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## A switchable sensor and scavenger: detection and removal of fluorinated chemical species by a luminescent metal–organic framework†

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Fluorosis has been regarded as a worldwide disease that seriously diminishes the quality of life through skeletal embrittlement and hepatic damage. Effective detection and removal of fluorinated chemical species such as fluoride ions ( $F^-$ ) and perfluorooctanoic acid (PFOA) from drinking water are of great importance for the sake of human health. Aiming to develop water-stable, highly selective and sensitive fluorine sensors, we have designed a new luminescent MOF In(tcp) using a chromophore ligand 2,3,5,6-tetrakis(4-carboxyphenyl)pyrazine ( $H_4tcp$ ). In(tcp) exhibits high sensitivity and selectivity for turn-on detection of  $F^-$  and turn-off detection of PFOA with a detection limit of  $1.3 \mu\text{g L}^{-1}$  and  $19 \mu\text{g L}^{-1}$ , respectively. In(tcp) also shows high recyclability and can be reused multiple times for  $F^-$  detection. The mechanisms of interaction between In(tcp) and the analytes are investigated by several experiments and DFT calculations. These studies reveal insightful information concerning the nature of  $F^-$  and PFOA binding within the MOF structure. In addition, In(tcp) also acts as an efficient adsorbent for the removal of  $F^-$  ( $36.7 \text{ mg g}^{-1}$ ) and PFOA ( $980.0 \text{ mg g}^{-1}$ ). It is the first material that is not only capable of switchable sensing of  $F^-$  and PFOA but also competent for removing the pollutants via different functional groups.

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## Introduction

Fluoride ion ( $F^-$ ) is a kind of essential element of the human body and promote the formation of fluorapatite on the dental and skeletal surfaces at low concentration.<sup>1,2</sup> However, when  $F^-$  intake is beyond  $0.05 \text{ mg per kg per day}$ , it will become one of the most toxic and dangerous elements.<sup>3–6</sup> Excessive

consumption of  $F^-$  has been proven to be the main cause of dental and bone fluorosis.<sup>7–10</sup> Chronic absorption of an excess amount of  $F^-$  would cause gastric and kidney disorders and even death.<sup>11–13</sup>  $F^-$  can be easily absorbed by the human body when in contact with water, earth, and even air.<sup>14,15</sup> As the industrialization process continues to accelerate, more and more areas are suffering from fluorine pollution.<sup>16,17</sup> Perfluorooctanoic acid (PFOA), on the other hand, is a surfactant containing fifteen fluorine atoms. It has a high surface activity and is used in a variety of products, from surface protectants to polytetrafluoroethylene and insecticides.<sup>18–20</sup> Because of the high binding energy of carbon–fluorine (C–F) bonds, PFOA shows great resistance to natural biodegradation.<sup>21–25</sup> Via the food chain, PFOA can accumulate in the human body and shows immunological and neurological toxicity.<sup>26,27</sup> Hence, PFOA is regarded as one of the emerging contaminants by various organizations.<sup>28–30</sup> To avoid  $F^-$  and PFOA from being ingested by the human body, their monitoring, identification, and removal from drinking water are essential yet remain challenging.<sup>31</sup>

Some molecular compounds demonstrate turn-on/off and ratiometric luminescence detection for  $F^-$ .<sup>4,32,33</sup> While they show relatively higher sensitivity, complex procedures are usually required for the synthesis of these compounds. Metal–organic frameworks (MOFs), built by self-assembly processes, are regarded as an ideal platform for such sensing applications

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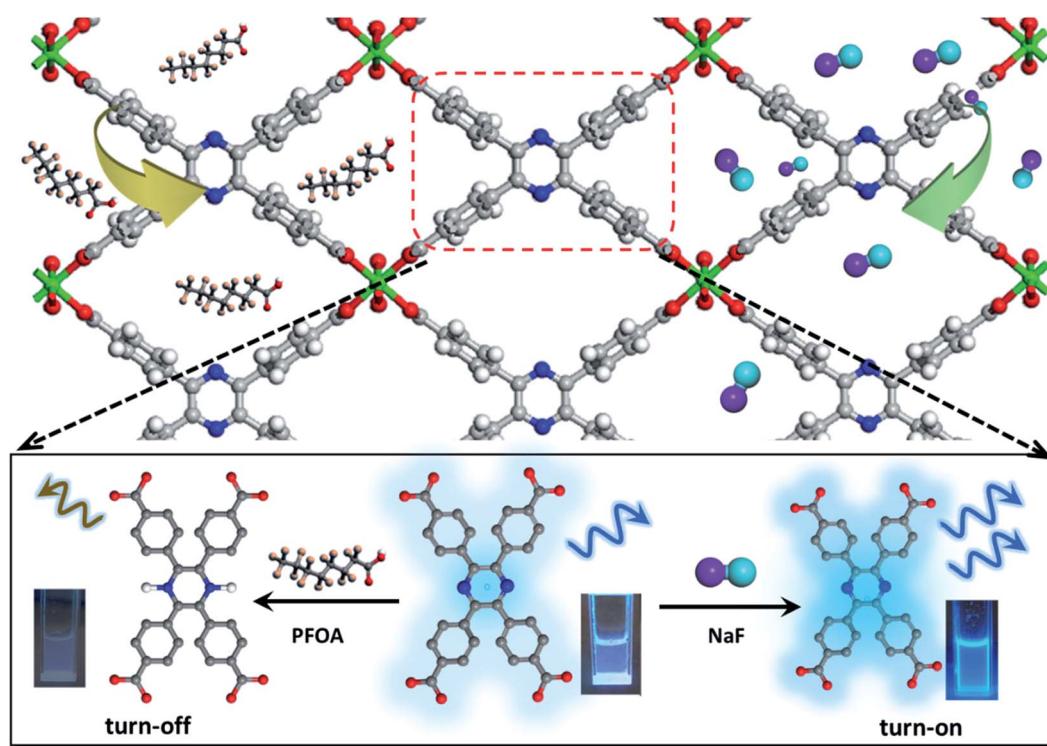
† Electronic supplementary information (ESI) available: Experimental details, DFT calculation details, PXRD, BET, TG, UV-vis data and luminescence spectra, tables as well as crystal structures and crystallographic data. CCDC 2057106 and 2057107. For ESI and crystallographic data in CIF or other electronic format see DOI: [10.1039/d1sc04070g](https://doi.org/10.1039/d1sc04070g)



due to their facile synthesis, high porosity and surface tunability which conventional molecular sensors are usually lacking.<sup>34</sup> To this date, there are only a few reports on  $F^-$  detection and adsorption by MOFs. For example, an europium-based MOF, Eu-MOF-1, with dual-emission is investigated for ratiometric detection of  $F^-$  ions with high selectivity and the limit of detection (LOD) is 2  $\mu\text{M}$ .<sup>35</sup> Another MOF, SION-105 with boron as the receptor site, shows a good response to  $F^-$  with high selectivity and a LOD of 0.1 ppm.<sup>1</sup> SION-105 was also used to develop an  $F^-$  detection device for convenient use. A guest encapsulated MOF,  $\text{NH}_2\text{-MIL-101(Al)}$  @ DCF, reaches a lower LOD for  $F^-$  (0.05  $\mu\text{M}$ ).<sup>36</sup> However, the  $F^-$  response of  $\text{NH}_2\text{-MIL-101(Al)}$  @ DCF requires addition of  $\text{H}_2\text{O}_2$ , making the process inconvenient. The dominant technique for PFOA analysis at the present time is HPLC-MS with high sensitivity, but it is time- and energy-consuming and must be carried out in professional laboratories with cumbersome instruments.<sup>37</sup> Recent studies have shown that luminescent materials are promising candidates for convenient and real-time PFOA detection. For example, a photoluminescence-based sensor made of molecularly imprinted polymer-coated quantum dots has been used for the enrichment and determination of PFOA in water with the LOD of 11.8 nM.<sup>22</sup> Being highly porous and luminescent, luminescent MOFs (LMOFs) have considerable potential for simultaneous pre-concentration, detection and removal of fluorine containing species. Often a single LMOF may be capable of detecting different kinds of pollutants with different response behaviors.

