

Electronic Supplementary Information (ESI)

Light-Emitting Conjugated Microporous Polymers Based on Excited-State Intramolecular Proton Transfer Strategy and Selective Switch-off Sensing of Anions

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Section 1. Instrumental characterization

The infrared spectra were recorded from 500 to 3500 cm^{-1} on an Avatar FT-IR 360 spectrometer by using KBr pellets. Elemental analyses were measured by an Elementar model vario EL cube analyzer. Field emission scanning electron microscopy was recorded on a SU8020 model HITACHI microscope. Powder X-ray diffraction data were performed on a PANalytical BV Empyrean diffractometer by depositing powder on glass substrate, from $2\theta = 4.0^\circ$ to 35° with 0.02° . Thermogravimetric analysis (TGA) was performed on a TA Q500 thermogravimeter with the heating at a rate of $10^\circ\text{C min}^{-1}$ from 35°C to 800°C under nitrogen. Nitrogen sorption isotherms were measured at 77 K with a JW-BK 132F analyzer. The absolute fluorescence quantum yields were measured on Edinburgh FLS920 by using an integrating sphere. Photoluminescence spectra were recorded on a Cary Eclipse Fluorescence Spectrophotometer. Frontier molecular orbital (FMO) plots of model-1 and model-2 at the level of B3LYP/6-31G(d,p). CMPs was dispersed in THF to yield a nearly transparent dispersion which was immediately subjected to fluorescence spectroscopy. After the addition anions, time-dependent fluorescence spectra were recorded.

Section 2. ^{13}C NMR spectra

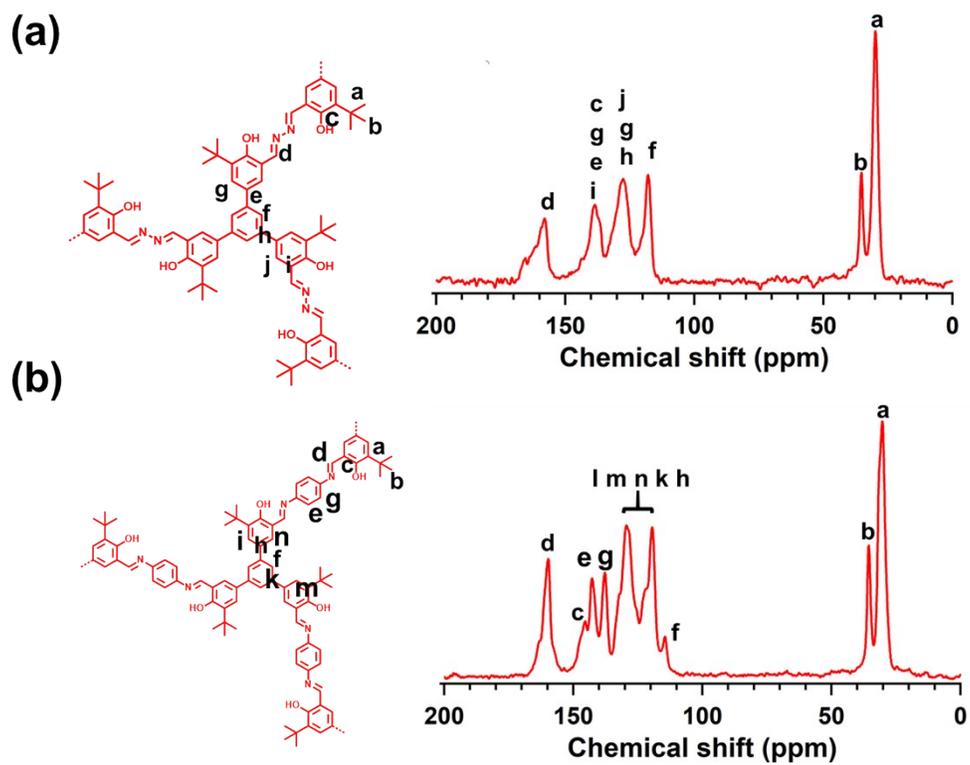


Fig. S1. ^{13}C NMR spectra (a) CMP-A and (b) CMP-B.

