Supporting Information

Synthesis of a 3D Photochromic Coordination Polymer with Interpenetrating Arrangement: Crystal Engineering for Electron Transfer between Donor and Acceptor Units
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Experimental Section

All the reagents were purchased from commercial channels and used without further purification; N-(3-carboxyphenyl)-4,4′-bipyridinium chloride was synthesized as reported. UV-Visible spectral measurements were carried out using a HITACHI U-3010 spectrometer. The ESR spectra were recorded at room temperature with a Bruker EMX-10/12 Electron Spin Resonance Spectrometer. IR spectra were characterized by a Bruker Tensor 27 FTIR spectrometer in the range of 4000-400 cm⁻¹ using a KBr disk.

Synthesis of 1: $Zn(NO_3)_2 \cdot 6H_2O$ (29.7mg, 0.1mmol) was added to a mixture of p-H₂BDC (16.6mg, 0.1mmol), N-(3-carboxyphenyl)-4,4'-bipyridinium chloride (31mg, 0.1mmol) in H₂O (2ml), C₂H₅OH (4ml) and DMF (4ml). The mixture was sealed in a 25ml Teflon-lined steel bomb and heated at 85°C for 48h. Yellow block-like crystals were collected by filtration, washed by water and ethanol, and dried at room temperature (0.018mmol, 23mg, 54% yield based on $Zn(NO_3)_2 \cdot 6H_2O$).

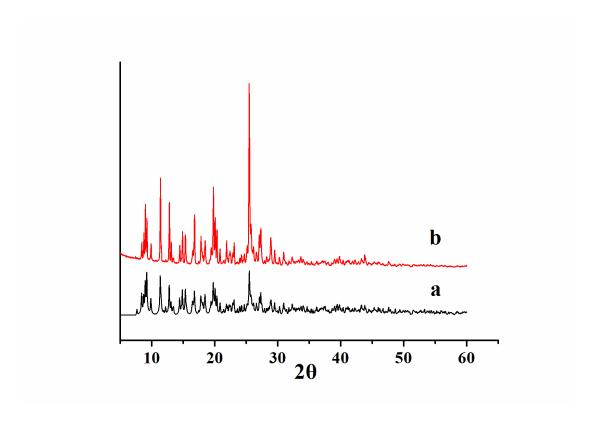


Figure S1 PXRD patterns for compound **1**: (a) simulated; (b) of a sample at room temperature.

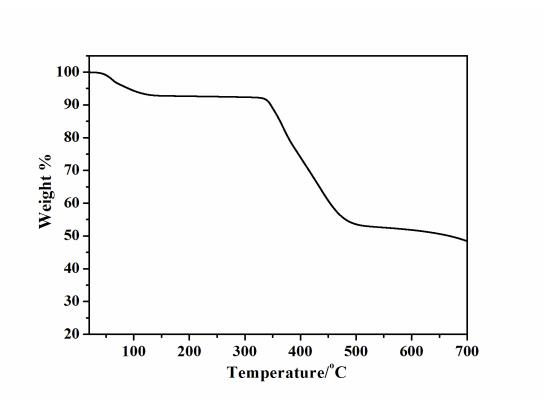


Figure S2. Thermal gravimetric curve of 1

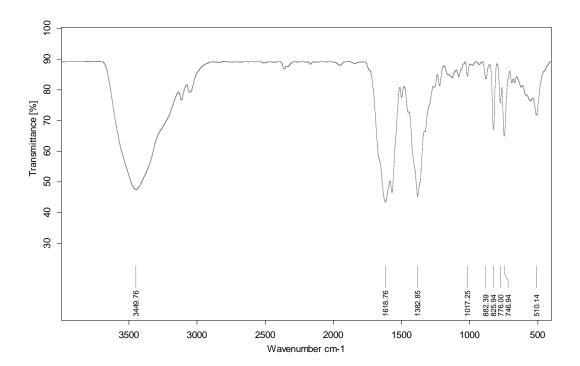


Figure S3. IR spectrum of 1