

## RESEARCH ARTICLE

View Article Online  
View Journal | View IssueCite this: *Inorg. Chem. Front.*, 2024,  
11, 5709Supracluster assembly of an  $Al_4$  precursor toward  
the study of enhanced optical properties†Fan Yang,<sup>‡a,b</sup> Rui-Yan Chen,<sup>‡a,c</sup> San-Tai Wang,<sup>a,b</sup> Yan-Ping He,<sup>ID \*a</sup>  
Wei-Hui Fang<sup>ID \*a,b</sup> and Jian Zhang<sup>ID a</sup>

Supracluster assembly offers a unique opportunity for the isolation of customized materials using functional cluster precursors. Herein, for the first time, we demonstrate supracluster assembly based on an  $Al_4^{4-}$  precursor through both non-covalent interactions and coordination bonds. By reacting readily available, ultrastable, and soluble  $Al_4^{4-}$  as an anionic precursor with a metal center and neutral chelated chromophores, we successfully synthesized an extensive library of supracluster assembled crystalline solids, ranging from ion-pair structures, surface-modified clusters, and chains, to layered compounds. The advantage of this method is that it achieves stepwise performance regulation and enhancement using given functional precursors. These materials from supracluster assembly possess a broad range of tunable photophysical properties, including photoluminescence and third-order nonlinear optical (NLO) properties. Such extraordinary supracluster assembly will be widely applicable for creating a new generation of rationally organized solid materials with tailorable and customized properties.

Received 5th May 2024,  
Accepted 17th July 2024

DOI: 10.1039/d4qi01103a

rsc.li/frontiers-inorganic

## Introduction

Post-synthetic modification is a useful technique to precisely prepare new target materials by introducing desired properties into pre-designed structures.<sup>1–3</sup> Such a method involves introducing small organic molecules or polymers, metal ions, *etc.* into pre-synthesized materials without destroying the structure of the material itself so as to obtain new functional materials. Therefore, it is prevalent and widely used in a broad range of materials, including metal- and covalent organic frameworks,<sup>1,3–5</sup> porous- and metal-organic cages,<sup>2,6</sup> and interlocked molecules.<sup>7</sup>

The molecular nature of discrete metal-organic cages and cluster compounds endows them with easy solution-processability unattainable with traditional framework materials.<sup>8</sup> Thus, these discrete compounds can be adapted for stepwise assembly through solution processing in addition to the solid state. In this regard, the stepwise approach involves dissolving these discrete supramolecular entities in an appropriate

solvent and then reacting them with the desired functional element in covalent, coordinated and noncovalent ways.<sup>6</sup> Typical representatives are tetrahedral Ti-,<sup>9–13</sup> Zr-,<sup>8,14–19</sup> Fe-,<sup>20–22</sup> and Ln-based cages,<sup>23,24</sup> cuboctahedral Cr-based cages, *etc.*<sup>25–27</sup> The most distinguished representative in cluster science is polyoxometalates.<sup>28–33</sup> However, the stepwise assembly method based on main group metal clusters has not been effectively developed. Typical examples include the assembly of anionic  $[V_{10}O_{28}]^{6-}$ / $[\alpha-1,2,3-\alpha-1,2,3-SiV_3W_9O_{40}]^{7-}$  and cationic  $[AlO_4Al_{12}(OH)_{24}(H_2O)_{12}]^{7+}$  clusters.<sup>34,35</sup>

Recently, we reported the supracluster assembly of Archimedean  $\{Al_6M_3\}_8$  cages with 72 hydrogen bonds.<sup>36</sup> The supracluster assembly fills the gap between “supramolecular”<sup>37</sup> and “superparticle”<sup>38</sup> systems. The word “supracluster” is rarely used but was once used to describe and emphasize the size of clusters, where Teo and Dale named compounds  $[(Ph_3P)_{10}Au_{12}Ag_{12}PtCl_7]Cl$  and  $[Ru_6C(CO)_{17}]$  in 1993 and 1995,<sup>39,40</sup> whereas our proposed supracluster assembly aims to achieve “chemistry beyond the cluster”, in that attention should be paid to the interactions between clusters, such as hydrogen bonds,  $\pi \cdots \pi$  interactions and covalent bonds, with the interactions between the vertex and edge-sharing supracluster assemblies based on Archimedean  $\{Al_6M_3\}_8$  cages being hydrogen bonds. In our previous work, we investigated the coordination-driven self-assembly of aluminum molecular rings, which can be viewed as supracluster assembly *via* coordination bonds.

Herein, for the first time, we demonstrate supracluster assembly based on an  $Al_4$  precursor through both hydrogen

<sup>a</sup>State Key Laboratory of Structural Chemistry, Fujian Institute of Research on the Structure of Matter, Chinese Academy of Sciences, 350002 Fuzhou, P.R. China. E-mail: hyp041@163.com, fwh@fjirsm.ac.cn

<sup>b</sup>University of Chinese Academy of Sciences, Beijing 100049, China

<sup>c</sup>MOE Key Laboratory of Cluster Science, School of Chemistry and Chemical Engineering, Beijing Institute of Technology, Beijing 100081, China

† Electronic supplementary information (ESI) available. CCDC 2303391–2303400. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4qi01103a>

‡ These authors contributed equally to this work.

bonds and coordination bonds. Based on our recent efforts and progress in aluminum oxo clusters (AIOCs),<sup>41–47</sup> we believe that  $(\text{Me}_2\text{NH}_2)_4[\text{Al}_4(\text{L})_4(\text{Cat})_2]$  (AIOC-13, a cluster denoted as  $\text{Al}_4$  hereinafter)<sup>42</sup> is a suitable precursor for further study of supracluster assembly of target materials with photo-related properties (Scheme 1) where (1) it fulfills the requirements of high solubility/stability and accessibility for scale-up synthesis; and (2) it provides one successful example of incorporating lanthanides<sup>42</sup> proving the possibility of assembly and that the abundance of interaction sites (aromatic ring, hydroxyl group and bridging oxo) will bring more interesting modes of assembly. The presence of aromatic rings and an oxo moiety may lead to weak interaction with guest molecules. The suspended oxo group may result in coordination with oxyphilic metal ions. Such assembly is feasible from the point of view of charge matching. We selected a series of chelating ligands of different lengths, and the distance between the two chelating sites plays a key role in assembling the compound type to form ion pairs and coordination-modified compounds, respectively. With subsequent assembly, the photo-related properties will also be tuned. Accordingly, we probe an avenue toward the enhancement of intended optical properties like photoluminescence and third-order nonlinear optical (NLO) effects. For example, the employment of chiral ligands and heavy metal ions enhances third-order NLO performance.<sup>48,49</sup>

## Results and discussion

### Synthesis

The facile synthesis of large amounts of AIOC-13 as the source of the  $\text{Al}_4$  clusters is described in detail in the Experimental section (Scheme S1 and Tables S1–S3†). It is a monodisperse anionic cluster with  $(\text{Me}_2\text{NH}_2)^+$  counteranions, which can be soluble and stable in water and common solvents. There are eight uncoordinated carboxyl oxygen atoms and six phenyl rings in each  $\text{Al}_4$  cluster, making it an excellent precursor for assembly with cationic units. The reasons we chose the N,N-

chelated (phenanthroline-type) and P,P-chelated (phenylphosphine-type) ligands are that they meet the requirements of both ligand chelate and photo-functional conjugated groups (Fig. 1). By investigating the supracluster assembly behaviors of  $\text{Al}_4$  clusters with metal ions (such as  $\text{Ag}^+$ ,  $\text{Fe}^{2+}$  and  $\text{Cu}^{2+}$ ), we successfully obtained a variety of novel ionic pair compounds *via* short chelating ligands (hydrogen bonds and  $\pi\cdots\pi$  interactions) (AIOC-168 to AIOC-172) and coordination-modified compounds through relatively long chelating ligands (AIOC-173 to AIOC-176) (Table 1). Rapid crystal growth of these materials can be achieved by solvent-thermal synthesis in one day.

