

Supporting Information

Preparation of Multifunctional Hollow Microporous Organic Nanospheres via One-pot Hyper-cross-linking Mediated Self-assembly Strategy

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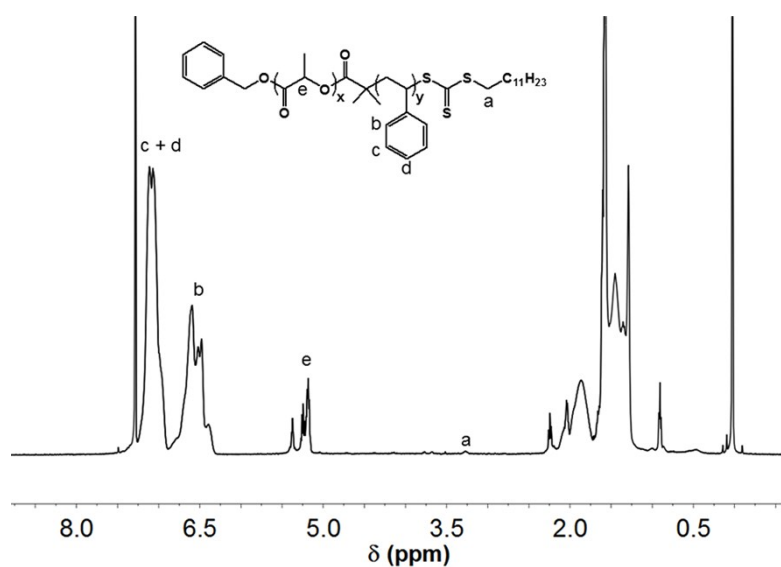


Fig. S1 ¹H NMR characterization of PLA-b-PS intermediate (x=60, y=264).

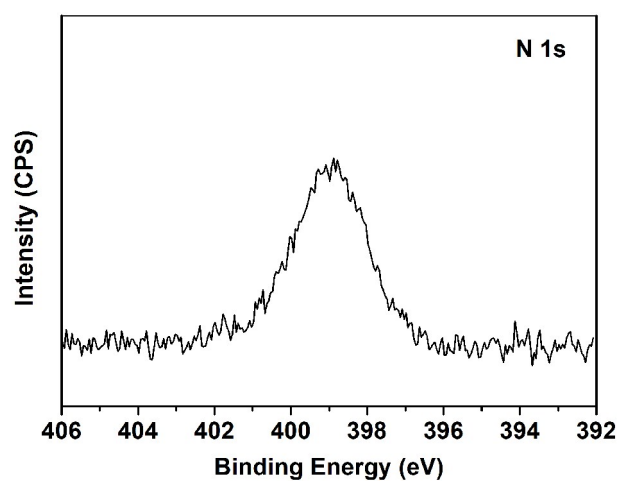


Fig. S2 XPS N 1s spectra of H-MONs-Ru.

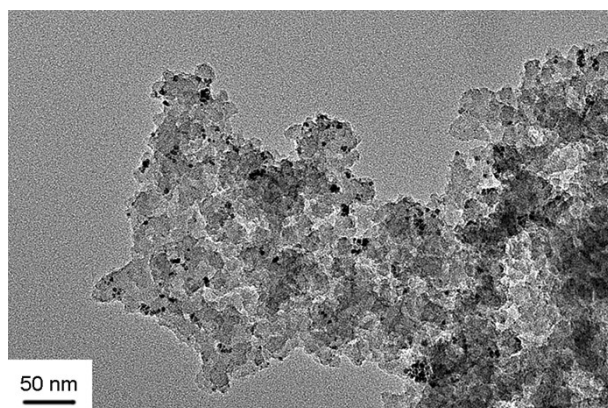


Fig. S3 TEM image for MOP-Ru.

