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Feature article

Block copolymer template-directed synthesis of well-ordered metallic nanostructures

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3D nanostructured inorganic materials appear as promising candidates for various practical applications. Here we focus on metal nanofoams, a class of 3D nanomaterials uniquely combining the properties of metals and nanoporous materials, and review the recent developments in their preparation methods. Common approaches, such as dealloying, sol–gel synthesis, nanosmelting, combustion synthesis, etc., render metallic nanostructures with highly disordered architectures which might have adverse effects on their mechanical properties. In contrast, block copolymers have the ability to self-assemble into bicontinuous ordered nanostructures that can be applied as templates for the preparation of well-ordered metal nanofoams. Several examples of block copolymer template-directed synthesis of continuous metallic nanostructures will be described and the prospects of this approach will be discussed.

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1. Introduction

To fulfill the constant quest for novel functional materials, the cooperation and development of research areas, such as material science, chemistry, physics, and biology, is required. Nowadays, much attention is given to the field of nanostructured materials that, by definition, have at least one dimension in the range from 1 to 100 nm. In comparison to macroscopic systems, nanomaterials have advanced properties arising from their nanoscale dimensions. Zero-dimensional (0D) nanostructures, nanoparticles, are widely applied in biology and medicine for probing of DNA structure [1], MRI contrast enhancement [2], tissue engineering [3,4], cancer therapy [5], etc. Furthermore, they find application in food packaging and food safety [6], as catalysts with enhanced performance [7], for hydrogen storage [8,9], data storage [10], etc. One-dimensional (1D) nanostructures (e.g., wires, rods, tubes) are expected to play an important role as both interconnects and functional units in fabricating electronic, optoelectronic, electrochemical, and electromechanical nanodevices [11,12]. Several methods have been used to fabricate 1D nanostructures including template-directed synthesis, self-assembly of 0D nanostructures, size reduction of 1D microstructures, electrospinning and others. For example, Park et al. demonstrated the preparation of metallic line patterns using functional block copolymer templates [13]. Polystyrene-block-poly(2-vinylpyridine) (PS–b–P2VP) and poly(2-vinylpyridine)-block-poly(-methyl methacrylate) (P2VP–b–PMMA) block copolymers were spin-coated onto Ag-coated silicon substrate and annealed under solvent vapor to achieve the desired surface morphology. Subsequently, porous block copolymers were obtained by either reconstruction of the P2VP block in ethanol or removal of the PMMA block by UV irradiation followed by rinsing with acetic acid. Finally, silver line patterns were generated using electrochemical etching or direct metal evaporation and lift-off processes. 2D nanostructures, like thin films, can be grown by vapor-phase (sputtering, chemical vapor deposition, atomic layer deposition, etc.) or liquid-phase deposition (electrochemical deposition, Langmuir–Blodgett films, self-assembled monolayers, etc.) [14–16]. Finally, 3D nanostructures represent the ultimate form factor for a number of materials. However, their creation is a rather delicate task [17]. The available top-down approaches are extremely challenging: lithography techniques allow precise patterning, but can only be implemented in an inherently 2D manner, whilst the available 3D microscale techniques cannot be easily translated to the nanoscale. Currently, the attention is focused on bottom-up approaches that are usually based on self-assembling processes often similar to those occurring in nature.

In this feature article we discuss 3D nanostructured metals. First, we describe the established methods for metal nanofoam preparation and their limitations in terms of ordering at the
nanoscale. Then, we discuss the application of block copolymers and block copolymer based systems as precursors to metallic nanostructures and suggest the possibilities to overcome the aforementioned limitations with examples of block copolymer template-directed synthesis of metal nanofoams. Finally, we end with a brief outlook on the future trends in this research area.

2. Metal nanofoams: properties and preparation methods

A nanoporous metal foam is defined as a three-dimensional structure comprised of interconnected metallic features with a porosity greater than 50 vol% and in which sub-micron pores significantly contribute to the specific surface area of the foam [18]. Metal nanofoams have, on the one hand, properties characteristic of metals such as electrical and thermal conductivity, catalytic activity and ductility/malleability whilst, on the other hand, also display properties characteristic of nanostructured materials such as high surface area, ultralow density and high strength-to-weight ratio. Due to this unique combination, metal nanofoams appear as candidates for a number of technological applications. Several examples are described in the following paragraph.

