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Multifunctional Sensing Ability of a new Pt/Zn-Based Luminescent Coordination Polymer

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We synthesized a new Pt/Zn-based coordination polymer, $\{Zn[Pt(CN)_2(5,5'-dcbpy)] \cdot 4H_2O\}_n$, (5,5'-H₂dcbpy = 5,5'-dicarboxy-2,2'-bipyridine), which exhibits reversible colour changes in response to temperature change or exposure to chemical vapours and liquids. Such chromic behaviour shows promise for sensing not only changes in temperature but also for detecting chemical solvents and vapours. The single crystal X-ray structure indicates that one-dimensional coordination polymeric chains formed by an alternating arrangement of $[Zn(H_2O)_3]^{2+}$ and $[Pt(CN)_2(5,5'-dcbpy)]^{2-}$ stacked to produce moderate metallophilic interactions between the Pt(II) ions. Thermogravimetric analysis and water vapour adsorption measurements show that both the crystal water and water coordinated to Zn(II) ions can be removed and re-adsorbed reversibly by heating or under vacuum. Emission spectra at various temperatures and/or in the presence of vapours or liquids reveal that the complex exhibits thermochromic and solvatochromic-like behaviours, with the emission band shifting between 616 and 671 nm. IR spectroscopy and powder X-ray diffraction measurements suggest that this multichromic behaviour is a result of the cooperative phenomena of water adsorption/desorption around the Zn(II) ions and the modification of the metallophilic interaction.

1 Introduction

Luminescent square-planar platinum(II)-diimine complexes have long attracted much attention owing to their interesting optical properties.¹⁻² Some of the complexes exhibit bright phosphorescence from a triplet metal-to-ligand charge transfer (³MLCT) transition state.¹ In the case of one-dimensionally stacked Pt(II) complexes, the electronic interactions between d_{z^2} orbitals of Pt(II) ions give rise to the metal-metal-to-ligand charge transfer (MMLCT) emission state.² The energy of the MMLCT state strongly depends on the distance between adjacent Pt ions.³ Thus, the colour and luminescence of these Pt(II) complexes are controllable by changing the metallophilic interaction between Pt ions. Recently, taking advantage of this characteristic, some groups have reported the ability of Pt(II)-diimine complexes to detect chemical vapours.⁴ This ability, the so-called 'electronic nose', is mainly based on the adsorption of vapour into the crystal lattice that induces significant change in the metallophilic interaction.⁵ As a result, the colour of the complex changes by exposure to chemical vapours; this is called as *vapochromism*. Most Pt(II)-diimine based vapochromic materials are composed of the Pt(II)-polypyridine molecule, which acts as the chromophore, and interactions between these chromophoric molecules are usually based on van der Waals interaction, hydrogen bonding and Coulombic interaction.^{4,5} These interactions give some flexibility to solid-state structures that enable them to recognize chemical vapours and change the structure accompanied by an adsorption of the vapours. However, from the viewpoint of crystal engineering, these interactions make it difficult to control the structure and

vapour selectivity.

Coordination polymers (CPs), on the other hand, have recently attracted increasing attention because of their interesting gas adsorption capacity, controllable frameworks and other properties.⁶ It is well known that the framework of CPs can be controlled by substitution of the metal ion and bridging ligand; that is, the coordination geometry of the metal ion mainly determines the shape, and the bridging ligand contributes to the size of the crystal lattice.⁷ This structural controllability may enable systematic control of both structure and vapour selectivity. Most CPs are insoluble in common solvents such as water, alcohols and hydrocarbons. Considering sensor durability, the insolubility of the sensing material may result in high stability and reversible chromism of these sensing materials by preventing the elution and/or decomposition of the functional molecules. Therefore, with the aim of improving structural control of the vapochromic complexes and maintaining insolubility in solvents, we have designed a new series of coordination polymers (CPs) based on the vapochromic Pt^{II}-diimine complex $[Pt(CN)_2(4,4'-H_2dcbpy)]$ (4,4'-H₂dcbpy = 4,4'-dicarboxy-2,2'-bipyridine) that can act as a complex ligand through two carboxylate groups.⁸ In this paper, we report the crystal structure, adsorption properties and chromic behaviour of a newly synthesized Pt/Zn-based CP, $\{Zn[Pt(CN)_2(5,5'-dcbpy)] \cdot 4H_2O\}_n$ (**1·4H₂O**; 5,5'-H₂dcbpy = 5,5'-dicarboxy-2,2'-bipyridine), and demonstrate that it exhibits reversible colour change in response to temperature change or exposure to chemical vapours and liquids. The chromic behaviour of this CP makes it a promising material for multi-functional environmental sensor applications.

