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# Adaptive coordination-driven supramolecular syntheses toward new polymetallic Cu(I) luminescent assemblies.

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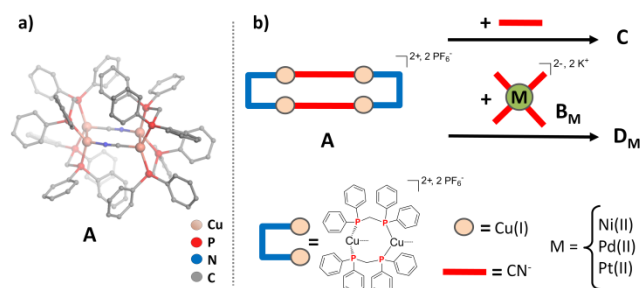
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**ABSTRACT:** A TADF tetrametallic Cu(I) metallacycle **A** behaves as a conformationally adaptive pre-organized precursor to afford, through straightforward and rational coordination-driven supramolecular processes, a variety of room-temperature solid-state luminescent polymetallic assemblies. Reacting various cyano-based building blocks with **A**, a homometallic Cu(I) 1D-helical coordination polymer **C** and  $\text{Cu}_8\text{M}$  discrete circular hetero-bimetallic assemblies  $\text{D}_\text{M}$  ( $\text{M} = \text{Ni}, \text{Pd}, \text{Pt}$ ) are obtained. Their luminescence behaviours are studied, revealing notably the crucial impact of the spin-orbit coupling offered by the central M metal center on the photophysical properties of the hetero-bimetallic  $\text{D}_\text{M}$  derivatives.

## INTRODUCTION

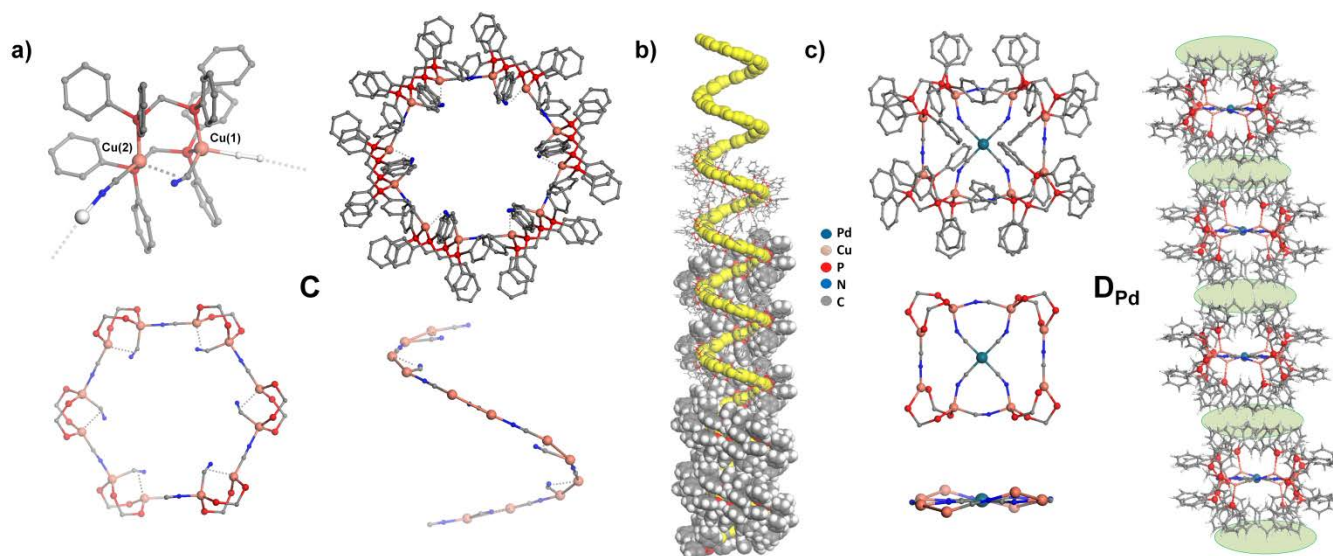
Coordination-driven supramolecular synthesis (CDS) has emerged in the last decade as one of the most powerful modern synthetic strategies for providing an impressive number of increasingly complex supramolecular assemblies.<sup>1</sup> CDS uses the robust, reversible and directional metal-ligand coordinative bond to selectively build high-order functional metallo-supramolecular architectures via the self-assembly of discrete individual components (metal centers and multitopic linkers) having pre-defined geometries and connectivities. Recently, thermally activated delayed fluorescence (TADF) Cu(I) coordination complexes have witnessed impressive developments<sup>2</sup> due to their exalted photophysical performances, that can now compete with those of phosphorescent derivatives based on expensive heavy metal complexes.<sup>3</sup> Nevertheless, no studies have focused up to now on the use of such cheap TADF Cu(I) complexes as pre-organized precursors for the general CDS of emissive coordination-driven supramolecular assemblies. This might be due to the fact that Cu(I) complexes are expected to present labile and non-directional coordination spheres which is *a priori* biased misperception to allow selective CDS toward discrete well-defined metallo-supramolecular architectures. Herein we report for the first time the selective one-step CDS of a series of multimetallic luminescent Cu(I) supramolecular assemblies using the Cu(I) derivative **A**<sup>4</sup> (Figure 1) as pre-organized adaptive supramolecular precursor. We also show that these new derivatives present a variety of emissive behaviours, opening up new perspectives in luminophor design and in the manifold fields of applications that these derivatives can address (lighting and visual displays, telecommunication, sensors etc.).



