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# Tailored nanotopography of photocurable composites for control of cell migration†

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External mechanical stimuli represent elementary signals for living cells to adapt to their adjacent environment. These signals range from bulk material properties down to nanoscopic surface topography and trigger cell behaviour. Here, we present a novel approach to generate tailored surface roughnesses in the nanometer range to tune surface properties by particle size and volume ratio. Time-resolved local mean-squared displacement (LMSD) analysis of amoeboid cell migration reveals that nanorough surfaces alter effectively cell migration velocities and the active cell migration phases. Since the UV curable composite material is easy to fabricate and can be structured *via* different light based processes, it is possible to generate hierarchical 3D cell scaffolds for tissue engineering or lab-on-a-chip applications with adjustable surface roughness in the nanometre range.

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## 1 Introduction

The ultimate goal in the field of biofabrication and tissue engineering is to imitate the physiological environment of living cells in order to provide a controlled artificial environment as natural as possible. For the development of 3D cell scaffolds as basic modules in regenerative medicine and lab-on-chip applications, production methods including 3D printing,<sup>1</sup> stereolithography,<sup>2</sup> nanoimprint,<sup>3</sup> and direct laser writing<sup>4,5</sup> are used. These can be applied to various material classes ranging from hydrogels<sup>6</sup> to glass<sup>7</sup> and metals.<sup>8</sup> Simultaneously, manifold structure designs from 2D<sup>9</sup> to 3D<sup>10</sup> as well as chemical<sup>11</sup> and mechanical<sup>12</sup> surface altering procedures are under investigation, all with the aim to mimic the native extracellular 3D matrix and to influence cell behaviour in defined ways. This is an extremely challenging task, since many factors are involved when it comes to biophysicochemical interactions at the interface of the cells' membrane and the scaffold surface.<sup>13</sup> Besides the biochemical influence of the extracellular matrix, mechanical stimuli of the surrounding scaffold induce inter- and intracellular responses like proliferation,<sup>14</sup> cell differentiation<sup>15</sup> and gene expression.<sup>16</sup>

Several mechanical bulk- and surface properties, *e.g.* the elastic modulus<sup>17</sup> or surface topography in the micro-<sup>18</sup> and

nanometre<sup>19</sup> range as well as different geometrical surface features,<sup>20</sup> have been investigated over the last years. Recently the surface roughness, which is adjustable in multiple ways *via* additive<sup>21</sup> or subtractive<sup>22</sup> methods, raised special interest due to its potential to significantly influence proliferation, cell adhesion,<sup>23</sup> and stem cell differentiation.<sup>24,25</sup> To generate the desired surface roughness, different techniques are used like photolithography, nanoimprint, or self assembling polymer films.<sup>26</sup> They offer a high resolution and design freedom, however, for most of them the production process is expensive, due to the costly equipment and multiple processing steps. An easier way is the utilization of particles to create an adjustable surface roughness.<sup>27</sup> Besides the cheap and easy processing, several bulk and surface functionalities can be tuned for every specific requirement. In this way, it is easy to add and vary properties of a 3D scaffold, with custom shape and custom surface roughness. Furthermore, optimizations in terms of surface chemistry, intrinsic fluorescence for imaging purposes up to incorporated drug-delivery systems are possible. To enhance the scaffold with antibacterial or actuatoric characteristics, silver or magnetic particles can be used.<sup>28</sup>

In many physiological processes cell migration plays a crucial role, from embryogenesis to immune response and wound healing. To study amoeboid cell migration influenced by external cues, *Dictyostelium discoideum* (*D. discoideum*) is a popular cell model organism in the biophysical field,<sup>29</sup> since many properties can be transferred to mammalian cells.<sup>30</sup> During cell migration a distinct cell-surface interaction is necessary in order to transmit forces generated by the cytoskeleton to deform the cell body and to move it along the substrate. Simultaneously, the cell explores its surrounding *via* mechanosensing by its randomly arranged protrusions, where

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mechanical stimuli induce biochemical reactions. This can lead to cell polarization followed by directed cell migration and can be exploited for directed cell guidance.<sup>31,32</sup> Compared to mammalian cells, *D. discoideum* do not carry genes for integrins or most extracellular matrix proteins and therefore do not develop focal adhesion points on the substrate.<sup>33</sup> However, the so-called actin foci transmembrane protein *SadA* is found to mediate force transmission in surface adhesion and phagocytosis,<sup>34,35</sup> among other involved proteins like *phg1A*, *sibA* and talin A. Apart from that, adhesion is based on van der Waals forces,<sup>36</sup> which depends on the polarizability of the involved molecules and can be either attractive or repulsive and originate from permanent charges or ions inside the cell medium and are apparent at any contact site.

In this work, we present a novel approach to fabricate tailored surface roughnesses in the nanometre range by incorporating silica particles into a photo-structurable matrix, which is part of the hybrid polymer class ORMOCER®. It is widely used as smoothing<sup>37</sup> or passivation layer<sup>38</sup> in different applications and known for its excellent processability ranging from bulk material down to sub-micron resolution structures and from simple surface coating<sup>39</sup> to the creation of custom freeform 3D micro structures *via* two-photon polymerization.<sup>40</sup> This matrix material is processable with every light curing method and its biocompatibility has been proven with different cell types<sup>41–43</sup> making it ideally suited for future biomedical applications. Under normal conditions, ORMOCER® I (OC-I) – like many other light curable polymers – is polymerized under exclusion of oxygen, since it acts as a radical scavenger suppressing radical curing.<sup>44</sup> Here, we exploit this behaviour of OC-I forming a thin, oxygen-induced inhibition layer, when cured under ambient air. When dissolving this layer after curing, the surface exhibits a topography in the nanometre range corresponding to the particles protruding out of the matrix surface. The surface nanoroughness is tunable by adjusting the diameter and filling degree of incorporated particles. Here, tailored nanorough surfaces have been tested for biological application by their influence on the cell migration behaviour of *D. discoideum* amoebae. The living cells were seeded onto the sample surface and monitored *via* time-lapse fluorescent microscopy. The cell locomotion paths were analysed using a local mean-squared displacement (LMSD) algorithm in order to distinguish between directed and quasi-random migration modes depending on the surface roughness of the sample.

