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Syntheses, structural diversities and characterization of a series of coordination polymers with two isomeric oxadiazol-pyridine ligands†

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In this work two positional-isomeric oxadiazol-pyridine ligands 3-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L_1) and 4-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L_2) have been designed and synthesized. A series of novel coordination polymers, namely $[\text{Cd}_2(\mu_2-L_1)_2(\mu_2-\text{NCS})_4]_n$ (**1**), $[\{\text{Cd}(L_1)(\mu_2-\text{dca})_2(\text{H}_2\text{O})\} \cdot \text{H}_2\text{O}]_n$ (**2**), $[\{\text{Cu}(\mu_2-L_1)_2(\text{NCS})_2\} \cdot 0.5\text{H}_2\text{O}]_n$ (**3**), $[\{\text{Ag}_2(\mu_2-L_1)(\mu_3-L_1)_2\} \cdot 2\text{PF}_6]_n$ (**4**), $[\{\text{Ag}_3(\mu_2-L_1)_4(\mu_2-\text{CF}_3\text{SO}_3)(\text{CF}_3\text{SO}_3) \cdot \text{CF}_3\text{SO}_3\}]_n$ (**5**), $[\{\text{Cd}(L_2)_2(\mu_2-\text{NCS})_2\}]_n$ (**6**), $[\text{Ag}(\mu_2-L_2)(\mu_2-\text{CF}_3\text{SO}_3)]_n$ (**7**) and $[\{\text{Ag}(\mu_2-L_2)\} \cdot \text{BF}_4]_n$ (**8**) have been isolated. Both **1** and **2** are 2D Cd^{II} coordination polymers containing infinite $\{\text{Cd}-\text{NCS}-\text{Cd}\}$ chains (for **1**) or infinite $\{\text{Cd}-\text{dca}-\text{Cd}\}$ layers (for **2**), respectively. **3** is a 2D Cu^{II} coordination polymer, in which central metal ions are bridged via a bidentate bridging L_1 ligand. While when different Ag^{I} salts were introduced into the reaction system, 1D Ag^{I} coordination polymers **4** and **5** with diverse coordination modes can be isolated. Furthermore, when the isomeric oxadiazol-pyridine L_2 is used to replace L_1 in the reaction system, **6–8** can be isolated. **6** is a 2D Cd^{II} coordination polymer containing $\{\text{Cd}-\text{NCS}-\text{Cd}\}$ layers. **7** is a 2D neutral Ag^{I} coordination polymer while **8** is a 2D cationic Ag^{I} coordination polymer. Variable temperature magnetic susceptibility measurements (2–300 K) reveal anti-ferromagnetic interactions between central copper(II) ions for **3**. Solid-state luminescent properties of **1**, **2** and **4–8** have been investigated indicating strong fluorescent emissions. Additionally, luminescent measurements illustrate that complex **8** exhibits highly sensitive luminescence sensing for $\text{Cr}_2\text{O}_7^{2-}$ ions in aqueous solutions with high quenching efficiency $K_{\text{sv}} = 2.08 \times 10^4 \text{ L mol}^{-1}$ and low detection limit (0.19 μM ($S/N = 3$)). **8** also represents the first report of a coordination polymer based on oxadiazol-pyridine derivatives with a luminescence response to $\text{Cr}_2\text{O}_7^{2-}$ anion pollutants in aqueous solutions.

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Introduction

In the past several decades, metal organic coordination polymers have attracted much interest due to their promising applications for gas storage/separation, catalysis, luminescence, drug delivery, sensors and magnetic materials.^{1–9} Among these coordination materials, luminescent metal organic coordination polymers have received great interest because they can

be applied in the selective and sensitive detection of environmental pollutants, which can further protect human health and the environment from harm by these pollutants.¹⁰ Their tunable channel size, good structural stability, and large surface area of luminescent coordination polymers make these coordination materials more competitive over other luminescent materials because more analytes can effectively interact with the channel surface areas of the coordination polymers, which can ultimately improve the rate of luminescence response, decreasing the fluorescence detection limit and improving the detection sensitivity.¹¹

In the crystal engineering of functional coordination polymers, the variable selection of metal ions and organic ligands provides endless possible outcomes of the final products. Although these porous coordination polymers with targeted secondary building units (SBUs) and topologies can be synthesized by careful selection of the metal ions and organic ligands, the final structures of the products are still highly influenced by several factors including metal-ligand ratio, pH, solvent,

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temperature, and soon. In particular, judicious selection of ligands is important for deliberate structural changes and functional properties of coordination polymers.¹² Therefore many new flexible or rigid ligands have been explored and assembled into metal–organic polymers *via* coordination bonds or supramolecular interactions such as hydrogen bonds, π – π stacking, and host–guest ionic interactions *et al.*^{13–16}

1,3,4-Oxadiazoles can be employed as one kind of efficient bridging ligands and offer a range of charge-balance requirements, alternative linking modes and different orientations of donor groups.^{17,18} Currently oxadiazole-pyridine and their derivatives have gained more interest as ligands to bridge different metal ions to form functional coordination polymers because of their intriguing bridging fashions.¹⁹ It is noted that investigation of the substituent position isomerization influence of oxadiazole-pyridine ligands on the formation of coordination polymers is still scarcely reported. Especially, to the best of our knowledge, 3-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (**L**₁) and 4-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (**L**₂) are never reported in the coordination chemistry. The development of new ligand systems is continuously an important aspect for the chemistry of metal–organic coordination polymer. Previously we also reported a series of functional coordination polymers using versatile N-containing ligands.²⁰ In this work two position-isomeric oxadiazol-pyridine ligands **L**₁ and **L**₂ have been employed, a series of novel coordination polymers, namely [Cd₂(μ_2 -**L**₁)₂(μ_2 -NCS)₄]_n (**1**), {[Cd(**L**₁)(μ_2 -dca)₂(H₂O)]·H₂O}_n (**2**), {[Cu(μ_2 -**L**₁)₂(NCS)₂]·0.5H₂O}_n (**3**), {[Ag₂(μ_2 -**L**₁)(μ_3 -**L**₁)₂]·2PF₆]_n (**4**), {[Ag₃(μ_2 -**L**₁)₄(μ_2 -CF₃SO₃)(CF₃SO₃)·CF₃SO₃]_n (**5**), {[Cd(**L**₂)₂(μ_2 -NCS)₂]_n (**6**), [Ag(μ_2 -**L**₂)(μ_2 -CF₃SO₃)]_n (**7**) and {[Ag(μ_2 -**L**₂)]·BF₄]_n (**8**) have been isolated. Variable temperature magnetic susceptibility measurements (2–300 K) reveal anti-ferromagnetic interactions between central copper(II) ions for **3**. Solid-state luminescent properties of **1**, **2** and **4–8** have been investigated indicating strong fluorescent emissions. Additionally, luminescent measurements illustrate that **8** exhibits highly sensitive luminescence sensing for Cr₂O₇^{2–} in aqueous solutions with high quenching efficiency $K_{sv} = 2.08 \times 10^4$ L mol^{–1} and low detection limit (0.19 μ M (S/N = 3)), which make it a promising candidate for sensing Cr₂O₇^{2–} ions in practical. Luminescent measurements illustrate that **8** also represents the first report of coordination polymers based on oxadiazol-pyridine derivatives as luminescent response to Cr₂O₇^{2–} ions in the water solutions. Different coordination modes of **L**₁ and **L**₂ are also briefly discussed, which indicates these isomeric oxadiazol-pyridine building blocks have great potential in the construction of these unique metal organic coordination polymers.

Experimental section

General methods

Ligands **L**₁ and **L**₂ were prepared by literature methods.²¹ All the other reagents were purchased commercially and used without further purification. Deionized water was used as solvent in this work. C, H, and N microanalyses were carried out with a Perkin-Elmer 240 elemental analyzer. FT-IR spectra were recorded from KBr pellets in the range 4000–400 cm^{–1} on a Bio-Rad FTS 135

spectrometer. Variable-temperature magnetic susceptibilities were measured using a MPMS-7 SQUID magnetometer. Diamagnetic corrections were made with Pascal's constants for all constituent atoms. Photoluminescence spectra were measured by MPF-4 fluorescence spectrophotometer with a xenon arc lamp as the light source.

Synthesis of 3-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L**₁).** 35 mmol of the nicotinohydrazide were suspended in 95 mmol (2.7 eq.) triethyl orthoacetate. The reaction mixture was heated under reflux for 12 hours. After cooling to ambient temperature the formed solid was filtered off, washed twice with 15 mL of diethylether and dried in air. Yield: 70% of theory. ¹H-NMR (400 MHz, DMSO-*d*₆): δ = 2.61 ppm (s, 3H, CH₃); 7.88–7.90 (m, 2H, arom. C–H); 8.80–8.82 (m, 2H, arom. C–H). ¹³C-NMR (100 MHz, DMSO-*d*₆): δ = 10.7 ppm (s, 1C, CH₃); 119.9 (s, 2C, arom. C–H); 130.5 (s, 1C, C_q); 150.9 (s, 2C, arom. C–H); 162.4 (s, 1C, C_q=N); 165.0 (s, 1C, C_q=N). FT-IR (cm^{–1}, KBr): 1565(m), 1549(w), 1465(m), 1430(m), 1349(w), 1247(w), 1131(m), 1086(m), 1044(m), 1014(m), 823(m), 705(m).

Synthesis of 4-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L**₂).** 35 mmol of the 4-pyridinecarboxylic hydrazide were suspended in 95 mmol (2.7 eq.) triethyl orthoacetate. The reaction mixture was heated under reflux for 12 hours. After cooling to ambient temperature the formed solid was filtered off, washed twice with 15 mL of diethylether and dried in air. Yield: 55% of theory. ¹H-NMR (400 MHz, DMSO-*d*₆): δ = 2.60 ppm (s, 3H, CH₃); 7.61–7.64 (m, 1H, arom. C–H); 8.32–8.34 (m, 1H, arom. C–H); 8.78–8.79 (m, 1H, arom. C–H); 9.13 (s, 1H, arom. N=C–H). ¹³C{¹H}-NMR (100 MHz, DMSO-*d*₆): δ = 10.6 ppm (s, 1C, CH₃); 120.0 (s, 1C, C_q); 124.3 (s, 1C, C_q); 133.9 (s, 1C, arom. C–H); 146.9 (s, 1C, arom. C–H); 152.2 (s, 1C, arom. C–H); 164.4 (s, 1C, arom. C–H); 162.1 (s, 1C, C_q=N). FT-IR (cm^{–1}, KBr): 1585(m), 1538(w), 1413(m), 1430(m), 1354(w), 1271(m), 1142(m), 1033(m), 834(m), 770(m).

Preparation of compounds 1–8

[Cd₂(μ_2 -**L**₁)₂(μ_2 -NCS)₄]_n (**1**). A mixture of CdCl₂ (18.33 mg, 0.10 mmol), **L**₁ (17.71 mg, 0.10 mmol), NH₄NCS (15.23 mg, 0.20 mmol) and water (20 mL) was refluxed and stirred for 12 h, then the resulting solutions were filtered. The filtrate was kept in a CaCl₂ desiccator. After a few days suitable colorless single crystals were obtained for X-ray diffraction study, yield 55%. Elemental analysis calcd (%) for C₂₀H₁₄Cd₂N₁₀O₂S₄: C 30.82%, H 1.81%, N 17.97%; found: C 30.97%, H 1.93%, N 18.16%. FT-IR (cm^{–1}, KBr): 2090(s), 1572(m), 1558(w), 1465(m), 1263(w), 1131(m), 1098(m), 1033(m), 1014(m), 814(m), 719(m).

{[Cd(μ_2 -**L**₁)(μ_2 -dca)₂(H₂O)]·H₂O}_n (**2**). A mixture of CdCl₂ (18.33 mg, 0.10 mmol), **L**₁ (17.71 mg, 0.10 mmol), Nadca (17.81 mg, 0.20 mmol) and water (20 mL) was refluxed and stirred for 12 h, then the resulting solutions were filtered. The filtrate was kept in a CaCl₂ desiccator. After a few days suitable colorless single crystals were obtained for X-ray diffraction study, yield 57%. Elemental analysis calcd (%) for C₂₁H₁₁CdN₉O₃: C 45.88%, H 2.02%, N 22.93%; found: C 46.15%, H 2.16%, N 23.12%. FT-IR (cm^{–1}, KBr): 3500(m), 2170(s), 1570(m), 1550(w), 1352(m), 1260(w), 1129(m), 1080(m), 1035(m), 1015(m), 815(m), 720(m).



$\{[\text{Cu}(\mu_2\text{-L}_1)_2(\text{NCS})_2] \cdot 0.5\text{H}_2\text{O}\}_n$ (3). A mixture of CuCl_2 (17.05 mg, 0.10 mmol), L_1 (17.71 mg, 0.10 mmol), NH_4NCS (15.23 mg, 0.20 mmol) and water (20 mL) was refluxed and stirred for 12 h, then the resulting solutions were filtered. The filtrate was kept in a CaCl_2 desiccator. After a few days suitable green single crystals were obtained for X-ray diffraction study, yield 37%. Elemental analysis calcd (%) for $\text{C}_{18}\text{H}_{14.5}\text{CuN}_8\text{O}_{2.5}\text{S}_2$: C 42.35%, H 2.86%, N 21.95%; found: C 42.46%, H 2.97%, N 22.13%. FT-IR (cm^{-1} , KBr): 3410(m), 2090(s), 1557(m), 1420(m), 1359(w), 1268(m), 1093(m), 816(m), 719(m).

