

**Octavinylsilsesquioxane-based luminescent nanoporous
inorganic-organic hybrid polymers constructed by Heck
coupling reaction**

Libo Sun, Zhiqiang Liang,* Jihong Yu*

State Key Laboratory of Inorganic Synthesis & Preparative Chemistry, Jilin

University, Changchun, 130012, P. R. China

Email: liangzq@jlu.edu.cn; jihong@jlu.edu.cn

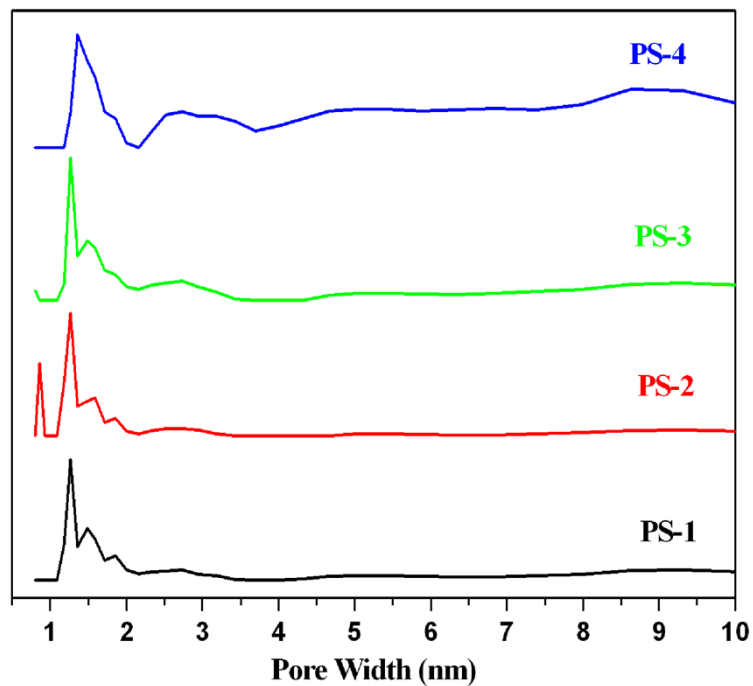


Fig. S1 The pore size distribution of **PS-n** (1-4) calculated by the NLDFT method.

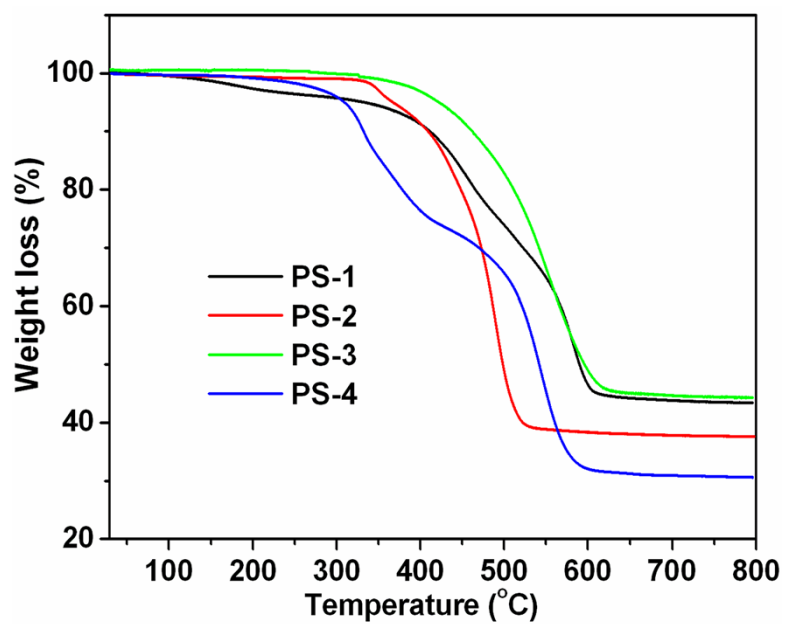


Fig. S2 TG curves for **PS-n** (1-4) under air atmosphere.