## 2 Materials and methods

### 2.1 Sample preparation

**2.1.1 Hybrid polymer matrix ORMOCER® OC-I.** OC-I is synthesized *via* an anorganic condensation reaction of the silane precursors 3-methacryloxypropyltrimethoxysilane (MEMO) and diphenylsilanediol (DPD, both ABCR GmbH, Germany), mixed in a molar ratio of 1 : 1.<sup>45</sup> Adding 2 wt% of the photoinitiator Irgacure® 369 (2-benzyl-2-dimethylamino-1-(4-morpholinophenyl)-butanone-1), Ciba Geigy, Switzerland) results in a UV-curable resin.<sup>46,47</sup>

**2.1.2 Silica particles.** All incorporated particles were synthesized *via* a modified Stoeber process according to Dembski *et al.*<sup>48</sup> and Milde *et al.*<sup>49</sup> In the synthesis process of silica particles, 225 g of the precursor tetraethoxysilane (TEOS, 98%, Sigma-Aldrich Chemie GmbH, Germany) is hydrolyzed in 4.5 L ethanol (Sigma-Aldrich Chemie GmbH, Germany) and 225 mL of the catalyst ammonium hydroxide (NH<sub>4</sub>OH 25%, Fluka®, Honeywell International, Inc., USA). During aging at RT for 3 days, hydrolysis and condensation reaction of the silanol groups enable the formation of colloidal silica particles. Before further use, the solvent is evaporated and the particles washed three times with ethanol.

**2.1.3 Preparation of composite and cured layers.** A silica particle butanone suspension was added to OC-I and mixed for 36 hours on a 40 °C hotplate allowing the solvent to evaporate.

A drop of the composite was placed onto a 1.5 H high precision cover slip (Marienfeld, Germany), doctor-bladed to a 30 µm layer and cured in an iron-light source UV-Chamber (BK 850, Beltron®, Germany) under ambient air for 5 minutes. Oxygen leads to an inhibition surface layer<sup>50</sup> of several micrometers, which was removed during a 15 min development step in a mixture (1 : 1) of isopropanol and 4-methyl-2-pentanone (Sigma-Aldrich Chemie GmbH, Germany). The samples were UV-cured again for 5 minutes within the solvent bath to cross-link remaining carbon double bonds. This was followed by a 24 hours post heating step at 60 °C under a vacuum of 100 mbar to ensure complete evaporation of the solvent and post-curing of the anorganic network of the ORMOCER® matrix. Table 1 shows the produced surface samples with the respective particle diameter and filling ratio. Only particles with diameters smaller than 0.5 µm were utilized for the creation of nanorough surfaces to achieve surface roughness values in the nanometre range distinct from a truly three-dimensional environment.

### 2.2 Sample characterization

**2.2.1 Scanning electron microscopy.** All scanning electron microscopy (SEM) samples were sputter coated with platinum (Med010, former Balzers Union, Lichtenstein, now Baltec, Germany) before imaging with a scanning electron microscope using Inlens- and SE2-detectors (Supra25, Carl Zeiss AG, Germany).

To analyze the silica particle size distribution, a drop of highly diluted silica particle butanone suspension was applied onto a SEM sample holder and dried before sputter coating. A watershed separation was performed on the SEM images using ImageJ 1.51 h (National Institutes of Health, USA) in order to enhance the particle detection, before determining the particle size using the ImageJ plug-in “Analyse Particle”. The error in particle diameter is based on the standard deviation of this analysis.

For imaging of cured composite layers, the coated cover slip was cut and the layer surface and layer cross section were examined under a 60 and 90 degree angle, respectively.

**2.2.2 Atomic force microscopy.** Surface roughness parameters were determined using an atomic force microscope (AFM, Alpha 300A, WITec, Germany). To neutralize electrostatic charges, the samples were cleaned with ionized compressed air (Haug, Germany) directly before the AFM measurements, which



**Table 1** Overview of all sample variations with specifics on particle diameter and volume fraction as well as the gas environment during UV-curing

Sample	Particle diameter (nm)	Particle filling ratio (vol%)	Curing environment
OC-I AR	—	—	Argon
OC-I	—	—	Ambient air
52–20	52 ± 8	20	Ambient air
165–5	165 ± 11	5	Ambient air
165–10	165 ± 11	10	Ambient air
165–20	165 ± 11	20	Ambient air
421–20	421 ± 22	20	Ambient air
421–20 AR	421 ± 22	20	Argon

were performed in tapping mode with non-contact cantilevers (42 N m<sup>-1</sup>, 285 kHz, WITec, Germany). For each surface type one sample was investigated at, five 10 μm × 10 μm spots (approximately corresponding to the cell-substrate contact area), which were scanned with a resolution of 512 px × 512 px and a velocity of 1 s per line. A first order slope correction and a first order line correction were performed on each topography (WITec Project 2.10, WITec, Germany) followed by a 2D FFT filter correction in order to suppress noise (Gwyddion 2.48). The surface roughness was distinguished from the sample waviness by a 2 μm L-filter and the surface roughness parameters were calculated with the software SPIP 6.7.2 (Image Metrology A/S, Denmark). The arithmetical mean deviation of the two-dimensional profile  $S_a$  is defined as:

$$S_a = \frac{1}{MN} \sum_{k=0}^{M-1} \sum_{l=0}^{N-1} |z(x_k, y_l)| \quad (1)$$

where  $M$ ,  $N$  are the length of the 2D image in pixels and  $x$ ,  $y$  and  $z$  the 3D values for each pixel and describes the mean height deviation of all pixels from the mean height of the 2D area.

The peak-to-peak height  $S_z$  describes the height difference between the highest  $z_{\max}$  and lowest  $z_{\min}$  pixel value in the image:

$$S_z = z_{\max} - z_{\min} \quad (2)$$

The error bars of arithmetic mean values of the roughness parameters are based on the standard deviation of five measurements for each sample type and are below 5% and 10% for  $S_a$  and  $S_z$ , respectively.

## 2.3 Cell culture and microscopy imaging

**2.3.1 Cell culture.** For all live cell experiments an axenic *D. discoideum* strain expressing freeGFP (obtained from Dr Günther Gerisch, MPI for Biochemistry, Martinsried, Germany) was used. The cells were cultivated in HL5 medium (Formedium™, Hunstanton, UK) at 21 °C and pH = 6.7, complemented by the antibiotic gentamycin at a concentration of 20 μg mL<sup>-1</sup> (G-418, Biochrom AG, Berlin, Germany). The cell confluence was kept below 40%.

For microscopy experiments, HL5 medium was exchanged by phosphate buffer (PB), adjusted to pH = 6.0. Cell suspension was kept in a polytetrafluoroethylene (PTFE) frame, placed onto

the cured composite layer. Image acquisition was started after at least 45 min resting time to ensure all cells are settled down onto the sample surface.

