}$	Ref. 41
$[\text{O}_2]_{\text{eq}}$	1.5	$\text{mol m}^{-3}$	Ref. 42
$\phi$	0.6	—	Ref. 41

prepolymer mixture obeys Henry's law, with a proportionality constant  $k_H$  of  $140 \text{ L atm mol}^{-1}$ ; and (2),  $[\text{O}_2]_{\text{eq}} = 1.5 \text{ mol m}^{-3}$  under standard ambient conditions (20.9%  $\text{O}_2$ ). Simulation results for  $\tau = 0.1$  are shown in Fig. 3. Physically, this corresponds to  $\sim 100 \text{ ms}$  for a  $2 \mu\text{m}$  tall device, similar to exposure times used during synthesis. Keeping the exposure time short helps to generate more uniform particles.<sup>45</sup>

At high chamber oxygen concentrations ( $\geq 10\%$ ), transport of oxygen near the channel interfaces occurs rapidly and prevents particle formation. Here, we adopt the convention that gelation occurs when monomer conversion first reaches 2% ( $\xi_c = 0.98$ ), similar to what was done previously.<sup>25</sup> Below 10%  $\text{O}_2$ , the propagation reaction dominates the inhibition reaction between radical species and dissolved oxygen. As a result, particles grow from the channel center, where the dissolved oxygen concentration is at a minimum, outward. The 10%  $\text{O}_2$  threshold satisfies the critical Da criterion for effective synthesis ( $\text{Da} \geq 4$ ) developed by Dendukuri *et al.*<sup>25</sup>

To confirm the validity of the simulations, we synthesized  $10 \mu\text{m}$  discs in a  $2 \mu\text{m}$  tall tiered microfluidic device under a range

of chamber oxygen concentrations. We imaged the resulting gel structures *in situ*. When  $[\text{O}_2]$  was above  $\sim 10\%$ , particle could not be made even using extended exposure durations ( $\sim 1 \text{ s}$ ), in agreement with model predictions. However, as  $[\text{O}_2]$  in the purge chamber was decreased, discs formed according to the mask pattern with *increasing* height (Fig. 3(c)). A similar trend was qualitatively forecast by simulations (Fig. 3(b)). In practice, performing synthesis with chamber oxygen level below 4% is not ideal, as it leads to excessive particle sticking after  $\sim 10 \text{ min}$  of operation.

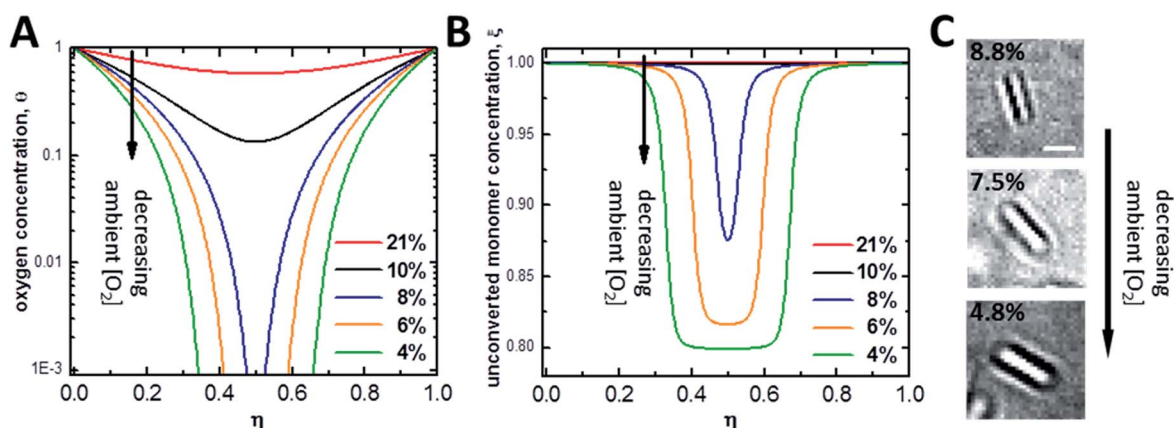
Synthesis of thinner particles in microfluidic channels with height  $H < 2 \mu\text{m}$  will be difficult for two reasons (1) the window of opportunity for effective synthesis is small (Da scales quadratically with  $H$  but only linearly with  $[\text{O}_2]_{\text{eq}}$ ), and (2) the pressure drop needed to drive flow scales as  $H^{-3}$ .<sup>46</sup> Therefore, device modifications may be required to prevent channels from bursting.

