

Supporting Information:

**Palladium (Pd) catalyst imbedded in polymers of intrinsic
microporosity for Suzuki-Miyaura coupling reaction**

Junwen Xu,^{†,‡} Junjie Ou,^{*,†} Lianfang Chen,^{†,‡} Haiyang Zhang,[†] Shujuan Ma,[†] Mingliang
Ye^{*,†}

[†]Key Laboratory of Separation Science for Analytical Chemistry, Dalian Institute of
Chemical Physics, Chinese Academy of Sciences, Dalian 116023, China

[‡]University of Chinese Academy of Sciences, Beijing 100049, China

Corresponding Author

*E-mail: junjieou@dicp.ac.cn (J. Ou)

*E-mail: mingliang@dicp.ac.cn (M. Ye)

Table of Contents

1. Chemicals and Instruments

2. Supplementary Figures and Tables

Figure S1. ^1H NMR spectra of monomer NBTFD.

Figure S2. ^1H NMR spectra of monomer PBTFD.

Figure S3. ^{13}C CP/MAS NMR spectra of PIM-N (c) and catalysts Pd/PIM-N (d).

Figure S4. SEM image of PIM-N.

Figure S5. SEM image of Pd/PIM-N.

Figure S6. SEM image of Pd/PIM-N after one cycle.

Figure S7. TEM image of PIM-N.

Figure S8. TEM image of Pd/PIM-N.

Figure S9. TEM image of Pd/PIM-N after one cycle.

Figure S10. Nitrogen adsorption isotherms PIM-N (circular) and Pd/PIM-N (rhombus).

Figure S11. The XPS image of Pd/PIM-P and Pd/PIM-N.

1. Chemicals and Instruments

p-Phenylenediamine, 1,5-naphthalenediamine, tetrafluorophthalic anhydride 5,5',6,6'-tetrahydroxy-3,3,3',3'-tetramethyl spirobisindane were purchased from TCI chemical (Shanghai, China). Palladium (II) acetate and anhydrous DMF was purchased from Sigma-Aldrich (St Louis, Mo, USA). Dichloromethane, glacial acetic acid, ethanol, acetonitrile and potassium carbonate anhydrous (K_2CO_3) were obtained from Kemiou Chemical Regent Co., Ltd. (Tianjin, China). The water was doubly distilled and purified by Milli-Q system (Millipore Inc., Milford, MA, USA). The C18 AQ beads (5 μm , 120 \AA) were obtained from MichromBioResources (Auburn, CA).

FT-IR spectra were obtained on a spectrometer with KBr pellets containing polymer sample (Thermo Scientific, USA). Elemental analysis was carried out on EMIA-8100H and EMG-930 (HORIBA, Japan). Solid-state ^{13}C cross-polarization magnetic angle spinning (CP-MAS) NMR spectra and ^1H NMR spectra were recorded on a Bruker Advance III 500 NMR spectrometer (Germany). X-ray photoelectron spectroscopy (XPS) data were acquired using an ESCALAB 250Xi XPS spectrometer with an Al $K\alpha$ X-ray source (Thermo Scientific, USA). The morphology study of porous organic polymer was carried out on a scanning electron microscopy (SEM, Gemini SEM 300, Zeiss, Germany). Transmission electron microscopy (TEM) was conducted on a JEOL 2000 EX electronic microscope (Japan) with an accelerating voltage of 120 keV. Nitrogen adsorption-desorption measurements were performed on a Micromeritics ASAP 2460 (USA) surface area analyzer and pore size analyzer. The samples were outgassed under vacuum for 7 h at 120 $^\circ\text{C}$ before measurements. The surface area was calculated via the Brunauer-Emmett-Teller (BET) method. Chromatography analysis was performed on UPLC (Water, USA).