### Supracluster assembly of structures *via* non-covalent bonds

In the beginning,  $\text{AgClO}_4$  and the 1,3-bdpp ligand were added to a DMF/EtOH/ $\text{H}_2\text{O}$  solution of the  $\text{Al}_4$  precursor, which was kept at 80 °C for one day, yielding faint yellow block crystals of AIOC-168. In the structure of AIOC-168, the Ag(I) center is P,P-chelated by two 1,3-bdpp ligands, giving rise to an  $[\text{Ag}(1,3\text{-bdpp})_2]^+$  ion (Fig. S1†). The supracluster assembly of the

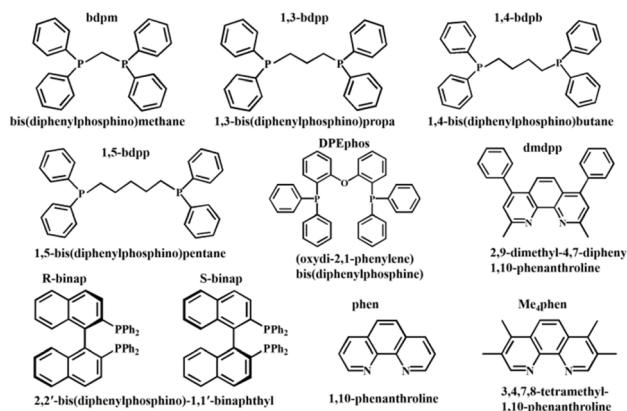
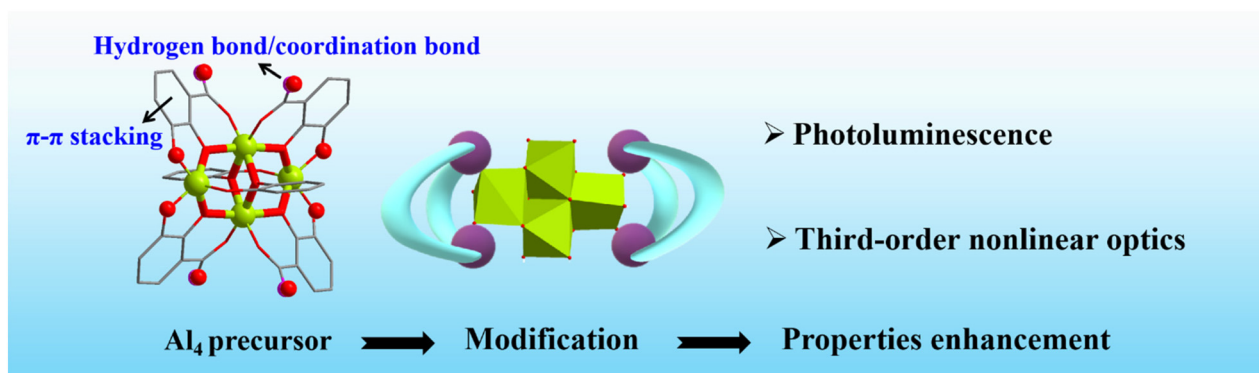


Fig. 1 The N,N-chelated or P,P-chelated ligands used in the supracluster assembly based on  $\text{Al}_4$  clusters.



**Scheme 1** Illustration of the selection of the  $\text{Al}_4$  precursor and its supracluster assembly towards enhancement of photo-related properties. The advantages of using the  $\text{Al}_4$  precursor include its high solubility/stability and accessibility for scale-up synthesis, as well as abundant binding sites. The supracluster assembly method includes non-covalent interaction assembly and coordination bond interaction assembly. The dark purple balls represent metal centers like  $\text{Ag}^+$ ,  $\text{Fe}^{2+}$  and  $\text{Cu}^{2+}$ , while the cyan loops represent flexible ligands.

**Table 1** Summary of the compounds resulting from the supracluster assembly of the Al<sub>4</sub> precursor

Complex	Composition	Space group	Structure dimensionality	R <sub>1</sub> value	CCDC
<b>Supracluster assembly of structures via non-covalent bonds</b>					
<b>AIOC-168</b>	(Me <sub>2</sub> NH <sub>2</sub> ) <sub>3</sub> [Ag(1,3-bdpp) <sub>2</sub> ][Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>P</i> $\bar{1}$	0D	0.053	2303391
<b>AIOC-169</b>	(Me <sub>2</sub> NH <sub>2</sub> ) <sub>3</sub> [Ag(dmdpp)(DPEphos)][Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>Pbca</i>	0D	0.138	2303392
<b>AIOC-170(R)</b>	(Me <sub>2</sub> NH <sub>2</sub> ) <sub>3</sub> [Ag( <i>R</i> -binap) <sub>2</sub> ] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ] <sub>2</sub>	<i>P2</i> <sub>1</sub>	0D	0.077	2303393
<b>AIOC-170(S)</b>	(Me <sub>2</sub> NH <sub>2</sub> ) <sub>3</sub> [Ag( <i>S</i> -binap) <sub>2</sub> ] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ] <sub>2</sub>	<i>P2</i> <sub>1</sub>	0D	0.053	2303394
<b>AIOC-171</b>	[Fe(phen) <sub>3</sub> ] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>I2/a</i>	0D	0.041	2303395
<b>AIOC-172</b>	[Cu(Me <sub>4</sub> phen) <sub>2</sub> (H <sub>2</sub> O)] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>P</i> $\bar{1}$	0D	0.051	2303396
<b>Supracluster assembly of structures via coordination bonds</b>					
<b>AIOC-173</b>	[Ag <sub>2</sub> (bdpm) <sub>2</sub> ] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>P</i> $\bar{1}$	0D	0.053	2303397
<b>AIOC-174</b>	[Ag <sub>3</sub> (1,5-bdpp) <sub>3</sub> ] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ](ClO <sub>4</sub> ) <sub>2</sub>	<i>P</i> $\bar{1}$	0D	0.043	2303398
<b>AIOC-175</b>	[Ag <sub>4</sub> (1,4-bdpp) <sub>3</sub> ] <sub>3</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>I2/a</i>	1D	0.036	2303399
<b>AIOC-176</b>	[Ag <sub>4</sub> (bdpm) <sub>2</sub> (1,4-bdpp)] <sub>2</sub> [Al <sub>4</sub> (L) <sub>4</sub> (Cat) <sub>2</sub> ]	<i>P</i> $\bar{1}$	2D	0.042	2303400

H<sub>3</sub>L = 2,3-dihydroxybenzoic acid; cat = catechol; bdpm = bis(diphenylphosphino)methane; 1,3-bdpp = 1,3-bis(diphenylphosphino)propane; 1,4-bdpp = 1,4-bis(diphenylphosphino)butane; 1,5-bdpp = 1,5-bis(diphenylphosphino)pentane; DPEphos = (oxydi-2,1-phenylene)bis(diphenylphosphine); *R*-binap = (*R*)-(+)-2,2'-bis(diphenylphosphino)-1,1'-binaphthyl; *S*-binap = (*S*)-(-)-2,2'-bis(diphenylphosphino)-1,1'-binaphthyl; phen = 1,10-phenanthroline; dmdpp = 2,9-dimethyl-4,7-diphenyl-1,10-phenanthroline; Me<sub>4</sub>phen = 3,4,7,8-tetramethyl-1,10-phenanthroline.

