



 Cite this: *RSC Adv.*, 2017, 7, 50035

Highly sensitive and selective fluorescent probe for Fe³⁺ and hazardous phenol compounds based on a water-stable Zn-based metal–organic framework in aqueous media†

 Huijun Li, Yaling He, Qingqing Li, Shaojie Li, Zhihao Yi, Zhouqing Xu * and Yuan Wang*

A novel two-fold interpenetrating network, namely, {[Zn(DPTMIA)]·(H₂O)₂(DMF)_{0.5}}_n (HPU-1), with rectangular channels was synthesized and structurally characterized. HPU-1 was shown to be potentially applicable as a multi-functional fluorescent probe. It was shown to specifically detect Fe³⁺, through a cation exchange mechanism, and hazardous phenol compounds, using a mechanism involving a decrease in the efficiency of the transfer of electrons and energy from the ligands to the metal ions. The detection limits of HPU-1 for Fe³⁺ and the phenol compounds were found to be lower than those of MOFs reported in the literature. This HPU-1 probe was shown to be functional in aqueous solution and to be affected neither by the type of solvent nor the pH.

Received 31st July 2017

Accepted 11th October 2017

DOI: 10.1039/c7ra08427g

rsc.li/rsc-advances

Introduction

Fe³⁺ is ubiquitous in cells, and thus plays a crucial role in the human body and other biological tissues. A deficiency and excess of Fe³⁺ from the normal permissible limit may induce serious disorders.¹ Therefore, the detection of iron has gained increasing interest in recent years.² Recently, the design of fluorophore-based Fe³⁺-sensing materials has been actively pursued because of the safety, operational simplicity, humanitarian implications and non-destructive character of these materials.³ However, these materials do show some problems, such as low sensitivity and poor stability. Therefore, synthesizing novel fluorescent probes for the detection of metal ions or small organic molecules is a challenging yet important task.

Fluorescent probes based on the metal–organic framework (MOF) have been widely investigated for the selective sensing of metal ions or toxic organic complexes on account of their visible luminescence and designable architectures, which have allowed for improved host–guest interactions and pre-concentrators for target analytes.⁴ Major research has focused on the lanthanide MOFs due to their high luminous efficiency, long-wavelength emission and lifetime.⁵ However, there are some defects existing in the synthesis process of Ln-MOFs, such as: expensive raw materials, secondary pollution and so on. Therefore, MOFs

based on transition metal ions, particularly Zn²⁺, have captured our interest because of their advantages, including their environmental compatibility, low cost, accessibility and unimodal emission.⁶ In addition, transition metal-based MOFs could provide facile Lewis acid active sites and potential space within the MOF channel for sensing an ion or molecule.⁷ However, to the best of our knowledge, only a few Fe³⁺ probes based on transition MOFs have been developed.⁸

Phenol compounds are important raw materials that are frequently used in synthesizing plastic, adhesives, dyes, medicine, and pesticides, but are also hazardous. Their long-term residual accumulation in the ecosystem has caused serious environmental problems because of their intrinsic high toxicity and non-degradability.⁹ Thus, the detection of phenol compounds through simple and inexpensive methods is an international concern. However, less effort has been focused on the detection of these compounds than has been expended on the detection of other hazardous substances. MOFs, due to their intrinsic traits including their tunable luminescence and exposed active sites, generally sense nitro-aromatic compounds more efficiently than do conventional sensors.¹⁰ The combination of advantages of MOFs enable them to be promising candidates as probes for phenol compounds. Furthermore, such a probe that can operate in an aqueous medium is highly desirable for in-field selective detection of phenol compounds present in soil and ground water. To develop fluorescent MOFs that work in aqueous media for the selective detection of Fe³⁺ and phenol compounds, a water-stable Zn-based MOF, {[Zn(DPTMIA)]·(H₂O)₂(DMF)_{0.5}}_n (HPU-1) with strong luminescence intensity and rectangle-type tubular channels was

College of Chemistry and Chemical Engineering, Henan Polytechnic University, Jiaozuo, 454000, China. E-mail: zhqxu@hpu.edu.cn; wangyuan08@hpu.edu.cn

† Electronic supplementary information (ESI) available: Selected bond lengths and bond angles, additional figures, powder X-ray patterns for the complex. CCDC 1542239 for HPU-1. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7ra08427g



synthesized. **HPU-1** was shown to be able to rapidly detect Fe^{3+} from a mixture of metal ions through an ion exchange mechanism and to exhibit high sensitivity for phenol compounds in aqueous media through an energy transfer mechanism.