$\{[\text{Ag}_2(\mu_2\text{-L}_1)_3] \cdot 2\text{PF}_6\}_n$ (4). A mixture of AgPF_6 (25.29 mg, 0.10 mmol), L_1 (17.71 mg, 0.10 mmol) and water (20 mL) was refluxed and stirred for 12 h, then the resulting solutions were filtered. The filtrate was kept in a CaCl_2 desiccator. After a few days colorless suitable single crystals were obtained for X-ray diffraction study, yield 59%. Elemental analysis calcd (%) for $\text{C}_{24}\text{H}_{21}\text{Ag}_2\text{F}_{12}\text{N}_9\text{O}_3\text{P}_2$: C 29.14%, H 2.14%, N 12.74%; found: C 29.34%, H 2.38%, N 12.97%. FT-IR (cm^{-1} , KBr): 1589(m), 1420(m), 1270(m), 1110(m), 838(s), 704(m).

$\{[\text{Ag}_3(\mu_2\text{-L}_1)_4(\mu_2\text{-CF}_3\text{SO}_3)(\text{CF}_3\text{SO}_3)] \cdot \text{CF}_3\text{SO}_3\}_n$ (5). A mixture of AgCF_3SO_3 (25.69 mg, 0.10 mmol), L_1 (17.71 mg, 0.10 mmol) and water (20 mL) was refluxed and stirred for 12 h, then the resulting solutions were filtered. The filtrate was kept in a CaCl_2 desiccator. After a few days suitable single crystals were obtained for X-ray diffraction study, yield 39%. Elemental analysis calcd (%) for $\text{C}_{35}\text{H}_{28}\text{Ag}_3\text{F}_9\text{N}_{12}\text{O}_{13}\text{S}_3$: C 29.70%, H 1.99%, N 11.87%; found: C 29.96%, H 2.06%, N 11.96%. FT-IR (cm^{-1} , KBr): 1589(m), 1410(m), 1275(m), 1168(m), 820(m), 1031(bm), 827(m), 720(m).

$\{[\text{Cd}(\text{L}_2)_2(\mu_2\text{-NCS})_2]\}_n$ (6). 6 was prepared through the similar synthetic method of 3 only using L_2 (17.71 mg, 0.10 mmol) to replace L_1 (17.71 mg, 0.10 mmol) in the reaction system. After a few days suitable colorless single crystals were obtained for X-ray diffraction study, yield 41%. Elemental analysis calcd (%) for $\text{C}_{18}\text{H}_{14}\text{CdN}_8\text{O}_2\text{S}_2$: C 39.24%, H 2.56%, N 20.34%; found: C 39.57%, H 2.79%, N 20.59%. FT-IR (cm^{-1} , KBr): 2100 (s), 1617(m), 1567(m), 1251(m), 728(m).

$[\text{Ag}(\mu_2\text{-L}_2) \cdot (\mu_2\text{-CF}_3\text{SO}_3)]_n$ (7). 7 was prepared through the similar synthetic method of 5 only using L_2 (17.71 mg, 0.10 mmol) to replace L_1 (17.71 mg, 0.10 mmol) in the reaction system. After a few days suitable colorless single crystals were obtained for X-ray diffraction study, yield 36%. Elemental analysis calcd (%) for $\text{C}_9\text{H}_7\text{Ag}_2\text{F}_3\text{N}_3\text{O}_4\text{S}$: C 20.55%, H 1.34%, N 7.99%; found: C 20.76%, H 1.57%, N 8.16%. FT-IR (cm^{-1} , KBr): 1616(m), 1565(m), 1418(m), 1272(m), 1142(m), 1033(m), 834(m), 707(m).

$\{[\text{Ag}(\mu_2\text{-L}_2)] \cdot \text{BF}_4\}_n$ (8). A mixture of AgBF_4 (0.10 mmol), L_2 (0.10 mmol) and water (20 mL) was refluxed and stirred for 12 h, then the resulting solutions were filtered. The filtrate was kept in a CaCl_2 desiccator. After a few days suitable single crystals were obtained for X-ray diffraction study, yield 46%. Elemental analysis calcd (%) for $\text{C}_{16}\text{H}_{14}\text{AgBF}_4\text{N}_6\text{O}_2$: C 37.17%, H 2.73%, N 16.26%; found: C 37.34%, H 2.97%, N 16.46%. FT-IR (cm^{-1} , KBr): 1616(m), 1566(m), 1416(m), 1266(m), 1051(bm), 836(m), 725(m).

X-ray crystallography

Diffraction intensities for complexes 1–8 were collected on a Bruker SMART 1000 CCD diffractometer with graphite-

monochromated Mo- $\text{K}\alpha$ radiation ($\lambda = 0.71073 \text{ \AA}$) by using the ω - ϕ scan technique. Lorentz polarization and MULTI-SCAN absorption corrections were applied. The structures were solved by direct methods and refined with the full-matrix least-squares technique using the SHELXS-97 and SHELXL-97 programs.^{22,23} Anisotropic thermal parameters were assigned to all non-hydrogen atoms. The positions of hydrogen atoms of the water molecules were located from difference maps and refined with isotropic temperature factors. The positions of other hydrogen atoms were generated geometrically. Analytical expressions of neutral-atom scattering factors were employed, and anomalous dispersion corrections were incorporated. The crystallographic data and details of refinements for compounds 1–8 are summarized in Table 1. Selected bond lengths and angles are listed in Table 2. Corresponding hydrogen bonds lengths and angles for 1–8 are listed in Table S1.†

Results and discussion

The crystal samples of complexes 1–8 can retain their crystalline integrity at room temperature for two weeks. As shown in Schemes 1 and 2, only nitrogen atoms of oxadiazole and pyridine rings can participate in coordination and the steric effect prevents oxadiazole oxygen atoms from metal coordination. There are five different coordination modes in 1–8: for 1, 3 and 5, these L_1 ligands in these coordination polymers contain bridging coordination modes a. For 2, these L_1 ligands contain mono-dentate coordination mode b, while in 4 two different coordination (bidentate bridging mode a and tridentate bridging mode c) for L_1 can be observed. For 6, mono-dentate coordination mode d for L_2 can be observed while for 7–8 bidentate bridging coordination mode for L_2 can be observed. These various different coordination modes also draw the self-assembly of coordination polymers to form different coordination polymers.

Structure of $[\text{Cd}_2(\mu_2\text{-L}_1)_2(\mu_2\text{-NCS})_4]_n$ (1)

As shown in Fig. 1, the asymmetric structural unit of $[\text{Cd}_2(\mu_2\text{-L}_1)_2(\mu_2\text{-NCS})_4]_n$ (1) contains two central Cd^{II} ions (Cd1 and Cd2), two μ_2 -bridging L_1 ligands and two μ_2 -bridging SCN^- groups. The central Cd^{II} atom is six-coordinated by two nitrogen atoms from L_1 , two SCN^- sulfur atoms and two SCN^- nitrogen atoms forming CdN_4S_2 donor set. The Cd–N distances vary from 2.302(4) to 2.431(2) Å, Cd–S distances are 2.7140(15) Å, all the N–Cd–N and N–Cd–S angles fall in the range of 83.24(12)–94.27(9)°, such coordination mode is in line with an octahedrally coordinated Cd^{II} center.

As shown in Fig. 2, these thiocyanate ligands adopt bidentate bridging coordination mode, which link these neighboring Cd^{II} centers forming $\{\text{Cd} \cdots \text{NCS} \cdots \text{Cd}\}$ chains. The shortest Cd \cdots Cd distance within the $\{\text{Cd} \cdots \text{NCS} \cdots \text{Cd}\}$ chains is 5.714(20) Å. On the other hand, as listed in Scheme 2(a), these L_1 ligands in 1 adopt μ_2 -bridging coordination mode. These $\{\text{Cd} \cdots \text{NCS} \cdots \text{Cd}\}$ chains are further interlinked *via* bi-dentate L_1 ligands forming the 2D coordination polymer. These L_1 carbon atoms (C(4) and C(8))





Table 1 Crystal data and structure refinement information for compounds 1–8

	1	2	3	4	5	6	7	8
Empirical formula	C ₂₀ H ₁₄ Cd ₂ N ₁₀ O ₂ S ₄	C ₂₁ H ₁₄ CdN ₉ O ₃	C ₁₈ H _{14.5} CuN ₈ O _{2.5} S ₂	C ₂₄ H ₂₁ Ag ₂ F ₁₂ N ₉ O ₃ P ₂	C ₃₅ H ₂₈ Ag ₃ F ₉ N ₁₂ O ₁₃ S ₃	C ₁₈ H ₁₄ CdN ₈ O ₂ S ₂	C ₉ H ₇ Ag ₂ F ₃ N ₃ O ₄ S	C ₁₆ H ₁₄ AgBF ₄ N ₆ O ₂
<i>F_w</i>	779.45	441.70	510.54	989.18	1415.48	550.89	418.11	517.01
Crystal syst.	Monoclinic	Monoclinic	Monoclinic	Triclinic	Triclinic	Monoclinic	Monoclinic	Monoclinic
Temperature (K)	299(2)	298(2)	300(2)	296(2)	296(2)	298(2)	296(2)	127.3(6)
Space group	C2/c	C2/c	P2(1)/n	P1	P1	P2(1)/c	P2(1)/n	P2/a
Space group number	15	12	11	2	2	14	14	15
Flack parameter	—	0	—	—	—	—	—	—
<i>a</i> (Å)	12.402(4)	25.8009(18)	8.9845(5)	9.0183(13)	8.6773(11)	14.0427(13)	8.1723(5)	13.7987(5)
<i>b</i> (Å)	15.8257(5)	8.5402(6)	10.6321(6)	10.8981(17)	15.4665(19)	10.0230(10)	19.7588(12)	11.4490(5)
<i>c</i> (Å)	14.732(5)	7.8195(5)	11.8557(7)	19.524(3)	18.997(2)	7.5702(7)	8.6182(5)	12.1306(6)
α (°)	90	90	90	104.313(3)	77.458(3)	90	90	90
β (°)	112.998(11)	100.454(2)	107.885(2)	91.093(3)	82.510(3)	102.733(3)	107.0410(10)	96.794(4)
γ (°)	90	90	90	112.549(3)	76.521(2)	90	90	90
<i>V</i> (Å ³)	2661.6(15)	1694.4(2)	1077.78(11)	1703.2(4)	2411.6(5)	1039.30(17)	1330.52(14)	1902.95(15)
<i>Z</i>	4	4	2	2	2	2	4	4
<i>F</i> (000)	1520	872	519	968	1392	548	816	1024
ρ (mg m ^{−3})	1.945	1.731	1.573	1.929	1.949	1.760	2.087	1.805
Abs coeff (mm ^{−1})	1.952	1.322	1.243	1.353	1.444	1.286	1.727	1.124
Data/restraints/params	2596/0/176	3295/1/227	2104/0/153	1281/30/97	11794/0/676	2025/0/143	2753/0/190	1685/49/160
GOF	1.168	1.011	1.045	1.005	1.016	1.056	1.069	1.094
<i>R</i> ₁ ^a (<i>I</i> = 2 σ (<i>I</i>))	0.0292	0.0220	0.0340	0.0549	0.0482	0.0297	0.0289	0.0414
w <i>R</i> ₂ ^a (all data)	0.0363	0.0231	0.0445	0.0858	0.0956	0.0363	0.0343	0.0439

^a $R_1 = \Sigma ||F_o| - |F_c|| / \Sigma |F_o|$, $wR_2 = [\Sigma w(F_o^2 - F_c^2)^2 / \Sigma w(F_o^2)]^{1/2}$.