For solution-based sensing applications, LMOFs must possess high stability. The stability of MOFs is mainly determined by thermodynamic and kinetic factors.<sup>38</sup> The thermodynamic stability is consistent with the hard/soft acid/base (HSAB) principle. Hard bases, such as carboxylate-based ligands, can react with high-valence (hard acid) metals (*e.g.*  $\text{Ti}^{4+}$ ,  $\text{Zr}^{4+}$ ,  $\text{Al}^{3+}$ ,  $\text{Fe}^{3+}$  and  $\text{In}^{3+}$ ) to form stable MOFs.<sup>39</sup> Soft bases such as pyrazole can combine with low-valence (soft acid) metals such as  $\text{Ni}^{2+}$ ,  $\text{Co}^{2+}$  and  $\text{Zn}^{2+}$  to afford highly water stable MOFs.<sup>40</sup> For example, Li and co-workers developed a series of BUT-32 and BUT-33 (BUT = Beijing University of Technology) MOFs with remarkable water stability as a result of combination of soft base pyrazole and soft acid  $\text{Ni}^{2+}$ .<sup>41,42</sup> By use of hard acid ( $\text{Hf}^{4+}$  and  $\text{Zr}^{4+}$ ) and hard base carboxylate ligands PCN MOFs developed by Zhou and co-workers also demonstrate high water and acid/base stability.<sup>43-45</sup>

This work presents, for the first time, a new water stable LMOF for the sensitive detection of  $F^-$  and PFOA *via* a distinctly different signal change, as well as effective adsorption/removal of both species. The LMOF, simplified as  $\text{In}(\text{tcpp})$  or LMOF-651, was prepared by reacting  $\text{InCl}_3$  and 2,3,5,6-tetrakis(4-carboxyphenyl)pyrazine ( $\text{H}_4\text{tcpp}$ ) under solvothermal conditions.  $\text{In}(\text{tcpp})$  shows high porosity, water stability, and strong luminescence-based turn-on detection of  $F^-$  and turn-off detection of PFOA in aqueous solutions. LODs of 1.3  $\mu\text{g L}^{-1}$  and 19  $\mu\text{g L}^{-1}$  were achieved for  $F^-$  and PFOA, respectively (Scheme 1). The  $F^-$  detection process was fully recyclable. Furthermore,  $\text{In}(\text{tcpp})$  adsorbs a relatively large amount of  $F^-$  (36.7  $\text{mg g}^{-1}$ ) and PFOA (980.0  $\text{mg g}^{-1}$ ), demonstrating its



**Scheme 1** The structure of  $\text{In}(\text{tcpp})$  (top) and schematic presentation of its luminescence switchable turn-off response toward PFOA (bottom left) and turn-on response toward  $F^-$  ions (bottom right). Color codes: C grey; N blue; F light blue; O red; In green; Na purple; and H white.



strong potential for treating drinking water. IR spectroscopic analysis and DFT calculations were carried out to explore the mechanism of  $F^-$  and PFOA detection and adsorption.

## Results and discussion

### Synthesis, crystal structure and characterization

$H_4\text{tcpp}$  was synthesized according to the former work.<sup>46</sup>  $H_4\text{tcpp}$  and  $\text{InCl}_3$  were used to prepare  $\text{In}(\text{tcpp})$  by a solvothermal method in the presence of formic acid at 100 °C for 48 h (Fig. 1a and b). As a comparison,  $\text{In}(\text{tcpb})$  or LMOF-652 was also constructed with  $\text{InCl}_3$  and 1,2,4,5-tetrakis(4-carboxyphenyl)benzene ( $H_4\text{tcpb}$ ) using the same synthetic method (ESI† for details). The two In-MOFs are isostructural and differ only by the two atoms in the central ring of the ligands, which allowed us to compare and understand their similarities and differences in the sensing behavior toward fluorine-based chemical species.

The crystals of the resulting  $\text{In}(\text{tcpp})$  and  $\text{In}(\text{tcpb})$  are transparent, hexagonal-shaped rods (Fig. S1†). Single crystal X-ray diffraction analysis (Table S1†) reveals that  $\text{In}(\text{tcpp})$  crystallizes in an orthorhombic space group of  $Cmmm$  ( $a = 7.2195(3)$  Å,  $b = 22.5222(10)$  Å,  $c = 15.7802(7)$  Å) with a chemical formula of  $[\text{In}_2(\text{tcpp})(\text{OH})_2]$ . Each tcpp is connected to eight  $\text{In}^{3+}$  ions through  $\text{In}-\text{O}$  bonds to form an ordered one-dimensional rhombic channel, with a cross-section of about  $9.2 \times 13.4$  Å (Fig. 1b and c). The hexacoordinated  $\text{In}^{3+}$  ion bonds to four carboxylate oxygen atoms from different tcpp ligands, and two hydroxo oxygen atoms that connect the adjacent  $\text{In}^{3+}$  ions through the  $\text{In}-\text{OH}-\text{In}$  bonds (Fig. 1d).<sup>47,48</sup>  $\text{In}^{3+}$  ions are linked to each other through the tcpp ligands and bridging oxygen atoms to form a 1D chain. The PXRD analysis confirmed the

phase purity of  $\text{In}(\text{tcpp})$  (Fig. S2†). A comparison of the single crystal and PXRD data of  $\text{In}(\text{tcpb})$  verified that it is isoreticular to  $\text{In}(\text{tcpp})$  with the same space group topology (Fig. S3 and Table S2†).

$\text{In}(\text{tcpp})$  samples were activated by exchanging the solvents with acetone and then evacuated at 60 °C under vacuum for 24 h. No obvious differences were observed in the PXRD patterns of the as-made and activated  $\text{In}(\text{tcpp})$  samples, indicating high framework stability (Fig. S2†). Similar procedures were used to activate  $\text{In}(\text{tcpb})$ . Porosity characterization reveals type I adsorption behavior for both  $\text{In}(\text{tcpp})$  and  $\text{In}(\text{tcpb})$ , illustrating their microporous structures (Fig. S4a and S5a†). The Brunauer–Emmett–Teller (BET) surface area of  $\text{In}(\text{tcpp})$  and  $\text{In}(\text{tcpb})$  was estimated to be  $535 \text{ m}^2 \text{ g}^{-1}$  and  $534 \text{ m}^2 \text{ g}^{-1}$ , respectively. The pore size distribution analysis yielded a pore size  $\sim 6$  Å for both  $\text{In}(\text{tcpp})$  and  $\text{In}(\text{tcpb})$  (Fig. S4b and S5b†).