Section 3. Solid-state electronic absorption spectra

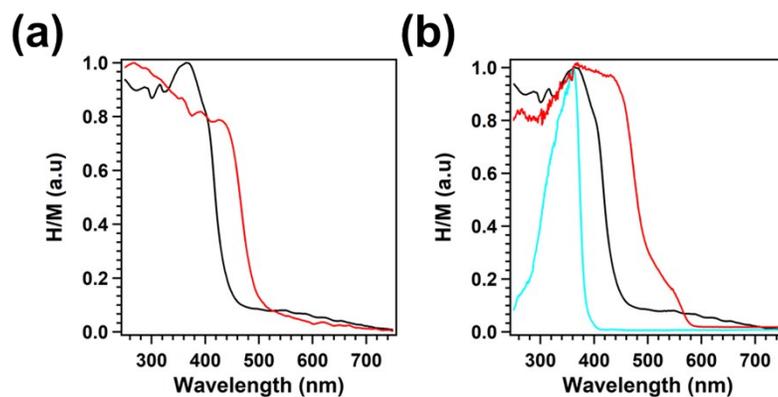


Fig. S2. Solid-state electronic absorption spectra of (a) CMP-A (red) and monomer C (black); (b) CMP-B (red), monomer C (black) and monomer B (sky-blue).

Section 4. elemental analysis

Table 1

| Samples | Element | C% | H% | N% |
|---------|------------|--------|------|------|
| CMP-A | Observed | 77.15 | 7.61 | 6.91 |
| | Calculated | 75.72 | 6.99 | 6.68 |
| CMP-B | Observed | 80.20 | 7.01 | 5.75 |
| | Calculated | 80.64. | 6.67 | 5.88 |

Section 5. TGA curves

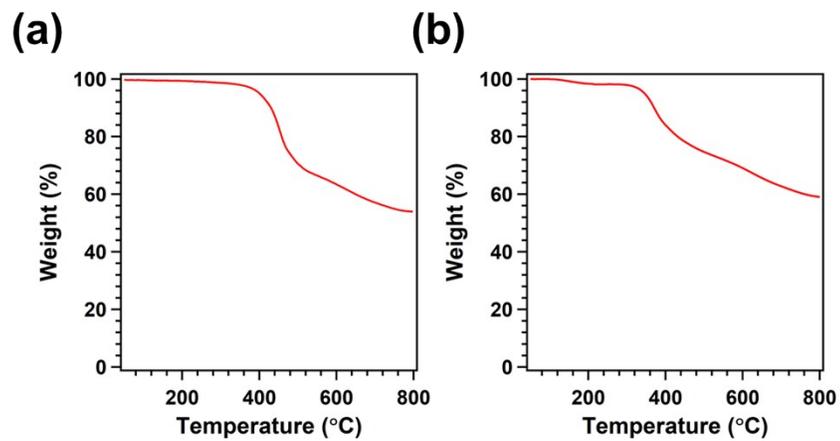


Fig. S3. TGA curves of (a) CMP-A and (a) CMP-B.

Section 6. PXRD patterns

(a)

(b)

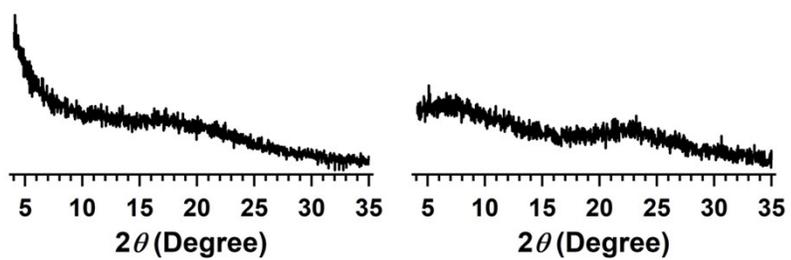


Fig. S4. PXRD patterns of (a) CMP-A and (a) CMP-B.