Table 2 Selected bond lengths (Å) and angles (°) for 1–8^a

1					
Cd(1)–N(4A)	2.302(4)	Cd(1)–N(4)	2.302(4)	Cd(2)–N(1)	2.402(3)
Cd(1)–N(3A)	2.431(3)	Cd(1)–N(3)	2.431(3)	Cd(2)–N(1A)	2.402(3)
Cd(1)–S(2A)	2.7140(15)	Cd(1)–S(2)	2.7140(15)	Cd(2)–N(5B)	2.291(4)
Cd(2)–N(5)	2.291(4)	N(4)–Cd(1)–S(2A)	93.70(10)	N(3A)–Cd(1)–S(2A)	81.44(9)
N(4A)–Cd(1)–N(4)	102.64(18)	N(4A)–Cd(1)–N(3A)	170.09(13)	N(4A)–Cd(1)–S(2)	93.69(10)
N(4)–Cd(1)–N(3A)	83.24(12)	N(4A)–Cd(1)–N(3)	83.24(12)	N(3A)–Cd(1)–S(2)	94.27(9)
N(4)–Cd(1)–N(3)	170.09(13)	N(3A)–Cd(1)–N(3)	92.13(16)	S(2A)–Cd(1)–S(2)	173.86(6)
2					
Cd(1)–N(4)	2.261(3)	Cd(1)–N(6A)	2.264(4)	Cd(1)–N(1)	2.358(3)
Cd(1)–N(7)	2.326(4)	Cd(1)–N(9A)	2.326(3)	Cd(1)–O(2)	2.363(2)
N(4)–Cd(1)–N(6A)	98.40(13)	N(4)–Cd(1)–N(7)	92.38(14)	N(4)–Cd(1)–N(1)	90.76(11)
N(6A)–Cd(1)–N(7)	91.66(17)	N(4)–Cd(1)–N(9A)	168.22(14)	N(7)–Cd(1)–N(1)	173.45(13)
N(6A)–Cd(1)–N(9A)	93.34(14)	N(7)–Cd(1)–N(9A)	86.39(16)	N(4)–Cd(1)–O(2)	84.10(11)
N(6)–Cd(1)–N(1)	93.57(14)	N(7)–Cd(1)–O(2)	86.59(13)	N(6A)–Cd(1)–O(2)	177.01(13)
N(9A)–Cd(1)–N(1)	89.37(13)	N(1)–Cd(1)–O(2)	88.01(9)	N(9A)–Cd(1)–O(2)	84.13(12)
3					
Cu(1)–N(4)	1.974(2)	Cu(1)–N(4A)	1.974(2)	Cu(1)–N(1A)	2.0578(19)
Cu(1)–N(1)	2.0578(19)	N(4A)–Cu(1)–N(1A)	88.99(8)	N(1)–Cu(1)–N(1A)	180.0
N(4)–Cu(1)–N(4A)	180.0	N(4)–Cu(1)–N(1)	88.99(8)	N(4)–Cu(1)–N(1A)	91.01(8)
N(4A)–Cu(1)–N(1)	91.01(8)	N(4)–Cu(1)–N(1A)	91.01(8)		
4					
Ag(1)–N(2)	2.277(4)	Ag(1)–N(4A)	2.307(4)	Ag(2)–N(7)	2.421(4)
Ag(1)–N(8)	2.309(4)	Ag(1)–N(3)	2.406(4)	Ag(2)–N(1)	2.554(4)
Ag(2)–N(6)	2.225(4)	Ag(2)–N(9A)	2.231(4)	N(4A)–Ag(1)–N(3)	98.30(17)
N(2)–Ag(1)–N(4A)	124.79(16)	N(2)–Ag(1)–N(8)	111.05(14)	N(6)–Ag(2)–N(9A)	152.34(16)
N(4A)–Ag(1)–N(8)	109.87(16)	N(2)–Ag(1)–N(3)	102.49(14)	N(8)–Ag(1)–N(3)	108.07(14)
5					
Ag(1)–N(4)	2.209(4)	Ag(1)–N(12A)	2.211(4)	Ag(3)–N(10)	2.154(3)
Ag(1)–N(1)	2.458(4)	Ag(2)–N(7)	2.284(3)	Ag(3)–N(9)	2.152(3)
Ag(2)–N(3)	2.297(4)	Ag(2)–N(6)	2.352(4)	Ag(2)–O(5)	2.591(4)
N(4)–Ag(1)–N(1)	103.68(13)	N(12A)–Ag(1)–N(1)	104.13(14)	N(3)–Ag(2)–O(5)	77.79(13)
N(7)–Ag(2)–N(3)	143.19(14)	N(7)–Ag(2)–N(6)	109.20(13)	N(9)–Ag(3)–N(10)	175.45(15)
N(3)–Ag(2)–N(6)	104.96(13)	N(7)–Ag(2)–O(5)	94.06(14)	N(6)–Ag(2)–O(5)	123.73(15)
6					
Cd(1)–N(4)	2.325(3)	Cd(1)–N(4A)	2.325(3)	Cd(1)–S(1A)	2.7253(9)
Cd(1)–N(1)	2.364(2)	Cd(1)–N(1A)	2.364(2)	Cd(1)–S(1B)	2.7253(9)
N(4)–Cd(1)–N(4A)	180.00(15)	N(4)–Cd(1)–N(1)	90.51(9)	N(4)–Cd(1)–S(1A)	91.75(8)
N(4A)–Cd(1)–N(1)	89.49(9)	N(1)–Cd(1)–S(1B)	90.37(6)	N(1)–Cd(1)–S(1A)	89.63(6)
7					
Ag(1)–N(1)	2.165(2)	Ag(1)–N(3A)	2.184(2)	Ag(1)–O(2A)	2.623(2)
Ag(1)–O(4)	2.710(2)	N(1)–Ag(1)–N(3A)	164.72(9)	N(1)–Ag(1)–O(4)	94.19(1)
N(1)–Ag(1)–O(2A)	99.27(1)	N(3A)–Ag(1)–O(4)	83.54(1)		
8					
Ag(1)–N(1)	2.282(3)	Ag(1)–N(1)	2.282(3)	Ag(1)–N(3A)#1	2.349(3)
Ag(1)–N(3B)	2.349(3)	N(1)–Ag(1)–N(3A)	125.45(12)	N(3A)–Ag(1)–N(3B)	94.65(17)
N(1)–Ag(1)–N(1)	121.29(18)	N(1)–Ag(1)–N(3)	125.45(12)	N(1)–Ag(1)–N(3)	94.85(12)
N(1)–Ag(1)–N(3)	94.85(12)				

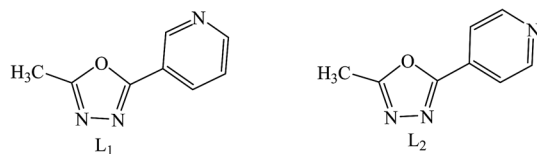
^a Symmetry transformations used to generate equivalent atoms: for 1: (A) $-x, y, -z + 1/2$; (B) $x + 1, y, z$. For 2: (A) $x, y, z - 1$. For 3: (A) $-x, -y + 1, -z + 1$. For 4: (A) $x + 1, y, z$. For 5: (A) $x + 2, y - 1, z + 1$. For 6: (A) $-x + 2, -y + 1, -z$; (B) $x, -y + 1/2, z - 1/2$. For 7: (A) $x - 1, -y + 3/2, z + 1/2$. For 8: (A) $3/2 - x, y, 2 - z$; (B) $1 - x, 1/2 + y, 3/2 - z$.

and SCN[−] nitrogen and sulfur atoms (N(4) and S(1)) generate these non-classical hydrogen bonds (C(4)–H(4)⋯N(4), 3.448(6) Å; C(8)–H(8B)⋯S(1), 3.652(5) Å), which further stabilize the 2D coordination polymer 1.^{24,25}

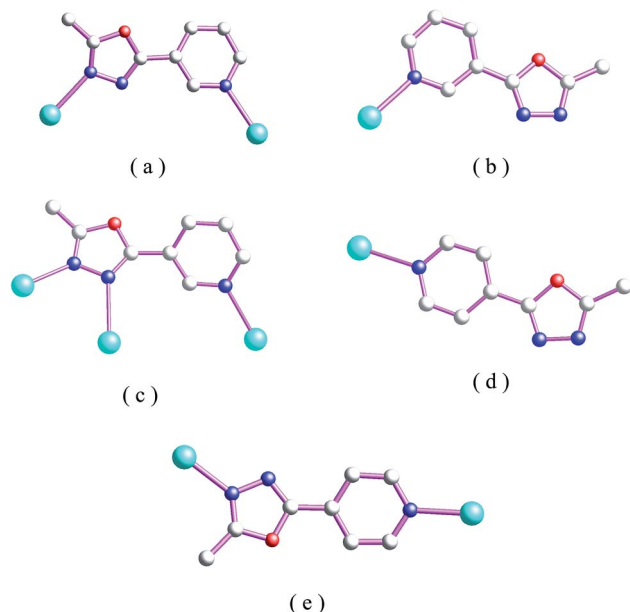
Structure of $\{[\text{Cd}(\text{L}_1)(\mu_2\text{-dca})_2(\text{H}_2\text{O})] \cdot \text{H}_2\text{O}\}_n$ (2)

The asymmetric structural unit of $\{[\text{Cd}(\mu_2\text{-L}_1)(\mu_2\text{-dca})_2(\text{H}_2\text{O})] \cdot \text{H}_2\text{O}\}_n$ (2) contains one central Cd^{II} ion (Cd1), one μ_2 -bridging L₁ ligand, two μ_2 -bridging dca[−] groups and one coordinated water





Scheme 1 Two isomeric oxadiazol-pyridine ligands 3-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L_1) and 3-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L_2).



Scheme 2 Five different coordination modes of L_1 and L_2 in **1–8** (color code: cyan, metal; blue, N; light grey, C).

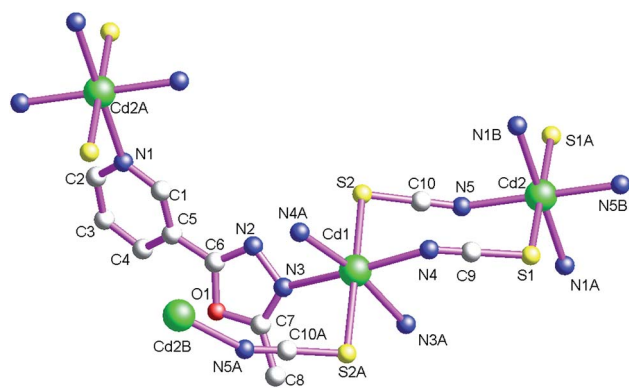


Fig. 1 The fundamental structural unit of $[\text{Cd}_2(\mu_2\text{-L})_2(\mu_2\text{-NCS})_4]_n$ (**1**).

molecule (O2) and one lattice aqua molecule (O3) (Fig. 3). The central Cd^{II} atom is six-coordinated by one nitrogen atom (N1) from one L₁, four dca[−] nitrogen atoms (N4, N7, N6A and N9A) and one terminal aqua molecule (O2) forming CdN₅O donor set. The Cd–N distances are 2.261(3)–2.326(4) Å, Cd–O distances are 2.363(2) Å. All the N–Cd–N and N–Cd–O angles fall in the range of 84.10(11)–98.40(13)°, such coordination mode is in accordance with an octahedrally coordinated Cd^{II} center.

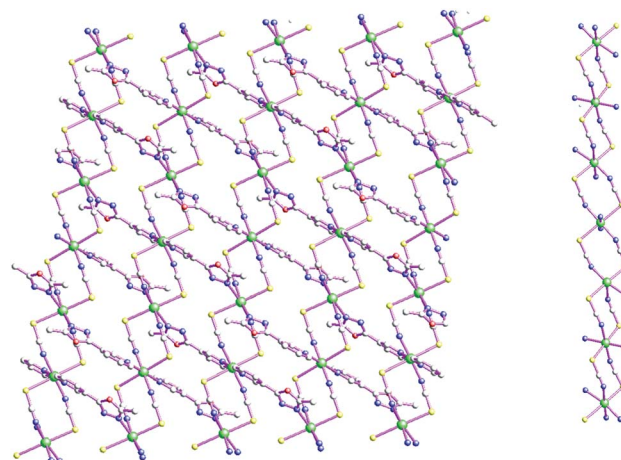


Fig. 2 2D coordination polymer $[\text{Cd}_2(\mu_2\text{-L}_1)_2(\mu_2\text{-NCS})_4]_n$ (**1**) containing infinite $\{\text{Cd}\text{-NCS}\text{-Cd}\}$ chains.

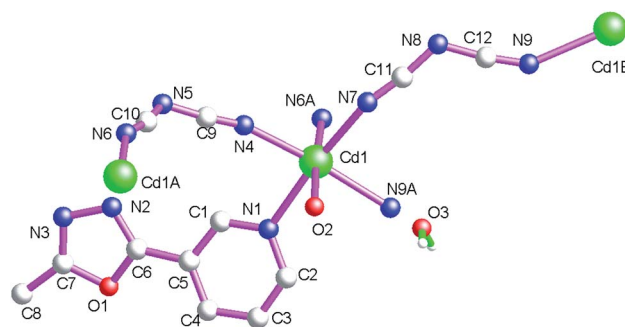


Fig. 3 The fundamental structural unit of $\{[\text{Cd}(\text{L}_1)(\mu_2\text{-dca})_2(\text{H}_2\text{O})] \cdot \text{H}_2\text{O}\}_n$ (2).

As shown in Fig. 4, these bidentate bridging dca^- ligands connect these neighboring Cd^{II} centers forming $\{\text{Cd}-\text{dca}-\text{Cd}\}$ layers. The shortest $\text{Cd}\cdots\text{Cd}$ distances within the $\{\text{Cd}-\text{dca}-\text{Cd}\}$ layers are 7.819(1) Å and 8.575(4) Å. On the other hand, as listed in Scheme 2(b), these L_1 ligands in **2** adopt mono-dentate coordination mode. Therefore these $\{\text{Cd}-\text{NCS}-\text{Cd}\}$ layers are further decorated *via* these mono-dentate L_1 ligands. These coordinated aqua molecules (O(2)), lattice aqua molecules (O(3)), L_1 nitrogen atoms (N(3)) and dca^- nitrogen atoms (N(8)) generate these classical hydrogen bonds $\text{O}(2)-\text{H}(2\text{A})\cdots\text{O}(3)$, 2.740(4) Å; $\text{O}(2)-\text{H}(2\text{B})\cdots\text{N}(3)$, 2.854(4) Å; $\text{O}(3)-\text{H}(3\text{A})\cdots\text{O}(2)$, 2.965(4) Å; $\text{O}(3)-\text{H}(3\text{B})\cdots\text{N}(8)$, 2.970(6) Å, which further stabilize the 2D coordination polymer **2**.