**Figure 1.** (a) Molecular structure of the dicationic precursor **A**; (b) Syntheses of the derivatives **C** and  $\text{D}_\text{M}$ .

## RESULTS AND DISCUSSION

Due to specific molecular design, some Cu(I) bimetallic precursors bearing short intermetallic distances and conformationally stable coordination directions have been identified as versatile pre-organized precursors.<sup>5</sup> Very specific CDS toward compact supramolecular metallacycles could thus be conducted, including the one-step quantitative preparation<sup>6</sup> of the blue-luminophor consisting of tetrametallic Cu(I) compact metallacycle **A** (Figure 1a).<sup>4</sup> **A** is a result of the selective self-association of two  $[\text{Cu}_2(\mu_2\text{-dppm})_2]$  fragments with two cyano ditopic linkers and undergoes very efficient TADF processes in the solid state.<sup>4</sup> In **A**, the four Cu(I) metal centers possess trigonal planar coordination spheres that could allow additional coordination of donor ligands to gain access to new supramolecular assemblies. In order to probe this assumption, **A** was reacted (Figure 1b) with KCN and the tetracyanometalated  $\text{K}_2[\text{M}(\text{CN})_4]$  metallo-ligands  $\text{B}_\text{M}$  ( $\text{B}_\text{Ni}$ ,  $\text{M} = \text{Ni}$ ;  $\text{B}_\text{Pd}$ ,  $\text{M} = \text{Pd}$ ;  $\text{B}_\text{Pt}$ ,  $\text{M} = \text{Pt}$ ), playing the role of, respectively, potential linear and square planar polytopic



**Figure 2.** a) Molecular X-ray structure of the bimetallic repetition unit of **C** (left top), view of the 1D-helicoidal coordination polymer **C** along the *c* axis showing six repetition units' pitch (right top), simplified 'top' and side views of the showing six repetition units' pitch of **C** (bottom); b) view perpendicular to the *c* axis of the 1D helicoidal polymer **C** with a gradual CPK style to stick and ball-style to a removal of the dppm ligands, the  $(\text{CuCN})_n$  helicoidal chain located in the wall of the 1D-channels formed being shown in CPK style in yellow; c) Molecular X-ray structure, simplified 'top' and side views of the derivative  $\mathbf{D}_{\text{Pd}}$  and columnar solid state stacking of the derivative  $\mathbf{D}_{\text{Pd}}$ , the green ellipses represent the intermolecular contact areas involving  $\pi$ -CH interactions (Counteranions, H atoms and solvent molecules have been omitted for clarity; location of the C and N atoms of the cyano ligands has been chosen arbitrarily in the X-ray structure views except for the  $\text{M}(\text{CN})_4$  fragments).