2. Supplementary figures and tables

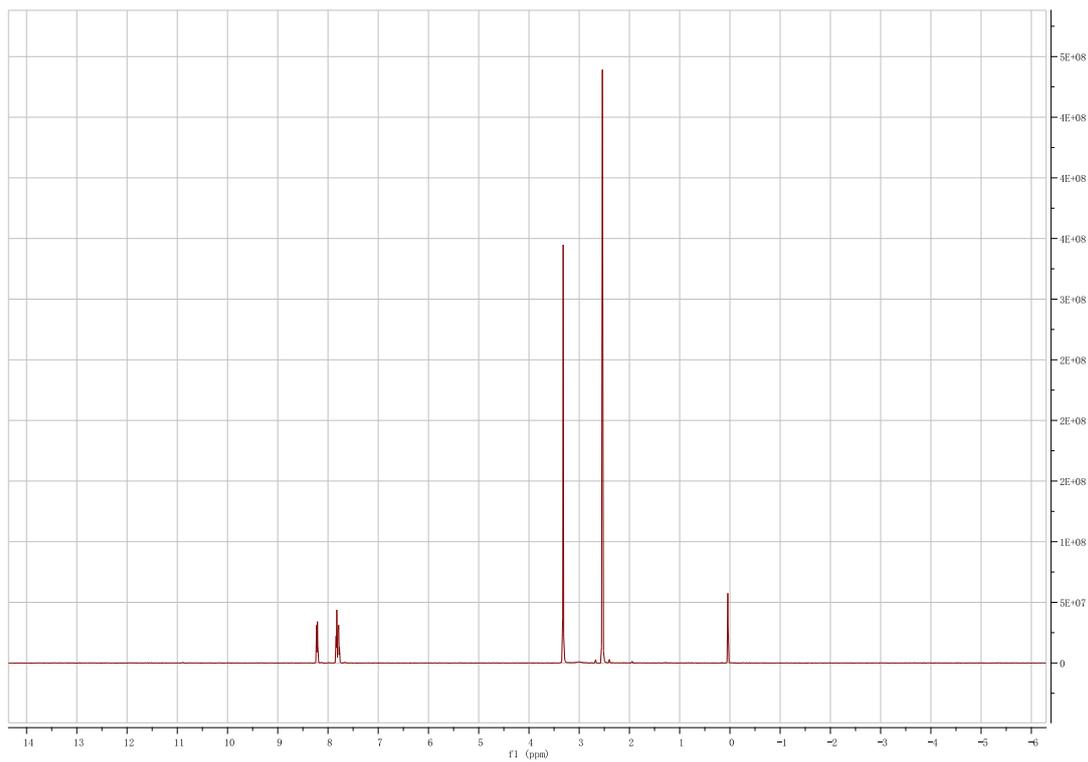


Figure S1. ^1H NMR spectra of monomer NBTFD.