Structure of $\{[\text{Cu}(\mu_2\text{-L}_1)_2(\text{NCS})_2] \cdot 0.5\text{H}_2\text{O}\}_n$ (3)

As shown in Fig. 5, the asymmetric structural unit of **3** contains one central Cu^{II} ion (Cu1), one μ_2 -bridging L₁ ligand and two terminal bridging SCN[−] groups. The central Cu^{II} atom is six-coordinated by four nitrogen atoms (N1, N1A, N3A and N3B) from four L₁, two thiocyanate nitrogen atoms (N4 and N4A) forming the CuN₆ donor set. The Cu–N distances are 1.974(2)–2.0578(19) Å, all the N–Cu–N angles fall in the range of 88.99(8)–

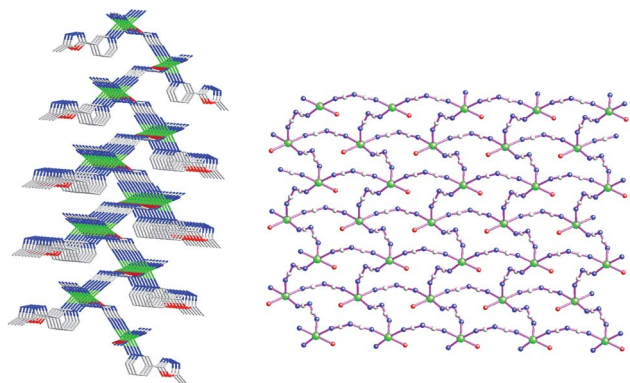


Fig. 4 Side view of 2D coordination polymer $\{[\text{Cd}(\text{L}_1)(\mu_2\text{-dca})_2(\text{H}_2\text{O})]\cdot\text{H}_2\text{O}\}_n$ (2) containing infinite $\{\text{Cd-dca-Cd}\}$ layers.

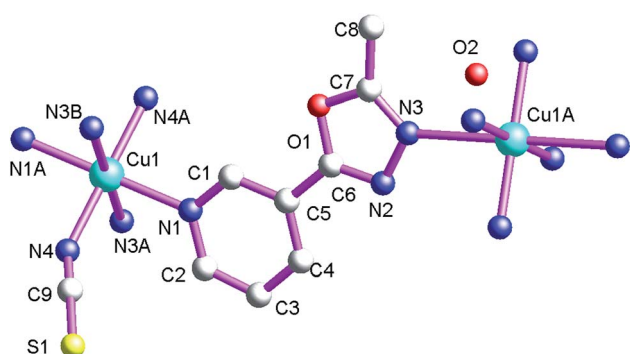


Fig. 5 The fundamental structural unit of $\{[\text{Cu}(\mu_2\text{-L}_1)_2(\text{NCS})_2]\cdot 0.5\text{H}_2\text{O}\}_n$ (3).

$91.01(8)^\circ$, such coordination mode is in accordance with an octahedrally coordinated Cu^{II} center.

As shown in Fig. 6 and Scheme 2(a), these L_1 ligands in 3 also adopt μ_2 -bridging coordination mode. These L_1 ligands connect neighboring Cu^{II} centers forming the 2D coordination polymer 3. The corresponding $\text{Cu}\cdots\text{Cu}$ distance within the 2D coordination polymer is $9.997(5)$ Å. These L_1 carbon molecules ($\text{C}(1)$) and L_1

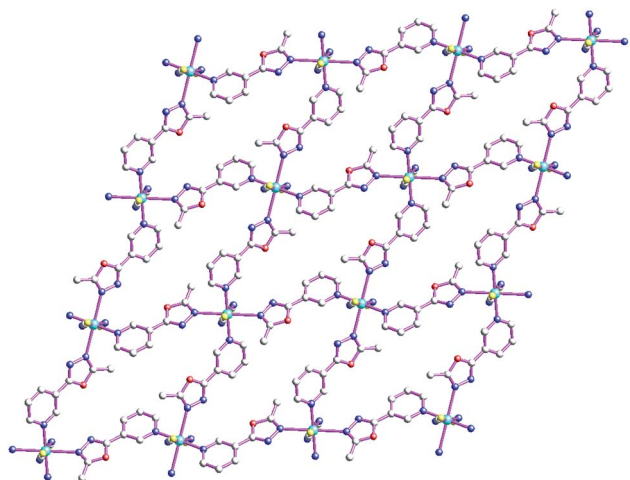


Fig. 6 2D coordination polymer $\{[\text{Cu}(\mu_2\text{-L}_1)_2(\text{NCS})_2]\cdot 0.5\text{H}_2\text{O}\}_n$ (3).

nitrogen atoms ($\text{N}(2)$) generate these non-classical hydrogen bonds ($\text{C}(1)\text{-H}(1)\cdots\text{N}(2)$, $3.304(3)$ Å), which also further extend 3 into a 3D supramolecular architecture. Such weak intermolecular interactions are key for the stabilization of the 2D coordination polymer 3.

Structure of $\{[\text{Ag}_2(\mu_2\text{-L}_1)(\mu_3\text{-L}_1)_2]\cdot 2\text{PF}_6\}_n$ (4)

When AgBF_4 are used, 1D $\text{Ag}(\text{I})$ coordination polymers, $\{[\text{Ag}_2(\mu_3\text{-L}_1)_2(\mu_2\text{-L}_1)]\cdot 2\text{PF}_6\}_n$ (4), can be isolated. For 4, the coordination polymer crystallizes in triclinic $P\bar{1}$ space group with one formula unit in the asymmetric unit. The asymmetric unit contains two Ag^{I} atoms (Ag1 and Ag2), two tri-dentate bridging L_1 , one bi-dentate bridging L_1 and two free PF_6^- anions (Fig. 7). The central Ag^{I} atom (Ag1) is four-coordinated by four nitrogen atoms (N2 , N4 , N3A and N4A) from four L_1 forming AgN_4 donor set. The other central Ag^{I} atom (Ag2) is also four-coordinated by four nitrogen atoms (N1 , N6 , N7 and N9A) from four L_1 forming AgN_4 donor set. The Ag-N distances are $2.225(4)$ – $2.554(4)$ Å, all the N-Ag-N angles are in the range of $98.30(17)$ – $152.34(16)^\circ$, such coordination mode is in accord with a tetrahedrally coordinated Ag^{I} center.

As shown in Fig. 8, Ag1 and Ag2 are inter-linked *via* oxadiazole nitrogen atoms forming di-nuclear $\text{Ag}(\text{I})$ clusters, neighboring $\text{Ag}\cdots\text{Ag}$ distances within di-nuclear $\text{Ag}(\text{I})$ clusters are $3.845(6)$ Å. These bridging L_1 ligands connect adjacent Ag^{I} ions forming a cluster-based 1D double chain coordination polymer, which is shown along the crystallographic a -axis. These non-classical hydrogen bonds ($\text{C}(8)\text{-H}(8\text{B})\cdots\text{F}(12)$, $3.341(9)$ Å; $\text{C}(17)\text{-H}(17)\cdots\text{F}(7)$, $3.190(10)$ Å; $\text{C}(17)\text{-H}(17)\cdots\text{F}(8)$, $3.347(10)$ Å; $\text{C}(21)\text{-H}(21)\cdots\text{N}(5)$, $3.504(8)$ Å) also can be observed, which further extend 4 into a 3D supramolecular architecture. Such weak intermolecular interactions are important for the stabilization of the 1D cluster-based Ag^{I} coordination polymer 4.

Structure of $\{[\text{Ag}_3(\mu_2\text{-L}_1)_4(\mu_2\text{-CF}_3\text{SO}_3)(\text{CF}_3\text{SO}_3)]\cdot \text{CF}_3\text{SO}_3\}$ (5)

When AgCF_3SO_3 is used, 1D $\text{Ag}(\text{I})$ coordination polymer, $\{[\text{Ag}_3(\mu_2\text{-L}_1)_4(\mu_2\text{-CF}_3\text{SO}_3)(\text{CF}_3\text{SO}_3)]\cdot \text{CF}_3\text{SO}_3\}$ (5), can be isolated.

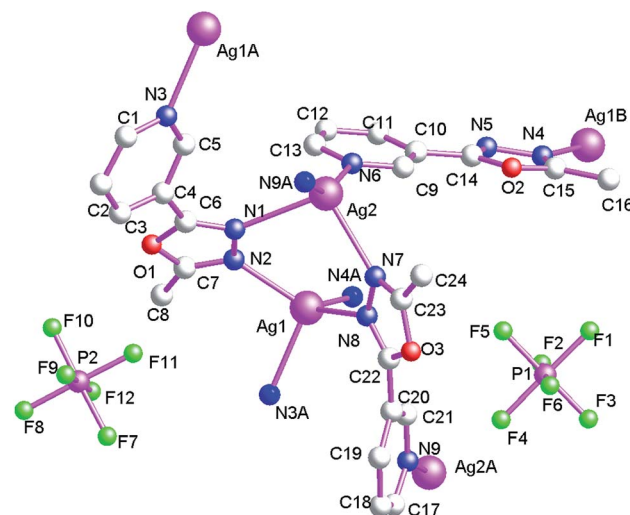


Fig. 7 The fundamental structural unit of $\{[\text{Ag}_2(\mu_2\text{-L}_1)(\mu_3\text{-L}_1)_2]\cdot 2\text{PF}_6\}_n$ (4).



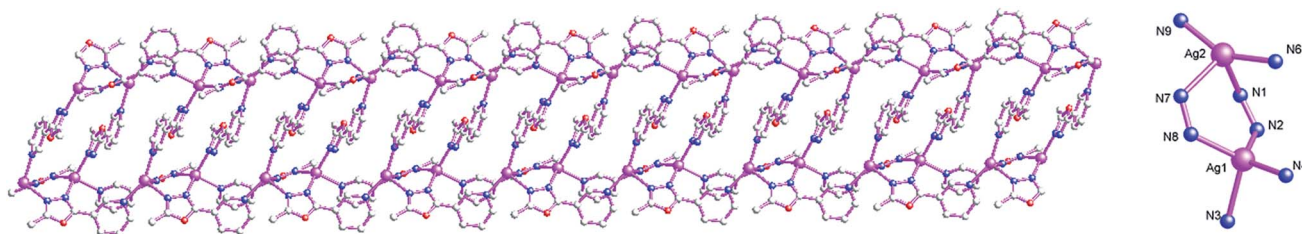


Fig. 8 1D Ag^I coordination polymer $\{[Ag_2(\mu_2-L_1)(\mu_3-L_1)_2] \cdot 2PF_6\}_n$ (4).

For 5, the coordination polymer also crystallizes in triclinic $P\bar{1}$ space group with one formula unit in the asymmetric unit. The asymmetric unit contains three Ag^I atoms (Ag1, Ag2 and Ag3), four bi-dentate bridging L₁, two terminal CF₃SO₃[−] anions and one free CF₃SO₃[−] anion (Fig. 9). The central Ag^I atom (Ag1) is three-coordinated by four nitrogen atoms (N4, N1 and N12A) from three L₁ forming AgN₃ donor set. The central Ag^I atom (Ag2) is four-coordinated by three nitrogen atoms (N3, N6 and N7) from three L₁ and one CF₃SO₃[−] oxygen atom (O5) forming AgN₃O donor set. The central Ag^I atom (Ag3) is three-coordinated by two nitrogen atoms (N9 and N10) from two L₁ and one CF₃SO₃[−] oxygen atom (O9) forming AgN₂O donor set. The Ag–N distances vary from 2.209(4) Å to 2.458(4) Å, Ag–O distances are 2.591(4) Å, all the N–Ag–N and N–Ag–O angles fall in the range of 77.79(13)–143.19(14)°, All the bond distances and bond angles all fall into the normal range.

As shown in Fig. 10, L₁ adopt bi-dentate bridging coordination mode, which connect neighboring Ag1, Ag2 and Ag3, forming 1D double chain coordination polymer 5. The shortest neighboring Ag⋯Ag distances within 1D double chain are 4.478(2) and 4.858(8) Å. These non-classical hydrogen bonds (C(1)–H(1A)⋯O(13), 3.348(7) Å; C(4)–H(4)⋯N(2), 3.259(6) Å; C(12)–H(12)⋯N(5), 3.241(6) Å; C(15)–H(15)⋯O(11), 3.393(7) Å) also can be observed,

which further extend 5 into a 3D supramolecular architecture. Such weak intermolecular interactions are key for the stabilization of 1D Ag^I double chain coordination polymer 5.

Structure of $\{[Cd(L_2)_2(\mu_2-NCS)_2]\}_n$ (6)

As shown in Fig. 11, the asymmetric structural unit of $\{[Cd(L_2)_2(\mu_2-NCS)_2]\}_n$ (6) contains one central Cd^{II} ions (Cd1), two terminal coordinated L₂ ligands and two μ₂-bridging NCS[−] groups. The central Cd^{II} atom is six-coordinated by two nitrogen atoms (N1 and N1A) from two L₂, two nitrogen atoms (N4 and N4A) and two sulfur atoms (S1A and S1B) form four NCS[−] groups forming CdN₄S₂ donor set. The Cd–N distances vary from 2.325(3) to 2.364(2) Å, Cd–S distances are 2.7253(9) Å. All the N–Cd–N and N–Cd–O angles are in the range of 89.63(6)–91.75(8)°, such coordination mode is in accordance with an octahedrally coordinated Cd^{II} center.

