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# Dual-ligand-functionalized dodeca-nuclear lanthanide–tungsten-cluster incorporated selenotungstates and fluorescence detection of dipicolinic acid (an anthrax biomarker)†

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A series of rigid–flexible-ligand functionalized dodeca-nuclear lanthanide–tungsten heterometal cluster incorporated ring-shaped selenotungstates  $[\text{H}_2\text{N}(\text{CH}_3)_2]_{16}\text{Na}_2[\text{Ln}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\text{O}_{21}][\text{B}-\alpha\text{-SeW}_9\text{O}_{33}]_4\cdot 29\text{H}_2\text{O}$  [ $\text{Ln} = \text{Ce}^{3+}$  (**1**),  $\text{Pr}^{3+}$  (**2**),  $\text{Nd}^{3+}$  (**3**);  $\text{H}_2\text{PZDA} = 2,3$ -pyrazine dicarboxylic acid,  $\text{H}_2\text{FMA} = \text{fumaric acid}$ ] and  $[\text{H}_2\text{N}(\text{CH}_3)_2]_{14}\text{Cs}_2\text{Na}_2[\text{Eu}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\text{O}_{21}][\text{B}-\alpha\text{-SeW}_9\text{O}_{33}]_4\cdot 30\text{H}_2\text{O}$  (**4**) were prepared by a cooperative strategy of rigid and flexible ligands. The polyanion skeletons of **1–4** comprise an inorganic–organic hybrid dodeca-nuclear  $[\text{Ln}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\text{O}_{21}]^{14+}$  heterometallic cluster surrounded by four trivalent Keggin  $[\text{B}-\alpha\text{-SeW}_9\text{O}_{33}]^{8-}$  subunits in a rectangle-like arrangement. Based on its fluorescence properties, **4** was used as a fluorescent probe for detecting an anthrax biomarker dipicolinic acid (DPA) with a low limit of detection, and excellent selectivity and anti-interference. The possible fluorescence enhancement mechanism of **4** in the process of detecting DPA may stem from energy transfer from DPA to  $\text{Eu}^{3+}$  ions in **4**, which can be confirmed by decay lifetime measurements of **4** in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  with and without DPA, and a positive signal of the partial substitution of  $\text{H}_2\text{O}$  ligands on  $\text{Eu}^{3+}$  ions by DPA ligands. This work proposes a feasible cooperative strategy of rigid and flexible ligands for constructing new structural types of polyoxometalate hybrids, and also provides some meaningful insights into efficient biomolecule detection by utilizing polyoxometalate-based luminescence materials.

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## Introduction

The design and fabrication of inorganic–organic hybrid fluorescence materials have drawn considerable attention in view of their wide applications in optical devices, displays, cell imaging, sensing, laser lighting, and so on.<sup>1,2</sup> Such materials combining inorganic and organic components can overcome the drawbacks of a single component and have a synergistic effect between two components. For example, the sensitization of lanthanide emissions by organic chromophores can play an important role in developing function-oriented structural assembly, functional complementation and performance optimization.<sup>3–5</sup>

Polyoxometalates (POMs) are a class of inorganic metal–oxo clusters and have exhibited their rich architectural diversity

and fascinating properties in optics, catalysis, magnetism, biomedicine, nanotechnology, *etc.*<sup>6–11</sup> As the most widely researched branch of POMs, polyoxotungstates (POTs), particularly inorganic–organic hybrid POT materials (IOHPMs), have attracted ongoing research interest.<sup>12,13</sup> In this respect, the persistent pursuit of IOHPMs was driven by the strong combining capacity of organic ligands towards metal centers or POTs, where organic ligands can provide abundant binding sites and modes for their coordination assembly by subtly acclimating themselves to the whole geometrical requirements.<sup>14–17</sup> Flexible organic ligands feature alkyl carbon-chain backbones with carboxyl, hydroxyl or amino groups and can bend or rotate freely around single bonds, whereas rigid organic ligands usually possess an aromatic or heterocyclic backbone that makes them spatially limited and oriented.<sup>18–20</sup> Notably, flexible or rigid ligands serving as chelating or bridging linkers have been extensively applied to a POT system and have demonstrated their outstanding potential for the fabrication of high-nuclear metal-substituted or extended IOHPMs.<sup>5,21,22</sup> However, mixed-ligand-participated IOHPMs remain underexplored. For instance, the great potential of organic ligands applied to the POT system has been shown in the design and

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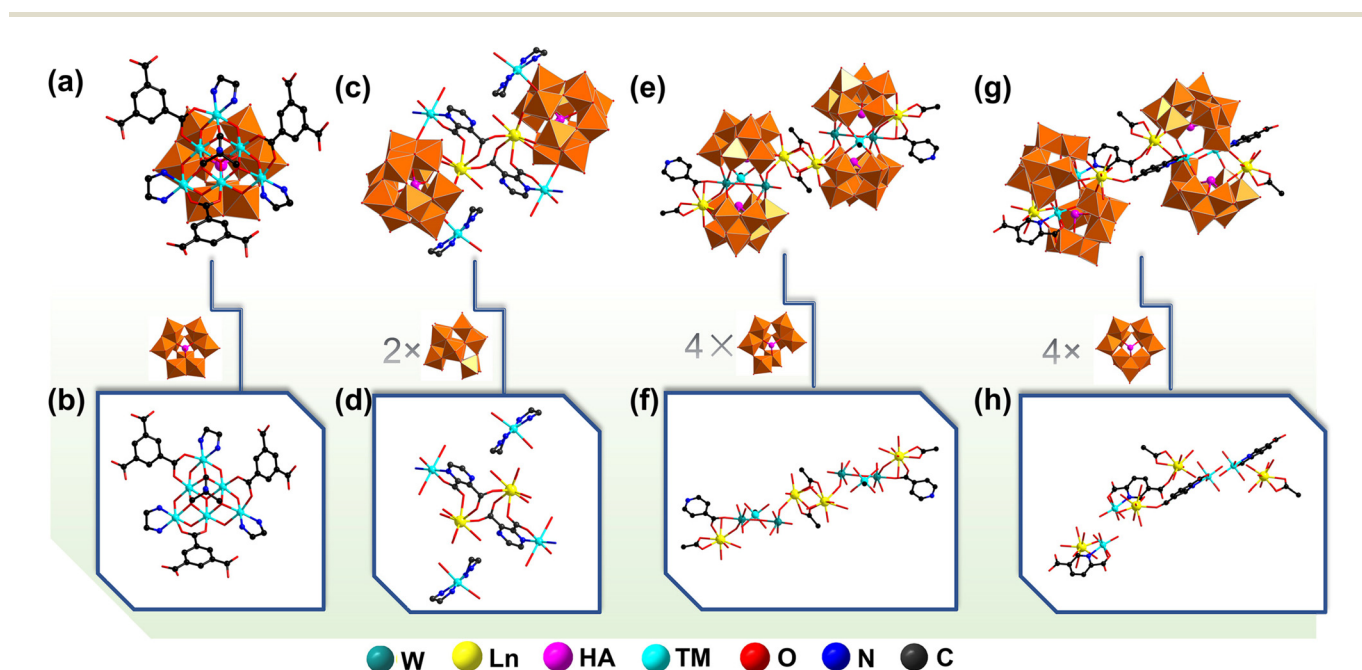
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assembly of POT–organic frameworks based on inorganic–organic Ni<sub>6</sub>-substituted {Ni<sub>6</sub>(μ<sub>3</sub>-OH)<sub>3</sub>(H<sub>2</sub>O)<sub>6</sub>Ln(B-α-PW<sub>9</sub>O<sub>34</sub>)} building blocks (BBs) (Fig. 1a and b).<sup>23</sup>

