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COLLOIDAL ORGANOSILICA SPHERES THROUGH SPONTANEOUS EMULSIFICATION

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Abstract

Colloidal particles of controlled size are promising building blocks for self-assembly of functional materials. Here, we systematically study a method to synthesize monodisperse, micrometer-sized spheres from 3-(trimethoxysilyl)propyl methacrylate (TPM) in a bench-top experiment. Their ease of preparation, smoothness, and physical properties provide distinct advantages over other widely employed materials such as silica, polystyrene, and poly(methyl methacrylate). We describe that the spontaneous emulsification of TPM droplets in water is caused by base-catalysed hydrolysis, self-condensation, and deprotonation of TPM. By studying the time-dependent size evolution, we find that the droplet size increases without any detectable secondary nucleation. Resulting TPM droplets are polymerized to form solid particles. The particle diameter can be controlled in the range of 0.4–2.8 μm by adjusting the volume fraction of added monomer and the pH of the solution. Droplets can be grown to diameters of up to 4 μm by adding TPM monomer after the initial emulsification. Additionally, we characterize various physical parameters of the TPM particles and we describe methods to incorporate several fluorescent dyes.

7.1 Introduction

Colloidal particles are widely used in industrial applications and also serve as model systems for fundamental studies, such as crystallization,^{47,198} the packing of small clusters,⁷¹ and the glass transition.⁵³ Many of these applications rely on the availability of spherical particles that have an accurately controlled size: for instance, crystallization requires spheres with a polydispersity below 6%.¹⁹⁹ Monodisperse colloidal spheres can also be employed as basic building blocks for fabricating more complex anisotropic structures, such as lock-and-key particles, colloidal molecules, and Janus particles.^{43,45,50,200}

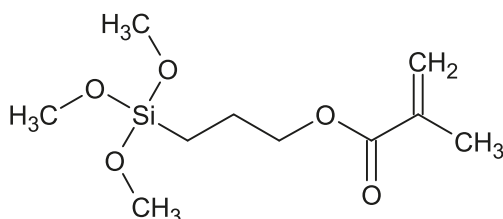


Figure 7.1. Structural formula of 3-(trimethoxysilyl)propyl methacrylate (TPM).

Therefore, the synthesis of micrometer-sized monodisperse spheres has become an indispensable technique in colloidal science. Synthesis protocols for a select range of widely used materials are well established, in particular for silica,²⁰¹ polystyrene,²⁰² and poly(methyl methacrylate).²⁰³ Each of these materials has its own advantages: for example, silane chemistry allows for a wide range of possible surface coatings, polystyrene particles can be density matched with water in order to exclude gravitational effects, and poly(methyl methacrylate) particles can be refractive index and density matched in an organic solvent, enabling accurate three-dimensional confocal imaging.

Despite the well-documented and mostly straightforward synthesis methods, many scientists refrain from synthesizing these particles themselves due to the basic chemistry infrastructure that is required. Here, we study a bench-top preparation of monodisperse colloidal particles that can be implemented with as little as a stirred beaker and thus is readily accessible to a wide range of scientists. This synthesis method employs the alkoxy silane 3-(trimethoxysilyl)propyl methacrylate (TPM, see Figure 7.1). The advantage of this material is threefold: First, the preparation can be performed without extensive chemical equipment and can be completed within four hours. Second, we show that tuning the available synthesis parameters allows control over the particle size from 0.45–1.70 μm . Third, the organosilica material of the resulting colloidal spheres has the chemical versatility of silica, while the material is stable in water, its density is lower than that of silica, and its refractive index is similar to that of poly(methyl methacrylate). This combination of features opens up new experimental possibilities, for instance the manipulation and three-dimensional tracking of dense suspensions²⁰⁴.

The preparation of TPM spheres begins with the spontaneous formation of TPM emulsion droplets in water. This phenomenon has been described previously for different alkoxy silanes in water under alkaline conditions.²⁰⁵ Since TPM can be polymerized at its methacrylate moiety, these emulsion droplets can also be used to fabricate solid colloidal particles, which has been used in the past to image Pickering emulsions^{51,206} as well as to make complex and anisotropic particles.^{50,200,204} However, the preparation method of TPM spheres has not been studied systematically to date. In order to provide a reference for the TPM microparticle preparation, we here first review previous studies on the emulsification of trialkoxy silanes. Then we summarize the preparation method and several ways of including fluorescent dyes, measurements of material properties, a time-resolved measurement of the synthesis, and a study of the effect of various synthesis parameters. Finally, we describe and characterize a seeded growth method for creating TPM droplets with sizes of up to 4 μm .

7.1.1 Review of Previous Work

Obey and Vincent [205] first showed that a monodisperse emulsion can be formed spontaneously by mixing an alkoxy silane, dimethyldiethoxy silane, with water or ethanol mixtures at alkaline conditions. They proposed a mechanism that required the base-catalysed nucleation and growth of poly-alkoxy silane in water from an initial single-phase system. The emulsions formed were thought to be kinetically stabilized against aggregation by negative charges on the droplet interface.

The same method was adapted to create emulsions of TPM stabilized by various inorganic nanoparticles and tetramethylammonium hydroxide (TMAH) ions, so-called Pickering emulsions.^{206,207} Stable emulsion droplets of diameters ranging from 10–100 nm were observed in the presence of these nanoparticles. For Pickering emulsions stabilized by magnetite particles,²⁰⁶ the droplet size increased with increasing TPM volume fraction and decreased when more of the magnetite particles were added at constant TPM volume fraction. The emulsion is also formed when the TPM is mixed with other monomers such as methyl methacrylate and styrene, indicating that only relatively small amounts of TPM are needed to stabilize the droplets.²⁰⁸

Later it was found that TPM emulsions can also be obtained in the absence of nanoparticles.^{50,51} These emulsions had diameters of approximately 1 μm , which is much larger than the Pickering emulsions. The initially polydisperse emulsions were formed after 1 day using a mixture of TPM, water, and various quaternary ammonium salts at pH 9 to 10.⁵¹ By adding pre-hydrolysed TPM drop-wise to an ammonia solution until the droplets reached their target size, Sacanna et al. [50] succeeded in making these TPM-in-water emulsions monodisperse. The advantage of using TPM is that these droplets could then be polymerized using a radical initiator, which resulted in solid dimpled spheres if the polymerization was initiated from the water phase. Furthermore, the droplets could be fluorescently labelled by the addition of a rhodamine dye covalently linked to the alkoxy silane APS (3-aminopropyl triethoxy silane) after droplet formation.²⁰⁰

negatively charged.²¹⁷ Sacanna et al. [214] proposed a mechanism where the TPM will produce its own charged surfactant consisting of deprotonated TPM at the TPM/water interface which could contribute to the stabilization of the emulsion droplets. By analysing bulk TPM/water surface tensions, Kraft et al. [218] found that emulsions are formed if the bulk surface tension is below 10 mN m^{-1} . Below this surface tension, an increasingly larger droplet surface area is created with decreasing surface tension. Increasing the concentration of hydrolysed TPM lowers the interfacial tension, pointing again toward the stabilizing properties of deprotonated TPM molecules.