2 Results and Discussion

Crystal structure. The reaction between $[\text{Pt}(\text{CN})_2(5,5'\text{-H}_2\text{dcbpy})]$ and $\text{Zn}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ under basic conditions led to immediate formation of the insoluble coordination polymer $\mathbf{1} \cdot 4\text{H}_2\text{O}$. This very fast reaction made it difficult to form single crystals of $\mathbf{1}$. Although most of the obtained crystals were very thin and twinned or polycrystalline, we successfully determined the crystal structure with enough quality by using a synchrotron light source (see Experimental).[‡] Figure 1 (a) shows the structure of $\mathbf{1} \cdot 4\text{H}_2\text{O}$ at 90 K viewed along the b axis. Complex $\mathbf{1}$ crystallized in the monoclinic $P2_1/m$ space group. The $[\text{Pt}(\text{CN})_2(5,5'\text{-dcbpy})]^{2-}$ complex anion acts as a bridging ligand to produce an infinite 1-D linear polymeric chain along the $a+c$ axis (Figure 1(b)). None of the cyano groups of $[\text{Pt}(\text{CN})_2(5,5'\text{-dcbpy})]^{2-}$ were directly bound to Zn ions. The Zn ion has a trigonal bipyramidal coordination structure with three water molecules in the equatorial plane and two carboxylate oxygens of the dcbpy ligand at the axial positions. The Pt-dcbpy units are uniformly stacked along the b axis to form a metallophilic interaction between adjacent Pt ions with 3.309(1) Å (Figure 1(c)). The distance between adjacent Pt ions observed for $\mathbf{1} \cdot 4\text{H}_2\text{O}$ is longer than that of $[\text{Pt}(\text{CN})_2(4,4'\text{-H}_2\text{dcbpy})] \cdot 4\text{H}_2\text{O}$ (3.2358(1) Å)⁸, suggesting the existence of

a weak metallophilic interaction in $\mathbf{1} \cdot 4\text{H}_2\text{O}$. There are one crystallographically independent, non-coordinated water molecule (O7 atom) that was hydrogen bonded with the coordinated water and the cyano group.

Water adsorption. As described in the Introduction, there are several vapochromic materials based on the Pt(II)-diimine chromophore.^{4,5} In these complexes, the chromic behaviour usually originates from reversible changes in metallophilic interactions induced by vapour adsorption. Our complex $\mathbf{1} \cdot 4\text{H}_2\text{O}$ also shows a moderate metallophilic interaction and includes water molecules in the crystal structure, suggesting a possibility for chromic behaviour. To clarify the reversibility of water vapour adsorption and desorption, thermogravimetric (TG) and differential thermal analyses (DTA) were performed. Figure 2 shows the TG-DTA plot of $\mathbf{1} \cdot 4\text{H}_2\text{O}$. Upon heating, the sample weight gradually decreased, and two steep weight losses were observed at 345 (4.0% loss) and 371 K (9.9% loss). These weight losses were close to the amounts of $2\text{H}_2\text{O}$ (5.8%) and $4\text{H}_2\text{O}$ (11.5%) in $\mathbf{1} \cdot 4\text{H}_2\text{O}$, respectively. The small disagreement between the calculated and observed weight losses may be due to the partial desorption of water at room temperature. At the weight-loss temperatures, endothermic peaks were also observed. These results indicate that the

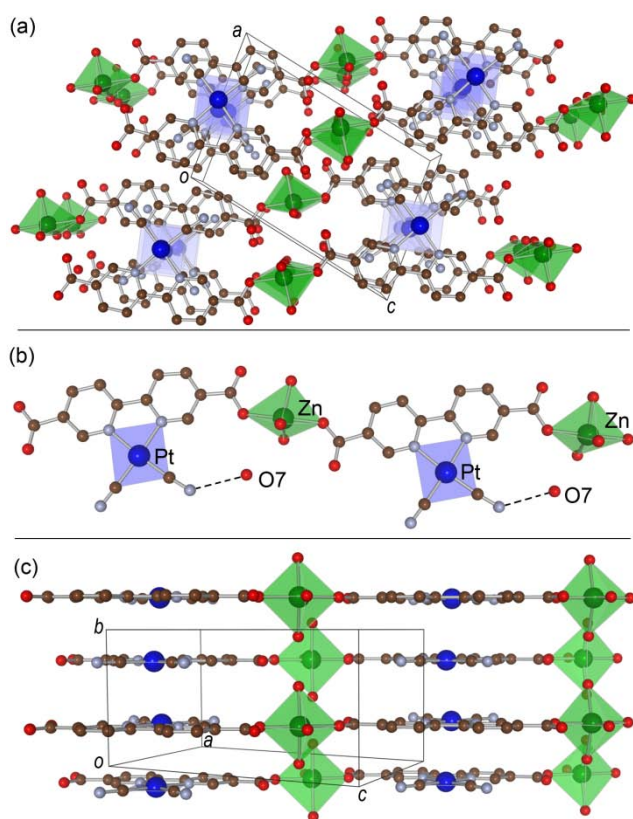


Fig. 1 (a) Perspective view of the unit cell of $\mathbf{1} \cdot 4\text{H}_2\text{O}$ along b axis. (b) One-dimensional chain structure of $\mathbf{1}$. Dotted lines show the hydrogen bonds. (c) Stacked structure with metallophilic interaction of $\mathbf{1}$. Pt, Zn, C, N and O atoms are indicated as blue, green, dark brown, light blue and red filled circles, respectively. Blue plates show the coordination environments of Pt, and green octahedrons show those of the Zn ions. H atoms are omitted for clarity.¹⁶