Experimental

Materials and physical measurements

All chemicals used in our work were commercially available and used as purchased without further purification. The other reagents and solvents referred to were also commercially available and were employed without further purification. Elemental analyses (C, H, and N) were carried out on a FLASH EA 1112 elemental analyzer. Powder X-ray diffraction (PXRD) experiments were recorded using Cu $K\alpha$ 1 radiation on a PANalytical X'Pert PRO diffractometer. Thermal analyses were performed on a Netzsch STA 449C thermal analyzer from room temperature at a heating rate of $10^\circ\text{C min}^{-1}$ in air. The luminescence spectra for the powdered solid samples were acquired at room temperature on a Hitachi F-4500 fluorescence spectrophotometer. The widths of the excitation and emission slits were each 5 nm. The UV spectra were recorded on a Purkinje General TU-1800 spectrophotometer.

Synthesis of $[\{\text{Zn}(\text{DPTMIA})\} \cdot (\text{H}_2\text{O})_2(\text{DMF})_{0.5}]_n$ (**HPU-1**)

The colorless block crystal of **HPU-1** was easily produced *via* a hydrothermal reaction of $\text{Zn}(\text{NO}_3)_2$ (14.85 mg, 0.03 mmol) and DPTMIA (5-(3,5-di-pyridin-4-yl-[1,2,4]triazol-1-ylmethyl)-isophthalic acid) (12.03 mg, 0.03 mmol) in a solution of H_2O , CH_3CN and DMF (2 + 2 + 1 mL) carried out at 80°C for 72 h. Yield: 47%. Elemental analysis data calcd for $\text{C}_{22.50}\text{H}_{20.50}\text{N}_{5.50}\text{O}_{6.50}\text{Zn}$: C 50.29%, H 3.84%, N 14.33%. Found: C 50.44%, H 3.28%, N 14.66%.

Crystal data collection and refinement

The crystallographic diffraction data for **HPU-1** were obtained on a Siemens Smart CCD single-crystal X-ray diffractometer with a graphite monochromatic MoK α radiation ($\lambda = 0.71073 \text{ \AA}$) at 296 K. The structure was solved by using direct methods with the SHELXS-2014 program of the SHELXTL package and refined on F^2 by using full-matrix least-squares techniques with SHELXL-2014. All empirical absorption corrections were applied using the SADABS program. All non-hydrogen atoms were refined anisotropically, while hydrogen atoms were located and refined geometrically. A SQUEEZE/PLATON technique was applied to remove disordered solvent that could not be satisfactorily modeled. The crystallographic data and structural refinement parameters of the complex are summarized in Table S1,[†] and the selected bond lengths and angles are collated in Table S2.[†]

Results and discussion

Crystal structure of $[\{\text{Zn}(\text{DPTMIA})\} \cdot (\text{H}_2\text{O})_2(\text{DMF})_{0.5}]_n$

The structure of **HPU-1** is shown in Fig. 1 and was determined using single-crystal X-ray diffraction (XRD) studies. The **HPU-1**

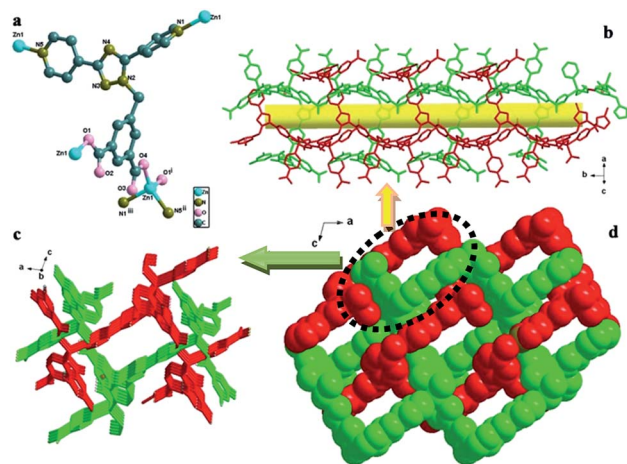


Fig. 1 (a) Linking of the DPTMIA ligand to four Zn^{2+} atoms in the title complex. (b) The rectangular channel. (c) The 2-fold interpenetrating framework. (d) 3D coordination framework with rectangle-type tubular channels.