In this study, we describe a preparation method of micrometer-sized TPM spheres as well as methods of including several fluorescent dyes. Then we characterize the material properties of TPM spheres. To further understand the spontaneous emulsification of TPM, we then study the time evolution of the TPM droplets during a synthesis as well as the effects of various synthesis parameters on the particle size distribution. Finally, we describe a seeded growth method to obtain particles with diameters of up to $4 \mu\text{m}$.

7.2 Methods

Materials 3-(Trimethoxysilyl)propyl methacrylate (TPM, 98 %), ammonium hydroxide (28 % NH_3 in water), tetramethylammonium hydroxide (TMAH, 25 % in water), azobis(isobutyronitrile) (AIBN, ≥ 98 %), potassium peroxydisulfate (KPS, ≥ 99 %), 2-hydroxy-2-methylpropiophenone (≥ 97 %), rhodamine B isothiocyanate (RITC, mixed isomers), fluorescein isothiocyanate isomer I (FITC, ≥ 90 %), 3-(aminopropyl)triethoxysilane (APS, ≥ 98 %), dimethyl sulfoxide (DMSO, ≥ 99.9 %), 1,3,5,7-tetramethyl-8-phenyl-4,4-difluoroboradiazaindacene (BODIPY, 97 %), perylene (≥ 99.5 %), pyridine (≥ 99.9 %), and 2-ethylhexyl 4-methoxycinnamate (98 %) were purchased from Sigma-Aldrich and used as received. Sodium hydroxide (NaOH, 98.5 %) was obtained from Acros Organics. All solutions were prepared from deionized water with $18.2 \text{ M}\Omega \text{ cm}$ resistivity, using a Millipore Filtration System (Milli-Q Gradient A10).

Emulsification To prepare TPM emulsions in water, we used ammonium hydroxide as a catalyst. 15 mL of ammonia solution of 1–1000 mM was added to a 100 mL plastic (polypropylene) beaker with a PTFE coated stir bar. The use of glassware was avoided as TPM reacts with the glass surface²¹⁰ and leaves visible patches that cannot be cleaned easily (see Fig. 7.3). The pH of the ammonia solution was measured using a HACH HQ40d pH meter with a calibrated IntelliCAL PHC101 standard gel filled pH electrode. pH measurements were not performed after the addition of TPM because we suspect TPM reacts with the glass surface of the pH probe. Throughout, the beaker was covered with Parafilm to prevent pH change due to ammonia evaporation. While the solution was stirred at 350 rpm (medium-sized vortex), an accurate volume of TPM monomer was injected with a plastic pipette. During the first 20 min of the reaction, the solution gradually becomes more turbid and the TPM droplets observed at the start of the reaction

disappear. At this point, the stirring speed was lowered and stirring continued at 200 rpm until a total stirring time of 2.5 h.

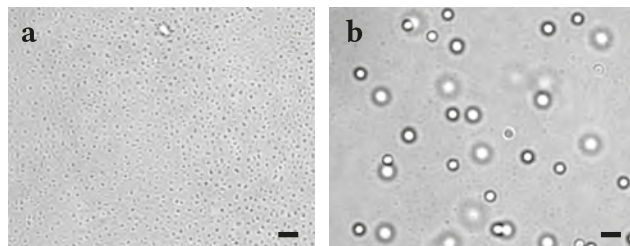


Figure 7.3. Bright field images of a glass cover slide containing the reaction mixture 10 min (a) and 45 min (b) after TPM injection. TPM reacts to glass cover slides, resulting in a visible speckle pattern. In (b), the TPM emulsion itself is also visible. Samples that were taken after 60 min of reaction did not develop this pattern.

Radical polymerization After the spontaneous emulsification of TPM, the emulsion droplets can be polymerized in a radical polymerization reaction. During this process, the methacrylate moieties of the TPM polymerize, yielding a rigid particle that can be stored at room temperature for years.

To polymerize a TPM emulsion, we transferred the emulsion to a capped vial and added 10 mg of the radical initiator AIBN per 15 mL of the emulsion. To initiate polymerization, the solution was heated to 80 °C for 2 h. To prevent sedimentation, the vial was tumbled by hand at 30 min intervals. In the case in which droplets sedimented too quickly, leading to coalescence and a polydisperse dispersion, polymerization in an oil bath under continuous tumbling was employed.

Alternatively, 0.5 mM potassium persulfate (KPS) can be used to initiate polymerization. Similar to AIBN, polymerization of the TPM droplets with KPS at 80 °C for 2 h yields spherical and solid colloids. Here we note that KPS can also yield dimpled spheres through shell buckling.²¹⁹ If heating the sample during polymerization is undesirable, then a photo-initiator (0.1 vol % 2-hydroxy-2-methylpropiophenone) can be employed. After illumination with ultraviolet light (a Rayonet photochemical reactor operating at 350 nm wavelength) for 1.5 h, rigid particles are obtained. See Figure 7.9 on page 116 for SEM pictures of colloidal particles prepared with these alternative initiators.

To corroborate the radical polymerization mechanism, we monitored the change in the FTIR spectrum in samples of emulsified TPM droplets and polymerized TPM spheres (see Fig. 7.11 on page 117). We found that the carbon-carbon double-bond vibration decreased by 63 % after polymerization. Therefore, we estimate the unreacted monomer fraction of methacrylate moieties at 37 %.

Fluorescent dyes To enable fluorescence microscopy of the TPM particles, several fluorophores are available for integration into the particles. All dyes were added after droplet formation. The dispersion was stirred for an additional 15 min to ensure a good distribution throughout the droplets. Then the emulsion was polymerized as described in the previous section.

Silane chemistry was used to covalently link the dyes to the organosilica network of TPM.^{220,221} For example, RITC (576 nm emission) and FITC (525 nm emission) can be covalently attached to the amino group in 3-(aminopropyl)triethoxysilane (APS), as follows: 10.2 mg RITC and 8.6 μL of APS were dissolved in 10 mL of DMSO and stirred overnight. Typically, 10 μL of this RITC-APS solution was added per 1 mL of TPM emulsion. The dye solution can be stored at room temperature under light exclusion for months without degradation.

FITC is linked to APS as follows: 3 mg FITC and 22.5 μL APS were dissolved in 250 μL ethanol and stirred overnight. Typically, 2.3 μL of this FITC-APS solution was added per 1 mL of TPM emulsion. For long-term storage, water must be excluded to prevent hydrolysis and self-condensation of the APS molecules.²²⁰

Because of the hydrophobic nature of TPM, it is possible to also include hydrophobic dyes: both BODIPY (508 nm emission, also available in other colours) and perylene (447 nm emission) were dissolved at 1 g L^{-1} in DMSO. Typically, 10 μL of one of these dye solutions is added per 1 mL of TPM emulsion in order to obtain fluorescent particles. These non-covalently incorporated dyes migrate from the particles when they are transferred to a hydrophobic solvent.