**2.3.2 Live cell imaging.** The microscopy measurements were carried out on a Nikon Eclipse Ti microscope (Nikon Instruments Inc., Japan). Images were recorded using a 20×, objective with a numerical aperture of 0.7 (Nikon Instruments Inc., Japan) and a 1000 px × 1000 px EM-CCD camera (C9100, Hamamatsu Photonics K.K., Japan) at temperatures of 19–21 °C. Fluorescence and bright field images were acquired every 8 s for 60–90 min with exposure times below 150 ms.

**2.3.3 Image processing and cell tracking.** The fluorescent image time series were employed for the cell migration analysis while the bright field images were used as a control during image processing and analysis, which was carried out using the open source software ImageJ 1.51 h (National Institutes of Health, USA). After binarization, the time series was analysed by the ImageJ plug-in “Cell Evaluator”, which determines and correlates the centre-of-mass for every cell throughout the time sequence yielding time-dependent centre-of-mass trajectories. For migration analysis, only trajectories exceeding 100 time points have been taken into account. To get a statistically reliable data set, each sample type has been measured on at least two different days, for which a new sample with freshly seeded cells was used each time. Every sample was measured twice in different regions with a time interval of several hours. Every time two cell migration time-lapse videos were recorded using the multi-point method. This resulted in at least 100 analysed cells for each sample type (Table S1†).

A high number of cells is important in order to minimize biological related noise. However, even similarly treated and genetically identical cells exhibit a significant and inevitable variability in cell behaviour due to intrinsic biological noise.<sup>51,52</sup>

## 2.4 Cell migration analysis

The obtained cell centre-of-mass trajectories have been evaluated with the MATLAB® (The MathWorks, Inc., USA) algorithm Cell Motion Analysis, which is based on an algorithm designed for intracellular particle motion<sup>53</sup> and was adjusted to distinguish different modes of cell migration.<sup>31,54</sup> For amoeboid migrating cells, like *D. discoideum*, the migration behaviour consists of two distinct types: directed runs with higher velocities



and only small changes in direction in contrast to phases of diffusion-like motion, during which the cell probes its environment. For this study a third classification was implemented additionally, which separates cells that are moving less than a given distance from their starting position over the whole time sequence. These are classified as “non-migrating” and not taken into account for the migration analysis. Here, we used 5  $\mu\text{m}$  as a minimum distance for cells to move, which estimates half of an average cell diameter. The percentage of time points categorised as “non-migrating” are listed in Table S2† for every sample type.

The cell motion analysis algorithm is based on the local mean-squared displacement (LMSD)  $\langle \Delta R^2(\tau) \rangle_i$  where the mean-squared displacement (MSD) is calculated in a rolling time window of  $T$ :

$$\langle \Delta R^2(\tau) \rangle_i = \langle (R(t_i + \tau) - R(t_i))^2 \rangle_{t_i - \frac{T}{2} < t_i < t_i + \frac{T}{2}} \quad (3)$$

where  $R(t_i)$  is the initial cell position and  $R(t_i + \tau)$  the cell position after the passed lag time  $\tau$ . This calculation is done for every time point and the obtained lag time-dependent LMSD is fitted by a power law for every time point.

$$\frac{\langle \Delta R^2(\tau) \rangle_i}{\tau^2} = A_i \left( \frac{\tau}{\tau_0} \right)^{\alpha_i} \quad (4)$$

By choosing  $l = 1 \mu\text{m}$  and  $\tau_0 = 1 \text{ s}$ , the pre-factor  $A_i$  carries no physical dimensions.<sup>55</sup> Depending on the obtained value for  $\alpha$ , every time point is classified as either “directed” for  $1.75 \leq \alpha \leq 2.00$  or “quasi-random” for  $\alpha < 1.75$ . This is based on previous work<sup>31,54</sup> and adjusted for this experiment.

The total of the median values of the alpha and velocity distributions were subjected to a statistical analysis, in comparison to the flat reference surface without particles. Since the median values are not normally distributed, the non-parametric Mann–Whitney test was used. In order to check whether the distributions differ significantly from each other, a Kruskal–Wallis analysis was performed. All calculations were conducted using the software OriginPro 2019 9.6.0.172, (OriginLab Corporation, USA).

For each analyzed cell measurement video, the percentage of active migration phases was determined. From all videos of a sample variation a mean value with corresponding standard deviation was calculated.

### 3 Results and discussion

In this work, we developed a novel type of fabrication process to create tailored surface roughnesses in the nanometre range utilizing particle–polymer composites under exploitation of an oxygen inhibition layer. These nanorough surfaces are made of a biocompatible UV curable composite based on a hybrid-polymer ORMOCER® with incorporated silica particles. The polymer matrix material was successfully used as cell scaffold with different mammalian cell types, like rabbit muscle-derived myogenic stem cells<sup>41</sup> and human microvascular endothelial cells.<sup>43</sup> The influence of the acquired nanoroughness on the

migration behaviour of *D. discoideum* cells has been acquired by time-lapse microscopy and analysed by the local mean-squared displacement of the cell motion.