### 3.3 Brownian motion of microparticles

Achieving the colloidal (or cellular) size regime is a necessary synthesis milestone for probing the interactions of designer microparticles with biological or biomimetic environments. It is well known that one of the hallmarks of colloidal objects is they undergo spontaneous, random walk when suspended in a viscous fluid. A common statistical measure for particle dynamics in an inherently stochastic process is mean-squared displacement (MSD),  $\langle \Delta R^2(\tau) \rangle$ , calculated from particle trajectories:

$$\langle \Delta R^2(\tau) \rangle = \langle (R(t + \tau) - R(t))^2 \rangle \quad (8)$$

where  $R(t)$  is the position of the particle at time  $t$ , and  $\tau$  is the lag time between the two positions occupied by the same particle.  $\langle \dots \rangle$  denotes a time-average. In a Newtonian fluid, such as



**Fig. 3** Steady state profiles of oxygen ( $\theta$ ) and unconverted monomer concentration ( $\xi$ ) as a function of non-dimensionalized channel height ( $\eta$ ). The results shown in (A) and (B) were obtained by numerically integrating eqn (6) and (7) using the parameter values listed in Table 1 for a range of ambient oxygen concentrations. Decreasing ambient  $\text{O}_2$  concentration leads to a reduction in oxygen penetration depth near the channel walls, where rapid oxygen diffusion prevents particle formation. Cross-linked structures start to grow from the channel center as  $\theta$  and  $\xi$  fall below a critical threshold value ( $\theta_c \sim 10^{-3}$ ,  $\xi_c = 0.98$ ) for gelation. Modulating the chamber  $[\text{O}_2]$  allows particle height to approach the sub-micron regime. (C) A series of bright-field micrographs showing the change in height of  $10 \mu\text{m}$  diameter discs for a range of chamber  $[\text{O}_2]$  (noted in the upper left corner). Above  $\sim 10\%$ , no particles can be made, in agreement with model predictions. Scale bar is  $5 \mu\text{m}$ .

water, the MSD of particles moving in a plane (2D) grows linearly with lag time,

$$\langle \Delta R^2(\tau) \rangle = 4D_i\tau \quad (9)$$

where  $D_i$  is the translational diffusion coefficient in bulk ( $i = \infty$ ) or near the surface ( $i = 0$ ).

To demonstrate the colloidal nature of the microparticles generated using the modified SFL technique, we synthesized 5  $\mu\text{m}$  and 8  $\mu\text{m}$  diameter discs in the presence of 8.5%  $\text{O}_2$ . These synthesis experiments would not be feasible without lowering the oxygen concentration in the vicinity of the synthesis microfluidic device. Rhodamine B was covalently conjugated to the hydrogel backbone during synthesis to facilitate particle tracking. We recorded the motion of the microdiscs at least 4 particle diameters away from the rigid bottom surface of the observation chamber. The experimental buffer solution was density-matched by adding 90% v/v deuterium oxide in order to prolong the observation period, in which the particles remain in the focal plane of the microscope objective. We did not observe any particle swelling after  $\text{D}_2\text{O}$  addition. The choice to synthesize discs rather than more complex shapes enables the use of a public-domain algorithm to first locate the particles, then extract the particle trajectories, and finally calculate the MSDs<sup>47</sup> (Fig. 4). Trajectories were constructed by linking of the brightness centroid locations of candidate particles found on at least 10 consecutive frames, until they fell out of focus. The resulting MSD curves are linear in  $\tau$ , consistent with Brownian motion. The corresponding translational diffusivity coefficients in bulk ( $D_\infty$ ) were calculated from lines of best fit to the data using eqn (9) (Table 2). The values of  $D_\infty$  decrease as the size of the discs increase, as one would intuitively expect.

We can use the bulk diffusivity measurements to estimate the height of the colloidal particles. This method is valuable because the out-of-plane dimension of non-spherical colloids is

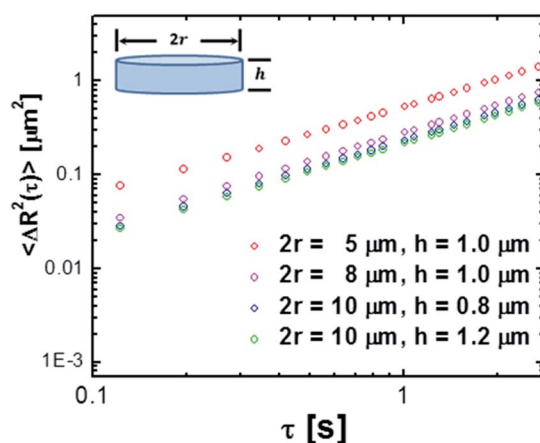


Fig. 4 Mean-square displacement ( $\langle \Delta R^2 \rangle$ ) in bulk as a function of lag time,  $\tau$  for various colloidal discs suspended in 90% v/v  $\text{D}_2\text{O}$ . The MSD curves show a power-law slope of 1, which is consistent with Brownian motion, revealing the colloidal nature of these disc-shaped particles. For Brownian motion in 2-dimensions,  $\langle \Delta R^2 \rangle = 4D\tau$ , where  $D$  is the translational diffusivity. The value of  $h$  for each particle sample was calculated from eqn (11) given the diffusivity measurement.

