Magnetic Nanowires Generated via the Waterborne Desalting Transition Pathway

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ABSTRACT: We report a simple and versatile waterborne synthesis of magnetic nanowires following the innovative concept of electrostatic “desalting transition”. Highly persistent super-paramagnetic nanowires are generated from the controlled assembly of oppositely charged nanoparticles and commercially available polyelectrolytes. The wires have diameters around 200 nm and lengths between 1 μm and 0.5 mm, with either positive or negative charges on their surface. Beyond, we show that this soft-chemistry assembly approach is a general phenomenon independent of the feature of the macromolecular building blocks, opening significant perspectives for the design of multifunctional materials.

KEYWORDS: superparamagnetic nanoparticles, polyelectrolytes, nanostructured wires, electrostatic complexation, desalting transition

1. INTRODUCTION

Magnetic nanowires have received considerable attention during the past years because of their potential applications in cell manipulation,1–5 microfluidics,6 and micromechanics.7–9 These objects are anisotropic colloidial particles with submicronic diameters and lengths in the range 1–100 μm. For specific applications, elongated magnetic structures appear as promising alternatives to the spherical magnetic beads developed so far. At the application of a magnetic field, torques may be applied to the wires and their orientational behavior be monitored, providing additional degrees of freedom for investigating complex fluids. Wires can also be made hollow with an internal surface in contact with the bulk phase and be used for separation and release of actives.10

As far as the synthesis of magnetic nanowires is concerned, two main strategies are put forward in the literature. The first approach consists in the electrodeposition of magnetic atoms such as nickel or iron into thin alumina-based porous templates with cylindrical holes.11,12 At the dissolution of the template, ferromagnetic nanowires are produced and dispersed in water-based solvents. These wires are used in cell guidance, cell separation, and micro rheology experiments.2–4,6,12 One major drawback with nickel nanowires is that these structures carry a permanent magnetic moment, and thus aggregate spontaneously because of magnetic dipolar interactions.3,5 The second strategy is based on the coassembly of polymer microbeads loaded with magnetic nanoparticles.7–9,13,14 With this technique, permanent chains are formed by the application of a magnetic field for alignment and by the addition of binding agents for connecting the beads. Filaments and chains with micronic diameters and length 1–200 μm are fabricated, either in bulk solutions8,13 or tethered on substrates.7,14,15 The two synthesis routes end up, however, with different magnetic structures: the nickel nanowires are stiff and ferromagnetic, whereas the bead assemblies remain paramagnetic and semiflexible. Attempts to build supracolloidal assemblies using sub-10 nm particles were also carried out, resulting in isotropic clusters16–18 or in tortuous pearl-necklace filaments.19–21

We report here a simple and versatile waterborne synthesis of magnetic nanowires following the innovative concept of electrostatic “desalting transition” put forward recently on a sub-10 nm nanoparticles/charged-neutral block copolymers system.22,23 We generalize this electrostatic coassembly pathway to strong and weak homopolyelectrolytes (homoPEs) and demonstrate that the transition between a disperse and an aggregated state of particles observed as a function of the ionic strength is a general phenomenon independent of the nature of the polyelectrolytes considered. Furthermore, under the presence of a magnetic field

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this soft chemistry approach enables the fabrication on-demand of highly persistent superparamagnetic nanowires with lengths comprised between 1 μm and 0.5 mm bearing either positive or negative charges on their surface (Figure 1).

2. MATERIALS AND METHODS

Iron oxide nanoparticles were synthesized according to the Massart technique by alkaline coprecipitation of iron(II) and iron(III) salts, oxidation of the magnetite (Fe3O4) into maghemite (γ-Fe2O3) nanoparticles and by size-sorting by subsequent phase separations. At the end of the synthesis, the particles were positively charged and dispersed in water at a weight concentration ε ≈ 10 wt % and pH 1.8. The magnetic dispersions were characterized by transmission electron microscopy (TEM) and diffraction, vibrating sample magnetometry (VSM), magnetic sedimentation and light scattering (DLS). For the dispersions investigated here, the size distribution was found to be log-normal with median diameter 8.3 nm and a polydispersity 0.26. The polydispersity index is defined as the ratio between the standard deviation and the average diameter. To improve their stability, the particles were coated with anionic charges carried by the particles at their surfaces. With this notation, Z = 1 describes the isoelectric solution, characterized by the same number densities of positive and negative chargeable ions. For the desalting transition pathway, ammonium chloride (Sigma) was used to adjust the ionic strength up to 3 M.