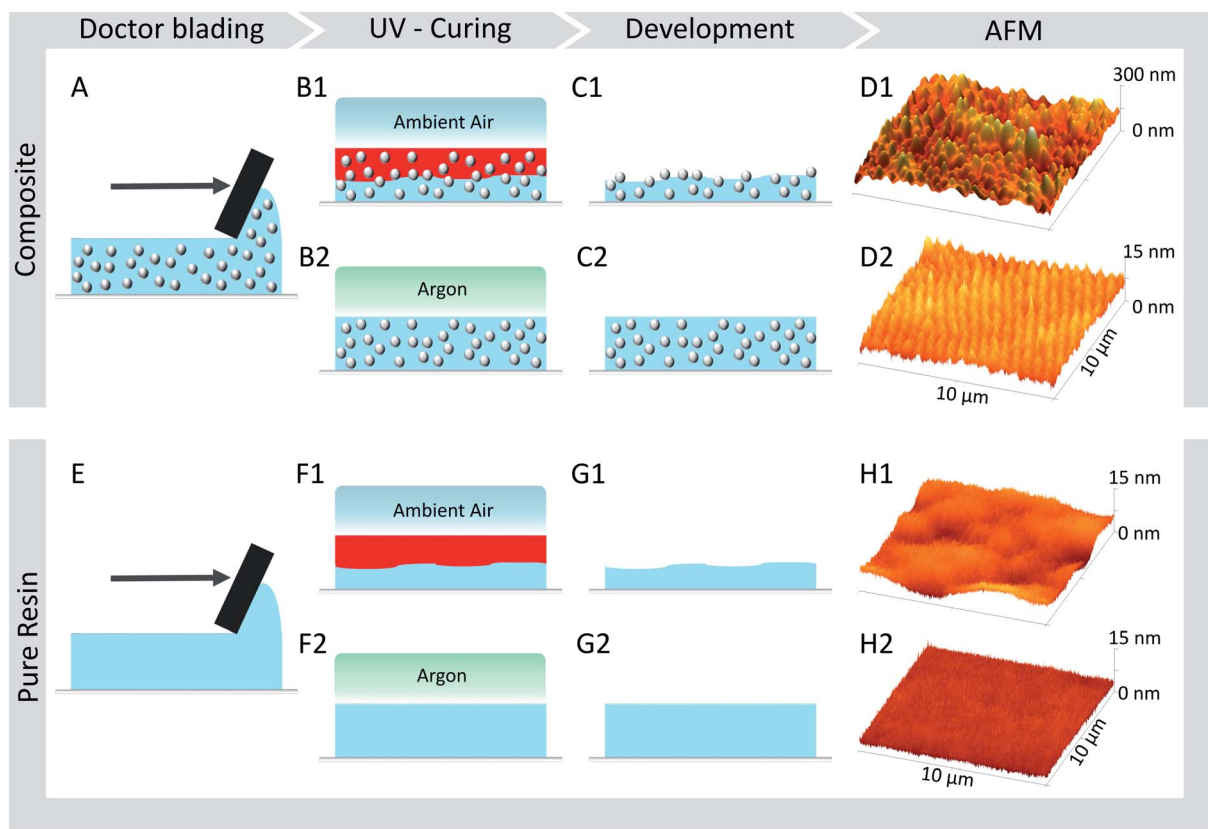
To fabricate the surfaces, the composite (Fig. 1A) or the reference material of pure resin (Fig. 1E) were applied onto a cover slip *via* doctor blading. According to the standard procedure, this is followed by a UV curing step under argon atmosphere (composite: Fig. 1B2, pure resin: Fig. 1F2), resulting in a smooth surface for both surface types, which is confirmed by AFM measurements of the final surfaces (composite: Fig. 1D2, pure resin: Fig. 1H2). The particles (here: 421 nm) are hidden under the polymer surface, proving the excellent surface smoothing properties of the ORMOCER® matrix. In contrast to this standard procedure, we achieve tailored nanorough surfaces by exploiting the fact, that an oxygen related inhibition layer is created at the sample surface when cured under ambient conditions (composite: Fig. 1B1, pure resin: Fig. 1F1). The monomer conversion increases with increasing distance to the air boundary due to the decreasing oxygen concentration. At a given layer depth the degree of polymerization is high enough to endure the development step while the uncured inhibition layer is dissolved (composite: Fig. 1C1, pure resin: Fig. 1G1). Since the particles are not affected by the solvent, they protrude from the polymer surface generating roughness values in the nanometre range (Fig. 1D1).

Tailored surface roughnesses has been achieved by (1) utilizing differently sized particles (marked blue in Fig. 2) and by (2) varying the volume fraction of the particles in the polymeric matrix (marked green in Fig. 2). For (1) three different surface roughnesses – each with a particle volume fraction of 20 vol% – have been manufactured with particles of diameters: 52 nm, 165 nm and 421 nm (see Table 1). To investigate the contribution of the particle volume fraction on the surface roughness, for (2) three different surfaces have been fabricated, each containing particles with a diameter of 165 nm and a varying particle volume fraction of 5 vol%, 10 vol% and, 20 vol%. Flat surfaces made of pure OC-I polymer without any particles serve as a reference. Additionally, two surface types, one of the reference material (OC-I polymer without particles) and one composite of the largest particle diameter (421 nm) have been prepared *via* the standard curing process under argon atmosphere, thus without oxygen inhibition layer. Table 1 lists all prepared surfaces, which are named by the used particle diameter and the particle volume fraction.

#### 3.1 Homogeneity of prepared composites

SEM imaging and image analysis (Fig. 2A–C) were used to verify the quality and particle size distribution of the spherical silica particles. For each particle type, the arithmetic mean diameter was calculated as well as the standard deviation. Particle sizes range from  $(52 \pm 8) \text{ nm}$ ,  $(165 \pm 11) \text{ nm}$  to  $(421 \pm 22) \text{ nm}$ . The particle size distributions were fitted with a log-normal function which reveals a FWHM of 31% in respect to the mean value of 52 nm which drops to 15% and 12% for 165 nm and 421 nm particles, respectively.





**Fig. 1** Work flow of sample production and characterization *via* AFM, for the case of particle filled resin 421–20 (A–D), and pure resin OC-I (E–H). (A and E) A thin material layer is applied on a coverslip *via* doctor blading, which is cured under ambient (B1 and F1) or argon atmosphere (B2 and F2) creating an inhibition layer in the case of ambient air, which is dissolved during a development step (C and G). This results in particles protruding from the sample surface after removal of the inhibition layer (D1) whereas the OC-I is smoothing the surface in the case of UV curing under argon gas (D2) (note the different z-scale). Without particles the OC-I polymer creates smooth surfaces for the standard argon curing (H2), while a waviness in the micrometer regime occurs after development of the inhibition layer when curing under ambient atmosphere (H1).

The OC-I polymer has a silicate (Si–O–Si) backbone and shows a very good processability with the incorporated silica particles. Their highly matched refractive indices lead to a low opacity for a composite material and enables observation *via* optical methods. To obtain surfaces with isotropic roughness, the homogeneity of the particle dispersion is crucial. However, particles tend to agglomerate with decreasing particle diameter. This effect is caused by the increasing surface-to-volume ratio and thus increasing van der Waals- and surface charge forces. Moreover, the surrounding media, adsorbed molecules and other surface interactions are essential for the particle–particle interaction and hence for a stable particle dispersion.<sup>56</sup> To circumvent particle agglomeration, which is enhanced in composites with high filling degree, several particle surface modifications, depending on the dispersion media, were reported.<sup>57,58</sup> To minimize the amount of additional processing steps, we overcame this problem even for a particle volume density up to 20 vol% without any surface modification by choosing an appropriate ORMOCER® matrix with a high inorganic fraction of 25 wt%. Additionally, the absence of any solvents and therefore the high viscosity of the resin suppresses particle diffusion and a similar mass density of particles and matrix leads to a homogeneous particle distribution and