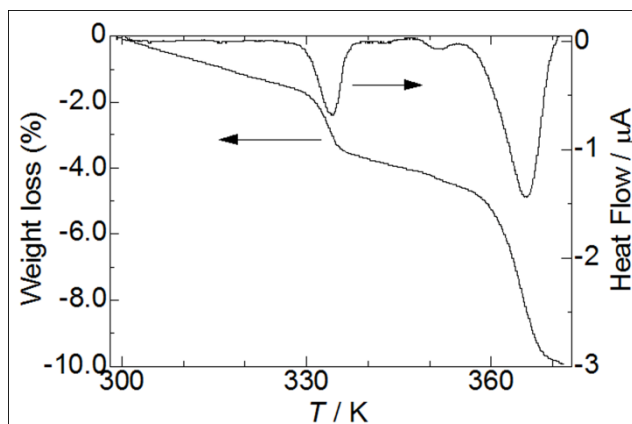


Fig. 2 TG-DTA curves of $\mathbf{1} \cdot 4\text{H}_2\text{O}$. (1 K/min heating, Ar flow rate: 300 ml/min)

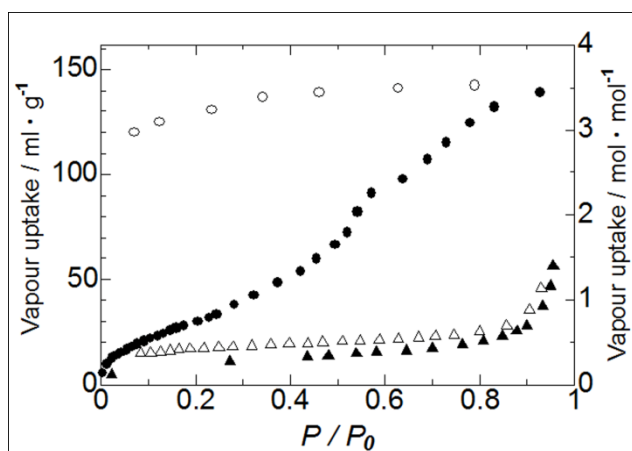


Fig. 3 Adsorption isotherms for water (circle) and methanol (triangle) vapour of $\mathbf{1}$ at 293 K. The open and closed symbols represent the adsorption and desorption process, respectively.

four water molecules included in $1 \cdot 4\text{H}_2\text{O}$ could be removed easily by heating. The chemical compositions at 298, 343 and 373 K, as estimated from the weight loss, represent approximately tetrahydrate $1 \cdot 4\text{H}_2\text{O}$, dihydrate $1 \cdot 2\text{H}_2\text{O}$ and anhydrous **1**, respectively. The adsorption isotherms of water and methanol vapour of **1** were measured (Figure 3). Before each measurement, the sample was heated at 373 K under vacuum to remove all crystal and coordinated water molecules. The dried complex **1** adsorbed water vapour at $3.5 \text{ mol} \cdot \text{mol}^{-1}$ saturated amount, corresponding to the number of coordinated water molecules in $1 \cdot 4\text{H}_2\text{O}$. The adsorption amount at the first step in the lower pressure region up to $P/P_0 = 0.57$ was approximately half the saturated amount (*ca.* $2 \text{ mol} \cdot \text{mol}^{-1}$). In the desorption process, the adsorption amount barely decreased to $P/P_0 = 0.1$, suggesting that the adsorbed water molecules are tightly bound in the pores. This result is consistent with almost all water molecules being bound to Zn(II) ions. On the other hand, only a very small amount of physisorption was observed for methanol vapour. In addition, note that **1** could not adsorb N_2 gas, suggesting the lack of permanent porosity (Figure S1). This water vapour selective adsorption is related to either the window size of the pore or the coordination ability of the adsorbates.

Thermochromic behaviour driven by water vapour. It is well known that some of the Pt(II)-diimine complexes show bright emission from the triplet metal-metal-to-ligand charge transfer ($^3\text{MMLCT}$) state generated by the effective metallophilic interaction between Pt ions.^{3–5} Because the coordination polymer $1 \cdot 4\text{H}_2\text{O}$ shows reversible water adsorption/desorption and has effective metallophilic interaction, we examined the possibility of chromic behaviour.