Table 2 Bulk diffusivity of the microdiscs measured using multiple particle tracking experiments. Values of  $h$  were calculated using eqn (11).  $r_s$  is related to  $h$  by  $r_s = \sqrt[3]{3/4r^2h}$

$r$ ( $\mu\text{m}$ )	$r_s$ ( $\mu\text{m}$ )	$h$ ( $\mu\text{m}$ )	$D_\infty$ ( $\mu\text{m}^2 \text{s}^{-1}$ )
2.5	1.67	1.0	0.105
4	2.29	1.0	0.070
5	2.47	0.8	0.059
5	2.82	1.2	0.055

often difficult to directly quantify using conventional optical microscopy techniques. In general, bulk diffusivities are affected by particle geometry and size. In the simplest case, translational diffusion coefficient,  $D_s$ , of a sphere is related to its radius,  $r_s$ , by the Stokes–Einstein equation:

$$D_s = \frac{kT}{6\pi\eta r_s} \quad (10)$$

where  $k$  is the Boltzmann constant,  $T$  is the absolute temperature,  $\eta$  is the fluid viscosity.

For thin discoids, the bulk diffusivity,  $D_\infty$ , is a function of the aspect ratio,  $p = h/2r$ , where  $h$  is the height, and  $r$  is the radius of the disc. Hansen<sup>48</sup> developed the following correlation that fits results from Monte Carlo simulations:

$$D_s/D_\infty = 1.0304 + 0.0193 \ln p + 0.0623(\ln p)^2 + 0.0048(\ln p)^3 + 0.0017(\ln p)^4 \quad (11)$$

where  $D_s$  is the bulk diffusion coefficient of a sphere with the same volume as the disc. Similar empirical expressions for  $D_\infty$  also exist (see for example, ref. 49); the impact of choosing one equation over another to estimate the height of the discs is minimal ( $\pm 1$ –3%) over the narrow regime of aspect ratio under consideration. It is readily seen that deviation from spherical symmetry reduces particle mobility. But as the particle aspect ratio  $p \rightarrow 1$ ,  $D_\infty \cong D_s$ .

To estimate the height of colloidal discs using eqn (10) and (11), we assumed the diameter of the particles to be the size of the mask pattern. This is a good assumption for two reasons: (1) the coefficient of variation for SFL is no more than a few percent,<sup>50</sup> and (2) the high PEGDA concentration minimizes any swelling effects. High resolution confocal imaging of a 10  $\mu\text{m}$  disc supports this assumption (Fig. 5(c)). Particle height calculated from diffusivity measurements and eqn (11) are shown in Table 2. Error introduced in  $h$  through the uncertainties in the bulk diffusivity measurement ( $\pm 0.002 \mu\text{m}^2 \text{s}^{-1}$ ) is less than 0.1  $\mu\text{m}$  in all cases (Fig. S4†). We note that the 5  $\mu\text{m} \times 1 \mu\text{m}$  discs are the smallest particles synthesized to date using SFL.

While size and geometry of the in-plane feature is defined by the photomask, height of the particles can be tuned by adjusting the chamber oxygen concentration.<sup>3</sup> And more importantly, sub-micron-tall particles can be generated by lowering the chamber  $\text{O}_2$  concentration close to the critical threshold of 10%. To show this, we synthesized 10  $\mu\text{m}$  discs using 8% and 9.5%  $\text{O}_2$ , and measured their bulk diffusivities in a 90% (v/v)  $\text{D}_2\text{O}$  aqueous mixture (Fig. 4). We estimated the out-of-plane