anionic Al<sub>4</sub> cluster and cationic [Ag(1,3-bdpp)<sub>2</sub>]<sup>+</sup> unit forms an interesting ion-pair structure (Fig. 2a). However, in the above synthetic system of **AIOC-168**, we replaced the 1,3-bdpp ligand with DPEphos and added the dmdpp phenanthroline-type ligand, and we readily obtained another ion-pair structure

(**AIOC-169**) composed of an Al<sub>4</sub> cluster and [Ag(dmdpp)(DPEphos)]<sup>+</sup> unit (Fig. S4<sup>†</sup>), in which the Ag(I) center is chelated by one DPEphos ligand and one dmdpp ligand. When we replaced the organic ligand with a chiral binap ligand, we successfully synthesized a pair of pure enantiomers (**AIOC-170(R)**)



**Fig. 2** A family of structures from supracluster assembly based on pre-designed Al<sub>4</sub> clusters including (a) supracluster assembly of structures via non-covalent bonds and (b) supracluster assembly of structures via coordination bonds. Atom color code: green, Al; orange, Ag; pink, P; red, O; blue, N; gray, C; off-white, H.

and **AIOC-170(S)** (Fig. S7 and S10†). In their structures, two *R*- or *S*-binap ligands coordinate to one Ag(I) center, producing a chiral  $[\text{Ag}(\text{R-binap})_2]^+$  or  $[\text{Ag}(\text{S-binap})_2]^+$  unit, which further forms chiral ion-pair structures with the  $\text{Al}_4$  cluster. Their absolute configuration and enantiomeric nature have been confirmed by circular dichroism (CD) spectra, whose CD peaks exhibit obvious mirror images (Fig. S61†). Considering the chiral and fluorescent properties of **AIOC-170R/S**,<sup>50,51</sup> we tested their circularly polarized luminescence, and the results were not as expected. In the 3D packing structures of **AIOC-168** to **AIOC-170**, there are multiple supramolecular interactions, including  $\pi\cdots\pi$  stacking and C–H $\cdots\pi$ , C–H $\cdots\text{O}$ , O–H $\cdots\text{O}$ , and N–H $\cdots\text{O}$  interactions (Fig. S3, S6 and S9†). It is particularly necessary to point out that the Ag units form a cationic layer, while the  $\text{Al}_4$  clusters form an anionic layer (Fig. S2, S5 and S8†), and they take the ABAB alternating stacking mode, which is similar to that of hydrocalcite. Furthermore, the  $(\text{Me}_2\text{NH}_2)^+$  counteranions are located in the anionic  $\text{Al}_4$  cluster layer. By replacing the above  $\text{AgClO}_4$  with  $\text{FeSO}_4\cdot 7\text{H}_2\text{O}$  and  $\text{Cu}_2\text{O}$  and adding a small amount of ammonium hydroxide, we synthesized ion-pair structures **AIOC-171** and **AIOC-172** in the presence of phen and  $\text{Me}_4\text{phen}$  ligands, respectively. The Fe(II) and Cu(II) centers are N,N-chelated by two phen and  $\text{Me}_4\text{phen}$  ligands (Fig. S11 and S13†), respectively, and there is also an  $\text{H}_2\text{O}$  molecule that coordinates to the Cu(II) center. Through weak interactions, the  $\text{Al}_4$  clusters and the  $[\text{Fe}(\text{phen})_3]^{2+}$  or  $[\text{Cu}(\text{Me}_4\text{phen})_2(\text{H}_2\text{O})]^{2+}$  cations stack alternately into a 3D dense structure (Fig. S12 and S14†).

### Supracluster assembly of structures *via* coordination bonds

Encouraged by the above results, more assembly experiments were explored (Fig. 2b). When increasing the dosage of  $\text{Ag}^+$  ions and using the shorter bdpm ligand, we synthesized **AIOC-173**. In the structure of **AIOC-173** (Fig. S15 and S16†), two Ag(I) centers are connected by two bdpm ligands, *in situ* forming a binuclear  $[\text{Ag}_2(\text{bdpm})_2]^{2+}$  cycle (named the  $\text{Ag}_2$  cycle). Interestingly, each  $\text{Al}_4$  cluster links two such  $\text{Ag}_2$  cycles *via* the Ag–O coordination bond on both sides of the cluster, in which the carboxyl O atoms of the  $\text{Al}_4$  cluster coordinate to the Ag(I) atom of the  $\text{Ag}_2$  cycle. The Ag $\cdots$ Ag length is about 2.9 Å, and the Ag–O bond lengths are about 2.5 Å, which are within the normal bond-length range. All the Ag(I) centers are three-coordinated. As we continued to increase the dosage of  $\text{Ag}^+$  ions and used a longer 1,5-bdpp ligand, we obtained **AIOC-174**. Structural analyses show that each  $\text{Al}_4$  cluster in **AIOC-174** traps two *in situ* formed  $[\text{Ag}_3(1,5\text{-bdpp})_3]^{3+}$  cycles (each named as an  $\text{Ag}_3$  cycle). As shown in Fig. S17,† the Ag1 atom is four-coordinated, while Ag2 and Ag3 atoms are three-coordinated. The  $(\text{ClO}_4)^-$  counteranions are present in the structure of **AIOC-174** (Fig. S18†). Herein, through coordination interactions, we modified the  $\text{Ag}_2$  and  $\text{Ag}_3$  cycles onto the  $\text{Al}_4$  cluster and obtained two functional ligand-modified heterometallic clusters.

From the above results, it can be seen that the length of the organic ligand also affected the supracluster assembly behavior of the  $\text{Al}_4$  cluster, so we continued to change the type of

ligand and tried using mixed organic ligands. When the 1,4-bdpp ligand was used to react with  $\text{Ag}^+$  ions and  $\text{Al}_4$  clusters, **AIOC-175** was synthesized. In **AIOC-175**, three 1,4-bdpp ligands link with four Ag(I) centers to generate a tetranuclear  $[\text{Ag}_4(1,4\text{-bdpp})_3]^{4+}$  unit (named the  $\text{Ag}_4$  unit) (Fig. S19†). Such an *in situ* generated  $\text{Ag}_4$  unit further connects two adjacent  $\text{Al}_4$  clusters through Ag–O coordination bonds, giving rise to an infinite chain structure. The chains are stacked in ABAB mode to form a dense supramolecular structure (Fig. S20†), and there are rich interactions between adjacent chains in **AIOC-175**, including  $\pi\cdots\pi$  stacking, C–H $\cdots\pi$  and C–H $\cdots\text{O}$  interactions. However, when longer 1,4-bdpp and shorter bdpm ligands were added to the reaction system simultaneously, we obtained compound **AIOC-176**. Structural analysis shows that **AIOC-176** has a layered structure of  $\text{Al}_4$  clusters linked together by  $[\text{Ag}_2(\text{bdpm})_2]^{2+}$  cycles (named the  $\text{Ag}_2$  cycle) and  $[\text{Ag}_2(1,4\text{-bdpp})]^{2+}$  units (named the  $\text{Ag}_2$  unit) (Fig. S21†). In the structure of **AIOC-173**, the  $\text{Ag}_2$  cycle is only modified at both ends of the cluster, while it acts as a linker in **AIOC-176**, collaborating with the  $\text{Ag}_2$  unit to further connect with and form the  $\text{Al}_4$  clusters into a layer (Fig. S22†).

Herein, we demonstrated an extraordinary supracluster assembly including both non-covalent and coordination bonds. These assembled compounds are challenging to synthesize in one pot, so through supracluster assembly, we can indeed obtain new materials that cannot be obtained by direct synthesis. In theory, all kinds of metal cation units can form ion-pair structures with the anionic  $\text{Al}_4$  cluster. Similarly, coordination bonds can also modify different metal cation units at the face of the cluster. On the other hand, the  $\text{Al}_4$  cluster can also be organized into different dimensional structures, including chain and layer structures, as well as 3D frameworks. Through functional modification, we can obtain the desired applications.

