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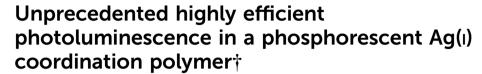


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A luminescent three-dimensional coordination polymer (CP) of [Cd<sup>II</sup>(pmd){Ag<sup>I</sup>(CN)<sub>2</sub>}<sub>2</sub>] (1; pmd = pyrimidine) comprising two different coordination modes of Ag+ ions was synthesised herein. 1 exhibited thermochromic luminescence, accompanied by positive thermal elongation of the Ag. Ag distance. Moreover, 1 showed a bright phosphorescence with the highest photoluminescence quantum yield ( $\Phi_{\rm em}$ ), approximately 60% at room temperature, among previously reported phosphorescent Aq-based CPs or metal-organic frameworks.

Emission-switchable luminescent materials are at the forefront of technological innovations owing to their potential applications in sensitive sensors, image displays, and fluorescent thermometers. 1-5 However, the practical application of emission materials requires the fulfilment of multiple criteria, including structural stability, facile and cost-effective fabrication, and high quantum efficiency. Notably, Ag-based luminescent materials have received considerable attention because of their attractive characteristics: (i) affordability, (ii) low toxicity and biocompatibility, (iii) outstanding luminescence properties, and (iv) sensing abilities. 6-10 For instance, an Ag clusterassembled material (SCAM) of [Ag<sub>12</sub>(StBu)<sub>6</sub>(CF<sub>3</sub>COO)<sub>6</sub>(CPPP)<sub>2</sub>  $(dimethylacetamide)_{12}$ <sub>n</sub> (CPPP = 2,5-bis(4-cyanophenyl)-1,4-b(pyridine-4-yl)-phenyl)-1,4-dihydropyrrolo[3,2-b]pyrrole) exhibited an enhancement of luminescence efficiency compared with that of the CPPP ligand owing to rigidifying methodology through matrix

Luminescent coordination polymers (CPs) and metalorganic frameworks (MOFs), a new class of crystalline porous and functional materials, are being extensively developed in the field of materials science. 13-22 In particular, Ag-CPs/MOFs have great potential as platforms for novel photofunctional materials. For example, a study reported the advantages of MOFs in device applications such as light-emitting diodes, including the facile preparation and highly designability of the frameworks. 23 However, Ag-CPs/MOFs still have an unresolved challenge of low photoluminescence quantum yield ( $\Phi_{\rm em}$ ) at RT, which hinders their wide applications at this stage. Therefore, development of high-performance Ag-based emission materials is required to improve luminescence efficiency.

Herein, we report a novel phosphorescent Ag-CP that exhibits an intense light-blue emission with the highest  $\Phi_{\rm em}$  value of 60% at RT. The luminescence features of the resultant CP were studied by variable-temperature (VT) X-ray diffraction, photoluminescence measurements, and density functional theory (DFT) calculations.

 $[Cd^{II}(pmd)[Ag^{I}(CN)_{2}]_{2}$  (1; pmd = pyrimidine) was synthesised by a slow-diffusion method using an aqueous solution of Cd(NO<sub>3</sub>)<sub>2</sub>·4H<sub>2</sub>O, pmd, and K[Ag(CN)<sub>2</sub>], yielding colourless single crystals (Fig. S1(d)-(f), ESI†). A detailed synthesis of 1 is provided in the ESI.† Single-crystal X-ray diffraction (SCXRD) analysis revealed that compound 1 crystallised in the monoclinic space group C2/c, and the crystal parameters are listed in Table S1 (ESI†). The framework of 1 contains Cd-CN-Ag linkages, which are bridged by bidentate pmd ligands and Ag+ ions, forming a highly dense 3D structure (Fig. 1, Table S1 and

coordination induced emission effects. 11 More recently, the development of highly emissive Ag-thermally activated delayed fluorescence (TADF) complexes was demonstrated, which was attributed to the narrow singlet (S<sub>1</sub>)-triplet (T<sub>1</sub>) energy gaps, hybrid local excitation and charge transfer (CT) characters facilitating inverse intersystem crossing, and molecular rigidity.12 Hence, Ag-based compounds are promising candidates for fabricating sustainable and high-performance luminescent materials.