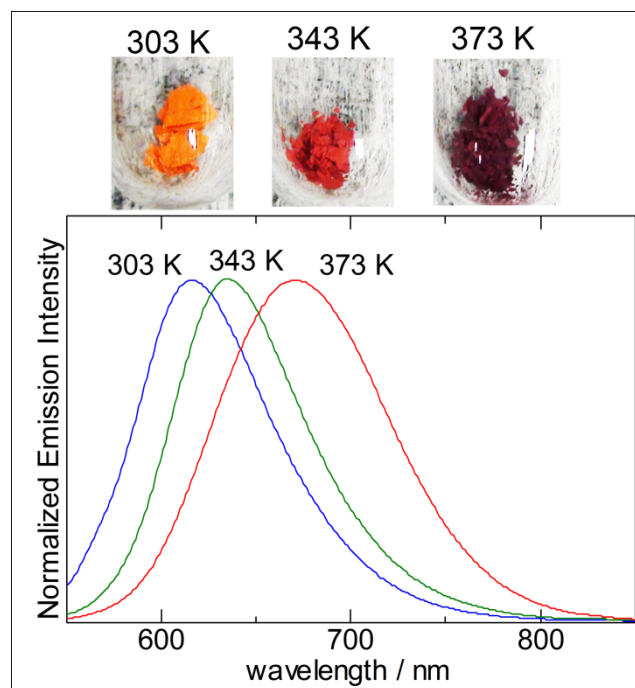


Fig. 4 (Top) Bright field images of **1** at 303, 343 and 373 K. (Bottom) Temperature dependence of the emission spectra of **1**. $\lambda_{\text{exc.}} = 519 \text{ nm}$.

Figure 4 shows the temperature dependence of the emission spectra of **1**. The emission maximum of $1 \cdot 4\text{H}_2\text{O}$ was observed at 614 nm at 303 K, that is, shorter by about 26 nm than that of $[\text{Pt}(\text{CN})_2(4,4'\text{-H}_2\text{dcbpy})] \cdot 4\text{H}_2\text{O}$ (640 nm).⁸ This difference as discussed in the *crystal structure* section, suggesting that the emission of $1 \cdot 4\text{H}_2\text{O}$ is attributable to the $^3\text{MMLCT}$ state. Upon heating, the emission band shifted to a longer wavelength, up to 636 at 343 K and to 671 nm at 373 K. In addition to the red shift of the emission band, the colour of the complex gradually changed from bright orange to red, and finally, to dark purple (see Figure 4). The red shifts observed in emission spectra imply that the shrinkage of the Pt-Pt distance occurs upon heating. In addition, the emission band and colour reverted to the original colours when the temperature was lowered in humid air, indicating the reversibility of the colour change, probably originates from the difference of the Pt-Pt distance.

To investigate the origin of thermochromism, powder X-ray diffraction (PXRD) patterns of $1 \cdot 4\text{H}_2\text{O}$ were measured at various temperatures (Figure 5). The patterns observed at 305 and 343 K were almost identical to the simulation pattern calculated from the structure determined by single crystal X-ray diffraction, showing that the structure of a dihydrate form $1 \cdot 2\text{H}_2\text{O}$ is very similar to that of tetrahydrate $1 \cdot 4\text{H}_2\text{O}$. On the other hand, the pattern at 373 K, where **1** should be anhydrous, was also similar to those at lower temperatures, but it should be noted that the 020 reflection observed at 20.73 deg. shifted about 0.3 deg to a higher angle. This result is evidence of the shrinkage of the crystal lattice along the *b* axis, which corresponds to the direction of metallophilic interaction between Pt ions. The observed shift of the 020 reflection to higher angle corresponds to the shortening of the Pt-Pt distance about 0.05 Å. Upon cooling to 305 K in humid air, the 020 reflection reverted to the original position as well as undergoing a colour change from dark purple to bright orange.

To obtain more detailed structural information about the thermochromism, the temperature dependence of IR spectrum