framework formed from Zn^{2+} ions bridged by DPTMIA ligands (Fig. 1). Each ligand was observed to coordinate with four Zn^{2+} atoms using its two carboxylate groups and two pyridine N atoms. Every Zn^{2+} ion showed the same tetrahedral coordination environment, with the ion being surrounded by two O atoms and two N atoms. As shown in Fig. S1,[†] the extension of the structure into a 3D framework was accomplished by the connections of the ligands and metal ions. The vacant space in **HPU-1** was calculated using Platon to be approximately 31.1%. Other potential voids were found to be filled through mutual interpenetration of another independent equivalent framework in a normal mode, producing a two-fold interpenetrating network (Fig. 1). Further analysis indicated the presence of long rectangular channels with cross-sectional dimensions of $11.4362 \times 8.8427 \text{ \AA}^2$ (Fig. 1).

Sensing of metal ions

As shown in Fig. S2,[†] thermal analyses indicated **HPU-1** to have high thermal stability. The water stability of **HPU-1** was also investigated, as shown in Fig. S3.[†] Powder XRD patterns of the as-synthesized and water-treated samples (**HPU-1** samples were immersed in water for one month) both agreed with the simulated pattern from single-crystal analysis, indicating the water stability of the framework of **HPU-1**, and hence showing its potential to function in an aqueous environment. To investigate the potential ion recognition of **HPU-1**, the samples of **HPU-1** were ground in various 0.01 mol L^{-1} aqueous solutions containing different metal ions (Ag^+ , Al^{3+} , Ba^{2+} , Ca^{2+} , Co^{2+} , Cr^{3+} , Cu^{2+} , Fe^{3+} , Mn^{2+} , Ni^{2+} , Pb^{2+} , Cd^{2+} , K^+ , Li^+ , Mg^{2+} , Na^+ , Eu^{3+} and Tb^{3+}). The suspension-state photoluminescence properties of the various $\text{M}^{n+}@\text{HPU-1}$ samples were determined, and are compared in Fig. 2. The emission intensity of **HPU-1** was nearly unchanged upon being combined with nearly any of the metal ions other than Fe^{3+} . In contrast, Fe^{3+} completely quenched the emission of **HPU-1** (Fig. S4[†]). The ability of **HPU-1** to sensitively



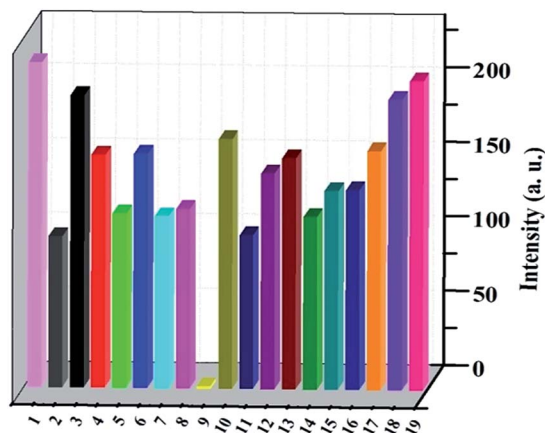


Fig. 2 Fluorescence spectra of HPU-1 (2 mg) in aqueous solutions of various metal ions (3 mL, 10 mM). (1) HPU-1, (2) Ag⁺, (3) Al³⁺, (4) Ba²⁺, (5) Ca²⁺, (6) Co²⁺, (7) Cr³⁺, (8) Cu²⁺, (9) Fe³⁺, (10) Mn²⁺, (11) Ni²⁺, (12) Pb²⁺, (13) Cd²⁺, (14) K⁺, (15) Li⁺, (16) Mg²⁺, (17) Na⁺, (18) Eu³⁺, (19) Tb³⁺.

