Bottom-Up Synthesis of Polymeric Micro- and Nanoparticles with Regular Anisotropic Shapes

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INTRODUCTION

The synthesis of shape-anisotropic polymeric nano- and microparticles has been of utmost interest for the development of new functional materials and drug carriers, as well as for various applications in coating, paint production, and building industries. In spite of the high cost and low throughput for many current methods, such particles have demonstrated superior properties that promise creation of hierarchical superstructures. The synthesis procedure has great potential in efficient nanomanufacturing as it can achieve scalable production of the above shapes in a wide range of sizes, with minimum infrastructure and process requirements and little maintenance of the equipment.

There are several major techniques for the synthesis of shape-anisotropic particles. Probably the most common method among them is the emulsion polymerization technique coupled with phase separation. The method is based on emulsification of a monomer or monomers mixture and functional modifications of the particles (e.g., composites with magnetic nanoparticles, oil-soluble additives, etc.). We also describe postsynthetic surface modifications that lead to hierarchical superstructures. The synthesis procedure has great potential in efficient nanomanufacturing as it can achieve scalable production of the above shapes in a wide range of sizes, with minimum infrastructure and process requirements and little maintenance of the equipment.
Modification of this method is the directed precipitation of dissolved polymers via evaporation or, more recently, by means of a directed flow gradient. Commonly, this modification of the method leads to the formation of polymeric fibers with different aspect ratios.

More complex shapes could be obtained via assembly of colloidal particles and physically connecting these assemblies via sintering or growth of an outer shell. Virtually any shape could be obtained upon elaborate planning, provided the Brownian motion of the colloids is negligible. These techniques need either nonscalable manual manipulation or templates to achieve complex shapes.

State-of-the-art techniques for control over both particles shape and dimensions were developed by means of microfluidic devices and various lithography techniques. They all provide exceptional control but unfortunately include drawbacks which hinder their commercialization: (very) low productivity, demanding maintenance procedures, and/or high infrastructure and production costs. There is clearly a need for the development of economically scalable and high throughput methods to make such particles that would open not only the currently envisioned but also many other applications where cost has prohibited the innovative imagination.

Here, we describe a method for bottom-up synthesis of shape-anisotropic, polymeric particles with controlled shape and size. The method is based on monomer drops, which self-shape upon mild cooling and polymerize upon UV irradiation. Because the process is driven by an internal phase transition, it requires very little infrastructure to guide the shape of the droplets. It is scalable with the volume of the emulsion used and compatible with continuous process implementations. The self-shaping allows the deformation of spherical drops into liquid polyhedra; platelets with three, four, and six sides; particles with protruding asperities; and fibrils with different aspect ratios (see Figure 1). The following UV-irradiation in the presence of an initiator permanently fixes the particle shape, which allows us to obtain a palette of anisotropic polymeric particles with regular shape. Here, by regular we mean that the lateral projection of the platelets presents a regular polygon (triangle, tetragon, or hexagon).

The currently debated mechanism of self-shaping most probably includes the combined effect of surfactant surface freezing and subsequent templating of a subsurface rotator phase. Upon cooling, a series of selected surfactants can undergo a two-dimensional surface transition or surface freezing. This freezing initiates the formation of subsurface rotator phase which in turn deforms the liquid drop. The monomer molecules in the rotator phase (aka plastic crystal) have positional order while maintaining rotational freedom around at least one axis. A requirement for the surface freezing and templating of this rotator phase is...
that the surfactant’s hydrophobic tail is to have the same or longer length, compared to that of the oil, or up to 3 hydrocarbon units shorter. \(^{33}\)

So far the rotator phase transition has been identified in alkanes, \(^{32,33}\) alkenes, \(^{33}\) cycloalkanes, \(^{35}\) alcohols, \(^{35}\) light-responsive amines, \(^{39}\) triglycerides, \(^{33}\) and a wide range of multi-component mixtures. \(^{35}\) Mixtures might include two or more components that both yield rotator phases or a combination of one substance that forms a rotator phase and one that does not. The main requirement for the selection of mixtures is that the substance with higher melting temperature is more than 15 vol % and forms a rotator phase. \(^{35}\)

The method allows scalable and efficient bottom-up production of these shapes in batch and continuous modes, with minimum equipment requirements and maintenance. \(^{32,33}\) Major benefits of the method include the lack of external constraints to create a shape and utilization of all the material in a droplet.