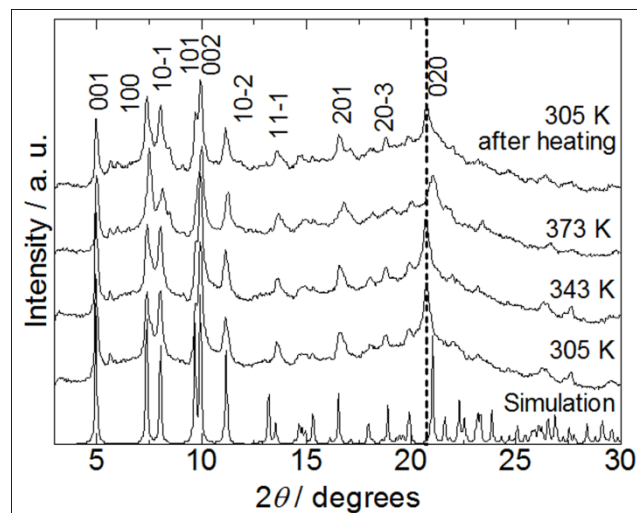


Fig. 5 Temperature dependence of PXRD patterns of **1**. Indexes for main observed reflections are shown at the top. The pattern at the bottom is the simulation calculated from the crystal structure of $1 \cdot 4\text{H}_2\text{O}$ at 90 K. The dotted line is drawn as a guide to the position of 020 reflection at 305 K.

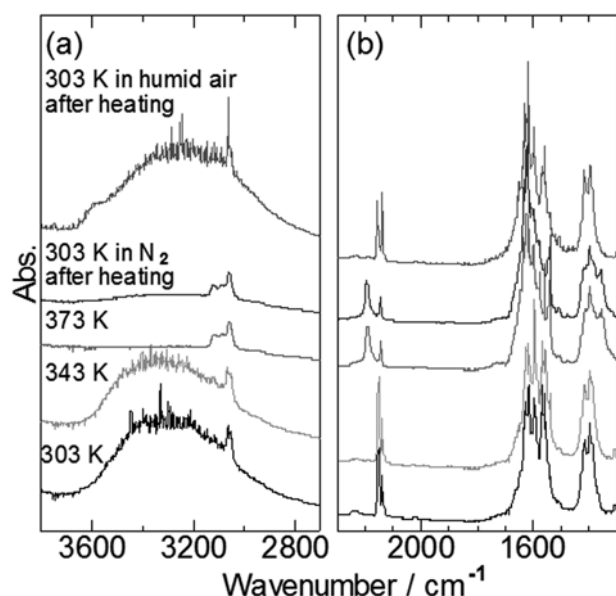


Fig. 6 Temperature dependence of IR spectrum of $1 \cdot 4\text{H}_2\text{O}$. (a) Region of O-H stretching mode in water. (b) Region of CN and CO stretching modes in cyanide ligand and carboxyl group.

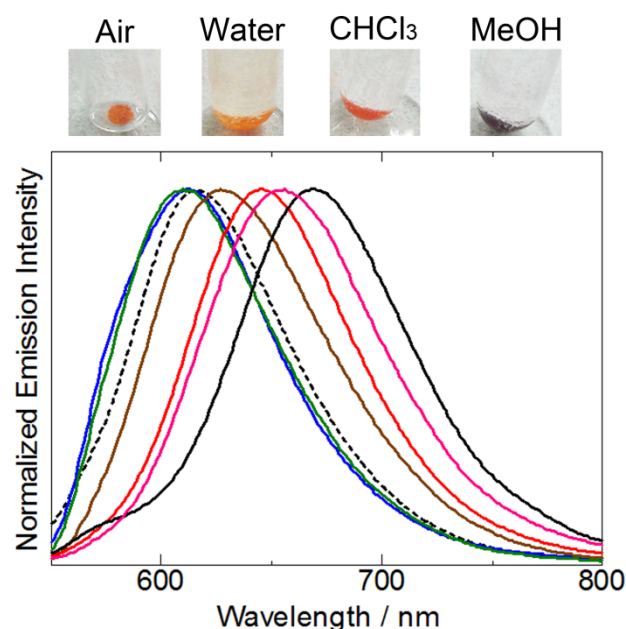


Fig. 7 Bright field images of $1 \cdot 4\text{H}_2\text{O}$ in air, soaked in water, CHCl_3 and MeOH, respectively (top), and emission spectra of $1 \cdot 4\text{H}_2\text{O}$ soaked in various liquid media at room temperature (bottom). $\lambda_{\text{ex}} = 519 \text{ nm}$. Dotted line shows the spectrum of $1 \cdot 4\text{H}_2\text{O}$ in air. Blue, green, brown, red, purple and black solid lines show the spectra of the sample soaked in water, hexane, benzene, ethanol, 1-pentanol, and DMF, respectively.