As shown in Fig. 12, these bidentate bridging NCS[−] ligands connect neighboring Cd^{II} centers forming the {Cd–NCS–Cd} layers. The shortest distances of Cd⋯Cd within the {Cd–NCS–Cd} layers are 6.280(1) and 7.570(1) Å. On the other hand, as listed in Scheme 2(b), these L₂ ligands in 6 adopt mono-dentate coordination mode. Therefore these {Cd–NCS–Cd} layers are

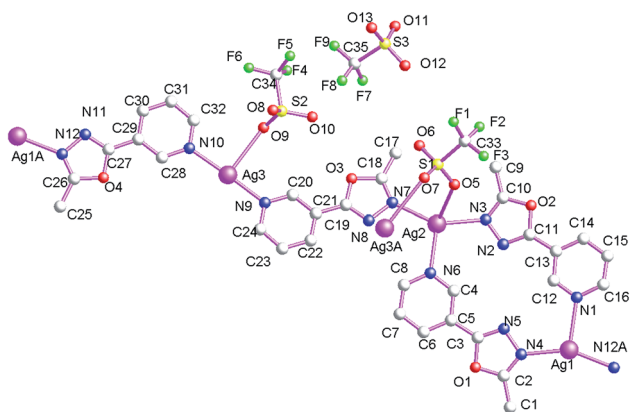


Fig. 9 The fundamental structural unit of $\{[Ag_3(\mu_2-L_1)_4(\mu_2-CF_3SO_3)(CF_3SO_3)] \cdot CF_3SO_3\}_n$ (5).

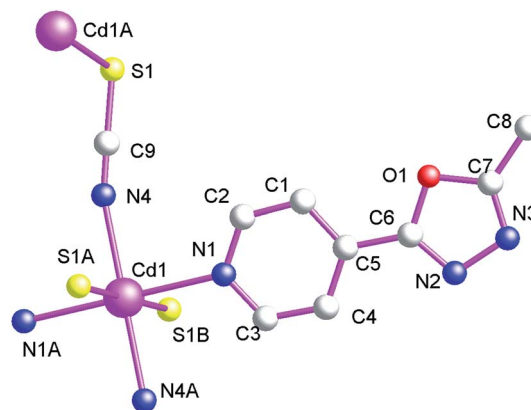


Fig. 11 The fundamental structural unit of $\{[Cd(\mu_2-L_2)_2(\mu_2-NCS)_2]\}_n$ (6).

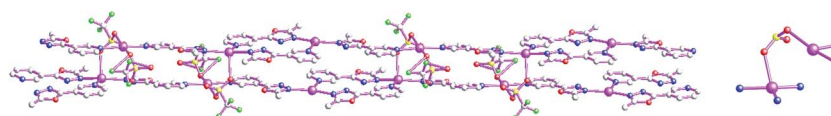


Fig. 10 1D Ag^I coordination polymer $\{[Ag_3(\mu_2-L_1)_4(\mu_2-CF_3SO_3)(CF_3SO_3)] \cdot CF_3SO_3\}_n$ (5).



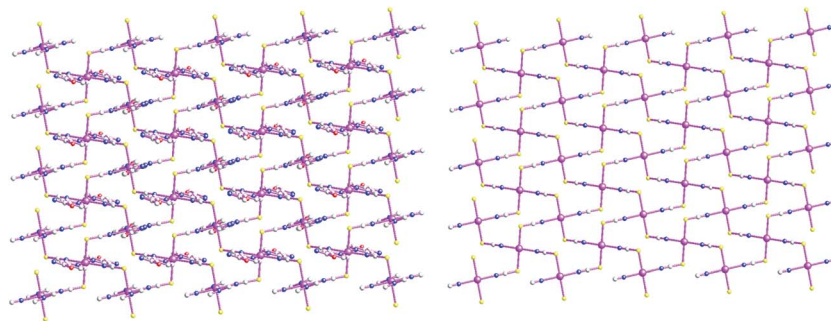


Fig. 12 2D coordination polymer $\{Cd(\mu_2-L_2)_2(\mu_2-NCS)_2\}_n$ (6) containing infinite $\{Cd-NCS-Cd\}$ layers.

further decorated *via* these mono-dentate L_2 ligands. These L_2 carbon atoms (C(1)) and nitrogen atoms (N(3)) generate non-classical hydrogen bonds (C(1)–H(1)⋯N(3), 3.528(4) Å), which further stabilize the 2D coordination polymer 6.

Structure of $[Ag(\mu_2-L_2)(\mu_2-CF_3SO_3)]_n$ (7)

As shown in Fig. 13, the asymmetric structural unit of $[Ag(\mu_2-L_2)(\mu_2-CF_3SO_3)]_n$ (7) contains one central Ag^I ions (Ag1), one μ_2 -bridging L_2 ligand, one μ_2 -bridging $CF_3SO_3^-$ group. For 7, the coordination polymer crystallizes in monoclinic $P2_1/n$ space group with one formula unit in the asymmetric unit. The central Ag^I atom (Ag1) is four-coordinated by two nitrogen atoms (N1 and N3A) from two L_1 and two $CF_3SO_3^-$ oxygen atoms (O2A and O4) forming AgN_2O_2 donor set. Ag–N distances are 2.165(2)–2.184(2) Å, Ag–O distances are 2.623(2)–2.710(2) Å, all the N–Ag–N and N–Ag–O angles are in the range of 83.54(1)–94.19(1)°, such coordination mode is in accord with a tetrahedrally coordinated Ag^I center.

As shown in Fig. 14, these $CF_3SO_3^-$ anions adopt bi-dentate bridging coordination modes *via* its two oxygen atoms (O2 and O4). On the other hand, these L_2 ligands also adopt bridging coordination mode as listed in Scheme 2(e). These bridging $CF_3SO_3^-$ anions and L_2 ligands link these neighboring $Ag(I)$ centers, which ultimately forming the 2D coordination polymer 7. These L_2 carbon atoms (C(2) and C(6)) and $CF_3SO_3^-$ oxygen and fluoride atoms (O(4) and F(1)) generate these non-classical

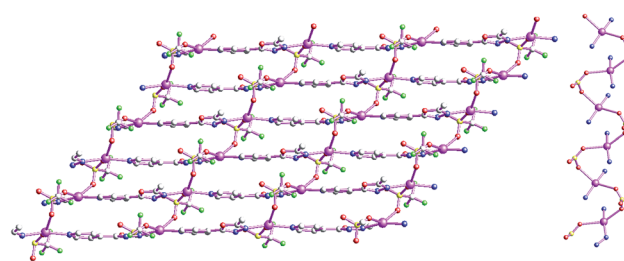


Fig. 14 2D coordination polymer $[Ag(\mu_2-L_2)(\mu_2-CF_3SO_3)]_n$ (7).

hydrogen bonds (C(2)–H(2C)⋯F(1), 3.338(5) Å; C(6)–H(6)⋯O(4), 3.308 Å), which further extend 7 into a 3D supramolecular architecture. Such weak intermolecular interactions are important for the stabilization of the 1D cluster-based Ag^I coordination polymer 7.

Structure of $\{[Ag(\mu_2-L_2)] \cdot BF_4\}_n$ (8)

As shown in Fig. 15, the fundamental structural unit of $\{[Ag(\mu_2-L_2)] \cdot BF_4\}_n$ (9) contains one central Ag^I ions (Ag1), one μ_2 -bridging L_2 ligand, one free BF_4^- group. For 8, the coordination polymer crystallizes in monoclinic I_2/a space group with one formula unit in the asymmetric unit. The central Ag^I atom (Ag1) is four-coordinated by four nitrogen atoms (N1, N3A, N1A and N3B) from four L_2 forming AgN_4 donor set. The Ag–N distances are 2.282(3)–2.349(3) Å, all the N–Ag–N angles are in the range of 94.65(17)–125.45(12)°, such coordination mode is in accord with a tetrahedrally coordinated Ag^I center.

As shown in Fig. 16, these L_2 ligands also adopt bridging coordination mode as listed in Scheme 2(e). These bridging L_2

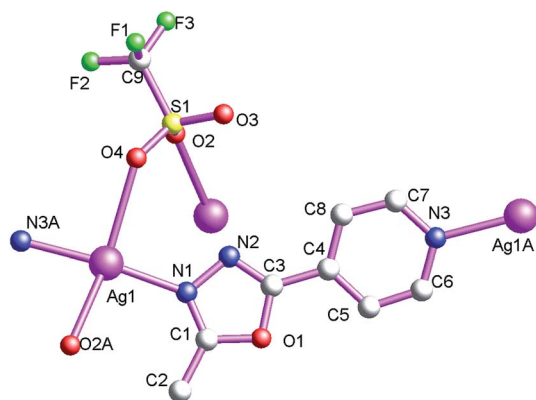


Fig. 13 The fundamental structural unit of $[Ag(\mu_2-L_2)(\mu_2-CF_3SO_3)]_n$ (7).

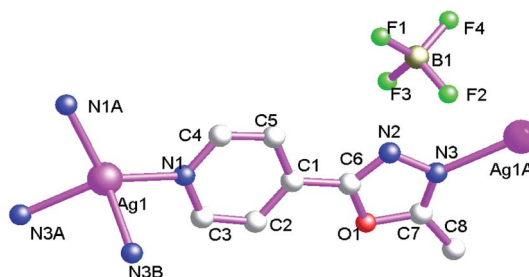


Fig. 15 The fundamental structural unit of $\{[Ag(\mu_2-L_2)] \cdot BF_4\}_n$ (8).



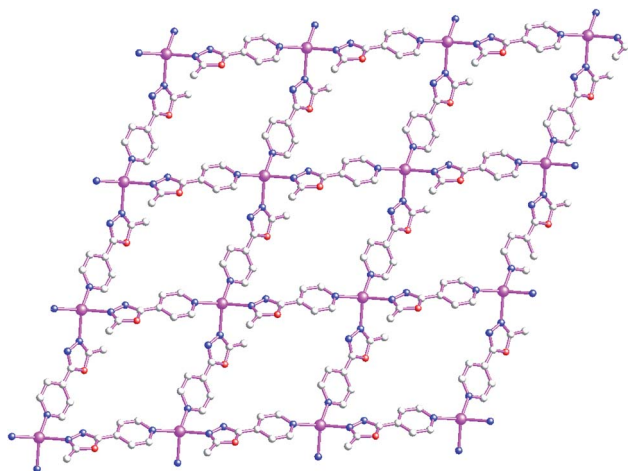


Fig. 16 2D cationic Ag^I coordination polymer {[Ag(μ_2 -L₂)]·BF₄]_n (8).

ligands link these neighboring Ag(I) centers, which ultimately forming the 2D grid-like cationic coordination polymer. These non-classical hydrogen bonds (C(4)–H(4)⋯F(2), 3.085(9) Å) also can be observed, which further extend **8** into a 3D supramolecular architecture. Such weak intermolecular interactions are important for the stabilization of the 2D Ag^I coordination polymer.

Magnetic properties of **3**

Variable temperature dc magnetic susceptibility data were collected on a microcrystalline powder sample of **3** in a 1000 Oe field in the temperature range 1.8–300 K. The temperature-dependent magnetic susceptibility data of complexes **3** have been measured in the temperature range of 2–300 K. The diamagnetic correction was evaluated by using Pascal's constants. As shown in Fig. 17, The $\chi_{\text{M}}T$ values are ca. 0.416 cm³ kmol^{−1} (for **3**) at room temperature, which are expected spin-only values for an isolated Cu(II) ion. When the temperature is lowered, $\chi_{\text{M}}T$ values remain constants and then smoothly decrease and drop rapidly below ca. 20 K reaching the

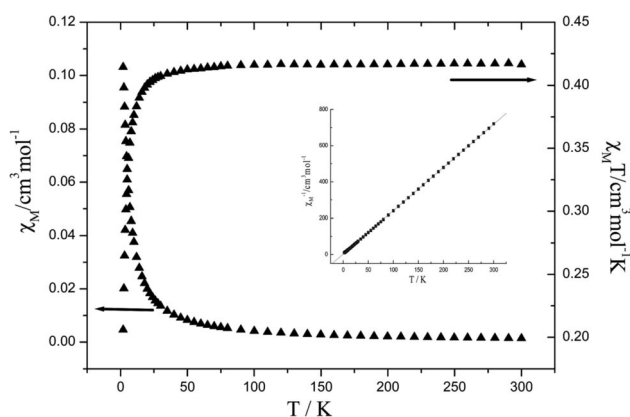


Fig. 17 Variable-temperature magnetic susceptibility data for **3** in the 2–300 K temperature range.

minimum values at 2 K. Fitting the magnetic susceptibility data with the Curie–Weiss law $\chi_{\text{M}} = C/(T - \theta)$, where C and θ represent the Curie constant and the Weiss temperature, it can give θ values of −1.05 K. These θ values reveal anti-ferromagnetic interactions between central Cu^{II} ions. The result of weak anti-ferromagnetic interactions can be attributed to can be attributed to very weak antiferromagnetic interactions as indicated by large metal–metal distances or the presence of zero-field splitting or both in **3**.²⁶

Luminescence properties

Coordination polymers have been investigated for potential applications as luminescent materials, such as light-emitting diodes (LEDs).²⁷ Owing to the ability of affecting the emission wavelength and strength of organic materials, syntheses of coordination polymers by judicious choice of conjugated organic spacers and transition metal centers can be an efficient method for obtaining new types of luminescent materials, especially for d¹⁰ or d¹⁰–d¹⁰ systems.²⁸ In the present work, we have explored the luminescent properties of L₁, L₂ ligands and organic/inorganic coordination polymers **1**, **2** and **4–8** based on the ligands in the solid state.

