RSC Advances

PAPER

Cite this: RSC Adv., 2017, 7, 38210

Received 4th June 2017 Accepted 21st July 2017 DOI: 10.1039/c7ra06238a rsc.li/rsc-advances

1. Introduction

In recent years, stimuli-responsive gels, $1-8$ as a type of smart supramolecular materials, have become one of the most significant realms due to their wide application prospects in chemosensors, $9-12$ biomaterials, $13-16$ displays, 17 physical materials, $18-22$ chemical engineering²³⁻²⁶ and other relevant areas.²⁷⁻³² Taking advantage of the dynamic and reversible nature of noncovalent interactions, such as strong van der Waal's forces, hydrogen bonds and π – π stacking,³³ the stimuli-responsive supramolecular gels can sense, process and actuate a response to an external change without assistance.³⁴⁻³⁶ Moreover, metal ionscoordinated supramolecular gels³⁷ have become a focus due to their tunable coordination binding strength and fascinating redox, optical, electronic, as well as magnetic properties of the metal ions.^{38,39} These excellent properties would bring benefit to the applications of smart materials.

An efficient iodide ion chemosensor and a rewritable dual-channel security display material based on an ion responsive supramolecular gel†

Qi Lin[,](http://orcid.org/0000-0001-5722-1347) \mathbf{D}^{*a} Lu Liu,^a Juan Liu,^{*b} Feng Zheng,^a You-Ming Zhang, \mathbf{D}^{a} Hong Yao^a and Ta[i](http://orcid.org/0000-0001-6673-0602)-Bao Wei^{D*a}

By introducing multi-self-assembly driving forces, coordination binding sites and signal groups into the same molecule, a well designed functional gelator G1 was synthesized. The gelator G1 could form a stable Pb^{2+} -coordinated supramolecular metallogel (PbG) accompanied with aggregation-induced fluorescence emission (AIE). PbG shows the reversible selective fluorescent response for I⁻ under a gelgel state. The detection limit of PbG for I^- is 1.0×10^{-7} M. The AIE fluorescence of PbG could be reversibly switched "on–off–on" under gel–gel states via alternatively adding I^- and Pb²⁺ water solution into PbG. Other anions could not induce similar stimuli-response for PbG. Interestingly, when a writing brush dipped in I^- water solution was used to write on the xerogel film of PbG, the film did not show any color changes. However under UV at 365 nm, a clear dark writing image appeared. This dark writing could be erased by brushing Pb²⁺ on the film. More interestingly, when the PbG film containing the invisible I⁻ writing was exposed to iodine vapor, a clear brown writing appeared on the film. However, when this film was placed under the room atmosphere for one minute, the brown writing gradually disappeared. Therefore, the PbG film could act as not only a convenient reversible Γ detection test kit, but also an erasable dual-channel security display material. **PAPER**
 (a) Check for updates

Che this exc. Adv. 2017.7, 38/10

Che this exc. Adv. 2017.7, 38/10

Che this exc. Adv. 2017.7, 38/10

Distributed **by Che this exc. Adv.** 2017.7, 38/10

Distributed **by Channel Securi**

However, although a significant amount of effort has been devoted to the development of metal ions coordinated supramolecular gels,⁴⁰ it is still a big challenge to design and synthesize supramolecular gels that can optically sense a given chemical stimulus with a specific selectivity. In view of this, as a part of our research interest in supramolecular chemistry,⁴¹⁻⁴⁵ we attempted to control the stimuli-response properties of supramolecular gels through the competitive coordination between supramolecular gelators, metal ions and guest compounds.

Herein, we designed and synthesized a gelator G1 based on multi-assembly driving forces,⁴⁶⁻⁴⁹ fluorescent signal groups⁵⁰ and coordination binding sites.^{51,52} The gelator $G1$ could form a stable supramolecular organogel in various solvents at very low critical gelation concentrations (CGCs). After the addition of Pb^{2+} to the G1 ethanol organogel (OG), OG could form a stable Pb^{2+} -coordinated supramolecular metallogel PbG accompanied by the pale blue aggregationinduced fluorescence emission (AIE).⁵³⁻⁵⁶ The AIE of PbG could be reversibly controlled by iodide anions with a specific selectivity in gel–gel states. PbG could act as not only a convenient reversible I^- detection test kit, but also an erasable dual-channel security display material. It is worth mentioning that the security display materials have become of increasing importance.57,58

a Key Laboratory of Eco-Environment-Related Polymer Materials, Ministry of Education of China, Key Laboratory of Polymer Materials of Gansu Province, College of Chemistry and Chemical Engineering, Northwest Normal University, Lanzhou, Gansu, 730070, P. R. China. E-mail: linqi2004@126.com; weitaibao@126.com