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<sup>†</sup> Electronic supplementary information (ESI) available: Experimental, crystallographic data, SEM, PXRD patterns, FT-IR spectra, TGA curve, UV-vis reflectance spectra and computational analysis. CCDC 2235241, 2235242 and 2235244-2235247. For ESI and crystallographic data in CIF or other electronic format see DOI: https://doi.org/10.1039/d3cc00179b

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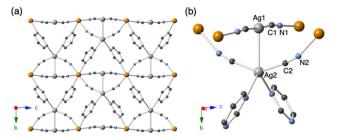


Fig. 1 Crystal structure of 1 at 300 K. (a) The view is along the a-axis. (b) Coordination environment of Ag sites in 1. Atomic code: Cd, orange; Ag, light grey; C, grey; N, blue. H atoms are omitted for clarity.

Fig. S1, ESI†). The Cd<sup>2+</sup> ions in 1 are hexa-coordinated with octahedral geometry, where two pmd ligands coordinated to the axial sites of Cd2+, and the other four were bridged by the cyanides of the [Ag(CN)<sub>2</sub>] species (Fig. S1(a), ESI†). More specifically, half of the Ag<sup>+</sup> ions in 1 adopt a linear geometry (labelled as Ag1 in Fig. 1(b)), whereas the others have tetrahedral structures (labelled as Ag2). Furthermore, the distance between the Ag1···Ag2 atoms  $(d_{\text{Ag1···Ag2}})$  at 300 K is 3.0033(5) Å, which is shorter than twice the van der Waals radius of the Ag+ ion (3.44 Å), indicating the presence of argentophilic interactions in the framework.<sup>24,25</sup> The chemical formula and phase purity of 1 were confirmed by elemental analysis, powder X-ray diffraction (PXRD) patterns (Fig. S2(a), ESI†), and thermogravimetric analysis (TGA) (Fig. S2(c), ESI†). Infrared (IR) spectra showed the two stretching modes of bridging CN<sup>-</sup> ligands at 2126 cm<sup>-1</sup> and 2160 cm<sup>-1</sup> (Fig. S2(b), ESI†), corresponding to the two different evanide coordination modes in 1 (Fig. 1(b)).

The thermal expansion (TE) of 1 was investigated using VT-SCXRD (Fig. 2 and Table S1, ESI†) and VT-synchrotron PXRD measurements (Fig. S3, ESI†) because the luminescent properties of the d<sup>10</sup> and d<sup>8</sup> systems (Au<sup>I</sup>, Ag<sup>I</sup>, Cu<sup>I</sup>, and Pt<sup>II</sup>) tend to be closely related to changes in the metallophilicity of their emission centres with temperature variation. 26-32 The obtained TE

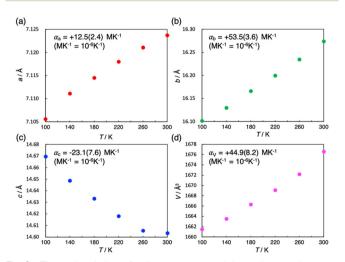


Fig. 2 Thermal variation of cell parameters and thermal expansion constants (MK $^{-1}$  = 10 $^{-6}$  K $^{-1}$ ) with standard deviation (SD): (a) a-, (b) b-, and (c) c-axis and (d) volume calculated using VT-SCXRD results (100-300 K).