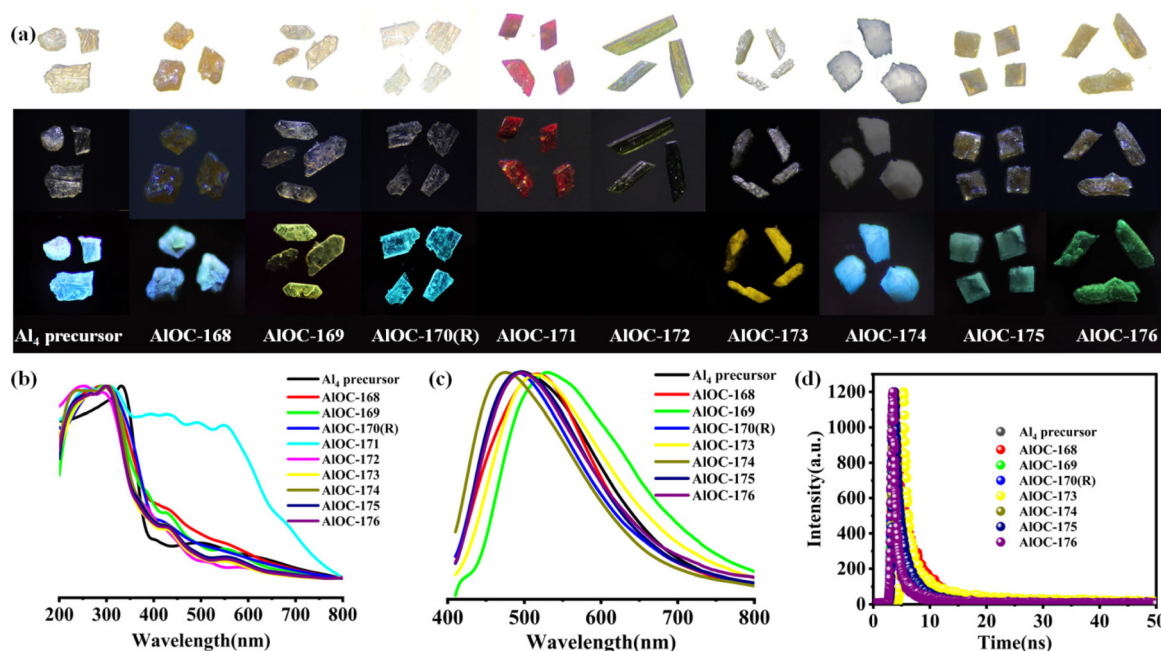
The powder X-ray diffraction (XRD) patterns of **AIOC-168** to **AIOC-176** are shown in Fig. S23–S31.† To ensure the high purity of all structures, we performed Le Bail fitting. The low  $R_{\text{wp}}$  and  $R_p$  values (<10%) in the Le Bail fittings show that these compounds have high phase purity, indicating that the self-assembly of  $\text{Al}_4$  clusters with metal cation units into these cluster-based compounds under certain conditions is quite stable and directed. The thermogravimetric analysis (TGA) curves of these materials were recorded, as shown in Fig. S32–S40 in the ESI.† Their TG curves do not have a relatively apparent weightless platform before 300 °C, which is consistent with their dense supramolecular packing structures. The solvents or counterions ( $(\text{Me}_2\text{NH}_2)^+$  or  $(\text{ClO}_4)^-$ ) were gradually lost with increasing temperature, followed by decomposition of their structures. The residues after decomposition might be  $\text{Ag}_2\text{O}$  and  $\text{CuO}$  or  $\text{Fe}_3\text{O}_4$ . Diffuse reflectance spectroscopy was used to study the UV/vis absorption of these compounds. According to the Kubelka–Munk function, yellowish  $\text{Ag}$ -cluster- and green  $\text{Cu}$ -cluster-based compounds exhibit relatively low bandgaps (2.6–3.1 eV) compared to that of the  $\text{Al}_4$  cluster raw material (3.2 eV), but that of the deep red **AIOC-171**, containing the  $[\text{Fe}(\text{phen})_3]^{2+}$  unit, is about 1.56 eV,

which is significantly lower than that of other compounds (Fig. 3a, b and S42–S50<sup>†</sup>). Their homogeneous elemental distribution was confirmed by their spectra from energy dispersive spectroscopy (EDS) as shown in Fig. S51–S58.<sup>†</sup>

### Studies of photoluminescence properties

Because the phenanthroline- and phenylphosphine-type ligands possess strong absorbing chromophores, and the Ag-based compounds themselves have good photoluminescent properties, we decided to study the solid-state excitation and emission spectra of compounds **AIOC-168** to **AIOC-176** at room temperature (Fig. 3c and d). For convenience, their detailed parameters are also summarized in Table 2. As shown in Fig. S59,<sup>†</sup> upon excitation at 400 nm, the  $\text{Al}_4$  precursor raw material displays an emission band at 400–700 nm and the

highest emission peak at 487 nm, which may belong to the  $\pi \rightarrow \pi^*$  transition of organic ligands. Under the same excitation conditions, these cluster-assembled compounds **AIOC-168** to **AIOC-170** and **AIOC-173** to **AIOC-176** show similar emission peaks, which can be observed at 500, 520, 489, 533, 477, 505 and 510 nm, respectively. However, **AIOC-171** and **AIOC-172** have no obvious photoluminescence under ultraviolet light (360 nm). Compound **AIOC-174** exhibits a blue-shift (*ca.* 10 nm) in comparison with the  $\text{Al}_4$  cluster, and it shows blue light. However, **AIOC-173** and **AIOC-176** display distinct red-shifts of *ca.* 20–50 nm, and their luminescent colours are viridescent and yellow, respectively (Fig. S60<sup>†</sup>). In these case, the ligand-based luminescence is dominant and the blue-shifted or red-shifted phenomenon may be attributed to metal–ligand/cluster coordinative interactions. Additionally, the rigidity of



**Fig. 3** The fluorescence properties of the pristine  $\text{Al}_4$  precursor and the compounds from supracluster assembly. (a) Pictures of  $\text{Al}_4$  precursor to **AIOC-176** crystals. The three images in each column from top to bottom show crystals in natural light, crystals in the dark without fluorescence irradiation, and crystals in the dark with fluorescence irradiation. (b) The UV/vis wavelengths of the  $\text{Al}_4$  precursor to **AIOC-176**. (c) The emission wavelengths of the  $\text{Al}_4$  precursor to **AIOC-176** except for those of **AIOC-171** and **AIOC-172**. (d) The transient fluorescence lifetime curves of the  $\text{Al}_4$  precursor to **AIOC-176** except for those of **AIOC-171** and **AIOC-172**.

**Table 2** The crucial photophysical parameters of the  $\text{Al}_4$  precursor and compounds from supracluster assembly

Compound	UV/vis wavelength (nm)	Excitation wavelength (nm)	Emission wavelength (nm)	Fluorescence lifetime (ns)	$\beta$ (cm $\text{GW}^{-1}$ )
$\text{Al}_4$ precursor	332	395	495	2.45	98
<b>AIOC-168</b>	288	395	515	3.24	200
<b>AIOC-169</b>	292	395	525	4.33	120
<b>AIOC-170(R)</b>	302	395	490	4.01	220
<b>AIOC-171</b>	306	395	455	0	130
<b>AIOC-172</b>	254	395	455	0	700
<b>AIOC-173</b>	301	395	515	4.25	750
<b>AIOC-174</b>	306	395	475	2.07	730
<b>AIOC-175</b>	300	395	495	2.02	710
<b>AIOC-176</b>	299	395	500	2.07	700

the ligands in the structure may also account for the fluorescence differences. The rigidity of the ligands in **AIOC-173** and **AIOC-169** is greater compared to that in other structures, and hence their fluorescence lifetimes are somewhat superior. In a word, the luminescent colour changes can be realized by different supracluster assembly behaviours.