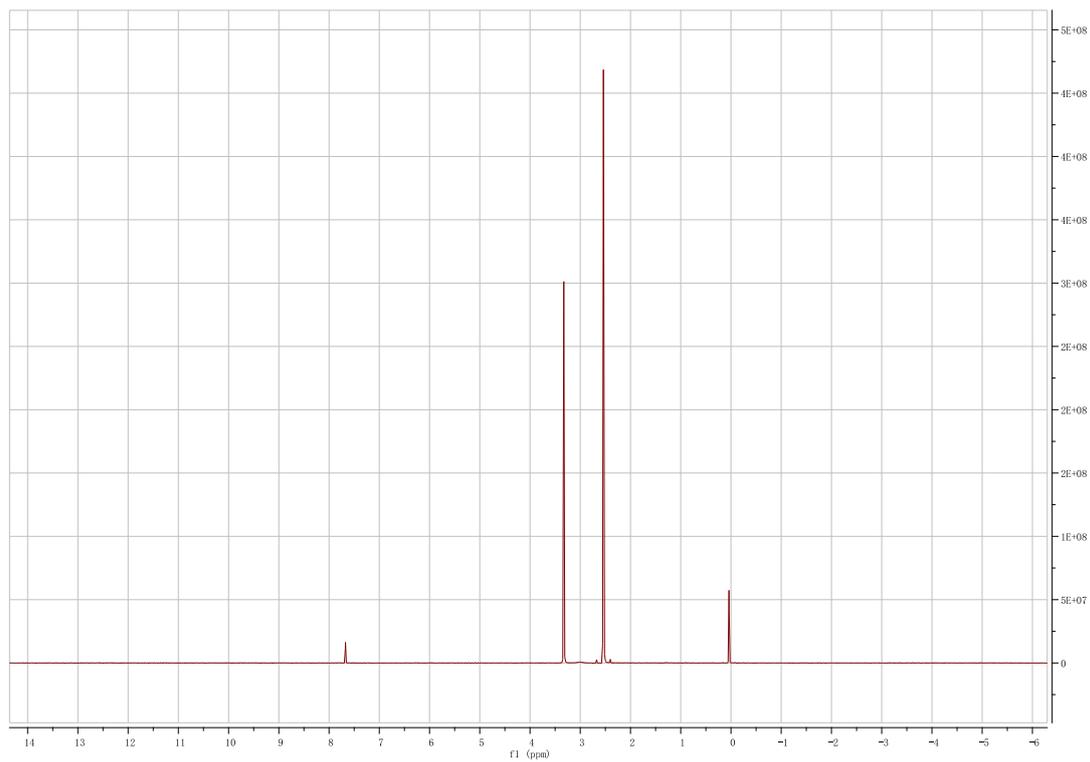


Figure S2. ^1H NMR spectra of monomer PBTFD.

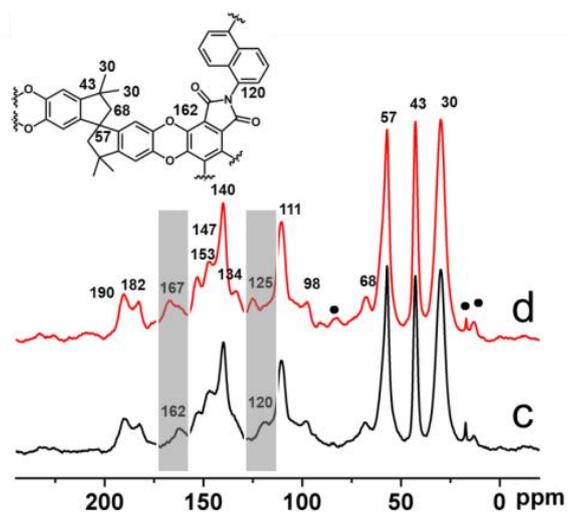


Figure S3. ¹³C CP/MAS NMR spectra of PIM-N (c) and catalysts Pd/PIM-N (d).

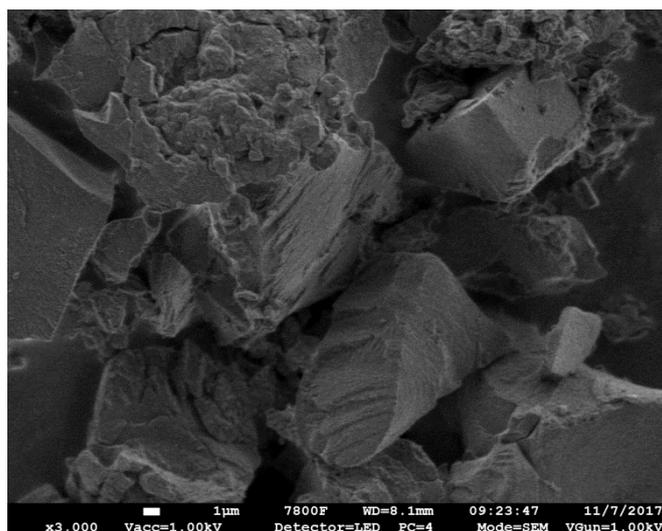


Figure S4. SEM image of PIM-N.

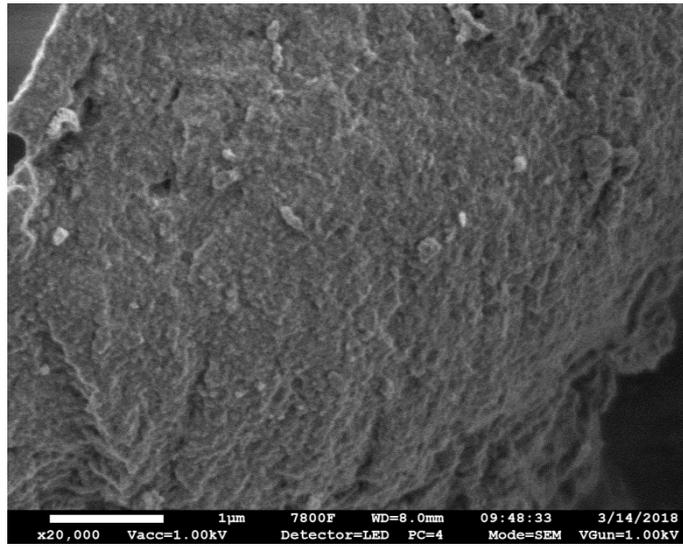


Figure S5. SEM image of Pd/PIM-N.