As shown in Fig. 18, at room temperature, free L₁ and L₂ ligands in the solid state are luminescent, which show the broad emission maximum centered at 360 nm ($\lambda_{\text{ex}} = 280$ nm) and 361 nm ($\lambda_{\text{ex}} = 280$ nm), respectively. For the free L₁ and L₂ ligands, the chromophores are aromatic pyridine and oxadiazol rings, the observed emission is due to π – π^* transition. In comparison with that of free ligand, upon excitation of solid samples of **1**, **2**, **6** and **8** show emission bands with maximum at ca. 360 nm (for **1**), ca. 361 nm (for **2**), ca. 363 nm (for **6**) and ca. 362 nm (for **8**) respectively. The emissions of **1**, **2**, **6** and **8** should be assigned to intraligand π – π^* transitions.²⁹ The slight shifts of the emission bands relative to free L₁ and L₂ ligands can be attributed to the coordination action of ligand to central metal ions.³⁰

On the other hand, for these 1D or 2D cluster-based Ag^I coordination polymers **4**, **5** and **7**, it is noted that red-shift of

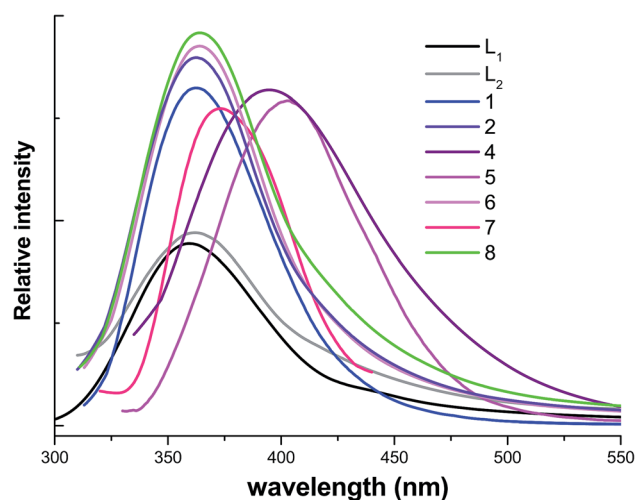


Fig. 18 Solid-state fluorescence spectra of **1–2**, **4–8** and L₁, L₂ ligand molecules.



luminescent emissions can be observed (emission bands with maximum centered at 395 nm for **4** ($\lambda_{\text{ex}} = 300$ nm), 403 nm for **5** ($\lambda_{\text{ex}} = 300$ nm) and 372 nm for **7** ($\lambda_{\text{ex}} = 300$ nm)). The photoluminescence origin of the emission bands should be attributed to metal-to-ligand charge transfer (MLCT) as discussed in the previous literature.¹ Because **5** and **6** contain similar 1D structural motifs, different band shapes in the luminescent emissions of **5** and **6** can be ascribed to different guest $\text{PF}_6^-/\text{CF}_3\text{SO}_3^-$ anions, which can greatly affect luminescence emission bands.³¹ These cluster-based Ag^{I} coordination polymers increase the ligand conformational rigidity, thereby reducing the non-radiative decay of the intraligand ($\pi-\pi^*$) excited state. The syntheses of new $\text{Cd}(\text{II})$ and $\text{Ag}(\text{I})$ complexes with these isomeric oxadiazol-pyridine ligands can be an efficient method for obtaining new types of luminescent materials.^{32,33}

Luminescent sensing properties of $\text{Cr}_2\text{O}_7^{2-}$ anion pollutants for **8**

In general, water pollutions are a global environmental issue, and considerable research attention has been paid to the removal of pollutants from waste water. With the development of modern industry, $\text{Cr}_2\text{O}_7^{2-}$ anions have been widely used in the field of chromium plating, metallurgy, pigment manufacturing, leather tanning, and wood preservation. These anion pollutants have been a focus of concern because they cause serious damage to human health and the environment. Therefore, the exploration of new materials for the efficient detection of these anion pollutants is highly important.³⁴

In this work, the product of **8** as the luminescent probes was carefully explored for sensing diverse ions, owing to its strong visible blue light when excited by ultraviolet light. The photoluminescence emission bands of **8** in aqueous solutions are centered at around 360 nm. These emission bands should be more likely to be due to intra-ligand transitions because similar photo-luminescent bands can be observed for free L_2 ligands in aqueous solutions. On the other hand, water stability of **8** was also investigated, the solid state samples were suspended in aqueous solutions for 12 h, then water solutions were filtrated and solid state samples were dried by the vacuum oven. Then solid state samples were characterized by PXRD patterns. As shown in Fig. S1,† PXRD patterns of solid state samples suspended in the water solutions and these calculated patterns deduced from single-crystal X-ray data are in good agreement. The result indicates that the coordination polymer structure is unchangeable and can retain stable in aqueous solutions.

In order to study the luminescent responses to different anions, the products of **8** were ground into powder and suspended in the water solution containing different salt solutions of the same concentration (10^{-4} M) of SiF_6^{2-} , CO_3^{2-} , HCO_3^{2-} , BF_4^- , NO_3^- , SCN^- , Ac^- , SO_4^{2-} , Br^- , Cl^- , F^- , I^- , BrO_3^- , IO_3^- , CrO_4^{2-} , HPO_4^{2-} and PO_4^{3-} and $\text{Cr}_2\text{O}_7^{2-}$. Aqueous solutions of SiF_6^{2-} , CO_3^{2-} , HCO_3^{2-} , BF_4^- , NO_3^- , SCN^- , Ac^- , SO_4^{2-} , Br^- , Cl^- , F^- , I^- , BrO_3^- , IO_3^- , CrO_4^{2-} , HPO_4^{2-} and PO_4^{3-} and $\text{Cr}_2\text{O}_7^{2-}$ were prepared from these corresponding potassium salts, respectively. As shown in Fig. 19(a), $\text{Cr}_2\text{O}_7^{2-}$ gave significant fluorescence quenching effect, while there was only

a negligible effect on the luminescence intensity for other anions. PXRD pattern of $\text{8-Cr}_2\text{O}_7^{2-}$ was measured and remains well consistent with the simulated one of **8**, indicating that **8** still remains stable. The result indicates the high selectivity of **8** for the detection and specific recognition of $\text{Cr}_2\text{O}_7^{2-}$ anions in aqueous solutions. Therefore **8** may be chosen as a candidate for the selective sensing of $\text{Cr}_2\text{O}_7^{2-}$.

Generally, waste water contains more than one type of pollutant anions, and therefore, it is essential to investigate the influence of mixed anions on the luminescence of **8**. The detailed experiments are as follows: 0.4 mL of $\text{Na}_2\text{Cr}_2\text{O}_7$ (10^{-4} M) and 0.4 mL of other anion solutions (10^{-4} M) were slowly dropped into a 3.2 mL suspension of **8**, respectively. The luminescence measurements of resultant solutions, containing $\text{Cr}_2\text{O}_7^{2-}$ and other anions, were carried out at once. It is also noted that there was only a negligible effect on the luminescence intensity for $\text{Cr}_2\text{O}_7^{2-}$ anions. This selective detection of $\text{Cr}_2\text{O}_7^{2-}$ anions has no interfering effects from other anions such as SiF_6^{2-} , CO_3^{2-} , HCO_3^{2-} , BF_4^- , NO_3^- , SCN^- , Ac^- , SO_4^{2-} , Br^- , Cl^- , F^- , I^- , BrO_3^- , IO_3^- , CrO_4^{2-} , HPO_4^{2-} and PO_4^{3-} , suggesting that **8** can selectively detect $\text{Cr}_2\text{O}_7^{2-}$ anions among the above anions.

Moreover, to explore the detection limit of **8** as a luminescent probe for detecting $\text{Cr}_2\text{O}_7^{2-}$, as shown in Fig. 19(b), a series suspension of $\text{8-Cr}_2\text{O}_7^{2-}$ (5 μM to 100 μM) were prepared by dropping different concentrations of $\text{Cr}_2\text{O}_7^{2-}$ solutions into the suspension of **8**. The luminescence intensity of **8** gradually decreases with increasing the concentration of $\text{Cr}_2\text{O}_7^{2-}$. As shown in Fig. 19(c) and (d), the luminescence intensity linearly decreases with the concentration of $\text{Cr}_2\text{O}_7^{2-}$ ranging from 0.5 μM to 70 μM . The detection limit of **8** as a luminescent probe for detecting $\text{Cr}_2\text{O}_7^{2-}$ is supported by the calculated results based on the equation: detection limit = $3\sigma/k$ (σ is the standard deviation of blank measurement; k is the slope between the luminescence intensity vs. $\log[\text{Cr}_2\text{O}_7^{2-}]$). To further explore the relationship between the quenching effect and $\text{Cr}_2\text{O}_7^{2-}$ concentration, the linear luminescence intensity vs. $\text{Cr}_2\text{O}_7^{2-}$ concentration plot was made, which can be fitted into $I_0/I = 1 + K_{\text{sv}}[\text{Cr}_2\text{O}_7^{2-}]$ (I_0 and I represent the luminescence intensity of **8** before and after adding $\text{Cr}_2\text{O}_7^{2-}$, respectively; $[\text{Cr}_2\text{O}_7^{2-}]$ represents the concentration of $\text{Cr}_2\text{O}_7^{2-}$, and K_{sv} represents the quenching rate constant). The K_{sv} value is calculated to be $2.08 \times 10^4 \text{ L mol}^{-1}$, indicating the high quenching efficiency of $\text{Cr}_2\text{O}_7^{2-}$ in the emission of **8**.^{35–37} The high quenching efficiency and low detection limit 0.19 μM ($\text{S/N} = 3$) also reveal that **8** can act as luminescent probes for discrimination and detection of $\text{Cr}_2\text{O}_7^{2-}$.

8 is a cationic coordination polymer, in which free BF_4^- anions are located between these 2D cationic layers, while in other Ag^{I} coordination polymers these anions are tightly linked to main frameworks through coordination bonds. As illustrated in the previous literature, these cationic coordination polymers can effectively capture $\text{Cr}_2\text{O}_7^{2-}$ anions through anion exchange process,³⁸ which may effectively lead to the fluorescent quenching of **8**. On the other hand, in comparison with other anions, only $\text{Cr}_2\text{O}_7^{2-}$ exhibits two wide absorption bands from 230 to 413 nm.³⁹ The bands almost cover the whole ranges of



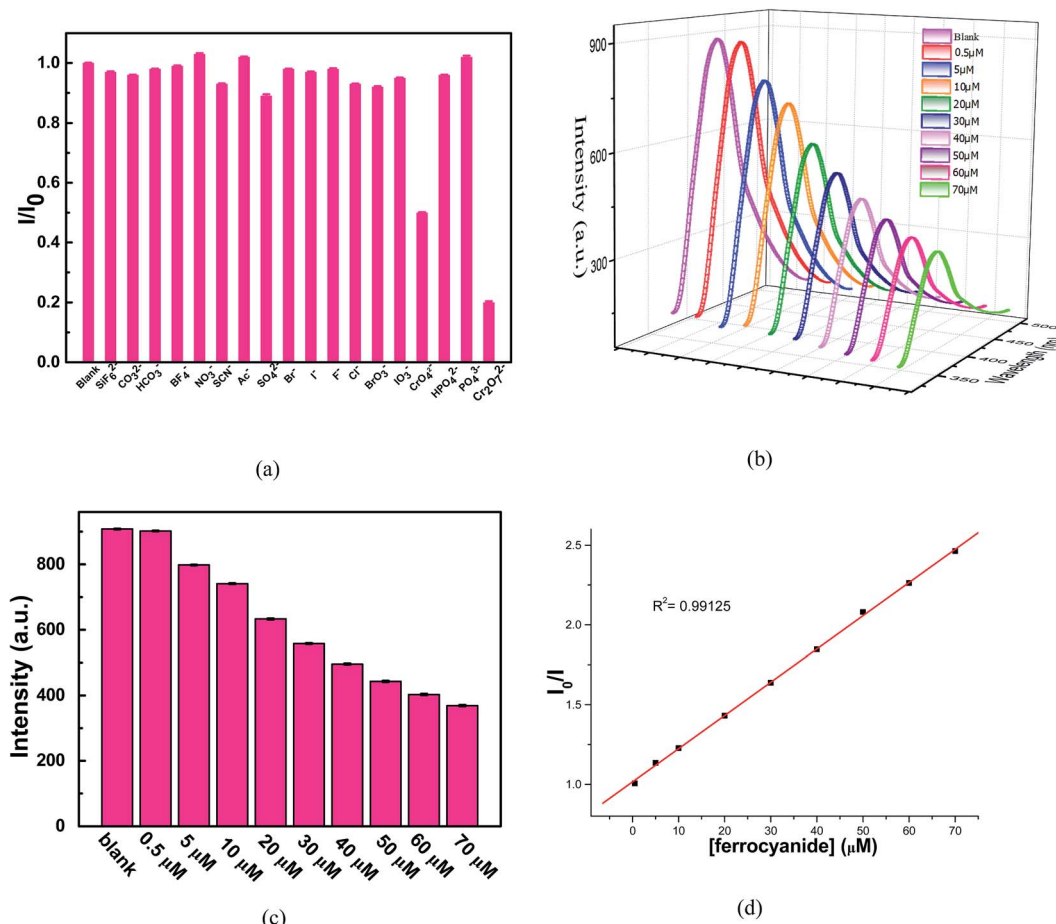


Fig. 19 (a) Comparison of the luminescence intensity of **8** at 360 nm incorporating aqua solutions (10^{-4} M) upon the addition of various anions; (b) liquid luminescence spectra of **8** under different concentrations of $\text{Cr}_2\text{O}_7^{2-}$ aqueous solution; (c) comparison of the luminescence intensity of **8** under different concentrations of $\text{Cr}_2\text{O}_7^{2-}$ anion pollutants aqueous solution. (d) The linear luminescence intensity vs. $\text{Cr}_2\text{O}_7^{2-}$ anion pollutants concentration plot.

absorption bands that arise from **8**. Thus, the luminescence quenching mechanism also further causes by the competition of excitation energy between $\text{Cr}_2\text{O}_7^{2-}$ anions and **8**. Upon illumination, the adsorption of $\text{Cr}_2\text{O}_7^{2-}$ ions for excitation energy hinders the UV/vis absorption of the target coordination polymer, thus resulting in a decrease, or even full quenching, of the luminescence intensities. This luminescent sensing mechanism for $\text{Cr}_2\text{O}_7^{2-}$ is consistent with those previously proposed by previous literature.^{40,41} To the best of our knowledge, work on these metal-organic polymers as luminescent probes for discrimination and detection of $\text{Cr}_2\text{O}_7^{2-}$ is important and still limited.⁴² **8** also represents the first report of coordination polymers based on oxadiazol-pyridine derivatives as luminescent response to $\text{Cr}_2\text{O}_7^{2-}$ in the water solutions.