of $1 \cdot 4\text{H}_2\text{O}$ was measured (Figure 6). At 303 K, the bands assignable to O-H and $\text{C}\equiv\text{N}$ stretching modes were observed at 3300 and 2150 cm^{-1} , respectively. The main asymmetric and symmetric C=O stretching modes were observed around 1596 and 1396 cm^{-1} , respectively. The energy difference between these two modes was about 200 cm^{-1} , suggesting that the bonding mode of the carboxylate group is mainly monodentate.⁹ At 343 K, where the complex changed from tetrahydrate to dihydrate, neither the IR spectrum nor the PXRD pattern changed significantly. On the other hand, above 373 K, several drastic changes were observed. The O-H stretching mode disappeared, and the $\text{C}\equiv\text{N}$ stretching mode was split into two peaks separated by about 50 cm^{-1} , suggesting that all water molecules were removed from the crystal lattice and the cyano groups may coordinate to the Zn ions in high temperature region.⁹ The C=O bond stretching modes also changed with increasing temperature. The two bands of the asymmetric C=O mode merged and shifted to lower energy at 1550 cm^{-1} , and a new band at 1360 cm^{-1} appeared, which is lower than that of the symmetric C=O bond. These changes provide clear evidence of the remarkable changes in the coordination mode of the carboxy group. In other words, the coordination environments of Zn(II) ions dramatically changed at this temperature. From the viewpoint of the coordination environment of Zn(II) ions, the number of coordinated atoms is five at 303 K (as discussed above). However, in anhydrous form **1**, there are no water molecules in the framework, which may limit the number of coordination atoms to no more than six, likely, four. This structural transformation around the Zn(II) ions makes the metallophilic interaction more effective, resulting in the red shift of the emission and the colour of complex **1**. In addition, the spectrum at 373 K was not changed by lowering the temperature to 303 K in a dry nitrogen atmosphere, whereas it

reverted to the spectrum at 303 K in humid air. These results indicate that water adsorption/desorption processes are key factors in the thermochromism of this complex.

Solvatochromic like behaviours in liquids. As described in the Introduction, most vapochromic behaviours of Pt complexes occur based on a molecular assembly involving weak intermolecular interactions, such as hydrogen bonds or van der Waals interactions.⁵ Those weak interactions may enable these materials to undergo the structural change induced by vapour adsorption. Our complex $1 \cdot 4\text{H}_2\text{O}$ is composed of the polymeric chain formed by coordination bonds, which are stronger than hydrogen bonds. To examine whether the complex $1 \cdot 4\text{H}_2\text{O}$ shows vapochromic behaviour, we measured the emission spectra under various conditions. Although the colour of $1 \cdot 4\text{H}_2\text{O}$ changed gradually when exposed to organic vapours, the corresponding changes occurred rapidly when the solid sample was immersed in the liquids. To estimate simply the effects of liquids, emission spectrum of $1 \cdot 4\text{H}_2\text{O}$ was measured for the sample soaked in various liquid media. Figure 7 shows the emission spectra of $1 \cdot 4\text{H}_2\text{O}$ soaked in various liquids; the observed emission maxima of each spectrum are summarized in Table S1. The complex $1 \cdot 4\text{H}_2\text{O}$ is essentially insoluble unless a strong base is added. The emission maximum of $1 \cdot 4\text{H}_2\text{O}$ in air was observed at 614 nm. When $1 \cdot 4\text{H}_2\text{O}$ was soaked in non-polar liquids such as hexane and toluene, the emission band barely changed, or slightly shifted to a longer wavelength (about 15 nm or less). Soaking $1 \cdot 4\text{H}_2\text{O}$ in highly polar liquids shifted the emission band to a significantly longer wavelength, over 30 nm, except in the case of water. Along with the red shift of the emission band,

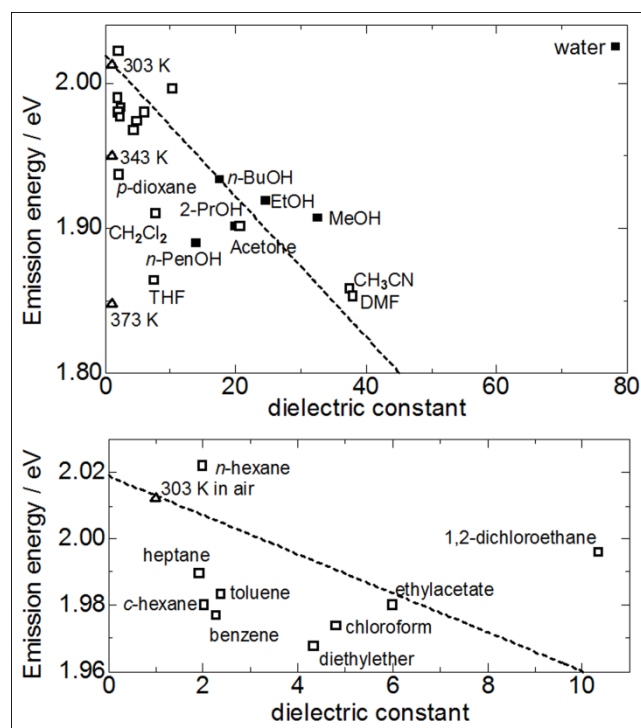
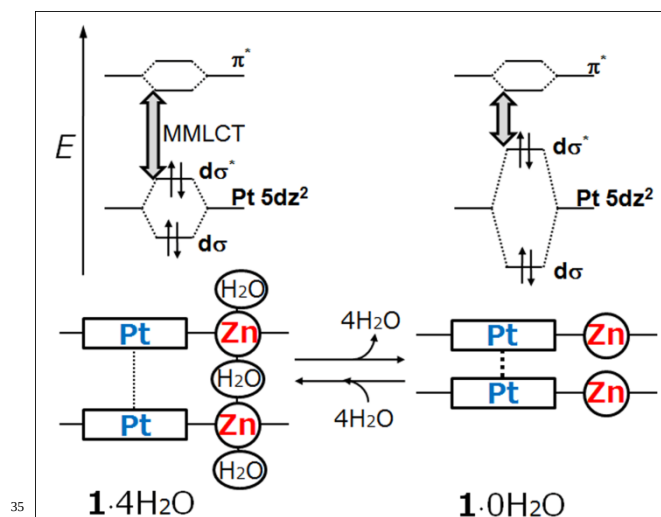