coefficients with standard deviation (SD) for the respective axes and volumes are summarised in Fig. 2. 1 displays anisotropic TE behaviour, which comprises positive TE and negative TE with coefficients of  $\alpha_a = +12.5(2.4) \text{ MK}^{-1} \text{ (MK}^{-1} = 10^{-6} \text{ K}^{-1}),$  $\alpha_{\rm b} = +53.5(3.6) \text{ MK}^{-1}, \ \alpha_{\rm c} = -23.1(7.6) \text{ MK}^{-1}, \ \text{and} \ \alpha_{\rm volume} =$ +44.9(8.2) MK<sup>-1</sup>, where the coefficient of thermal expansion  $\alpha = dl/ldT$  (l = lattice parameter). Such an anisotropic TE behaviour is often observed in cyanide-bridged CPs/MOFs due to relatively weak interactions, framework topologies, and various external stimuli.33-40 It is noted that the TE coefficient in the b axis related to  $d_{Ag1\cdots Ag2}$  is the largest of the values in these three axes. Additionally, the  $d_{Ag1\cdots Ag2}$  value systematically decreased from 3.0033(5) Å at 300 K to 2.9877(4) Å at 100 K during cooling (Fig. S4, ESI†).

Photophysical measurements of 1 were conducted to explore the relationship between its structure and luminescence characteristics. Fig. 3(a) shows the VT-emission spectra. At 300 K, 1 exhibited a broad emission with a maximum of  $\lambda_{em}$  = 481 nm upon excitation at  $\lambda_{ex}$  = 330 nm. In addition, the VT-emission spectra gradually red-shifted from 481 to 501 nm, accompanied by the enhancement of the luminescence intensity with cooling. As shown in Fig. 3(b), the VT-excitation spectra measured at  $\lambda_{\rm em}$  = 481–501 nm were similar in shape and rapidly fell around 340 nm, indicating the same luminescence origin in all temperature ranges. The VT-emission decay curves display exponential behaviour with lifetimes in the order of microseconds of 14.1 µs (300 K), 22.1 µs (250 K), 26.5 µs (200 K), 28.7 µs (150 K), 29.6 µs (100 K) and 30.1 µs (77 K) (Fig. 3(c)). These results imply that the resultant emission profiles have a spinforbidden triplet origin, in which the nonradiative decay process was diminished with cooling. 41,42 Moreover, the emission maximum energy of 1 exhibited a good relationship with  $d_{Ag1\cdots Ag2}$ 

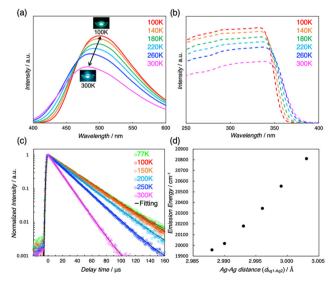


Fig. 3 (a) VT-emission spectra ( $\lambda_{\rm ex}$  = 330 nm). Photographs of **1** at 100 K and 300 K are shown in the inset. (b) VT-excitation spectra ( $\lambda_{em}$  = 481-501 nm) of **1**. (c) VT-emission decay curves ( $\lambda_{ex}$  = 350 nm) of **1**. (d) Plot of emission maximum energy (cm<sup>-1</sup>) vs.  $d_{Aq1\cdots Aq2}$  (Å).

(Fig. 3(d)), indicating the contribution of Ag-Ag sites to the luminescence properties.