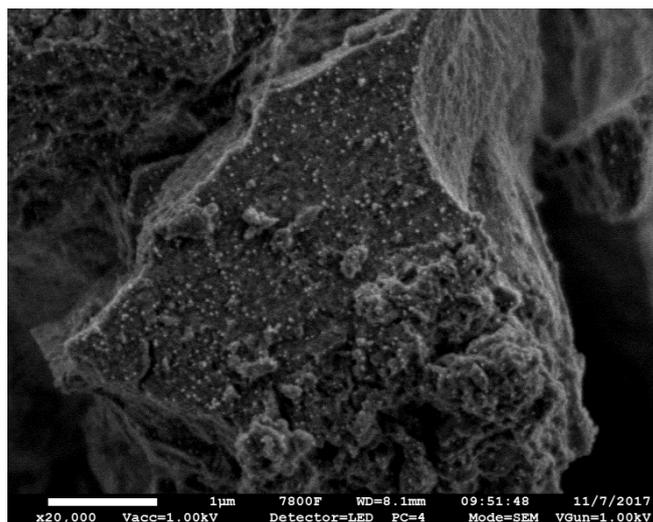


Figure S6. SEM image of Pd/PIM-N after one cycle.

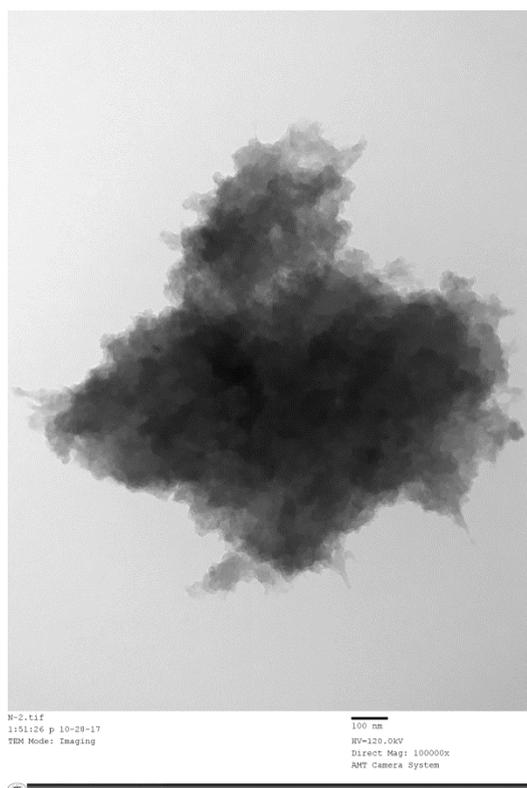


Figure S7. TEM image of PIM-N.

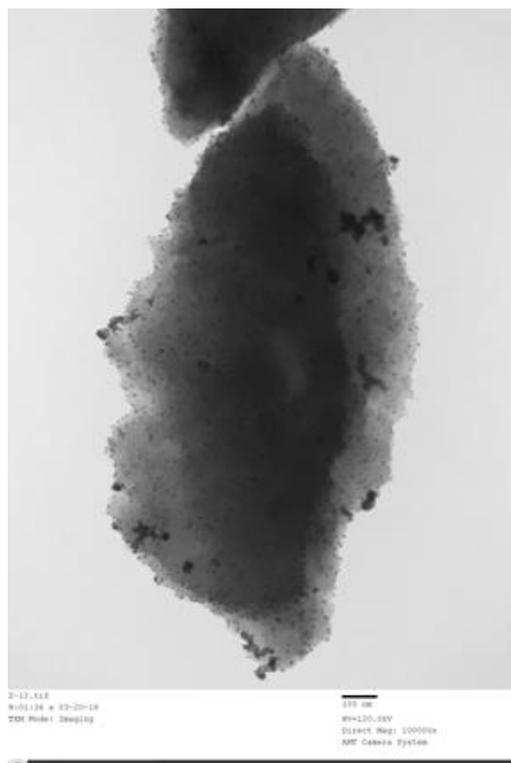


Figure S8. TEM image of Pd/PIM-N.