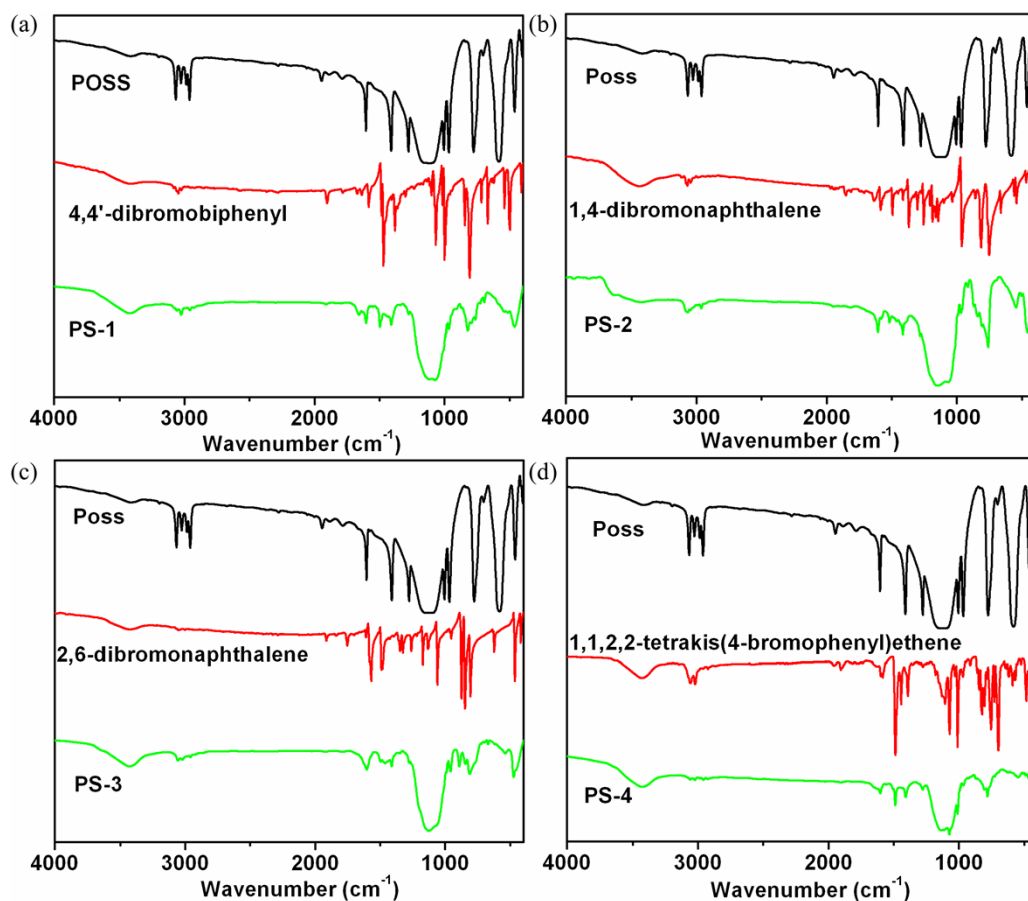


Fig. S3 FT-IR spectra of the monomers and **PS-n**: (a) monomers and **PS-1**; (b) monomers and **PS-2**; (c) monomers and **PS-3**; (d) monomers and **PS-4**.

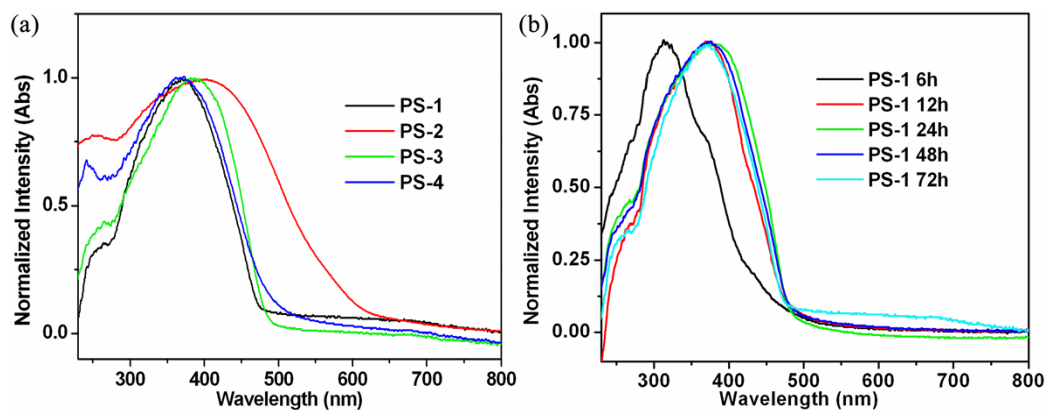


Fig. S4 (a) Solid state UV-vis absorption spectra for **PS-n** (1–4); (b) Solid state UV-vis absorption spectra for **PS-1-n** ($n = 6, 12, 24, 48, 72$ h).

