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Colour-tunable asymmetric cyclometalated Pt(II) complexes and STM-assisted stability assessment of ancillary ligands for OLEDs†

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We combined an asymmetric cyclometalated C[^]N[^]N Pt(II) complex with three different classes of monodentate ancillary ligands, namely a bulky phosphine, a tilted pyridine, and either a tilted or a coplanar isocyanide. The controlled planarization of the coordination environment yields green, orange or red phosphors. Electronic structure calculations using DFT reveal the excimeric intermolecular interactions in the excited triplet state, resulting in tunable photophysical properties. Finally, we assess the suitability of the ancillary ligands for optoelectronic applications by investigating the molecular integrity upon sublimation with the aid of scanning tunneling microscopy (STM). The ancillary ligand controls the surface-induced chiral self-assembly of the molecules on metallic substrates, as well as the availability of loose coordination sites. The concordance between the photoluminescence spectra and electroluminescence profiles for solution-processed and vapour-deposited devices confirm the predictions attained by STM regarding molecular integrity and processability.

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Introduction

Triplet emitters have found application as dopants in optoelectronic devices, such as organic light-emitting diodes¹ (OLEDs) and light-emitting electrochemical cells (LEECs).² Besides Ir(III),³ Rh(III),⁴ Re(I)⁵ and Cu(I),⁶ coordination compounds featuring Pt(II)⁷ and Au(III)⁸ cations have been described displaying strong phosphorescence, fluorescence or thermally-activated delayed fluorescence. The latter possess a d⁸ electronic configuration and a mostly square-planar coordination environment, and have found applications in photocatalysis, bioimaging and therapy.⁹

Both bidentate¹⁰ as well as tridentate¹¹ luminophores have been employed, yielding highly (electro)luminescent coordination compounds. We have also recently shown that neutral Pt(II) complexes bearing dianionic tridentate N[^]N[^]N ligands can be used as triplet emitters, particularly in electroluminescent devices. Depending on the substitution pattern, they can be found either as monomeric¹² or as aggregated phosphorescent species.¹³ For the monomeric species, the emission occurs from metal-perturbed ligand-centered triplet states (³MP-LC), whereas the aggregates emit from excimeric or metal-metal-to-ligand charge-transfer states (³MMLCT).^{11a} Planar species can be also coupled with metallic substrates by hybridization of the protruding d_{z²} orbitals with the surface. Such behavior can be accurately unraveled with the aid of scanning tunneling microscopy (STM) and spectroscopy (STS), as we have recently shown.¹⁴ In supramolecular chemistry, intermolecular interactions give rise to new properties.¹⁵ For instance, M. Kato *et al.*¹⁶ have investigated the capabilities of supramolecular arrays consisting of interacting Pt(II) complexes. De Cola *et al.* have elegantly shown that biomimetic self-assembly of Pt complexes can be monitored in real time and correlated with kinetic and thermodynamic parameters.¹⁷ We have previously shown that tridentate bis-diazole, bis-triazole- or bis-tetrazole-substituted pyridine luminophores (or asymmetric combinations thereof) with pyridine ancillary ligands give rise to fully planar coordination environments, whereas the bulky triphenylphosphine

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† Electronic supplementary information (ESI) available: Synthetic procedures, NMR and x-Ray data, STM and DFT procedures and data, photophysics and OLED fabrication. CCDC 1416852 (C2), 1416853 (C3) and 1416854. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6tc00093b



completely suppresses the intermolecular interactions, due to their out-of-plane phenyl units.^{12,14,18}

We have recently also described phosphorescent organo- and hydrogelators,^{13a,b,e} as well as crystalline arrays in which π -accepting ancillary ligands can switch-on Pt··Pt interactions, regardless of the presence of bulky adamantyl substituents.^{19a} Moreover, 2D-confinement on insulating or semiconducting substrates can also be used to force the interaction between adjacent metallic centers.^{19b} However, a deeper understanding and a defined control are still required, as stacking most frequently causes lowered solubility, undesired triplet–triplet annihilation, lowered quantum yields and unpredictable red-shifts.¹⁵