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Surprisingly, the emission intensity of 1 was quite strong even at RT, and the photoluminescence quantum yield  $(\Phi_{\rm em})$ was approximately 60%. To the best of our knowledge, the resultant  $\Phi_{\rm em}$  of 1 is the highest value at RT for phosphorescent Ag-based CPs or MOFs. To further investigate the deeper insights into the emission properties of 1, we carried out DFT calculations using the SCXRD data at 300 K as a structural model (details of the DFT calculations are summarized in the ESI†). The optimised infinite structure of 1 at the ground state (S<sub>0</sub>) provided reasonable cell parameters compared to the SCXRD data of 1 at 300 K (Fig. 4(a)); a discrete model structure cut from the optimised infinite form, where the pmd ligands were terminated by hydrogen atoms, was used for the S<sub>0</sub> calculation (Fig. 4(b) and Table S2, ESI†). The calculated molecular orbitals (MOs) related to the electronic transitions suggest that the luminescent feature of 1 is derived from both the Ag...Ag units and pmd ligands, where, for the  $S_0$  state, the highest occupied MO (HOMO) possesses a clear  $\sigma_{\mathrm{Ag-Ag}}^*$  character formed by Ag d orbitals, and the lowest unoccupied MO (LUMO) is the  $\pi^*$  orbital of the pmd ligands (Fig. 4(b)). Moreover, a roughly calculated T1 state also agreed with this orbital configuration along with a contraction of the M···M distance at the excited state (Fig. S5 and Table S2, ESI†), although the incorporation of the effects from surrounding charges and the resulting electric field through the charge embedding method should be required to discuss properly the relaxation of the T<sub>1</sub> states. 43 Thus, the luminescence origin of 1 can be ascribed to metal-metal-to-ligand charge transfer (MMLCT) transitions, which are sometimes observed in d10 or d8 multinuclear luminescence compounds. 44-46 Generally, MMLCT-based emissions are driven by the collaboration of close metallophilic interactions and ligands with appropriate  $\pi^*$  orbitals.<sup>47–50</sup> Hence, the resultant computational analysis indicated that 1 also exhibited a similar trend to the previously reported MMLCT character.

Finally, we studied the effect of MMLCT on the luminescence efficiency of **1**. Compound **1** shows a significantly high  $\Phi_{\rm em}$  value of approximately 60% at RT compared with the previous highest  $\Phi_{\rm em}$  of 22% for the recently reported phosphorescent Ag-MOF of type {[Ag<sub>2</sub>L<sub>2</sub>(CH<sub>3</sub>CN)<sub>2</sub>](BF<sub>4</sub>)<sub>2</sub>}<sub>n</sub> (2; L = diphenyl(2-pyrazyl)phosphine) with the previous highest  $\Phi_{\rm em}$  of 22% at RT.<sup>51</sup> Compound **2** is a 3D framework involving

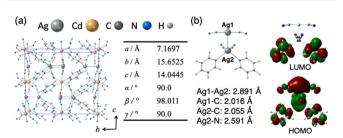


Fig. 4 (a) Infinite optimised structure and cell parameters of  ${\bf 1}$  at the ground  $S_0$  state by DFT calculations. (b) Calculated MOs of the model for  ${\bf 1}$  at the  $S_0$  state.

**Table 1** Comparison of photophysical properties for **1** and  $\{[Ag_2L_2(CH_3CN)_2](BF_4)_2\}_n$  **(2)**<sup>51</sup> in the solid state at 300 K

	1 (300 K)	2 (300 K) <sup>51</sup>
$\lambda_{\rm em}^a/{\rm nm}$	487	545
$\lambda_{\mathrm{em}}^{}a}/\mathrm{nm} \ \Phi_{\mathrm{em}}^{}b}$	0.596	0.22
$\tau_{\rm em}^{c}/\mu s$	14.1	139
$k_{\rm r}^{d}/{\rm s}^{-1}$	$4.23 \times 10^{4}$	$1.58 \times 10^{3}$
$\frac{{{ au_{ m em}}^c}/{\mu  m s}}{{k_{ m r}}^d/{ m s}^{-1}} \ {k_{ m nr}}^e/{ m s}^{-1}$	$2.87 \times 10^{4}$	$5.61 \times 10^3$
$k_{\rm r}/k_{ m nr}$	1.48	0.282

<sup>&</sup>lt;sup>a</sup> Emission maximum. <sup>b</sup> Photoluminescence quantum yields. <sup>c</sup> Emission lifetime. <sup>d</sup> Radiative decay rate constants  $(k_r)$  were estimated using the equation:  $\Phi_{\rm em}/\tau_{\rm em}$ . <sup>e</sup> Nonradiative decay rate constants  $(k_{\rm nr})$  were estimated using the equation:  $k_{\rm r}(1-\Phi_{\rm em})/\Phi_{\rm em}$ .