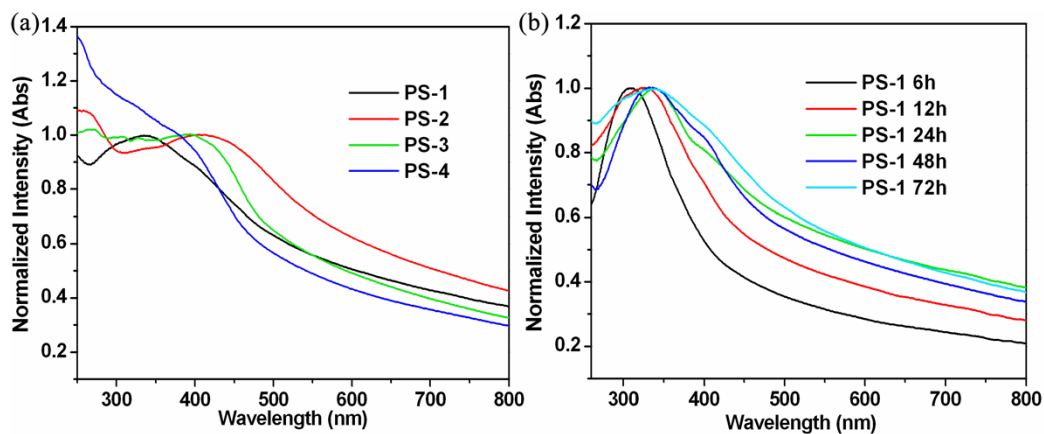


Fig. S5 (a) UV-vis absorption spectra for **PS-n** (1–4) in ethanol; (b) Solid state UV-vis absorption spectra for **PS-1-n** ($n = 6, 12, 24, 48, 72$ h) in ethanol.

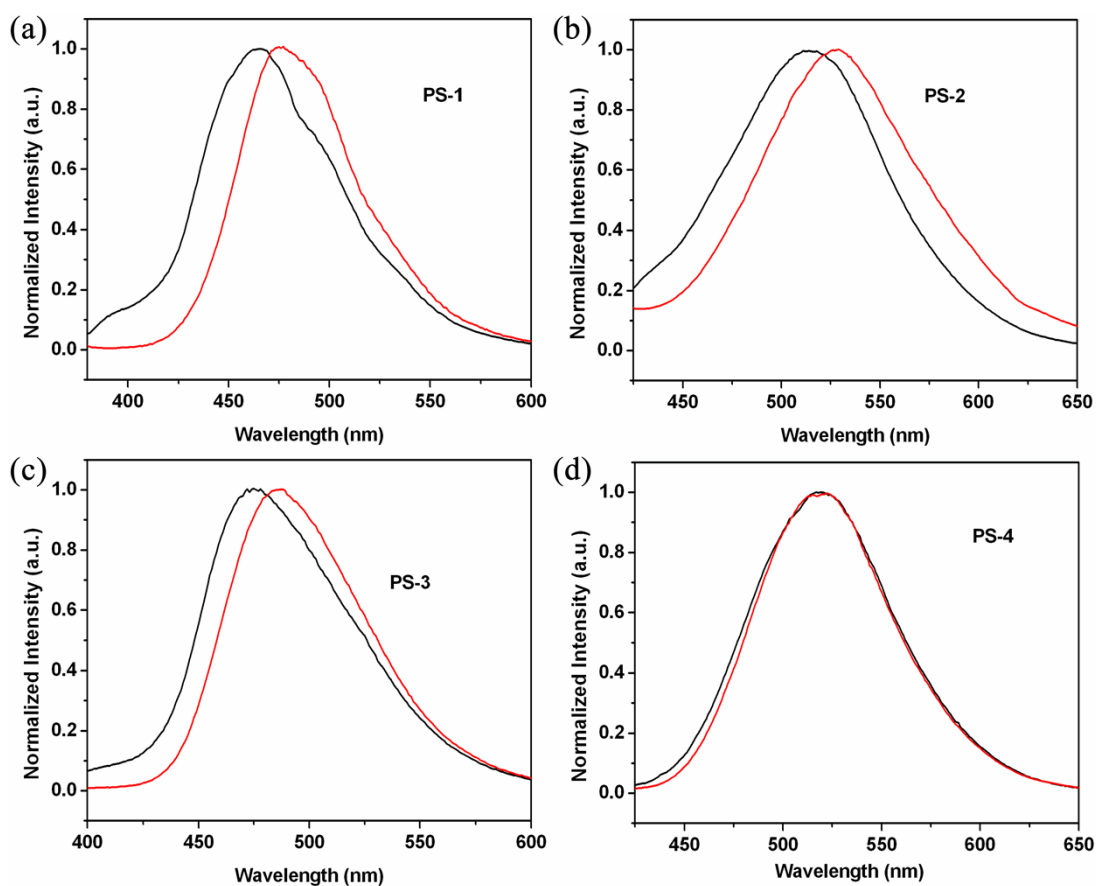


Fig. S6 Luminescent spectra of **PS-n** (1–4) in the solid state (red) and ethanol (black).