### Studies of third-order nonlinear optical (NLO) properties

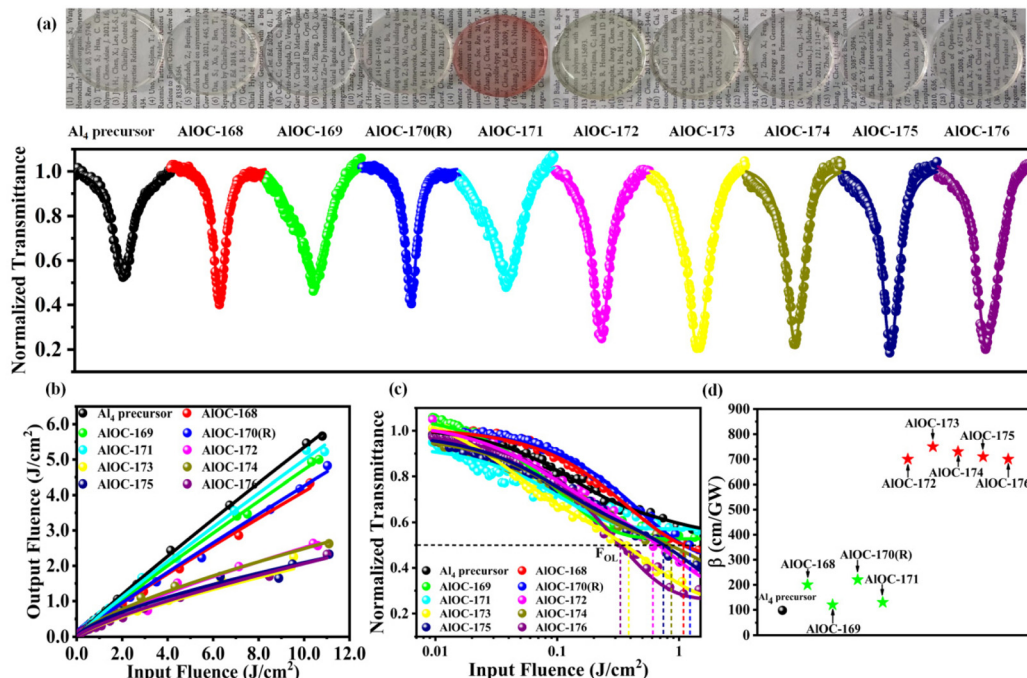
Third-order nonlinear optical (NLO) materials are of particular importance for various applications including the optical limiting effect. It is reported that effects of larger conjugated systems and heavy metals have been proven to have positive influences on third-order nonlinear optical and optical limiting behavior.<sup>49,52–54</sup> Considering the presence of heavy metals and large conjugated chromophores, we then decided to study their third-order NLO properties. In order to ensure the stability of compounds **AIOC-168** to **AIOC-176**, the crystals of these materials are effectively dispersed into polydimethylsiloxanes (PDMSs) and then made into a thin film (named **AIOCs@PDMS** film), respectively. (The Experimental section provides detailed experimental information.) For better comparison, we also tested the third-order NLO properties of the starting  $\text{Al}_4$  precursor by the same film-making method ( $\text{Al}_4$  precursor@PDMS). The third-order NLO properties of these films were studied using a typical open-aperture Z-scan system with a nanosecond laser at 532 nm.

Precise control of the intramolecular coordination bonds and intermolecular non-covalent forces of clusters tunes

charge transfer and band gap enhancement and ultimately improves the third-order nonlinear optical performance. Under an input laser pulse energy of 80  $\mu\text{J}$ ,  $\text{Al}_4$  precursor@PDMS to **AIOCs@PDMS** display obvious reverse-saturated absorption (RSA) behaviors and exhibit a sequentially enhanced optical limiting response from **AIOC-168** to **AIOC-176** (Fig. 4a and Table S4†). In order to quantitatively evaluate the NLO performance of the sample, the nonlinear absorption coefficient ( $\beta$ ) (eqn (1)) of the sample was obtained from the best-fit curve:

$$T(z, s = 1) = \frac{1}{\sqrt{\pi}q_0(z, 0)} \int_{-\infty}^{\infty} \ln[1 + q_0(z, 0)e^{-r^2}] dr \quad (1)$$

Here,  $q_0(z, 0) = \beta I_0 L_{\text{eff}}$ , where  $I_0$  is the on-axis peak intensity at the focus ( $z = 0$ ),  $L_{\text{eff}} = [1 - \exp(-\alpha l)]/\alpha$  denotes the effective thickness of the sample,  $\alpha$  is the linear absorption coefficient, and  $l$  indicates the sample thickness. The values of  $\beta$  for  $\text{Al}_4$  precursor@PDMS and **AIOCs@PDMS** composite films are 98, 200, 120, 220, 130, 700, 750, 730, 710, and 700  $\text{cm GW}^{-1}$ , respectively (Fig. 4d). Compared with the  $\text{Al}_4$  precursor, the introduced large conjugated ligands facilitate charge transfer between anion and cation groups, thus improving the third-order NLO performance. In the non-covalently assembled compounds (**AIOC-168** to **AIOC-172**), **AIOC-172** has relatively good nonlinear absorption due to its low bandgap and heavy metal effect. As for supracluster assembly *via* coordination bonds, compounds **AIOC-173** to **AIOC-176** exhibit better optical limiting responses, which is attributed to the coordination inter-



**Fig. 4** The third-order NLO of the pristine  $\text{Al}_4$  precursor and compounds from supracluster assembly. (a) The opening Z-scan curve at 532 nm of the samples. At the top of the Z-scan curves are photographs of the  $\text{Al}_4$  precursor@PDMS and **AIOCs@PDMS** before (upper panel) and after (lower panel) applying 360 nm UV light. (b) Curves of output fluence versus input fluence for the  $\text{Al}_4$  precursor, and **AIOC-168** to **AIOC-176**. (c) Plots of normalized transmittance versus input fluence for the  $\text{Al}_4$  precursor, and **AIOC-168** to **AIOC-176**. (d) Comparison of the nonlinear absorption coefficients at a laser intensity of 80  $\mu\text{J}$  for  $\text{Al}_4$  precursor@PDMS and **AIOCs@PDMS**.

action significantly enhancing charge transfer between anion and cation groups, thus improving the third-order NLO performance. The supercluster assembly strategy may serve as a reference for future performance enhancement. It is a direction worth exploring further. It is comparable to that of rare-earth metal or transition metal cluster-based materials,<sup>55,56</sup> although they are not comparable with metal-porphyrinic frameworks or porphyrin covalent organic frameworks (Table S5†).<sup>57,58</sup>

The curves of output fluence *versus* input fluence of such samples show that the output fluence linearly increases at low-incident fluence. At high-incident fluence, the output fluence deviates from linearity and shows a typical OL response (Fig. 4b). Their optical limiting threshold values ( $F_{OL}$ , defined as the input fluence at 50% linear transmittance) are 5.12 J cm<sup>-2</sup> ( $F_{OL}$  Al<sub>4</sub> precursor), 1.09 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-168), 3.45 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-169), 1.23 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-170(R)), 3.59 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-171), 0.61 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-172), 0.39 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-173), 0.86 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-174), 0.75 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-175), and 0.33 J cm<sup>-2</sup> ( $F_{OL}$  AIOC-176), respectively (Fig. 4c). In addition, we have carried out cycling experiments to verify the laser stability of AIOCs@PDMS materials. The results show that the RSA profiles of all third-order NLO materials have the same reproducibility after 4 or 8 cycles (Fig. S63–S72†).