Hydrogen is an environmentally friendly energy carrier and its storage is one of the key challenges in the hydrogen economy. It can be stored as pressurized gas, cryogenic liquid or solid fuel as a combination with other materials such as metal hydrides. Due to safety reasons, metal hydrides are the preferable option for hydrogen storage and it has been demonstrated that in comparison to bulk materials, nanostructured metals significantly improve the reaction kinetics, reduce the enthalpy of metal hydrides formation, lower the hydrogen absorption and release temperatures [19–21]. Nanoporous metals and their derivatives can also be employed for the production of novel 3D-structured batteries. Open porous networks and thin ligaments enable fast diffusion of ions into and out of electrodes, thus, creating fast discharging and recharging batteries [22]. Due to the size-effect-enhanced catalytic properties, nanostructured metals, such as nickel, could be a low-cost substitute for precious-metal catalysts [18]. Furthermore, literature reports the tunability of the physical properties of nanoporous metals through an applied voltage [23,24]. Injection of the surface charges into a porous metal nanofoam enhances the surface stresses leading to its expansion or contraction. Actuation properties of metallic nanofoams have been proven experimentally [25]. A bilayer foil consisting of solid and nanoporous gold layer was immersed in aqueous electrolyte and the electrochemical potential varied. The nanoporous layer expanded or shrunk whilst the solid one maintained its original dimensions, and thus, the bilayer foil bent with a tip displacement of several millimeters. The observed strain was orders of magnitude larger than in conventional cantilever bending experiments. Metal nanofoams can also be applied as implant materials as they have a high strength-to-weight ratio and assist osseointegration. The high porosity and interconnected nanochannels enhance bone ingrowth and insure the long-term stability of the implant [26,27].

Several approaches for the preparation of nanoporous metal foams have been established. Erlebacher et al. reported the formation of nanoporous gold by the dealloying process from a binary alloy of silver and gold [28]. Silver, as a less-noble metal, can be etched away either electrochemically or by using acid, resulting in a nanoporous gold network with a ligament spacing of ca. 10 nm (Fig. 1). The morphology of nanoporous gold in terms of sizes of ligaments and pores can be controlled by varying the dealloying potential [29]. An interesting approach in which the dealloying is combined with templating to render hierarchically porous gold has been reported by Nyce et al. [30] First, Ag–Au alloy was electroless plated onto micron-sized spheres of polystyrene and then the polymer was removed by pyrolysis. Finally, silver was etched away from Ag–Au alloy using nitric acid leaving behind hierarchically porous gold.

Sol–gel techniques are well-developed for the synthesis of metal nanoparticles, but their applicability to assemble nanoparticles into metal foams appears to be very challenging. It has
been found that the solubility of metal nanoparticles and the removal of the stabilizing ligand are the most important criteria for successful preparation of porous metals [31]. In one example of gold nanofoam preparation via the sol–gel route, highly water-soluble Au nanoparticles (solubility up to 30 wt%), stabilized with a mixture of hydrophilic and hydrophobic ligands, were dissolved in water and the solution was oxidized with hydrogen peroxide. Nanostructured gold precipitates were formed after 48 h (Fig. 2) [18].

Leventis et al. have recently reported a viable route towards metal nanofoams through nanosmelting of hybrid polymer/metal aerogels [32]. Using the sol–gel approach, an aerogel consisting of resorcinol-formaldehyde (RF) polymer and iron oxide was prepared. Subsequently, the aerogel was exposed to high temperatures (800–1000 °C) under argon atmosphere, which led to simultaneous polymer degradation and iron oxide reduction to metallic iron (Fig. 3). Similarly, the preparation of porous Cu, Ni, Sn, and Co networks by nanosmelting has been demonstrated [33].

Combustion synthesis represents another pathway for the nanostructured metal production, in which metal is mixed with the energetic precursor that releases energy during its decomposition and drives the formation of metal nanofoam. Tappan et al. have demonstrated the preparation of nanoporous Fe, Cu, Co, and Ag via combustion synthesis [34].

As described, utilizing previously presented techniques, one can prepare highly porous metal foams with nano-sized pores. However, the prepared foams are typically highly disordered which disables the effective stress transmission from the ligament nanoscale to the overall macroscale and results in poor mechanical properties [35–37]. The ordering issue can be overcome in the alternative block copolymer template-directed approach for metal nanofoam synthesis.

3. Self-assembled block copolymers and supramolecular complexes

Chemically distinct polymers generally do not mix, but macro-phase separate, in a similar fashion to oil and water. In a block copolymer, two or more chemically different homopolymers are covalently linked and this linkage prevents macrophase separation. However, due to the unfavorable interactions between the different blocks, phase separation still occurs but, because of their connectivity, only on the nanometer scale and this process is called microphase separation. The unfavorable interactions between chemically distinct species induce stretching of the polymer chains (i.e., to minimize the interaction enthalpy) and the entropic elasticity (of the entropic spring) resists this stretching (i.e., to maximize the conformational entropy). This enthalpy-entropy balance governs the microphase separation in block copolymers. Experimentally, the block copolymer phase behavior can be controlled by the selection of the composition and the architecture of the block copolymer, the overall degree of polymerization and the segment–segment interaction parameter [38–41]. The schematic illustrations of the most common, classical and some complex phases are depicted in Fig. 4, illustrating the domains of the minority phase. In the enlarged view of the lamellar morphology, the self-assembly of individual molecules within the domains is shown where blue and red parts represent the two chemically distinct blocks [42]. The phase diagram of diblock copolymers was theoretically predicted using self-consistent mean-field theory (Fig. 5a) [43,44]. Experimental phase diagrams are found to generally be similar to the calculated one, except that they appear more asymmetric. The experimentally determined phase diagram of polystyrene-block-polysoprene (PS–b–PI) is represented in Fig. 5b [45,46]. As the number of blocks in the copolymer increases, possibilities for novel, complex structures are created. Several examples include three-domain lamellae, core–shell or alternating spheres, cylinders, and gyroid, perforated structures and others [46–50]. Similarly, complex polymer architectures give rise to new, intriguing morphologies. For instance, a star-shaped architecture results in Archimedean tiling pattern morphologies that are unique for this type of copolymers [51].