inorganic linkers. As the colorless  $\text{CH}_2\text{Cl}_2$  solutions of the metallacycle **A** were mixed with methanol solutions of KCN and of precursors  $\mathbf{B}_M$ , neither color changes nor precipitates were observed. Yet, upon UV light excitation at 365 nm, an instantaneous change of the color of the light emitted by the solution occurred. For the reactions with KCN,  $\mathbf{B}_{\text{Ni}}$  and  $\mathbf{B}_{\text{Pd}}$ , the eye-perceived emitted color changed from blue (typical of  $\text{CH}_2\text{Cl}_2$  solutions of the metallacycle **A**) to orange, and to yellow for the reaction conducted with  $\mathbf{B}_{\text{Pt}}$ . Accordingly, the emission spectra recorded for these crude solutions (Figure S19) revealed a red-shift of the large emission band observed for **A**, suggesting that the reaction occurs immediately between **A** and KCN or  $\mathbf{B}_M$  to afford new derivatives whose electronic structures are altered compared to that of **A**. After crystallization, these crude solutions afforded respectively the new derivatives **C** and  $\mathbf{D}_M$  ( $\mathbf{D}_{\text{Ni}}$ , *M* = Ni;  $\mathbf{D}_{\text{Pd}}$ , *M* = Pd;  $\mathbf{D}_{\text{Pt}}$ , *M* = Pt) as air-stable colorless polycrystalline derivatives in good overall yields (Figure 1b).<sup>6</sup> These derivatives are moderately soluble in chlorinated solvents. Their  $^{31}\text{P}\{^1\text{H}\}$  NMR spectra<sup>6</sup> recorded in  $\text{CD}_2\text{Cl}_2$  reveal one broad singlet at -14.8 ppm for **C** and two broad singlets for  $\mathbf{D}_M$  complexes ( $\mathbf{D}_{\text{Ni}}$ ,  $\delta = -17.0$  and -19.1 ppm;  $\mathbf{D}_{\text{Pd}}$ ,  $\delta = -16.3$  and -18.7 ppm;  $\mathbf{D}_{\text{Pt}}$ ,  $\delta = -16.3$  and -18.6 ppm) that are shifted to low frequencies compared to the two broad singlets observed for **A** ( $\delta = -8.5$  ppm and -10.5 ppm).<sup>4</sup> Their  $^1\text{H}$  NMR spectra presented similar patterns in which the  $^1\text{H}$  NMR resonances of the dppm methylene protons are split into two groups of broad signals indicating their non-equivalence. These data suggest that the  $1\kappa\text{P}:2\kappa\text{P}$ -coordination of the dppm ligands observed in the precursor **A** is retained in these new derivatives, in which the gross molecular structure of the  $\text{Cu}_2(\mu_2\text{-dppm})_2$  fragment serves as the building unit. In addition, solid-state infrared spectra show two different  $\nu(\text{C}\equiv\text{N})$  stretches (2071 and 2120  $\text{cm}^{-1}$ ) for **C** (cf.

$\nu(\text{C}\equiv\text{N})$ : 2117  $\text{cm}^{-1}$  for **A**),<sup>4</sup> suggesting two different coordination modes for the cyano ligands, while for  $\mathbf{D}_M$  complexes it reveals  $\nu(\text{C}\equiv\text{N})$  modes corresponding to Cu-CN-Cu ( $\mathbf{D}_{\text{Ni}}$ , 2124  $\text{cm}^{-1}$ ;  $\mathbf{D}_{\text{Pd}}$ , 2121  $\text{cm}^{-1}$ ;  $\mathbf{D}_{\text{Pt}}$ , 2121  $\text{cm}^{-1}$ ) and Cu-CN-*M* ( $\mathbf{D}_{\text{Ni}}$ , 2138  $\text{cm}^{-1}$ ;  $\mathbf{D}_{\text{Pd}}$ , 2141  $\text{cm}^{-1}$ ;  $\mathbf{D}_{\text{Pt}}$ , 2151  $\text{cm}^{-1}$ ) fragments.

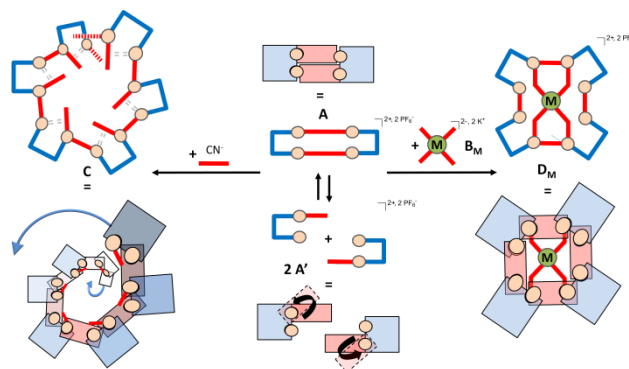
Finally, the molecular structures of derivatives **C** and  $\mathbf{D}_M$  complexes were established by X-ray diffraction studies<sup>6</sup> performed at low temperature on single crystals obtained at room temperature (RT) from pentane diffusion into  $\text{CH}_2\text{Cl}_2$  solutions of the respective complexes. The derivative **C** crystallizes as a neutral 1D-coordination polymer (which is likely to dissociate in solution). The repeating unit is based on a  $\text{Cu}_2(\mu_2\text{-dppm})_2$  fragment, on which two different cyano ligands are coordinated to the metal centers (Figure 2a, left top).<sup>7</sup> One of these cyano ligands acts as  $\mu_2$ -ditopic linker between two Cu(I) metal centers of two neighboring repeating units while the second cyano ligand bridges the two metal centers within each  $\text{Cu}_2(\mu_2\text{-dppm})_2$  fragment in an unusual 'end-on/side-on' coordination mode. As a result, the Cu(1) metal center bears a distorted tetrahedral coordination sphere while the Cu(2) atom presents a distorted trigonal planar coordination sphere (the 'side-on' interaction is not taken into account in this description). In these dimers, the intermetallic distance ( $d(\text{Cu}-\text{Cu}) = 3.48$  Å) is larger than in **A** ( $d(\text{Cu}-\text{Cu}) = 2.87$  Å) and precludes cuprophilic interactions in **C**. The  $\mu_2$ -cyano ligands present almost linear coordination angle on the metal centers ( $\text{Cu}-\text{C}-\text{N} = 177.8(6)^\circ$  and  $177.9(5)^\circ$ ) while the bridging cyano ligands show a significantly more bent coordination to the Cu(I) ions ( $\text{Cu}-\text{C}-\text{N} = 64(1)^\circ$  and  $157(1)^\circ$ ).<sup>7</sup> Five highly disordered  $\text{CH}_2\text{Cl}_2$  molecules complete the asymmetric unit. The solid-state infinite structure of **C** is formed by the connection of these bimetallic repeating units to afford a remarkable neutral 1D-helicoidal coordination polymer whose pitch is based on