## Conclusions

In summary, by employing the Al<sub>4</sub> cluster as a precursor, a supracluster assembly strategy (including both non-covalent interaction and coordination bonds) was applied to construct a series of interesting cluster-based architectures. By introducing large  $\pi$ -conjugated phenanthroline- or phenylphosphine-type organic ligands, these cluster-assembled compounds showed different photoluminescent colors. The results showed that supracluster assembly *via* coordination bonds facilitates charge transfer and can effectively enhance the third-order NLO properties of cluster-based materials compared with those based on non-covalent bonds. This work not only demonstrates a general synthesis strategy based on a pre-designed precursor but also provides important guidance for stepwise tuning and optimization toward specific properties.

## Experimental section

### Synthesis of the Al<sub>4</sub> precursor

The synthesis of (Me<sub>2</sub>NH<sub>2</sub>)<sub>4</sub>[Al<sub>4</sub>(L)<sub>4</sub>(Cat)<sub>2</sub>] (AIOC-13) is based on our previous work.<sup>41</sup> A mixture of Al(O<sup>i</sup>Pr)<sub>3</sub> (0.2 g, 0.98 mmol), 2,3-dihydroxybenzoic acid (H<sub>3</sub>L, 0.4 g, 2.60 mmol), pyrazole (2 g, 29.38 mmol), and DMF (5 mL) was sealed in a 25 mL Teflon reactor and transferred to a preheated oven at 160 °C for 3 days. When cooled to room temperature, light brown crystals of AIOC-13 were obtained (yield: 75% based on Al(O<sup>i</sup>Pr)<sub>3</sub>). Amplification synthesis is also possible:

using a 25 mL Teflon reactor, 0.27 g of AIOC-13 can be obtained, and using a 50 mL Teflon reactor, 0.68 g of AIOC-13 can be obtained.

### Supracluster assembly based on the Al<sub>4</sub> precursor

By investigating the supracluster assembly behavior of the Al<sub>4</sub> cluster with metal ions (such as Ag<sup>+</sup>, Fe<sup>2+</sup> and Cu<sup>2+</sup>) in the presence of chelated ligands (like N,N-chelated (phenanthroline-type) or P,P-chelated (phenylphosphine-type) ligands) with diverse binding distances and angles, we successfully obtained a variety of cluster-based solids. A typical solvothermal synthesis of AIOC-168 (Me<sub>2</sub>NH<sub>2</sub>)<sub>3</sub>[Ag(1,3-bdpp)<sub>2</sub>][Al<sub>4</sub>(L)<sub>4</sub>(Cat)<sub>2</sub>]·guests is described as follows: the Al<sub>4</sub> precursor (55 mg, 0.05 mmol) was dissolved in 2 mL of H<sub>2</sub>O at 100 °C for 6 h. A mixture of 2 mL of precursor, silver perchlorate (AgClO<sub>4</sub>, 14 mg, 0.07 mmol), 1,3-bis(diphenylphosphino)propane (1,3-bdpp, 45 mg, 0.11 mmol), EtOH (2 mL), and DMF (2 mL) was sealed in a 20 mL vial at 80 °C for 24 h. When cooled to room temperature, light brown crystals were obtained (yield: 75% based on the Al<sub>4</sub> precursor). The crystals were rinsed with EtOH and preserved in a sealed and dry environment. The synthetic procedure for other compounds is similar to that of AIOC-168 but with different ligands and solvents. For the detailed synthesis, please refer to the ESI†.

### Fabrication of Al<sub>4</sub> precursor@PDMS and AIOCs@PDMS

Al<sub>4</sub> precursor@PDMS and AIOCs@PDMS were prepared using Sylgard 184 (Dow Corning) by thoroughly mixing 10 parts base with 1 part curing agent. The coordinate crystals were mixed with the PDMS solution to form coordinate-dispersed PDMS suspension, and then the mixture suspension was added to a template and the template was placed in a vacuum oven at 60 °C for 6 hours. Finally, transparent and flexible Al<sub>4</sub> precursor@PDMS and AIOCs@PDMS were obtained.

### Z-Scan measurements

The third-order NLO properties of the sample were evaluated by using the Z-scan technique. The excitation light source was an Nd:YAG laser with a repetition rate of 5 Hz. The laser pulse (period, 5 ns; wavelength, 532 nm) was split into two beams with a mirror. The pulse energies at the front and back of the samples were monitored using energy detectors 1 and 2. All of the measurements were conducted at room temperature. The sample was mounted on a computer-controlled translation stage that shifted each sample along the z-axis.

## Author contributions

All authors contributed extensively to the work presented in this paper. J. Zhang and W.-H. Fang conceived the research project. F. Yang and R.-Y. Chen performed the synthesis, and characterization and optical property studies. T.-S. Wang and Y.-P. He helped with the crystallographic data refinement. W.-H. Fang and Y.-P. He wrote the manuscript and ESI† with input from the other authors.

## Data availability

X-ray crystallographic data have been deposited at the Cambridge Crystallographic Data Centre (<https://www.ccdc.cam.ac.uk/>) with CCDC reference numbers 2303391 to 2303400† (single-crystal AIOC-168 to AIOC-176).

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The research reported in this publication was supported by the National Natural Science Foundation of China (U23A20950, 22371278, and 92261108), the Science Fund for Distinguished Young Scholars of Fujian Province (2021J06035), the Funding of the Fujian Provincial Chemistry Discipline Alliance (50025401), and the Youth Innovation Promotion Association of the Chinese Academy of Sciences (Y2021081).