[Ag<sub>2</sub>L<sub>2</sub>(MeCN)<sub>2</sub>] building units (Fig. S6, ESI†), in which the emission mechanism of 2 was assigned to intraligand phosphorescence (<sup>3</sup>IL). By employing 2, we discuss the effects of different emission mechanisms on the photoluminescence quantum yield. Photophysical parameters such as  $\Phi_{\rm em}$ , emission lifetimes ( $\tau_{\rm em}$ ), radiative decay rate constants ( $k_{\rm r}$ ), and nonradiative decay rate constants ( $k_{\rm nr}$ ) of 1 and 2 at 300 K are summarised in Table 1.

Notably, the  $\tau_{\rm em}$  value of **1** was significantly smaller than that of **2**, indicating that, due to heavy-atom effects, the radiative deactivation of **1**, including intersystem crossing, is more efficient (Table 1). Moreover, the  $k_{\rm r}$  and  $k_{\rm nr}$  values of **1** were approximately 26 and 5 times larger than those of **2**, respectively. The relatively small  $k_{\rm nr}$  of **1** might be attributed to the coordination environment of the Ag ions, as well as the kind of Ag-surrounding ligands and the rigidity of the framework. Importantly, d<sup>10</sup> metal ions with a closed-shell electronic configuration can form a variety of coordination geometries, <sup>10,52</sup> which significantly influences the emission properties as demonstrated in an Au<sup>I</sup>-based complex. <sup>53</sup> Thus, the Ag···Ag-related emission path and structural rigidity of the tetrahedral Ag2 ions in **1** would be key factors for the highest  $\Phi_{\rm em}$  value.

In summary, we have reported a new phosphorescent Ag-CP of type  $[Cd^{II}(pmd)\{Ag^{I}(CN)_{2}\}_{2}]$  (1; pmd = pyrimidine) with an intensive phosphorescence originating from MMLCT transitions. The high luminescence quantum efficiency  $(\Phi_{em} = 59.6\%)$  of Ag-based CPs at RT might be due to the effective luminescent path involving Ag-Ag contacts and the structural rigidity of the framework, including the coordination mode of Ag<sup>+</sup> ions. We highlight that luminescent CPs with the  $^{3}$ MMLCT character can be promising for novel photofunctional materials because of their attractive possibilities such as highly efficient luminescence and favourable emission lifetimes.

H. Y. designed the project and performed all lab experiments and VT-synchrotron PXRD at SAGA-LS. M. S., K. M., and K. O. carried out photoluminescence lifetime measurements. J.P. and Y. H. performed the computational analysis. W. K. assisted in VT-emission and excitation spectroscopy measurements. H. Y. and H. M. edited the manuscript.

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## Conflicts of interest

There are no conflicts to declare.