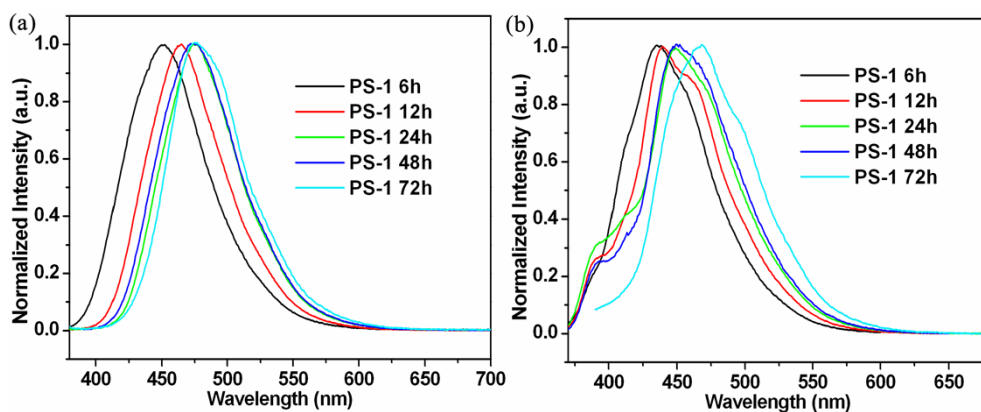


Fig. S7 Luminescent spectra of **PS-1-n** (n = 6, 12, 24, 48, 72 h) in the solid state (left) and ethanol (right).

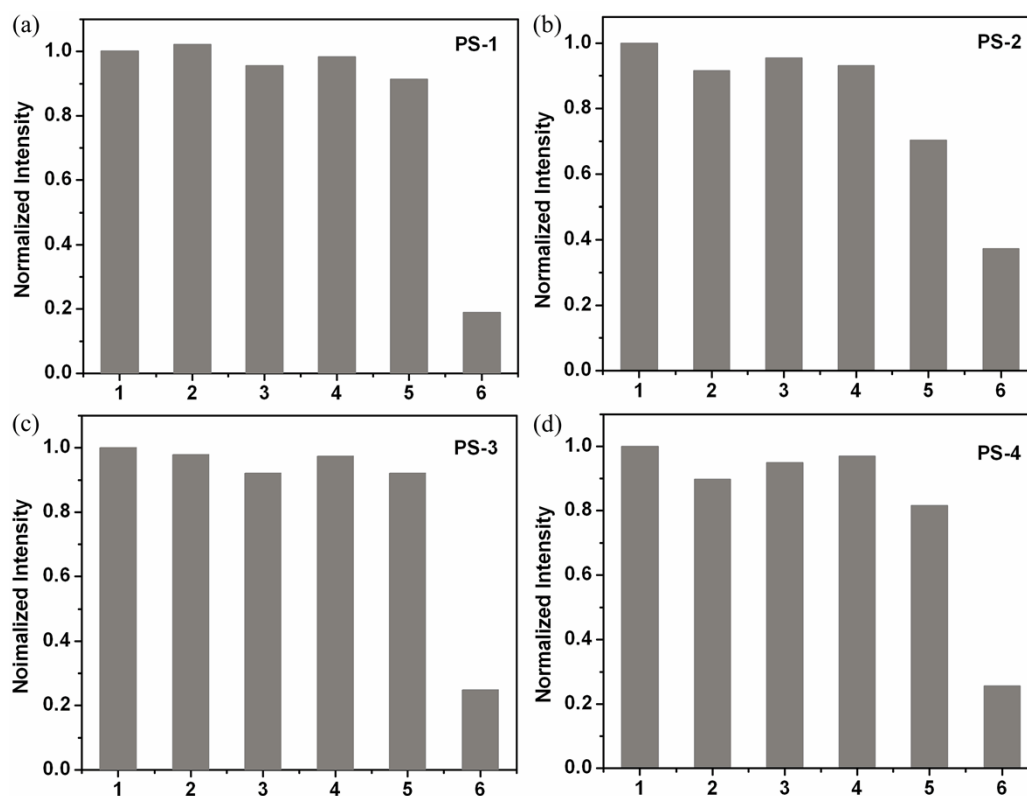


Fig. S8 The normalized luminescent intensity of **PS-n** (a: **PS-1**, b: **PS-2**, c: **PS-3**, d: **PS-4**) in ethanol (0.05 mg/mL) upon addition of ca. 46.5 μ M different analytes (1: original, 2: 2,4-dinitrochlorobenzene, 3: 2,4-dinitrotoluene, 4: 4-nitrotoluene, 5: 4-nitrophenol, 6: picric acid).