^bCollege of Chemical Engineering, Northwest University for Nationalities, Lanzhou, 730000, P. R. China. E-mail: liujuan656@126.com

[†] Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra06238a

2. Experimental

As show in Scheme 1, the compound 3,4,5-tris(hexadecyloxy) benzohydrazide was synthesized according to the literaturereported methods.⁵⁹ G1 was synthesized as follow: p-nitrobenzaldehyde (1 mmol), 3,4,5-tris(hexadecyloxy) benzohydrazide (1 mmol) and acetic acid (0.1 mL, as a catalyst) were added to ethanol (20 mL). Then, the reaction mixture was stirred under refluxed conditions for 24 hours. After the solvent was removed, the precipitated G1 was yielded and recrystallized with CHCl₃–EtOH to get the solid G1. Yield: 60%, m.p.: 106–110 °C, ¹H NMR (400 MHz, CDCl₃, Fig. S1†) δ 9.30 (s, $-NH$, 1H), 8.26 (d, $J = 8.7$ Hz, $-ArH$, $-CH$, 3H), 7.90 (s, $-ArH$, 2H), 7.07 (s, ArH, 2H), 4.01 (d, $J = 3.8$ Hz, $-CH_2$, 6H), 1.95–1.71 (m, $-CH_2$, 6H), 1.36 (m, $-CH_2$, 78H), 0.88 (t, $J = 6.5$ Hz, $-CH_3$, 9H). 13 C NMR (150 MHz, CDCl₃, Fig. S2†) δ 152.76, 123.99, 107.96, 77.19, 76.98, 76.77, 73.59, 73.46, 69.44, 69.16, 60.93, 31.91, 30.30, 29.70, 29.64, 29.62, 29.55, 29.40, 29.38, 29.35, 29.30, 26.06, 22.67, 14.38, 14.08; IR $(KBr, \, \text{cm}^{-1})$ v: 3455 $(-NH),$ 1715 (C=O), 1650 (CH=N). Anal. calcd. for $C_{62}H_{107}N_3O_6$: C 75.03, H 10.40, N 3.98; found: C 75.12, H 10.56, N 3.85. MS: ESI $[M + H]$ ⁺ m/z (Fig. S3†) found: 990.8284, calcd: 990.8233. Paper

Solution in Schwaring Mathematic School (1) and access Article is article in the symbol of the model on the symbol on 2017. The model on 13/10/2024 11:23. The model on 13/10/2024 11:23. The model on 13/10/2024 11:2

3. Results and discussion

First, we carefully investigated the gelation properties of G1. As shown in Table S1,† the gelator G1 could form a stable supramolecular organogel in various solvents at very low critical gelation concentrations. Among these solvents, the gelator G1 showed the lowest CGC (0.40%, wt/v%, 10 mg mL⁻¹ = 1%) and the highest gel–sol transition temperature (78 $^{\circ}$ C) in ethanol. The G1-based supramolecular organogel OG in ethanol is more stable than the gel in other solutions. Therefore, we investigated the influence of metal ions on the G1 organogel in ethanol.

Interestingly, after the addition of Pb^{2+} to the G1 ethanol organogel $[OG]$, OG could form a stable Pb^{2+} -coordinated supramolecular metallogel PbG. As shown in Fig. 1, PbG has no fluorescence emission in hot ethanol solution ($T > T_{gel}$). However, when the temperature of this hot ethanol solution dropped below the T_{gel} of PbG, the emission intensity at 340 nm showed an evident increase and reached a steady state, which indicated that the fluorescence emission of metallogel PbG was aggregation-induced emission (AIE).

The anion response capability of the supramolecular metallogel PbG was primarily investigated by adding various anions in water solutions (AcO⁻, HSO₄⁻, H₂PO₄⁻, F⁻, Cl⁻, Br⁻, I⁻, N₃⁻,

Scheme 1 The synthetic route of organogelator G1.