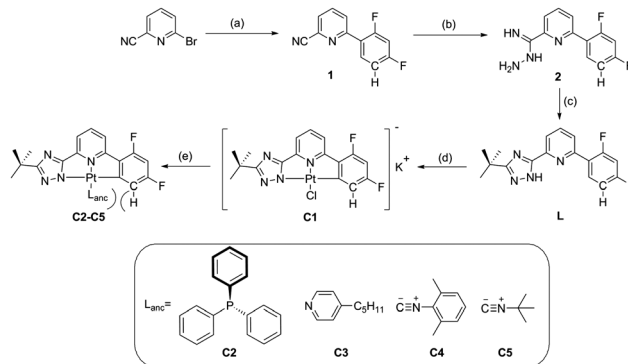
In order to achieve high photoluminescence quantum yields, good σ -donors as well as strong π -acceptors with large ligand field splitting are required to destabilize the otherwise thermally accessible, dark metal-centred states.^{11a} Yersin and co-workers have recently published an interesting concept for bright Pt(II) complexes bearing doubly cyclometallating carborane-substituted luminophores and isocyanide ancillary ligands,²⁰ whereas combinations of C^{^N}N^{^N} luminophores and isocyanides have been used previously by Che *et al.*²¹

In a previous work, we have shown that it is possible to find submolecular electronic set-screws with the aid of scanning tunnelling microscopy (STM) and scanning tunnelling spectroscopy (STS), enabling the realization of deep-blue phosphorescent Pt(II) complexes.^{14c} Herein we present a notably straightforward strategy towards fully tunable intermolecular interactions on the basis of controllable planarization of the coordination environment of Pt(II) complexes. We demonstrate that the introduction of a cyclometallating phenyl ring on a strongly σ -donating 2-(1,2,4-triazol-5-yl)-pyridine chelate yields an outstanding dianionic C^{^N}N^{^N} ligand for luminescent Pt(II) complexes. Moreover, the introduction of a C–H bond adjacent to the ancillary ligand (L_{anc}) forces the out-of-plane tilt of the monodentate unit. Depending on the bulkiness of the fourth ligand, variable tilt angles are achievable, which in turn hinder intermolecular interactions to different extents, yielding green, orange and red phosphors. The excimeric nature of the emissive states of monomeric and dimeric species was assessed by density functional theory (DFT). Finally, we elucidated the integrity of the complexes upon sublimation with the aid of STM, in order to consider the suitability of each of the ancillary ligands, and corroborated the concordance between the photoluminescence spectra and electroluminescence profiles for solution-processed and vacuum-deposited devices.

Results and discussion

Synthesis and structural characterization

The synthetic route (Scheme 1) starts with a Suzuki coupling, leading to a 2-cyanopyridine substituted with a difluorophenyl moiety. The triazole ring was introduced by reaction with hydrazine monohydrate followed by an acylation with trimethylacetyl chloride and a heat-aided ring closure. Cyclometallation with K_2PtCl_4 in acetic acid yielded the anionic complex **C1**, which upon reaction



Scheme 1 Synthetic route towards the C^{^N}N^{^N} ligand and the corresponding Pt(II) complex. (a) 2,4-Difluorophenylboronic acid, Pd(PPh₃)₄, K₂CO₃, THF, N₂, reflux, 24 h. (b) N₂H₄·H₂O, EtOH, rt. (c) (1) *t*-BuC(O)Cl, K₂CO₃, DMF, 0 °C to rt, (2) ethylene glycol, 160 °C. (d) K₂PtCl₄, acetic acid, reflux. (e) L_{anc} , THF, 12 h.

with a neutral monodentate ligand precursor led to complexes **C2–C5**. A detailed description of the conditions, purification methods as well as ¹H, ¹³C, ¹⁹F, ³¹P and ¹⁹⁵Pt NMR characterization can be found in the ESI† (Sections S1.1 and S1.2). **C2** and **C3** yielded crystals suitable for X-ray diffractometric analysis (Section S1.3, ESI†). Complex **C2** presents the formation of dimers through π ·· π -interactions involving only the phenylpyridine moieties of the luminophore (the shortest distance between aromatic rings is 3.437 Å). No interaction between adjacent metallic centers of the dimeric structures was observed (the Pt··Pt distance is 7.33 Å). These dimers are further stabilized by C–H··F and C–H··N interactions in linear chains along the *ac*-plane (2.586 Å and 2.709 Å, respectively, Fig. 1 (left)). In contrast, the packing diagram of **C3** shows the formation of dimers with clear Pt··Pt contacts and stronger π ·· π -interactions involving the entire luminophore (3.485 Å and 3.347 Å, respectively). Also for **C3**, C–H··F and C–H··N interactions were observed in zig-zag chains along the *ac*-plane (2.427 Å and 2.686 Å, respectively, Fig. 1 (right)).