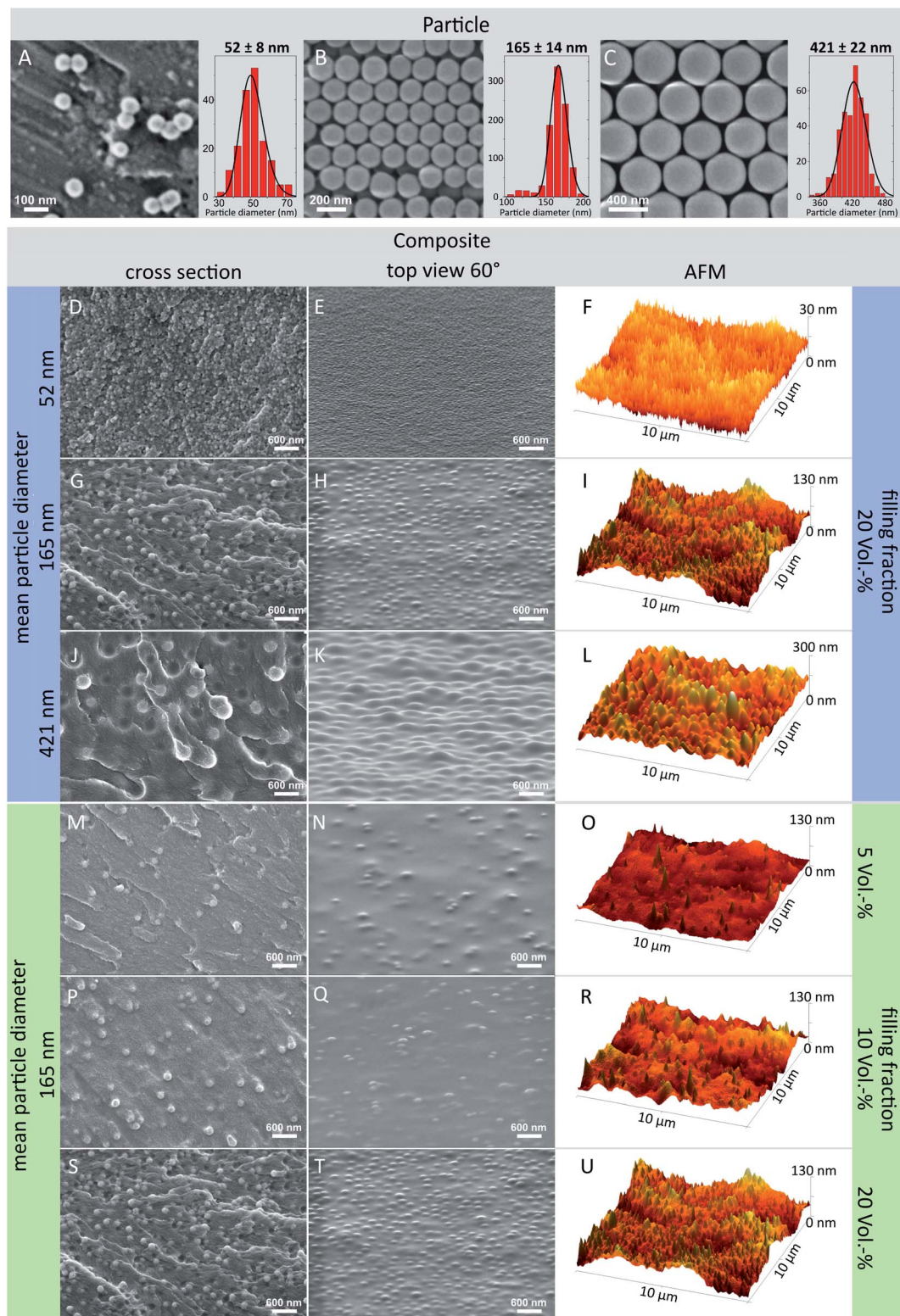
therefore stable composite resin. The particles are inclosed by the matrix material during polymerization, resulting in a homogeneously cured particle–polymer composite.

To verify the homogeneous distribution of the particles across the surface and within the bulk material, SEM investigation of cross sections (Fig. 2D, G, J, M, P and S) and the layer surface (Fig. 2E, H, K, N, Q and T) were performed. For all prepared samples, the particle distribution was random and no agglomeration was visible neither inside nor on top of the composite layers, including for the smallest particles with 52 nm in diameter. Overall, homogeneous surface roughness was achieved for all particle sizes and filling degree variations.

### 3.2 Oxygen inhibition layer and resulting surface roughness

The widely used photoinitiator Irgacure® 369 is a type I initiator and exhibits a high UV absorption with a high free radical initiation rate and therefore short curing times.<sup>59</sup> After excitation *via* photon absorption the initiator molecule decays and forms free radicals, which react with the carbon double bond of the 3-methacryloxypropyltrimethoxysilane (MEMO), which itself attacks another monomer forming a growing polymer radical. Depending on the photoinitiator and the monomer,





**Fig. 2** (A–C) Scanning electron microscopy images of silica particles used for ORMOCER®–particle composites. The insets show the particle size distribution and the particle mean diameters of 52 nm (A), 165 nm (B) and 421 nm (C), which were determined via SEM image analysis. (D–U) Scanning electron microscopy images of the cross section and 60° tilted sample as well as atomic force microscopy images of UV-cured ORMOCER®–silica particle composites, cured under ambient air, after dissolving the inhibition layer. (D–L) Samples with constant particle volume concentration of 20 vol% and increasing particle diameter from 52 nm (D–F), 165 nm (G–I) to 421 nm (J–L). (M–U) Samples with a constant particle diameter of 165 nm and increasing particle volume concentration from 5 vol% (M–O), 10 vol% (P–R) to 20 vol% (S–U).



several side and termination reactions are possible, finally leading to a finish of the chain reaction and a polymerized resin. However, cured under ambient conditions, oxygen molecules are solved inside the resin surface. There, they react with the initiator and growing polymer radicals, forming peroxy radicals, which leads to low monomer conversion even up to an uncured layer. This process depends on the oxygen concentration at its diffusion characteristics inside the resin, on the photoinitiator activation rate and on the chain reaction kinetics.<sup>50</sup>

During the development step the inhibition layer is dissolved while the particles are not affected by the solvent. Therefore, they are washed away if their contact area to the remaining polymerized matrix is not sufficient, while embedded particles remain inside the composite and protrude from the sample surface (Fig. 1D1). During drying, the polymerized matrix material covers the enclosed particles due to the high adhesion between silica particles and siloxane based ORMOCER®. The coating of the particles can also be observed in the SEM images of the surface which exhibit a smoothed cover (Fig. 2E, H, K, N, Q and T) compared to the rough layer cross section (Fig. 2D, G, J, M, P and S). Consequently the surface chemistry is homogeneous and similar to the pure polymer reference as all particles are covered by OC-I but surface roughness is still existent. In the ESI† close-up SEM images of the top (Fig. S3A†) and cross-section (Fig. S3B†) are provided, which illustrate that the particles are completely embedded beneath an ORMOCER® layer. When using polymerization techniques at which no inhibition layer is formed a much more pronounced surface roughness can be accomplished. In this case, however the sample surface is not chemically homogeneous, since the particles are not covered by the polymer. As an example Fig. S3C† shows a 3D structure created *via* direct laser writing, which exhibits a distinct boundary between the particle surface and the ORMOCER® matrix.