When additional components are combined with copolymer systems, the obtained complexes exhibit interesting phase behavior. Supramolecular complexes are defined as assemblies of molecules held together by non-covalent interactions. A specific example of a supramolecular complex is the so-called comb-coil block copolymer in which one of the blocks, namely the comb block, contains side chains that are non-covalently linked to the
main chain, usually via hydrogen bonds. A supramolecular complex of polystyrene-block-poly(4-vinylpyridine) (PS-b-P4VP) and pentadecylphenol (PDP) has been extensively studied in the group of Ikkala and our group[52–55]. Each PDP molecule contains a hydroxyl group which can act as a hydrogen-bond promoting group and each repeating unit of P4VP contains a nitrogen atom that can act as a hydrogen-bond accepting group. Therefore, if PDP is added to P4VP, hydrogen bonds are formed between the nitrogen of the pyridine ring and the hydrogen on the phenol ring. The chemical structure and the schematic representation of the PS-b-P4VP(PDP) supramolecule is illustrated in Fig. 6a. Non-polar PDP alkyl tails microphase separate from P4VP in such a way that PDP molecules align perpendicularly to the P4VP homopolymer which gives rise to the formation of two length scale structures-within-structures (Fig. 6b). The variation in the PDP content allows for the simple tuning of the long-length scale morphology of the complex[56–58]. Furthermore, PDP can be selectively removed by an appropriate solvent (e.g., ethanol) resulting in nanoporous polymer material [59,60].

4. What do block copolymers and butterflies have in common?

The gyroid morphology was first identified in 1967 in strontium soaps by Luzzatti and Spegt [61]. Three years later, this bicontinuous triple periodic minimal surface was mathematically described by the NASA physicist Alan Schoen and named “gyroid” [62]. The single gyroid contains the percolated network and matrix phases and it is characterized by the $I4_32$ symmetry, whilst the double gyroid with $l2_3d$ symmetry has two parallel network phases that are related by inversion (Fig. 7) [63]. The transmission electron micrographs typical for a bicontinuous polymer phase were published already in 1976, but the authors did not comment on the bicontinuous nature of the system [64]. The bicontinuous ordered polymer structure was reported for the first time in 1986 by Alward et al. in a star-block copolymer comprised of PS-b-PI arms [65]. Thomas et al. described the observed morphology as the ordered bicontinuous double diamond (OBDD) with $Pn3m$ symmetry [66]. However, theoretical calculations that followed did not provide the evidence that the OBDD is an equilibrium structure in AB diblock copolymers [67–70]. The gyroid morphology in block copolymers has been described for the first time in 1994 [71,72]. Hajduk et al. analyzed a linear diblock copolymer PS-b-PI containing 37 wt% PS, using small angle X-ray scattering (SAXS) and transmission electron microscopy (TEM). TEM images suggested the bicontinuous morphology, but SAXS data were inconsistent with the OBDD structure. Peaks in the SAXS pattern were found at the positions $2\theta/6, 2\theta/8, 2\theta/20, 2\theta/22, 2\theta/32$ and $2\theta/38$ which are allowed by twelve cubic space groups. One of them, the $la3d$ space group, has additional five allowed reflections that were
not present in the experimental Bragg pattern, whilst the others have seven or more. A space filling model with $Ia\overline{3}d$ symmetry was generated and used to predict SAXS and TEM results. They found a good agreement between the predicted and the experimental data [71]. Theoretical calculations that followed confirmed that the double gyroid can be an equilibrium structure in AB diblock copolymers [43]. In more complicated block copolymer systems, such as block copolymer blends, variations of double gyroid morphology have been discovered. The core–shell double gyroid phase is one example in which gyroid networks are built out of two different polymer blocks, one being a core and the other a corona of gyroid networks that are embedded in the third polymer block that represents a gyroid matrix [73].