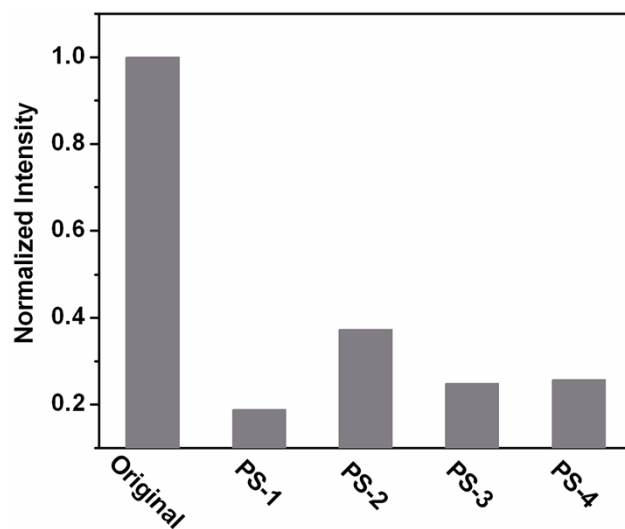


Fig. S9 The normalized luminescent intensity of **PS-n** (1–4) in ethanol (0.05 mg/mL) upon addition of ca. 46.5 μM PA.

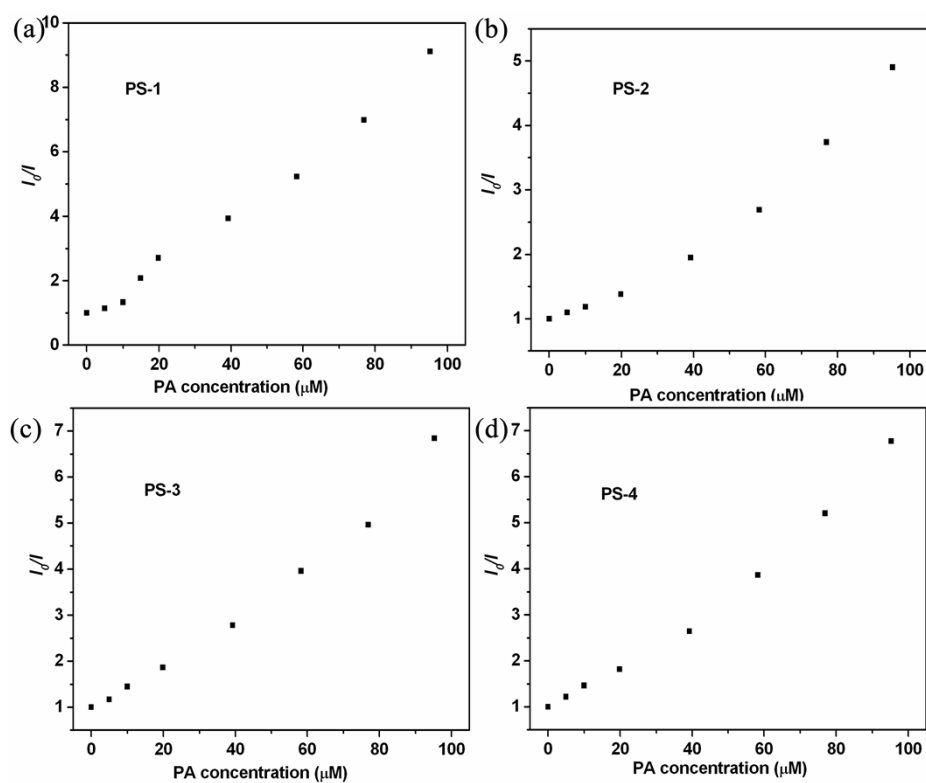


Fig. S10 The relative Stern-Volmer plots of relative PL intensity (I_0/I) versus TNP.

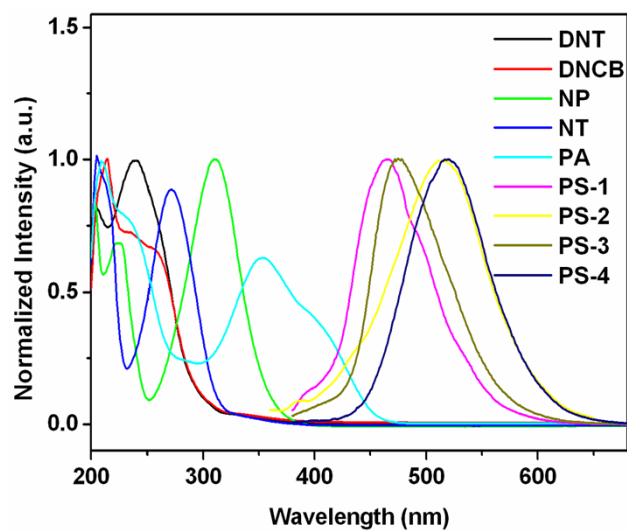


Fig. S11 Spectral overlap between the absorption spectra of analytes and the emission spectra of PS-n in ethanol.