## References

- J. L. Segura, S. Royuela and M. Mar Ramos, Post-synthetic modification of covalent organic frameworks, *Chem. Soc. Rev.*, 2019, **48**, 3903–3945.
- H. Wang, Y. Jin, N. Sun, W. Zhang and J. Jiang, Well-Defined, Versatile and Recyclable Half-Sandwich Nickelacarborane Catalyst for Selective Carbene-Transfer Reactions, *Chem. Soc. Rev.*, 2021, **50**, 8874–8886.
- S. Mandal, S. Natarajan, P. Mani and A. Pankajakshan, Post-Synthetic Modification of Metal-Organic Frameworks Toward Applications, *Adv. Funct. Mater.*, 2021, **31**, 2006291.
- T. Chen and D. Zhao, Post-synthetic modification of metal-organic framework-based membranes for enhanced molecular separations, *Coord. Chem. Rev.*, 2023, **491**, 215259.
- H. Ding, A. Mal and C. Wang, Tailored covalent organic frameworks by post-synthetic modification, *Mater. Chem. Front.*, 2020, **4**, 113–127.
- J. Liu, Z. Wang, P. Cheng, M. J. Zaworotko, Y. Chen and Z. Zhang, Post-synthetic modifications of metal-organic cages, *Nat. Rev. Chem.*, 2022, **6**, 339–356.
- P. Waeles, M. Gauthier and F. Coutrot, Challenges and Opportunities in the Post-Synthetic Modification of Interlocked Molecules, *Angew. Chem., Int. Ed.*, 2021, **60**, 16778–16799.
- F. A. El-malek and A. Steinbuchel, Post-synthetic Enzymatic and Chemical Modifications for Novel Sustainable Polyesters, *Front. Bioeng. Biotechnol.*, 2022, **9**, 817023.
- E. M. El-Sayed, Y.-D. Yuan, D. Zhao and D. Yuan, Zirconium Metal-Organic Cages: Synthesis and Applications, *Acc. Chem. Res.*, 2022, **55**, 1546–1560.
- Y.-P. He, L.-B. Yuan, J.-S. Song, G.-H. Chen, Q. Lin, C. Li, L. Zhang and J. Zhang, Optical Resolution of the Water-Soluble  $Ti_4(\text{embonate})_6$  Cages for Enantioselective Recognition of Chiral Drugs, *Chem. Mater.*, 2018, **30**, 7769–7775.
- G.-H. Chen, H.-Z. Li, Y.-P. He, S.-H. Zhang, X. Yi, F.-P. Liang, L. Zhang and J. Zhang,  $Ti_4(\text{embonate})_6$  Based Cage-Cluster Construction in a Stable Metal–Organic Framework for Gas Sorption and Separation, *Cryst. Growth Des.*, 2019, **20**, 29–32.
- Y.-P. He, L.-B. Yuan, G.-H. Chen, Q.-P. Lin, F. Wang, L. Zhang and J. Zhang, Water-Soluble and Ultrastable  $Ti_4L_6$  Tetrahedron with Coordination Assembly Function, *J. Am. Chem. Soc.*, 2017, **139**, 16845–16851.
- R.-Y. Chen, G.-H. Chen, Y.-P. He and J. Zhang, Coordination Assembly of Tetrahedral  $Ti_4(\text{embonate})_6$  Cages with Alkaline-Earth Metal Ions, *Chin. J. Struct. Chem.*, 2022, **41**, 2201001–2201006.
- G.-H. Chen, Y.-P. He, Y. Yu, H. Lv, S. Li, F. Wang, Z.-G. Gu and J. Zhang, Post-Assembly Modification of Homochiral Titanium–Organic Cages for Recognition and Separation of Molecular Isomers, *Angew. Chem., Int. Ed.*, 2023, **62**, e202300726.
- A. J. Gosselin, A. M. Antonio, K. J. Korman, M. M. Deegan, G. P. A. Yap and E. D. Bloch, Elaboration of Porous Salts, *J. Am. Chem. Soc.*, 2021, **143**, 14956–14961.
- E. J. Gosselin, G. E. Decker, A. M. Antonio, G. R. Lorzing, G. P. A. Yap and E. D. Bloch, A Charged Coordination Cage-Based Porous Salt, *J. Am. Chem. Soc.*, 2020, **142**, 9594–9598.
- G. Liu, Y. Di Yuan, J. Wang, Y. Cheng, S. B. Peh, Y. Wang, Y. Qian, J. Dong, D. Yuan and D. Zhao, Process-Tracing Study on the Postassembly Modification of Highly Stable Zirconium Metal–Organic Cages, *J. Am. Chem. Soc.*, 2018, **140**, 6231–6234.
- G. Liu, X. Zhang, Y. Di Yuan, H. Yuan, N. Li, Y. Ying, S. B. Peh, Y. Wang, Y. Cheng, Y. Cai, Z. Gu, H. Cai and D. Zhao, Thin-Film Nanocomposite Membranes Containing Water-Stable Zirconium Metal–Organic Cages for Desalination, *ACS Mater. Lett.*, 2021, **3**, 268–274.
- J. Liu, W. Duan, J. Song, X. Guo, Z. Wang, X. Shi, J. Liang, J. Wang, P. Cheng, Y. Chen, M. J. Zaworotko and Z. Zhang, Self-Healing Hyper-Cross-Linked Metal–Organic Polyhedra (HCMOPs) Membranes with Antimicrobial Activity and Highly Selective Separation Properties, *J. Am. Chem. Soc.*, 2019, **141**, 12064–12070.
- D. Nam, J. Huh, J. Lee, J. H. Kwak, H. Y. Jeong, K. Choi and W. Choe, Cross-linking Zr-based metal-organic polyhedra via postsynthetic polymerization, *Chem. Sci.*, 2017, **8**, 7765–7771.
- B. S. Pilgrim, D. A. Roberts, T. G. Lohr, T. K. Ronson and J. R. Nitschke, Signal transduction in a covalent post-assembly modification cascade, *Nat. Chem.*, 2017, **9**, 1276–1281.
- D. A. Roberts, B. S. Pilgrim, J. D. Cooper, T. K. Ronson, S. Zarra and J. R. Nitschke, Post-assembly Modification of Tetrazine-Edged  $Fe(II)_4L_6$  Tetrahedra, *J. Am. Chem. Soc.*, 2015, **137**, 10068–10071.