According to the soft–hard acid–base rule, ligands with N/O donors have close affinity with hard lanthanide (Ln) cations. Thus, the introduction of Ln ions can provide more opportunities for assembling structures and discovering properties.<sup>24–29</sup> As can be seen in a dual-ligand-functionalized Ln-incorporated POT {[Cu(en)<sub>2</sub>][Cu(en)<sub>2</sub>(H<sub>2</sub>O)][(α-H<sub>2</sub>SiW<sub>11</sub>O<sub>39</sub>)Dy(H<sub>2</sub>O)(PZDA)]<sub>2</sub>}<sup>6-</sup> (en = ethylenediamine) (Fig. 1c and d), each tetradentate PZDA<sup>2-</sup> ligand coordinates to two Dy<sup>3+</sup> ions and one [Cu(en)<sub>2</sub>]<sup>2+</sup> ion by one N and three carboxyl O atoms.<sup>30</sup> In 2017, Zhao *et al.* reported a series of novel Ln–organometal co-encapsulated tellurotungstates {[Sn(CH<sub>3</sub>)W<sub>2</sub>O<sub>4</sub>(IN)][(B-α-TeW<sub>8</sub>O<sub>31</sub>)Ln(H<sub>2</sub>O)(CH<sub>3</sub>COO)]<sub>2</sub>}<sup>20-</sup> (Ln = Ce<sup>3+</sup>–Tb<sup>3+</sup>; CH<sub>3</sub>COOH = acetic acid, HIN = isonicotinic acid), in which the IN<sup>-</sup> ligand is combined with one {W<sub>2</sub>O<sub>4</sub>} fragment and one CH<sub>3</sub>COO<sup>-</sup>-decorated Ln<sup>3+</sup> ion (Fig. 1e and f).<sup>31</sup> In 2019, Song *et al.* reported rarely seen four-ligand-participating Fe–Ln-heterometallic tellurotungstates [Ln<sub>2</sub>(CH<sub>3</sub>COO)<sub>2</sub>(H<sub>2</sub>O)<sub>4</sub>Fe<sub>2</sub>(HPDCA)<sub>2</sub>(B-β-TeW<sub>9</sub>O<sub>33</sub>)<sub>2</sub>][Ln<sub>2</sub>(H<sub>2</sub>O)<sub>8</sub>Fe<sub>2</sub>(HPDCA)<sub>2</sub>(B-β-TeW<sub>9</sub>O<sub>33</sub>)<sub>2</sub>]<sup>14-</sup> (Ln = Eu<sup>3+</sup>, Tb<sup>3+</sup>, Dy<sup>3+</sup>, Er<sup>3+</sup>; H<sub>2</sub>PDCA = 2,5-pyridine-dicarboxylic acid). Fig. 1g and h display a one-dimensional chain arrangement assembled from tetrameric structural units bridged by HPDCA<sup>-</sup> linkers.<sup>32</sup> Overall, reports on metal-functionalized IOHPMs (MFIOHPMs) with mixed flexible and rigid ligands are relatively less. Therefore, preparing new-type high-nuclear or extended MFIOHPMs modulated by flexible–rigid co-ligands and exploring their properties are of great

significance for the sustainable development of POM chemistry.

Based on the above considerations, we are very interested in the synthetic probability by utilizing *in situ* formed vacant POT units to react with Ln ions in the presence of rigid and flexible ligands to construct novel mixed-organic ligand functionalized MFIOHPMs with innovative structures and fascinating properties. Firstly, the stereo-directed [SeO<sub>3</sub>]<sup>2-</sup> heteroanion with a lone pair of electrons in the presence of tungstates is used to stimulate the generation of negatively charged lacunary selenotungstate (ST) units, which can grasp more positive Ln ions for polymeric assembly. Secondly, flexible fumaric acid (H<sub>2</sub>FMA) with two carboxyl groups, and rigid 2,3-pyrazine dicarboxylic acid (H<sub>2</sub>PZDA) with two carboxyl groups and two pyrazine-N atoms possess rich coordination modes and strong coordination preference to Ln and W centers, and are also conducive to building up hybrid aggregates. In addition, *in situ* formed POT units and introduced organic ligands are generally used as inorganic and organic chromophores to sensitize the emission of Ln ions, which can provide the optical application potentiality of the desired mixed-organic ligand functionalized MFIOHPMs.<sup>5,31</sup> As expected, a series of unique dual-ligand functionalized dodeca-nuclear Ln–W-cluster incorporated [H<sub>2</sub>N(CH<sub>3</sub>)<sub>2</sub>]<sub>16</sub>Na<sub>2</sub>[Ln<sub>4</sub>(H<sub>2</sub>O)<sub>6</sub>(HPZDA)<sub>2</sub>(HFMA)<sub>2</sub>W<sub>8</sub>O<sub>21</sub>][B-α-SeW<sub>9</sub>O<sub>33</sub>]<sub>4</sub>·29H<sub>2</sub>O [Ln = Ce<sup>3+</sup> (1), Pr<sup>3+</sup> (2), Nd<sup>3+</sup> (3)] and [H<sub>2</sub>N(CH<sub>3</sub>)<sub>2</sub>]<sub>14</sub>Cs<sub>2</sub>Na<sub>2</sub>[Eu<sub>4</sub>(H<sub>2</sub>O)<sub>6</sub>(HPZDA)<sub>2</sub>(HFMA)<sub>2</sub>W<sub>8</sub>O<sub>21</sub>][B-α-SeW<sub>9</sub>O<sub>33</sub>]<sub>4</sub>·30H<sub>2</sub>O (4) are successfully prepared, whose polyanions are constructed from four trivacant Keggin [B-α-SeW<sub>9</sub>O<sub>33</sub>]<sup>8-</sup> subunits encapsulating an inorganic–organic



**Fig. 1** Summary of representative structures of IOHPMs. (a and b) View of {Ni<sub>6</sub>(Tris)(en)<sub>3</sub>(BTC)<sub>3</sub>(B-α-PW<sub>9</sub>O<sub>34</sub>)} (Tris = tris(hydroxymethyl)amino-methane, en = ethylenediamine) and its remaining part without {B-α-PW<sub>9</sub>O<sub>34</sub>}. (c and d) View of {[Cu(en)<sub>2</sub>]<sub>2</sub>[Cu(PZDA)<sub>2</sub>][(α-H<sub>2</sub>SiW<sub>11</sub>O<sub>39</sub>)Ce(H<sub>2</sub>O)]<sub>2</sub>}<sup>6-</sup> and its remaining part without {α-H<sub>2</sub>SiW<sub>11</sub>O<sub>39</sub>}. (e and f) View of {[Sn(CH<sub>3</sub>)W<sub>2</sub>O<sub>4</sub>(IN)][(B-α-TeW<sub>8</sub>O<sub>31</sub>)Ln(H<sub>2</sub>O)(CH<sub>3</sub>COO)]<sub>2</sub>}<sup>20-</sup> and its remaining part without {B-α-TeW<sub>8</sub>O<sub>31</sub>}. (g and h) View of [Ln<sub>2</sub>(CH<sub>3</sub>COO)<sub>2</sub>(H<sub>2</sub>O)<sub>4</sub>Fe<sub>2</sub>(HPDCA)<sub>2</sub>(B-β-TeW<sub>9</sub>O<sub>33</sub>)<sub>2</sub>][Ln<sub>2</sub>(H<sub>2</sub>O)<sub>8</sub>Fe<sub>2</sub>(HPDCA)<sub>2</sub>(B-β-TeW<sub>9</sub>O<sub>33</sub>)<sub>2</sub>]<sup>14-</sup> and its remaining part without {B-β-TeW<sub>9</sub>O<sub>33</sub>}.

hybrid dodeca-nuclear  $[\text{Ln}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\text{O}_{21}]^{14+}$  heterometal cluster. Intriguingly, the coexistence of rigid  $\text{HPZDA}^-$  and flexible  $\text{HFMA}^-$  ligands allows diverse coordination of pyrazine-N and carboxy-O sites to both or either of Ln and W centers, leading to the versatile generation of dual-ligand-functionalized high-nuclear heterometallic clusters, accompanied by the formation of five-, six- and seven-membered rings in the structures. Interestingly, **4** emits remarkable fluorescence (FL) and can be applied as a FL probe to detect a typical anthrax biomarker dipicolinic acid (DPA); it exhibits a low detection limit (3.83  $\mu\text{M}$ ), a rapid respond time (10 s) and outstanding selectivity and anti-interference. When it is applied in a practical complex environment, the **4**-based FL sensor still exhibits good detection activity and acceptable recovery ranges. Moreover, the possible mechanism of recognizing DPA is studied. This research shows the good application potentiality of rigid and flexible-ligand functionalized MFIOHPMs in the aspect of FL detection of the anthrax biomarker DPA.