Imaging methods TPM emulsions were imaged using a Nikon Eclipse Ti microscope with a 100 \times oil-immersion objective (NA=1.4). Dried samples of polymerized particles were sputter coated with Pt/Pd and imaged using a FEI nanoSEM 200 scanning electron microscope (SEM) using an acceleration voltage of 15 kV. The particles obtained through the seeded growth process were imaged on a Zeiss Merlin field emission SEM using an acceleration voltage of 1 kV. The SEM images were analysed using ImageJ software and a Python algorithm²²² to determine the particle size distribution.

7.3 Results and Discussion

7.3.1 Physical properties

This section summarizes some physical properties of TPM. The density and refractive index measurements were performed by our collaborators at *New York University*, and the dielectric measurements by Dr. Rob Kortschot at *Utrecht University*.

The supplier-reported density of TPM monomer is 1.045 g cm^{-3} . We measured the density of TPM in dispersion with an Anton-Paar DMA 4500M density meter. From the total dispersion density at four different particle mass fractions, the density of self-emulsified TPM was determined to be $1.235 \pm 0.010 \text{ g cm}^{-3}$. After polymerization, the density increased by approximately 7% to $1.314 \pm 0.004 \text{ g cm}^{-3}$. Polymerized particles that were calcined for 2 h at $700 \text{ }^\circ\text{C}$ further increased in density to $2.094 \pm 0.016 \text{ g cm}^{-3}$. All density measurements were performed at $20.0 \text{ }^\circ\text{C}$.

The refractive index of polymerized TPM particles was measured by a refractive index matching process. TPM particles were dispersed in a mixture of pyridine ($n_D = 1.509$) and 2-ethylhexyl 4-methoxycinnamate ($n_D = 1.545$) and the scattering intensity of the dispersion was measured at angles 60° , 120° , and 150° with a static light scattering instrument (Scitech Instruments ST-100). The refractive index of TPM particles was estimated to be between 1.512 and 1.513, assuming that the fluids are non-solvents for the particles. Refractive index measurements were performed with a standard Abbe refractometer (Edmund Optics) at room temperature ($22.5 \text{ }^\circ\text{C}$).

The dielectric constant of liquid TPM was measured using a calibrated parallel plate capacitor and an HP4192A impedance analyzer (Hewlett-Packard) in the frequency range of 10–100 kHz. The dielectric constant of liquid TPM was found to be 6.58 ± 0.04 at $22.3 \text{ }^\circ\text{C}$.

The zeta potential of the polymerized TPM particles was measured using a Zetasizer Nano ZS (Malvern Instruments) on dilute aqueous samples (pH 5.6) at $25.0 \text{ }^\circ\text{C}$. We determined that the TPM particles were negatively charged for all synthesis parameters, with a zeta potential of $-36 \pm 3 \text{ mV}$.

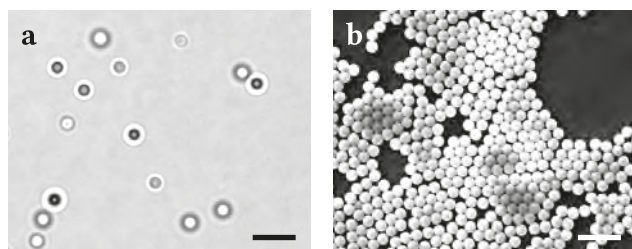


Figure 7.4. Microscopy images of TPM droplets. (a) Optical microscopy image of unpolymerized TPM droplets. (b) SEM micrograph of the same dispersion after polymerization. The scalebars denote $5 \mu\text{m}$.

7.3.2 Emulsification

In order to gain insight into the spontaneous emulsification of TPM in water, we studied the evolution of the droplet size during synthesis. At given intervals following the TPM monomer injection, we sampled 100 μL of the reacting dispersion and imaged a 10 μL aliquot with light microscopy. In order to freeze the emulsification process so that the size distribution could be determined accurately, we polymerized each sample directly as follows: 1 min before sampling, a vial with 1 mg of AIBN and 1 mL of water at 80 $^{\circ}\text{C}$ was prepared. Then, a 90 μL aliquot of the sample was added to the vial and polymerized for at least 2 h in an 80 $^{\circ}\text{C}$ oven.

With SEM, we observed that the particles remained spherical during polymerization (see Figure 7.4). Next to these TPM spheres, sub-200 nm objects were visible that decreased in number throughout the process. As we did not detect these before polymerization, we concluded that they are an artefact of the employed SEM sample preparation. Possibly, the incompletely reacted TPM species polymerize from solution. As these objects tended to precipitate on the growing droplets, the smoothness of the particles could be confirmed only at the end of the process. For the size measurements, we therefore washed the polymerized samples two times with water. Note that we measure the diameter of the polymerized TPM particles: as the density of polymerized TPM is 7 % higher with respect to the emulsion droplets, we expect the particles to be smaller accordingly.

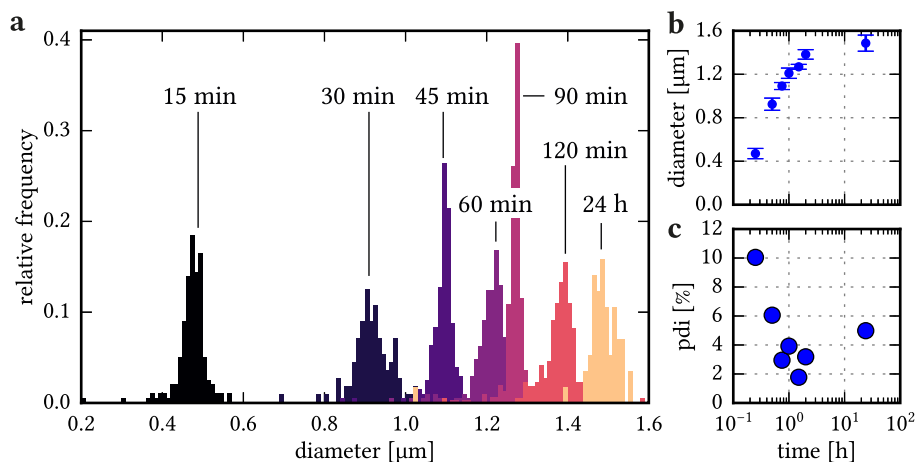


Figure 7.5. Temporal evolution of the size distribution of polymerized TPM droplets in a typical synthesis. The corresponding SEM images are shown in Fig. 7.12 and bright field microscopy videos are available online. (a) Size distributions at different time points in the synthesis. (b) The distributions show a strongly decreasing growth speed. (c) The polydispersity index (PDI) decreases initially and then slowly increases.



The temporal evolution of the size distribution during a synthesis in ammonia solution at initial pH 10.4 and TPM volume fraction $\phi = 0.005$ is shown in Figure 7.5. Corresponding SEM images can be found in Figure 7.12 on page 118. We define $t = 0$ at the time of TPM monomer injection. At $t = 10$ min, ca. 200 nm TPM droplets were observed by light microscopy. At $t = 15$ min, spherical particles with a unimodal diameter distribution of $0.47 \pm 0.05 \mu\text{m}$ were identified using SEM. After this initially fast nucleation step, droplets grow with a decreasing growth rate (see Figure 7.5b). As can be seen in Figure 7.5c, the polydispersity index decreases before plateauing to 2-4% at $t = 90$ min. After that period, the size distribution broadens slightly until the droplets reach a stable size of $1.49 \pm 0.07 \mu\text{m}$.