3. RESULTS AND DISCUSSION

The aggregate formation was monitored using different formulation pathways, including direct mixing, dialysis, and dilution. Only dialysis was performed in the presence of an external applied magnetic field, in agreement with the scheme of Figure 1. 3.1. Direct Mixing. With the direct mixing protocol, the complexation of oppositely charged species was controlled by pouring rapidly the magnetic dispersion into the polymer solution at the desired charge ratio Z. The samples were allowed to rest for a week before being investigated by light scattering. Figure 2a displays the hydrodynamic diameters (Dh) obtained for PAA2K-γ-Fe2O3 complexed with PTEA11K-b-PAM30K copolymers and with PDADMAC, PAH and PEI polycations. For the 4 systems, the Dh’s were found to pass through a sharp maximum around Z = 1. The measured diameters for PTEA11K-b-PAM30K diblocks (Dh = 100 nm) were lower than those of the homoPEs (Dh = 300 nm), indicating a softening of the assembly process when noninteracting neutral blocks are present. The intense peaks around the charge stoichiometry bear strong similarities with that observed for polyplexes and lipoplexes in transfection strategies, suggesting analogous complexation mechanisms. 3.2. Desalting Transition without Magnetic Field. Although appealing and simple, the direct mixing protocol was not appropriate to control the sizes and morphologies of hybrid nanostuctures. Indeed, under strong driving forces like electrostatic interactions, the key factor in the complex formation is the competition between the reaction time of the components and the homogenization time of the solutions. To overcome these limitations, alternative pathways inspired from molecular biology were developed. The protocols applied here consist first in the screening of the electrostatic interactions by bringing the dispersions to high ionic strength (Ib), and second in the progressive reduction of Ib by dialysis (Figure 1) or by dilution. With this technique, the oppositely charged species were intimately mixed in solution but did not interact owing to the

Figure 1. Schematic representation of the desalting transition protocol that controls the nanoparticle coassembly and wire formation.
hydrodynamic diameter measured on dispersions containing PAA2K obtained by direct mixing at concentration 0.1 wt % and temperature 25 °C. The polymer/particle dispersions were made of PAA2K and oppositely charged polymers (PEI; and green, PAH). The polymer/particle dispersions were obtained by direct mixing at concentration 0.1 wt % and temperature 25 °C. The samples were allowed to rest for a week before being investigated by light scattering. (b) Ionic strength dependence of the hydrodynamic diameter measured on dispersions containing PAA2K−γ-Fe2O3 particles and oppositely charged polymers (Z = 1). The ionic strength was monitored by addition of ammonium chloride (NH4Cl). The color code is the same as in a). With decreasing IS the desalting transition occurred at a critical ionic strength 0.37, 0.54, and 2.3 M for PTEA11K-b-PAM30K, PDADMAC, and PEI respectively. The empty symbols and hatched area indicate hydrodynamic sizes that could not be measured by light scattering. Optical microscopy was used instead.

Figure 2. (a) Hydrodynamic diameter $D_H$ as a function of Z for PAA2K−γ-Fe2O3 complexed with homo- and copolymers for charge ratios $1 \times 10^{-3}$ to 100 (black, PTEA11K-b-PAM30K; red, PDADMAC; blue; PEI; and green, PAH). The polymer/particle dispersions were obtained by direct mixing at concentration 0.1 wt % and temperature 25 °C. The samples were allowed to rest for a week before being investigated by light scattering. (b) Ionic strength dependence of the hydrodynamic diameter measured on dispersions containing PAA2K−γ-Fe2O3 particles and oppositely charged polymers (Z = 1). The ionic strength was monitored by addition of ammonium chloride (NH4Cl). The color code is the same as in a). With decreasing IS the desalting transition occurred at a critical ionic strength 0.37, 0.54, and 2.3 M for PTEA11K-b-PAM30K, PDADMAC, and PEI respectively. The empty symbols and hatched area indicate hydrodynamic sizes that could not be measured by light scattering. Optical microscopy was used instead.

electrostatic screening. In the dilution process, deionized water was added stepwise to the polymer/particle salted dispersion and the scattered intensity and hydrodynamic diameter were determined by light scattering. For a dispersion containing both anionic particles and cationic PTEA11K-b-PAM30K diblocks, $D_H$ versus IS exhibited an abrupt upturn (Figure 2b). Found at 0.37 M, this upturn sets the limit between two states: above, the particles remain dispersed and unaggregated; below, the particles are retained within dense spherical clusters thanks to the cationic polymer “glue”. No transition could be evidenced for the particles or for the copolymers taken separately (see the Supporting Information, SI-3).