Section 7. Isostatic heat of carbon dioxide adsorption

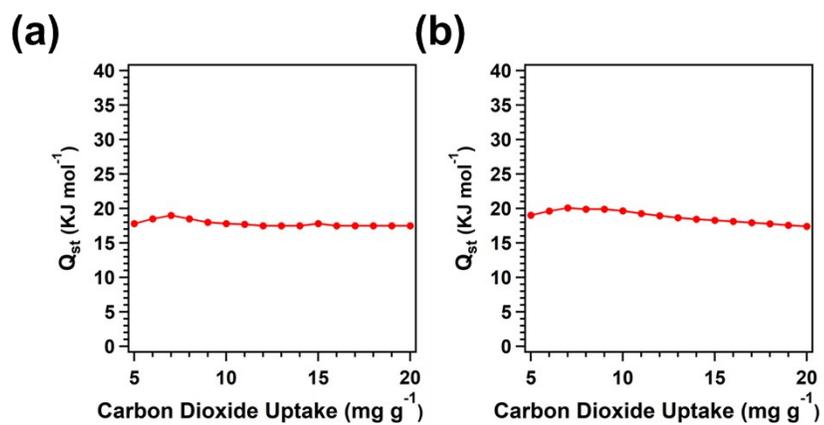


Fig. S5. The isosteric heat of adsorption of CO_2 for (a) CMP-A and (b) CMP-B.

Section 8. Fluorescence spectra

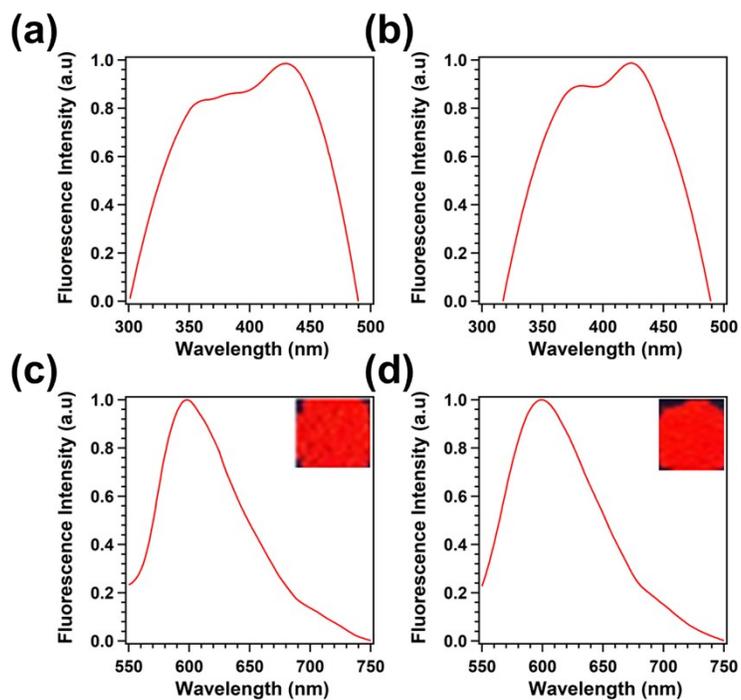


Fig. S6. Normalized fluorescence excitation spectra of (a) CMP-A and (b) CMP-B. Normalized fluorescence emission spectra of (c) CMP-A and (d) CMP-B.

Section 9. Dynamic light scattering measurement

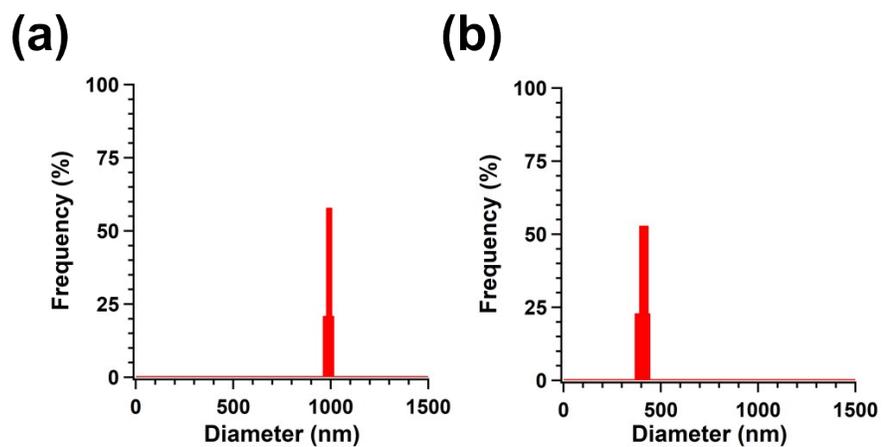


Fig. S7. Dynamic light scattering profile of (a) CMP-A and (b) CMP-B.