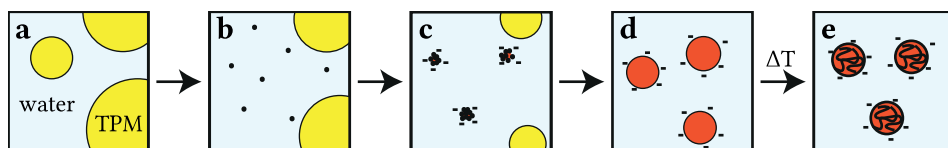


Figure 7.6. Schematic representation of the spontaneous emulsification and subsequent polymerization of TPM. (a) The process starts with TPM monomer injection with a two-phase system consisting of liquid TPM monomer (yellow) and water. (b) Hydrolysis and self-condensation of TPM result in oligomers of several monomer units (black dots). (c) Oligomers nucleate emulsion droplets (red) that are eventually stabilized by charge. The dispersion becomes turbid. (d) The droplets grow by acquiring more TPM-species. In this stage, additional TPM monomer or dye solutions can be added that also transfer to the emulsion droplets. (e) In a second synthesis step, the methacrylate moiety of the TPM is polymerized by adding a radical initiator at elevated temperature.

On the basis of these observations, we propose the following mechanism for the spontaneous emulsification of TPM droplets (see Figure 7.6). Although our mechanism does not capture the details of the nucleation process, it provides a qualitative description of the particle synthesis. Directly after monomer injection, TPM starts to hydrolyse and self-condensate (see Fig. 7.2 on page 104) into oligomers of several monomer units due to a well-established cascade of base-catalysed reactions.^{209,211,212} The gradual shrinkage of bulk monomer droplets has also been observed directly in the preparation of magnetite-covered TPM droplets.²⁰⁷ Because the oligomers become increasingly less soluble in water, they may self-nucleate or aggregatively nucleate (or both) to form the initial emulsion droplets. The accumulation of surface charge on the droplets due to silanol deprotonation prevents or limits the agglomeration of the small particles from forming larger ones.^{205,215,216} The influence of deprotonation will be further investigated in the next section by varying the pH.

At this point, the number of droplets does not appear to change appreciably: we hypothesize that any insoluble oligomeric species with insufficient charge is taken up by the droplets. The dispersion gradually becomes turbid, and we observed the formation of an emulsion with an average droplet diameter of approximately $0.5 \mu\text{m}$. Particles

grow with decreasing growth rate by taking up TPM species in hydrolysed, condensed and/or coagulated form. The macroscopic monomer reservoirs are ultimately consumed during the growth of the droplets. Droplet growth is faster for smaller particles so that the size distribution of emulsion droplets will narrow with time. At a later stage the polydispersity increases slightly, possibly due to Ostwald ripening.

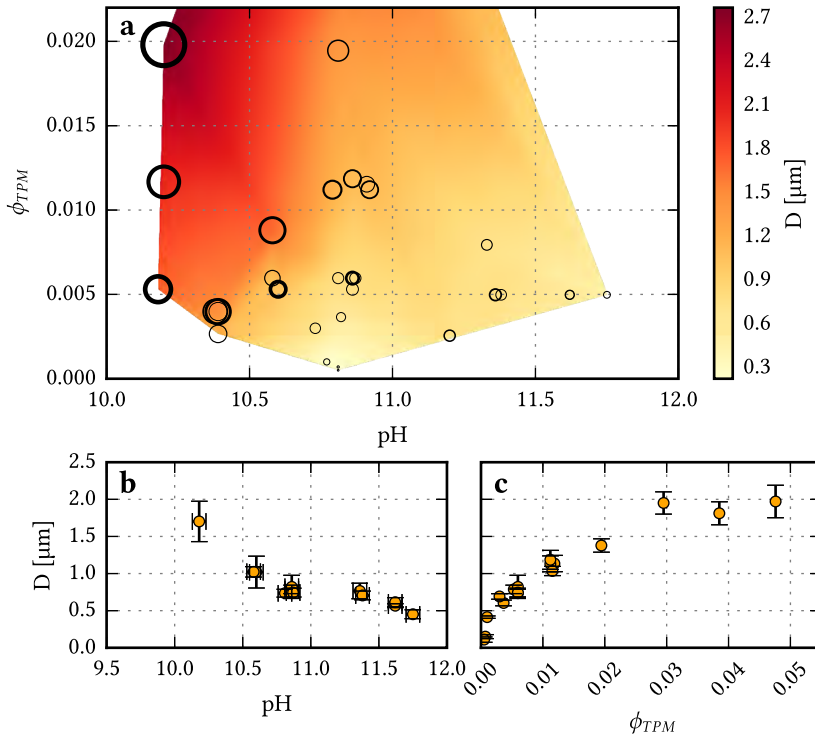


Figure 7.7. Plots of the influence of pH and TPM volume fraction on the diameter of TPM particles. (a) The diameter and line thickness of the plotted circles represent the diameter and polydispersity of the polymerized particles, respectively. The colours are linear interpolations between the measured points. (b) Particle diameter D as function of pH at a fixed TPM monomer fraction of 0.005 to 0.006. The particle size decreases as the pH increases. (c) Particle diameter as function of TPM monomer fraction ϕ at a fixed initial pH of 10.80 ± 0.05 . The particle size increases as the TPM monomer fraction increases.

7.3.3 Change of pH and monomer fraction

The synthesis of TPM particles, as described in the preceding section, has only a few parameters, of which the most important ones are the pH and TPM monomer fraction. We demonstrate their effect on the resulting particle size and polydispersity by a systematic variation and polymerization after 2.5 h of emulsification. See Figure 7.7. Monodisperse emulsions formed only when the pH exceeded 9.7; below that value, highly polydisperse emulsions with sizes ranging from 2–20 μm after 24 h of stirring were observed (see Fig. 7.10 on page 116). Apparently, the hydrolysis and self-condensation reactions proceed too slowly below pH 9.7. Also, colloidal stability is compromised under this condition due to insufficient deprotonation of the TPM species. Above pH 12, we observed the formation of emulsions with a large spread of diameters ranging from 0.2 to 0.5 μm , which limits the synthesis conditions at the other end of the pH range.

For reaction conditions with initial pH values between 10.2 and 11.8, we found that the particle diameter decreased monotonically for increasing pH (see Figure 7.7b). The observed trend is supported by our proposed particle formation mechanism: at higher pH, TPM oligomerization occurs more rapidly.^{211,212,217} Silanol deprotonation is enhanced under these conditions as well,²¹⁶ resulting in a larger number of primary droplets. Therefore, if the TPM volume fraction is kept constant, then smaller particles are produced as we observed. A shorter nucleation period is further supported by the observed narrowing of the size distribution with increasing pH.