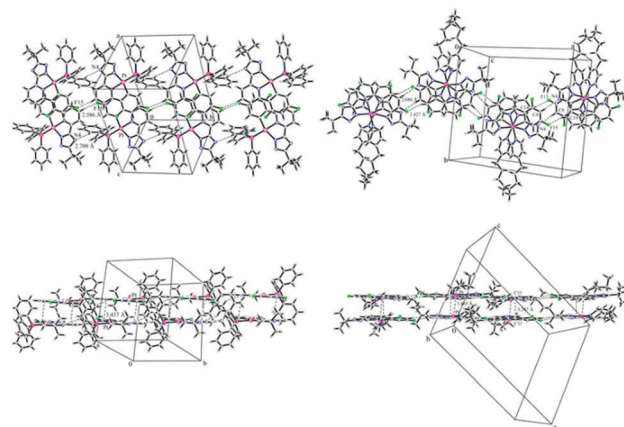


Fig. 1 Left: Packing diagram presenting the C–H··F and C–H··N interactions along the *ac*-plane (top) and π ·· π stacking involving only the phenylpyridine moiety (bottom) in complex **C2**. Right: Packing diagram presenting the C–H··F and C–H··N interactions along the *ac*-plane (top) and Pt–Pt interactions and π ·· π stacking (bottom) in complex **C3**.



Table 1 Representative photophysical properties, binding energies and intermetallic distances for compounds **C2–C5**

Complex	C2	C3	C4	C5
$\lambda_{\text{abs}}/\text{nm}$	217, 324 ^a	253, 323 ^b	262, 327 ^b	256, 313, 338 ^b
$\lambda_{\text{em,solid}}^c/\text{nm}$	503	599	622	698
$\lambda_{\text{em,deacr}}^c/\text{nm}$	508 ^a	504 ^b	511 ^b	507 ^b
$\Phi_{\text{L,solid}}^d (\pm 0.02)$	0.84	0.10	0.14	0.31
$\Phi_{\text{L,deacr}}^d (\pm 0.02)$	0.70 ^a	0.08 ^b	0.71 ^b	0.73 ^b
$\tau_{\text{deacr}}^e/\mu\text{s} (\pm 0.1 \mu\text{s})$	15.2 ^a	2.0 ^b	13.6 ^b	16.5 ^b
$\tau_{77\text{K}}^e/\mu\text{s} (\pm 0.1 \mu\text{s})$	15.2	12.2	10.6	13.5
ΔE^g (eV) S_0/T_1	−0.107/0.513	−0.358/−0.353	−0.255/−0.212	−0.236/−0.167
Pt··Pt distance ^h (Å) S_0/T_1	5.952/2.867	3.927/3.849	3.625/2.822	4.492/2.818

^a In tetrahydrofuran. ^b In dichloromethane. ^c Powder, measured at room temperature. ^d Measured at room temperature in an integrating sphere. ^e Obtained from the bluest emission maximum. Amplitude-averaged lifetimes are given when multiexponential decays are observed. ^f In frozen glassy matrices (2-MeTHF). ^g Dimer binding energies calculated as the difference between the S_0 dimer energy and the energy of two S_0 monomers, or as the difference between the T_1 dimer energy and the sum of the T_1 energy and the S_0 energy of the monomer. ^h Calculated for a S_0 dimer or for a T_1 dimer.