We further measured the contact angle of water and ethylene glycol on the composite surfaces with a filling ratio of 20 vol% and calculated the surface energy. We did not find a significant difference in both values compared the flat reference surface (Fig. S4†), which is a strong indicator that the surface chemistry does not change on the different investigated surfaces. Further, the added surface roughness does not significantly change the contact angle. The fabrication process is finished with a temperature treatment inducing a condensation reaction of the unfunctionalized silica particles and the silicate fraction of the hybrid polymer matrix, ensuring a strong particle–matrix interaction.

The Young's modulus of the cell environment can trigger cell behaviour in certain ways. The bulk OC-I exhibits a value of  $(1.01 \pm 0.03)$  GPa, while silica particles are in the order of 45 GPa, depending on their porosity.<sup>60</sup> Both values are beyond those of the natural cell environment<sup>43</sup> and the particles are covered with OC-I. Therefore we might conclude that cell behaviour is not directly affected by the silica particle's Young's modulus.

Due to the inhomogeneous diffusion of the oxygen inside the resin before and during the polymerization step, OC-I cured

under ambient conditions reveals an additional waviness over a length scale of several micrometers (Fig. 1H1). In the case of the particle composite, this surface waviness is superimposed with the nanotopography, determined by the particle radius and volume fraction. In order to distinguish the surface waviness from the particle induced surface features, the AFM data was separated with a  $2\ \mu\text{m}$  L-filter to calculate the surface roughness parameters  $S_a$  (arithmetical mean deviation, Fig. 3A) and  $S_z$  (height difference of the highest and lowest point of the image, Fig. 3B).

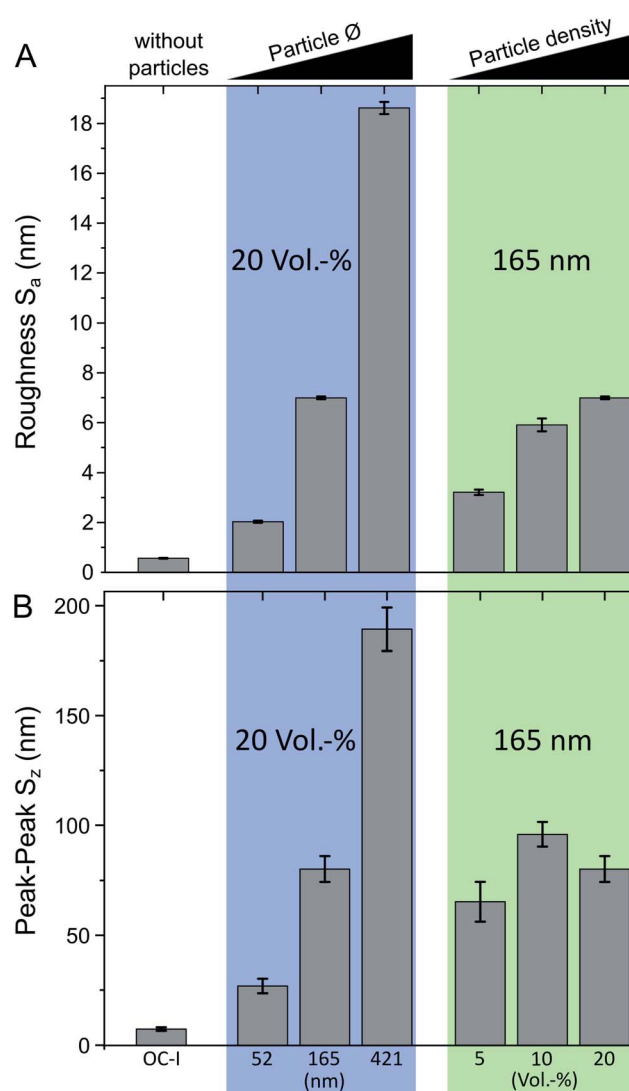


Fig. 3 Surface roughness parameters based on AFM measurements. (A) Arithmetical mean deviation of the assessed 2D profile  $S_a$ , (B) peak-to-peak value of highest and lowest point in 2D profile  $S_z$  (the error bars of arithmetic mean values are based on the standard deviation of five measurements for each sample type). Sample group with a constant particle volume concentration of 20 vol% and increasing particle diameter from 52 nm, 165 nm to 421 nm is highlighted in blue. Sample group with a constant particle diameter of 165 nm and increasing particle volume concentration from 5 vol%, 10 vol% to 20 vol% is highlighted in green.



For both curing cases, under argon and ambient air, the  $S_a$  values of pure OC-I (without particles) just differ slightly with  $S_a(\text{OC-I AR}) = 0.5 \text{ nm}$  and  $S_a(\text{OC-I}) = 0.6 \text{ nm}$ . In the case of standard curing under argon, the particle composite with the largest particle diameter exhibits a  $S_a$  value with  $S_a(421-20 \text{ AR}) = 0.9 \text{ nm}$ , comparable to pure OC-I, indicating that the smoothing properties of ORMOCER® hinder the formation of nanotopography. As expected, the  $S_a$  values increase with the particle size at a constant filling degree of 20 vol% (Fig. 3A, marked blue) from  $S_a(52-20) = 2.0 \text{ nm}$ ,  $S_a(165-20) = 7.0 \text{ nm}$  to  $S_a(421-20) = 18.6 \text{ nm}$ . For a constant particle diameter of 165 nm (Fig. 3A, marked green), the surface roughness can be adjusted *via* the volume fraction as the  $S_a$  value increases with a higher filling degree from  $S_a(165-5) = 3.2 \text{ nm}$  to  $S_a(165-10) = 5.9 \text{ nm}$  to  $S_a(165-20) = 7.0 \text{ nm}$ .