Similarly to previous experiments on magnetite-covered TPM emulsions,²⁰⁷ we found that the particle diameter increased with increasing monomer volume fraction. For a fixed pH of 10.8 (see Figure 7.7c) the diameter of the emulsion droplets could be continuously tuned from 0.1 to 2.0 μm by simply adjusting the TPM volume fraction from 0.05 to 5 vol %. Below a volume fraction of 0.1 vol %, the relative polydispersity increased to 20 %. This effectively sets a lower limit of the TPM monomer fraction at 0.1 vol %, which corresponds to a droplet diameter of 0.4 μm . Above a monomer volume fraction of 3 vol %, the droplets did not grow significantly, while the droplet polydispersity increased. Therefore, we conclude that at pH 10.8 the practical range for the TPM volume fraction is 0.1 to 3 vol %.

7.3.4 TPM monomer storage

To investigate the importance of the TPM monomer storage conditions, we also performed the synthesis using a TPM stock that was exposed periodically to air during a period of roughly six months. It is known²¹¹ that TPM slowly forms oligomers when contacted with small amounts of moisture such as is present in the air. Using the air-exposed TPM, we synthesized particles and measured their average size as a function of monomer volume fraction and pH, see Fig. 7.13 on page 119. We found the particle size to depend non-linearly on the pH. Above pH 9.7 a stable emulsion was formed, comparable to the synthesis with fresh TPM. Similarly, from pH 9.7 to 10.7, the particle size decreases monotonically. However, from pH 10.7 to 12, the particle size was found

to increase again to a diameter of $1.47 \pm 0.14 \mu\text{m}$ (pH 12.0, $\phi = 0.005$), which contrasts sharply with the monotonous decrease to $0.45 \pm 0.06 \mu\text{m}$ (pH 11.8, $\phi = 0.005$) observed for fresh TPM. The particle size polydispersity did not increase significantly due to the uncontrolled moisture exposure. These results indicate that the initial droplet formation, but not the droplet growth, is affected by the ageing of TPM. Therefore, to ensure batch to batch reproducibility, we recommend handling the TPM monomer as a moisture-sensitive compound, taking care to exclude or minimize exposure to air, for example by storing under a nitrogen atmosphere.

7.3.5 Influence of base species

To evaluate other base catalysts, we performed the particle synthesis using three different basic compounds: ammonia, sodium hydroxide (NaOH) and tetramethylammonium hydroxide (TMAH). If the emulsification process were dependent on the base-catalysed processes alone, then the type of base used to adjust the pH should not be important.^{211,216} Indeed, we observed that for all three bases a monodisperse emulsion was formed under appropriate conditions, confirming that pH is the control parameter for stable emulsion formation.

The results for the particle size as a function of pH and TPM volume fraction are presented in Figure 7.14 (TMAH) and Figure 7.15 (NaOH) on page 120. For fixed monomer volume fraction ($\phi = 0.005$), the resulting particles were $0.61 \pm 0.06 \mu\text{m}$ when using ammonia at pH 11.6, $0.72 \pm 0.03 \mu\text{m}$ when using TMAH at pH 11.7, and $0.78 \pm 0.08 \mu\text{m}$ when using NaOH at pH 11.8. Because of the differences in base dissociation constants, the respective concentrations of these solutions were 0.9 M ammonia ($\text{p}K_{\text{b}} = 4.75$), 0.4 M TMAH ($\text{p}K_{\text{b}} = 4.2$), and 6.3 mM NaOH ($\text{p}K_{\text{b}} = 0.20$). Although there is a large difference in base concentration, the resulting particle sizes are very similar. This observation supports our claim that the synthesis process and resulting particle size depend mainly on pH and less on the type of base employed.

The fact that TMAH can be used to form monodisperse emulsions of TPM in water also sheds new light on previous experiments performed with TPM droplets that were stabilized by various inorganic colloids in the presence of TMAH.^{206–208,214,218,223} It was proposed that the formed TPM emulsions were stabilized by inorganic colloids at the interface, deprotonated TPM molecules, and TMAH that functions as a peptizing agent. Our study shows that the TMAH-induced deprotonation of TPM might be more important than the stabilization by TMAH itself. This suggests that TPM droplet formation is governed primarily by the volume fraction of TPM monomer and the pH of the solution.

7.3.6 Seeded growth

TPM droplets can be grown to larger sizes through a seeded growth process. This work was performed by Rohit Bhan, Dr. Andrew Hollingsworth, and Dr. Stefano Sacanna (*New York University*). After the initial emulsification of the droplets, additional TPM monomer is added to the stirred system to increase the diameter of the particles. A period

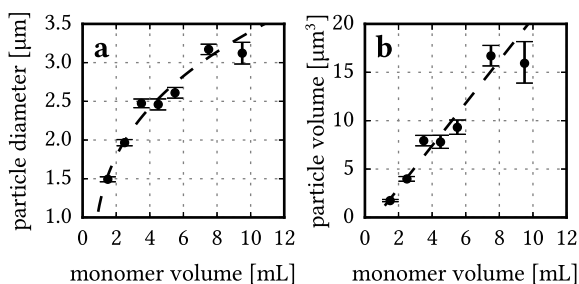


Figure 7.8. Dependence of TPM particle size on added monomer in a seeded growth process. (a) Particle diameter as a function of total added monomer volume. (b) Particle volume as function of total added monomer volume. The dashed line in (b) is obtained with linear regression. The same line is also displayed in (a). Corresponding SEM images can be found in Figure 7.16 on page 121.

of 1 h is necessary for complete hydrolysis and condensation of the first addition of oil-monomer, resulting in visibly larger emulsions. This process can be repeated multiple times, allowing growth to a desired particle size.

In a typical procedure, we add 500 μL TPM monomer ($\geq 98\%$) to 100 mL of ammonia solution at pH 11 (1 mL of 2.8% ammonia and of 99 mL water in a stirred and covered glass beaker). Every hour, we added an additional 1 mL of TPM monomer to the emulsion. When using this technique to grow large droplets (above 2.5 μm), both the ratio of water-to-oil and interparticle spacing decrease. At low water-to-oil ratios, a phase separation can occur, driving emulsion droplets back into the oil phase. This situation can be avoided easily by increasing the volume of the solution. Adding 100 mL of ammonia solution every fifth addition of monomer sustained particle growth. Larger particles can be grown in fewer intervals by increasing the amount of monomer added. We recommend adding no more than four times the original amount of monomer (2 mL in our procedure). The addition of too much monomer can cause a phase separation between water and TPM monomer and thus a loss of particles.

This process results in very low particle polydispersity, as can be seen from the crystal lattice formation in the SEM images (see Figure 7.16 on page 121). The resulting particle diameters are shown in Figure 7.8. Clearly, the particle volume increases linearly with the added monomer volume, indicating a pure growth process. This behaviour is unlike the behaviour for the initial emulsification step. The absence of secondary nucleation further corroborates the conclusion that the added TPM is distributed evenly over the TPM droplets.

7.4 Conclusion

We have systematically studied a straightforward particle synthesis to prepare monodisperse colloids based on the spontaneous emulsification of a polymerizable trialkoxysilane. Also, we characterized the density, refractive index, dielectric constant, and zeta potential of the particles as well as several ways to include dyes for use in fluorescence microscopy. We found that the TPM particle size can be controlled via the pH and the volume fraction of added monomer to produce particles of 0.4–2.8 μm in diameter. The particle size was observed to increase with increasing monomer volume fraction while the particle size decreased with increasing pH. Furthermore, we showed that these trends hold regardless of which base is used as a catalyst.