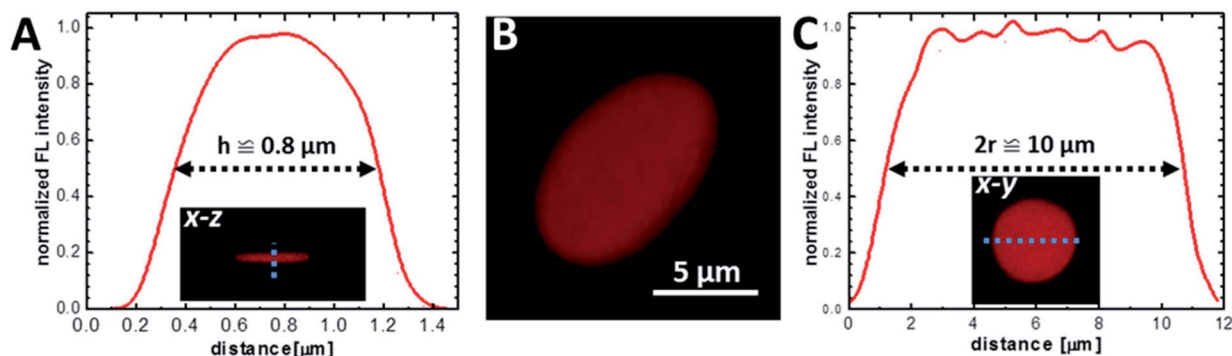


Fig. 5 Laser scanning confocal image of a representative 10  $\mu\text{m}$  disc synthesized using 9.5%  $\text{O}_2$  (B). Normalized fluorescence intensity profiles taken across the  $x$ - $z$  (A, blue dotted line) and  $x$ - $y$  (C, blue dotted line) planes of the particle. Particle height is estimated to be  $\sim 0.8 \pm 0.1 \mu\text{m}$ . Full width at half-maximum (FWHM), as indicated black dotted lines, was used to define the particle boundaries.

dimension of the particles to be 1.2 and 0.8  $\mu\text{m}$ , respectively (Table 2).

To verify the latter result, a laser scanning confocal microscope equipped with a high numerical aperture oil immersion objective ( $\text{NA} = 1.3$ ) was used to directly visualize the thinnest particle sample (10  $\mu\text{m} \times 0.8 \mu\text{m}$  discs). Confocal microscopy relies on point illumination through a pinhole to eliminate out-of-focus light in the region of interest (ROI) within the sample, resulting in increased optical resolution. Using an excitation wavelength appropriate for rhodamine B (568 nm), the in-plane resolution is  $\sim 0.25 \mu\text{m}$ . The particles were immersed in 2% alginate solution to arrest their orientation, and imaged on the bottom surface. The image stack was assembled using ImageJ to construct the 3D structure of the microgels (Fig. 5(b)). Dimensions of the particle were gauged at full width at half-maximum (FWHM) from spatially-resolved intensity profiles (Fig. 5). Accordingly, the particle height was estimated to be  $\sim 0.8 \mu\text{m}$ , in agreement with predictions made based on diffusivity measurements. Error introduced by Brownian motion during imaging acquisition is estimated to be  $\sim 0.1 \mu\text{m}$ .

Next, we examine the Brownian motion of discoids near a planar solid surface. Studying the transport behaviour of non-spherical particles near physical boundaries has proven to be a difficult task both theoretically and experimentally. Analytical expressions from theoretical treatment of hindered diffusion are limited to the simplest particle geometries (*i.e.*, spheres<sup>51</sup> and discs). Techniques which are capable of creating exotic particle morphologies (*i.e.*, lock-key particles,<sup>52</sup> boomerangs,<sup>53</sup> *etc.*), have only recently been introduced.

To measure the hindered diffusion coefficients of discs experimentally, we suspended the particles in an aqueous buffer and allowed ample time for them to sediment to the bottom surface of the observation chamber prior to imaging. Up to 2500 frames were collected per movie in order to gather at least 20 000 particle trajectories in total and extract the in-plane translational diffusion coefficient,  $D_0$ . Fig. 6 and Table 3 show MSDs and hindered diffusivities, respectively, in two different buffers. Unlike their bulk counterparts (Table 2), mobility of microparticles is reduced by as much as  $\sim 80\%$  owing to hydrodynamic interactions with the wall. For thin discs

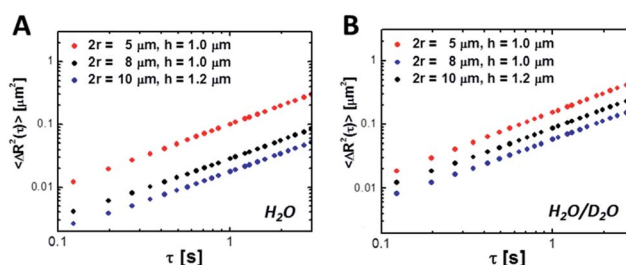


Fig. 6 Mean-square displacement  $\langle \Delta R^2 \rangle$  near a wall as a function of lag time,  $\tau$  for various colloidal discs in (A) water and (B) a mixture of deuterium oxide and water (9 : 1 ratio by volume).