## References

- 1 M. Irie, T. Fukaminato, T. Sasaki, N. Tamai and T. Kawai, Nature, 2002, 420, 759.
- 2 X. Zhang, S. Rehm, M. M. Safont-Sempere and F. Würthner, Nat. Chem., 2009, 1, 623.
- 3 P. Kumar, S. Singh and B. K. Gupta, Nanoscale, 2016, 8, 14297.
- 4 D. Jariwala, V. K. Sangwan, L. J. Lauhon, T. J. Marks and M. C. Hersam, Chem. Soc. Rev., 2013, 42, 2824.
- 5 C. D. S. Brites, S. Balabhadra and L. D. Carlos, Adv. Opt. Mater., 2019,
- 6 S. Medici, M. Peana, G. Crisponi, V. M. Nurchi, J. I. Lachowicz, M. Remelli and M. A. Zoroddu, Coord. Chem. Rev., 2016, 327, 349.
- 7 R. A. D. Arancon, A. M. Balu, A. A. Romero, M. Ojeda, M. Gomez, J. Blanco, J. L. Domingo and R. Luque, Environ. Res., 2017, 154, 204.
- 8 K. Kennes, C. Martin, W. Baekelant, E. Coutino-Gonzalez, E. Fron, M. B. J. Roeffaers, J. Hofkens and M. Van der Auweraer, ACS Appl. Mater. Interfaces, 2019, 11, 12179.
- 9 S. Horiuchi, S. Moon, A. Ito, J. Tessarolo, E. Sakuda, Y. Arikawa, G. H. Clever and K. Umakoshi, Angew. Chem., Int. Ed., 2021, 60, 10654.
- 10 V. W. Yam and K. K. Lo, Chem. Soc. Rev., 1999, 28, 323.
- 11 Z. Wei, X.-H. Wu, P. Luo, J.-Y. Wang, K. Li and S.-Q. Zang, Chem. -Eur. J., 2019, 25, 2750.
- J.-H. Jia, D. Liang, R. Yu, X.-L. Chen, L. Meng, J.-F. Chang, J.-Z. Liao, M. Yang, X.-N. Li and C.-Z. Lu, Chem. Mater., 2020, 32, 620.
- 13 M. J. Katz, T. Ramnial, H.-Z. Yu and D. B. Leznoff, J. Am. Chem. Soc., 2008, 130, 10662.
- 14 A. Lan, K. Li, H. Wu, D. H. Olson, T. J. Emge, W. Ki, M. Hong and J. Li, Angew. Chem., Int. Ed., 2009, 48, 2334.
- 15 N. Yanai, K. Kitayama, Y. Hijikata, H. Sato, R. Matsuda, Y. Kubota, M. Takata, M. Mizuno, T. Uemura and S. Kitagawa, Nat. Mater., 2011. 10, 787.
- 16 Y. Takashima, V. M. Martínez, S. Furukawa, M. Kondo, S. Shimomura, H. Uehara, M. Nakahama, K. Sugimoto and S. Kitagawa, Nat. Commun., 2011, 2, 168.
- 17 N. B. Shustova, A. F. Cozzolino, S. Reineke, M. Baldo and M. Dincă, J. Am. Chem. Soc., 2013, 135, 13326.
- 18 R.-B. Lin, S.-Y. Liu, J.-W. Ye, X.-Y. Li and J.-P. Zhang, Adv. Sci., 2016, 3. 1500434.
- 19 W. P. Lustig, S. Mukherjee, N. D. Rudd, A. V. Desai, J. Li and S. K. Ghosh, Chem. Soc. Rev., 2017, 46, 3242.