## Results and discussion

### Synthesis

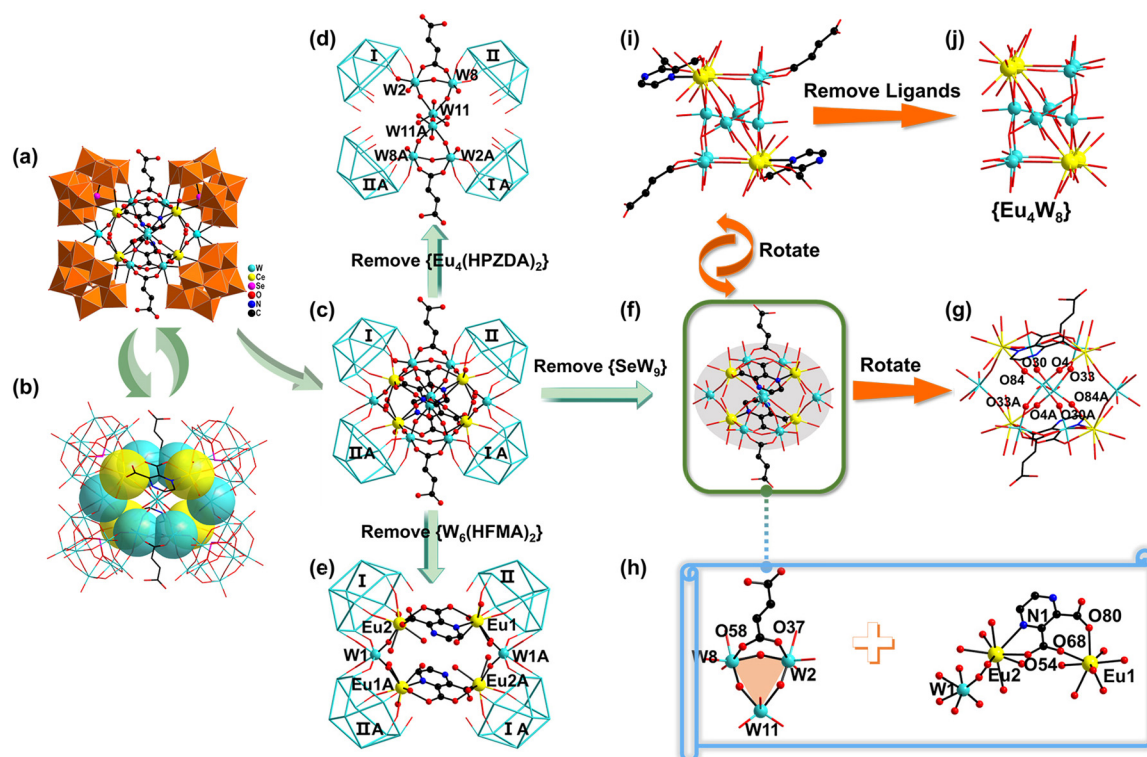
Through relevant investigations on STs, it is found that the one-pot self-assembly reaction is a suitable and effective synthetic strategy for preparing novel ST derivatives.<sup>33,34</sup> Also, besides the delicate regulation on reaction parameters such as the molar ratio of reactants and reaction temperature, the acidic pH window from 3 to 5 is more favorable for the generation of “open” lacunary ST units.<sup>35,36</sup> On the basis of a previous work, **1–4** were obtained by the reaction of  $\text{Na}_2\text{WO}_4 \cdot 2\text{H}_2\text{O}$ ,  $\text{Na}_2\text{SeO}_3$ ,  $\text{Ln}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$  ( $\text{Ln} = \text{Ce}^{3+}$ ,  $\text{Pr}^{3+}$ ,  $\text{Nd}^{3+}$ ,  $\text{Eu}^{3+}$ ),  $\text{H}_2\text{PZDA}$ ,  $\text{H}_2\text{FMA}$ ,  $\text{CH}_3\text{COOH}$  and dimethylamine hydrochloride in aqueous solution ( $\text{pH} = 3$ ). For **4**,  $\text{CsCl}$  was added to the reaction system for growing good-quality crystals. However, parallel experiments revealed that the introduction of  $\text{CH}_3\text{COOH}$  guarantees a relatively clear reaction system and the formation of high-quality crystals with high yield. Otherwise, a large amount of amorphous precipitate was formed during the crystal growth process, so the crystal yield was greatly reduced. We speculated that  $\text{CH}_3\text{COOH}$  could not only work as an acid regulator, but also work as an organic solubilizer to ameliorate the reactivity of Ln and tungstates.<sup>37</sup> Thus,  $\text{CH}_3\text{COOH}$  is an indispensable starting material in the synthesis of **1–4**. Besides, much attention was also paid to other reaction factors including pH, the ratio of W : Se and the usage of  $\text{Ln}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ . Comparative experiments showed that the pH range of 2.8–3.5 is conducive to crystal generation; in particular, a pH value of 3 is the optimal value. The ratio of W : Se between 17.5 and 32.8 favors the generation of **1–4**, and the optimum ratio is 25.97. In addition, the dosage of  $\text{Ln}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$  between 0.450 and 0.700 g results in the successful formation of **1–4** and the highest yield was achieved when the amount was around 0.500 g. Moreover, the addition of  $\text{CsCl}$  greatly promoted the good crystallinity of **4**.

### Structural description

The experimental XRD patterns coincide well with their corresponding simulated patterns, illustrating the good phase purity of **1–4** (Fig. S1†). The IR and TG tests of **1–4** have been recorded for structural characterization (Fig. S2 and S3†).

The crystallographic structural parameters of **1–4** are summarized in Table S1.† As the structures of **1–4** are similar, only the structure of **4** will be discussed in detail here. **4** consists of an unusual rigid-flexible-ligand functionalized multi-metal-stabilized tetrameric polyanion  $[\text{Eu}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\text{O}_{21}][\text{B-}\alpha\text{-SeW}_9\text{O}_{33}]_4^{18-}$  (**4a**, Fig. 2a). In the tetrameric **4a** polyanion, four trivalent Keggin subunits  $[\text{B-}\alpha\text{-SeW}_9\text{O}_{33}]^{8-}$  ( $\{\text{SeW}_9\}$ ) are aligned in a rectangle-like mode and connected by a dodeca-nuclear heterometal cluster  $[\text{Eu}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\text{O}_{21}]^{14+}$  ( $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$ ) through W–O–W and W–O–Eu linkers into a novel ring-shaped architecture. The heterometal cluster is situated at the center of **4a** (Fig. 2b), whilst the O2 atom is the center of symmetry of the whole polyanion (Fig. S4†). Remarkably, the exposed vacancies of each trivalent  $\{\text{SeW}_9\}$  subunit are filled with W and Eu centers. For better understanding the assembly of peripheral  $\{\text{SeW}_9\}$  subunits (labelled as I, II, IA and IIA) and the internal  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster, the tetrameric **4a** polyanion can be simplified and displayed in two more perspicuous ways by breaking up the intricate  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  core (Fig. 2c–e). One way is the connection of I, II, IA and IIA  $\{\text{SeW}_9\}$  subunits by the  $\text{HFMA}^-$ -decorated hourglass-like hexa-W cluster  $[\text{W}_6\text{O}_{17}(\text{HFMA})_2]$  ( $\{\text{W}_6(\text{HFMA})_2\}$ ), where the adjacent I and II  $\{\text{SeW}_9\}$  subunits are grasped together by W2 and W8 ions into a dimer, and two dimers are further concatenated with each other through O–W11–O bonds to generate the tetramer (Fig. 2d). The other way is the fusion of I, II, IA and IIA  $\{\text{SeW}_9\}$  subunits by the  $\text{HPZDA}$ -bridged annular hexa-nuclear heterometal cluster  $[\text{Eu}_4(\text{H}_2\text{O})_6(\text{HPZDA})_2\text{W}_2\text{O}_4]^{14+}$  ( $\{\text{Eu}_4(\text{HPZDA})_2\text{W}_2\}$ ), where  $\text{Eu}^{3+}$  and  $\text{Eu}^{2+}$  ions are respectively encapsulated into the pockets of II and I  $\{\text{SeW}_9\}$  subunits into a C-shaped dimer and two dimers directly coordinate with rigid  $\text{HPZDA}^-$  ligands with the support of pyrazine N and carboxylic O sites into a tetramer (Fig. 2e). Thus, the double bridging roles of the hexa-nuclear  $\{\text{W}_6(\text{HFMA})_2\}$  cluster and hexa-nuclear  $\{\text{Eu}_4(\text{HPZDA})_2\text{W}_2\}$  ring lead to the aggregation of four  $\{\text{SeW}_9\}$  subunits, ultimately forming the neoteric and stable assembly of the **4a** polyanion.