the connection of six repeating units (Figure 2a). Finally, in the extended solid state, helices are oriented parallel to the *c* axis affording a molecular solid in which infinite channels (with cross-sectional diameter: *ca.* 10 Å, filled by the CH<sub>2</sub>Cl<sub>2</sub> molecules) having inner walls bearing (CuCN)<sub>n</sub> helical wire are observed (Figure 2b). It is worth to note that since **C** crystallizes in a non-centrosymmetric space group, only one kind of helical structure (P or M) is found in a single crystal. Nevertheless, the crystallization process being not enantioselective, single crystals bearing both right- and left-handed helices are collected.

Isostructural derivatives **D<sub>M</sub>** are discrete polymetallic assemblies. The asymmetric units contain one dicationic Cu<sub>8</sub>M<sub>1</sub>(CN)<sub>8</sub>dppm<sub>8</sub> discrete unit, two hexafluorophosphate counter-anions and ten disordered CH<sub>2</sub>Cl<sub>2</sub> solvent molecules. These Cu<sub>8</sub>M<sub>1</sub>(CN)<sub>8</sub>dppm<sub>8</sub> complexes (Figure 2b) bear a central square-planar 'M(CN)<sub>4</sub>' unit surrounded by a 'Cu<sub>8</sub>(CN)<sub>4</sub>' metallacycle which can be described as the connection of four Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub> fragments by four peripheral μ<sub>2</sub>-CN ligands.<sup>7</sup> In each Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub> fragment, one of the metal centers has a distorted trigonal planar coordination sphere resulting from the coordination of two 1κP:2κP dppm ligands and the peripheral cyano ligands. The second Cu(I) metal center presents a distorted tetrahedral coordination sphere due to an additional link with a central μ<sub>2</sub>-CN ligand, which connects the 'Cu<sub>8</sub>(CN)<sub>4</sub>' metallacycle to the central 'M(CN)<sub>4</sub>' fragment. Within the 'Cu<sub>8</sub>(CN)<sub>4</sub>' metallacycle, the distorted trigonal planar Cu(I) ions as well as the distorted tetrahedral Cu(I) ions are connected together two by two by a peripheral μ<sub>2</sub>-CN ligand (Figure 2c). The resulting 'Cu<sub>8</sub>M(CN)<sub>8</sub>' inorganic cores show almost planar gross rectangular shapes (Figure 2c; maximum deviation from the mean plane defined by the 'Cu<sub>8</sub>M(CN)<sub>8</sub>' fragment = 0.531 Å (**D<sub>Ni</sub>**); 0.665 Å (**D<sub>Pd</sub>**); 0.640 Å (**D<sub>Pt</sub>**)) of similar size despite the different atomic radii of the central metal ions (10.7 \* 9.6 Å<sup>2</sup> (**D<sub>Ni</sub>**); 10.6 \* 9.7 Å<sup>2</sup> (**D<sub>Pd</sub>**); 10.6 \* 9.6 Å<sup>2</sup> (**D<sub>Pt</sub>**), Figure S18). Intermetallic Cu(I) distances in the Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub> fragments are too large for cuprophilic interactions (d(Cu-Cu) = 3.35 - 3.48 Å (**D<sub>Ni</sub>**); 3.36 - 3.46 Å (**D<sub>Pd</sub>**); 3.36 - 3.44 Å (**D<sub>Pt</sub>**)). The peripheral cyano ligands present slightly tilted coordination angle with the metal centers (Cu-C-N = 158.3°-175.9°). The central cyano ligands are almost linear at the M metal centers (M-C-N = 175.2°-177.8°) but present much more bent coordination angles on the Cu(I) ions (Cu-C-N = 145.4°-153.3°). Finally, **D<sub>M</sub>** derivatives stack on the top of each other in the crystalline solid state along infinite columns (Figure 2c), presumably allowing the maximization of cumulative stabilizing CH-π interactions between most of the phenyl rings of the dppm ligands of neighboring assemblies.