Table 3 Hindered diffusivity estimates for colloidal discs from experiment and theory (eqn (13)) in two particle storage buffers. Addition of  $\text{D}_2\text{O}$  decreases density mismatch between the buffer and the particle, causing the average gravitational height to increase

$r$ ( $\mu\text{m}$ )	Buffer	$h_g$ (nm)	$D_0$ ( $\mu\text{m}^2 \text{s}^{-1}$ )	
			Experiment	Model
2.5	$\text{H}_2\text{O}$	213.4	0.0285	0.0271
2.5	$\text{D}_2\text{O}/\text{H}_2\text{O}$	1067	0.0381	0.0814
4	$\text{H}_2\text{O}$	83.3	0.0069	0.0056
4	$\text{D}_2\text{O}/\text{H}_2\text{O}$	416.7	0.0210	0.0199
5	$\text{H}_2\text{O}$	44.5	0.0042	0.0029
5	$\text{D}_2\text{O}/\text{H}_2\text{O}$	266.7	0.0129	0.0093

diffusing close to a wall, the preferred mode of motion is in the plane of the particle, parallel to the glass slide. Addition of 90% deuterium oxide by volume reduced the density mismatch ( $\Delta\rho$ ) between the aqueous buffer and the particles from 0.10 to 0.02  $\text{g mL}^{-1}$ . The particle density was estimated using the weighted sum of PEGDA and the photoinitiator densities to be 1.1  $\text{g mL}^{-1}$ . This assumes 100% conversion, and represents the density upper bound. As a result, the average gravitational height of the particles,

$$h_g = \frac{kT}{\Delta\rho V_{pg}} \quad (12)$$



increases, as does the value of  $D_0$  (Table 3). In eqn (12),  $V_p$  is the particle volume, and  $g$  is the constant of gravitational acceleration ( $9.81 \text{ m s}^{-2}$ ). The elevation-dependent Brownian dynamics are clearly seen for the 8 and  $10 \mu\text{m}$  discs, for which viscous drag forces opposing edgewise translation decreases most rapidly.<sup>54</sup>

In reality, the equilibrium separation distance between the particle and the surface continuously fluctuates as a result of thermal motion. It is difficult to isolate the dynamics of a particle sliding parallel to the glass slide from movement in the transverse direction without resorting to specialized techniques, such as total internal reflection microscopy.<sup>55</sup> As a result, the experimentally-determined  $D_0$  values are in fact apparent diffusion coefficients that encompass the cumulative effect of sampling a range of gap heights on particle motion:

$$D_0 = \langle D_0 \rangle = \int_0^\infty kTf^{-1}(a)p(a)da \quad (13)$$

where  $p(a)$  is the normalized probability of finding the particle at a distance  $a$  from the bottom surface,  $f(a)$  is the friction coefficient of the particle in close proximity to the wall.

Kim *et al.* derived an analytical expression for  $f(a)$ , characterizing the edgewise translation of a thin circular plate parallel to an infinite plane:<sup>54</sup>

$$f(a) = \eta \left( \frac{\pi r}{a} + 3 \ln \frac{r}{a} + 6.294 \right) r \quad (14)$$

At equilibrium,  $p(a)$  is the Boltzmann distribution, with  $h_g$  as the average lubricating gap distance, where potential and thermal energies balance:

$$p(a) \propto \exp(-a/h_g) \quad (15)$$

Substituting eqn (13) and (14) into eqn (15), we can calculate the hindered diffusivity as long as the particle dimensions are known *a priori*. This simple model contains no adjustable parameters, and captures the essential physics. It can predict  $D_0$

to within 25% when  $h_g \ll r$  (Table 3). This can be attributed, at least in part, to overestimating the conversion and therefore, particle density. As the separation distance grows, however, this model breaks down, as the colloidal discs can no longer be approximated as infinitely thin (Table 3,  $2.5 \mu\text{m}$  radius discs in  $\text{D}_2\text{O}/\text{H}_2\text{O}$ ). We note that a similar approach was used to understand the Brownian fluctuations of charge-stabilized colloidal spheres located near a flat plate.<sup>56</sup> The close agreement between the model predictions and experimental data gave us confidence regarding the accuracy with which we estimated the particle dimensions using multiple particle tracking. This model can be used to guide the self-assembly of non-spherical particles near a solid boundary.

### 3.4 Flow of hydrogel colloids through microvascular networks

The ability of SFL to generate anisotropic colloidal particles allows us to begin to probe the flow behaviour of particles in a biological environment. Past experiments revealed interesting shape<sup>4</sup> and elasticity<sup>5,7</sup> induced passage properties, both *in vitro* and *in vivo*. A novel method for understanding particle flow characteristics involves the use of engineered, microfluidic vascular networks ( $\mu\text{VNs}$ ).  $\mu\text{VNs}$  can be formed by endothelial cells (HUVECs) undergoing the process of vasculogenesis. In the presence of human lung fibroblasts, HUVECs gradually develop stable networks of interconnected, perfusable vessels after  $\sim 4$  days in non-contact cell culture.<sup>26</sup> Phenotypic state of the network, such as the number of branches, and average diameter or length of the vessels, can be tuned by adjusting initial input parameters during cell culture (*e.g.*, concentration of growth factors). To demonstrate the passage of fluorescent PEG-based colloids through  $\mu\text{VNs}$ , we flowed  $10 \mu\text{m} \times 1.2 \mu\text{m}$  discs by applying a small ( $\sim 5 \text{ mmH}_2\text{O}$ ) hydrostatic driving pressure across the vascularized region on the microfluidic device. Particles entered the network *via* vessel openings to the adjacent medium channel (Fig. S5†), and navigated through by exploiting the path of least fluidic resistance. A representative