We proposed a mechanism for the spontaneous emulsification of TPM droplets. First, the hydrolysis and self-condensation of TPM produce oligomers that subsequently nucleate ca. 200 nm primary droplets. Deprotonation of the TPM species provides charge to these droplets. When the droplet charge is sufficiently large to ensure colloidal stability, the process enters a growth step in which the particle size increases without secondary nucleation. The relative polydispersity decreases in this growth step, although at longer times the particle size distribution broadened slightly.

For increasing synthesis pH, we obtain smaller droplets. As a higher pH induces more charges in the droplet surface, deprotonation suppresses TPM oligomer coagulation at an earlier stage by means of electrostatic repulsion between droplets, so that a (colloidally) stable emulsion with more droplets is obtained. Because the available TPM monomer is distributed over the droplets, a smaller size is obtained. After the initial emulsification step, particles can be grown to sizes of up to 4 μm by repeated addition of monomeric TPM. In this way, particles with a narrow size distribution can be obtained over a large size range from 0.4–4 μm .

TPM particles have already been proven to be a versatile template for the bulk synthesis of more complex particles.^{47,50,200,204} The organosilica nature of the TPM particles provides handles for many chemical functionalizations, and the fact that the size of the particles can be tuned by changing easily accessible parameters such as the pH or the amount of added monomer provides flexibility. In this method, no extensive chemical equipment is required: if for instance a dilute sodium hydroxide stock is used, then the synthesis can be performed as a tabletop experiment. The simplicity of this colloidal synthesis makes it easily accessible to a wide range of applications.

7.5 Supporting Figures

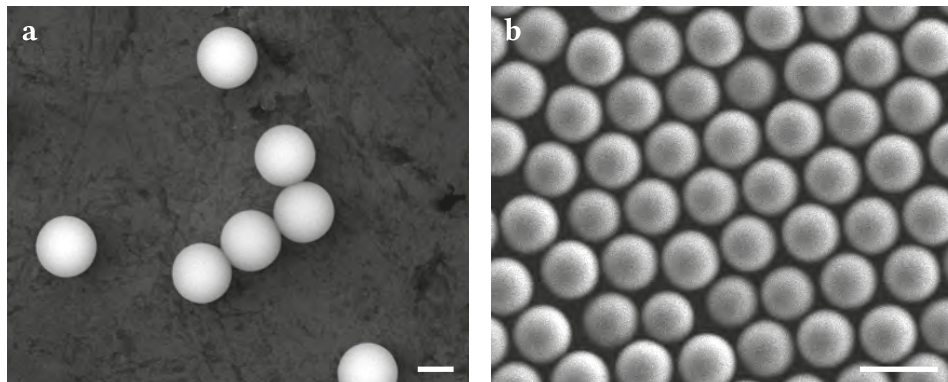


Figure 7.9. SEM pictures of TPM particles with two alternative radical initiators. Different batches of emulsion droplets were polymerized (a) thermally with potassium persulfate (KPS) or (b) photochemically with 2-hydroxy-2-methylpropiophenone. The scalebars denote 1 μm .

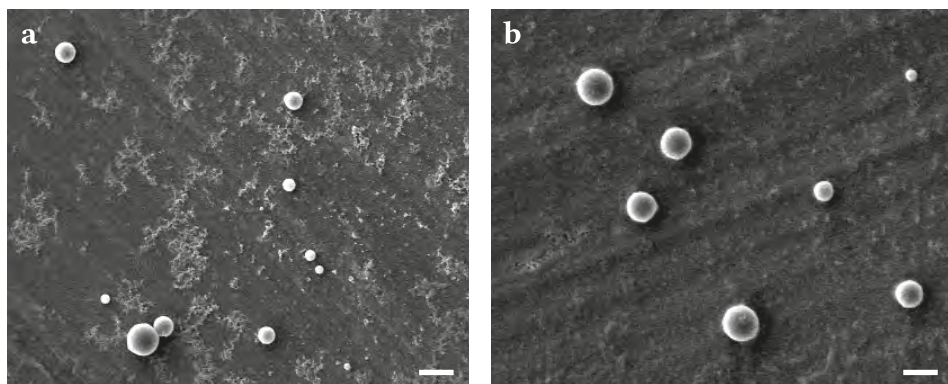


Figure 7.10. SEM micrographs showing TPM droplets that were stirred for 24 h at (a) pH 8.7 and (b) pH 9.4. Both samples contain highly polydisperse TPM droplets with diameters ranging from 2–20 μm . The scalebars denote 5 μm .