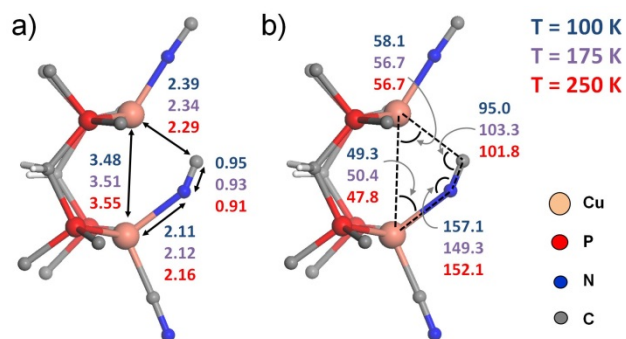
At first glance, the discrepancy observed between the supramolecular structures characterized (1D-helicoidal coordination polymer **C** versus discrete polymetallic assemblies **D<sub>M</sub>**) might seem inappropriate to consider **A** as a directional pre-organized precursor in CDS. Nevertheless, it can be rationalized by considering that the derivative **A** is the result of the dimerization of reactive and adaptable "Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub>CN" fragments **A'** bearing coordinatively highly unsaturated Cu(I) metal centers (Scheme 1). Indeed, in such an hypothesis, a reversible equilibrium is likely to occur in solution between **A** and **A'** allowing reaction of the latter with additional free cyano ligands or M(CN)<sub>4</sub><sup>2-</sup> fragments (Scheme 1). In the first case, the cyano ligand can be considered as formally inserted within the metal centers of **A'** affording "Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub>CN"

**Scheme 1. Suggested mechanism of self-assembling for the syntheses of **C** and **D<sub>M</sub>** from the precursor **A**.**



fragments that self-assemble in the 1D-helicoidal structure of **C**. In the second case, the square planar M(CN)<sub>4</sub><sup>2-</sup> building block acts as a template to connect four **A'** units in the discrete assemblies **D<sub>M</sub>** (Scheme 1). Importantly, such self-assembly processes do not strictly follow the basic guiding rules of the CDS chemistry, since the pre-organized bimetallic building blocks do not present rigid geometric constraints that will impose the geometry of the supramolecular assemblies obtained. However, the lability of the cyano ligands and the conformational flexibility of the "Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub>CN" building block have allowed the selective and rational design of new supramolecular assemblies **C** and **D<sub>M</sub>** (Scheme 1) based on an adaptable pre-organized stable molecular clip.

Temperature-dependent X-ray crystal structure analyses<sup>6</sup> were performed on single crystals of **C**, **D<sub>Pd</sub>** and **D<sub>Pt</sub>** in order to get insights about the structural rigidity of these derivatives upon cooling from 250 K to 100 K. Besides a moderate and progressive shrinkage of the unit cell volume upon cooling (*ca.* 3.0 %, 3.8 % and 4.1 % volume contraction for **C**, **D<sub>Pd</sub>** and **D<sub>Pt</sub>** respectively), these measurements reveal a significant temperature-dependent molecular arrangement in **C** (Figure 3 and S13, Table S6) while the molecular geometry in **D<sub>Pd</sub>** and **D<sub>Pt</sub>** are mostly unchanged.<sup>8</sup> Indeed, in **C**, the intermetallic distances within the Cu<sub>2</sub>(μ<sub>2</sub>-dppm)<sub>2</sub> fragment are decreased upon cooling (d(Cu-Cu), 3.55 Å (250 K); 3.51 Å (175 K); 3.48 Å (100 K)). Concomitantly, within this fragment, the 'end-on/side-on' coordinated cyano ligand is found to be