The thermal stability of both MOFs was evaluated. From the thermogravimetric analysis (TGA) curve, the as-made  $\text{In}(\text{tcpp})$  shows 21 wt% mass loss at 200 °C, which comes from the uncoordinated solvent molecules (Fig. S6†). For the activated  $\text{In}(\text{tcpp})$  sample, there was no weight loss until 200 °C, illustrating that the guest molecules were fully removed from the MOF channels.  $\text{In}(\text{tcpb})$  showed 25 wt% loss of guest solvent molecules at 210 °C (Fig. S7†). Both compounds were stable up to 400 °C. Upon heating at 200 °C for 2 h, the characteristic diffraction peaks of  $\text{In}(\text{tcpp})$  remained (Fig. S8†). The PXRD patterns kept unchanged after soaking the  $\text{In}(\text{tcpp})$  sample in water, in pH 2 (HCl) and pH 12 (NaOH) aqueous solutions for 24 h, demonstrating its excellent chemical stability. Also,  $\text{N}_2$  adsorption isotherms were collected after the sample was heated at 200 °C or treated under different acidity conditions at pH 2 (HCl) and pH 12 (NaOH), and the porosity remains the same (Fig. S9†), which further confirms the high stability of  $\text{In}(\text{tcpp})$ . The thermal stability of  $\text{In}(\text{tcpb})$  was found to be similar to that of  $\text{In}(\text{tcpp})$ . However, its resistance to the chemical environment was less than the former as it collapsed after being treated in pH 11 for 24 h (Fig. S10†).

The UV-vis spectra of  $\text{In}(\text{tcpp})$  and  $H_4\text{tcpp}$  ligand were collected at room temperature (Fig. S11†).  $H_4\text{tcpp}$  and  $\text{In}(\text{tcpp})$  show very similar absorption with a characteristic  $\pi \rightarrow \pi^*$  transition centered at 325 nm. As a result, the emission of the  $\text{In}(\text{tcpp})$  originates from the ligand. 3D luminescence spectra of  $\text{In}(\text{tcpp})$  and  $H_4\text{tcpp}$  were measured and the optimal excitation was observed at 280 nm (Fig. S12†). Under 280 nm excitation,  $\text{In}(\text{tcpp})$  emits at 400 nm with 30 nm blue shift compared to the free  $H_4\text{tcpp}$  (Fig. S13†). The blue shift of  $\text{In}(\text{tcpp})$  was attributed to the different structural torsion of the free  $H_4\text{tcpp}$  and tcpp fixed in the  $\text{In}(\text{tcpp})$ . As shown in Fig. S14a,† in the free  $H_4\text{tcpp}$ , the dihedral angle between the benzene ring and pyrazine core is approximately 20°. However, the dihedral angle of tcpp increases to approximately 56° to cater the rigid structure of  $\text{In}(\text{tcpp})$  (Fig. S14b†). The torsion of the benzene rings changes the  $\pi$ -conjugation of the whole ligand, resulting in the blue-shift emission.<sup>46</sup> The optical properties of  $\text{In}(\text{tcpb})$  were also measured and the results are shown in Fig. S15–S17.† The optimal excitation of 270 nm was selected to obtain the emission at 380 nm for  $\text{In}(\text{tcpb})$ .

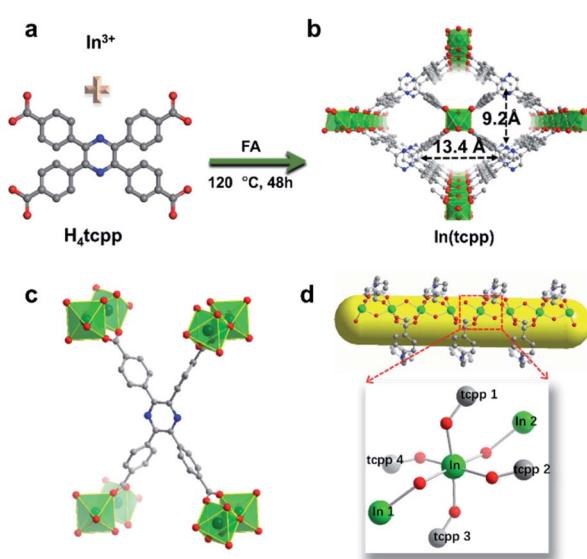


Fig. 1 Representation of the crystal structure of  $\text{In}(\text{tcpp})$ . (a) The organic and inorganic building units of  $\text{In}(\text{tcpp})$  and reaction conditions. (b) The one-dimensional channel in  $\text{In}(\text{tcpp})$ . (c) The coordination of the tcpp ligand. (d) The 6-coordinated  $\text{In}^{3+}$  ion and 1D  $\text{In}^{3+}$  chain along the  $a$ -axis. Color code: C grey; N blue; O red; In green; H atoms are omitted.

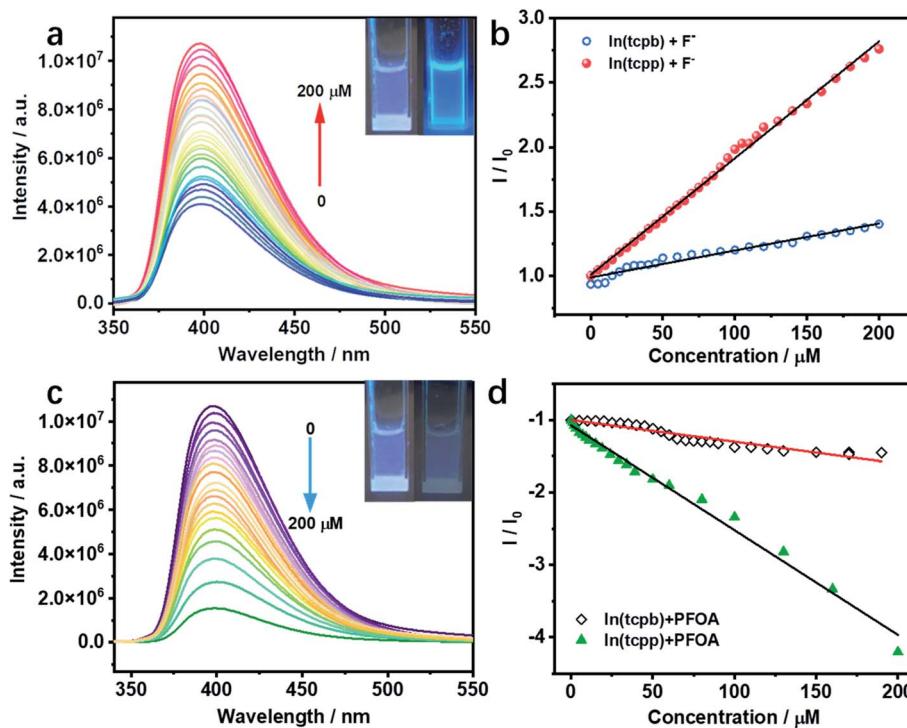


Fig. 2 (a) In(tcp) luminescence turn-on response to  $\text{F}^-$  at various concentrations. (b) The concentration-dependent luminescence response of In(tcp) and In(tcpb) to  $\text{F}^-$ . (c) In(tcp) luminescence turn-off response to PFOA at various concentrations. (d) The concentration-dependent luminescence response of In(tcp) and In(tcpb) to PFOA. Insets in (a) and (c): images of the In(tcp) solution samples before (left) and after (right) the addition of the analytes. The detection response time for (a)–(d) 5 min.

### Switchable detection and removal of $\text{F}^-$ and PFOA

Considering the interaction between guest analytes and functional groups of In(tcp), activated In(tcp) was selected for the detection experiments. In(tcp) remained stable after being soaked in 1 mM NaF aqueous solution for 24 h (Fig. S18†), illustrating that it is sufficiently stable for  $\text{F}^-$  detection. As shown in Fig. S19,† a drastic enhancement in the photoluminescence (PL) intensity was observed within  $\sim$ 5 min upon addition of NaF solution.