Fig. 8 Plot of the emission energy of **1**·4H₂O against dielectric constant of the soaking liquid in all region (upper) and low dielectric constant region below 10 (lower). Data without liquid are shown as open triangles, and with aprotic and protic liquids are shown as open and closed squares, respectively. The dotted lines are drawn as a guide.

the colour change from orange to dark purple induced by soaking is also very similar to that of the thermochromism (see images in Figure 7). As well as the red shifts observed in emission spectra, the absorption band in diffusion reflectance spectrum of **1** shifted to longer wavelength about 100 nm when the sample was soaked in DMF (see Figure S3 in supporting information). It is noteworthy that the emission band also shifted to a longer wavelength in liquid MeOH, in spite of the lack of adsorbability for MeOH. This result provides clear evidence showing that this *solvatochromic-like* behaviour originates from a structural transformation similar to that occurring in thermochromism. In fact, the emission band maximum of the sample soaked in DMF (669 nm) is almost the same as that observed at 373 K (671 nm). In addition, the emission energies seem to be proportional to the dielectric constant of the liquid as shown in Figure 8. The red shifts observed for THF, dichloromethane, and 1,4-dioxane are relatively larger than that for nonpolar liquids. THF and dichloromethane have large dipole moments, 1.670 and 1.364 debye, respectively, which are close to that of MeOH.¹⁰ 1,4-dioxane does not have a high dipole moment (about 0.002 debye) but has high affinity for water.¹⁰ It should be emphasised that these emission shifts in the soaking solvents were hardly observed in the solvents that included 5% or more water (Figure S2). Considering these results, we believe that this *solvatochromic-like* behaviour originates from the elution of the water molecules involved in **1**·4H₂O to liquid media, resulting in the red shift of emission with colour deepening.



Scheme 1 Possible mechanism of the chromic behaviour of the complex **1**. Upper and lower show schematic energy diagrams and structures of the two forms (**1**·4H₂O and **1**·0H₂O), respectively.

3 Conclusion

We have newly synthesised a Pt/Zn-based coordination polymer $\{Zn[Pt(CN)_2(5,5'\text{-dcbpy})]\cdot 4H_2O\}_n$, and found that this polymer exhibits chromic behaviour under various environmental conditions. The structure of the polymer comprises one-dimensional polymeric chains formed by alternating Zn(II) cations and $[Pt(CN)_2(5,5'\text{-dcbpy})]^{2-}$ complex anions. These chains are stacked to generate moderate metallophilic interaction between the Pt(II) ions. The temperature dependence shown by the powder X-ray diffraction and the IR spectrum reveal that the adsorption/desorption of water molecules coordinated to Zn(II) ion is the main driving force of the chromic behaviour. The possible mechanism of the chromic behaviour of **1** is summarized in scheme 1. The coordination geometry of Zn(II) strongly depends on the number of coordinated water molecules. As a result of changing the number of water molecules coordinated to the Zn(II) ion, the metallophilic interaction between Pt ions changes simultaneously. In other words, the Zn(II) ions act as the water-adsorbing site and the Pt units serve as both the chromophore and emitting site. These chromic behaviours driven by water adsorption/desorption should be applicable to a multifunctional environmental sensor that can sense temperature, pressure, humidity, chemical vapours, and liquids. Our preliminary experiments imply that the Zn(II) ion can be substituted by the other divalent metal ions, such as Ca(II) and Ni(II), and that the adsorption properties of these CPs seem to depend on the kind of the metal. Further work to develop a series of coordination polymers based on $[Pt(CN)_2(5,5'\text{-dcbpy})]^{2-}$ complex anion are now in progress.