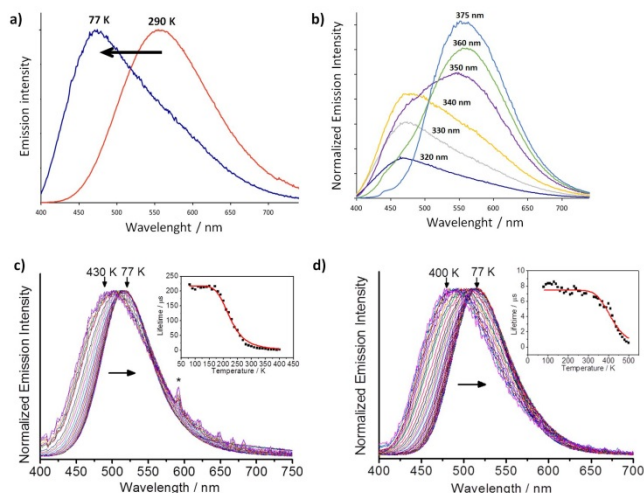


**Figure 3.** Temperature variation of a) selected lengths (Å) and b) angles (°) in the repetition unit of the X-ray structure of **C** (phenyl rings have been omitted for clarity).

significantly labile presenting shorter Cu-terminal N/C atom distances at low temperature and a tendency to partially de-coordinate and switch from the ‘end-on/side-on’ coordination mode to a lateral  $\mu_2$ -coordination mode at high temperature.

Derivatives **C** and **D<sub>M</sub>** display solid-state luminescence properties that are markedly different from those of the precursor **A**.<sup>6</sup> Crystalline samples of the neutral helix **C** display at 298 K an eye-perceived yellow luminescence under excitation at 320 nm characterized by a broad and featureless band in the emission spectrum with a maximum at 538 nm (Figure 4a) and an emission quantum yield (EQY) of 20 %. The emission band at 538 nm is typical of metal-perturbed dpmm ligand-centered phosphorescence.<sup>9</sup> Upon cooling from 290 K to 90 K under excitation at 320 nm,<sup>6</sup> a gradual drop of the emission band at 538 nm, together with a growth of a higher-energy band to the blue, is observed (Figure 4a) affording at 77 K an emission maximum of 468 nm ( $\Delta\lambda_{em} = 0.344$  eV ( $2780$  cm<sup>-1</sup>)) associated with an intense solid-state visually-perceived blue luminescence. Such a shift of emission maxima to the blue upon decreasing temperature is suggestive of the lack of a TADF mechanism in **C**. The higher-energy emission band is likely a result of an emission origin of metal-to-ligand charge transfer (MLCT) [ $d\sigma^*(Cu) \rightarrow \pi^*(CN)/\pi^*(dpmm)$ ] state with mixing of a copper-centered d-s/d-p excited state modified by the very weak Cu...Cu interaction. Emission bands of similar energy are also found in other related systems.<sup>10</sup> Interestingly, a change is also observed upon variation of the excitation wavelength at 77 K (Figure 4b). The shift in emission maxima and the change in the relative ratio of the high-energy and low-energy bands with excitation wavelength are typical of multiple emission arising from multiple species and site heterogeneity, typically found in polymeric species as is the case of the helicoidal polymeric nature of **C**.

Considering complex **D<sub>Ni</sub>**, only very weak emission is observed in the solid state, with a gradual red shift upon cooling from 400 K to 77 K (Figure S25). Its very weak luminescence and uncertainty in lifetime measurements have precluded its emission origin assignment. Conversely, in the solid state, the derivative **D<sub>Pd</sub>** displays at 298 K a moderate turquoise luminescence upon excitation at 350 nm with emission maximum  $\lambda_{em}$  at 500 nm (Figure 4c). This emission band is broad and featureless with an emissive lifetime  $\tau$  of 12  $\mu$ s at 298 K and an EQY of 2 %. In addition, at 298 K, **D<sub>Pt</sub>** exhibits intense turquoise luminescence upon excitation at 350 nm characterized also by a broad and featureless emission band ( $\lambda_{em} = 510$  nm,  $\tau = ca.$  8  $\mu$ s, EQY = 18 %). Upon cooling in the solid state with excitation at 350 nm, both emission spectra of derivatives **D<sub>Pd</sub>** and **D<sub>Pt</sub>** present gradual red-shifts associated with intense solid state green luminescence at low temperature (Figure 4c,d: for **D<sub>Pd</sub>** from  $\lambda_{em} = 493$  nm at 430 K to  $\lambda_{em} = 515$  nm at 77 K ( $\Delta\lambda_{em} = 0.108$  eV /  $870$  cm<sup>-1</sup>) and for **D<sub>Pt</sub>** from  $\lambda_{em} = 486$  nm at 400 K to  $\lambda_{em} = 510$  nm at 77 K nm ( $\Delta\lambda_{em} = 0.120$  eV /  $968$  cm<sup>-1</sup>)). It is worth to note that the energy shift in the **D<sub>Pd</sub>** derivative is relatively small compared to the one observed for its **D<sub>Pt</sub>** analog while the energy domain in which their thermochromic luminescence is observed is mostly the same (together with the thermochromic shift observed for the very weak emission spectrum of **D<sub>Ni</sub>**, Figure S25). However, there are much significant changes in their respective emission decay lifetime  $\tau$  upon cooling. Indeed, it varies from *ca.* 2.7  $\mu$ s at 400 K to *ca.* 222  $\mu$ s at 80 K for derivative **D<sub>Pd</sub>** while it varies