The results of the titration experiment are shown in Fig. 2a. Upon addition of  $\text{F}^-$ , the PL intensity of In(tcp) increased monotonically. The enhancement reached 250% at a concentration of 200  $\mu\text{M}$   $\text{F}^-$ . The turn-on efficiency was calculated by the Stern–Volmer (SV) eqn (1):

$$I_0/I = K_{\text{sv}}[\text{F}^-] + 1 \quad (1)$$

where  $I_0$  is the initial emission intensity of In(tcp),  $I$  is the emission intensity after the addition of  $\text{F}^-$ ,  $[\text{F}^-]$  is the molar concentration of  $\text{F}^-$ , and  $K_{\text{sv}}$  is the Stern–Volmer constant, which is used to evaluate the performance of In(tcp) as the sensor. The  $K_{\text{sv}}$  and LOD were determined to be  $9.06 \times 10^3 \text{ M}^{-1}$  and  $1.3 \mu\text{g L}^{-1}$ , respectively (Fig. 2b), which are better than those of most of the luminescent sensors (Table S3†). The response of low concentration  $\text{F}^-$  is illustrated in Fig. S20.† Compared to the optimal value of  $\text{F}^-$  concentration ( $1.5 \text{ mg L}^{-1}$ ) in drinking water set by the World Health Organization (WHO),

the LOD value achieved by In(tcp) has exceeded the WHO limit by three orders of magnitude ( $10^3$ ).<sup>49</sup>

In(tcp) is stable in PFOA solution as there was no obvious change in its PXRD patterns before and after exposure to a 1 mM PFOA solution for 24 h (Fig. S21†). Interestingly, In(tcp) shows monotonic quenching as a function of PFOA concentration (Fig. 2c). As depicted in Fig. 2d and S22,† In(tcp) responds to PFOA at very low concentrations with the  $K_{\text{sv}}$  and LOD of  $1.446 \times 10^4 \text{ M}^{-1}$  and  $19 \mu\text{g L}^{-1}$ , respectively.<sup>49</sup> It shows a comparative sensitivity as a PFOA sensor (Table S4†). In addition, it can act as an excellent scavenger for PFOA removal as discussed in the following section. However, the isostructural In(tcpb) shows little PL response to  $\text{F}^-$  and PFOA (Fig. 2b and d).

As stated earlier, excessive  $\text{F}^-$  is regarded as a water contaminant, while some common light ions, such as  $\text{Ca}^{2+}$ ,  $\text{Mg}^{2+}$  and  $\text{Cl}^-$ , are also found in drinking water and play important roles in biological processes. Therefore, selective detection of  $\text{F}^-$  ions over other harmless ions is of great importance. As shown in Fig. 3a and b, In(tcp) exhibits distinct selectivity toward  $\text{F}^-$  over other cations and anions although their concentrations are higher than  $\text{F}^-$ . The fast, selective, and turn-on response makes In(tcp) a promising sensor for easy and convenient detection of  $\text{F}^-$  ions. Further, the relative selectivity of In(tcp) was tested in different pH solutions (Fig. S23 and S24†). We found that at higher pH values (e.g. pH 10), strong luminescence enhancement was observed with addition of  $\text{F}^-$  and In(tcp) exhibits good selectivity toward  $\text{F}^-$ . While at lower pH (e.g. pH 3),  $\text{F}^-$  gives rise to very little change

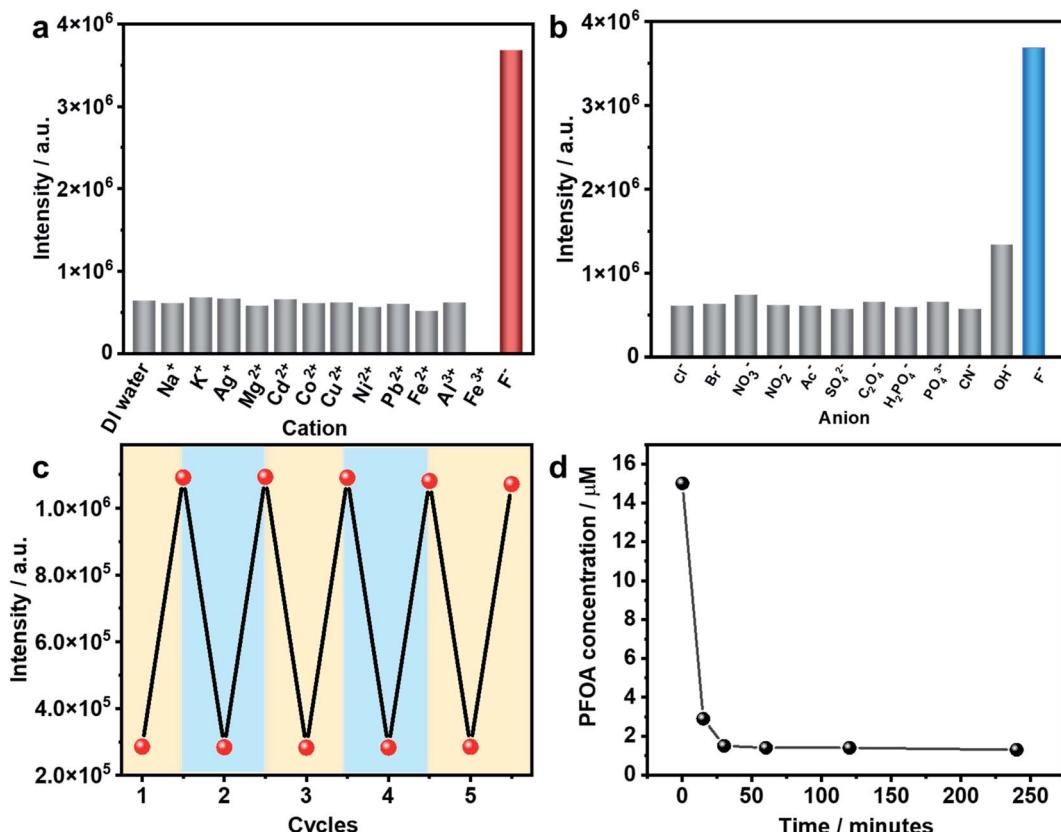


Fig. 3 Selectivity of In(tcpP) for (a) cations and (b) anions. In(tcpP)  $0.025 \text{ mg mL}^{-1}$ , the concentration of  $\text{F}^-$  is  $650 \mu\text{M}$  and other cations and anions are  $1 \text{ mM}$ . (c) The In(tcpP) luminescence turn-on and turn-off in 5 cycles. (d) The In(tcpP) PFOA removal efficiency in different pH

in the luminescence intensity. Moreover, in pH 10 solution (Fig. S24†), the sensitivity of  $\text{F}^-$  is similar to that in pure water.

Reusability is another important performance measure for a luminescence-based sensor. Considering the effective interaction between boric acid and  $\text{F}^-$ ,<sup>35</sup>  $\text{F}^-@\text{In(tcpP)}$  was dissolved in boric acid solutions of pH 3 to regenerate the sensor material. After being stirred and centrifuged three times, In(tcpP) was fully regenerated as its PL intensity reached the original level. The recovered In(tcpP) was then reused four times consecutively without apparent changes in its PL profile (Fig. 3c); the PXRD patterns show that In(tcpP) remained stable after 5 cycles (Fig. S25†), confirming its good recyclability. Taking into consideration its high porosity and stability, In(tcpP) was further used to remove  $\text{F}^-$  ions. An uptake capacity of  $36.7 \text{ mg g}^{-1}$  (original concentration  $100 \text{ ppb}$ , remaining concentration  $1.2 \text{ ppb}$ ) was achieved for this scavenger, a value that is higher than those of most of the  $\text{F}^-$  removal materials, such as graphene, mesoporous alumina, and hydrous ferric oxide.<sup>50</sup>

The PFOA removal performance was assessed in different pH solutions. As shown in Fig. S26,† In(tcpP) adsorbs  $980 \text{ mg g}^{-1}$  PFOA in acidic solution (pH 3, original concentration  $100 \text{ ppb}$ , remaining concentration  $2.14 \text{ ppb}$ ), which is higher than those of most of the porous materials (such as mesoporous silica, activated carbons, activated carbons, molecularly imprinted polymers, etc.) reported to date.<sup>51</sup> We find that 90% of PFOA (at

a concentration of  $15 \mu\text{M}$ ) can be removed within 30 min (Fig. 3d). Thus, strong evidence confirms that In(tcpP) is a good candidate not only for the switchable detection of PFOA and  $\text{F}^-$ , but also for their capture and removal.