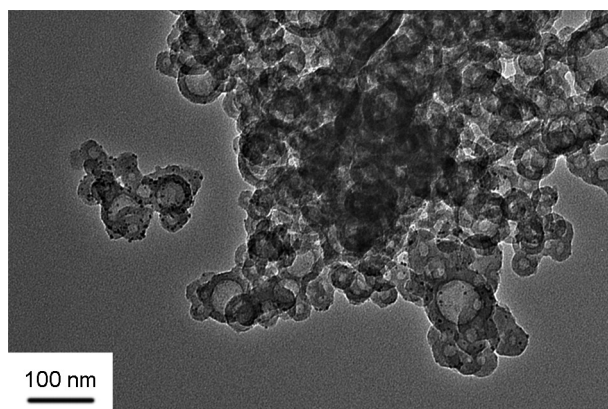


Fig. S4 TEM image for H-MONs-Ru after six runs.

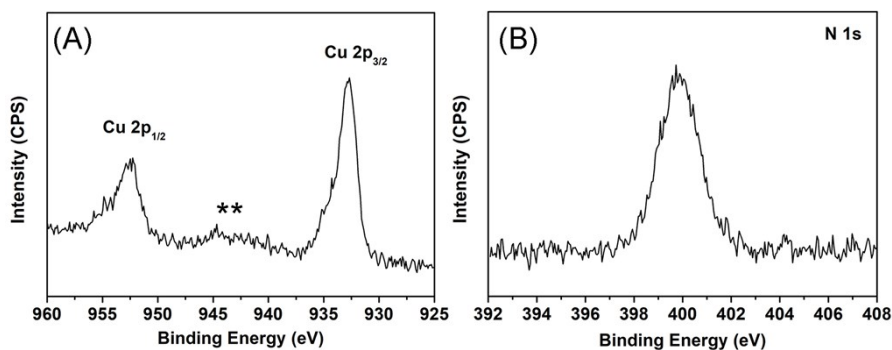


Fig. S5 XPS (A) Cu 2p and (B) N 1s spectra of H-MONs-Cu. The Cu 2p XPS spectrum of H-MONs-Cu showed shakeup satellite peaks at 942-945 eV, which confirmed the presence of Cu(II) species. Two intense peaks at 932.6 eV and 952.4 eV were attributable to $\text{Cu}^{2+} 2p_{3/2}$ and $\text{Cu}^{2+} 2p_{1/2}$, respectively.¹ In addition, the binding energy of the N 1s peak was shifted toward higher binding energy (at around 400 eV), which can be attributed to a decrease in the electron density of the N atom by the coordination between the N atoms in H-MONs-Cu and immobilized Cu atoms.²



Fig. S6 Iodine vapor adsorption apparatus set-up.

Table S1. Comparison of catalytic efficiency for the CTH of nitrobenzene with various catalysts.

We compared H-MONs-Ru with various catalysts reported in the literature for the CTH of nitrobenzene. Though in different reaction conditions, such as hydrogen-donor reagents, catalyst amount, and reaction conditions, as-prepared H-MONs-Ru catalyst shows good catalytic activity.

Catalyst	H ₂ source	Amount of cat.	Time/yield	Ref.
Ru-BBA-1 ^{a)}	NaBH ₄	0.17 mol%	30 min/100%	3
Rh NPs/SBA-NH ₂ ^{b)}	N ₂ H ₄	10 mg	5 min/100%	4
Ru/CMK-3 ^{c)}	N ₂ H ₄	20.9 mg	60 min/58%	5
Ni-PVAm/SBA-15 ^{d)}	NaBH ₄	0.12 g	20 min/98%	6
AuNPs-sPSB ^{e)}	NaBH ₄	0.1 mol%	60 min/99%	7
Cu(10%)-Meso-PANI ^{f)}	NaBH ₄	50 mg.	3 h/100%	8
Re/OMC ^{g)}	NaBH ₄	0.02 mg	25 s/98%	9
Au/rutile ^{h)}	FA	1 mol%	0.67 h/99%	10
O-MoS ₂ ⁱ⁾	N ₂ H ₄	20 mg	60 min/99%	11
H-MONs-Ru	NaBH ₄	0.025 mol%	45 min/100%	This work

^{a)} Reaction conditions: 0.5 mmol of nitrobenzene, 2.5 mmol of NaBH₄, THF/H₂O (1:3, v/v), RT.