Fig. 1 Temperature dependent fluorescent spectra of the PbG (1%, in ethanol, G1–Pb²⁺ = 1 : 1) during gelation process (λ_{ex} = 340 nm). The temperature was decreased from 85 \degree C to 25 \degree C and a fluorescence spectrum was recorded every 5 °C.

SCN $^-$, ClO₄ $^-$, CN $^-$, CO₃^{2 $^-$}, S^{2 $^-$} and SO₄^{2 $^-$}, 1 mol L⁻¹) to **PbG**. As shown in Fig. 2, when adding water solutions of various anions to the small amount of metallogel PbG on a spot plate, only I⁻ could quench the fluorescence of PbG, while other anions could not. These results indicated that **PbG** could selectively sense I^- , which was attributed to I^- competitively binding with Pb²⁺.

Moreover, the I^- response properties of PbG were investigated by fluorescence titrations. As shown in Fig. 3, with the addition of I^- into PbG, the spectra showed evident red shifts, which was attributed to the coordination of I^- with Pb²⁺. In **PbG**, Pb^{2+} coordinated with the gelator through the acylhydrazone moiety. When I^- was added into PbG, Pb^{2+} coordinated with I^- and the acylhydrazone moiety was released, which induced the fluorescence spectra of the gel to undergo the red shifts. In addition, the emission intensity at 474 nm decreased with increasing the concentration of $I⁻$. The detection limit of the fluorescence spectra changes, which was calculated on the basis of $3\delta/S, ^{60}$ was 2.037×10^{-6} M (Fig. S4†) for I⁻ anion.

Interestingly, after the addition of Pb^{2+} into the I⁻ containing PbG, the fluorescence of PbG recovered, which was attributed to the Pb^{2+} coordination with G1 again. These properties

Fig. 2 Photograph of PbG (1%, in ethanol, $GI-Pb^{2+} = 1:1$) selectively detects I^- in the presence of various anions (5 equiv.), from left to right and top to bottom are free **PbG**, $PbG + ACO^-$, HSO_4^- , $H_2PO_4^-$, F^- , Cl^- , Br⁻, I⁻, N₃⁻, SCN⁻, ClO₄⁻, CN⁻, CO₃²⁻, S²⁻ and SO₄²⁻, respectively, on a spot plate under UV at 365 nm.

Fig. 3 Fluorescence spectra of PbG (1%, in ethanol, $GI-Pb^{2+} = 1:1$) with increasing concentration of I^- (using 1 mol L^{-1} TBA in water solution as the I⁻ sources), $\lambda_{\rm ex} = 340$ nm.

make PbG act as an I^- and Pb²⁺ controlled "on-off-on" fluorescent switch. By alternate addition of I^- and Pb^{2+} , the switching could be performed reversibly at least for three cycles with a small fluorescent efficiency loss (Fig. 4).

In order to facilitate the use of the metallogel PbG, the $I^$ detection film based on PbG was prepared by pouring the heated ethanol solution of PbG onto a clean glass surface and then drying in air. The **PbG** film was white under natural light and showed a blue fluorescence emission under UV at 365 nm. When a writing brush dipped in I^- water solution was used to write on the film, the film did not show any color changes. However, under UV at 365 nm, a clear dark writing image appeared (Fig. 5). This dark writing image could be erased by brushing Pb^{2+} on the film. More interestingly, when the **PbG** film containing the invisible I^- writing was exposed to iodine vapor, a clear brown writing appeared on the film. However, when the film was put under the room atmosphere for one minute, the brown writing disappeared gradually. Therefore, the PbG film could act as not only a convenient reversible $I^$ detection test kit, but also an erasable dual-channel security display material. **EXCAPIONICES**
 EXCREMENT ACCESS ARTICLE ART

In order to investigate the self-assembly and stimuliresponse mechanism of PbG, a series of experiments was

Fig. 5 Writing, erasing and coloration of a natural light invisible image on a PbG supramolecular gel film (obtained from 1% ethanol metallogel, PbG, $G1-Pb^{2+} = 1 : 1$. Writing: write in I^- water solution; erasing: brush Pb^{2+} water solution; coloration: expose the PbG film into the iodine vapor ca. 5 s). The photographs were taken at room temperature under room light and exposed to a 365 nm UV light.