### EXPERIMENTAL SECTION

**Materials.** All chemicals used in this study were purchased from Sigma-Aldrich. In most experiments, we used stearyl methacrylate (SMA) for preparation of emulsions and polymerization. SMA has purity of \(\geq 98.5\%\) and up to 10.5% hexadecyl methacrylate admixture. In a separate series of experiments, we used a series of alkyl acrylates with 14, 16, and 18 carbon atoms in the alkyl chain as the main phase in the polymerizing drops. Tetradecyl and hexadecyl acrylate were purchased from TCI and had \(>95\%\) and \(>90\%\) purity, respectively, whereas stearyl acrylate was purchased from Sigma-Aldrich and had \(>97\%\) purity.

The nonionic surfactants Tween 40 and Tween 60 were used for stabilization of the emulsions and to assist the droplet self-shaping. According to the manufacturer, Tween 40 has a polyoxyethylene (20) sorbitan head with a mixture of C16 palmitic (~90%) and C18 stearic tails (~10%), whereas Tween 60 has 40–60% stearic acid tails and a total concentration of stearic and palmitic acid of \(\geq 90\%\). Both Tweens were dissolved (at e.g. 1.5 wt %) in deionized water (Elix module 5, Millipore).

\(\alpha\)-Ketoglutaric acid (\(\geq 99.0\%\)) was used for a photoinitiation of the monomer polymerization in the emulsion drops. Iron oxide (Fe\(_3\)O\(_4\)) magnetic nanoparticles solution with 20 nm average particle size and concentration of 5 mg/mL in toluene was purchased from Sigma-Aldrich (CAS 700304) and used for magnetic functionalization of the particles.

**Emulsion Preparation.** Micrometer-sized monodisperse emulsions were prepared using an Internal Pressure Microkit in buoyancy operated mode (SPG Technology Co., Ltd., Japan, http://www.spgrhochno.co.jp/english/product/internal.shtml). We used hydrophilic cylindrical glass membranes with pore sizes of 1, 3, and 10 μm for preparation of drops with average diameters 3, 10, and 33 μm, respectively. We dissolved 1.5 wt % surfactant in the water phase and started generating oil drops at the lowest transmembrane pressure (at which drop generation started) to yield monodisperse drops. Because we generated relatively large drops (which creamed) and because we did not need many drops for the capillaries, we generated emulsions with at least 1 vol % oil (SMA).

Submicrometer emulsion droplets were prepared with high-pressure homogenizer (PandaPLUS 2000, GEA). 1.5 g of SMA was dispersed in 148.5 g of 1.5 wt % Tween 40 solution. The emulsion was passed five times through the high-pressure homogenizer at 100 bar. The drop size distribution by volume and number is provided in Supporting Information section 6.

All emulsions were used within 2–3 weeks of their preparation and stored at temperatures around 20 °C.

**Shape Formation and Polymerization.** In a 1.5 mL Eppendorf tube, we mixed 0.9 mL of KGA (2.5 wt %), 0.1 mL of surfactant solution (1.5 wt %), and 0.05 mL of premade SMA emulsion (1 vol % or more). This premix was placed in a rectangular capillary, in an isothermally adjusted chamber with precise temperature control (±0.2 °C). \(^{32}\) Drop deformations were observed via optical microscopy in transmitted, cross-polarized white light (Axioplan, Zeiss, Germany). After self-shaping of the particles in the desired shapes, the initiator was activated via UV diode illumination (LTP-L034UVH365, Liteon), set at \(\leq 600\) mW. The illumination was held for 30 or 120 min.

**Surface Modification of the Particles.** To enable the surface modification of the particles, we used the following procedure: In a 1.5 mL Eppendorf tube, we mixed 0.5 mL of 5 wt % KGA, 0.5 mL of 10 wt % Tween 40 solution, and 0.05 mL of premade SMA emulsion (1 vol % or more). Afterward, we polymerized the particles with UV-irradiation and left them at a constant temperature for several hours.