One prominent characteristic of **4a** is the existence of the interesting dodeca-nuclear heterometallic  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster, comprising four  $\text{Eu}^{\text{III}}$  ions, eight  $\text{W}^{\text{VI}}$  centers, two  $\text{HPZDA}^-$  ligands and two  $\text{HFMA}^-$  ligands (Fig. 2f). The dodeca-nuclear cluster can be reckoned as the combination of a  $\{\text{W}_6(\text{HFMA})_2\}$  unit and a  $\{\text{Eu}_4(\text{HPZDA})_2\text{W}_2\}$  ring by four pairs of Eu–O–W bonds, where the  $\{\text{W}_6(\text{HFMA})_2\}$  cluster is interpenetrated with the ring (Fig. 2g). Interestingly, the  $\{\text{W}_6(\text{HFMA})_2\}$  cluster links to a “hourglass” by superimposing W11 and W11A atoms together, where the top and the bottom are two identical mono- $\text{HFMA}^-$ -chelating  $\{\text{W}_3\text{O}_8(\text{HFMA})\}$  units interconnected by sharing the O2 atom (Fig. S5†). For the



**Fig. 2** (a) Tetrameric **4a** polyanion. (b) Highlighted  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster in **4a**. (c) Simplified structure of **4a**. (d) Connection mode of four  $\{\text{SeW}_9\}$  subunits and the central  $\{\text{W}_6(\text{HFMA})_2\}$  cluster. (e) Connection mode of four  $\{\text{Eu}_2(\text{HPZDA})\}$  subunits and the central  $\{\text{Eu}_4(\text{HPZDA})_2\text{W}_2\}$  cluster. (f) Top view of the  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster. (g) Highlighted O bridges in the  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster. (h) View of  $\{\text{W}_3\text{O}_8(\text{HFMA})\}$  and  $\{\text{Eu}_2(\text{HPZDA})\}$  subunits. (i) Side view of the  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster. (j) The remaining part of the  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster by removing organic ligands.

$\{\text{W}_3\text{O}_8(\text{HFMA})\}$  unit, both O58 and O37 atoms in one carboxyl group of the flexible  $\text{HFMA}^-$  ligand participate in the coordination of W8 and W2 atoms, producing an uncommon “-C-O-W-O-W-O-” six-membered ring, almost perpendicular to the apex-O-linked- $\text{W}_3$ -based triangle (Fig. 2h). The  $\{\text{Eu}_4(\text{HPZDA})_2\text{W}_2\}$  ring is composed of two mono- $\text{HPZDA}^-$ -chelating  $\{\text{Eu}_2(\text{HPZDA})\}$  units alternately linked by two W atoms (W1 and W1A). In each  $\{\text{Eu}_2(\text{HPZDA})\}$  unit, the multidentate  $\text{HPZDA}^-$  ligand simultaneously chelates with  $\text{Eu}1^{3+}$  and  $\text{Eu}2\text{A}^{3+}$  ions *via* N and O coordination sites, where the  $\text{Eu}1^{3+}$  ion is bonded to two O atoms (O68 and O80) from two carboxyl groups to form an unusual “-C-C-O-Eu-O-C-C-” seven-membered ring, whilst the  $\text{Eu}2^{3+}$  ion is firmly grasped by one N (N1) and one neighboring carboxyl O atom (O54) to form a classical “-N-C-C-O-Eu-” five-membered ring (Fig. S6<sup>†</sup>). Thus, it can be seen that the employment of rigid and flexible organic ligands with abundant N- and O binding sites offers more possibilities for synthesizing organically functionalized Ln-incorporated POM hybrids and discovering more interesting and versatile coordination assembly modes.<sup>38</sup> Moreover, two crystallographically unique  $\text{Eu}^{3+}$  ions in **4a** have two different geometries. Specifically, the  $\text{Eu}1^{3+}$  ion has an octa-coordinate bi-capped triangular prism geometry that is formed by two O atoms (O28, O34) from one  $\{\text{SeW}_9\}$  subunit, three O atoms (O4, O22 and O33) from three central  $\{\text{WO}_6\}$  octahedra,

one water molecule (O1W), and two carboxyl O atoms (O68, O80) of the  $\text{HPZDA}^-$  ligand [ $\text{Eu}1\text{-O}$ : 2.327(11)–2.498(14) Å] (Fig. S7a<sup>†</sup>). Distinctly, the  $\text{Eu}2^{3+}$  ion is surrounded by two O atoms (O19, O52) belonging to one  $\{\text{SeW}_9\}$  subunit, three O atoms (O26, O30 and O84) from the central  $\{\text{WO}_6\}$  octahedra, two water molecules (O2W, O3W), and one N atom and one carboxyl O atom from the  $\text{HPZDA}^-$  ligand, adopting a nine-coordinate tri-capped triangular prism geometry [ $\text{Eu}2\text{-O}$ : 2.379(13)–2.543(14) Å,  $\text{Eu}2\text{-N}$ : 2.685(17) Å] (Fig. S7b<sup>†</sup>).

A further rotation of the dodeca-nuclear  $\{\text{Eu}_4(\text{HPZDA})_2(\text{HFMA})_2\text{W}_8\}$  cluster produces a visual “cuboid”, but an irregular polyhedron (Fig. 2i and S8<sup>†</sup>), offering a new perspective on this structure. By removing all ligands that hang on the top and the bottom of the visual “cuboid”, a  $\{\text{Eu}_4\text{W}_8\}$  unit consisting of an “N”-shaped  $\{\text{W}_6\}$  cluster and a slant  $\{\text{Eu}_4\text{W}_2\}$  ring by sharing eight  $\mu_2\text{-O}$  atoms is intuitively observed (Fig. 2j and Fig. S9<sup>†</sup>), and the additional two edge lengths of  $\text{W}1\cdots\text{Eu}1$  and  $\text{W}1\cdots\text{Eu}2$  are 4.162 and 4.268 Å. The  $\{\text{W}_6\}$  cluster can be seen as a dimer of the two apex-linked  $\{\text{W}_3\text{O}_3\}$  group, where the W2, W8 and W11 group and the W11A, W8A and W2A group respectively form two identical triangles. In the  $\{\text{Eu}_4\text{W}_2\}$  ring, four  $\text{Eu}^{3+}$  ions constitute a regular parallelogram with the edge lengths of 6.265 and 7.290 Å, and two adjacent angles of 88.187° and 91.883°, respectively (Fig. S10<sup>†</sup>). With further capping on the two opposite sides of the parallelogram by W1

and W1A centers, a pseudo-hexagon made up of four Eu and two W centers is ulteriorly constructed as the angle between the parallelogram and triangle (Eu1, Eu2, W1) is as small as  $1.342^\circ$  (Fig. S11†).