The desalting transition was shown to be a key feature in the fabrication process of the nanowires; its occurrence with widespread homoPEs had then to be investigated. The ionic strength of mixed dispersions containing PAA2K−γ-Fe2O3 and PDADMAC, PEI, and PAH was increased up to 3 M (NH4Cl) to verify the efficiency of the screening. Effective screening was found for PDADMAC and PEI, but not for PAH. For this later system, even at 3 M, the oppositely charged species interacted strongly and large aggregates were formed ($D_H = 400$ nm, see the Supporting Information, SI-3). Figure 2b displays the desalting transition for PAA2K−γ-Fe2O3/PDADMAC and PAA2K−γ-Fe2O3/PEI, at the critical ionic strengths 0.54 and 2.30 M respectively. With the homoPEs, the features of the desalting transition remained similar to those found with copolymers. The sizes of the clusters below the critical value increased however rapidly with decreasing IS. As anticipated, Figure 2b suggests that the interactions are stronger with homoPEs than with the diblock. Dispersions prepared apart from the charge stoichiometry, i.e., at $Z = 0.3$ and $Z = 7$ were found to undergo similar desalting transitions, with aggregates of smaller sizes and slightly shifted ionic strengths (see the Supporting Information, SI-4).

3.3. Desalting Transition in the Presence of a Magnetic Field. From the initial 3 cationic polymers, only PDADMAC and PEI exhibited a well-marked desalting transition. These polymers were tested with respect to dialysis in the presence of a 0.3 T magnetic field to stimulate the unidirectional growth of the aggregates (Figure 1). Once the ionic strength of the dialysis bath reached its stationary value, the magnetic field was removed and the solutions were studied by optical microscopy. Figures 3 show optical transmission microscopy images of aggregates made of PAA2K−γ-Fe2O3/PDADMAC dispersions at $Z = 0.3$ (Figure 3a), 1 (Figure 3b) and 7 (Figure 3c). Large and irregular aggregates in the 100 μm range were obtained at $Z = 1$, illustrating again that at the charge stoichiometry the interactions were strong and that electrostatic aggregates grew uncontrolled. With an excess of cationic ($Z = 0.3$) or anionic ($Z = 7$) charges however, regular nanostructured wires were readily fabricated. The wires grown apart from the charge stoichiometry ($Z = 0.3$ and 7) had a median length $L_0 = 90$ and 20 μm respectively, and exhibited the same mechanical and magnetic properties than the ones made from PTEA11K-b-PAM30K copolymers: they were all rigid threads with superparamagnetic properties inherited from the single particles. As an illustration of their magnetic properties, a movie of wires subjected to $\pi/2$-reorientations is shown in Supporting Information (Movie 1). The results obtained with PDADMAC were utterly reproduced with polyethyleneimine (see the Supporting Information, SI-5), showing in fine that the wire formation is a general phenomenon that does not depend on the nature of the polycations. For all polymers tested, the length distributions of the wires were found to be log-normal with median length $L_0 (= 10−100 \mu m)$ and polydispersity 0.5. With PDADMAC, the tail of the distribution contained few but very long wires of length 0.5 mm, suggesting the possible influence of the chain architecture (linear for PDADMAC and branched for PEI) on the wire formation (Figure 3d). To confirm that the wires made with copolymers and homoPEs had the same structures, TEM was monitored on PDADMAC and PEI-based dispersions. Figures 4 compare the internal structures of wires obtained with PTEA11K-b-PAM30K at $Z = 1$ (Figure 4a), PDADMAC at $Z = 0.3$ and 7 (Figure 4b,c), and PEI at $Z = 0.3$ and 7 (Figure 4d,e) as binding agents. Extremities or intermediate pieces reveal diameters comprised between 150 and 400 nm with similar particles densities. These findings are in good agreement with the optical microscopy images which displayed no major differences between wires at the micrometer size. Note finally that the desalting transition technique allows the high-throughput synthesis, typically $1 \times 10^{10}$ wires of length 10 μm within 10 min.