- 20 Y. Zhang, S. Yuan, G. Day, X. Wang, X. Yang and H.-C. Zhou, Coord. Chem. Rev., 2018, 354, 28.
- 21 B. R. Varju, S. A. Wollschlaeger and D. B. Leznoff, Chem. Eur. J., 2019. 25. 9017.
- 22 X.-Y. Liu, W. P. Lustig and J. Li, ACS Energy Lett., 2020, 5, 2671.
- 23 M. Gutiérrez, C. Martín, B. E. Souza, M. V. Auweraer, J. Hofkens and J.-C. Tan, Appl. Mater. Today, 2020, 21, 100817.
- 24 P. Pyykkö, Chem. Rev., 1997, 97, 597.
- 25 H. Schmidbaur and A. Schier, Angew. Chem., Int. Ed., 2015, 54, 746.
- 26 T. H. Kim, Y. W. Shin, J. H. Jung, J. S. Kim and J. Kim, Angew. Chem., Int. Ed., 2008, 47, 685.
- 27 S. Perruchas, X. F. Le Goff, S. Maron, I. Maurin, F. Guillen, A. Garcia, T. Gacoin and J.-P. Boilot, J. Am. Chem. Soc., 2010, 132, 10967.
- 28 I. O. Koshevoy, C.-L. Lin, A. J. Karttunen, M. Haukka, C.-W. Shih, P.-T. Chou, S. P. Tunik and T. A. Pakkanen, Chem. Commun., 2011,
- 29 B. Li, R.-W. Huang, J.-H. Qin, S.-Q. Zang, G.-G. Gao, H.-W. Hou and T. C. W. Mak, Chem. - Eur. J., 2014, 20, 12416.
- 30 M. Jin, T. S. Chung, T. Seki, H. Ito and M. A. Garcia-Garibay, J. Am. Chem. Soc., 2017, 139, 18115.
- 31 M. Jin, S. Yamamoto, T. Seki, H. Ito and M. A. Garcia-Garibay, Angew. Chem., 2019, 131, 18171.
- 32 D. Saito, T. Ogawa, M. Yoshida, J. Takayama, S. Hiura, A. Murayama, A. Kobayashi and M. Kato, Angew. Chem., Int. Ed., 2020, 59, 18723.
- 33 S. Margadonna, K. Prassides and A. N. Fitch, J. Am. Chem. Soc., 2004, 126, 15390.
- 34 K. W. Chapman and P. J. Chupas, J. Am. Chem. Soc., 2007,
- 129, 10090. 35 A. L. Goodwin, M. Calleja, M. J. Conterio, M. T. Dove, J. S. O. Evans,
- D. A. Keen, L. Peters and M. G. Tucker, Science, 2008, 319, 794. 36 J. L. Korčok, M. J. Katz and D. B. Leznoff, J. Am. Chem. Soc., 2009,
- 131, 4866. 37 J. Chen, L. Hu, J. Deng and X. Xing, Chem. Soc. Rev., 2015, 44, 3522.
- 38 J. S. Ovens and D. B. Leznoff, *Inorg. Chem.*, 2017, **56**, 7332–7343.
- 39 B. R. Mullaney, L. Goux-Capes, D. J. Price, G. Chastanet, J.-F. Létard and C. J. Kepert, Nat. Commun., 2017, 8, 1053.
- 40 R. Ohtani, H. Yoshino, J. Yanagisawa, H. Ohtsu, D. Hashizume, Y. Hijikata, J. Pirillo, M. Sadakiyo, K. Kato, Y. Shudo, S. Hayami, B. Le Ouay and M. Ohba, Chem. - Eur. J., 2021, 27, 18135.
- 41 M. D. Allendorf, C. A. Bauer, R. K. Bhakta and R. J. T. Houk, Chem. Soc. Rev., 2009, 38, 1330.
- 42 Y. Cui, Y. Yue, G. Qian and B. Chen, Chem. Rev., 2012, 112, 1126.
- 43 Z. Tu, G. Han, T. Hu, R. Duan and Y. Yi, Chem. Mater., 2019, 31, 6665.
- 44 A. Aliprandi, M. Mauro and L. De Cola, Nat. Chem., 2016, 8, 10.
- 45 S. Carrara, A. Aliprandi, C. F. Hogan and L. De Cola, J. Am. Chem. Soc., 2017, 139, 14605.
- 46 M. Yoshida and M. Kato, Coord. Chem. Rev., 2018, 355, 101.
- 47 J. C. Deaton, A. Chakraborty, R. Czerwieniec, H. Yersin and F. N. Castellano, Phys. Chem. Chem. Phys., 2018, 20, 25096.
- 48 M. Jin, T. Sumitani, H. Sato, T. Seki and H. Ito, J. Am. Chem. Soc., 2018, 140, 2875.
- 49 D. J. Shields, T. Elkoush, E. Miura-Stempel, C. L. Mak, G.-H. Niu, A. D. Gudmundsdottir and M. G. Campbell, Inorg. Chem., 2020, **59**, 18338.
- 50 T. Seki, K. Ida, H. Sato, S. Aono, S. Sakaki and H. Ito, Chem. Eur. J., 2020, 26, 735.
- 51 M. I. Rogovoy, A. S. Berezin, D. G. Samsonenko and A. V. Artem'ev, Inorg. Chem., 2021, 60, 6680.
- 52 C.-M. Che, M.-C. Tse, M. C. W. Chan, K.-K. Cheung, D. L. Phillips and K.-H. Leung, J. Am. Chem. Soc., 2000, 122, 2464.
- 53 K. Igawa, N. Yoshinari, M. Okumura, H. Ohtsu, M. Kawano and T. Konno, Sci. Rep., 2016, 6, 26002.