- 23 T. K. Ronson, B. S. Pilgrim and J. R. Nitschke, Pathway-Dependent Post-assembly Modification of an Anthracene-Edged  $M(II)_4L_6$  Tetrahedron, *J. Am. Chem. Soc.*, 2016, **138**, 10417–10420.
- 24 Q.-Q. Yan, L.-P. Zhou, H.-Y. Zhou, Z. Wang, L.-X. Cai, X.-Q. Guo, X. Q. and Q.-F. Sun, Metallopolymers cross-linked with self-assembled  $Ln(4)L(4)$  cages, *Dalton Trans.*, 2019, **48**, 7080–7084.
- 25 L.-L. Yan, C.-H. Tan, G.-L. Zhang, L.-P. Zhou, J.-C. Bunzli and Q.-F. Sun, Stereocontrolled Self-Assembly and Self-Sorting of Luminescent Europium Tetrahedral Cages, *J. Am. Chem. Soc.*, 2015, **137**, 8550–8555.
- 26 G. R. Lorz, A. J. Gosselin, B. A. Trump, A. H. P. York, A. Sturluson, C. A. Rowland, G. P. A. Yap, C. M. Brown, C. M. Simon and E. D. Bloch, Understanding Gas Storage in Cuboctahedral Porous Coordination Cages, *J. Am. Chem. Soc.*, 2019, **141**, 12128–12138.
- 27 G. R. Lorz, B. A. Trump, C. M. Brown and E. D. Bloch, Selective Gas Adsorption in Highly Porous Chromium(II)-Based Metal–Organic Polyhedra, *Chem. Mater.*, 2017, **29**, 8583–8587.
- 28 J. Park, Z. Perry, Y. P. Chen, J. Bae and H. C. Zhou, Chromium(II) Metal–Organic Polyhedra as Highly Porous Materials, *ACS Appl. Mater. Interfaces*, 2017, **9**, 28064–28068.
- 29 T. Zhang, J. Brown, R. J. Oakley and C. F. J. Faul, Towards functional nanostructures: Ionic self-assembly of polyoxometalates and surfactants, *Curr. Opin. Colloid Interface Sci.*, 2009, **14**, 62–70.
- 30 H. N. Miras, J. Yan, D.-L. Long and L. Cronin, Engineering polyoxometalates with emergent properties, *Chem. Soc. Rev.*, 2012, **41**, 7403–7430.
- 31 M.-P. Santoni, G. S. Hanan and B. Hasenknopf, Covalent multi-component systems of polyoxometalates and metal complexes: Toward multi-functional organic–inorganic hybrids in molecular and material sciences, *Coord. Chem. Rev.*, 2014, **281**, 64–85.
- 32 V. Das, R. Kaushik and F. Hussain, Heterometallic 3d-4f polyoxometalates: An emerging field with structural diversity to multiple applications, *Coord. Chem. Rev.*, 2020, **413**, 213271.
- 33 J. M. Cameron, G. Guillemot, T. Galambos, S. S. Amin, E. Hampson, K. Mall Haidaraly, G. N. Newton and G. Izzet, Supramolecular assemblies of organo-functionalised hybrid polyoxometalates: from functional building blocks to hierarchical nanomaterials, *Chem. Soc. Rev.*, 2022, **51**, 293–328.
- 34 N. Ogiwara, T. Iwano, T. Ito and S. Uchida, Proton conduction in ionic crystals based on polyoxometalates, *Coord. Chem. Rev.*, 2022, **462**, 214524.
- 35 H. Choi, Y. U. Kwon and O. H. Han, Nanocomposite gels between  $(V_{10}O_{28})^{-6}$  and  $[AlO_4(Al_{12}(OH)_{24}(H_2O)_{12})]^{7+}$  polyoxometalate clusters, *Chem. Mater.*, 1999, **11**, 1641–1643.
- 36 Y.-J. Liu, H.-F. Su, Y.-F. Sun, S.-T. Wang, C.-Y. Zhang, W.-H. Fang and J. Zhang, Supracluster Assembly of Archimedean Cages with 72 Hydrogen Bonds for the Aldol Addition Reaction, *Angew. Chem., Int. Ed.*, 2023, **62**, e202309971.
- 37 D. Xia, P. Wang, X. Ji, N. M. Khashab, J. L. Sessler and F. Huang, Functional Supramolecular Polymeric Networks: The Marriage of Covalent Polymers and Macrocyclic-Based Host-Guest Interactions, *Chem. Rev.*, 2020, **120**, 6070–6123.
- 38 Y. Xia, T. D. Nguyen, M. Yang, B. Lee, A. Santos, P. Podsiadlo, Z. Tang, S. C. Glotzer and N. A. Kotov, Self-assembly of self-limiting monodisperse supraparticles from polydisperse nanoparticles, *Nat. Nanotechnol.*, 2011, **6**, 580–587.
- 39 B. K. Teo, H. Zhang and X. Shi, The First Example of a Trimetallic Biicosahedral Supracluster and Its Implication for the Vertex-Sharing Polyicosahedral Growth, *J. Am. Chem. Soc.*, 1993, **115**, 8489–8490.
- 40 M. J. Dale, P. J. Dyson, B. F. G. Johnson, P. R. R. Langridge-Smith and H. T. Yates, Laser-desorption mass spectrometry of  $[Ru_6C(CO)_{17}]$  and its derivatives: Cluster aggregation in the gas phase, *J. Chem. Soc. Dalton*, 1996, 771–774.
- 41 L. Geng, C.-H. Liu, S.-T. Wang, W.-H. Fang and J. Zhang, Designable Aluminum Molecular Rings: Ring Expansion and Ligand Functionalization, *Angew. Chem., Int. Ed.*, 2020, **59**, 16735–16740.
- 42 S.-T. Wang, S.-H. Zhang, W.-H. Fang and J. Zhang, Stepwise Coordination Assembly Approach toward Aluminum-Lanthanide-based Compounds, *Inorg. Chem.*, 2020, **59**, 13760–13766.
- 43 S. Yao, W.-H. Fang, Y. Sun, S.-T. Wang and J. Zhang, Mesoporous Assembly of Aluminum Molecular Rings for Iodine Capture, *J. Am. Chem. Soc.*, 2021, **143**, 2325–2330.
- 44 Y.-J. Liu, Y.-F. Sun, S.-H. Shen, S.-T. Wang, Z.-H. Liu, W.-H. Fang, D. S. Wright and J. Zhang, Water-stable porous  $Al_{24}$  Archimedean solids for removal of trace iodine, *Nat. Commun.*, 2022, **13**, 6632.
- 45 Z.-H. Liu, S.-H. Shen, C.-Y. Zhang, J. Niu, Q.-H. Li, J. Zhang and W.-H. Fang, Design and synthesis of a deep-cavity aluminium-organic macrocycle to trap dyes and generate enhanced non-linear optical performance, *Inorg. Chem. Front.*, 2024, **11**, 3777–3785.
- 46 S.-T. Wang, X. Qi, R.-Q. Chen, W.-H. Fang and J. Zhang, Two solvent-dependent  $Al_{16}$  nanorings: design, synthesis and nonlinear optical limiting behavior, *Inorg. Chem. Front.*, 2024, **11**, 462–469.
- 47 Y. Zhang, Q.-H. Li, W.-H. Fang and J. Zhang, Aluminum molecular rings bearing amino-polyalcohol for iodine capture, *Inorg. Chem. Front.*, 2022, **9**, 592–598.
- 48 D. Dini, M. J. Calvete and M. Hanack, Nonlinear Optical Materials for the Smart Filtering of Optical Radiation, *Chem. Rev.*, 2016, **116**, 13043–13233.
- 49 W.-H. Zhang, Q. Liu and J.-P. Lang, Heterometallic transition metal clusters and cluster-supported coordination polymers derived from Tp- and Tp\*-based Mo(W) sulfido precursors, *Coord. Chem. Rev.*, 2015, **293**, 187–210.
- 50 K. Nakashima, R. Suizu, S. Morishita, N. Tsurumachi, M. Funahashi, H. Masu, R. Ozawa, K. Nakamura and K. Awaga, Enhanced Circularly Polarized Luminescence by

- a Homochiral Guest–Host Interaction in Gyroidal MOFs, [Ru(bpy)<sub>3</sub>] [M<sub>2</sub>(ox)<sub>3</sub>] (bpy = 2,2'-Bipyridyl, ox = Oxalate, M = Zn, Mn), *ACS Mater. Au*, 2023, **3**, 201–205.
- 51 M. Saqlain, H. M. Zohaib, S. Qamar, H. Malik and H. Li, Strategies for the enhancement of CPL properties, *Coord. Chem. Rev.*, 2024, **501**, 215559.
- 52 S.-J. Bao, Z.-M. Xu, Y. Ju, Y.-L. Song, H. Wang, Z. Niu, X. Li, P. Braunstein and J.-P. Lang, The Covalent and Coordination Co-Driven Assembly of Supramolecular Octahedral Cages with Controllable Degree of Distortion, *J. Am. Chem. Soc.*, 2020, **142**, 13356–13361.
- 53 Z.-K. Wang, M.-H. Du, P. Braunstein and J.-P. Lang, A Cut-to-Link Strategy for Cubane-Based Heterometallic Sulfide Clusters with Giant Third-Order Nonlinear Optical Response, *J. Am. Chem. Soc.*, 2023, **145**, 9982–9987.
- 54 Q. Liu, W.-H. Zhang and J.-P. Lang, Versatile thiomolybdate (thiotungstate)-copper-sulfide clusters and multidimensional polymers linked by cyanides, *Coord. Chem. Rev.*, 2017, **350**, 248–274.
- 55 H. Hou, Y. Wei, Y. Song, Y. Fan and Y. Zhu, First octameric ellipsoid lanthanide (III) complexes: crystal structure and nonlinear optical absorptive and refractive properties, *Inorg. Chem.*, 2004, **43**, 1323–1327.
- 56 P. Mathur, S. Ghose, M. M. Hossain, C. Satyanarayana, S. Banerjee, G. R. Kumar, P. B. Hitchcock and J. F. Nixon, Phosphaalkyne-bridged clusters: synthesis, characterization, and nonlinear optical properties of Fe<sub>4</sub>Se<sub>2</sub>(μ-Se<sub>2</sub>PCBut)(CO)<sub>11</sub>, *Organometallics*, 1997, **16**, 3815–3818.
- 57 D.-J. Li, Q.-H. Li, Z.-R. Wang, Z.-Z. Ma, Z.-G. Gu and J. Zhang, Interpenetrated Metal-Porphyrinic Framework for Enhanced Nonlinear Optical Limiting, *J. Am. Chem. Soc.*, 2021, **143**, 17162–17169.
- 58 B. P. Biswal, S. Valligatla, M. Wang, T. Banerjee, N. A. Saad, B. M. K. Mariserla, N. Chandrasekhar, D. Becker, M. Addicoat, I. Senkovska, R. Berger, D. N. Rao, S. Kaskel and X. Feng, Nonlinear Optical Switching in Regioregular Porphyrin Covalent Organic Frameworks, *Angew. Chem., Int. Ed.*, 2019, **58**, 6896–6900.