detect Fe³⁺ in an aqueous solution inspired us to further examine the effects of other metal ions on its Fe³⁺-sensing functions. For this purpose, the same number of equivalents of Fe³⁺ and each of the other metals ions were mixed with HPU-1. Here, in each case, the fluorescence was completely quenched (Fig. 3). This result showed that the interference from conventional metal ions can be neglected, confirming the superior selectivity of HPU-1 for Fe³⁺. Therefore, Fe³⁺ can be considered as a recognizable ion for studying its ability to quench the fluorescence of HPU-1 in suspension. The quenching of HPU-1 luminescence was examined for Fe³⁺ concentrations between 0 and 1000 μM. As shown in Fig. 4, the luminescence intensity of HPU-1 was essentially quenched upon the gradual addition of 100 μL Fe³⁺. The detection limit of HPU-1 for Fe³⁺ was calculated to be 1.09×10^{-3} mol L⁻¹ (Fig. S5†), lower than those of most of the reported

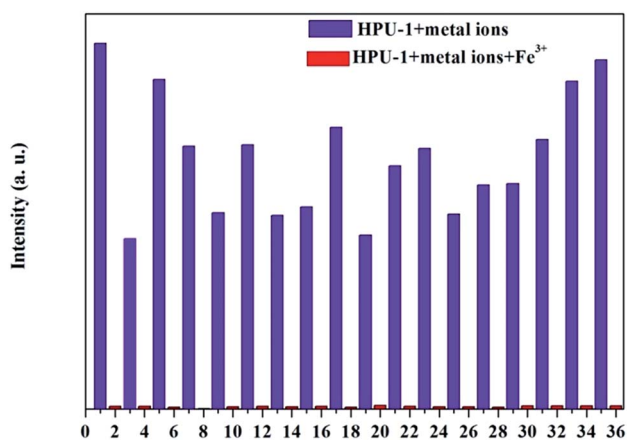


Fig. 3 Fluorescence response of HPU-1 (2 mg) at a wavelength of 430 nm in the absence and presence of various metal ions ((1) HPU-1, (2) Fe³⁺, (3) Ag⁺, (5) Al³⁺, (7) Ba²⁺, (9) Ca²⁺, (11) Co²⁺, (13) Cr³⁺, (15) Cu²⁺, (17) Mn²⁺, (19) Ni²⁺, (21) Pb²⁺, (23) Cd²⁺, (25) K⁺, (27) Li⁺, (29) Mg²⁺, (31) Na⁺, (33) Eu³⁺, (35) Tb³⁺). Purple bars: a free sensor or a sensor treated with the marked ions. Red bars: a sensor treated with the marked metal ion followed by an equivalent of Fe³⁺.

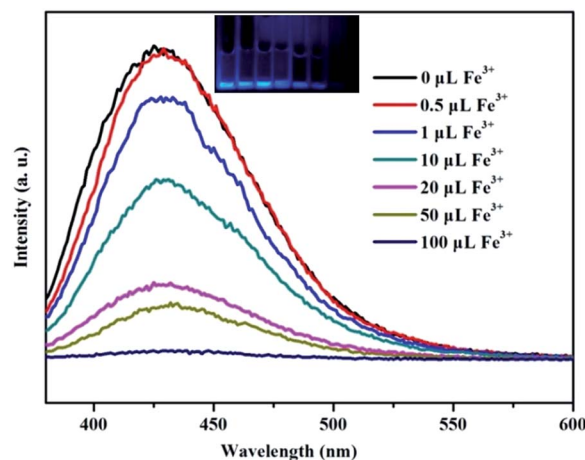


Fig. 4 Fluorescence emission spectra of HPU-1 (2 mg) in aqueous solutions of various concentrations of Fe³⁺.

Fe³⁺-sensing MOFs.¹¹ In addition, the quenching effect can be quantitatively explained by the Stern–Volmer equation,¹² written as

$$\frac{I_0}{I} = 1 + K_{SV}[M]$$

where I_0 and I are the luminescence intensities of the before-sensing and after-sensing products, respectively, $[M]$ is the Fe³⁺ concentration, and K_{SV} is the coefficient of quenching. The K_{SV} value was calculated from the luminescence data to be 1.0×10^4 L mol⁻¹ (Fig. S6†), indicating that Fe³⁺ had particularly strong quenching effect on the luminescence of HPU-1. Compared with the other reported Fe³⁺-sensing MOFs, HPU-1 was found to be a highly selective and sensitive probe for Fe³⁺ in aqueous solution (Table S3†).¹³