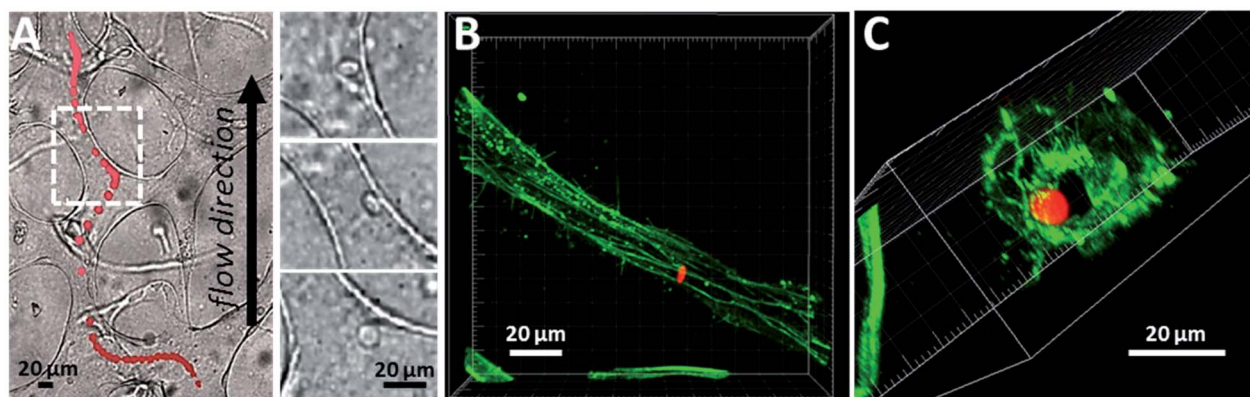


Fig. 7 (A) Trajectory of a colloidal disc ( $2r = 10 \mu\text{m}$ ,  $h = 1.2 \mu\text{m}$ ) through the microvascular network grown on a multiculture microfluidic device. The black arrow indicates the general direction flow driven by a small hydrostatic pressure difference across the network. The discs assume different orientations during their passage (center panel). The close-ups also show that they do not adhere to the microvessel walls. Confocal micrographs of the longitudinal (B) and the transverse (C) directions of a vessel segment (green) containing a residual particle (red).

particle trajectory through a portion of the network is shown in Fig. 7. The fluorescence signal from the candidate particle was isolated from consecutive frames of experimental movies (Video S1†) and superimposed on top of a bright-field image showing the layout of the microvessels in the ROI. To preserve mass conservation, particles accelerated (marked by the spreading of their fluorescence signals) when they encountered contractions, and *vice versa*. Importantly, particles do not adhere to the vessel walls (Fig. 7(a), right panel), showing signs of bio-inertness. To clearly show the size of a representative vessel in 3D relative to that of a particle, we fluorescently stained for actin with Alexa Fluor® 488 phalloidin after the flow experiment and imaged a residual particle lingering in the lumen. These results demonstrate the feasibility of using  $\mu$ VNs as a useful tool to study particle passage properties. Future experiments may offer mechanistic insight on shape or size effects on particle flow that ultimately result in distribution patterns seen in animal studies.

## 4. Conclusions

In summary, we have shown that controlling the oxygen concentration surrounding the synthesis microfluidic device allows us to synthesize colloidal particles using flow lithography. One of the key advantages of this technique is that geometry of the particles is mask-defined, and as a result, not limited to simple polygonal shapes. Morphologically-complex colloids are critical for efficient assembly of ordered functional materials. In addition, passage of non-spherical colloids through microfluidic vascular networks deserves further attention. This platform can be used to understand the effect of particle shape and deformability on passage properties in a physiological, yet controlled setting.

## Acknowledgements

HZA and LC are supported by the Institute for Collaborative Biotechnologies through grant W911NF-09-0001 from the U.S. Army Research Office. Additional support is provided by NSF grants CMMI-1120724 and DMR-1006147, and the Singapore MIT Alliance for Research and Technology SMART. The content of the information does not necessarily reflect the position or the policy of the Government, and no official endorsement should be inferred. HZA thanks Andrew Ryan for making the microscope purge chamber, Wendy Salmon for her assistance with confocal microscope training, Bashar Hamza, A. J. Aranyosi, and Octavio Hurtado for device microfabrication at the Charlestown Navy Yard Clean Room facility, and A. Wagner for helpful comments on the manuscript. MBC is supported by the National Science Foundation Graduate Fellowship.