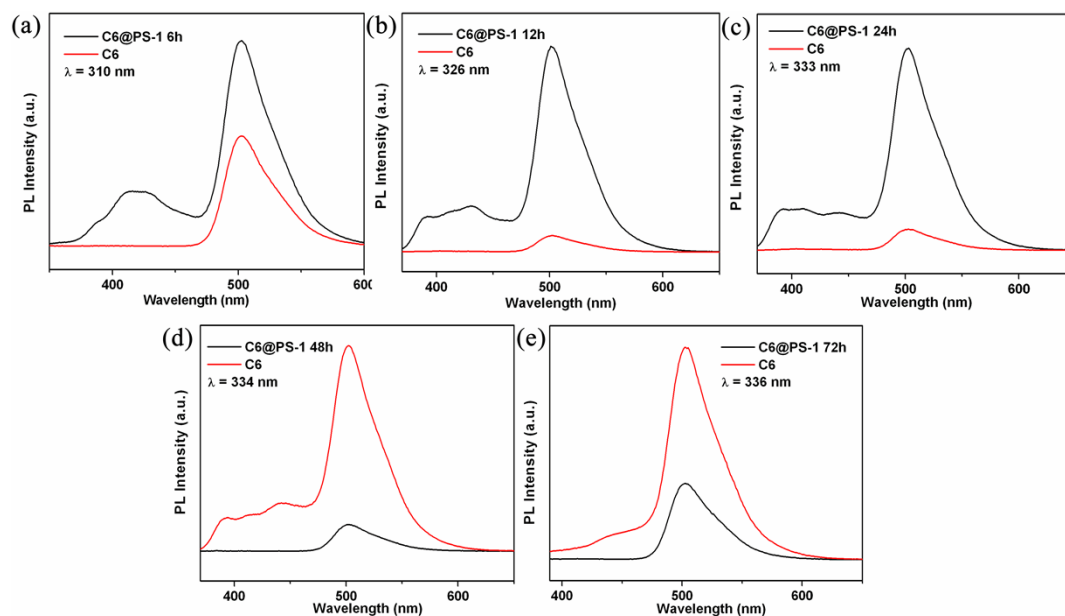


Fig. S12 Comparison emissions of C6 with and without the addition of PS-1-n in ethanol solution.

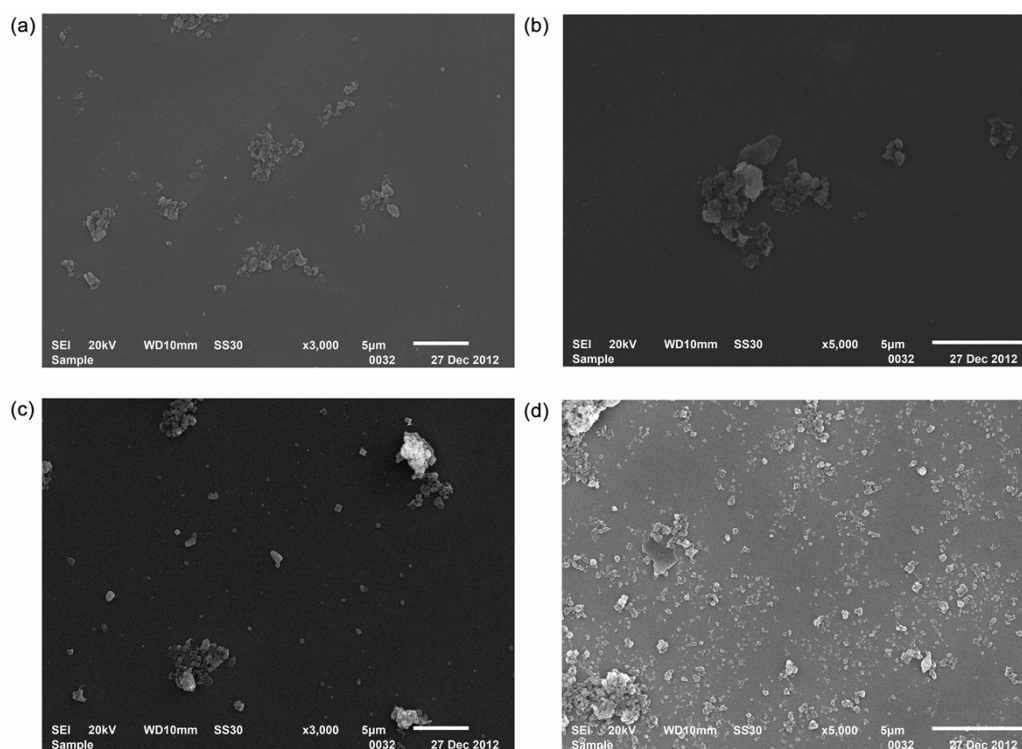


Fig. S13 Scanning electron microscope images for **PS-1** (a), **PS-2** (b), **PS-3** (c) and **PS-4** (d) at different magnifications. The samples were sputter-coated with gold before analysis.