The  $S_z$  value is an indicator for the maximum feature height cells are sensing during migration and for a particle fraction of 20 vol%, it ranges from  $S_z(52-20) = 27 \text{ nm}$  to  $S_z(165-20) = 80 \text{ nm}$  and to  $S_z(421-20) = 189 \text{ nm}$  (Fig. 3B). As these values of the maximum height in the surface profile are in the order of half the respective particle diameter, apparently particles that protrude more than half of the particle diameter from the surface are washed away during the development step. This hypothesis is also affirmed by the  $S_z$  values of the samples with a constant particle diameter of 165 nm, which show no direct correlation of the  $S_z$  value with increasing particle volume fraction. This means that more particles protrude from the sample surface with increasing filling degree and leading to higher  $S_a$  values, but the maximum protrusion height stays constant within a certain range.

### 3.3 Cell migration analysis

To verify the influence of surface topographies in the nanometer range on cell behaviour, we analysed amoeboid migration behaviour of *D. discoideum* cells on such nanorough samples, in comparison to a chemically equivalent flat reference surface. Bright field and fluorescent images of migrating cells were obtained every 8 s for at least one hour per live-cell measurement. The centre-of-mass of every cell was determined at all time points. All cells are exploring their adjacent environment intensely by forming protrusions on all investigated surface topographies (see exemplary Videos S1–S6†). Using a local mean-squared displacement algorithm with a rolling time window,<sup>31</sup> the trajectories of all migrating cells were separated into directed and quasi-random migration states based on the alpha value determined by the power law fit of the LMSD (eqn (4)). During directed migration ( $1.75 \leq \alpha \leq 2.00$ ), cells exhibit a directed and faster motion pattern as compared to the uniform search behaviour during a quasi-random phases ( $\alpha < 1.75$ ). Both cell migration modes occur alternately and can be influenced *via* different external cues.<sup>54</sup> However, compared to many other cell types, for *D. discoideum* the directed motion pattern even occurs without any external gradient or other cues, as amoeboid migration consists of alternating directed and quasi-random phases.<sup>54</sup> For all nanorough surfaces, approximately 30 000 time points of cell migration were acquired in order to assure a statistically reliable data set (Table S1†). Typical migration trajectories of a single cell migrating (Fig. 4A1

and A2) on a pure polymer surface (Fig. 4B1) and on a nanorough composite surface (Fig. 4B2) illustrate that cell migration is strongly confined on the nanorough surface (note the different z-scales in Fig. 4B1 and B2).

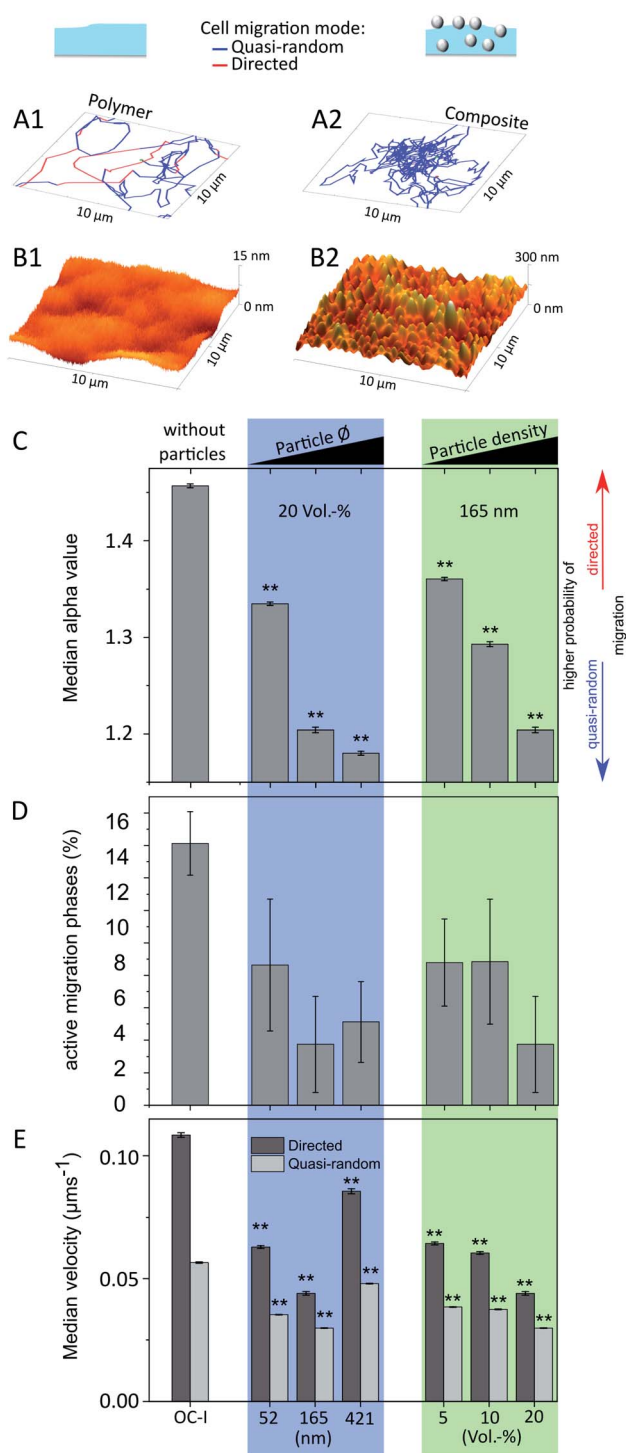
Regarding the surface roughness, there are three intertwined parameters: the particle diameter, the number of particles on the surface per area, and the resulting surface roughness. It is known that cellular behaviour can be influenced by these roughness parameters and cells are able to detect topographies down to 10 nm.<sup>61</sup> Therefore, nanorough surfaces with a constant filling degree and varying particle diameter (marked blue) affect cell migration *via* the mechanical stimuli, which are generated by the density as well as the curvature of the surface features. In the case of a constant particle diameter with varying filling degree (marked green) however, only the feature density changes, which makes it ideal for direct comparison of the cell migration parameters. Estimating this particle density reveals that one particle inside the 421–20 sample, corresponds to 17 in the 165–20 sample and 531 in the 52–20 composite, respectively (see Table S3† for details).

