Supporting Information



Figure S1 Typical SEM image of the bifunctional PMOs synthesized with P123 as SDA and BTET/BTEB as the organosilica precursors with CO_2 pressure of 5.90 MPa.



Figure S2 ²⁹Si MAS NMR spectra of the bifunctional PMOs synthesized with P123 as SDA and BTET/BTEB as the organosilica precursors with different CO₂ pressures.



Figure S3. Simulation of the ²⁹Si MAS NMR spectra of the bifunctional PMO materials according to Figure S2.

Table S1. ²⁹Si MAS NMR data of the bifunctional PMO materials synthesized with P123 as SDA and BTET/BTEB as organosilica precursors with different CO₂ pressures

Pressure (MPa)	T ⁿ (thiophene)[%]			T ⁿ (benzene)[%]			Ratio $\sum_{T^{n} \text{this share}}$
	T^1	T^2	T^3	T^1	T^2	T^3	$T^{n}_{benzene}$
5.90	10	27	13	12	25	13	1:1
4.90	10	27	13	12	25	13	1:1



Figure S4. ¹³C MAS NMR spectra of the bifunctional PMOs synthesized with P123 as SDA and BTET/BTEB as organosilica precursors with different CO₂ pressures.



Figure S5. (A) Absorption spectra of BPB (4×10^{-6} M) in citric acid buffers of pH 3.0 (a), 3.2 (b), 3.4 (c), 3.6 (d), and 3.8 (e) over the wavelength range of 350–660 nm. (B) Dependence of the ratio I_{430}/I_{590} on the hydrogen ion concentration of citric acid buffers.