^{b)} Reaction conditions: 1.0 mmol of nitrobenzene, 2.0 mmol of N₂H₄, H₂O (3.0 mL), RT.

^{c)} Reaction conditions: 8.1 mmol of nitrobenzene, 1.0 mL of N₂H₄, H₂O (1.0 mL), 30 °C.

^{d)} Reaction conditions: 2.0 mmol of nitrobenzene, 8.0 mmol of NaBH₄, H₂O (3.0 mL), RT.

^{e)} Reaction conditions: 2.54 mmol of nitrobenzene, 15.24 mmol of NaBH₄, methanol (6.0 mL), 25 °C.

^{f)} Reaction conditions: 1.0 mmol of nitrobenzene, 5.0 mmol of NaBH₄, ethanol/H₂O (1:1, 10 mL), 40 °C.

^{g)} Reaction conditions: nitrobenzene (1.8 mL, 0.1mM), NaBH₄ (0.5 mL, 0.1M), H₂O (3.0 mL), RT.

^{h)} Reaction conditions: 1.0 mmol of nitrobenzene, 3.0 mmol of FA, toluene (5.0 mL), N₂ (1 bar), 60 °C.

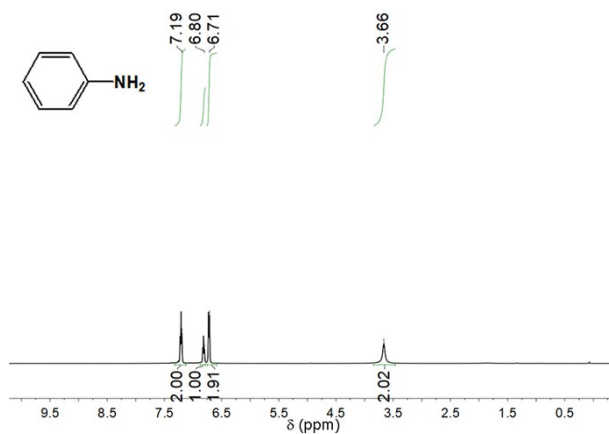
ⁱ⁾ Reaction conditions: 0.5 mmol of nitrobenzene, 1.5 mmol of N₂H₄, ethanol (2.0 mL), 50 °C.

Table S2. Comparison of I₂ Adsorption data in various adsorbents.

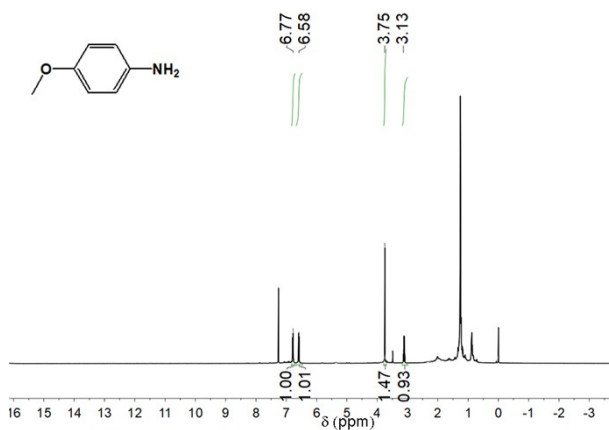
Adsorbent	Temperature (°C)	Pressure	Iodine Capacity (mg I₂/g)	Ref.
MFM-300(Sc)	80	N ₂ atmosphere	1540	12
SCMPs	75	1 bar	2220	13
NTP	75	1 bar	1800	14
PAF-1	25	40 bar	1860	15
JUC-Z2	25	40 bar	1400	15
ZIF-8	75	1 bar	1200	16
Activated carbon	75	1 bar	300	14
Ag@Zeolite Mordenites	95	1 bar	275	17
Ag@Mon-POF	70	1 bar	250	18
CC3	20	1 bar	364	19
H-MONs-TA	75	1 bar	1300	This work
H-MONs-CZ	75	1 bar	1100	This work

Analytical data for compounds of the CTH of nitroarenes.