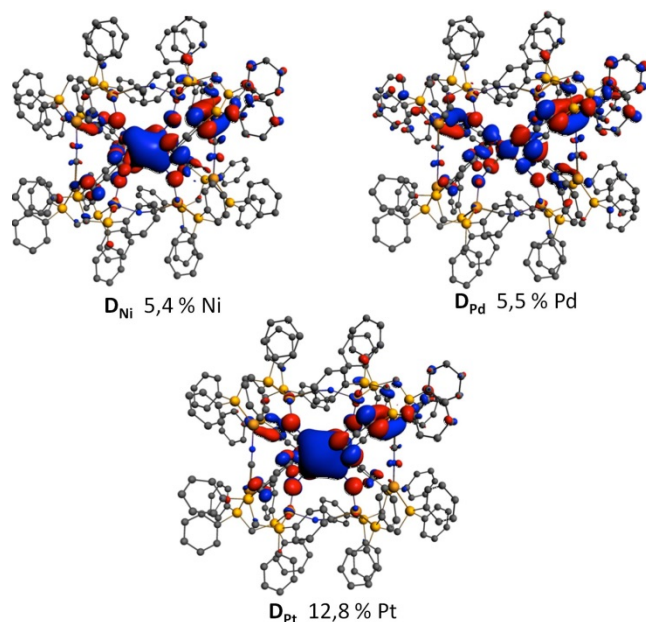


**Figure 4.** a) Normalized solid-state emission spectra of **C** at 77K and 290K ( $\lambda_{ex} = 320$  nm); b) Non-normalized solid-state emission spectra of **C** at 77K at different excitation wavelengths; c) Normalized solid-state emission spectra of **D<sub>Pd</sub>** at different temperatures between 430 K and 77 K upon excitation at 350 nm (full width at half maximum:  $3040$  cm<sup>-1</sup> (80 K);  $5580$  cm<sup>-1</sup> (430 K)); the asterisk represents an instrumental artifact; Insert: plot of emission decay lifetime against temperature (80 K to 400 K); red line represents the fit according to the TADF equation; d) Normalized solid-state emission spectra of **D<sub>Pt</sub>** at different temperatures between 400 K and 77 K upon excitation at 350 nm (full width at half maximum:  $2720$  cm<sup>-1</sup> (80 K);  $4140$  cm<sup>-1</sup> (400 K)); Insert: plot of emission decay lifetime against temperature (80 K to 500 K); red line represents the fit according to the TADF equation.

from *ca.* 0.5  $\mu$ s at 500 K to *ca.* 7.9  $\mu$ s at 80 K for **D<sub>Pt</sub>** (Figure 4c,d, inset).

The emission behavior of **D<sub>Pd</sub>** resembles those typically described for Cu(I) TADF derivatives.<sup>2</sup> Fits of the decay lifetime measured for **D<sub>Pd</sub>** were thus performed accordingly with Equation (1) derived from the TADF model.<sup>6</sup> Values of  $\Delta E(S_1-T_1) = 1290$  cm<sup>-1</sup>,  $\tau(S_1) = 18.6$  ns and  $\tau(T_1) = 218$   $\mu$ s were extracted. They are comparable to those obtained for the precursor **A** ( $\Delta E(S_1-T_1) = 1560$  cm<sup>-1</sup>,  $\tau(S_1) = 9.9$  ns and  $\tau(T_1) = 185$   $\mu$ s)<sup>4</sup> and support *a priori* a TADF mechanism. On the other hand, attempts to fit the decay lifetime for **D<sub>Pt</sub>** to Equation 1 are less than satisfactory given the lack of data points at the high temperature regime. Nevertheless, a rough estimate of a  $\Delta E(S_1-T_1)$  of  $2980$  cm<sup>-1</sup> and a  $\tau(T_1)$  of 7.4  $\mu$ s is obtained by fitting the data to the TADF equation.<sup>11</sup> The relatively large  $\Delta E(S_1-T_1)$  estimated for **D<sub>Pt</sub>** is still within the thermal energy barrier (0.4 eV or  $3230$  cm<sup>-1</sup>) possible for reverse intersystem crossing (RISC).<sup>12</sup> The shorter  $\tau(T_1)$  than those observed in the Cu(I) and **D<sub>Pd</sub>** TADF systems is not unreasonable given the larger spin-orbit coupling constant for Pt.<sup>2c</sup> Nevertheless, it is difficult to confirm the TADF nature for **D<sub>Pt</sub>** with certainty as one cannot exclude the possibility of the competition between ISC to give phosphorescence and RISC to give TADF in **D<sub>Pt</sub>**, i.e. platinum can provide significant spin-orbit coupling, resulting in efficient phosphorescence path in addition to the TADF path.<sup>13</sup> Therefore, considering that **D<sub>M</sub>** derivatives are isostructural, the differences observed along the series clearly suggest a strong impact of the nature of the central M atom in