carried out. First, in the concentration dependent ¹H NMR of G1 (Fig. 6), the –NH– (H_b) and –N=CH– (H_a) resonance signals showed significant downfield shifts as the concentration of G1 rose. These results revealed that in the gelation process, the –NH– (H_b) and –N=CH– (H_a) groups formed hydrogen bonds with the $-C=O$ groups on the adjacent gelators. On the other hand, with a gradual increase in concentration, the ¹H NMR signal of phenyl protons $(H_c, H_d, H_e,$ and H_f) showed an evident upfield shift, indicating that the $\pi-\pi$ stacking interactions between the phenyl groups involved in the gelation process.⁶¹ Therefore, the gelator G1 self-assembled to supramolecular organogel OG by the hydrogen bonds, $\pi-\pi$ stacking as well as the vdW existing in the long alkyl chains.

The formation mechanism of supramolecular metallogel was also investigated by ¹H NMR titrations, as shown in Fig. 7. With the addition of Pb²⁺, the -NH– (H_b) group on the gelator showed significant downfield shifts, which indicated that the gelator coordinated with Pb^{2+} via the acylhydrazone moiety. In addition, in the IR spectra (Fig. S5†) the stretching vibrations of -C=O and -C=N- of G1 showed obviously shifts from 3455 and 1650 cm^{-1} to 3583 and 1588 cm^{-1} , respectively. These phenomena indicated that in **PbG**, Pb^{2+} coordinated with the nitrogen and oxygen atoms on the acylhydrazone group.

This presumed self-assembly and coordination mechanism was also supported by the T_{gel} of OG and PbG. For instance, as shown in Fig. 8, under the same condition, the T_{gel} of OG was significantly higher than that of PbG. The large differences of

Fig. 4 Fluorescent "on–off–on" cycles of PbG, controlled by the alternative addition of I^- and Pb²⁺, $\lambda_{ex} = 340$ nm.

Fig. 6 Partial ${}^{1}H$ NMR spectra of G1 in CDCl₃ with different concentrations.

Fig. 7 Partial $^1\mathsf{H}$ NMR spectra of **G1** (2 mg mL $^{-1}$) and **G1** mixed with 1– 7 equiv. of Pb²⁺ in ethanol- d_6 .

 T_{gel} between OG and PbG were ascribed to the breakage of intermolecular hydrogen bonds between -N=C-H on one gelator and $-C=0$ on the other one in **OG**, which was caused by the coordination of Pb²⁺ with the gelator G1. Moreover, the Pb²⁺ coordination process reduced the distance of $\pi-\pi$ stacking between the phenyls, which enhanced the aggregation induced emission of PbG.

This proposed mechanism was also supported by the XRD patterns (Fig. S6†). The XRD patterns of OG, PbG and the PbG treated with I⁻ showed the peaks at $2\theta = 18.62 - 27.70^{\circ}$, corresponding to the d spacing 3.5 Å, 3.45 Å, and 3.8 Å, respectively. As shown in Scheme 2 and Fig. S6,† in OG, the peaks at 21.28 and the d spacing 3.5 Å were attributed to the π - π stacking, which existed in the phenyl groups of OG. While, in PbG, the d spacing changed to 3.45 Å, which was attributed to the Pb^{2+} coordination process reducing the distance of $\pi-\pi$ stacking between the phenyls. Meanwhile, the d spacing 3.8 Å was attributed to the interlamellar spacing between the supramolecular chains. In addition, after the formation of PbG, the peaks of OG (at $2\theta = 21.28$, 23.22, 23.90, 25.30°) disappeared and these peaks reappeared after **PbG** was treated with $I⁻$. These phenomena confirmed that Pb^{2+} coordinated with OG and induced a change in the XRD pattern of OG, while the addition of I^{$-$} into PbG induced the competitive coordination of I $-$ with

Fig. 8 Plots of T_{gel} against the concentrations of organogel OG and metallogels PbG $(G1-Pb^{2+} = 1: 1)$ in ethanol.

Scheme 2 Chemical structure of the G1 and the presumed self assembly and reversible stimuli-response mechanism.

 Pb^{2+} and led to the recovery of XRD peaks. According to ${}^{1}H$ NMR, IR and XRD, the ion response mechanism of OG and PbG could be presumed as in Scheme 2.