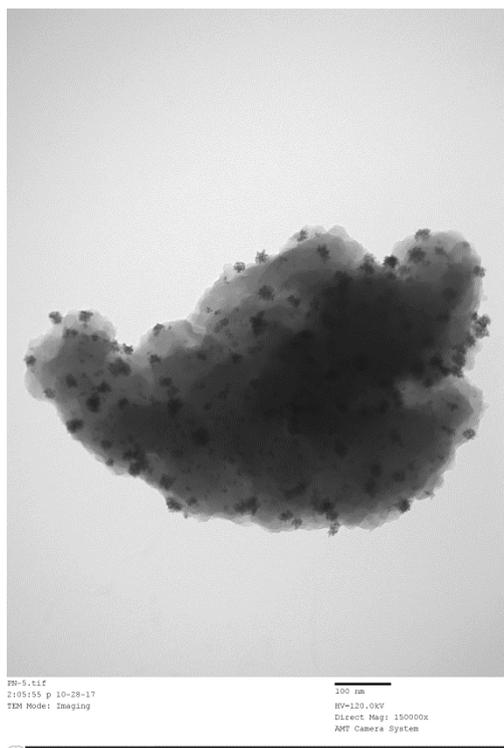


Figure S9. TEM image of Pd/PIM-N after one cycle.

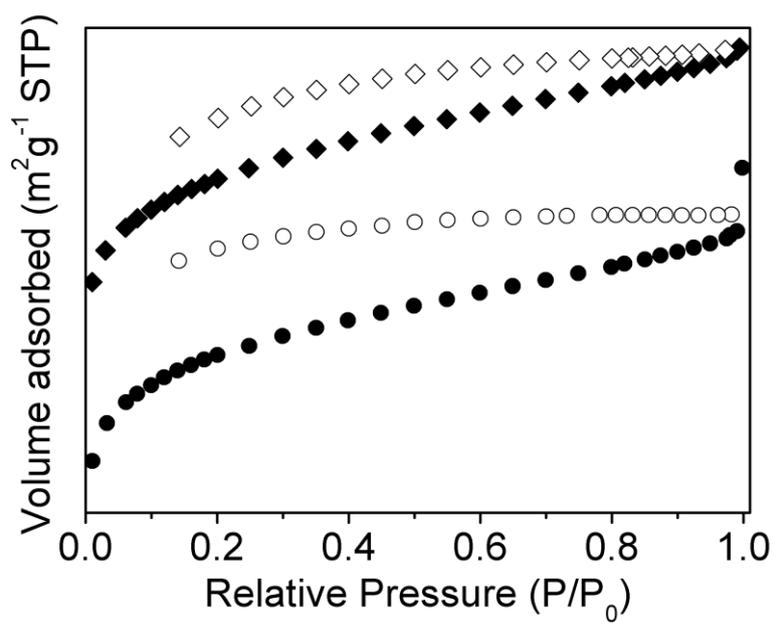


Figure S10. Nitrogen adsorption isotherms PIM-N (circular) and Pd/PIM-N (rhombus).

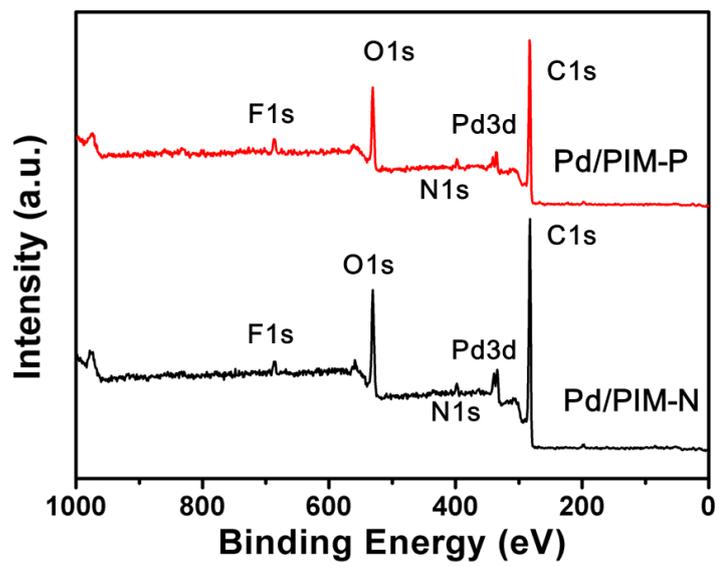


Figure S11. The XPS image of Pd/PIM-P and Pd/PIM-N.