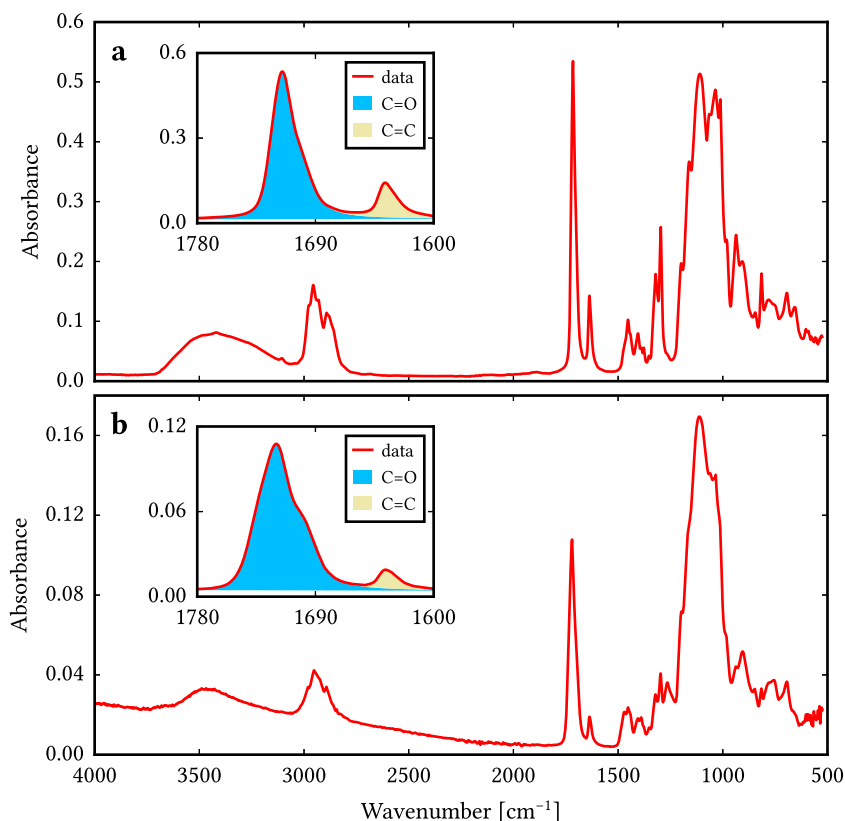


Figure 7.11. Infrared spectra of (a) unpolymerized TPM droplets and (b) polymerized TPM particles. The emulsification was performed in 12.4 mM ammonia (pH 10.7) at a TPM monomer volume fraction $\phi = 0.005$. After emulsification, an aliquot the droplets was first washed with water to remove the excess of ammonia. Then the droplets were coalesced by centrifugation at 3500 rcf and the spectrum (top) was recorded. Another aliquot of the emulsion droplets was polymerized with 0.67 g L⁻¹ AIBN at 80 °C for 2 h. The polymerized particles were washed two times with water, dried overnight at 90 °C and then the FTIR spectrum was recorded. To estimate the degree of polymerization, we compared the area under the peaks at 1636 cm⁻¹ and 1720 cm⁻¹, which respectively are the C=C and C=O stretch vibrations. As shown in the inset graphs, we numerically integrated the baseline-corrected absorbance, separating the two peaks by a local fit of two Lorentzians. As the C=O bonds are unaffected during polymerization, we may use the ratio between these areas to compute the decrease in the number of C=C bonds. We found that this area ratio (C=C:C=O) decreased from 0.195 to 0.072, corresponding to a polymerization degree of 63%. This corroborates that the TPM droplets have polymerized through a radical polymerization in the second step. These data were acquired by Zhe Gong and Dr. Stefano Sacanna at *New York University*.

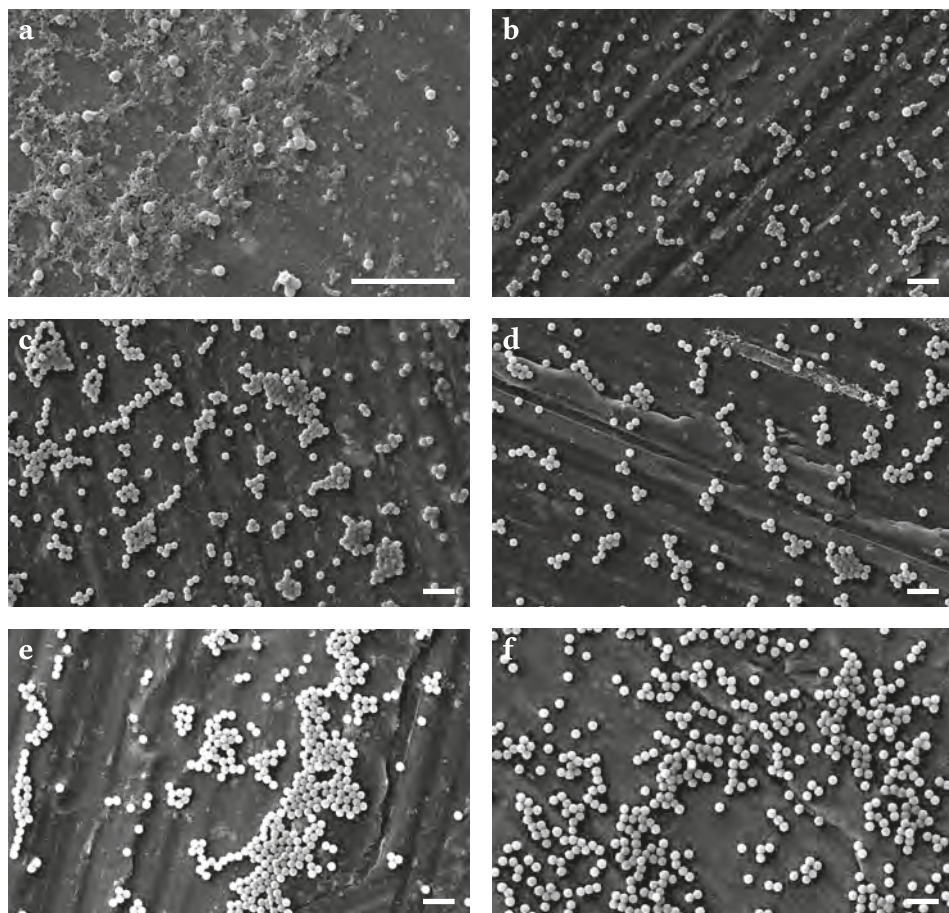


Figure 7.12. Representative SEM micrographs showing the time evolution of polymerized TPM droplets in a single synthesis. The shown pictures were obtained from the polymerized samples that were taken at 15 min (a), 30 min (b), 45 min (c), 60 min (d), 90 min (e), and 120 min (f) after TPM monomer injection. There are sub-200 nm objects visible that decrease in number throughout the process. We also observed these objects by light microscopy in the polymerized samples, but not in the unpolymerized samples. Therefore we concluded that they are an artefact of the employed sample preparation. Note that the samples were washed for measuring the particle diameters; these images are not shown. Scalebars denote 5 μm .

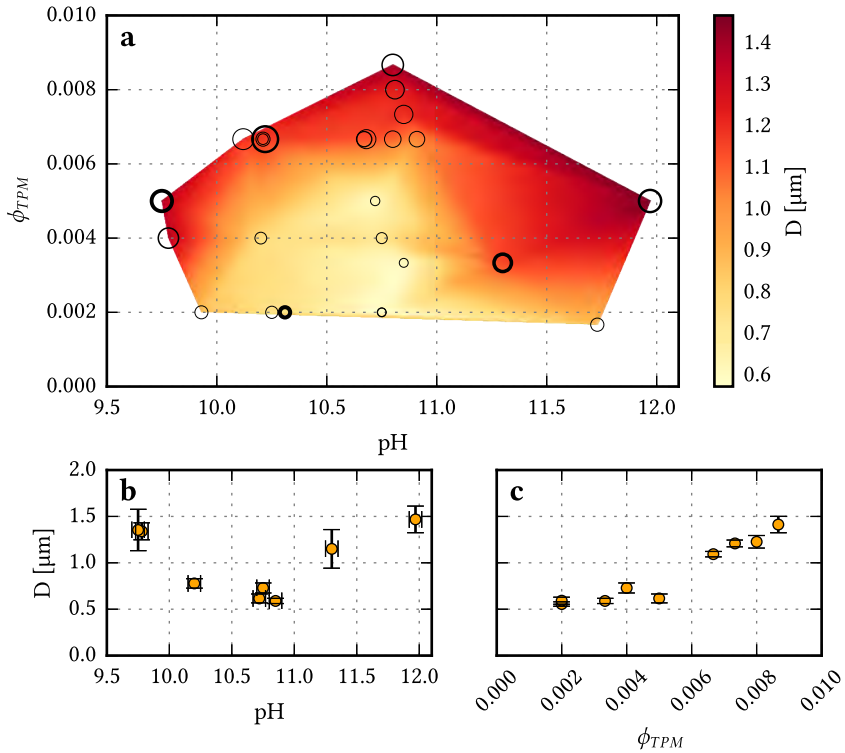


Figure 7.13. Plots of the influence of pH and TPM volume fraction on the particle diameter, in the case TPM is used that was subject to ambient conditions for approximately 6 months. Ammonia was used to adjust the pH. (a) The diameter and line thickness of the plotted circles represent the diameter and polydispersity of the polymerized particles, respectively. The colours are linear interpolations between the measured points. (b) Particle diameter D as function of pH at a fixed TPM monomer fraction of 0.003 to 0.005. The particle size decreases as the pH increases and at high pH increases again. The minimum particle size occurs at pH 10.75. The size polydispersity has a minimum between pH 10 and 11. (c) Particle diameter as function of TPM monomer fraction ϕ at a fixed pH of 10.7 to 10.9. The particle diameter increases with increasing monomer volume fraction.