Conclusions and perspectives

In conclusion, in this work two position-isomeric oxadiazol-pyridine ligands 3-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L_1) and 4-(5-methyl-1,3,4-oxadiazol-2-yl)pyridine (L_2) have been designed and synthesized. A series of novel coordination polymers, namely $[\text{Cd}_2(\mu_2-L_1)_2(\mu_2-\text{NCS})_4]_n$ (**1**), $[\text{Cd}(L_1)(\mu_2-\text{dca})_2(\text{H}_2\text{O})] \cdot$

H_2O_n (**2**), $[\text{Cu}(\mu_2-L_1)_2(\text{NCS})_2] \cdot 0.5\text{H}_2\text{O}_n$ (**3**), $[\text{Ag}_2(\mu_2-L_1)(\mu_3-L_1)_2] \cdot 2\text{PF}_6_n$ (**4**), $[\text{Ag}_3(\mu_2-L_1)_4(\mu_2-\text{CF}_3\text{SO}_3)(\text{CF}_3\text{SO}_3)] \cdot \text{CF}_3\text{SO}_3$ (**5**), $[\text{Cd}(L_2)_2(\mu_2-\text{NCS})_2]_n$ (**6**), $[\text{Ag}(\mu_2-L_2)(\mu_2-\text{CF}_3\text{SO}_3)]_n$ (**7**) and $[\text{Ag}(\mu_2-L_2)] \cdot \text{BF}_4_n$ (**8**) have been isolated. Both **1** and **2** are 2D Cd^{II} coordination polymers containing infinite $\{\text{Cd}-\text{NCS}-\text{Cd}\}$ chains (for **1**) or infinite $\{\text{Cd}-\text{dca}-\text{Cd}\}$ layers (for **2**). **3** is a 2D Cu^{II} coordination polymer, in which central metal ions are bridged *via* bidentate bridging L_1 ligand. While when different Ag^{I} salts were introduced into the reaction system, 1D cluster-based Ag^{I} coordination polymers **4** and **5** with diverse coordination modes can be isolated. Further when the isomeric oxadiazol-pyridine L_2 is used to replace L_1 in the reaction system, **6–8** can be isolated. **6** is a 2D Cd^{II} coordination polymer containing 2D $\{\text{Cd}-\text{NCS}-\text{Cd}\}$ layers. **7** is a 2D neutral Ag^{I} coordination polymer while **8** is a 2D cationic Ag^{I} coordination polymer. Variable temperature magnetic susceptibility measurements (2–300 K) reveal anti-ferromagnetic interactions between central copper(II) ions for **3**. Solid-state luminescent properties of **1**, **2** and **4–8** have been investigated indicating strong fluorescent emissions. Additionally, luminescent measurements illustrate that complex **8** also exhibits highly sensitive luminescence sensing for $\text{Cr}_2\text{O}_7^{2-}$ anion pollutants in aqueous solutions with high quenching



efficiency $K_{sv} = 2.08 \times 10^4 \text{ L mol}^{-1}$ and low detection limit (0.19 μM ($S/N = 3$)). **8** also represents the first report of coordination polymers based on oxadiazol-pyridine derivatives as luminescent response to $\text{Cr}_2\text{O}_7^{2-}$ anion pollutants in the water solutions. On the basis of this work, further syntheses, structures and properties studies of these coordination polymers using these isomeric oxadiazol-pyridine derivatives as basic building blocks are also under way in our laboratory.

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References

- (a) D. Venkataraman, S. Lee, J. S. Moore, P. Zhang, K. A. Hirsch, G. B. Gardner, A. C. Covey and C. L. Prentice, *Chem. Mater.*, 1996, **8**, 2030; (b) O. M. Yaghi, T. J. Pinnavaia and M. F. Thorpe, in *Access in Nanoporous Materials*, ed. Plenum, New York, 1995, p. 111; (c) A. Stein, S. W. Keller and T. E. Mallouk, *Science*, 1993, **259**, 1558; (d) P. J. Fagan and M. D. Ward, *Sci. Am.*, 1992, **267**, 48; (e) *Inclusion Compounds*, ed. T. Iwamoto, J. L. Atwood, J. E. D. Davies and D. D. MacNicol, Oxford, New York, 1991, vol. 5, p. 177; (f) B. F. Hoskins and R. Robson, *J. Am. Chem. Soc.*, 1990, **112**, 1546; (g) J. Lu, W. T. A. Harrison and A. J. Jacobson, *Angew. Chem., Int. Ed.*, 1995, **34**, 2557.
- (a) B. Moulton and M.-J. Zaworotko, *Chem. Rev.*, 2001, **101**, 1629; (b) O. M. Yaghi, M. O'Keeffe, N. W. Ockwig, H. K. Chae, M. Eddaoudi and J. Kim, *Nature*, 2003, **423**, 706; (c) S. Kitagawa, R. Kitaura and S. I. Noro, *Angew. Chem., Int. Ed.*, 2004, **43**, 2334; (d) G. Kubas, *Chem. Rev.*, 2007, **107**, 4152.
- (a) M. Eddaoudi, J. Kim, N. Rosi, D. Vodak, J. Wachter, M. O'Keeffe and O. Yaghi, *Science*, 2002, **295**, 469; (b) S. Bourrelly, P. Llewellyn, C. Serre, F. Millange, T. Loiseau and G. Férey, *J. Am. Chem. Soc.*, 2005, **127**, 13519; (c) B. Panella, M. Hirscher, H. Puetter and U. Mueller, *Adv. Funct. Mater.*, 2006, **16**, 520; (d) Y. Liu, J. F. Eubank, A. J. Cairns, J. Eckert, V. C. Kravtsov, R. Luebke and M. Eddaoudi, *Angew. Chem., Int. Ed.*, 2007, **46**, 3278; (e) A. Phan, C. J. Doonan, F. J. Uribe-Romo, C. B. Knobler, M. O'Keeffe and O. M. Yaghi, *Acc. Chem. Res.*, 2010, **43**, 58.
- (a) R.-B. Getman, Y. S. Bae, C.-E. Wilmer and R.-Q. Snurr, *Chem. Rev.*, 2011, **112**, 703; (b) P. Horcajada, R. Gref, T. Baati, P.-K. Allan, G. Maurin, P. Couvreur, G. Férey, R.-E. Morris and C. Serre, *Chem. Rev.*, 2011, **112**, 1232; (c) L.-E. Kreno, K. Leong, O.-K. Farha, M. Allendorf, R.-P. Van Duyne and J.-T. Hupp, *Chem. Rev.*, 2011, **112**, 1105; (d) J.-R. Li, J. Sculley and H.-C. Zhou, *Chem. Rev.*, 2011, **112**, 869.
- (a) J. R. Li, R. J. Kuppler and H. C. Zhou, *Chem. Soc. Rev.*, 2009, **38**, 1477; (b) R. Vaidhyanathan, S. S. Iremonger, G. K. H. Shimizu, P. G. Boyd, S. Alavi and T. K. Woo, *Science*, 2010, **330**, 650; (c) P. Nugent, Y. Belmabkhout, S. D. Burd, A. J. Cairns, R. Luebke, K. Forrest, T. Pham, S. Ma, B. Space, L. Wojtas, M. Eddaoudi and M. J. Zaworotko, *Nature*, 2013, **495**, 80; (d) H. Yin, J. Wang, Z. Xie, J. Yang, J. Bai, J. Lu, Y. Zhang, D. Yin and J. Y. S. Lin, *Chem. Commun.*, 2014, **50**, 3699.
- (a) R. Kitaura, K. Fujimoto, S. Noro, M. Kondo and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2002, **41**, 133; (b) R. Kitaura, G. Onoyama, H. Sakamoto, R. Matsuda, S. Noro and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2004, **43**, 2684; (c) B. Chen, F. R. Fronczek and A. W. Maverick, *Inorg. Chem.*, 2004, **43**, 8209; (d) P. A. Maggard, B. Yan and J. Luo, *Angew. Chem., Int. Ed.*, 2005, **44**, 2; (e) R. Custelcean and M. G. Gorbunova, *J. Am. Chem. Soc.*, 2005, **127**, 16362; (f) Y. Yoshida, K. Inoue and M. Kurmoo, *Inorg. Chem.*, 2009, **48**, 267; (g) Y. Inokuma, T. Arai and M. Fujita, *Nat. Chem.*, 2010, **2**, 780; (h) M. J. Manos, E. J. Kyprianidou, G. S. Papaefstathiou and A. J. Tasiopoulos, *Inorg. Chem.*, 2012, **51**, 6308.
- (a) M. Kurmoo, *Chem. Soc. Rev.*, 2009, **38**, 1353; (b) P. Mahata, S. Natarajan, P. Panissod and M. Drillon, *J. Am. Chem. Soc.*, 2009, **131**, 10140; (c) G. Rogez, N. Virat and M. Drillon, *Angew. Chem., Int. Ed.*, 2010, **49**, 1921; (d) D. D. Sante, A. Stroppa, P. Jain and S. Picozzi, *J. Am. Chem. Soc.*, 2013, **135**, 18126; (e) A. Stroppa, P. Barone, P. Jain, J. M. Perez-Mato and S. Picozzi, *Adv. Mater.*, 2013, **25**, 2284; (f) G. Beobide, W. Wang, O. Castillo, A. Luque, P. Román, G. Tagliabue, S. Galli and J. A. R. Navarro, *Inorg. Chem.*, 2008, 5267.
- (a) Y. Takashima, V. M. Martinez, S. Furukawa, M. Kondo, S. Shimomura, H. Uehara, M. Nakahama, K. Sugimoto and S. Kitagawa, *Nat. Commun.*, 2011, **2**, 168; (b) D. N. Dybtsev, H. Chun and K. Kim, *Angew. Chem., Int. Ed.*, 2004, **43**, 5033.
- (a) E. Y. Lee and M. P. Suh, *Angew. Chem., Int. Ed.*, 2004, **43**, 2798; (b) C. L. Chen, A. M. Goforth, M. D. Smith, C. Y. Su and H. C. zur Loye, *Angew. Chem., Int. Ed.*, 2005, **44**, 6673; (c) Q. Chu, D. C. Swenson and L. R. MacGillivray, *Angew. Chem., Int. Ed.*, 2005, **44**, 3569; (d) J. P. Ma, Y. B. Dong, R. Q. Huang, M. D. Smith and C. Y. Su, *Inorg. Chem.*, 2005, **44**, 6143; (e) J. Y. Lee, S. Y. Lee, W. Sim, K. M. Park, J. Kim and S. S. Lee, *J. Am. Chem. Soc.*, 2008, **130**, 6902; (f) Z. Wang and S. M. Cohen, *J. Am. Chem. Soc.*, 2007, **129**, 12368.
- (a) Y. Han, X. Li, L. Li, C. Ma, Z. Shen, Y. Song and X. You, *Inorg. Chem.*, 2010, **49**, 10781; (b) S. Dang, J.-H. Zhang, Z.-M. Sun and H. Zhang, *Chem. Commun.*, 2012, **48**, 11139; (c) S. Dang, J.-H. Zhang and Z.-M. Sun, *J. Mater. Chem.*, 2012, **22**, 8868; (d) W.-H. Zhu, Z.-M. Wang and S. Gao, *Inorg. Chem.*, 2007, **46**, 1337.
- (a) J. Rocha, L. D. Carlos, F. A. A. Paz and D. Ananias, *Chem. Soc. Rev.*, 2011, **40**, 926; (b) D. Banerjee, Z. Hu and J. Li, *Dalton Trans.*, 2014, **43**, 10668; (c) L. Zhang, Z. Kang, X. Xin and D. Sun, *CrystEngComm*, 2016, **18**, 193; (d) D. Zhao, Y. Cui, Y. Yang and G. Qian, *CrystEngComm*, 2016, **18**, 3746.