The quenching of the luminescence of HPU-1 by Fe³⁺ may have been due to (a) interactions between the sensed metal ions and organic ligands, (b) the collapse of the crystal structure, and/or (c) cation exchange of the central metal ions and the sensed metal ions.¹⁴ Therefore, to elucidate the possible mechanism for such photoluminescence quenching by Fe³⁺, we acquired powder XRD patterns of the samples of HPU-1 immersed in a 0.1 mol L⁻¹ Fe³⁺ solution for 30 min. As shown in Fig. S7,† the PXRD patterns differed little from the original one, illustrating the remarkable quenching effect results from the transformation of the parent HPU-1 framework to Fe-MOF through the cation exchange of Zn²⁺ with Fe³⁺. To further test this hypothesis, ground powder samples of HPU-1 were immersed in a 0.01 mol L⁻¹ Fe³⁺ aqueous solution. After carrying out this immersion for 0.5, 1, 2, 5, 16, and 24 h, the resulting filtrates were introduced for ICP determination. Table S4† shows the concentrations of Zn²⁺ and Fe³⁺ in the filtrate. The concentration of Zn²⁺ in the filtrated solution gradually increased as the level of Fe³⁺ was decreased. In addition, as shown in Fig. S8,† the PXRD patterns of the products of immersion for various amounts of time indicated that the structures gradually changed as the concentration of Fe³⁺ was changed in the products. Moreover, the colors of the products gradually deepened as the immersion time was increased. The

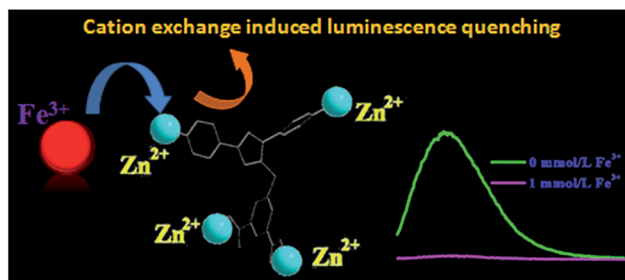


Fig. 5 Schematic representation of the sensing mechanism.

SEM and EDS results showed the presence of Fe^{3+} in the products of 30 minutes of immersion (Fig. S9 and S10[†]). Combining these observations with the above experimental results, we proposed that the fluorescence quenching resulted from the cation exchange mechanism, where Fe^{3+} entered into the **HPU-1** framework and replaced Zn^{2+} , which lowered the energy transfer from the ligands to Zn^{2+} , and thus led to the quenching effect (Fig. 5). Moreover, when Fe^{3+} was added into the aqueous solution of DPTMIA containing Zn^{2+} , the fluorescence intensity was entirely quenched, which also verified that Fe^{3+} can exchange with Zn^{2+} (Fig. S11[†]).

Sensing for hazardous organic analytes

To determine the ability of **HPU-1** to sense toxic organics and explosives, the **HPU-1** powder was immersed in 0.01 mol L^{-1} aqueous solutions of 4-NP, 2,4-DNP, 1,3-DNB, 3-NP, 2,4-DNT, 4-NB, 1,4-DMB, MB, 1,3,5-TMB, and NB for luminescence studies. As shown in Fig. 6 and S12,[†] 4-NP, 2, 4-DNP, and 3-NP showed large quenching effects compared with the other organic analytes, which indicated that **HPU-1** selectively sensed phenolic compounds. In addition, the responses were not disturbed by other organic analytes (Fig. S13 and S14[†]). To examine the sensing sensitivity toward 4-NP or 2,4-DNP, a batch of suspensions of **HPU-1** dispersed in aqueous solution with gradually increasing contents of 4-NP or 2,4-DNP was prepared and their emission spectra were recorded (Fig. S15 and S16[†]). The

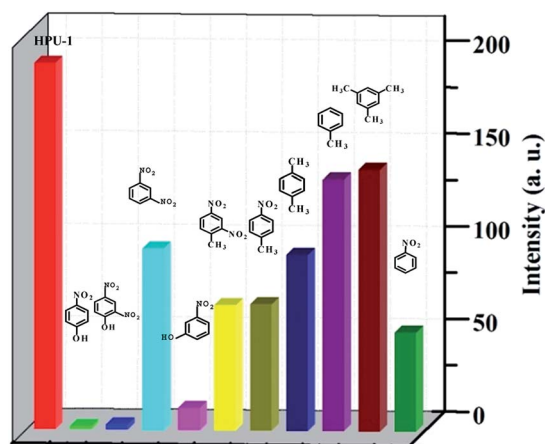


Fig. 6 Fluorescence intensities of **HPU-1** (2 mg) in aqueous solutions of various organic analytes (3 mL, 0.01 mol L^{-1}).