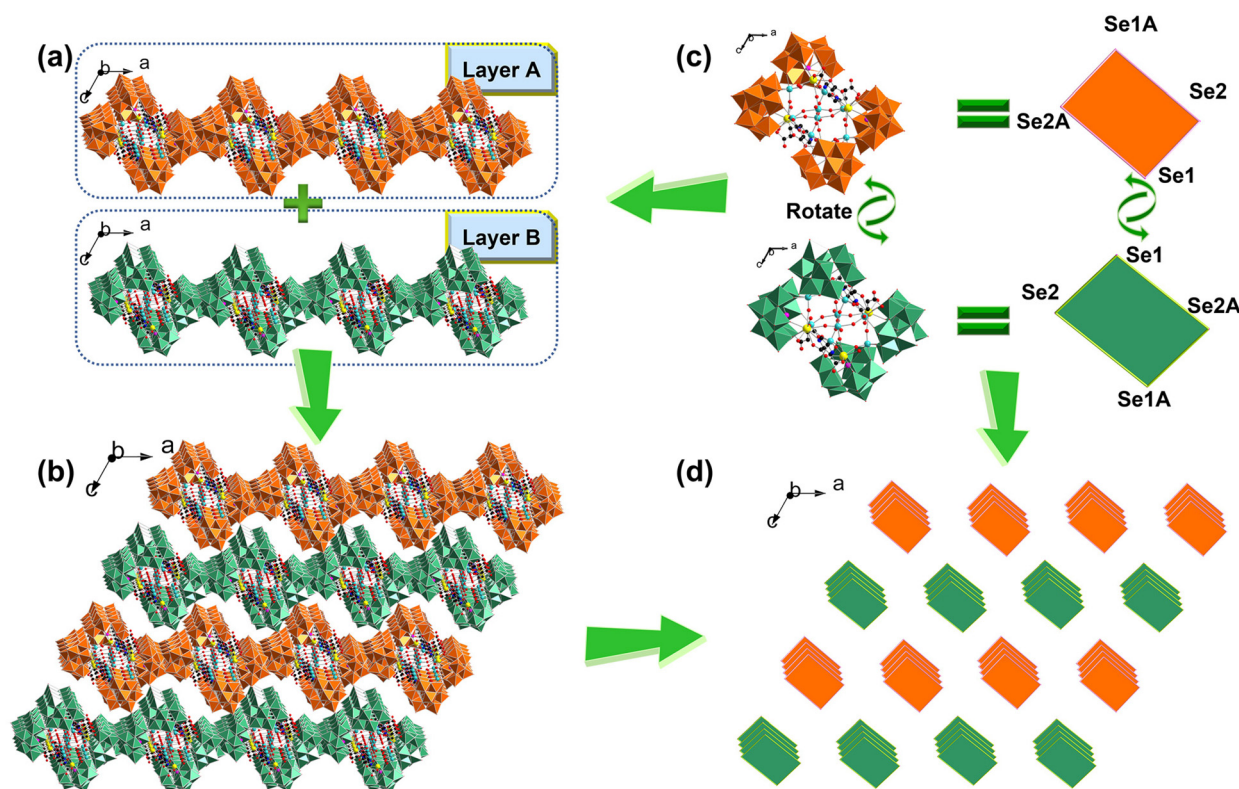
Finally, the 3D stacking pattern of **4a** polyanions is also described. It can be observed in Fig. 3 that the **4a** polyanions in the 3D stacking are aligned in an -ABAB- mode (layer A and layer B) along the *b* axis (Fig. 3a and b). Every layer in the *ac* plane features a fascinating wave-shaped arrangement. By visualizing each **4a** polyanion as a rectangle with four vertices as Se1, Se2, Se1A and Se2A atoms, a simplified 3D stacking is presented, from which we can see that layer B can be obtained by rotating **4a** polyanions in layer A by  $180^\circ$  along the *b* axis (Fig. 3c and d). Besides, the 3D stacking along the *c* axis is illustrated in Fig. S12.†

### FL properties

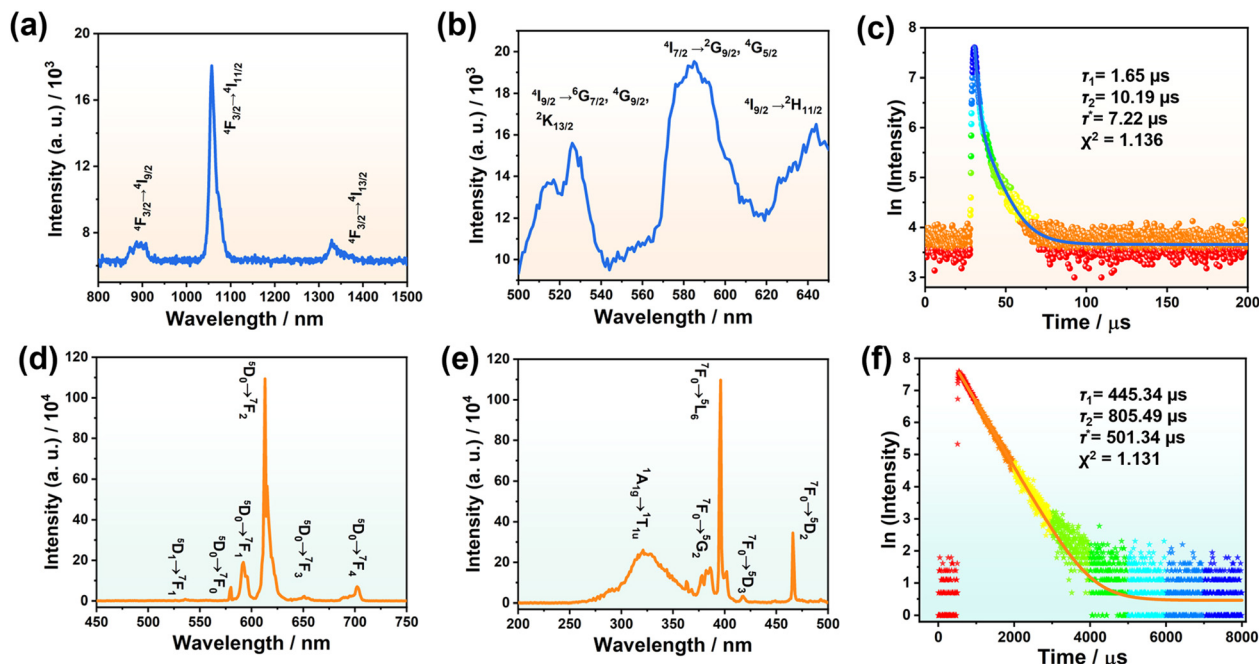
Owing to the rich spectroscopic terms of Ln ions, Ln-based materials have gained wide interest in virtue of their unique superiority in FL such as long-lived narrow emission lines, high color purity and quantum efficiencies, as well as significant applications in light-emitting diodes, displays and imaging, optical sensors and devices, *etc.*<sup>28,39–42</sup> The incorporation of the appropriate fluorescent Ln centers into the POM frameworks can endow POMs with remarkable luminescence

properties. Here, the solid-state FL properties of **3** and **4** at ambient temperature were investigated.

Under excitation at 585 nm, the emission spectrum of **3** was recorded between 800 and 1500 nm in the near-infrared region (Fig. 4a), which shows three peaks at 893, 1057 and 1329 nm, respectively, corresponding to the  ${}^4F_{3/2} \rightarrow {}^4I_{9/2}$ ,  ${}^4F_{3/2} \rightarrow {}^4I_{11/2}$  and  ${}^4F_{3/2} \rightarrow {}^4I_{13/2}$  transitions of  $\text{Nd}^{3+}$  ions.<sup>43</sup> Notably, the peak at 1057 nm ( ${}^4F_{3/2} \rightarrow {}^4I_{11/2}$ ) dominates over the whole spectrum. Meanwhile, by monitoring the emission at 1057 nm, the excitation spectrum of **3** containing several peaks at 526, 585 and 631 nm was recorded (Fig. 4b), and these peaks can be identified as  ${}^4I_{9/2} \rightarrow {}^4G_{7/2}$ ,  ${}^4G_{9/2}$ ,  ${}^2K_{13/2}$ ,  ${}^4I_{9/2} \rightarrow {}^2G_{7/2}$ ,  ${}^4G_{5/2}$ , and  ${}^4I_{9/2} \rightarrow {}^2H_{11/2}$  transitions, respectively.<sup>44</sup> In addition, the FL lifetime of **3** was measured under the conditions of the strongest excitation ( $\lambda_{\text{ex}} = 584$  nm) and emission ( $\lambda_{\text{em}} = 1057$  nm), where the lifetime curve obeys a double exponential function with the fitting results of  $\tau_1 = 1.65$   $\mu\text{s}$  (34.14%) and  $\tau_2 = 10.19$   $\mu\text{s}$  (65.86%), and the average lifetime  $\tau^*$  was calculated to be 7.22  $\mu\text{s}$  according to the equation  $\tau^* = (A_1\tau_1^2 + A_2\tau_2^2)/(A_1\tau_1 + A_2\tau_2)$  (Fig. 4c).<sup>45</sup> The FL emitting spectrum of **4** was recorded under excitation at 396 nm and exhibits six peaks at 536, 580, 592, 613, 651 and 703 nm, which correspond to the  ${}^5D_1 \rightarrow {}^7F_1$ ,  ${}^5D_0 \rightarrow {}^7F_0$ ,  ${}^5D_0 \rightarrow {}^5F_1$ ,  ${}^5D_0 \rightarrow {}^5F_2$ ,  ${}^5D_0 \rightarrow {}^5F_3$  and  ${}^5D_0 \rightarrow {}^5F_4$  transitions of  $\text{Eu}^{3+}$  ions (Fig. 4d).<sup>46</sup> The excitation spectrum of **4** was obtained through monitoring the



**Fig. 3** (a) View of layer A and layer B in the 3D stacking of **4a** polyanions. (b) The 3D stacking view of **4a** along the *b* axis. (c) Simplified models of **4a** polyanions in layer A and layer B. (d) Simplified 3D stacking of **4a** along the *b* axis. Because spatial orientations of **4a** polyanions in layer A and layer B are different, the  $\{\text{WO}_6\}$  octahedra in layer A are shown in orange while the  $\{\text{WO}_6\}$  octahedra in layer B are shown in sea green.