The alpha exponent of the power law fits to the local mean-squared displacement values is a significant parameter to characterize migration behaviour, with values of 1 for a diffusion-like, quasi-random migration behaviour and values of 2 for a straight, directed cell motion. Here, we find lower median values for the alpha exponent for cell migration on all nanorough surfaces (Fig. 4C, for detailed distributions see Fig. S1†) as compared to the migration values on the polymer reference surface without particles. This means that cell migration is hindered on nanorough surfaces. For both sample groups, increasing particle diameter (blue) and increasing particle density (green), a trend towards smaller alpha values is observable with increasing surface roughness. Cell migration on both samples with the highest surface roughnesses 165–20 ( $S_a = 7.0 \text{ nm}$ ) and 421–20 ( $S_a = 18.6 \text{ nm}$ ) yields median alpha values in the diffusive range of  $\alpha = 1.14$  and  $\alpha = 1.12$ , respectively, compared to a much higher median alpha value of  $\alpha = 1.4$  for cell migration on the smooth reference surface.

Regarding the proportion of directed migration phases, it is not possible to establish a correlation among the composite samples, due to its standard deviation (Fig. 4D). Yet, all nanorough samples provoke a significant reduction of the directed migration phases, which drops from more than  $(16 \pm 2)\%$  for the smooth polymer surface down to only  $(4 \pm 2)\%$  and  $(5 \pm 2)\%$  for the roughest surfaces 165–20 and 421–20, respectively. The ratio of directed and quasi-random migration phases depends upon other cues on the mechanical stimulus of the surface and can be altered for instance by aligned surface structures to induce cell guidance. In our case however, the surface features are randomly distributed and might trigger isotropical mechanical cues leading preferably to quasi-random locomotion or even non-migrating cells. A directed motion is therefore interrupted with a much higher probability compared to a surface with no or much less surface features.

Analogously to the median alpha values, the median values of the instantaneous velocities of directed and quasi-random migration states (Fig. 4E, for detailed distributions see





**Fig. 4** Typical migration trajectories (A1 and A2) of *D. discoideum* on pure polymer surfaces (B1) and on nanorough composite surfaces (B2) (note the different z-scale in the AFM images). The migration path is separated in directed (red) and quasi-random migration phases (blue) depending on the alpha value, determined via a local mean-squared displacement analysis, which was used to compare the median alpha value (C) (see eqn (4)), ratio of migration phases classified as active (D) and the cells median velocity (E) on different nanorough surfaces and on a similarly treated pure polymer surface as reference. Sample group with a constant particle volume concentration of 20 vol% and increasing particle diameter from 52 nm, 165 nm to 421 nm is highlighted in blue. Sample group with a constant particle diameter of

Fig. S2†) are significantly lower compared to cell migration on the reference surface. For increasing feature density, e.g. a higher filling degree and constant particle diameter (green), a clear trend towards lower migration velocities is observable. However, for increasing particle size and a constant filling degree (blue) the sample 421–20 does not follow this trend. Its particle diameter is the highest and in the order of the diameter of bacteria (0.2–10  $\mu\text{m}$  (ref. 62)), which is the standard prey of *D. discoideum*,<sup>30</sup> thus other pathways regarding phagocytosis might be triggered and change the migration behaviour severely for this surface type.

Cell-surface interaction of *D. discoideum* is based on van der Waals forces and, among others, the transmembrane protein SadA.<sup>34–36</sup> Both adhesion mechanisms depend strongly on the interaction area between cell body and substrate, which increases with higher surface roughness. This might explain our finding that cell migration is strongly suppressed by nanorough surfaces, since the nanotopography increases the contact area between cell membrane and the substrate, leading to higher overall adhesive forces and suppressing the ability to migrate fast and in a directed way. However, further studies regarding the adhesion mechanism are needed to elucidate these underlying phenomena by immunocytochemistry. For several cell types, an optimum in adhesive strength is observed in order to gain maximal migration rates<sup>63</sup> providing a high adhesion to generate high cell forces as well as enabling easy detachment to move the cell body. These here presented nanorough surfaces can be used to adjust the contact area to optimize cell adhesion and migration or to manufacture heterogeneous samples combining pure resin with composite to create selective areas with fast and slow migration rates.

## 4 Conclusions

In this study, we developed a novel procedure to fabricate tailored surface roughnesses in the nanometre range based on a biocompatible and UV curable particle–polymer composite. This composite consists of the hybrid polymeric matrix ORMOCER® OC-I with stable incorporated silica particles. We demonstrated a tunable surface roughness *via* two different approaches: (1) variation of the particle diameter or (2) by changing the particle filling degree. Nanoroughness was achieved by exploiting the property of the composite matrix to create an oxygen-related polymerization inhibition layer at the surface, when cured under ambient air. After dissolving this uncured inhibition layer, the particles protrude from the surface and yield the nanorough topography. This was analyzed *via* AFM measurements in order to verify the homogeneous particle distribution and to determine the arithmetic mean values of the surface roughness  $S_a$ , which is adjustable from 2.0 nm to 18.6 nm compared to 0.6 nm of the flat reference.

165 nm and increasing particle volume concentration from 5 vol%, 10 vol% to 20 vol% is highlighted in green. Significance level (\*\*:  $p \leq 0.01$ ) is based on non-parametric Mann–Whitney test. Error bars in (C), (E) represents standard error and in (D) standard deviation.



These customizable nanorough surfaces have been investigated influencing the cell migration behaviour of *D. discoideum* cells, which was analysed by a time-resolved local mean-squared displacement analysis. This evaluation allows for the distinction of two amoeboid migration patterns: fast, directed runs versus slow, random-like migration phases. Without exception the migration was hindered on nanorough surfaces compared to smooth reference samples lacking particles. The amount of time spent in directed migration phases, the instantaneous migration velocities, and the alpha exponent, derived from power law fits to the local mean-squared displacement values, were significantly reduced on nanorough surfaces as compared to the flat surface reference. In particular for nanorough surfaces with a constant particle diameter and varying filling degree, a direct correlation between all investigated migration parameters and the particle volume fraction exhibited that increasing surface roughness suppresses cell migration. In this case the alpha exponent can be tuned within a range of 1.4 and 1.1. As a reason for this behaviour, we assume that the randomly allocated particles perturb the cell polarization and thus hinder phases of direct migration. Additionally, the higher contact surface between cell body and substrate leads to higher adhesion forces, which slow down the overall migration velocity.

The manufactured nanorough composite material is photocurable and 3D-structurable with any light-based fabrication method, like lithography or two-photon polymerization. This material class is therefore well suited as base material for hierarchical structures in the order of centimetres and cell-relevant features in the micrometer range with an adjustable surface nanotopography. Besides that, the general incorporation of functionalized nanoparticles to the composite opens up a wide range of tuneable material characteristics in order to generate multifunctional structures and devices as cell scaffolds for lab-on-a-chip or tissue engineering applications.

## Conflicts of interest

There are no conflicts of interest to declare.

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