Photophysical characterization

All the complexes were characterized in terms of UV-Vis absorption and emission spectroscopy in the solid state (powders, neat films and doped into PMMA), as well as in solution at room temperature and in frozen glassy matrices at 77 K, including absolute photoluminescence quantum yields (Φ_{L}), time-resolved luminescence decays as well as radiative and non-radiative rate constants. The detailed results are shown in the (Section S4, ESI†). A summary of the most relevant photophysical data is presented in Table 1. The complexes show absorption bands around 220–260 nm, which can be assigned to π – π^* transitions, as well as bands around 320 nm, which can be ascribed to a transition with mixed MLCT/LC character. Furthermore, the long excited state lifetimes ($>10 \mu\text{s}$) and vibrational progressions observed in frozen matrices at 77 K indicate that the emission occurs mainly from $^3\text{MP-LC}$ excited states. All the complexes (**C2–C5**) show green phosphorescence in diluted solutions (10^{-5} M) with high Φ_{L} for deaerated samples (up to 73%, **C5**). Their emission maxima shift only slightly from 504 nm for **C3** to 511 nm for **C4**, indicating a minor contribution of the ancillary ligand to the ground- and the excited states of the chromophore. These experimental results also correlate with the DFT calculations (ESI,† Section S3), showing that the highest occupied molecular orbital (HOMO) as well as the lowest unoccupied molecular orbital (LUMO) are mostly located on the chelating luminophore, with marginal participation of the metallic center (see ESI,† Section S3). In the solid state, complex **C2** shows a blue-shifted, monomeric ($^3\text{MP-LC}$) emission with a slightly blue-shifted maximum at 503 nm, due to the lack of stabilizing solvent molecules (Fig. 1). The bulky ancillary P(Ph)_3 ligand suppresses aggregation, avoiding bimolecular quenching processes, and yielding a high Φ_{L} (84%). On the other hand, complexes **C3–C5** show a characteristic broad, red-shifted $^3\text{MMLCT}$ emission band that is attributed to stacked species. The emission wavelength red-shifts from 599 nm (**C3**) and 622 nm (**C4**) down to 698 nm (**C5**) in powders and neat films, depending on the bulkiness of the ancillary ligand (**C2** vs. **C5**) and its tilt angle with respect to the coordination plane minimizing the steric hindrance with the tridentate moiety (**C3** vs. **C4**). Interestingly, the highest Φ_{L} for the aggregated species was achieved for the deepest red-emitting compound (**C5**, 31%), whereas **C3** and

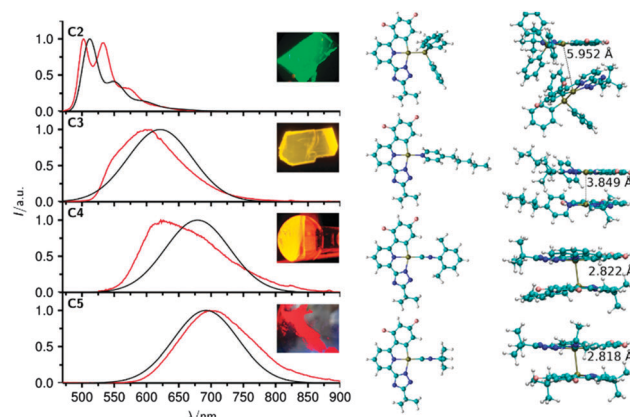
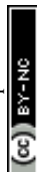


Fig. 2 Left: Normalized emission spectra of complexes **C2–C5** in the solid state (red: experimental; black: calculated). Right: DFT-calculated geometries of monomeric species and Pt··Pt distances of triplet dimers.

C4 have rather low Φ_{L} (10% and 14%, respectively). Pt··Pt interactions occur mainly in the excited triplet state ($^3\text{MMLCT}$ excimer), as indicated by DFT calculations (ESI,† Section S3). Consequently, the binding energies and intermetallic distances of singlet and triplet dimers show that the Pt··Pt distance significantly shrinks in the excited triplet state with respect to the ground state (Table 1). This shortening also nicely correlates with the observed bathochromic shift, and the calculated emission spectra are in good agreement with the experimental observations. The lowest triplet states of the dimers actually show the electron transfer from an intermetallic antibonding (d_{z^2} – d_{z^2}) to a bonding interligand (π^* – π^*) orbital ($^3\text{MMLCT}$). As a consequence of the bulky P(Ph)_3 , however, the **C2** dimer is less stable by 0.51 eV than two monomers (*i.e.*, one triplet plus one ground state), whereas all the other dimers are stable with respect to dissociation. For all the complexes, the binding of the dimers in the ground state can be attributed to van der Waals interactions, as the Pt··Pt distances lie above 3.5 Å (Fig. 2).