#### MOF-analyte interaction mechanism

To unveil the MOF-analyte interaction mechanism, we turned to DFT calculation and infrared/X-ray photoelectron spectroscopy to examine the binding of PFOA and NaF in In(tcpP). We calculated the removal efficiency based on XPS and HPLC-MS methods, which allow for the determination of the quantity of adsorbed  $\text{F}^-$  and PFOA. The EPR and NMR experiments were carried out before and after adding NaF into the free ligand  $\text{H}_4\text{tcpP}$ .<sup>52</sup> As shown in Fig. S27 and S28,† there are no obvious differences before and after the NaF addition, suggesting that no charge/electron transfer interactions occurred between NaF and  $\text{H}_4\text{tcpP}$ . Based on the UV-vis and PL spectra (Fig. S29†), small and gradual increase in the absorption intensity was observed as a function of time, while an appreciable enhancement in the PL intensity was noted after 30 min. These data indicate that the strong basicity of NaF promotes the dissolution of the ligand giving rise to the increases in the UV-vis and PL intensity.

We then studied the configuration of  $\text{F}^-$  in In(tcpP) via XPS. The content of  $\text{F}^-$  is much higher than that of  $\text{Na}^+$  as shown

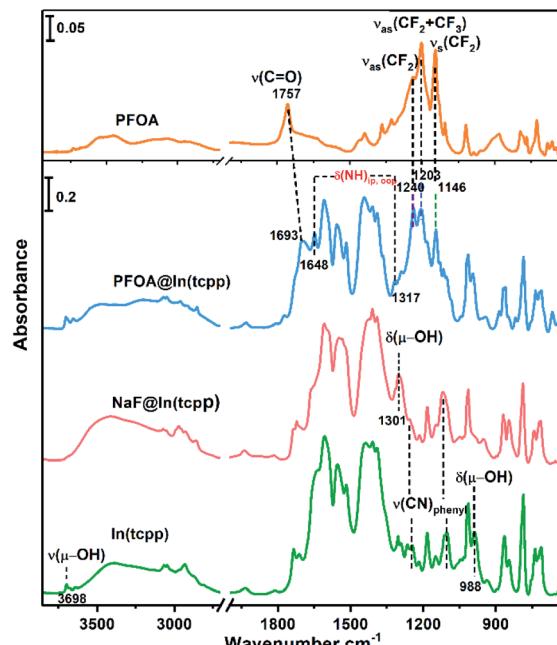


Fig. 4 FTIR spectra of free PFOA, PFOA@In(tcp), NaF@In(tcp), and In(tcp) (from top to bottom), referenced to the blank KBr pellet.

from XPS results (Fig. S30 and Table S5†). We speculate that  $\text{F}^-$  exists in the form of both  $\text{NaF}$  and  $\text{HF}$  within the sample, but the latter is the majority species. We thus analyzed the favorable binding site of  $\text{HF}$  within In(tcp), which is near  $\text{N}_{\text{tcp}}$  (Fig. S31a†). The  $\text{H}^+$  of  $\text{HF}$  is  $1.733 \text{ \AA}$  away from the  $\text{N}_{\text{tcp}}$  and the  $\text{H}-\text{F}$  bond length is  $0.975 \text{ \AA}$ , which is slightly elongated compared to our calculated gas-phase bond length of  $0.938 \text{ \AA}$ . The binding configuration predicted computationally is in good agreement with what we observed in the IR spectra (Fig. 4), namely, significant perturbations of the ring mode, *e.g.*, blue shift of the  $\nu(\text{CN})_{\text{pyrazine}}$  band is observed in IR spectra, indicating the direct interaction of  $\text{HF}$  with the tcp linker. In addition, no amine deformation bands  $\delta(\text{NH})_{\text{ip,oop}}$  were detected because  $\text{H}$  stays closer to  $\text{F}$  without protonating the  $\text{N}$  site on the organic linker, in perfect agreement with our theoretical predictions. Moreover, the blue shift of the  $\nu(\text{CN})_{\text{pyrazine}}$  band as observed in our IR spectra is mimicked with the induced charge density calculations where a strong interaction of  $\text{H}@\text{HF}$  with the  $\text{N}@\text{tcp}$  linker is clearly visible (Fig. S31b†).

We further assessed the possibility of the minority species  $\text{NaF}$  interacting directly with the  $\text{N}$  sites in In(tcp), and found that they are unfavorable based on the estimated binding energetics ( $1.35 \text{ eV}$  between  $\text{Na}^+$  and  $\text{N}_{\text{tcp}}$ ) (Fig. S32†). Further, numerous starting configurations of  $\text{Na}^+$ ,  $\text{F}^-$ , and  $\text{Na}^+\text{F}^-$  interacting with the tcp ligand or In node were considered. We found that after the  $\text{F}^-$  adsorption, the coordination of  $\text{F}^-$  at the axial position of In *via*  $\mu_{\text{H-O}}$  displacement is energetically unfavourable. Instead, the most energetically favourable model for  $\text{NaF}$  binding is that  $\text{Na}^+$  interacts with oxygen on the  $\text{In-O}_{\text{tcp}}$ , and the  $\text{F}^-$  interacts with both  $\text{OH}$  and  $\text{Na}$ , based on their charge rearrangement (Fig. 5a and b). This is consistent with the infrared spectroscopic results, which suggest that  $\text{NaF}$  most

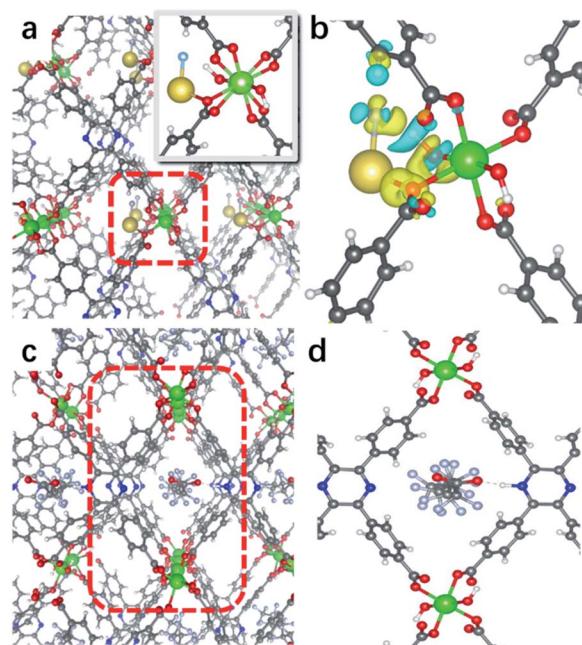


Fig. 5 (a) Optimized structure of  $\text{NaF}@\text{In}(\text{tcp})$ , highlighting  $\text{Na}^+$  and  $\text{F}^-$  interactions. (b) Charge density rearrangement upon formation of the bond at an isolevel of  $0.002 \text{ eV \AA}^{-3}$ . (c) Binding of PFOA and In(tcp) in  $\text{PFOA}@\text{In}(\text{tcp})$ . (d) Configuration of  $\text{PFOA}@\text{In}(\text{tcp})$  (atoms are colored as follows: C grey, O red, N blue, In green, F light blue, Na yellow, H white).