**Fig. 4** (a) The NIR emission spectrum of **3** ( $\lambda_{\text{ex}} = 584$  nm). (b) The excitation spectrum of **3** ( $\lambda_{\text{em}} = 1057$  nm). (c) The lifetime curve of **3** obtained by monitoring the emission at 1057 nm and the excitation at 584 nm. (d) The emission spectrum of **4** ( $\lambda_{\text{ex}} = 396$  nm). (e) The excitation spectrum of **4** ( $\lambda_{\text{em}} = 613$  nm). (f) The lifetime curve of **4** obtained by monitoring the emission at 613 nm and the excitation at 396 nm.

emission at 613 nm, and the highest intensity peak is located at 396 nm ( ${}^7F_0 \rightarrow {}^5L_6$ ) while the other peaks are situated at 320, 386, 417 and 466 nm ( ${}^1A_{1g} \rightarrow {}^1T_{1u}$ ,  ${}^7F_0 \rightarrow {}^5G_2$ ,  ${}^7F_0 \rightarrow {}^5D_3$  and  ${}^7F_0 \rightarrow {}^5D_2$ ) (Fig. 4e).<sup>47</sup> Besides, the lifetime curve of **4** obeys a double exponential function with the fitting results of  $\tau_1 = 445.34 \mu\text{s}$  (84.45%) and  $\tau_2 = 805.49 \mu\text{s}$  (15.55%), and the average  $\tau^*$  is 501.34  $\mu\text{s}$  (Fig. 4f). As shown in the CIE 1931 chromaticity diagram, **4** exhibits red emission (Fig. S13†).

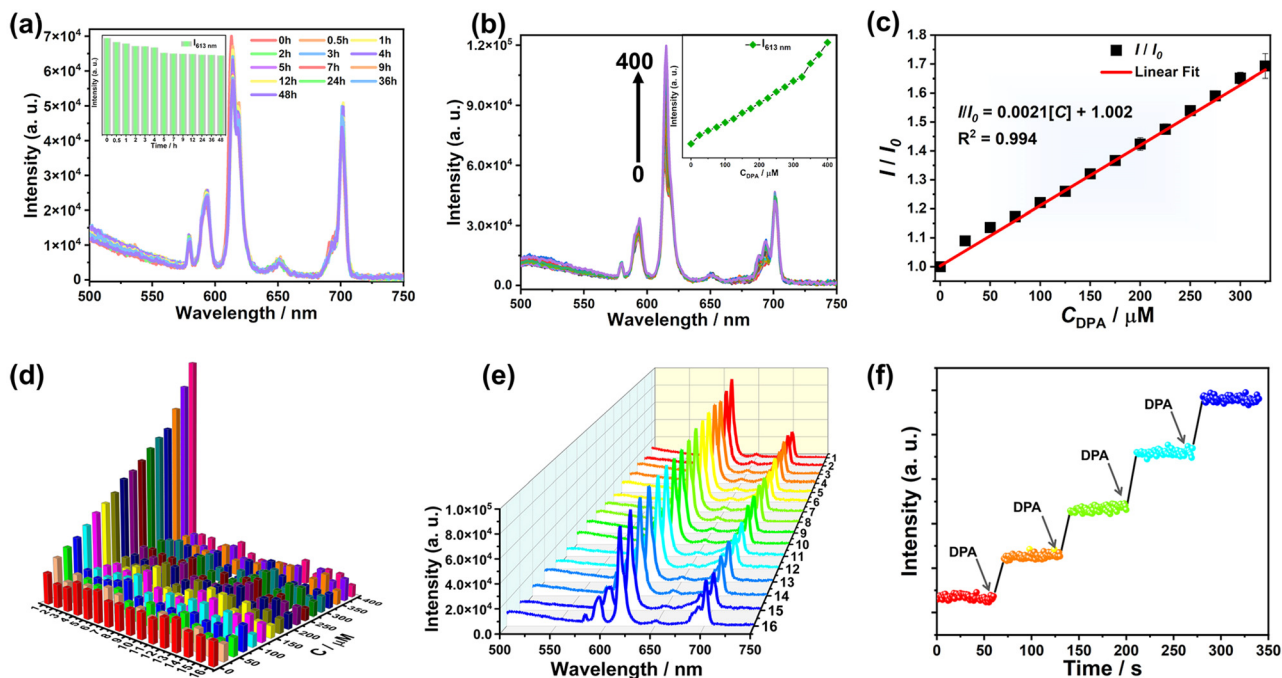
### FL sensing for DPA detection

*Bacillus anthracis* is a very dangerous bacterium that can cause a serious disease called anthrax through the inhalation of spores, and anthrax spores are also a potential biological warfare agent.<sup>48,49</sup> Therefore, it is quite indispensable to develop a rapid, accurate, and sensitive detection method. As is known, dipicolinic acid (DPA) is the main marker of anthrax spores and various analytical methods have been applied to detect DPA, such as immunological analysis, electrochemical method, mass spectrometry, biological methods, and so on.<sup>50–52</sup> However, most of these methods are limited by time/cost consumption and difficult operation of equipment. Recently, a FL sensing technique with rapid and highly efficient detection capability has offered a significant platform for DPA detection.<sup>53</sup> Here, we use **4** to construct a FL sensor to detect DPA in aqueous solution and it is necessary to take the structural stability of the **4a** polyanion in aqueous solution into account. Firstly, as revealed by electrospray ionization mass spectrometry (ESI-MS) of **4** in water, three distinct ESI-MS signals appearing at  $m/z$  1183.01 (5<sup>-</sup>), 1479.21 (4<sup>-</sup>) and 1972.32 (3<sup>-</sup>) can be respectively assigned to  $\{\text{H}_6(\text{Eu}_2\text{W}_4$

$(\text{H}_2\text{O})_3\text{O}_{11}(\text{HFMA})[\text{SeW}_9\text{O}_{33}]_2\}^{5-}$  (5915.45),  $\{\text{H}_7(\text{Eu}_2\text{W}_4(\text{H}_2\text{O})_3\text{O}_{11}(\text{HFMA})[\text{SeW}_9\text{O}_{33}]_2)^{4-}$  (5916.45) and  $\{\text{H}_8(\text{Eu}_2\text{W}_4(\text{H}_2\text{O})_3\text{O}_{11}(\text{HFMA})[\text{SeW}_9\text{O}_{33}]_2)^{3-}$  (5917.45), suggesting the comparatively good structural stability of the **4a** polyanion with the loss of HPZDA<sup>-</sup> ligands in water (Fig. S14a†). Moreover, the UV-visible spectral result of **4** in aqueous solution in 48 h can also support the comparatively good structural stability of the **4a** polyanion (Fig. S14b†). Therefore, the benign FL performance together with the water stability drives us to employ **4** as a promising FL probe in water.

The time-varying FL emission spectra of **4** within 48 h in water (3 mg mL<sup>-1</sup>) were monitored (Fig. 5a). It is evident that the intensity at 613 nm derived from the  ${}^5D_0 \rightarrow {}^7F_2$  transition gradually descends in the first 5 h, which is probably caused by dynamic exchange of the water ligands of **4a** polyanions with water molecules in aqueous solution. Then, the emission intensity remains almost unchanged in the following long time from 5 to 48 h, indicating that **4** is suitable to work as a FL probe.