- 12 (a) D. J. Tranchemontagne, J. L. Mendoza-Cortés, M. O'Keeffe and O. M. Yaghi, *Chem. Soc. Rev.*, 2009, **38**, 1257; (b) L. Ma, C. Abney and W. Lin, *Chem. Soc. Rev.*, 2009, **38**, 1248; (c) J. Lee, O. K. Farha, J. Roberts, K. A. Scheidt, S. T. Nguyen and J. T. Hupp, *Chem. Soc. Rev.*, 2009, **38**, 1450; (d) M. D. Allendorf, C. A. Bauer, R. K. Bhakta and R. J. T. Houk, *Chem. Soc. Rev.*, 2009, **38**, 1330.
- 13 (a) H. Kajiro, A. Kondo, K. Kaneko and H. Kanoh, *Int. J. Mol. Sci.*, 2010, **11**, 3803; (b) L. Carlucci, G. Ciani, D. M. Proserpio, T. G. Mitina and V. A. Blatov, *Chem. Rev.*, 2014, **114**, 7557; (c) A. Schneemann, V. Bon, I. Schwedler, I. Senkovska, S. Kaskel and R. A. Fischer, *Chem. Soc. Rev.*, 2014, **43**, 6062.
- 14 (a) R. Kitaura, K. Seki, G. Akiyama and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2003, **42**, 428; (b) A. Kondo, H. Noguchi, S. Ohnishi, H. Kajiro, A. Tohdoh, Y. Hattori, W. C. Xu, H. Tanaka, H. Kanoh and K. Kaneko, *Nano Lett.*, 2006, **6**, 2581; (c) A. Kondo, T. Nakagawa, H. Kajiro, A. Chinen, Y. Hattori, F. Okino, T. Ohba, K. Kaneko and H. Kanoh, *Inorg. Chem.*, 2010, **49**, 9247; (d) D. Xiao, H. Chen, D. Sun, J. He, S. Yan, J. Yang, X. Wang, R. Yuan and E. Wang, *CrystEngComm*, 2012, **14**, 2849; (e) A. Kondo, N. Kojima, H. Kajiro, H. Noguchi, Y. Hattori, F. Okino, K. Maeda, T. Ohba, K. Kaneko and H. Kanoh, *J. Phys. Chem. C*, 2012, **116**, 4157; (f) Y. Jiang, J. Huang, B. Kasumaj, G. Jeschke, M. Hunger, T. Mallat and A. Baiker, *J. Am. Chem. Soc.*, 2009, **131**, 2058.
- 15 (a) Y. B. Dong, Q. Zhang, L. L. Liu, J.-P. Ma, B. Tang and R. Q. Huang, *J. Am. Chem. Soc.*, 2007, **129**, 1514; (b) Y. F. Han, W. G. Jia, Y. J. Lin and G. X. Jin, *Angew. Chem., Int. Ed.*, 2009, **34**, 6234.
- 16 (a) J.-G. Haasnoot, *Coord. Chem. Rev.*, 2000, **200–202**, 131; (b) J. P. Zhang, S. Horike and S. Kitagawa, *Angew. Chem., Int. Ed.*, 2007, **46**, 889.
- 17 (a) C. H. Lee, H. Y. Huang, Y. H. Liu, T. T. Luo, G. H. Lee, S. M. Peng, J. C. Jiang, I. Chao and K. L. Lu, *Inorg. Chem.*, 2013, **52**, 3962; (b) G.-M. Sun, F. Luo, Y.-M. Song, X.-Z. Tian, H.-X. Huang, Y. Zhu, Z.-J. Yuan, X.-F. Feng, M.-B. Luo, S.-J. Liu and W.-Y. Xu, *Dalton Trans.*, 2012, **41**, 11559.
- 18 B.-C. Tzeng and T.-Y. Chang, *Cryst. Growth Des.*, 2009, **9**, 5343.
- 19 Y. B. Dong, T. Sun, J. P. Ma, X. X. Zhao and R. Q. Huang, *Inorg. Chem.*, 2006, **45**, 10613.
- 20 J.-P. Ma, C.-W. Zhao, S.-Q. Wang, J.-P. Zhang, X. Niu and Y.-B. Dong, *Chem. Commun.*, 2015, **51**, 14586.
- 21 (a) B. Ding, L. Yi, Y. Wang, P. Cheng, D.-Z. Liao, S.-P. Yan and Z.-H. Jiang, *Dalton Trans.*, 2006, 665; (b) B. Ding, L. Yi, P. Cheng, D. Z. Liao and S. P. Yan, *Inorg. Chem.*, 2006, **45**, 5799; (c) B. Ding, Y.-Y. Liu, Y. Q. Huang, W. Shi, P. Cheng, D.-Z. Liao and S.-P. Yan, *Cryst. Growth Des.*, 2009, **9**, 593; (d) E.-C. Yang, H.-K. Zhao, B. Ding, X.-G. Wang and X.-J. Zhao, *Cryst. Growth Des.*, 2007, **7**, 2009; (e) B. Ding, P. Yang, Y.-Y. Liu, Y. Wang and G.-X. Du, *CrystEngComm*, 2013, **15**, 2490; (f) X.-X. Wu, Y.-Y. Wang, P. Yang, Y.-Y. Xu, J.-Z. Huo, B. Ding, Y. Wang and X.-G. Wang, *Cryst. Growth Des.*, 2014, **14**, 477; (g) J. Y. Liu, Q. Wang, L. J. Zhang, B. Yuan, Y. Y. Xu, X. Zhang, C. Y. Zhao, D. Wang, Y. Yuan, Y. Wang, B. Ding, X. J. Zhao and M. M. Yue, *Inorg. Chem.*, 2014, **53**, 5972; (h) Y. Wang, B. Yuan, Y. Y. Xu, X. G. Wang, B. Ding and X. J. Zhao, *Chem.-Eur. J.*, 2015, **21**, 2107.
- 22 G. Navarrete-Vázquez, G. M. Molina-Salinas, Z. V. Duarte-Fajardo, J. Vargas-Villarreal, S. Estrada-Soto, F. González-Salazar, E. Hernández-Núñez and S. Said-Fernández, *Bioorg. Med. Chem.*, 2009, **46**, 309.
- 23 G. M. Sheldrick, *SHELXS-97, Program for X-ray Crystal Structure Solution*, Göttingen University, Göttingen, Germany, 1997.
- 24 G. M. Sheldrick, *SHELXL-97, Program for X-ray Crystal Structure Refinement*, Göttingen University, Göttingen, Germany, 1997.
- 25 (a) B. Salignac, S. Riedel, A. Dolbecq, F. Sécheresse and E. Cadot, *J. Am. Chem. Soc.*, 2000, **122**, 10381; (b) A. Hijazi, J. C. Kemmagne-Mbouguen, S. Floquet, J. Marrot, J. Fize, V. Artero and E. Cadot, *Dalton Trans.*, 2013, **42**, 4848.
- 26 (a) A. Barba-Bon, A. M. Costero, S. Gil, M. Parra, J. Soto, R. Martinez-Manezac and F. Sancenon, *Chem. Commun.*, 2012, **48**, 3000; (b) G. S. Yang, M. N. Li, S. L. Li, Y. Q. Lan, W. W. He, X. L. Wang, J. S. Qin and Z. M. Su, *J. Mater. Chem.*, 2012, **22**, 17947.
- 27 Z. G. Gu, Y. F. Xu, X. H. Zhou, J. L. Zuo and X. Z. You, *Cryst. Growth Des.*, 2008, **8**, 1306.
- 28 (a) Z. Wang, K. Hu, S. Gao and H. Kobayashi, *Adv. Mater.*, 2010, **22**, 1526; (b) G. Rogez, N. Viart and M. Drillon, *Angew. Chem., Int. Ed.*, 2010, **49**, 1921; (c) D. Di Sante, A. Stroppa, P. Jain and S. Picozzi, *J. Am. Chem. Soc.*, 2013, **135**, 18126; (d) M. Mączka, A. Ciupa, A. Gągor, A. Sieradzki, A. Pikul, B. Macalik and M. Drozd, *Inorg. Chem.*, 2014, **53**, 5260.
- 29 (a) X. Li, X. W. Wang and Y. H. Zhang, *Inorg. Chem. Commun.*, 2008, **11**, 832; (b) C. A. Bauer, T. V. Timofeeva, T. B. Settersten, B. D. Patterson, V. H. Liu, B. A. Simmons and M. D. Allendorf, *J. Am. Chem. Soc.*, 2007, **129**, 7136.
- 30 (a) S. H. Rahaman, R. Ghosh, G. Mostafa and B. K. Ghosh, *Inorg. Chem. Commun.*, 2005, **8**, 1137; (b) Y. Lui and R. T. Tiekink, *CrystEngComm*, 2005, **7**, 20; (c) G. Singha, S. Girdhara and M. Garga, *Synth. React. Inorg., Met.-Org., Nano-Met. Chem.*, 2013, **43**, 1107–1111.
- 31 (a) J. M. Hao, B. Y. Yu, K. V. Hecke and G. H. Cui, *CrystEngComm*, 2015, **17**, 2279; (b) G. H. Cui, C. H. He, C. H. Jiao, J. C. Geng and V. A. Blatov, *CrystEngComm*, 2012, **14**, 4210; (c) X. X. Wang, B. Y. Yu, K. V. Hecke and G. H. Cui, *RSC Adv.*, 2014, **4**, 61281; (d) L. Qin, J. Zheng, S. L. Xiao, X. H. Zheng and G. H. Cui, *Inorg. Chem. Commun.*, 2013, **34**, 71; (e) X. X. Wang, M. X. Zhang, B. Y. Yu, K. V. Hecke and G. H. Cui, *Spectrochim. Acta, Part A*, 2015, **139**, 442.
- 32 (a) C. M. Che, C. W. Wan, K. Y. Ho and Z. Y. Zhou, *New J. Chem.*, 2001, **25**, 63; (b) J. H. Luo, M. C. Hong, R. H. Wang, R. Cao, L. Han and Z. Z. Lin, *Eur. J. Inorg. Chem.*, 2003, 2705; (c) X. L. Wang, C. Qin, E. B. Wang and Z. M. Su, *Chem.-Eur. J.*, 2006, **12**, 2680.
- 33 (a) C. X. Chen, Q. K. Liu, J. P. Ma and Y. B. Dong, *J. Mater. Chem.*, 2012, **22**, 9027; (b) C. Fang, Q. K. Liu, J. P. Ma and Y. B. Dong, *Inorg. Chem.*, 2012, **51**, 3923.
- 34 (a) Q. G. Zhai, X. Y. Wu, S. M. Chen, C. Z. Lu and W. B. Yang, *Cryst. Growth Des.*, 2006, **6**, 2126; (b) C. Ren, Y. N. Zhang,



- W. J. Shi, B. Liu, Y. Y. Wang and Q. Z. Shi, *CrystEngComm*, 2011, **13**, 5179; (c) J. Y. Gao, N. Wang, X. H. Xiong, C. J. Chen, W. P. Xie, X. R. Ran, Y. Long, S. T. Yue and Y. L. Liu, *CrystEngComm*, 2013, **15**, 3261.
- 35 K. L. Wong, G. L. Law, Y. Y. Yang and W. T. Wong, *Adv. Mater.*, 2006, **18**, 1051.
- 36 (a) C. X. Chen, Q. K. Liu, J. P. Ma and Y. B. Dong, *J. Mater. Chem.*, 2012, **22**, 9027; (b) C. Fang, Q. K. Liu, J. P. Ma and Y. B. Dong, *Inorg. Chem.*, 2012, **51**, 3923.
- 37 (a) Q. G. Zhai, X. Y. Wu, S. M. Chen, C. Z. Lu and W. B. Yang, *Cryst. Growth Des.*, 2006, **6**, 2126; (b) C. Ren, Y. N. Zhang, W. J. Shi, B. Liu, Y. Y. Wang and Q. Z. Shi, *CrystEngComm*, 2011, **13**, 5179; (c) J. Y. Gao, N. Wang, X. H. Xiong, C. J. Chen, W. P. Xie, X. R. Ran, Y. Long, S. T. Yue and Y. L. Liu, *CrystEngComm*, 2013, **15**, 3261.
- 38 P. F. Shi, B. Zhao, G. Xiong, Y. L. Hou and P. Cheng, *Chem. Commun.*, 2012, **48**, 8231.
- 39 K. L. Wong, G. L. Law, Y. Y. Yang and W. T. Wong, *Adv. Mater.*, 2006, **18**, 1051.
- 40 Z. Xie, L. Ma, K. E. deKrafft, A. Jin and W. Lin, *J. Am. Chem. Soc.*, 2010, **132**, 922.
- 41 (a) W. S. Liu, T. Q. Jiao, Y. Z. Li, Q. Z. Liu, M. Y. Tan, H. Wang and L. F. Wang, *J. Am. Chem. Soc.*, 2004, **126**, 2280; (b) X. Yang, Z. Huang, J. Dang, C. L. Ho, G. Zhou and W. Y. Wong, *Chem. Commun.*, 2013, **49**, 4406; (c) H. X. Zhao, L. Q. Liu, Z. D. Liu, Y. Wang, X. J. Zhao and C. Z. Huang, *Chem. Commun.*, 2011, **47**, 2604.
- 42 (a) Z. C. Hu, B. J. Deibert and J. Li, *Chem. Soc. Rev.*, 2014, **43**, 5815; (b) H. Xu, C. S. Cao, X. M. Kang and B. Zhao, *Dalton Trans.*, 2016, **45**, 18003.