The double gyroid as an isotropic structure with continuous triple periodic percolated domains appears to be superior to classical morphologies (lamellar, cylindrical or spherical) for a number of technological applications. For instance, the gyroid structure, due to the high interfacial area per specimen volume, could be exploited for gas separation purposes [74]. Shefelbine et al. prepared ABC triblock copolymer with the core–shell double gyroid morphology. Glassy polystyrene was chosen as a middle block that separated two rubbery end blocks — polyisoprene and polydimethylsiloxane. A gas mixture would enter one rubbery block, characterized by high permeability and low selectivity. Further, it would diffuse selectively through a glassy layer and would, finally, be collected from the second rubbery layer. Furthermore, due to the ordering and periodicity in three dimensions, the gyroid structure is an excellent candidate for photonic crystal applications [75–77]. Urbas et al. [76] examined the optical properties of the gyroid forming PS-b-PI block copolymer with spacing comparable to the wavelength of visible light. Furthermore, the gyroid forming block copolymers could be used for templating purposes: one block could be selectively removed which results in a porous gyroid template that can be, for example, backfilled with an inorganic material. This will be further elaborated in Section 5 and supported with examples in Section 6. For templating purposes, the gyroid is preferable over anisotropic block copolymer phases (e.g., cylindrical) which often require laborious alignment procedures prior to the application [78–80].

The gyroid structure appears also in nature, namely in butterfly wing scales, and it is responsible for the shimmering wing colors of several butterfly species. Saranathan et al. [81] investigated 3D

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**Fig. 7.** (a) The single gyroid morphology with $I4\overline{1}32$ symmetry, (b) the double gyroid morphology with $Ia\overline{3}d$ symmetry. Adapted with permission from Ref. [116].
photonic nanostructures in five butterfly species by SAXS, TEM and SEM and identified their morphology as the single gyroid with \(I4_132\) symmetry (Figs. 8 and 9). The butterflies initially develop the thermodynamically favorable double gyroid phase that consists of two networks, different in composition, embedded in a matrix. Chitin is then deposited on one of the two networks and after the cell dies, the chitin-air structure is left behind. The morphology of the porous chitin structure is single gyroid which is optically more efficient than the double gyroid one. This structure could be mimicked in gyroid forming ABC terpolymers, with two chemically distinct networks [82]. A single gyroid template could be provided by the selective removal of one of the networks.

5. Block copolymer template directed preparation of metal nanofoams

The nanoscale ordering in block copolymer structures can be extended to inorganic materials with more appealing properties, e.g., metals. This process generally encompasses several steps (Fig. 10):

1. Selective removal of the sacrificial polymer block,
2. Backfilling of the porous structure with inorganic material (e.g., metal),
3. Removal of the remaining polymer template.

The first two steps can be combined into one by selective inclusion of inorganic compounds, e.g., metal ions, in one of the copolymer blocks [83].

5.1. Selective removal of the sacrificial polymer block

In recent years, many studies successfully demonstrated the selective degradation of polymer blocks from a self-assembled block copolymer structure. Polydiene blocks, such as polyisoprene (PI) or polybutadiene (PBD), can be etched away via ozonolysis [84–86]. Chen et al. [86] examined the ozonolysis efficiency in blends of block copolymer PS-\(b\)-PI and homopolymer PS and found a strong dependence on the morphology of the PI microdomains. The degree of ozonization after 24 h reaction demonstrated a decreasing trend from 90, 80, 70 to 50% if the PI microdomains transformed from lamellae, to gyroid network, to cylinders, to spheres, respectively, in the PS matrix, which was attributed to the decrease of the contact area PI/ozone and the accessible volume of PI microdomains.

UV irradiation can be employed for selective removal of PMMA blocks. The exposure of PS-\(b\)-PMMA block copolymer to UV light leads to degradation of the PMMA phase and simultaneous cross-linking of the PS phase [87]. If a blend of block copolymer PS-\(b\)-PMMA and homopolymer PMMA is used as a precursor, the nanoporous structure can be generated in two ways: (1) all PMMA (PMMA block and PMMA homopolymer) can be removed by UV irradiation or (2) only PMMA homopolymer can be removed using a selective solvent, namely acetic acid [88].

Polyacetic acid (PLA) blocks can be readily removed from a block copolymer (e.g., PS-\(b\)-PLA) by a mild aqueous base [89–91]. From crystalline-amorphous block copolymers, such as polyethylene-\((b\)-block-poly(styrene-\((b\)-polystyrene-\((b\)-poly(vinylidene fluoride)-block-poly(styrene-\((b\)-polystyrene-\((b\)-PVDF-\((b\)-PS), the amorphous component can be selectively etched with fuming nitric acid [92,93]. Reactive ion etching can be used to generate porous structures from block copolymer precursors with one etch-resistant block [94–97]. Finally, amphiphilic molecule PDP can be removed from a supermolecular complex containing P4VP/PDP comb-block by selective dissolution in ethanol, as already mentioned in Section 3, resulting in porous polymer template [54,59,60].