## Notes and references

- 1 K. W. Bong, S. C. Chapin and P. S. Doyle, *Langmuir*, 2010, **26**, 8008.
- 2 R. L. Srinivas, S. C. Chapin and P. S. Doyle, *Anal. Chem.*, 2011, **83**, 9138.
- 3 H. Z. An, E. R. Safai, H. B. Eral and P. S. Doyle, *Lab Chip*, 2013, **13**, 4765.
- 4 R. Haghgoie, M. Toner and P. S. Doyle, *Macromol. Rapid Commun.*, 2010, **31**, 128.
- 5 T. J. Merkel, S. W. Jones, K. P. Herlihy, F. R. Kersey, A. R. Shields, M. Napier, J. C. Luft, H. L. Wu, W. C. Zamboni, A. Z. Wang, J. E. Bear and J. M. DeSimone, *Proc. Natl. Acad. Sci. U. S. A.*, 2011, **108**, 586.
- 6 K. Chen, T. J. Merkel, A. Pandya, M. E. Napier, J. C. Luft, W. Daniel, S. Sheiko and J. M. DeSimone, *Biomacromolecules*, 2012, **13**, 2748.
- 7 N. Doshi, A. S. Zahr, S. Bhaskar, J. Lahann and S. Mitragotri, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 21495.
- 8 C. L. Modery-Pawłowski, L. L. Tian, V. Pan, K. R. McCrae, S. Mitragotri and A. Sen Gupta, *Biomaterials*, 2013, **34**, 526.
- 9 N. Doshi, J. N. Orje, B. Molins, J. W. Smith, S. Mitragotri and Z. M. Ruggeri, *Adv. Mater.*, 2012, **24**, 3864.
- 10 H. Z. An, M. E. Helgeson and P. S. Doyle, *Adv. Mater.*, 2012, **24**, 3838.
- 11 S. H. Kim, H. C. Shum, J. W. Kim, J. C. Cho and D. A. Weitz, *J. Am. Chem. Soc.*, 2011, **133**, 15165.
- 12 J. A. Champion, A. Walker and S. Mitragotri, *Pharm. Res.*, 2008, **25**, 1815.
- 13 T. J. Merkel, K. Chen, S. W. Jones, A. A. Pandya, S. M. Tian, M. E. Napier, W. E. Zamboni and J. M. DeSimone, *J. Controlled Release*, 2012, **162**, 37.
- 14 J. A. Champion, Y. K. Katare and S. Mitragotri, *J. Controlled Release*, 2007, **121**, 3.
- 15 P. Kolhar and S. Mitragotri, *Adv. Funct. Mater.*, 2012, **22**, 3759–3764.
- 16 J. W. Yoo, N. Doshi and S. Mitragotri, *Macromol. Rapid Commun.*, 2010, **31**, 142.
- 17 K. Herlihy, T. Merkel, H. Y. An, W. L. Lin and J. DeSimone, *Nanotech Conference & Expo 2009, Vol. 2, Technical Proceedings*, 2009, pp. 60.
- 18 H. C. Shum, A. R. Abate, D. Lee, A. R. Studart, B. G. Wang, C. H. Chen, J. Thiele, R. K. Shah, A. Krummel and D. A. Weitz, *Macromol. Rapid Commun.*, 2010, **31**, 108.
- 19 J. P. Rolland, B. W. Maynor, L. E. Euliss, A. E. Exner, G. M. Denison and J. M. DeSimone, *J. Am. Chem. Soc.*, 2005, **127**, 10096.
- 20 S. Bhaskar, K. M. Pollock, M. Yoshida and J. Lahann, *Small*, 2010, **6**, 404.
- 21 J. A. Champion, Y. K. Katare and S. Mitragotri, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **104**, 11901.
- 22 Y. P. Wang, T. J. Merkel, K. Chen, C. A. Fromen, D. E. Betts and J. M. DeSimone, *Langmuir*, 2011, **27**, 524.
- 23 D. Dendukuri, S. S. Gu, D. C. Pregibon, T. A. Hatton and P. S. Doyle, *Lab Chip*, 2007, **7**, 818.
- 24 D. Dendukuri, D. C. Pregibon, J. Collins, T. A. Hatton and P. S. Doyle, *Nat. Mater.*, 2006, **5**, 365.
- 25 D. Dendukuri, P. Panda, R. Haghgoie, J. M. Kim, T. A. Hatton and P. S. Doyle, *Macromolecules*, 2008, **41**, 8547.
- 26 J. A. Whisler, M. B. Chen and R. D. Kamm, *Tissue Eng., Part C*, 2013, **20**(7), 543–552.
- 27 M. B. Chen, J. A. Whisler, J. S. Jeon and R. D. Kamm, *Integr. Biol.*, 2013, **5**, 1262.