quenching constants calculated from the Stern-Volmer (SV) equation for 4-NP and 2,4-DNP were 1.45×10^5 and 1.074×10^4 L mol^{-1} (Fig. S17 and S18[†]), better than the values in the limited literature. In addition, the detection limits of **HPU-1** for 4-NP and 2,4-DNP were calculated to be 6.31×10^{-4} and 9.86×10^{-4} mol L^{-1} (and about 1.67×10^{-3} mol L^{-1} in the literature) (Fig. S19 and S20[†]).^{15,9b} As mentioned above, **HPU-1** was observed to have large rectangular channels filled with guest molecules. Thus, the sensing mechanism likely involved guest molecules included in the channels exchanging with 4-NP or 2,4-DNP, resulting in the luminescence response. To test this assumption, samples of **HPU-1** were soaked in 1 mg mL^{-1} ethanol solution of 4-NP or 2,4-DNP for 0, 1, 2, 3, 4, 5, 6, 7, 12, 24 and 72 h. Then, UV/vis absorption spectra of the filtrates were acquired and analyzed. Results showed that the absorption peaks of 4-NP or 2,4-DNP were nearly unchanged, demonstrating that 4-NP or 2,4-DNP was not absorbed in the cavity of **HPU-1** (Fig. S21 and S22[†]). Thus, the quenching of the luminescence of Zn^{2+} may have been due to the hydrogen-bonding interactions between NP and the organic ligands resulting in a decrease in the energy transferred from the ligand to Zn^{2+} ions.¹⁶

Finally, the luminescence properties of **HPU-1** in various solvents were investigated for examining the practicality of using **HPU-1** in environmental systems. Well-ground samples of **HPU-1** (2 mg) were dispersed in 2 mL of different solvents, and the emission spectra of **HPU-1** excited at 350 nm revealed strong luminescence (Fig. S23[†]); these results showed these common solvents to have no effect on the luminescence of **HPU-1**. In addition, we also investigated the effect of pH on the fluorescence intensity of **HPU-1**. As shown in Fig. S24,[†] the fluorescence intensity was observed to be nearly the same in relatively acidic and alkaline environments. Therefore, the highly sensitive and selective sensing of Fe^{3+} and phenol compounds in aqueous solutions has provided the foundation for an efficient, straightforward, and real-time detection of trace amounts of this metal ion and hazardous class of compounds in environmental systems.

Conclusions

In conclusion, a water-stable two-fold interpenetrating network (**HPU-1**) with rectangular channels was synthesized and characterized. **HPU-1** was found to be a highly selective and sensitive probe of Fe^{3+} in aqueous solution. Based on our observations and experimental results, we proposed a cation exchange mechanism for the fluorescence quenching. **HPU-1** also showed highly selective and sensitive detection of phenol compounds in aqueous solution. Quenching of the fluorescence was proposed to result from a decrease in the efficiency of the transfer of electrons and energy from the ligands to the metal ions. This work demonstrated the potential application of a fluorescent Zn-MOF as a multi-responsive probe for the detection of Fe^{3+} and hazardous phenol compounds in aqueous media.

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

This work was supported by the National Natural Science Foundation of China (Nos. 21601050), NSFC – Henan region mutual funds (U1604124), the key scientific research project of Henan higher education (16A150010), Science and technology research project of Henan province (152102210314).