5.2. Backfilling of the porous structure with inorganic material

Metal oxides (e.g., SiO\(_2\) or TiO\(_2\)) can be introduced inside the pores of a polymer template via the sol–gel process. In the sol–gel process, the starting colloidal solution — sol, typically composed of metal alkoxides or metal salts, fills pores of a polymer template and subsequently, via hydrolysis and polycondensation reactions, transforms into gel — an integrated network composed of discrete colloidal particles or polymer networks [98]. Hsu et al. prepared a polystyrene-\((b\)-block-poly(-l-lactide) (PS-\(b\)-PLLA) block copolymer with double gyroid morphology and then selectively removed the PLLA block via hydrolysis [99,100]. The obtained porous PS network was filled with amorphous SiO\(_2\) via the sol–gel process. The SAXS pattern, TEM and electron tomography (Fig. 11a) images of the PS/SiO\(_2\) nanohybrid confirmed its double gyroid morphology. After the polymer removal, a gyroid forming SiO\(_2\) network was exposed (Fig. 11b) and its porosity was calculated from nitrogen adsorption.

Fig. 9. SAXS patterns of the nanostructures of five different butterfly species indicating the single gyroid morphology with \(I4_132\) symmetry. Reprinted with permission from Ref. [81].

Fig. 10. Block copolymer template directed preparation of ordered inorganic nanostructures: 1. selective removal of the sacrificial polymer block, 2. backfilling of the porous structure with inorganic material (e.g., metal), 3. removal of the remaining polymer template.
measurements to be 66%. The refractive index of dense amorphous SiO$_2$ is rather low and it further decreases in the porous sample. The refractive index of the prepared gyroid SiO$_2$ network was estimated to be about 1.1, thus it appeared suitable for optical applications such as antireflection structures [99]. The hydrolysis reaction rate in the sol–gel process towards SiO$_2$ is exceptionally low which allows for complete diffusion of sol into the template pores before the gelation starts. On the other hand, titanium alkoxides are not as stable and pores will be blocked by gel before sol completely penetrates the template. Therefore, to fabricate the gyroid PS/TiO$_2$ nanohybrid, one has to carefully control the reaction conditions. A thermal treatment was employed to remove PS and expose the free-standing, gyroid anatase TiO$_2$ (Fig. 12). Its photocatalytic efficiency was determined by evaluating the degradation of methylene blue under UV irradiation. It was found that the decomposition rate is about two times higher in comparison to commercial TiO$_2$ powder [100].

Electrochemical and electroless plating are commonly used techniques to deposit metals onto or into templates [101–104]. To electrochemically deposit metal inside the pores of a polymer template, the working electrode is usually coated by a template and, together with the counter and the reference electrode, placed in the plating bath and connected to an external source of current. To completely plate the sample, the surface of the underlying working electrode should be accessible to the plating bath, which may appear difficult for thicker polymer templates with many defects or highly complex structure. Also relatively simple polymer templates, such as the cylindrical one, require proper alignment prior to electrochemical plating. In electroless plating, a polymer template is immersed into the electroless plating bath and metal is deposited via an autocatalytic process on the surface of the template. Electroless plating does not require an external current source and allows plating onto complex shapes and through pores. Furthermore, non-conductive surfaces, such as polymers or ceramics, can undergo electroless plating after the appropriate treatment. The pretreatment usually consists of two steps: (1) sensitization in SnCl$_2$/HCl, when Sn$^{2+}$ ions adsorb onto the template surface, (2) activation in PdCl$_2$/HCl, when Pd sites are formed through the following redox reaction: $\text{Sn}^{2+} + \text{Pd}^{2+} \rightarrow \text{Sn}^{4+} + \text{Pd}^0$. The Pd sites act as catalysts during the electroless metal plating procedure. In electroless deposition, metal ions, M$^{n+}$, are reduced by the reducing agent, R$^n_-$, and the process can be simply described as: $\text{M}^{n+} + \text{R}^n_- \rightarrow \text{M}^0 + \text{R}^{n-}$. The process is autocatalytic, i.e., it is catalyzed by metal being deposited. Besides the source of metal ions and the reducing agent, the electroless plating bath usually contains the complexing agent, that acts as a buffer and reduces the concentration of free metal ions, and the stabilizer, that prevents the decomposition of the plating solution.

### 5.3. Removal of the remaining polymer template

There are several methods to remove polymer from the polymer/inorganic nanohybrid, such as pyrolysis [30, 60], dissolution [105, 106], UV degradation [99, 100], etc.

Pyrolysis involves the decomposition of polymer at elevated temperature, leaving behind an inorganic material with preserved morphology. Inorganic materials may as well undergo changes at elevated temperatures. For instance, thermal treatment will increase the crystallinity of TiO$_2$, thus, amorphous TiO$_2$ may transform to one of its crystalline forms: rutile, anatase, and brookite [100]. Another example is nanoporous gold that undergoes thermal coarsening due to the recrystallization in solid state at elevated temperatures [107]. Therefore, the applicability of pyrolysis for polymer template removal depends on the type of inorganic material and its purposes.