**Composite Particles Formation.** 0.5 g of suspension of Fe\(_3\)O\(_4\)-in-toluene was dispersed in 2.1 g of SMA. Afterward, toluene was evaporated yielding 0.12 wt % iron oxide, dispersed in the SMA. The Fe\(_3\)O\(_4\)-in-suspension was emulsified via membrane emulsification or ultrasound homogenization and then used for self-shaping and polymerization. The ultrasonic homogenization was made in the following way: 20 mL of emulsion containing 1 vol % SMA in 1.5 wt % Tween 40 was prepared via handshaking. This premix was then homogenized for 10 s at 400 W, using ultrasonic homogenizer (SKL650-HDN, Ningbo Haishu Sklon Development Co.).

**Dynamic Light Scattering (DLS).** The sizes of the submicrometer liquid drops and a reference sample of fibrilar particles were determined by using Malvern Zetasizer Nano ZS. The measurements were performed with \(\lambda = 633\) nm laser in backscattering mode at 173° scattering angle. The equipment was set to fully automated mode for measuring the hydrodynamic radius of the nanoemulsion drops, whereas the measurement time was set manually to 10 s per measurement for the fibrilar particles, and 16 measurements were performed per point. The apparent diffusion coefficient was measured, and the size of the rods was calculated as explained in Supporting Information section 6.

**Electron Microscopy.** Scanning electron microscopy (SEM, Tescan Lyra3, Czech Republic) was used for imaging the polymeric particles larger than 300 nm. Transmission electron microscopy (Jeol JEM-2100, USA) was used for measuring the particle sizes below 300 nm and to visualize the dispersed Fe\(_3\)O\(_4\) particles in micrometer-sized fibrils. In both cases, the polymerized suspensions were processed for imaging immediately after their preparation. For SEM imaging we blew out the polymerized suspensions from the capillaries onto porous, cellulose filters (with nominal pore size 400 nm). The surfactant and some of the smaller particles were absorbed within the filter pores, whereas larger particles remained on top of the filters. Afterward, the particles were rinsed with a few drops of deionized water to remove the excess of surfactant. In the end, the particles were left to dry for at least 24 h on the cellulose filters (in a clean closed box to avoid contamination with dust).

Samples for SEM were sputter-coated with 5–10 nm gold layer for imaging, whereas samples for TEM were prepared via scratching the porous filters with a spatula over a carbon-taped grid.

### RESULTS AND DISCUSSION

We first demonstrate the shaping and polymerization reaction with monodisperse emulsions of stearyl methacrylate (SMA) in an aqueous solution of Tween 40 surfactant, also containing \(\alpha\)-ketoglutaric acid as a photoinitiator. The cooling cell was connected to a cryo-thermostat, and the temperature was set to 18.9 °C. After the temperature of the cooling chamber was stabilized at 18.9 °C, the glass capillary containing the emulsion sample was carefully inserted into the cooling cell. Thus, the emulsion was subjected to a rapid cooling from room temperature down to 18.9 °C without temperature overshooting, and it was maintained at this temperature for a long period, during which the droplets undergo shape changes...
First, the spherical droplets deformed to polyhedra and then to hexagonal platelets. Afterward, they gradually changed their shape into a mixture of triangular platelets, tetragonal platelets, and rods or long fibers, the relative ratio of which could be controlled by the waiting time at the transformation temperature. For example, after 2 min of maintaining 18.9 °C, we obtained predominantly hexagonal particles, after 15 min triangular particles, and after 1 h fibrils. To polymerize the desired shape, we switched on a UV LED (365 nm), which triggered the reaction in the drop. Representative images of polymerized particles are shown (Figure 1, Stage 3). The shapes and yield of polymerized SMA particles could be controlled via changing the surfactant for the emulsion and the size of the initial drops (Supporting Information section 1).

The temperature of transformation at which the emulsions are held was selected to ensure self-shaping of more than 90% of the drops in the emulsion at relatively low rate, thus allowing us to polymerize them in the desired shape along their evolution. At lower temperatures (e.g., 18 °C) the self-shaping process is faster and more polydisperse shapes are obtained for the platelets, but the transition to fibers is reached in just a few minutes. At even lower temperatures, e.g., 16 °C, some of the drops freeze before we can polymerize them.

To demonstrate the versatility of the method, we show that we can control not only the shape but also particle size and chemical composition. Polymer shapes were obtained from three other polymerizable monomers (tetra-, hexa-, and octadecyl acrylates). We found conditions where in the presence of Tween 60 surfactant and temperatures near the melting point of the oil the droplets easily evolve in shape (e.g., Supporting Information section 2). We discuss the effects of monomer differences in the polymerization process.