- 28 H. Bow, I. V. Pivkin, M. Diez-Silva, S. J. Goldfless, M. Dao, J. C. Niles, S. Suresh and J. Y. Han, *Lab Chip*, 2011, **11**, 1065.
- 29 D. C. Appleyard, S. C. Chapin, R. L. Srinivas and P. S. Doyle, *Nat. Protoc.*, 2011, **6**, 1761.
- 30 K. W. Bong, S. C. Chapin, D. C. Pregibon, D. Baah, T. M. Floyd-Smith and P. S. Doyle, *Lab Chip*, 2011, **11**, 743.
- 31 V. Pelletier, N. Gal, P. Fournier and M. L. Kilfoil, *Phys. Rev. Lett.*, 2009, **102**, 188303.
- 32 T. Savin and P. S. Doyle, *Biophys. J.*, 2005, **88**, 623.
- 33 H. B. Eral, V. Lopez-Mejias, M. O'Mahony, B. L. Trout, A. S. Myerson and P. S. Doyle, *Cryst. Growth Des.*, 2014, **14**, 2073.
- 34 K. W. Bong, J. Xu, J. H. Kim, S. C. Chapin, M. S. Strano, K. K. Gleason and P. S. Doyle, *Nat. Commun.*, 2012, **3**, 805.
- 35 D. B. Papkovsky and R. I. Dmitriev, *Chem. Soc. Rev.*, 2013, **42**, 8700–8732.
- 36 H. Shiku, T. Saito, C. C. Wu, T. Yasukawa, M. Yokoo, H. Abe, T. Matsue and H. Yamada, *Chem. Lett.*, 2006, **35**, 234.
- 37 L. Brandao, L. M. Madeira and A. M. Mendes, *J. Membr. Sci.*, 2007, **288**, 112.
- 38 S. Kızılel, V. H. Pérez-Luna and F. Teymour, *Macromol. Theory Simul.*, 2006, **15**, 686.
- 39 C. Decker and A. D. Jenkins, *Macromolecules*, 1985, **18**, 1241.
- 40 H. Lin and B. D. Freeman, *Macromolecules*, 2006, **39**, 3568.
- 41 L. Lecamp, P. Lebaudy, B. Youssef and C. Bunel, *Polymer*, 2001, **42**, 8541.
- 42 M. D. Goodner and C. N. Bowman, *Chem. Eng. Sci.*, 2002, **57**, 887.
- 43 M. M. Denn, *Process fluid mechanics*, Prentice-Hall, Englewood Cliffs, N.J., 1980.
- 44 S. K. Suh, K. W. Bong, T. A. Hatton and P. S. Doyle, *Langmuir*, 2011, **27**, 13813.
- 45 S. Park, D. Kim, S. Y. Ko, J. O. Park, S. Akella, B. Xu, Y. Zhang and S. Fraden, *Lab Chip*, 2014, **14**, 1551.
- 46 T. Gervais, J. El-Ali, A. Gunther and K. F. Jensen, *Lab Chip*, 2006, **6**, 500.
- 47 J. C. Crocker and B. D. Hoffman, *Methods Cell Biol.*, 2007, **83**, 141.
- 48 S. Hansen, *J. Chem. Phys.*, 2004, **121**, 9111–9115.
- 49 A. Ortega and J. G. de la Torre, *J. Chem. Phys.*, 2003, **119**, 9914.
- 50 D. C. Pregibon, M. Toner and P. S. Doyle, *Science*, 2007, **315**, 1393.
- 51 P. Huang and K. S. Breuer, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2007, **76**, 046307.
- 52 S. Sacanna, W. T. Irvine, P. M. Chaikin and D. J. Pine, *Nature*, 2010, **464**, 575.
- 53 A. Chakrabarty, A. Konya, F. Wang, J. V. Selinger, K. Sun and Q. H. Wei, *Phys. Rev. Lett.*, 2013, **111**, 160603.
- 54 M. U. Kim, K. W. Kim, Y. H. Cho and B. M. Kwak, *Fluid Dynam. Res.*, 2001, **29**, 137.
- 55 S. G. Flicker, J. L. Tipa and S. G. Biko, *J. Colloid Interface Sci.*, 1993, **158**, 317.
- 56 M. A. Bevan and D. C. Prieve, *J. Chem. Phys.*, 2000, **113**, 1228.