4 Experimental

Syntheses. All starting materials, K_2PtCl_4 and 3-methylpyridine were used as received from commercial sources, and the solvents were used without any purification. $Pt(CN)_2$,¹¹ 5,5'-H₂dcbpy¹² and $[Pt(CN)_2(5,5'\text{-H}_2\text{dcbpy})]^{2-}$ complex anion are now in progress.

prepared according to published methods. Elemental analysis was performed at the analysis centre in Hokkaido University.

{Zn[Pt(CN)₂(5,5'-dcbpy)]·4H₂O}_n (1·4H₂O): KCl saturated aqueous solution (5 ml) was added to the solution of [Pt(CN)₂(5,5'-H₂dcbpy)] (40 mg, 82 μmol) in 0.1 M aqueous ammonia (8 ml). A yellow precipitate emerged immediately and was then filtered. The yellow precipitate was dissolved in water (15 ml). To this clear yellow solution, the solution of Zn(NO₃)·6H₂O (24 mg, 84 μmol) in water (5 ml) was added. The orange precipitate immediately formed. After subsequent stirring for 1 h at room temperature, the solids were filtered off from the reaction mixture. The isolated precipitate was washed thoroughly with water and then dried in air for 1 day to afford orange powder 1·4H₂O (42 mg, 67 μmol) in 82% yield. A single crystal suitable for X-ray structural determination of 1·4H₂O was obtained by the diffusion method of water/EtOH with ethyl acetate as an intermediate buffer at 323 K. Elemental analysis calcd. for C₁₄H₆N₄O₄PtZn·4H₂O: C 26.83, H 2.25, N 8.94; found: C 26.59, H 2.31, N 8.70.

Single crystal X-ray structural determination. Single crystal X-ray diffraction measurement was performed using a Rigaku MERCURY CCD diffractometer at NW2A beamline of the Advanced Ring, Photon Factory of KEK, Japan. The wavelength of the synchrotron X-ray was 0.6890(1) Å. A single crystal of 1·4H₂O (crystal dimension: 0.20 × 0.04 × 0.02 mm³) was mounted on a MicroMount with paraffin oil. A nitrogen gas flow temperature controller was used to cool the sample at 90 K. Diffraction data were collected using CrystalClear¹³ and processed by HKL2000.¹⁴ The structure was solved by a direct method using SIR2004 and refined by the full-matrix least-squares using SHELXL97.¹⁵ The non-hydrogen atoms were refined anisotropically. H atoms were refined using the riding model.

Powder X-ray diffraction. Powder X-ray diffraction measurements at various temperatures were performed using a Rigaku SPD diffractometer at beamline BL-8B, Photon Factory, KEK, Japan. The wavelength of the synchrotron X-ray was 1.200(1) Å. The sample was placed in a glass capillary of 0.5 mmφ diameter. The temperature of the sample was controlled using a nitrogen gas flow variable temperature controller.

Luminescence spectroscopy. Emission and excitation spectra were recorded under various conditions on a Jasco FP-6600 spectrofluorometer. The sample temperature was controlled by a JASCO ETC-273 peltier-type temperature controller. About 1 mg of the sample was placed in a glass capillary of 0.5 mmφ in diameter. Slit widths of excitation and emission light are 5 and 6 nm, respectively.

IR spectroscopy. Temperature-dependent IR spectra were recorded on a Nicolet 6700 FT-IR spectrometer using a Nicolet Continuum microscope. The sample temperature was controlled by a Linkam LK-600 hotstage.

Thermogravimetric analysis. Thermogravimetry and differential thermal analysis were performed using a Rigaku ThermoEvo TG8120 analyzer.

Adsorption Isotherms. The adsorption isotherms of water and MeOH vapours at 293 K were performed using an automatic

volumetric adsorption apparatus (BELSORP MAX; BEL Japan, Inc.)

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Notes and references

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- † Electronic Supplementary Information (ESI) available: Adsorption isotherms for N₂ at 77 K, emission spectrum in acetonitrile containing 5% water, diffusion reflectance spectra and the relationship between emission maxima, luminescent quantum efficiencies and dielectric constants of the liquids. See DOI: 10.1039/b000000x/
- ‡ Crystal data of 1·4H₂O: {Zn[Pt(CN)₂(5,5'-dcbpy)]·4H₂O}_n, *T* = 90(1) K, *P*2₁/*m* (#11), *a* = 9.495(17) Å, *b* = 6.577(11) Å, *c* = 14.10(2) Å, β = 101.38(3)°, *V* = 864(3) Å³, *Z* = 2, μ(MoKα) = 7.6 mm⁻¹, *D*_{calc} = 2.410 g·cm⁻³, *R*_{int} = 0.112. No. of collected and unique reflections were 9503 and 3055, respectively. The final *R*_i and w*R*(*F*²) were 0.0780 and 0.2296 (*I* > 2σ), respectively. Full Crystallographic details have been deposited with Cambridge Crystallographic Data Centre as supplementary publication No. CCDC-744577.
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