A series of FL titration experiments are helpful to illustrate the sensitivity of **4** (in the following experiments, the concentration of **4** in water is 3 mg mL<sup>-1</sup>) in the detection of DPA. As demonstrated in Fig. 5b, the consecutive addition of DPA results in the enhancement of the  ${}^5D_0 \rightarrow {}^7F_2$  emission intensity of Eu<sup>3+</sup> ions. In particular, the  ${}^5D_0 \rightarrow {}^7F_2$  emission intensity is more sensitive to the introduced DPA, and the intensity (when the concentration of DPA is up to 400  $\mu\text{M}$ ) is almost 2-fold higher than that in the absence of DPA. The ratio of  $I({}^5D_0 \rightarrow {}^7F_2)/I({}^5D_0 \rightarrow {}^7F_1)$  is a criterion to evaluate the distorted degree of the local symmetry around Eu<sup>3+</sup> ions. Here, a slight



**Fig. 5** (a) Time-varying FL emission spectra of **4** (inset: the change of the peak intensity at 613 nm with time). (b) Comparison of the emission spectra of **4** with the addition of different concentrations of DPA (inset: the change of the peak intensity at 613 nm with the DPA concentration). (c) Linear relationship between the  $I/I_0$  ratio and the DPA concentration. (d) Comparison of the FL  ${}^5D_0 \rightarrow {}^7F_2$  emission intensity of **4** by the addition of DPA or each interferent with the concentration varying from 0 to 400  $\mu\text{M}$  (in the order: (1) DPA, (2) 2,3-PDCA, (3) 2,5-PDCA, (4) 2-PA, (5) NA, (6) BA, (7) *p*-PA, (8) *m*-PA, (9)  $\text{H}_3\text{BTC}$ , (10) Gly, (11) GSH, (12)  $\text{K}^+$ , (13)  $\text{Mg}^{2+}$ , (14)  $\text{Ca}^{2+}$ , (15)  $\text{Cl}^-$ , (16)  $\text{CH}_3\text{COO}^-$ ). (e) Comparison of the FL spectra of **4** in the presence of DPA (325  $\mu\text{M}$ ) and each different interferent (325  $\mu\text{M}$ ) (in the order: (1) DPA, (2) 2,3-PDCA, (3) 2,5-PDCA, (4) 2-PA, (5) NA, (6) BA, (7) *p*-PA, (8) *m*-PA, (9)  $\text{H}_3\text{BTC}$ , (10) Gly, (11) GSH, (12)  $\text{K}^+$ , (13)  $\text{Mg}^{2+}$ , (14)  $\text{Ca}^{2+}$ , (15)  $\text{Cl}^-$ , (16)  $\text{CH}_3\text{COO}^-$ ). (f) Time scan measurement of the FL response of **4** with the addition of DPA.

alteration in the intensity ratio from 2.6 to 3.8 is observed after titration with DPA (Fig. S15 and Table S2<sup>†</sup>), implying that the local site-symmetry of  $\text{Eu}^{3+}$  ions slightly changes.<sup>54</sup> Additionally, a good linear correlation exists between the emission intensity ratio ( $I/I_0$ ) and the concentration of DPA in the range of 0 to 325  $\mu\text{M}$ , wherein,  $I_0$  and  $I$  respectively represent the FL intensity of the  ${}^5D_0 \rightarrow {}^7F_2$  transition before and after the addition of DPA. The correlation obeys the equation  $I/I_0 = 0.0021C_{\text{DPA}} + 1.002$  ( $R^2 = 0.994$ ) with a limit of detection (LOD) of 3.83  $\mu\text{M}$  (Fig. 5c and Table S3<sup>†</sup>). In addition, on comparing with previous literature reports (Table S4<sup>†</sup>), it is not difficult to find that the LOD (3.83  $\mu\text{M}$ ) obtained in this work is relatively higher than some of them, but is obviously below the infectious concentration of anthrax spores (60  $\mu\text{M}$ ).<sup>55</sup> Repeated trials show the good luminescence response and repeatability of **4** to DPA (Fig. S16<sup>†</sup>).

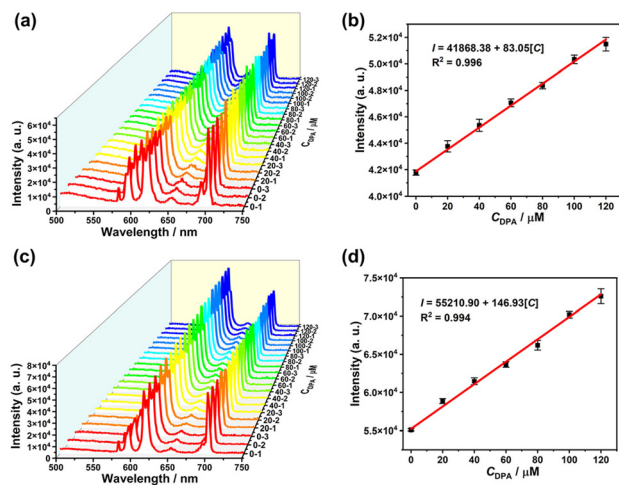
Furthermore, the selectivity and anti-interference ability of **4** for DPA recognition were explored. Some potential interferents including DPA analogues (2,3-pyridine dicarboxylic acid (2,3-PDCA), 2,5-pyridine dicarboxylic acid (2,5-PDCA), 2-picolinic acid (2-PA), nicotinic acid (NA), benzoic acid (BA), *p*-phthalic acid (*p*-PA), *m*-phthalic acid (*m*-PA), 1,3,5-benzenetricarboxylic acid ( $\text{H}_3\text{BTC}$ )), amino acids (glycine (Gly), glutathione (GSH)) (the structural formulae are listed in Scheme S2<sup>†</sup>) and anions/cations were selected. Remarkably, none of them inter-

feres with the recognition and detection of DPA because the  ${}^5D_0 \rightarrow {}^7F_2$  emission intensity of **4** is hardly or even negligibly changed by adding each interferent at various dosages (Fig. 5d and S17, S18a, b<sup>†</sup>). Moreover, markedly enhanced luminescence signals can still be acquired when DPA coexists with other interferents (325  $\mu\text{M}$ ) (Fig. 5e and S18c<sup>†</sup>). In addition, three interferents (2-PA, Gly and  $\text{Mg}^{2+}$ ) are respectively selected and the lifetime curves of **4** have been tested in the presence of each interferent at different concentrations (Fig. S19<sup>†</sup>). It is obviously observed that the addition of each interferent at different concentrations can hardly change the FL lifetime of **4**. These results clearly indicate that **4** shows outstanding selectivity and anti-interference ability towards the analytical determination of DPA.

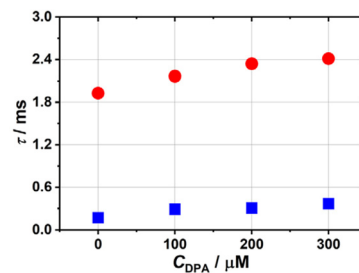
In addition, the response time of the FL sensor is also an extremely important parameter for practical use. Therefore, the time scan measurements of the FL response of **4** at the  ${}^5D_0 \rightarrow {}^7F_2$  transition were performed at an excitation wavelength of 396 nm. The emission intensity increases instantly and rapidly reaches a steady level within 10 s when DPA is added at a dosage of 100  $\mu\text{M}$  per time (Fig. 5f), which is faster than the majority of reported FL sensors applied to detect DPA (Table S4<sup>†</sup>). This result manifests that **4** can potentially work as a stable and efficient sensor for the quick detection of DPA in biological samples.

To evaluate the practical sensing application, **4** was utilized to detect DPA in tap water and a  $10^3$  times diluted human serum sample. The detection system was established by adding a sequence of concentrations of DPA in tap water and the human serum sample. It is worth mentioning that the characteristic emission intensity at 613 nm is enhanced by increasing the DPA concentration. The characteristic emission intensity at 613 nm and the DPA concentration in both systems exhibit a good linear relationship with the fitting equation  $I = 41\,868.38 + 83.05C_{\text{DPA}}$  ( $R^2 = 0.996$ ) for tap water (Fig. 6a and b) and  $I = 55\,210.90 + 146.93C_{\text{DPA}}$  ( $R^2 = 0.994$ ) for the human serum sample (Fig. 6c and d). Based on these equations, the added DPA concentration can be reasonably calculated, and the recoveries (defined as  $C_{\text{DPA,cal.}}/C_{\text{DPA,theor.}}$ ) for detecting DPA in tap water and the human serum sample were determined to be 94.98–99.83% and 96.68–104.35% with the relative standard deviation (RSD,  $n = 3$ ) in the range of 1.20–3.88% and 3.70–6.80%, respectively (Table S5<sup>†</sup>). These results suggest that **4** has great potential as a sensor for detecting DPA in real and complex environments.