References

- (a) J. Wang, M. Jiang, L. Yan, R. Peng, J. Meng, F. Huang, X. X. X. Guo, Y. Li and P. Y. Wu, *Inorg. Chem.*, 2016, **55**, 12660–12668; (b) L. W. He, X. L. Yang, K. X. Xu and W. Y. Lin, *Chem. Commun.*, 2017, **53**, 4080–4083; (c) L. Y. Wang, Y. Tian, L. M. Ding, B. Zhao, X. Y. He, B. Song and S. F. Liu, *RSC Adv.*, 2017, **7**, 16916–16923; (d) S. C. Zhou, M. Zhang, F. Y. Yang, F. Wang and C. Y. Wang, *J. Mater. Chem. C*, 2017, **5**, 2466–2473; (e) B. D. Wang, J. Hai, Z. C. Liu, Q. Wang, Z. Y. Yang and S. H. Sun, *Angew. Chem., Int. Ed.*, 2010, **49**, 4576–4579; (f) Q. H. Tan, Y. Q. Wang, X. Y. Guo, H. T. Liu and Z. L. Liu, *RSC Adv.*, 2016, **6**, 61725–61731.
- (a) D. M. Chen, N. N. Zhang, C. S. Liu and M. Du, *J. Mater. Chem. C*, 2017, **5**, 2311–2317; (b) J. L. Bricks, A. Kovalchuk, C. Trieflinger, M. Nofz, M. Buschel, A. L. Tolmachev, J. Daub and K. Rurack, *J. Am. Chem. Soc.*, 2005, **127**, 13522–13529; (c) A. Barba-Bon, A. M. Costero and S. Gil, *Chem. Commun.*, 2012, **48**, 3000–3002; (d) R. F. Bogale, Y. Z. Chen, J. W. Ye, Y. Y. Yang, A. Rauf, L. Y. Duan, P. Tian and G. L. Ning, *Sens. Actuators, B*, 2017, **245**, 171–178.
- (a) Q. Tang, S. X. Liu, Y. W. Liu, J. Miao, S. J. Li, L. Zhang, Z. Shi and Z. P. Zheng, *Inorg. Chem.*, 2013, **52**, 2799–2801; (b) X. Y. Dong, R. Wang, J. Z. Wang, S. Q. Zang and T. C. W. Mak, *J. Mater. Chem. A*, 2015, **3**, 641–647; (c) Y. T. Liang, G. P. Yang, B. Liu, Y. T. Yan, Z. P. Xi and Y. Y. Wang, *Dalton Trans.*, 2015, **44**, 13325–13330; (d) Y. Shi, S. L. Chou, J. Z. Wang, D. Wexler, H. J. Li, H. K. Liu and Y. P. Wang, *J. Mater. Chem.*, 2012, **22**, 16920–16926; (e) Y. J. Cui, Y. F. Yue, G. D. Qian and B. L. Chen, *Chem. Rev.*, 2012, **112**, 1126–1162.
- (a) J. M. Zhou, W. Shi, H. M. Li, H. Li and P. Cheng, *J. Phys. Chem. C*, 2014, **118**, 416–426; (b) X. Feng, Y. Q. Feng, N. Guo, Y. L. Sun, T. Zhang, L. F. Ma and L. Y. Wang, *Inorg. Chem.*, 2017, **56**, 1713–1721; (c) Y. Q. Xiao, Y. J. Cui, Q. Zheng, S. C. Xiang, G. D. Qian and B. L. Chen, *Chem. Commun.*, 2010, **46**, 5503–5505; (d) Z. M. Hao, X. Z. Song, M. Zhu, X. Meng, S. N. Zhao, S. Q. Su, W. T. Yang and H. J. Zhang, *J. Mater. Chem. A*, 2013, **1**, 11043–11050; (e) S. Mohapatra, B. Rajeswaran, A. Chakraborty, A. Sundaresan and T. K. Maji, *Chem. Mater.*, 2013, **25**, 1673–1679; (f) R. W. Huang, Y. S. Wei, X. Y. Dong, X. H. Wu, C. X. Du, S. Q. Zang and T. C. W. Mak, *Nat. Chem.*, 2017, **9**, 689–697.
- (a) Y. Zhou, H. H. Chen and B. Yan, *J. Mater. Chem. A*, 2014, **2**, 13691–13697; (b) X. Lian and B. Yan, *Inorg. Chem.*, 2016, **55**, 11831–11838; (c) Y. J. Li, Y. L. Wang and Q. Y. Liu, *Inorg. Chem.*, 2017, **56**, 2159–2164; (d) J. N. Hao and B. Yan, *J. Mater. Chem. A*, 2014, **2**, 18018–18025; (e) W. T. Xu, Y. F. Zhou, D. C. Huang, M. Y. Su, K. Wang and M. C. Hong, *Inorg. Chem.*, 2014, **53**, 6497–6499.