their photophysical properties. Taking into account the large size of the molecules (417 atoms, 4417 basis functions) and



**Figure 5.** Isocontour plots ( $\pm 0.02$  (e.bohr $^{-1}$ ) $^{1/2}$ ) of the lowest unoccupied molecular orbitals (LUMO) of  $\mathbf{D}_M$  with the associated Mulliken atomic character for M.

the influence of the solid state packing on the geometry, it was not possible to perform full DFT geometry optimization of the excited states to get deep insights about the electronic causes of these different photophysical properties. Nevertheless, the study of the molecular orbitals of the ground state and the nature of the first vertical singlet and triplet excited states of  $\mathbf{D}_M$  are significantly instructive.<sup>6</sup> It reveals that in all cases the vertical excitation to  $S_1$  is described by a combination of electronic transitions from the highest occupied orbitals to the lower-lying unoccupied orbitals. They include the LUMO at different scale depending on the central metal atom (7.5 % of the transitions for  $\mathbf{D}_{Ni}$ ; 43.7 % for  $\mathbf{D}_{Pd}$ ; 59.9 % for  $\mathbf{D}_{Pt}$ ).<sup>14</sup> Moreover, these LUMOs are partly localized on the central metal atom at different extent regarding the nature of M: less than 6% of M atomic character for  $\mathbf{D}_{Ni}$  and  $\mathbf{D}_{Pd}$ , while it reaches up to 12.8 % for  $\mathbf{D}_{Pt}$  (Figure 5). Consequently, the implication of the electronic density of the central metal atom in the overall electronic transition upon the excitation process is less than 1% for  $\mathbf{D}_{Ni}$ , is 2 % for  $\mathbf{D}_{Pd}$  but reaches up to 8 % for  $\mathbf{D}_{Pt}$ . This is nevertheless sufficient to imply enhancement of the spin-orbit coupling components both in the non-radiative and radiative processes involving  $S_1$ , with a magnitude order of Ni < Pd << Pt. This may lead to the stronger competition between efficient ISC and efficient RISC that give rise to a competition between spontaneous phosphorescence and TADF along this series in agreement with the measured photophysical properties. Therefore, these observations demonstrate the crucial and somehow unanticipated impact of the nature of the central metal atom (and more specifically of its spin-orbit coupling value) on the luminescence properties of the entire  $\mathbf{D}_M$  assemblies.

## CONCLUSIONS

The Cu(I) derivative **A** has been revealed to be a versatile pre-organized precursor that provides a straightforward and rational access to a variety of luminescent polymetallic assemblies thanks to its remarkable conformational flexibility. The origin and the diversity of both the supramolecular architectures obtained as well as the photophysical behaviour observed (with luminescence spectra spread across a large part of the visible spectra) highlight the unexplored potential of the use of such adaptive TADF Cu(I) molecular precursors in the fields of CDS toward new luminescent materials. Our current investigations intensively extend this synthetic route to other systems together with an in-depth understanding of their photophysical properties.

## ASSOCIATED CONTENT

### Supporting Information.

The Supporting Information is available free of charge on the ACS Publications website at DOI : xxx/xxx

Synthesis and characterization, X-ray crystallography, photophysical and computational studies (PDF)

Temperature dependent X-ray crystallography data for **C** (CCDC 184431 (100K), 184432 (175K) and 184434 (250K); CIF),  $\mathbf{D}_{Ni}$  (CCDC 184436 (150K); CIF),  $\mathbf{D}_{Pd}$  (CCDC 184442 (100K), 184433 (150K), 184443 (200K) and 184441 (250K); CIF) and  $\mathbf{D}_{Pt}$  (CCDC 184437 (100K), 184339 (150K), 184438 (200K) and 184440 (250K); CIF)

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### Notes

The authors declare no competing financial interests.

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