### 6. Examples of metal nanofoams prepared by block copolymer templating

There have been a few reports in the literature demonstrating the replication of complex block copolymer structures into the ordered metal phase. In the following paragraphs, we describe several prominent examples and focus on the fabrication of gyroid metal nanofoams.

Crossland et al. [105] investigated the thin film morphology of poly(4-fluorostyrene)-block-poly(D,L-lactide) (PFS-b-PLA) that in

![Fig. 11. The reconstructed 3D images of: (a) gyroid PS/SiO$_2$ nanohybrid, (b) gyroid SiO$_2$ composed of two parallel networks related by inversion (red and green). Reprinted with permission from Ref. [99]. Copyright 2010 American Chemical Society.](image-url)
bulk acquires the cylindrical morphology at 150 °C and the gyroid morphology at 180 °C. When the thin film was annealed for 35 h at 180 °C and then cooled down to room temperature over 3 h, it was found to adopt the gyroid phase in the entire sample volume. The PLA block was then selectively removed by hydrolysis in a mild aqueous base and platinum was deposited into the remaining pores via electrochemical deposition. Final removal of the PFS block, by dissolution in toluene or exposure to O₂ plasma, resulted in isolated mounds or a continuous freestanding array of a gyroid Pt replica (Fig. 13) depending on the properties of the underlying substrate. Smooth Au-coated silicon substrate favors the formation of isolated mounds of Pt replica (Fig. 13a and c). Chemical modification of the Au-coated silicon substrate using a self-assembled monolayer (SAM) of 1-undecanethiol induced much higher substrate coverage (Fig. 13b and d). Presumably, in comparison to Au, SAM is more energetically neutral to copolymer blocks and does not favor the PFS phase. Surface topography is also found to significantly influence the replication of gyroid Pt over large areas. FTO-glass or Au-coated FTO glass, with the amplitude of surface roughness (ca. 50 nm) comparable to the copolymer domain spacing, led to the complete surface coverage and formation of a continuous freestanding array of gyroid Pt replica.

Vignolini et al. [108] demonstrated the creation of a gold optical metamaterial starting from triblock copolymer polyisoprene-block-polystyrene-block-polyethylene oxide (PI-b-PS-b-PEO). The polymer adopted the double gyroid morphology with two chemically distinct networks (PI and PEO) in a matrix of polystyrene. The PI network was selectively removed by UV irradiation and subsequent rinsing with ethanol. Then, the polymer template was backfilled with gold by electrochemical deposition and, finally, the remaining template was removed by plasma etching resulting in a single gyroid, nanoscale ordered Au foam. The preparation procedure is schematically depicted in Fig. 14. The single gyroid Au replica was examined under linearly polarized light (Fig. 15). The tested area contained one grain boundary that divided the sample into two domains with the same orientation perpendicular to the substrate surface and the twinning in lateral direction. At the twinning boundary, the angle between [100] directions was found to be about 105° (Fig. 15a). Both domains reflected the identical color when the polarization of incident light was perpendicular to the grain boundary (Fig. 15b). If the polarization of incident light was parallel to the [100] direction of one of the domains (simultaneously, it is almost perpendicular to the [100] direction of the other domain), the largest optical contrast was observed (Fig. 15c). The absorption peak was found to tune sinusoidally between 525 and 630 nm with the polarization direction (Fig. 15e). For the polarization along the [100] direction in the single gyroid Au, the absorption peak appeared at short wavelengths and is red-shifted for polarization in perpendicular direction (Fig. 15d). Additionally,
the Au replica exhibits optical chirality. Transmission measurements with circularly polarized light were performed while the sample was rotated around the [110] surface normal. The maximum transmission was observed for two rotation angles in each full turn, i.e., when the optical axis coincides with the gyroid [111] chiral direction.

Warren et al. [109] demonstrated an alternative route towards ordered metal nanofoams via direct self-assembly of block copolymers with ligand-stabilized metal nanoparticles. Block copolymer polyisoprene-block-poly(dimethylaminoethyl methacrylate) PI-b-PDMAEMA and ligand-stabilized Pt nanoparticles were dissolved in chloroform. The solutions were transferred to Al dishes and heated at 50 °C for approximately 1 h until the solvent completely evaporated. After annealing at 130 °C for 2 days under vacuum, samples adopted the ordered morphology. A cylindrical sample was exposed to rapid pyrolysis and converted to an ordered porous Pt composite. Finally, carbon was removed using Ar plasma and the Pt nanofoam was exposed. The electrical conductivity of the Pt composite was found to be rather high (400 S/cm). This strategy can be extended to different metals and block copolymer morphologies.

In 2011, our group reported the formation of a freestanding gyroid nickel nanofoam via electroless plating of a block copolymer based template [60]. At the same time, Hsueh et al. [106] published the preparation of gyroid Ni replica starting from PS-b-PLLA block copolymers. The unique aspects of our approach will be described next.