In a recent paper, it was concluded that the mechanism for the wire formation proceeded in two steps: (i) the nucleation and
growth of spherical clusters of particles, and (ii) the alignment of the clusters induced by the magnetic dipolar interactions. For the kinetics, the clusters growth and their alignment occurred in parallel, leading to a continuous welding of the cylindrical structure. The results obtained with homoPEs indicate that the kinetics of coassembly at charge stoichiometry is extremely rapid. All three pathways explored, direct mixing (Figure 2a), dilution (Figure 2b), and dialysis (Figure 3) showed it: at $Z = 1$, the nanoparticle aggregates are the largest and their morphologies mainly uncontrolled. With block copolymers, the coassembly kinetics was slowed down by the presence of the inert neutral blocks. With homoPEs, the finding of regular nanowires apart from stoichiometry suggests a kinetics governed by the balance between electrostatic repulsions and dipolar attractions between the precursor aggregates. We thus anticipate that the wires made with an excess of polymers should be positively charged and those with an excess of nanoparticles negatively charged. Electrokinetic measurements were performed on the wire dispersions to determine their electrostatic charges, however the data remained inconclusive because of the large sizes of the aggregates. Smaller Brownian isotropic clusters ($D_H = 250$ nm) were thus fabricated using the same dialysis conditions but without external magnetic field. Electrophoretic mobility and $\zeta$-potential were determined for PDADMAC and PEI based complexes at $Z = 0.3, 1$, and $7$ using Zetasizer Nano ZS Malvern Instrument. The intensity distribution of the electrophoretic mobility shown in Figure 5 for the
three systems confirmed our hypothesis. At $Z = 0.3$, $\mu_0$ is centered around $+3 \times 10^{-4}$ cm$^2$ V$^{-1}$ s$^{-1}$ for both polyelectrolytes, whereas it is approximately 0 and $-2.2 \times 10^{-4}$ cm$^2$ V$^{-1}$ s$^{-1}$ at $Z = 7$ (see the Supporting Information, SI-6). A final test was performed to confirm these results. Positively ($L_0 = 90$ nm) and negatively ($L_0 = 20$ nm) charged PDADMAC wires were mixed together. The suspended colloids precipitated, resulting in the formation of large bottlebrush-like aggregates (Figure S, lower panel), where the short wires agglutinated onto the large ones thanks to attractive electrostatic interactions, suggesting oppositely charged wires.

4. CONCLUSION

In conclusion, we have shown that the previous copolymer based coassembly strategy could be generalized to strong and weak polyelectrolytes. In terms of cost and practicality, this represents a remarkable improvement. Beyond, it shows that specific polymer architectures are not essential for the elaboration of supracolloidal electrostatic assemblies of adaptable size, charge and morphology, the structural, mechanical and magnetic properties being independent of the nature of the binding agents. The enhanced stability of the nanostructures built up with anionic iron oxide nanoparticles and cationic homoPEs is interpreted as resulting from a nonequilibrium associating process. Once the aggregates are formed, their microstructure remained unchanged over a period longer than years. As a result, the wires can be moved in the solvent or rotated by the application of a field or gradient without changing their internal structure. Because the present strategy for electrostatic coassembly of nanoparticles is simple and versatile, it should open new perspectives for the design of nanodevices, such as tips, tweezers, actuators applicable in biomedicine for stimulating or sorting living cells or as novel contrast agents. We envision very exciting applications in these domains where the wires can be tailored on-demand according to the medium under scrutiny.

ASSOCIATED CONTENT

Supporting Information. Characterization of the iron oxide nanoparticles and that of the cationic polymers; evolution of the hydrodynamic diameter of particles and polymers as a function of ionic strength as well as the phenomenology of the desalting transition described in detail; magnetic nanowires obtained from PEI homopolyelectrolytes and results of the electrophoretic mobility and $\zeta$-potential (PDF). The movie shows a series of reorientations of wires following 90° flips of the magnetic field (AVI). This material is available free of charge via the Internet at http://pubs.acs.org/.

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