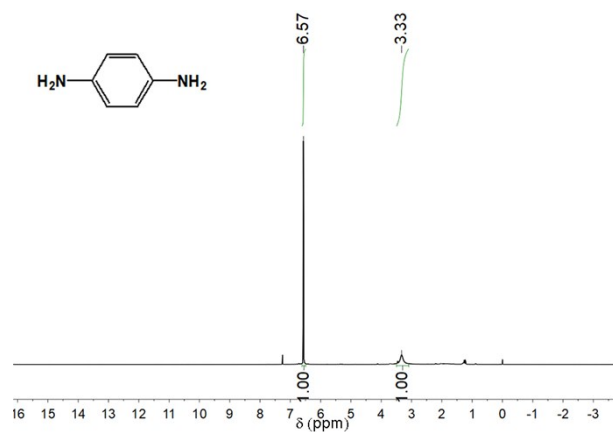
Aniline. ^1H NMR (500 MHz, CDCl_3): δ 7.19 (t, $J = 8.0$ Hz, 2H); 6.80 (t, $J = 7.5$ Hz, 1H); 6.71 (d, $J = 7.5$ Hz, 2H); 3.66 (s, 2H).



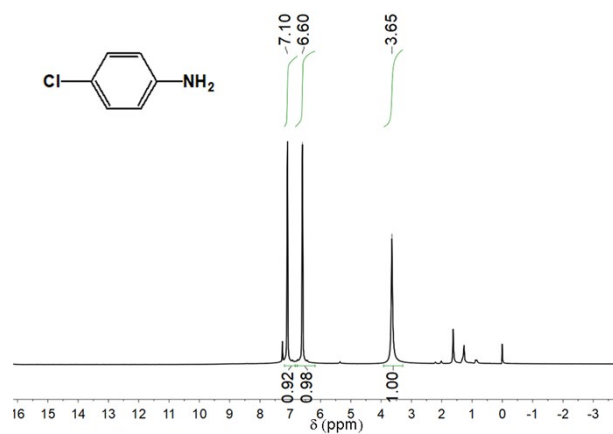
4-Aminoanisole. ^1H NMR (500 MHz, CDCl_3): δ 6.77 (d, $J = 8.0$ Hz, 2H); 6.58 (d, $J = 8.0$ Hz, 2H); 3.75 (s, 3H); 3.13 (bs, 2H).



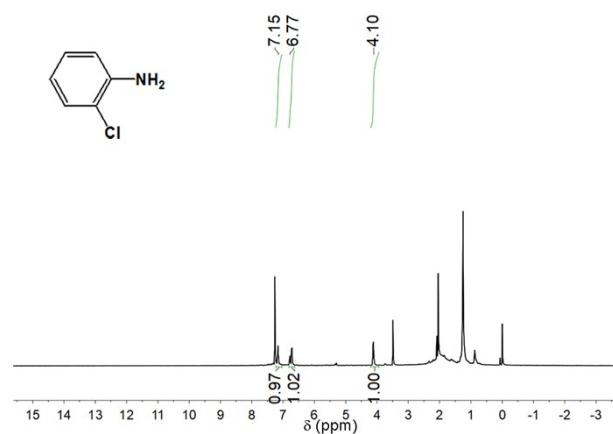
1, 4-Phenylenediamine. ^1H NMR (500 MHz, CDCl_3): δ 6.57 (s, 4H); 3.33 (s, 4H).



4-Chloroaniline. ^1H NMR (500 MHz, CDCl_3): δ 7.10 (d, $J = 6.5$ Hz, 2H); 6.60 (d, $J = 6.5$ Hz, 2H); 3.65 (s, 2H).



2-Chloroaniline. ^1H NMR (500 MHz, CDCl_3): δ 7.15 (m, 2H); 6.77 (d, $J = 7.5$ Hz, 2H); 4.10 (bs, 2H).



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