STM analysis upon sublimation

The STM images in Fig. 3 show the molecular arrangement of complexes **C2–C5** vapour-deposited onto Au(111) under ultrahigh



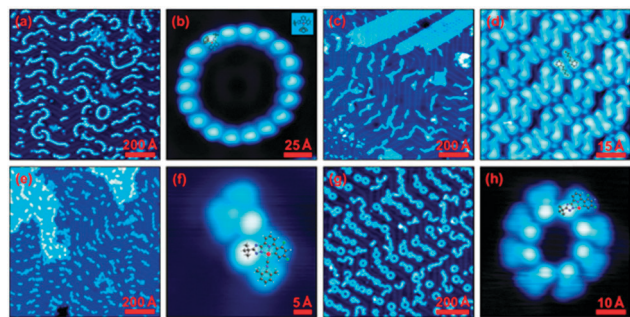


Fig. 3 Overview and close-up STM images of self-assembled monolayers of **C2** (a and b), **C3** (c and d), **C4** (e and f) and **C5** (g and h). While **C2** and **C4** remain stable upon sublimation, **C3** and **C5** have not been found intact on the surface. Larger images are found in the ESI† (Section S2).

vacuum conditions ($p < 10^{-10}$ mbar). **C2** (a and b) was evaporated at 436 K and self-assembles at low coverages into 2D-twisted chains with random length and direction. The submolecular details could not be resolved most likely due to the bulkiness of the $P(Ph)_3$ and therefore, the orientation of the complex remains unclear. However, we assume these complexes to be intact and the ancillary ligand to appear brightest in the images as the measured structures match in size with **C2**. Schneider and others have shown that planar asymmetric aromatic systems without a stereogenic centre become prochiral upon adsorption to a surface.²² This adsorption-induced chirality results in two possible stereoisomers for complex **C2**, which influence the orientation of the growing chain either clock- or anticlockwise (Fig. S5b, ESI†). As seen in the close-up image (b), also a closed ring can be formed, consisting of only one type of stereoisomer. The STM images of **C3** (c and d) show densely packed islands as well as irregularly arranged stripes on the substrate. A close-up image (d) reveals that the monolayer islands consist of paired elongated structures each containing a bright protrusion at one end. Based on the comparison of the features with molecular models and our finding with a similar Pt complex,^{14a} we conclude that the images show only the ancillary ligand (4-ampylpyridine), which indicates a decomposition of **C3**. In this case, the self-assembly is probably favoured by van der Waals interactions. One explanation for the decomposition is the lower dissociation energy of the Pt–N bond due to sterical hindrance (*vide supra*), but also the enhanced planarity maybe relevant, as it favours aggregation and hinders sublimation. However, the evaporation temperature is only slightly increased (443 K) as compared to **C2**. **C4** does not assemble in long-range regular structures at low coverages but seems to prefer pairing of a few complexes along the fcc regions of the Au(111) surface (e). Although the sublimation temperature is again slightly higher (453 K) due to further planarization, no decomposition seems to occur as we only found intact complexes (f). The increased stability is based on the strong isonitrile ligand (both a σ -donor and a π -acceptor). Complex **C5** (g and h) sublimates with the highest temperature (513 K) and assembles analogously to **C2** in rings or twisted chains with prochiral behavior. From the close-up images we deduce that the structures consist only of the tridentate ligand and the Pt-core. As we could not find any indications of the