7. Supramolecular route to gyroid Ni nanofoams

As mentioned in Section 3, supramolecular complexes of block copolymers containing the P4VP block and amphiphilic PDP molecules have been thoroughly examined in the group of Ikkala and our group. We recently attempted to use supramolecular complexes of triblock copolymer poly(tert-butoxystyrene)-block-poly(styrene)-block-poly(4-vinylpyridine) PtBOS-b-PS-b-P4VP and amphiphilic PDP molecules that self-assemble into a core–shell double gyroid morphology as precursors to well-ordered metal nanofoams [59]. The TEM image of the representative core–shell double gyroid PtBOS-b-PS-b-P4VP(PDP) complex (Fig. 17) displays a typical gyroid double-wave pattern. The core channels are emptied by ethanol treatment and subsequently electroless nickel plated. Channels located close to the surface are filled with nickel whilst ones in the middle of the film remain empty as shown in Fig. 18. This seems to be induced by narrow channels (average pore diameter is determined by mercury porosimetry to be 12 nm) and the fast-growing nickel deposit near the free surface that hinders the reagents’ diffusion towards the bulk [110].

Subsequently, we evaluated a supramolecular complex PS-b-P4VP(PDP) as a precursor to a gyroid nickel nanofoam. Unique to this approach is that the gyroid matrix, or more precisely, part of the matrix, was removed from the complex to render a nanoporous template. Our procedure is schematically summarized in Fig. 19.
Fig. 15. Linear dichroism of the Au gyroid: (a) SEM image of the examined sample area, (b, c) Optical reflection images obtained with linearly polarized light, perpendicular to the grain boundary (b) or parallel to the [100] gyroid direction of the domain 2 (c); (d) Reflection spectra for area 1 with the polarization perpendicular (red) and parallel (dark grey) to the [100] gyroid direction, (e) Dependence of the extinction (reflectivity) peak position and width on the angle between the [100] gyroid direction and incident light polarization. Reprinted from Ref. [108].

Fig. 16. The TEM image of the electroless nickel plated polymer template with the double gyroid morphology. The inset shows a 25 nm nanochannel coated with 10 nm nickel nanoparticles. Reprinted with permission from Ref. [85]. Copyright 1997 American Chemical Society.

Fig. 17. TEM micrograph of the PtBOS-b-PS-b-P4VP(PDP)x complex with x = 0.7, f_{BOS} = f_{PS} = 0.397 and M_{total} = 88 300 g mol^{-1}. P4VP(PDP)x domains are selectively stained with iodine and appear dark in the image. The double-wave pattern indicates the cross-section through the gyroid [211] plane. Adapted from Ref. [59].
parent block copolymer PS-b-P4VP and hydrogen bonding PDP molecules are dissolved in chloroform and solvent annealed for one week (Fig. 19a). The supramolecular PS-b-P4VP(PDP)_x complex (x represents the ratio between the number of PDP molecules and the number of P4VP monomer units) with the double gyroid morphology consisting of PS networks and a P4VP(PDP)_x matrix is obtained (Fig. 19b). Then, the complex is immersed in ethanol, a selective solvent for PDP, which leads to the formation of an open network structure consisting of a PS core and a P4VP corona (Fig. 19c). Electroless plating is employed to backfill the porous polymer template with nickel (Fig. 19d) and finally, the remaining polymer is removed by pyrolysis and a well-ordered gyroid nickel replica is exposed (Fig. 19e).

Contrary to previously described examples, our porous polymer template PS(core)-P4VP(corona) is obtained by partial removal of the gyroid matrix and represents a double network occupying more than 50 vol%. Removal of the matrix from the gyroid forming conventional diblock copolymers would result in a template occupying only ca. 35 vol%, which is not suitable for the formation of a metallic replica with porosity high enough to fulfill the general requirement for metal nanofoams (porosity higher than 50 vol%). Furthermore, the removal of the sacrificial block from conventional block copolymers, such as PS-b-PLA or PS-b-PEO [111], results in a hydrophobic PS surface that usually requires modification prior to electroless plating in order to achieve the uniform metal deposition [112]. In our approach, the presence of the P4VP hydrophilic corona facilitates the penetration of water-based plating reagents into the porous template and enables successful metal deposition.

The morphology of supramolecular complexes PS-b-P4VP(PDP)_x is examined by TEM and SAXS. Fig. 20a and b displays typical gyroid patterns of a representative supramolecular complex: the double-wave and the wagon-wheel patterns that are known to represent projections through the [211] and the [111] plane of the gyroid unit cell, respectively. The PS block domains appear bright whilst the P4VP(PDP)_x block domains appear dark due to iodine staining. Fig. 20c represents the double-wave pattern of a different gyroid sample of which the periodicity is decreased with a factor of 2. SAXS peaks at positions: √6q*, √8q*, √14q*, √22q*, and √50q* confirm the bicontinuous Ia3d morphology of the sample (Fig. 20d).
porosimetry. The BET specific surface area of 104 m² g⁻¹ is rather high, the template occupies almost 60 vol%, the average pore diameter is 40 nm, and pore size distribution is very narrow (Fig. 21).