To get further insight into the morphological features of the supramolecular organogel OG, metallogel PbG and PbG treated with I^- , SEM studies were carried out with their xerogels, respectively. As shown in Fig. S7,† the SEM images of OG showed an overlapped rugate layer structure. The metallogel PbG also showed overlapped rugate layer structures and the aggregation structure of layer was compacted. However, after I was added into the PbG xerogel, the micro states experienced obvious changes. There were a large number of micro cavities formed in the xerogel of PbG. These micro cavities provided the PbG xerogel with the properties for the adsorption of iodine vapor. Therefore, the mechanism of iodine vapor-caused color change could be attributed to the iodine vapor adsorption into these micro cavities.

4. Conclusions

In summary, a novel supramolecular gelator G1 has been designed and synthesized. The gelator G1 could form a stable supramolecular metallogel PbG. Through the competitive coordination of Pb^{2+} and I⁻ with the gelator **G1**, the aggregation-induced emission of the supramolecular metallogel PbG was controlled as "on–off–on". More interestingly, after Pb²⁺ competitive coordinated with I^- , there were a large number of micro cavities formed in the PbG xerogel, which enabled the PbG xerogel to absorb the iodine vapor and show a brown color. PbG could act as not only a convenient high selective and sensitive I^- detection test kit, but also an erasable dual-channel secret documentation medium.

Acknowledgements

This study was supported by the National Natural Science Foundation of China (NSFC) (No. 21662031; 21661028; 21574104; 21262032) and the Program for Changjiang Scholars and Innovative Research Team in University of Ministry of Education of China (IRT 15R56).

Notes and references

- 1 M. D. Segarra-Maset, V. J. Nebot, J. F. Miravet and B. Escuder, Chem. Soc. Rev., 2013, 42, 7086.
- 2 H. Tan, X. Ma and M. Fu, Bull. Mater. Sci., 2013, 36, 153.
- 3 L. Predoana, I. Stanciu, M. Anastasescu, J. M. Calderon-Moreno, M. Stoica, S. Preda, M. Gartner and M. Zaharescu, J. Sol-Gel Sci. Technol., 2016, 78, 589.
- 4 Z. X. Liu, Y. Feng, Z. C. Yan, Y. M. He, C. Y. Liu and Q. H. Fan, Chem. Mater., 2012, 24, 3751.
- 5 Q. Xia, Y. Mao, J. Wu, T. Shu and T. Yi, J. Mater. Chem. C, 2014, 2, 1854.
- 6 S. Basak, J. Nanda and A. Banerjee, Chem. Commun., 2014, 50, 2356.