- (a) K. C. Wang, X. Tian, Y. H. Jin, J. Sun and Q. H. Zhang, *Cryst. Growth Des.*, 2017, **4**, 1836–1842; (b) Y. Rachuri, B. Parmar, K. K. Bisht and E. Suresh, *Cryst. Growth Des.*, 2017, **17**, 1363–1372; (c) X. D. Zhu, K. Zhang, W. X. Zhou, Q. H. Li, Y. Q. Fu, R. M. Liu and G. X. Qian, *Inorg. Chem. Commun.*, 2016, **75**, 37–40.
- (a) A. K. Chaudhari, S. S. Nagarkar, B. Joarder and S. K. Ghosh, *Cryst. Growth Des.*, 2013, **13**, 3716–3721; (b) L. Wang, Z. Q. Yao, G. J. Ren, S. D. Han, T. L. Hua and X. H. Bu, *Inorg. Chem. Commun.*, 2016, **65**, 9–12.
- (a) L. J. Han, W. Yan, S. G. Chen, Z. Z. Shi and H. G. Zheng, *Inorg. Chem.*, 2017, **56**, 2936–2940; (b) B. Lv, X. F. Wang, H. M. Hu, Y. F. Zhao, M. L. Yang and G. L. Xue, *Inorg. Chim. Acta*, 2016, **453**, 771–778.
- (a) J. L. Yu, X. Y. Wang, Q. Kang, J. H. Li, D. Z. Shen and L. X. Chen, *Environ. Sci.: Nano*, 2017, **4**, 493–502; (b) Q. H. Tan, Y. Q. Wang, X. Y. Guo, H. T. Liu and Z. L. Liu, *RSC Adv.*, 2016, **6**, 61725–61731.
- (a) G. Y. Wang, L. L. Yang, Y. Li, H. Song, W. J. Ruan, Z. Chang and X. H. Bu, *Dalton Trans.*, 2013, **42**, 12865–12868; (b) S. S. Nagarkar, A. V. Desai and S. K. Ghosh, *Chem. Commun.*, 2014, **50**, 8915–8918; (c) W. Xie, S. R. Zhang, D. Y. Du, J. S. Qin, S. J. Bao, J. Li, Z. M. Su, W. W. He, Q. Fu and Y. Q. Lan, *Inorg. Chem.*, 2015, **54**, 3290–3296.
- M. Zheng, H. Q. Tan, Z. G. Xie, L. G. Zhang, X. B. Jing and Z. C. Sun, *Inorg. Chem.*, 2017, **56**, 2936–2940.
- (a) S. S. Nagarkar, A. V. Desai and S. K. Ghosh, *Chem. Commun.*, 2014, **50**, 8915–8918; (b) T. D. Gauthier, E. C. Shane, W. F. Guerin, W. R. Seitz and C. L. Grant, *Environ. Sci. Technol.*, 1986, **20**, 1162–1166; (c) J. Keizer, *J. Am. Chem. Soc.*, 1983, **105**, 1494–1498.
- (a) S. K. Sahoo, D. Sharma, R. K. Bera, G. Crisponi and J. F. Callan, *Chem. Soc. Rev.*, 2012, **41**, 7195–7227; (b) M. Zheng, H. Q. Tan, Z. G. Xie, L. G. Zhang, X. B. Jing and Z. C. Sun, *ACS Appl. Mater. Interfaces*, 2013, **5**, 1078–1108; (c) Y. Zhou, H. H. Chen and B. Yan, *J. Mater. Chem. A*, 2014, **2**, 13691–13697.
- L. H. Cao, F. Shi, W. M. Zhang, S. Q. Zang and T. C. W. Mak, *Chem.–Eur. J.*, 2015, **21**, 15705–15712.
- Y. Q. Wang, Q. H. Tan, H. T. Liu, W. Sun and Z. L. Liu, *RSC Adv.*, 2015, **5**, 86614–86619.
- (a) S. N. Zhao, X. Z. Song, M. Zhu, X. Meng, L. L. Wu, S. Y. Song, C. Wang and H. J. Zhang, *RSC Adv.*, 2015, **5**, 93–98; (b) B. W. Xu, X. F. Wu, H. B. Li, H. Tong and L. X. Wang, *Macromolecules*, 2011, **44**, 5089–5092.