Section 10. ^1H NMR spectra

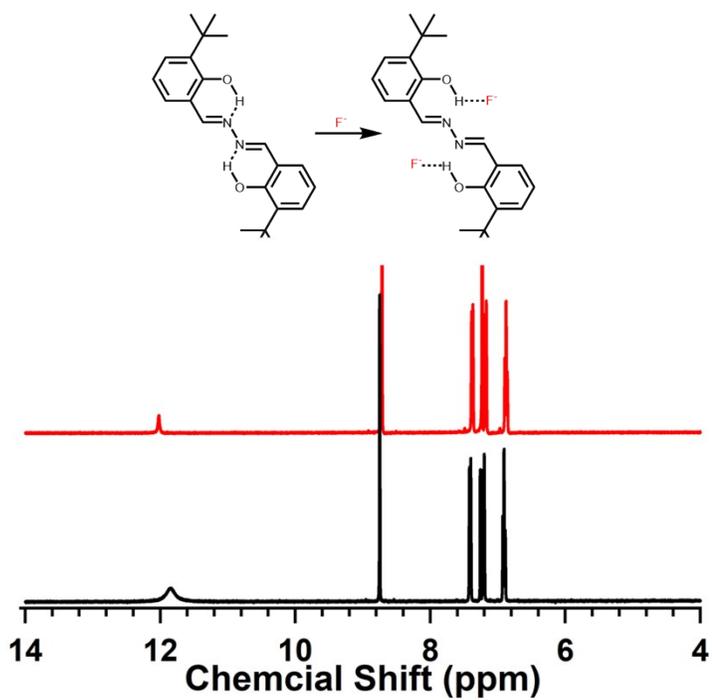


Fig. S8. NMR spectra of module before (black curve) and after fluoride anions (red curve).

Section 11. Plot of PL quenching efficiency

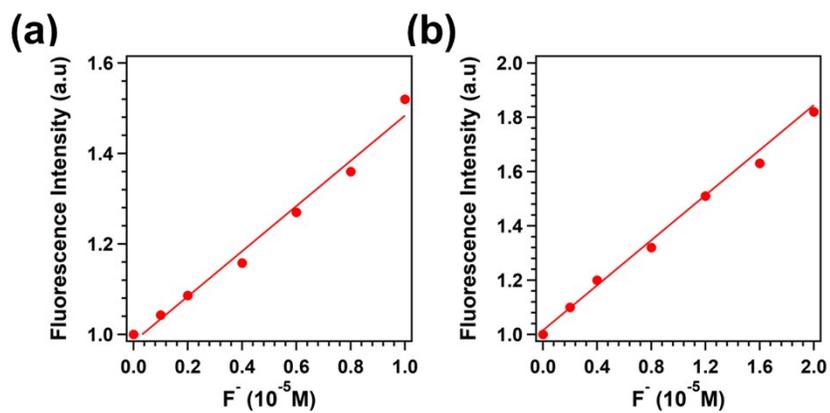


Fig. S9. Plot of PL quenching efficiency (I_0/I) as a function of fluoride anions concentration of (a) CMP-A and (b) CMP-B (fluoride anions: 0 to $10^{-4} M$).

Section 12. Fluorescence spectra

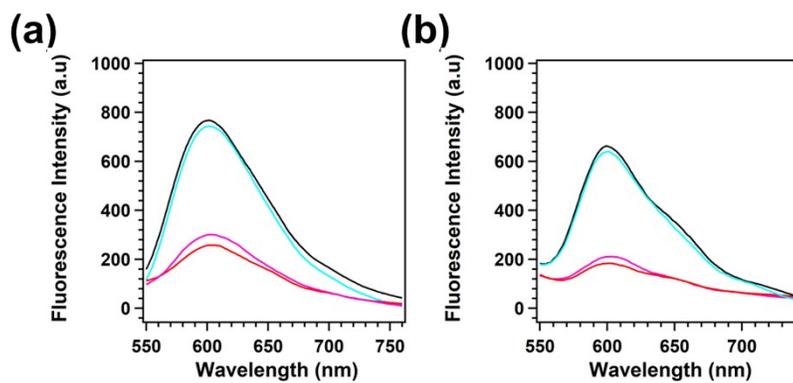


Fig. S10. Fluorescence spectra of (a) CMP-A and (b) CMP-B in THF (Original CMPs: black curves; CMPs@F⁻@H⁺: sky-blue curves; CMPs@F⁻@H⁺@F⁻: pink curves; CMPs@F⁻: red curves).