likely binds in close proximity to the  $\text{COO}$  and  $\text{OH}$  groups (Fig. 4). Firstly, the bridging  $\text{OH}$  stretching band  $\nu(\mu-\text{OH})$  at  $3698 \text{ cm}^{-1}$  and bending band  $\beta(\text{OH})$  at  $983 \text{ cm}^{-1}$  disappear. Additionally, a new band is observed at  $1301 \text{ cm}^{-1}$ . DFT calculation has ascribed this new band to the shifted  $\beta(\text{OH})$  vibration upon forming H-bonding interaction with  $\text{F}^-$  (Table S6†). Furthermore, a decrease of the intensity of stretching bands  $\nu(\text{COO})$  is detected in the  $\text{NaF}@\text{In}(\text{tcp})$  spectrum, giving an indication that  $\text{NaF}$  interacts with the  $\text{COO}$  group. The photoluminescence enhancement was not captured, possibly due to exciton formation upon the photoexcitation.

Based on the titration PFOA turn-off results, we tested the model of PFOA binding to the nitrogen of the linker through its  $\text{H}^+$ , forming  $\text{PFOA}^-$ . Several other binding sites were also examined but were found to be less favorable and, therefore, are not part of this discussion. In the  $\text{PFOA}@\text{In}(\text{tcp})$ , we found that hydrogen binds more favorably to the nitrogen of the linker, but is still close enough to the  $\text{PFOA}^-$  molecule to form a hydrogen bond (Fig. 5c and d) with a strong binding energy of  $1.57 \text{ eV}$ . This is in accordance with an approximately  $1:1$  ratio of  $\text{N}_{\text{tcp}}:\text{PFOA}$ , calculated based on our HPLC-MS results of  $980.0 \text{ mg g}^{-1}$  PFOA adsorption (see the characteristic absorption peak and standard curve of PFOA in Fig. S33†). The protonation of N is evidenced by the appearance of two new bands at  $1648$  and  $1317 \text{ cm}^{-1}$  as well as a red frequency shift of the  $\nu(\text{C=O})$  band from  $1757 \text{ cm}^{-1}$  to  $1693 \text{ cm}^{-1}$  (Fig. S34†). Frequency modeling by DFT shows that these two bands at  $1648$  and  $1317 \text{ cm}^{-1}$  can be assigned to in-plane and out-of-plane deformation bands of the  $-\text{NH}$  group, respectively. The so

formed  $-\text{NH}_{\text{tcpp}}$  group can further interact with the PFOA's carboxylate group by hydrogen bonding interaction, resulting in an observed frequency shift of the  $\nu(\text{C}=\text{O})$  band to  $1693\text{ cm}^{-1}$  (Fig. 4). The protonation of N is evident from the Raman spectra (Fig. S35†). After interacting with PFOA, the  $\nu(\text{N}-\text{H})$  band is observed at  $1568\text{ cm}^{-1}$ . Further, the hydrogen bonding between  $\text{N}_{\text{tcpp}}$  and PFOA blueshifts the  $\nu(\text{C}-\text{O})$  from  $1148\text{ cm}^{-1}$  to  $1145\text{ cm}^{-1}$ . Our theoretical and spectroscopic results show a distinct binding mechanism due to the presence of N in the organic component and the acid–base interaction between the PFOA molecule and In(tcpb) framework is the driving force for PFOA detection and adsorption (Fig. S36†). Using these models and comparing them to the pristine In(tcpb) MOF, we calculated the photoluminescence quenching of PFOA and  $\text{PFOA}^-$  using our previously developed methodology.<sup>53,54</sup> As can be seen in Fig. S37,† PFOA shows strong quenching of the photoluminescence in  $\text{PFOA}@\text{In}(\text{tcpb})$  (92%).

## Conclusions

In summary, we have designed and prepared a robust and porous LMOF, In(tcpb). In(tcpb) acts as a "switchable" sensor, showing a strong turn-on and turn-off PL signal when exposed to  $\text{F}^-$  anions and PFOA. Its limits of detection for  $\text{F}^-$  ions and PFOA are  $1.3\text{ }\mu\text{g L}^{-1}$  and  $19\text{ }\mu\text{g L}^{-1}$ , respectively, indicating high sensitivity towards both chemical species. In(tcpb) also shows excellent selectivity for  $\text{F}^-$  over other cations and anions commonly found in drinking water. The  $\text{F}^-$  detection process is fully recyclable and the sensor material is reusable. Moreover, In(tcpb) can be used as an effective scavenger to capture and remove  $\text{F}^-$  and PFOA from aqueous solution. FTIR analysis and DFT calculations were performed to explore the MOF–analyte interaction mechanism: in the case of PFOA, the  $\text{N}_{\text{tcpp}}$  is the site of interaction that is responsible for the quantitative adsorption of PFOA; the maximum loading amount is consistent with the number of available binding sites. The interaction between  $\text{F}^-$  and In(tcpb) occurs primarily at the bridging  $-\text{OH}^-$ , which is also consistent with the In/F ratio. This work demonstrates that switchable detection and removal of similar pollutants by the same MOF may be achieved and may serve as an effective approach in the future design of high performance MOF-based chemical sensors and scavengers.

## Experimental

### Preparation of compounds

**Synthesis of In(tcpb).**  $\text{InCl}_3$  (55 mg, 0.25 mmol) and  $\text{H}_4\text{tcpp}$  (50 mg, 0.09 mmol) were dissolved in a mixture solution of DMA (5 mL) and water (2.5 mL). And then 0.8 mL of formic acid was added to the above solution. The as-obtained mixture was transferred to a 25 mL glass vial. After being sonicated for 5 min, the mixture was maintained at  $100\text{ }^\circ\text{C}$  for 48 h under static conditions. After naturally cooling to room temperature, the solution was removed by filtration and the product was washed with DMA to remove the residual  $\text{H}_4\text{tcpp}$  and  $\text{In}^{3+}$  ions. Finally, the samples were dried at  $60\text{ }^\circ\text{C}$  for 12 h.

**Synthesis of In(tcpb).** The In(tcpb) synthesis method is similar to that of In(tcpb) except for replacing  $\text{H}_4\text{tcpp}$  with  $\text{H}_4\text{tcpb}$  (50 mg, 0.09 mmol). After cooling to room temperature, the solid products were filtered and washed with DMA to remove the residual  $\text{H}_4\text{tcpb}$  and  $\text{In}^{3+}$  ions. Finally, In(tcpb) was dried at  $60\text{ }^\circ\text{C}$  for 12 h.

**Detection of  $\text{F}^-$  or PFOA.** The activated In(tcpb) or In(tcpb) crystals were ground and dispersed in water and kept under sonication until getting the uniform suspension. Luminescence titrations were carried out by incrementally adding aliquots of  $\text{F}^-$  or PFOA solutions to the In(tcpb) or In(tcpb) suspension followed by sonication. The luminescence spectra were recorded 5 min after adding the appropriate amount of  $\text{F}^-$  or PFOA stock solution. This exposure time ensured that the analyte entered the LMOF pores. Each measurement was repeated three times, and the average value was used. Note: to get the obvious turn on/off response to  $\text{F}^-$  or PFOA, the concentration of In(tcpb) is  $0.05\text{ mg mL}^{-1}$  for turn-on, and In(tcpb)  $0.005\text{ mg mL}^{-1}$  for turn-off.