Ultimately, to clarify the FL enhancement mechanism of **4** with the addition of DPA, the emission decay dynamics has been studied. In accordance with the FL decay curves, the average lifetime of  $\text{Eu}^{3+}$  ions (the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition) significantly increases from 173.92 to 367.06  $\mu\text{s}$  with the incremental addition of DPA (0–300  $\mu\text{M}$ ) (Fig. S20<sup>†</sup>). Considering the strong coordination affinity of DPA towards  $\text{Eu}^{3+}$  ions owing to its unique coordination sites with three active chelating sites (two carboxyl and one pyridyl groups), we speculate that the FL enhancement mechanism might be attributed to the replacement of coordinate  $\text{H}_2\text{O}$  ligands around  $\text{Eu}^{3+}$  ions in **4** by DPA, which can effectively inhibit the FL quenching triggered by the O–H oscillators of coordinate  $\text{H}_2\text{O}$  molecules.<sup>56–58</sup> To verify this hypothesis, the number ( $q$ ) of coordinate  $\text{H}_2\text{O}$  ligands on the  $\text{Eu}^{3+}$  ions in **4** was determined by measuring the decay lifetimes in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  with and without DPA (Fig. 7) according



**Fig. 6** FL spectra and linear relationships between the emission intensity at 613 nm and the DPA concentration in tap water (a and b) and the human serum sample (c and d).



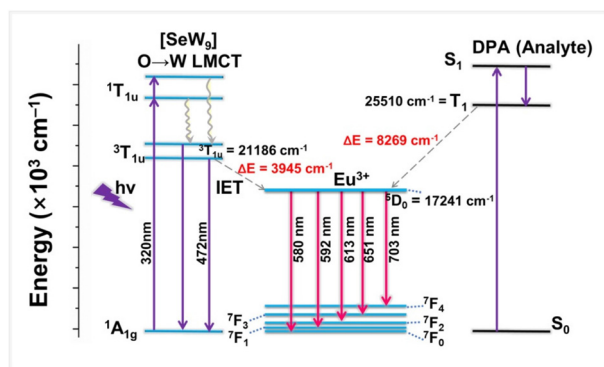
**Fig. 7** The lifetimes of **4** in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  with different DPA concentrations (blue dots represent the  $\text{H}_2\text{O}$  system and red dots represent the  $\text{D}_2\text{O}$  system).

to the equation  $q = 1.1 \times [\tau_{\text{H}_2\text{O}}^{-1} - \tau_{\text{D}_2\text{O}}^{-1} - 0.31]$ , where  $\tau_{\text{H}_2\text{O}}$  and  $\tau_{\text{D}_2\text{O}}$  are the lifetimes in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  environments, respectively, and  $q$  was found to be 5.4 in the absence of DPA ( $\tau_{\text{H}_2\text{O}} = 0.174$  ms,  $\tau_{\text{D}_2\text{O}} = 1.927$  ms), indicating that about five  $\text{H}_2\text{O}$  ligands coordinate with the  $\text{Eu}^{3+}$  centers (Table 1 and Fig. S21<sup>†</sup>).<sup>59</sup> With the addition of DPA (100, 200, 300  $\mu\text{M}$ ), the decay lifetime progressively increases while  $q$  inversely decreases to 2.20 when the DPA concentration is 300  $\mu\text{M}$  ( $\tau_{\text{H}_2\text{O}} = 0.367$  ms,  $\tau_{\text{D}_2\text{O}} = 2.411$  ms). This result indicates that DPA is more likely to coordinate with the  $\text{Eu}^{3+}$  ions. Such a coordination environment alteration of the  $\text{Eu}^{3+}$  ions leads to the enhancement of sensitization and energy transfer. In order to exclude the phenomenon that the  $\text{Eu}^{3+}$  ions separate from the **4a** polyanions to coordinate with DPA forming simple Eu–DPA complexes in the process of **4** detecting DPA, control experiments in which  $\text{Eu}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$  (3 mg  $\text{mL}^{-1}$ ) is used as a FL sensor to detect DPA at different concentrations in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  have been performed (Fig. S22 and S23a, b<sup>†</sup>). As illustrated in Fig. S22 and S23a, b,<sup>†</sup> the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_1$  emission at 592 nm is the strongest and the addition of DPA has a negligible effect on the emission intensity at 592 nm, but, in the process of **4** detecting DPA, the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  emission at 613 nm is the strongest. This comparison can confirm that the  $\text{Eu}^{3+}$  ions do not separate from the **4a** polyanions to form simple Eu–DPA complexes in the process of **4** detecting DPA. In the meantime, the lifetime curves of  $\text{Eu}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$  (3 mg  $\text{mL}^{-1}$ ) in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  with different DPA concentrations have been measured (Fig. S23c and d<sup>†</sup>). Thus, the number ( $q$ ) of coordinate  $\text{H}_2\text{O}$  molecules on the  $\text{Eu}^{3+}$  ions in  $\text{Eu}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$  has been calculated to be ca. 9 (Table S6<sup>†</sup>). Moreover, the number of coordinate  $\text{H}_2\text{O}$  molecules on the  $\text{Eu}^{3+}$  ions does not change with the increase of the DPA concentration. This observation can further confirm that  $\text{Eu}^{3+}$  ions indeed do not separate

**Table 1** Summary of the lifetimes of **4** in  $\text{H}_2\text{O}$  and  $\text{D}_2\text{O}$  with different DPA concentrations and the calculated  $q$  values

| $C_{\text{DPA}}/\mu\text{M}$ | $\tau_{\text{H}_2\text{O}}/\text{ms}$ | $\tau_{\text{D}_2\text{O}}/\text{ms}$ | $q$  |
|------------------------------|---------------------------------------|---------------------------------------|------|
| 0                            | 0.174                                 | 1.927                                 | 5.41 |
| 100                          | 0.292                                 | 2.165                                 | 2.92 |
| 200                          | 0.310                                 | 2.340                                 | 2.74 |
| 300                          | 0.367                                 | 2.411                                 | 2.20 |





**Fig. 8** The energy level diagram exhibiting the proposed energy transfer pathways from the excited triplet states ( $T_1$ ) of  $\{SeW_9\}$  and DPA ligands to the  $^5D_0$  state of the  $Eu^{3+}$  ions. IET is internal energy transfer.

from **4a** polyanions to form simple Eu–DPA complexes in the process of **4** detecting DPA. Based on the above analysis, a possible schematic diagram of energy transfer is provided to illustrate the possible mechanism (Fig. 8). Judging by Reinhoudt's empirical rule, energy transfer from the analyte to  $Eu^{3+}$  ions occurs when the energy gap between the lowest triplet energy level of the analyte and  $^5D_0$  of the  $Eu^{3+}$  ion is greater than  $3500\text{ cm}^{-1}$ .<sup>60</sup> In **4**, the emission of  $Eu^{3+}$  ions ( $^5D_0$ ,  $17241\text{ cm}^{-1}$ ) can be firstly sensitized by  $\{SeW_9\}$  building blocks ( $^3T_{1u}$ ,  $21168\text{ cm}^{-1}$ ).<sup>61</sup> After the addition of DPA, DPA can chelate with  $Eu^{3+}$  ions by substituting coordinate  $H_2O$  molecules, and the lowest triplet energy level ( $T_1$ ,  $25510\text{ cm}^{-1}$ ) of DPA can also allow the energy transfer to the  $Eu^{3+}$  ions, leading to the emission enhancement of  $Eu^{3+}$  ions.<sup>62</sup>

## Conclusions

In summary, a family of unique dual-ligand functionalized dodeca-nuclear Ln–W-cluster incorporated STs **1–4** have been successfully prepared by the cooperative strategy of rigid and flexible ligands. It is noteworthy that **4** can be used as a FL sensor to detect DPA in tap water and a human serum sample, which is also the first time to use the Ln-incorporated ST as a FL probe for the detection of DPA in tap water and a human serum sample. The results demonstrate that **4** can sensitively detect DPA with a low detection limit ( $3.83\text{ }\mu\text{M}$ ), good selectivity and anti-interference and quick response capability in the detection of DPA. This work not only provides a new synthetic strategy for the synthesis of rigid–flexible-ligand functionalized Ln-incorporated POMs, but also presents exploratory research on inorganic–organic hybrid Ln-incorporated POMs applied to fluorescence biological detection of an anthrax biomarker.

## Author contributions

Tiantian Gong, Sen Yang and Jiancai Liu performed synthesis, characterization, investigation of photoluminescence pro-

perties and wrote the manuscript. Zixu Wang, Mengyao Li and Siyu Zhang participated in PXRD, IR and TG characterization. Lijuan Chen and Junwei Zhao provided research ideas, determined crystal structures and revised the whole manuscript.

## Conflicts of interest

There are no conflicts to declare.

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