In addition to changing the polymer chemistry, we found that the drop self-shaping method is compatible with the incorporation of nanoparticles in the oil phase. This greatly extends the opportunities for synthesis of multifunctional composite micro- and nanoparticles with controlled shape (Figure 2). We show that by adding 0.12 wt % 20 nm Fe₃O₄ particles to SMA before emulsion preparation we can follow the same procedure for self-shaping pure SMA and obtain composite polymer particles with various shapes (Figure 2a).

TEM images show the incorporation of the particles throughout the composites, and we also demonstrate the magnetization and movement of hexagonal platelets in an external magnetic field (Video SV2). On the basis of our recent work with multicomponent oil droplets (see ref 35), we expect the process would allow for incorporation of other oils, mixed with the polymerizable monomer. Indeed, we successfully incorporated a wide range of oily additives (as shown in the Supporting Information section 4), demonstrating great potential to tune multiple chemical functionalities for applications, such as encapsulation, drug delivery, bio- and mechanical sensing, and mechanical reinforcement. The compounds tested here are mainly oil-miscible monomers that could copolymerize with SMA (methacrylic acid, vinyl acetate, and 2-ethylhexyl methacrylate), cross-linking agents (trimethylolpropane methacrylate and divinylbenzene) which give rise to polymer networks, and oil-soluble initiator...
Components that were initially dissolved in the oil phase but subsequently partition between oil and water were also used for particle modification—methacrylic acid was copolymerized with SMA, and ethylene glycol diacrylate was grafted on the surface of SA particles. The latter made the particles more rigid during heating at short time polymerizations due to cross-linking, wherein the core of the droplets remained liquid. The sizes of polymer particles fabricated by this method range from ~50 nm to ~1 mm scales, with various elongation ratios, L/2R (L being the longest path along the particle; see Figure 2A). For hexagons L/2R is between 2 and 4, for triangles between 2.5 and 5, and for fibers with initial radius R μm between 3 and 80 (Figure 2b). The average thickness of the platelets could be calculated based on the elongation of the drops, when volume conservation is assumed (e.g., negligible shrinkage and no breakage during deformation and polymerization). The thickness was estimated to be 1.56 ± 0.5 μm for hexagonal platelets, 1.5 ± 0.25 μm for the triangular platelets, and around 1.0 μm for fibers, upon longer elongation times. Therefore, the average ratios between the longest length × height for drops with R = 5 μm are approximately 23 μm × 1.6 μm for hexagons, 28 μm × 1.5 μm for triangular platelets, and between 16 μm × 7 μm and 800 × 1 μm for fibers, depending on the waiting time (larger aspect ratios are obtained at longer times).

The initial droplet size was previously shown to influence the kinetics and distribution of shapes obtained from a particular oil. Previous optical measurements had also given some evidence near the diffraction limit that smaller droplets gave rise to smaller fiber diameters. Having polymerization allowed us to study this effect using electron microscopy.

To clarify these effects for stearyl methacrylate (SMA) and to test the ultimate limits of the self-shaping method, we prepared a series of emulsions with drop radii ranging between 50 nm and 5 μm and polymerized them (see Supporting Information section S7). We studied quantitatively the sphere-to-fiber transition for drops of initial radius R. Figure 2a shows their elongation (with correspondent thinning due to volume conservation) up to very long times, into cylindrical fiber structures of length L and diameter 2r (aspect ratio L/2r). The shapes could be fixed and isolated at any stage by starting UV polymerization, and the fibril length could be controlled over more than 3 orders of magnitude (~250 nm to 800 μm).

Indeed, we found that smaller initial drops produced somewhat smaller in diameter fibers, but also the diameter 2r (50–1000 nm) continues to decrease (aspect ratio increase) with time spent in cooling deformation before polymerization. The sharp edges of the ends of the thinnest fibrils obtained show they are likely fragments of a longer polymer fiber.

The ~50 nm diameter of the fiber, observed in Figure 2b-i, sets experimentally the lower limit for the thickness of the rotator phase. Note that this thickness is somewhat smaller than the initially anticipated one and could depend significantly on the specific system studied.

