

Polymer Cubosomes: Synthesis, Self-Assembly and Conversion to Functional Materials

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The self-assembly of block copolymers is nowadays well understood. The resulting block copolymer morphology is mainly influenced by the packing parameter $p = \nu/a_0l_c$; where ν and l_c are the volume and the length of the hydrophobic block, and a_0 is the surface area of the hydrophilic block. If $p < 1/3$ spheres are formed, for $1/3 < p < 1/2$ cylinders are found, and a $1/2 < p < 1$ will result in vesicles.^{1,2} Exceeding a $p > 1$ results in the formation of cubosomes and hexosomes.² The polymer cubosomes have a bicontinuous pore network and a high surface area, therefore they can be used as templating platform for different materials like metal oxides and MOFs to introduce high surface area and ordered porous structure.^{3,4} Herein, we show the synthesis of poly(ethylene oxide)-based amphiphilic block copolymers via RAFT polymerization, discuss the influence of chain lengths and cosolvent on the self-assembly process. We also present an alternative formation route (redispersion) to the commonly used nanoprecipitation method. The resulting polymer cubosomes can be used as a templating platform for TiO₂, Co₃O₄ and porous carbon.

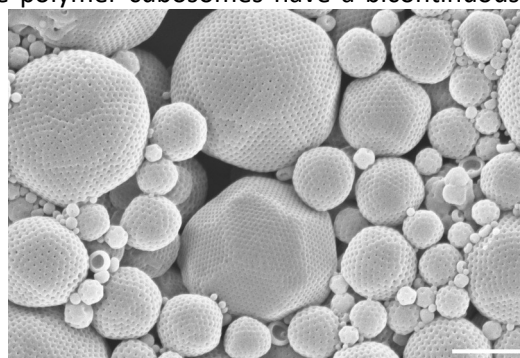


Figure 1. SEM image of block copolymer cubosomes. (scale bar 1 μ m).

Keywords: block copolymers, mesoporous materials, metal oxides, self-assembly, templating

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