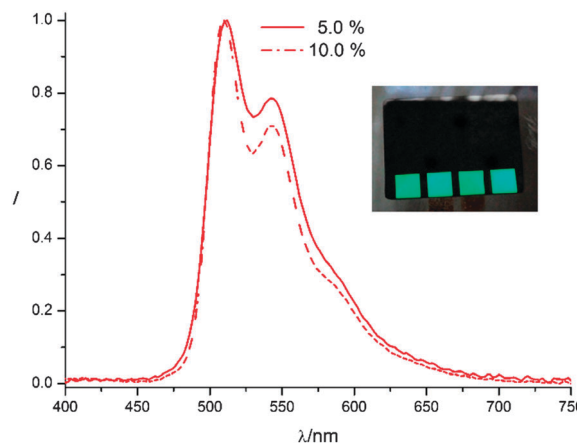


Fig. 4 Normalized electroluminescence spectra of complex **C2** in optimized solution processed (solid) and sublimation processed (dashed) OLEDs.

ancillary ligand adsorbed onto the surface, the decomposition in the evaporator during the sublimation process seems the most probable scenario. These results suggest that a free coordination site is available at the Pt centre, which probably loosely interacts with the triazole moiety of a neighbouring complex (Fig. 3).

OLED fabrication

Our findings indicate that only **C2** and **C4** would make adequate dopants for vapour-deposited devices. Since **C2** shows the highest color purity and Φ_L , it was chosen for a comparative test regarding its ability to perform in electroluminescent devices. In order to find a suitable host material for spin coated devices, cyclic voltammetry, UPS, UV-Vis absorption and photoluminescence emission spectra of neat films from the host material and complex **C2** were measured as well as the host materials doped with 5, 10 and 15 wt% of complex **C2**. The final device architecture and the optimization procedure are found in the ESI† (Section S5). The integrity of the complex itself was confirmed by comparing vapour-deposited and solution-processed OLEDs (Fig. 4), which clearly show an invariable electroluminescence that nicely overlaps with the photoluminescence in the solid state and in solution (see Fig. 1).

Conclusions

In summary, we demonstrated that the controllable intermolecular interaction gives rise to tunable Pt(II) complexes displaying green, orange or red phosphorescence. DFT calculations of monomeric and dimeric species showed the excimeric character of the excited state of red-emitting aggregates, and STM was used to verify the molecular integrity upon thermal sublimation. Combination of steady-state spectroscopic and STM measurements allowed us to assess the best candidate with the highest stability and color purity for the realization of both vapour-deposited and solution-processed electroluminescent devices. DFT calculations helped us to understand the nature of the intermolecular interactions within aggregated species, which is stronger in the excited state than in the ground state, and showed that bulky substituents



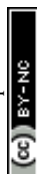
suppress aggregation. These results regarding green and red emitters complement our previous STM/STS-guided development of deep-blue triplet emitters, as required for RGB-based electroluminescent technologies. They also show that the synergetic interaction between synthesis, spectroscopy, theory and microscopy constitute a virtuous approach towards the design and optimization of improved optoelectronic materials. The implementation of phosphorescent stacks in electroluminescent devices by direct coupling with metallic substrates lies at hand, as we have recently observed the hybridization and charge injection at metallic substrates for Pt(II) complexes. The ligand-dependent prochiral assembly with a loose coordination site could be interesting for surface catalysis. A clear understanding of molecular orientation in electroluminescent films might also play a crucial role in the design and realization of new optoelectronic devices, as the outcoupling efficiency is influenced by the relative direction of the transition dipole moments, a phenomenon that has been recently shown by Brütting and Thompson.²³ Further studies regarding different substitution patterns on the luminophoric chelate are underway, as well as the synthesis of multinuclear complexes with enhanced intermolecular interactions. One goal to be achieved is an enhanced radiative rate constant and a reduced excited state lifetime, which would further enhance the efficiency of the devices by avoiding triplet-triplet annihilation especially at high current densities and luminances. Based on the tunable red-shifted wavelengths, applications in bioimaging can be envisaged as well.

Acknowledgements

J. S. wrote the manuscript, synthesized and characterized the ligand precursors and complexes, fabricated and characterized the OLED devices with the help of D. L., N. L. and D. Z.; L. S. carried out the photophysical characterization of the Pt-complexes; P. R. E. and D. W. rendered the STM measurements; C. D. performed the X-ray diffractometric analysis; C. S. and N. L. D. carried out the DFT calculations; C. A. S. conceived the experiments, discussed the results and wrote the manuscript. Financial support from the DFG (SFB-TRR 61, projects B13 and C07) is gratefully acknowledged.

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