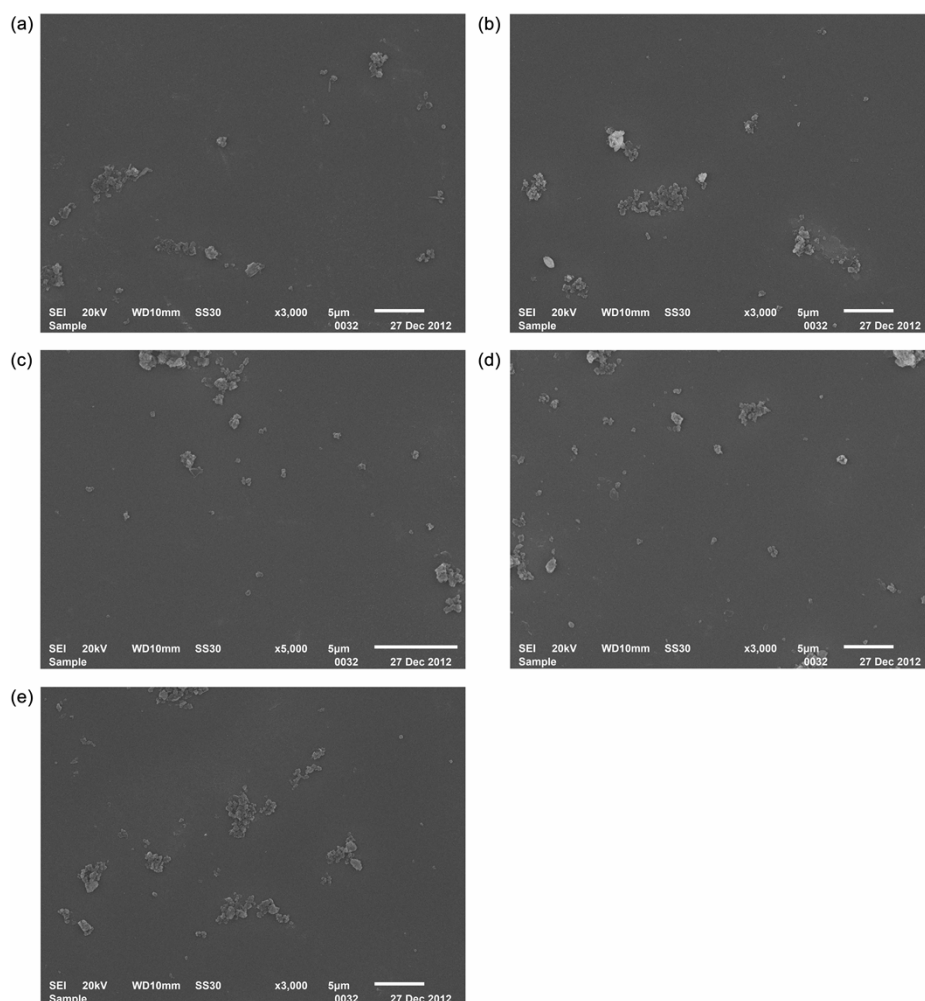


Fig. S14 Scanning electron microscope images for **PS-1** (6 h) (a), **PS-1** (12 h) (b), **PS-1** (24 h) (c), **PS-1** (48 h) (d) and **PS-1** (72 h) (e) at different magnifications. The samples were sputter-coated with gold before analysis.

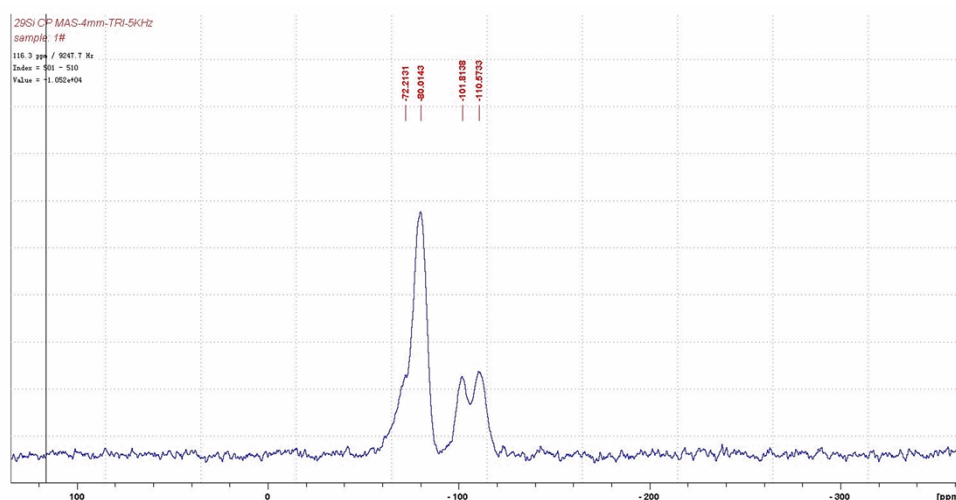


Fig. S15 The ^{29}Si NMR of **PS-1**.