- 7 A. S. Lee, S. S. Choi, K. Y. Baek and S. S. Hwang, Inorg. Chem. Commun., 2016, 73, 7.
- 8 S. S. Babu, V. K. Praveen and A. Ajayaghosh, Chem. Rev., 2014, 114, 1973.
- 9 V. Lozano, R. Hernández, A. Ardá, J. Jiménez-Barbero, C. Mijangos and M. J. Pérez-Pérez, J. Mater. Chem., 2011, 21, 8862.
- 10 Y. He, Z. Guo, P. Jin, C. Jiao, H. Tian and W. Zhu, Ind. Eng. Chem. Res., 2015, 54, 2886.
- 11 C. D. Jones and J. W. Steed, Chem. Soc. Rev., 2016, 45, 6546.
- 12 P. Xue, J. Ding, M. Jin and R. Lu, J. Mater. Chem. C, 2017, 5, 5299.
- 13 Q. Duan, Y. Cao, Y. Li, X. Hu, T. Xiao, C. Lin, Y. Pan and L. Wang, J. Am. Chem. Soc., 2013, 135, 10542.
- 14 G. Visioli, A. Comastri, D. Imperiale, G. Paredi, A. Faccini and N. Marmiroli, Food Anal. Methods, 2016, 9, 469.
- 15 J. Jielile, A. Jialili, G. Sabirhazi, N. Shawutali, D. Redati, J. Chen, J. Bai and K. Aldyarhan, Appl. Biochem. Biotechnol., 2012, 166, 521.
- 16 Y. Cao, Y. Li, X. Y. Hu, X. Zou, S. Xiong, C. Lin and L. Wang, Chem. Mater., 2015, 27, 1110.
- 17 J. Zhou, X. Du, Y. Gao, J. Shi and B. Xu, J. Am. Chem. Soc., 2014, 136, 2970.
- 18 G. Sun, J. Xiao, M. Lu, H. Wang, X. Chen, Y. Yu, Y. Pan and Y. Wang, Appl. Biochem. Biotechnol., 2015, 175, 400.
- 19 T. Kitahara, M. Shirakawa, S. Kawano, U. Beginn, N. Fujita and S. Shinkai, J. Am. Chem. Soc., 2005, 127, 14980.
- 20 X. Y. Hu, T. Xiao, C. Lin, F. Huang and L. Wang, Acc. Chem. Res., 2014, 47, 2041.
- 21 G. Mun, H. Choi, N. Im, J. Ahn, J. Park, H. Seo, Y. Choi, J. H. Lee and J. H. Jung, RSC Adv., 2017, 7, 26827.
- 22 C. Kim, K. Y. Kim, J. H. Lee, J. Ahn, K. Sakurai, S. S. Lee and J. H. Jung, ACS Appl. Mater. Interfaces, 2017, 9, 3799.
- 23 E. P. Simonenko, A. V. Derbenev, N. P. Simonenko, V. G. Sevastyanov and N. T. Kuznetsov, Russ. J. Inorg. Chem., 2015, 60, 1444.
- 24 Z. Cheng, J. Liao, B. He, F. Zhang, F. Zhang, X. Huang and L. Zhou, ACS Sustainable Chem. Eng., 2015, 3, 1677.
- 25 M. Jiang, J. Zhu, C. Chen, Y. Lu, Y. Ge and X. Zhang, ACS Appl. Mater. Interfaces, 2016, 8, 3473.
- 26 Y. Cao, X. Y. Hu, Y. Li, X. Zou, S. Xiong, C. Lin, Y. Z. Shen and L. Wang, J. Am. Chem. Soc., 2014, 136, 10762.
- 27 N. V. V. Satya Bhushan and R. N. Nayak, J. Oral Maxillofac. Surg., 2010, 9, 119.
- 28 J. A. Foster, K. K. Damodaran, A. Maurin, G. M. Day, H. P. G. Thompson, G. J. Cameron, J. C. Bernal and J. W. Steed, Chem. Sci., 2017, 8, 78. **SC Advances**

