

Supporting information

Preparation of structurally colored films assembled by Polystyrene@Silica, air@Silica and air@Carbon@Silica core-shell nanoparticles with enhanced color visibility

Fen Wang, Xin Zhang, Jianfeng Zhu and Ying Lin

*School of Materials Science and Engineering, Shaanxi University of Science and
Technology, Xi'an 710021, China*

Corresponding author: Fen Wang

Tel.: +8615114805183;

fax: +8602986168688.

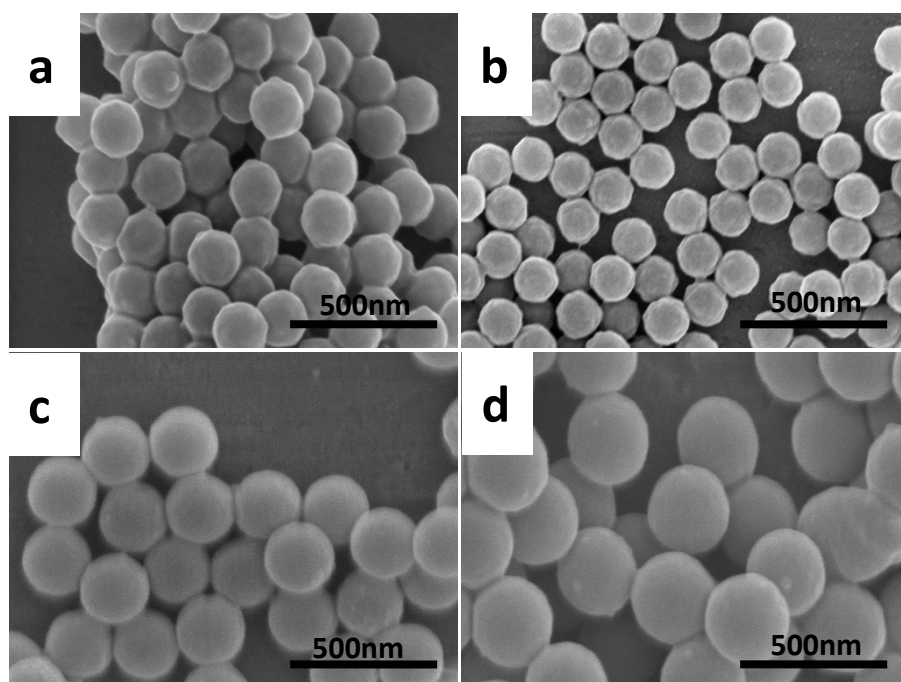


Figure S1. PS spheres of different size with a low polydispersity have been prepared by the emulsifier-free emulsion polymerization method: a) 180 ± 5 nm, b) 200 ± 5 nm, c) 240 ± 5 nm, and d) 290 ± 5 nm

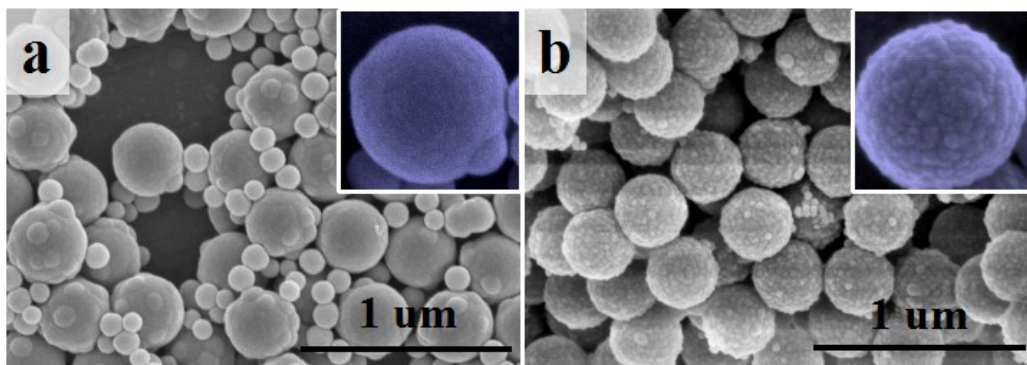


Fig. S2 (a) SiO₂ coated PS (PS was used without modification) (b) SiO₂ coated PS (PS was modified by NH₃·H₂O); inserts in (a) - (b) are high-magnification SEM images.