Prior to electroless plating, the Pd catalyst is deposited onto the gyroid template surface to direct the selective nickel ion reduction. Successively, nickel metal fills the pores of the polymer template. Fig. 22a represents the TEM micrograph of the unstained nickel plated gyroid sample and the contrast in the image originates from metal deposited in the nanochannels. Additionally, the characteristic wagon-wheel pattern confirms the preservation of the double gyroid morphology during the processing steps. HR TEM micrographs (Fig. 22b, c) show relatively large, interconnected Ni crystallites and EDX analysis (Fig. 22d) reveals the chemical composition of the plated sample. As expected, prominent carbon and nickel peaks are observed, together with the oxygen peak that indicates the oxidation of the nickel nanofoam when stored in air.

Finally, the remaining polymer is decomposed by heating isothermally at 350 °C for at least half an hour, leaving the nickel network intact. The exposed nickel replica preserves the inverse gyroid morphology, the shape of the gyroid matrix, as confirmed by SEM (Fig. 23). This shape is the negative of the gyroid networks shape, therefore their surface area is equal and their porosities sum up to 100%. Additionally, this a one-part shape whilst its negative consists of two interpenetrated parts.

In the work that followed, we performed a comprehensive study of the phase behavior of supramolecular complexes PS-b-P4VP(PDP)ₓ in the region of the phase diagram where the gyroid
morphology with the $P4VP(PDP)_x$ as a majority block was expected [57]. By varying the molar mass and the composition of the starting PS-$b$-$P4VP$ copolymer and the amount of PDP in the complex, double gyroid templates, and consequently well-ordered gyroid metal nanofoams, with tunable feature sizes and porosities can be obtained. The phase behavior of our supramolecular complexes PS-$b$-$P4VP(PDP)_x$ on the other side of the phase diagram, i.e., in the region where the gyroid phase with the $P4VP(PDP)_x$ as a minority block was expected, appeared to be quite surprising [58]. Instead of the expected GYR phase, a broad region of the hexagonally perforated layer (HPL) phase is found. The HPL is a metastable phase in conventional block copolymers, but an equilibrium phase in PS-$b$-$P4VP(PDP)_x$ complexes with the $P4VP(PDP)_x$ as a minority component. This untypical phase behavior can be explained by the presence of PDP "side chain" molecules in the minority component of the complex, where the HPL morphology relieves the packing frustrations.

8. Summary and outlook

In this feature article we focus on the preparation and utilization of a new class of materials, metal nanofoams, and discuss the possibility of using block copolymers as a tool for ordered metallic nanostructures. Metal nanofoams have the potential to outperform classical materials in a number of fields: hydrogen storage, production of novel batteries, catalysis, actuating, etc. Several
techniques for their preparation have been developed and most of them result in highly porous, but immensely disordered metal nanofoams. Block copolymers can microphase separate into well-ordered nanostructures from which one of the blocks can be selectively etched. The resulting porous template can be backfilled with metal and subsequently, the remaining polymer can be removed leading to well-ordered metal nanofoams. Even more appealing for this purpose than conventional block copolymers, are supramolecular complexes where side chains hydrogen-bonded to one of the blocks can be removed by simple dissolution. Additionally, the morphology of these complexes can be easily tailored by alteration of the amount of added side chains. The bicontinuous gyroid morphology appears as the most suitable nanoscale architecture to provide the interconnected, structurally isotropic metal nanofoams. Pt, Au and Ni gyroid replicas have already been reported in the literature and their application in different fields evaluated.

In the future, we expect metal nanofoam characteristics to be improved and adjusted to meet the requirements for desired applications. For instance, their structure can be tailored by employing alternative block copolymer phases as precursors, such as the orthorhombic Fdd network or plumber’s nightmare phase. Additionally, their chemical composition can be varied and alternative metals, such as Ag, Cu, Co, etc., can be deposited in the polymer template channels, leading to new ordered metal nanofoams. Furthermore, block copolymer template-directed synthesis can be combined with dealloying; from the alloy deposited in block copolymer template channels, a less noble metal can be removed, leading to a hierarchically porous ordered metal nanofoam. In addition, block copolymer templates can be oriented by, for instance, electric fields [113] or large amplitude oscillatory shearing [114,115]. Oriented block copolymer templates will lead to metal nanofoams with improved ordering which might have superior properties, as actuators, for example, in comparison to the existing metal nanofoams. The synthetic routes towards well-ordered metal nanofoams and their application potential are still poorly examined and we expect further investigation and exciting discoveries to emerge.

References

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