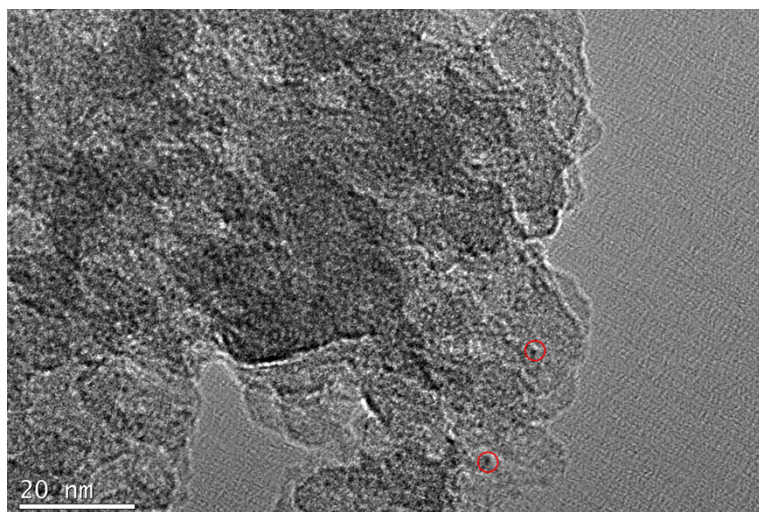


Fig. S16 The TEM image of PS-1. The red round cycles show the SiO₂ particles.

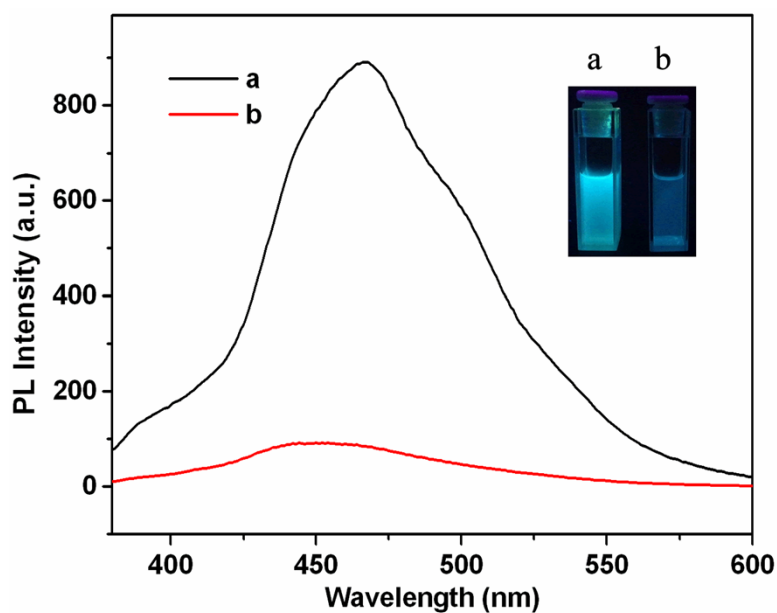


Fig. S17 Comparison emissions of PS-1 (in ethanol solution) synthesized using our condition (a) and Feng's condition (b).

Table S1 EDX analyses of PS-*n* (1-4).

PS- <i>n</i>	wt. %				at. %			
	C	Si	Br	Pd	C	Si	Br	Pd
PS-1	40.65	11.11	0.19	0.48	66.54	6.21	0.10	0.05
PS-2	44.20	5.22	13.68	0.14	71.58	3.62	3.33	0.03
PS-3	46.03	10.58	1.72	0.45	65.44	6.43	0.37	0.07
PS-4	39.65	14.28	3.8	0.18	62.78	9.67	0.90	0.06

Table S2 Porous properties of **PS-n** (1-4).

PS-n	$S_{\text{BET}}/\text{m}^2 \text{ g}^{-1}$	$S_{\text{Langmuir}}/\text{m}^2 \text{ g}^{-1}$	$S_{\text{micro}}/\text{m}^2 \text{ g}^{-1}$	$V_{\text{total}}^a/\text{cm}^3 \text{ g}^{-1}$	$V_{\text{micro}}^b/\text{cm}^3 \text{ g}^{-1}$
PS-1	550.61	752.32	212.28	0.543	0.093
PS-2	372.79	502.98	185.23	0.319	0.084
PS-3	685.63	938.08	182.85	0.760	0.077
PS-4	393.95	544.34	57.85	0.608	0.021

^a Total volume at $P/P_0 = 0.97$.

^b The micropore volume calculated from t -plot method.