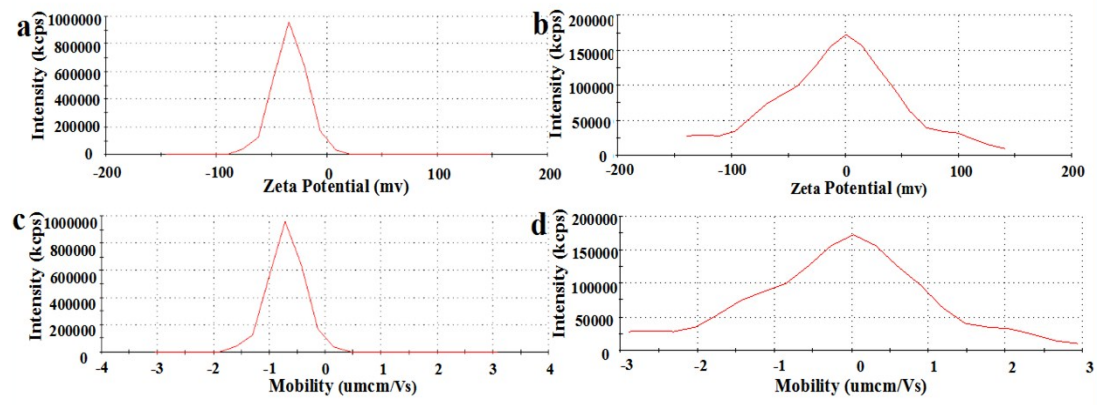


Fig. S3 Electrophoretic measurements on PS colloids: zeta potential for bare (a) and ammonium hydroxide modified (b) PS colloids; electrophoretic mobility of bare (c) and ammonium hydroxide modified PS (d) colloids.

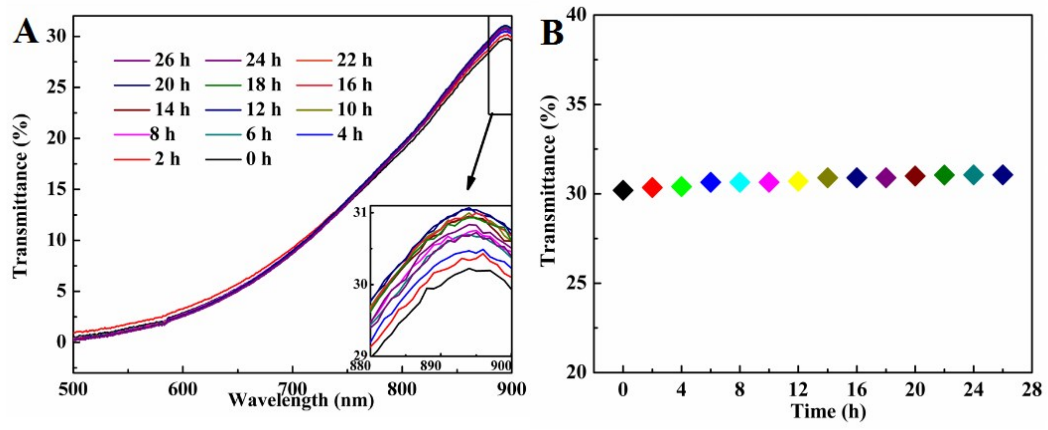


Fig. S4 Transmittance changes of the PS@SiO₂ solution over times.

Table S2. the size, and shell thickness of the core-shell particles before and after calcination

sample	Core particle	C _{TEOS} (uL)	D _{PS@SiO₂} (nm)	D _{Air@SiO₂} (nm)	Shell thickness (nm)	Ref peak(nm)
PS180	180±5	100	220 ± 5	210± 5	15	387
PS200	200±5	150	250 ± 5	240± 5	20	448
PS240	240±5	200	300 ± 5	290± 5	25	545
PS290	290±5	200	340 ± 5	330± 5	20	599