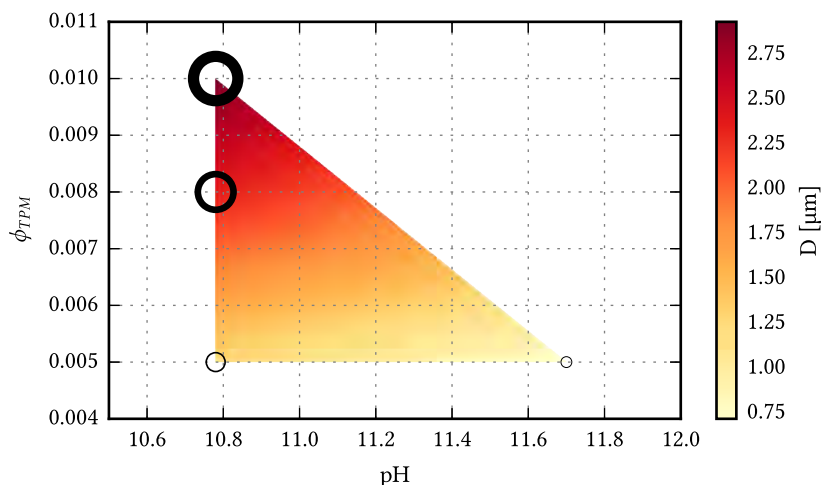


Figure 7.14. Plots of the influence of pH and TPM volume fraction on the particle diameter, using tetramethyl ammonium hydroxide (TMAH) to adjust the pH. The diameter and line thickness of the plotted circles represent the diameter and polydispersity of the polymerized particles, respectively. The particle diameter increases with increasing monomer volume fractions and decreases as the pH increases.

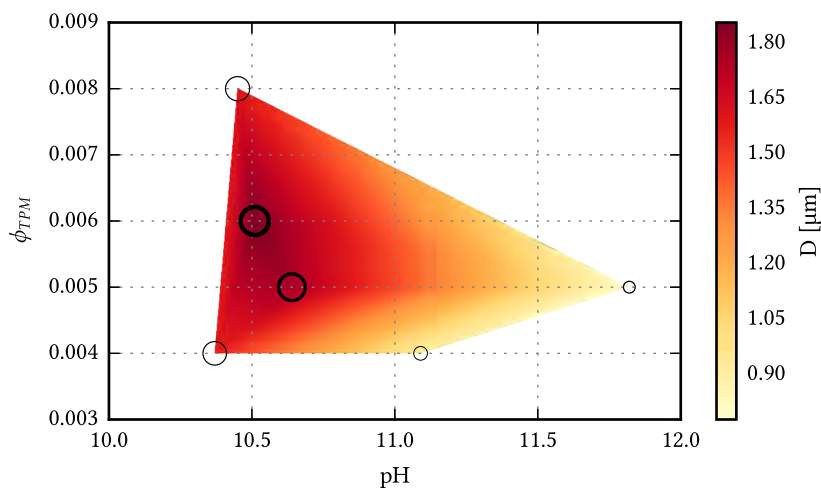


Figure 7.15. Plots of the influence of pH and TPM volume fraction on the particle diameter, using sodium hydroxide (NaOH) to adjust the pH. The diameter and line thickness of the plotted circles represent the diameter and polydispersity of the polymerized particles, respectively. The particle diameter increases with increasing monomer volume fractions, but less than when TMAH or ammonia are used as base catalyst. The particle size decreases as the pH increases.

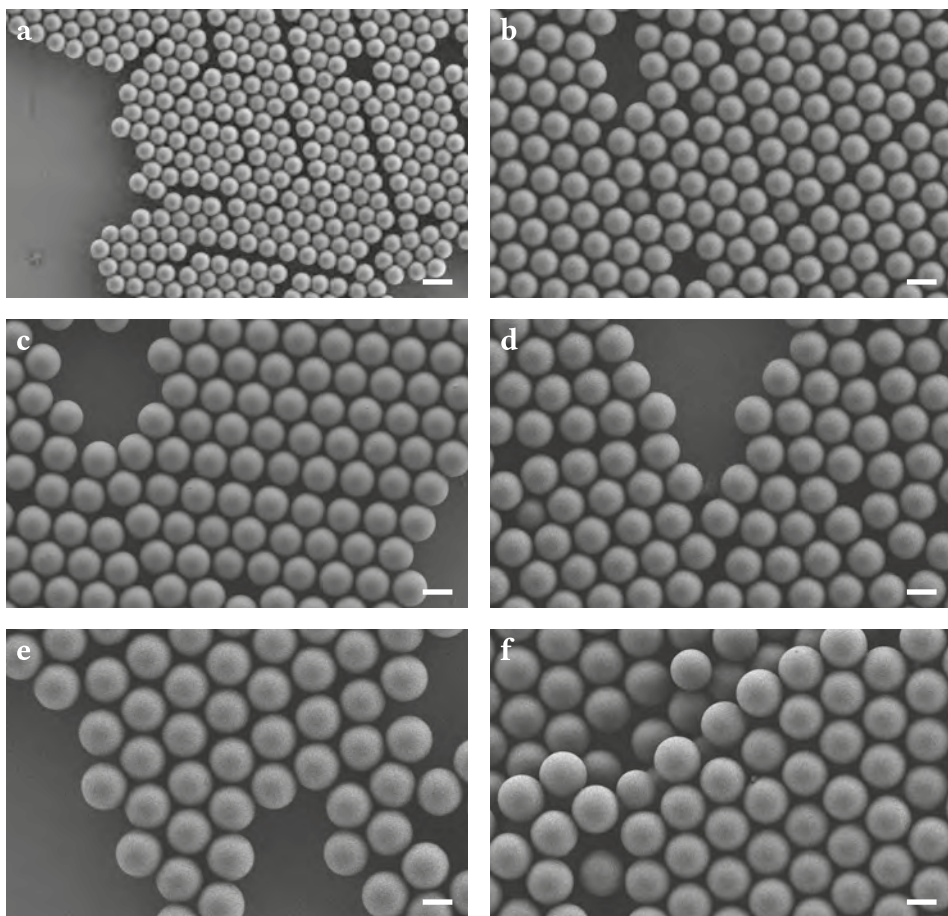


Figure 7.16. SEM micrographs of TPM particles obtained through a single seeded growth process. After nucleation with 500 μL TPM monomer in 100 mL diluted ammonia, samples were made after (a) a single addition of 1 mL monomer, (b) a second addition of 1 mL monomer, (c) a third addition of 1 mL monomer. Then a fourth addition of 1 mL monomer was done, the reaction volume was doubled by adding 100 mL diluted ammonia, and (d) the fifth addition of 1 mL monomer, (e) the sixth addition of 2 mL monomer and (f) the seventh addition of 2 mL monomer were done. Scalebars denote 2 μm .