**Limit of detection for  $\text{F}^-$  and PFOA.** According to the International Union of Pure and Applied Chemistry (IUPAC), the limit of detection was calculated using the following equation (eqn (2)).

$$\text{LOD} = kS_b/a \quad (2)$$

where LOD is the limit of detection,  $k = 3$  for spectral chemical analysis, and  $a$  is the sensitivity of the sensor (*i.e.* the slope of the calibration curve). The confidence level is 90% corresponding to  $k = 3$ .  $S_b$  is the standard deviation of the blank background.

**Recyclability test for  $\text{F}^-$  detection.** A  $200\text{ }\mu\text{M}$   $\text{F}^-$  was added into the In(tcpb) suspension ( $90.5\text{ mg mL}^{-1}$ ) and the luminescence intensity was recorded. The above suspension was washed with pH 3 boric acid and centrifuged three times to remove the  $\text{F}^-$ . Then the  $\text{F}^-$  was added into the In(tcpb) suspension for the second use. Before and after each cycle the In(tcpb) PXRD was measured to make sure that the In(tcpb) remained stable for further use.

**NaF removal efficiency analysis.** Stock solutions of  $0.5\text{ }\mu\text{M}$  NaF were prepared. 10 mg activated In(tcpb) was added into the above NaF aqueous solutions. After 30 min, In(tcpb) was centrifuged and washed three times with methanol. The XPS analysis for  $\text{F}^-$  and  $\text{Na}^+$  was performed after the above materials were dried for 12 h in a vacuum.

**PFOA removal efficiency analysis.** Stock solutions of  $15\text{ }\mu\text{M}$  PFOA were prepared by dissolving PFOA into 100 mL ultrapure deionized water. 3 mg activated In(tcpb) was added into the above PFOA aqueous solutions. The supernatant PFOA concentration was obtained at 0, 15, 30, 60, 120, and 240 min. The concentration of each sample was calculated according to the calibration curve obtained from HPLC-MS.

To estimate the PFOA removal efficiency of In(tcpb) in solutions of different pH, 55 mg In(tcpb) was added into 10 mL PFOA in different pH buffer (15 mM). The samples were sonicated to ensure good mixing. After 30 min, the concentration of supernatant PFOA was obtained after centrifugation.



**HPLC-MS analysis of PFOA.** The above supernatant PFOA concentration was determined by using an HPLC interfaced with an electrospray mass spectrometer. HPLC separation was performed on a Thermo Scientific UltiMate 3000 HPLC-QE high performance liquid chromatograph system. 20 micro liters of the supernatant was injected onto a Waters BEH C18 (100 mm length  $\times$  2.1 mm i.d., 1.7  $\mu$ m packing) with a 20 mm guard column of the same material. A deionized water solution containing 10 mM ammonium acetate and methanol as the mobile phase was delivered with a flow-rate of 300  $\mu$ L min $^{-1}$ . The gradient started at 90% methanol followed by 2 min ramp to 10% methanol, and then another 5.5 min ramp to 10% methanol, and then reverting to initial conditions allowing 6 min stabilization time. Detection was performed with an Agilent 1100 MSD ion-trap-SL system (Agilent, Palo Alto, CA, USA) operating in the electrospray negative mode. The optimized conditions were as follows: nebulizer gas pressure 30 psi, drying nitrogen gas temperature 350 °C, capillary voltage 3.2 kV. The capillary exit voltages were  $-122$  V for PFOA. Quantitation was monitored using  $[\text{M}-\text{H}]^-$  ions for PFOA in the single MS mode by relating the area of the analyte to the area of the standard line.

## Data availability

All experimental supporting data are provided in the ESI.†

## Author contributions

H. Yin and J. Li designed the experiments. H. Yin performed the syntheses, measurements and detection on the compounds. K. Tan, S. Jensen, S. Ullah, K. Oyekan, N. Meyer and T. Thonhauser performed the FTIR measurements and DFT calculations. K. Tan organized the detection and removal mechanism. S. Teat performed single X-ray diffraction analyses. X. Hei, E. Velasco and X. Wang gave suggestions and helped in the whole process. H. Yin wrote the first draft of the manuscript, which was subsequently edited by J. Li and X. Yin. All authors discussed the results and commented on the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 F. M. Ebrahim, T. N. Nguyen, S. Shyshkanov, A. Gladysiak, P. Favre, A. Zacharia, G. Itskos, P. J. Dyson and K. C. Stylianou, *J. Am. Chem. Soc.*, 2019, **141**, 3052–3058.
- 2 J. D. B. Featherstone, *Community Dent. Oral Epidemiol.*, 1999, **27**, 31–40.
- 3 Z. Kheradpisheh, M. Mirzaei, A. H. Mahvi, M. Mokhtari, R. Azizi, H. Fallahzadeh and M. H. Ehrampoush, *Sci. Rep.*, 2018, **8**, 2674.
- 4 T. K. Mandal, Y. Hou, Z. Gao, H. Ning, W. Yang and M. Gao, *Adv. Sci.*, 2016, **3**, 1600217.
- 5 O. Barbier, L. Arreola-Mendoza and L. M. Del Razo, *Chem.-Biol. Interact.*, 2010, **188**, 319–333.
- 6 A. Bontemps, L. Conquet, C. Elie, V. Magneron, C. Gloaguen, D. Kereselidze, K. Tack, O. C. Barbier and Y. Gueguen, *Int. J. Environ. Res. Public Health*, 2019, **16**, 1136–1156.
- 7 Y. Liu, A. Jiang, Q. Jia, X. Zhai, L. Liu, L. Ma and J. Zhou, *Chem. Sci.*, 2018, **9**, 5242–5251.
- 8 B. Ke, W. Chen, N. Ni, Y. Cheng, C. Dai, H. Dinh and B. Wang, *Chem. Commun.*, 2013, **49**, 2494–2496.
- 9 S. Ayoob and A. K. Gupta, *Crit. Rev. Environ. Sci. Technol.*, 2006, **36**, 433–487.
- 10 S. Roy and G. Dass, *Resour. Environ.*, 2013, **3**, 53–58.
- 11 M.-L. Cittanova, B. Lelongt, M.-C. Verpont, M. Geniteau-Legendre, F. Wahbe, D. Prie, P. Coriat and P. M. Ronco, *Anesthesiology*, 1996, **84**, 428–435.
- 12 M. I. Jimenez-Cordova, C. Gonzalez-Horta, J. C. Ayllon-Vergara, L. Arreola-Mendoza, G. Aguilar-Madrid, E. E. Villareal-Vega, A. Barrera-Hernandez, O. C. Barbier and L. M. Del Razo, *Environ. Res.*, 2019, **169**, 220–228.
- 13 Y. Tian, H. Che, J. Wang, D. Wang, L. Yang, L. Wang, Y. Nie and X. Tian, *J. Hazard. Mater.*, 2021, **411**, 125182.
- 14 A. Rasool, A. Farooqi, T. Xiao, W. Ali, S. Noor, O. Abiola, S. Ali and W. Nasim, *Environ. Geochem. Health*, 2018, **40**, 1265–1281.
- 15 K. J. Barmao, J. K. Cherutoi, C. Y. Mitei, M. L. L. Were, A. Kiprop and O. G. Achieng', *J. Environ. Manage.*, 2019, **8**, 15–24.
- 16 Q. Zhang, S. Bolisetty, Y. Cao, S. Handschin, J. Adamcik, Q. Peng and R. Mezzenga, *Angew. Chem., Int. Ed.*, 2019, **58**, 6012–6016.
- 17 F. Ahmadijokani, H. Molavi, M. Rezakazemi, T. M. Aminabhavi and M. Arjmand, *Coord. Chem. Rev.*, 2021, **445**, 214037.
- 18 L. Zheng, Y. Zheng, Y. Liu, S. Long, L. Du, J. Liang, C. Huang, M. T. Swihart and K. Tan, *Talanta*, 2019, **194**, 1–6.
- 19 K.-U. Goss, *Environ. Sci. Technol.*, 2008, **42**, 456–458.
- 20 Y. Wang, P. B. Groszewicz, S. Rosenfeldt, H. Schmidt, C. A. Volkert, P. Vana, T. Gutmann, G. Bunkowsky and K. Zhang, *Adv. Mater.*, 2017, **29**, 1702473.
- 21 R. C. Buck, J. Franklin, U. Berger, J. M. Conder, I. T. Cousins, P. de Voogt, A. A. Jensen, K. Kannan, S. A. Mabury and