We investigated several other effects and processes that shed light on potential ways to control and improve such bottom-up polymerization of particle shapes. The method is sensitive to molecular rearrangements during polymerization, and we observe differences between polymerizations of different monomers. For example, SMA emulsions yielded regular polymer particles, whereas acrylates yielded similar shape changes at first, but during polymerization, they exhibited various degrees of distortion. This is likely due to the coupling of volumetric shrinkage stresses developed along the polymerization and the much thinner plate geometries in the acrylate samples. Among the three acrylates, the faster rotational diffusion and reorientation ability of the smallest molecules (tetradecyl acrylate) resulted in the best preservation of the shapes in particles, as illustrated in Figure S3 of Supporting Information section 2, pointing out a potential route for optimization of regular shapes, by utilizing a series of monomers. Because of the elastic stresses, all the particles developed surface wrinkles during polymerization, perturbing the (initially smooth) particle surface.

Highly corrugated particles are sometimes desirable for adjusting rheological, adsorption, and adhesion properties in colloids. We found we could achieve another level of structural hierarchy by controlling a postsynthetic separate process for modifying the surface, which as to our knowledge has not been observed until now (see Figure 3).

Figure 3. (a) Morphological modification of particles surface: (a) Case of single anisotropic particle and hypothetical mechanism. Scale bars are 5 μm. (b) Kinetics of spikes growth in the presence of higher surfactant concentration and spherical particles, 5 wt % Tween 40, 2.5 wt % ketoglutaric acid, and 0.05 wt % SMA. The scale bars are 20 μm.

A “crown” of spikes was observed to gradually grow on the particle surface when particles were left to polymerize for a long time (Figure 3a). The spike length and surface density increased with time until the whole surface of the particles was covered by a dense layer of micrometer long asperities. We hypothesized this behavior may be related to surfactant–monomer aggregates (mixed micelles) formed in the aqueous phase. The monomers polymerize in the interior of the micelles, forming nanoclusters of poly-SMA, and later attach to the already polymerized micrometer particles. We tested this hypothesis by increasing the concentration of Tween 40 surfactant in the aqueous phase to 5 and 10 wt %, to increase the amount of monomers solubilized in micelles. Afterward, we polymerized the emulsion drops at 33 °C. As a result of the small size of the poly-SMA clusters formed in the micelles (e.g., in the range between 3 and 12 nm; see Supporting Information section S5), these clusters started self-assembling into large spikes upon supercooling to 18.9 °C. The size of the spikes increased with the increase of surfactant concentration, and their growth was faster at higher surfactant concentrations (see Figure 3b and Supporting Information section S5). The latter trends support our hypothesis that the spikes are formed by
self-assembly of nanoclusters of poly-SMA, formed inside the surfactant micelles.

**CONCLUSIONS**

We have introduced a novel bottom-up method for synthesizing nanometer to millimeter size particles of a variety of regular anisotropic shapes, sizes, and chemical compositions. Drops of reactive monomers were shaped into desired structures and polymerized to fix the shape permanently by UV-irradiation. We outline the factors for controlling the yield of different shapes (emulsion surfactant, initial droplet size, and time in cooling stage). Regularly shaped hexagonal and triangular platelets and fibrilar particles were produced from droplets as small as 50 nm and with control over size from 50 nm to 1 mm in at least one of their dimensions.

Functional composite particles were obtained by incorporation of (e.g., magnetic) nanoparticles in the monomer oil droplets and by copolymerizing mixtures of reactive oil monomers (e.g., with cross-linking agents), allowing chemical customization of the shaped particles. Postsynthetic modification of the particles surface could also be attained via the reported self-assembly of nanocrystallites or by the classical methods of polymer and/or surfactant adsorption. Such modifications could be used to tune the particle interactions with other particle/surfaces, opening the prospect for self-assembly of larger, macroscopic hierarchical structures.

**ASSOCIATED CONTENT**

* Supporting Information
The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.macromol.8b00529.

Additional experimental results for the preparation of anisotropic particles with different surfactants and different drops sizes, examples of polymerization of particles with different compositions, as well as modification of particle surface morphology (PDF) Video SV1: self-shaping of particles presented in Figure 1 (AVI)
Video SV2: hexagonal platelet movement in a magnetic field (AVI)
Video SV3: reversible modification of particle surface morphology upon heating (AVI)

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Notes
The authors declare no competing financial interest.

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