and Imovative Research Team in University of Minitery of Δt , Y.Com, Y. Weigh, A. Com, Y. W. New Article is licensed on of Climate (1980)

Note S and *Y* Creative Commons Attribution 3.0 Unported Unported
	- 29 H. Wang, X. Ji, Z. Li, C. N. Zhu, X. Yang, T. Li, Z. L. Wu and F. Huang, Materials Chemistry Frontiers, 2017, 1, 167.
	- 30 P. Hallegot, G. Hussler, V. Jeanne-Rose, F. Leroy, F. Pineau and H. Samain, J. Sol-Gel Sci. Technol., 2016, 79, 365.
	- 31 S. Lee and S. T. Nguyen, Macromolecules, 2013, 46, 9169.
	- 32 P. Xue, Q. Xu, P. Gong, C. Qian, A. Ren, Y. Zhang and R. Lu, Chem. Commun., 2013, 49, 5838.
	- 33 T. Ghane, D. Nozaki, A. Dianat, A. Vladyka, R. Gutierrez, J. P. Chinta, S. Yitzchaik, M. Calame and G. Cuniberti, J. Phys. Chem. C, 2015, 119, 6344.
	- 34 N. Song, D. X. Chen, Y. C. Qiu, X. Y. Yang, B. Xu, W. Tian and Y. W. Yang, Chem. Commun., 2014, 50, 8231.
	- 35 X. Yan, D. Xu, X. Chi, J. Chen, S. Dong, X. Ding, Y. Yu and F. Huang, Adv. Mater., 2012, 24, 362.
	- 36 Y. M. Zhang, B. B. Shi, H. Li, W. J. Qu, G. Y. Gao, Q. Lin, H. Yao and T. B. Wei, Polym. Chem., 2014, 5, 4722.
	- 37 Q. Lin, B. Sun, Q. P. Yang, Y. P. Fu, X. Zhu, T. B. Wei and Y. M. Zhang, Chem.–Eur. J., 2014, 20, 11457.
	- 38 K. Lalitha, V. Sridharan, C. U. Maheswari, P. K. Vemula and S. Nagarajan, Chem. Commun., 2017, 53, 1538.
	- 39 Y. Zhang, B. Zhang, Y. Kuang, Y. Gao, J. Shi, X. X. Zhang and B. Xu, J. Am. Chem. Soc., 2013, 135, 5008.
	- 40 S. S. Abolmaali, A. Tamaddon, H. Najafi and R. Dinarvand, J. Inorg. Organomet. Polym., 2014, 4, 977.
	- 41 Q. Lin, T. T. Lu, X. Zhu, B. Sun, Q. P. Yang, T. B. Wei and Y. M. Zhang, Chem. Commun., 2015, 51, 1635.
	- 42 Q. Lin, T. T. Lu, J. C. Lou, G. Y. Wu, T. B. Wei and Y. M. Zhang, Chem. Commun., 2015, 51, 12224.
	- 43 Q. Lin, Q. P. Yang, B. Sun, J. C. Lou, T. B. Wei and Y. M. Zhang, RSC Adv., 2015, 5, 11786.
	- 44 Q. Lin, F. Zheng, L. Liu, P. P. Mao, Y. M. Zhang, H. Yao and T. B. Wei, RSC Adv., 2016, 6, 11928.
	- 45 Q. Lin, P. P. Mao, L. Liu, J. Liu, Y. M. Zhang, H. Yao and T. B. Wei, RSC Adv., 2017, 7, 11206.
	- 46 M. H. Liu and Y. Fang, Chin. Sci. Bull., 2012, 57, 4245.
	- 47 C. Zhao, X. Zhang, K. Li, S. Zhu, Z. Guo, L. Zhang, F. Wang, Q. Fei, S. Luo, P. Shi, H. Tian and W. H. Zhu, J. Am. Chem. Soc., 2015, 137, 8490.
	- 48 Q. Lin, Q. P. Yang, B. Sun, Y. P. Fu, X. Zhu, T. B. Wei and Y. M. Zhang, Soft Matter, 2014, 10, 8427.
	- 49 Y. Gong, T. Jiao, Q. Hu, N. Cheng, W. Xu, Y. Bi and L. Yu, J. Phys. Chem. C, 2015, 119, 16349.
	- 50 X. Ji, Y. Yao, J. Li, X. Yan and F. Huang, J. Am. Chem. Soc., 2013, 135, 74.
	- 51 Q. Lin, B. Sun, Q. P. Yang, Y. P. Fu, X. Zhu, Y. M. Zhang and T. B. Wei, Chem. Commun., 2014, 50, 10669.
	- 52 D. H. Qu, Q. C. Wang, Q. W. Zhang, X. Ma and H. Tian, Chem. Rev., 2015, 115, 7543.
	- 53 Y. Hong, J. W. Y. Lam and B. Z. Tang, Chem. Soc. Rev., 2011, 40, 5361.

- 54 W. Lei, Y. Liu, J. Bai, Y. Dai, Y. Kan, T. Chen and B. Yin, Tetrahedron, 2015, 71, 5465.
- 55 K. Duraimurugan, J. Sivamani, M. Sathiyaraj, V. Thiagarajan and V. Siva, J. Fluoresc., 2016, 26, 1211.
- 56 X. Wu, H. Li, Y. Kan and B. Yin, Dalton Trans., 2013, 42, 16302.
- 57 A. Kishimura, T. Yamashita, K. Yamaguchi and T. Aida, Nat. Mater., 2005, 4, 546.
- 58 N. Tamaoki, A. V. Parfenov, A. Masaki and H. Matsuda, Adv. Mater., 1997, 9, 1102.
- 59 Y. P. Fu, Q. Lin, T. B. Wei, P. Chen, X. Zhu, X. Liu and Y. M. Zhang, Chem. Reagents, 2013, 35, 367.
- 60 Analytical Methods Committee, Analyst, 1987, 112, 199.
- 61 C. Po, Z. Ke, A. Y. Y. Tam, H. F. Chow and V. W. W. Yam, Chem.–Eur. J., 2013, 19, 15735.