## **Electronic Supplementary Information**

## Three-dimensional coordination polymers constructed from $C_{2}$ -

## symmetric linkers of pyridyl dicarboxylate ligands

## Synthesis of H<sub>2</sub>L<sub>2</sub>

4-(ethoxycarbonyl)phenylboronic acid (4.46 g, 23 mmol), 3,5-Dibromopyridine (2.36g, 10 mmol), and K<sub>2</sub>CO<sub>3</sub> (2.10 g, 10.0 mmol) were added to 1,4-dioxane (30 mL). After stirring, Pd(PPh<sub>3</sub>)<sub>4</sub> (0.05 g, 0.043 mmol) was added, then the mixture was heated to 80 °C for 3 days under N<sub>2</sub>. The resultant was evaporated to dryness and taken up in CH<sub>2</sub>Cl<sub>2</sub> which later had been dried over MgSO<sub>4</sub>. This CH<sub>2</sub>Cl<sub>2</sub> solution was evaporated to dryness and the residue was washed briefly with ethanol (20 mL). The crude product was hydrolyzed by refluxing in 2 M aqueous NaOH followed by acidification with 37% HCl to afford H<sub>2</sub>L<sub>2</sub>. Yield: 2.5 g, 78.3%. <sup>1</sup>H NMR (DMSO-*d*<sub>6</sub>, 500 MHz), H<sub>2</sub>L<sub>2</sub>: 9.03 (d, 2H), 8.51 (s, 1H) 8.09 (s, 2H), 8.08 (s, 2H) 8.04 (s, 2H), 8.02 (s, 2H). Anal. Calcd (Found) for H<sub>2</sub>L<sub>3</sub>, C<sub>19</sub>H13O<sub>4</sub>N: C, 71.47 (71.13); N, 4.39 (4.28); H, 4.10 (3.89) %.

Complex	1	2
Formula	C <sub>16</sub> H <sub>14</sub> N <sub>2</sub> O <sub>5</sub> Zn	C <sub>22</sub> H <sub>18</sub> N <sub>2</sub> O <sub>5</sub> Zn
Mr	379.66	455.75
Crystal system	monoclinic	monoclinic
Space group	P21/c	P2 <sub>1</sub> /c
a, Å	11.877(2)	15.3645(11)
b, Å	13.855(3)	15.1275(11)
c, Å	13.921(3)	8.3110(6)
α, deg	90	90
β, deg	114.403(2)	91.8500(10)
γ, deg	90	90
V (Å <sup>3</sup> )	2086.3(7)	1930.7(2)
Ζ	4	4
Dealed (g cm <sup>-3</sup> )	1.209	1.568
Absorption coefficient (mm <sup>-1</sup> )	1.199	1.311
F(000)	766	936
Reflections collected / unique	10661 / 3805	10927 / 3524
Data / restraints / parameters	3805 / 0 / 219	3524 / 6 / 274
Rint	0.0396	0.1037
Final R indices [I>2sigma(I)]	0.0311	0.0519
Final wR(F2) values [I>2sigma(I)]	0.0843	0.1652
Final R1 values (all data)	0.0371	0.0578
Final wR(F2) values (all data)	0.0861	0.1719
Goodness of fit on F2	1.023	1.029

Table S1 Crystallographic data and structural refinement summary for complexes 1 and 2.

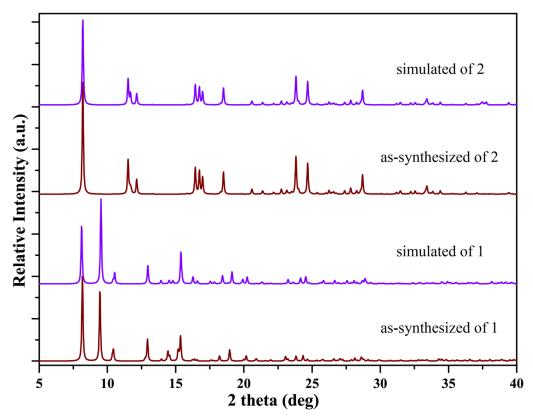


Fig. S1. The as-synthesized and simulated PXRD of 1 and 2.

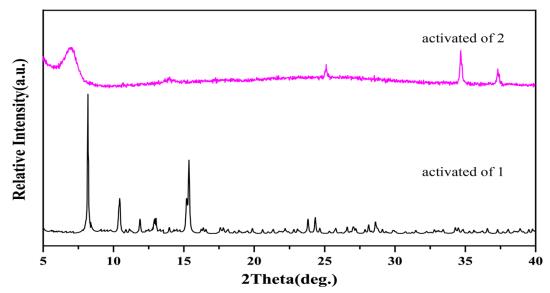


Fig. S2. The PXRD of activated sample of 1 and 2.

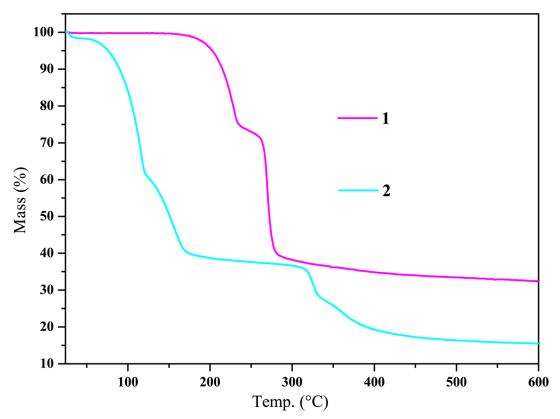


Fig. S3. The TGA curves of **1** and **2**.