S. P. van Leeuwen, *Integr. Environ. Assess. Manage.*, 2011, **7**, 513–541.

22 Z. Cheng, L. Du, P. Zhu, Q. Chen and K. Tan, *Spectrochim. Acta, Part A*, 2018, **201**, 281–287.

23 C. Fang, X. Zhang, Z. Dong, L. Wang, M. Megharaj and R. Naidu, *Chemosphere*, 2018, **191**, 381–388.

24 L. Liu, Y. Liu, N. Che, B. Gao and C. Li, *J. Hazard. Mater.*, 2021, **416**, 125866.

25 J. Wang, C. Cao, Y. Zhang, Y. Zhang and L. Zhu, *Appl. Catal. B*, 2021, **286**, 119911.

26 X. Li, L. W. Y. Yeung, S. Taniyasu, M. Li, H. Zhang, D. Liu, P. K. S. Lam, N. Yamashita and J. Dai, *Environ. Sci. Technol.*, 2008, **42**, 7078–7083.

27 L. Xiang, L. Chen, L. Y. Yu, P. F. Yu, H. M. Zhao, C. H. Mo, Y. W. Li, H. Li, Q. Y. Cai, D. M. Zhou and M. H. Wong, *Sci. Total Environ.*, 2018, **636**, 999–1008.

28 F. Riva, S. Castiglioni, E. Fattore, A. Manenti, E. Davoli and E. Zuccato, *Int. J. Hyg. Environ. Health*, 2018, **221**, 451–457.

29 J. M. Judge, P. Brown, J. G. Brody and S. Ryan, *J. Health Soc. Behav.*, 2016, **57**, 333–350.

30 B. Vrana, F. Smedes, R. Prokeš, R. Loos, N. Mazzella, C. Miege, H. Budzinski, E. Vermeirssen, T. Ocelka, A. Gravell and S. Kaserzon, *TrAC, Trends Anal. Chem.*, 2016, **76**, 153–165.

31 J. Zhang, J. Qian, Q. Mei, L. Yang, L. He, S. Liu, C. Zhang and K. Zhang, *Biosens. Bioelectron.*, 2019, **128**, 61–67.

32 F. M. Hinterholzinger, B. Ruhle, S. Wuttke, K. Karaghiosoff and T. Bein, *Sci. Rep.*, 2013, **3**, 2562.

33 M. Hirai and F. P. Gabbaï, *Chem. Sci.*, 2014, **5**, 1886–1893.

34 B. Chen, L. Wang, F. Zapata, G. Qian and E. B. Lobkovsky, *J. Am. Chem. Soc.*, 2008, **130**, 6718–6719.

35 Z.-R. Yang, M.-M. Wang, X.-S. Wang and X.-B. Yin, *Anal. Chem.*, 2017, **89**, 1930–1936.

36 Y. Sun, X. Xu, Y. Zhao, H. Tan, Y. Li and J. Du, *Talanta*, 2020, **209**, 120582.

37 P. Zhu, X. Ling, W. Liu, L. Kong and Y. Yao, *J. Chromatogr. B*, 2016, **1031**, 61–67.

38 X. Liu, X. Wang and F. Kapteijn, *Chem. Rev.*, 2020, **120**, 8303–8377.

39 S. Yuan, L. Feng, K. Wang, J. Pang, M. Bosch, C. Lollar, Y. Sun, J. Qin, X. Yang, P. Zhang, Q. Wang, L. Zou, Y. Zhang, L. Zhang, Y. Fang, J. Li and H. C. Zhou, *Adv. Mater.*, 2018, **30**, 1704303.

40 X. L. Lv, K. Wang, B. Wang, J. Su, X. Zou, Y. Xie, J. R. Li and H. C. Zhou, *J. Am. Chem. Soc.*, 2017, **139**, 211–217.

41 T. He, X. J. Kong, J. Zhou, C. Zhao, K. Wang, X. Q. Wu, X. L. Lv, G. R. Si, J. R. Li and Z. R. Nie, *J. Am. Chem. Soc.*, 2021, **143**, 9901–9911.

42 T. He, Z. Huang, S. Yuan, X. L. Lv, X. J. Kong, X. Zou, H. C. Zhou and J. R. Li, *J. Am. Chem. Soc.*, 2020, **142**, 13491–13499.

43 B. Wang, X. L. Lv, D. Feng, L. H. Xie, J. Zhang, M. Li, Y. Xie, J. R. Li and H. C. Zhou, *J. Am. Chem. Soc.*, 2016, **138**, 6204–6216.

44 B. Wang, P. Wang, L. H. Xie, R. B. Lin, J. Lv, J. R. Li and B. Chen, *Nat. Commun.*, 2019, **10**, 3861.

45 W. Bin, L. Xiu-Liang, F. Dawei, X. Lin-Hua, Z. Jian, L. Ming, X. Yabo, L. Jian-Rong and Z. Hong-Cai, *J. Am. Chem. Soc.*, 2016, **138**, 6204–6216.

46 H. Q. Yin, X. Y. Wang and X. B. Yin, *J. Am. Chem. Soc.*, 2019, **141**, 15166–15173.

47 M. D. DeFuria, M. Zeller and D. T. Genna, *Cryst. Growth Des.*, 2016, **16**, 3530–3534.

48 L. Liu, S.-M. Wang, Z.-B. Han, M. Ding, D.-Q. Yuan and H.-L. Jiang, *Inorg. Chem.*, 2016, **55**, 3558–3565.

49 A. H. Mahvi, M. A. Zazoli, M. Younecian, B. Nicpour and A. Babapour, *J. Med. Sci.*, 2006, **6**, 658–661.

50 G. K. Sarma and M. H. Rashid, *J. Chem. Eng. Data*, 2018, **63**, 2957–2965.

51 Z. Du, S. Deng, Y. Bei, Q. Huang, B. Wang, J. Huang and G. Yu, *J. Hazard. Mater.*, 2014, **274**, 443–454.

52 H. C. Wentz, G. Skorupskii, A. B. Bonfim, J. L. Mancuso, C. H. Hendon, E. H. Oriel, G. T. Sazama and M. G. Campbell, *Chem. Sci.*, 2019, **11**, 1342–1346.

53 S. Jensen, K. Tan, W. P. Lustig, D. S. Kilin, J. Li, Y. J. Chabal and T. Thonhauser, *Chem. Mater.*, 2019, **31**, 7933–7940.

54 S. Jensen, K. Tan, W. Lustig, D. Kilin, J. Li, Y. J. Chabal and T. Thonhauser, *J. Mater. Chem. C*, 2019, **7**, 2625–2632.

