

Self-assembly of singlet-emitting double-helical silver dimers: the curious coordination chemistry and fluorescence of bisquinolylpyridone

Received 00th January 20xx,
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

www.rsc.org/

Charlotte M. A. Farrow,^a Geoffrey R. Akien,^a Nathan R. Halcovitch,^a James A. Platts^{*b} and M. P. Coogan^{*a}

2,6-Bis(2-quinolyl)-4(1H)-pyridone **1**, a novel quinoline analogue of the well-known ligand 2-terpyridone, shows unusual fluorescence with a large Stokes shift and low energy emission. Pyridine-pyridone tautomerism is investigated by NMR and theoretical methods and indicates that the low energy emission is from the pyridine form. **1** reacts with Ag(I) salts to give a double helical Ag₂N₆ core showing a BLUE shift in fluorescence with respect to the free ligand, which has been characterised experimentally and theoretically as involving an unusual mixed MLCT/ILCT excited state and emission from a singlet state.

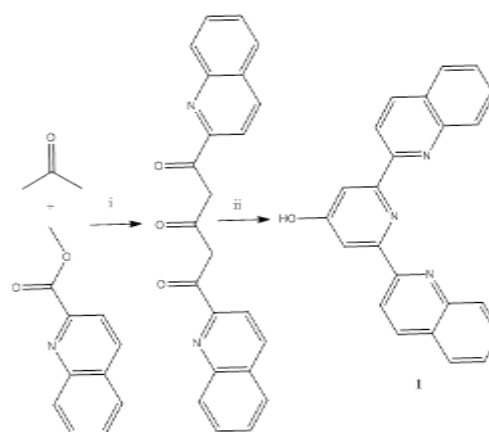
Introduction

Terpyridines have a rich and varied coordination chemistry¹ and amongst these a hydroxylated example 2,6-bis(2-pyridyl)-4(1H)-pyridone² (terpyridone hereafter) is widely applied in coordination chemistry. This ligand forms a variety of types of complex in which it bonds in a tridentate N[^]N[^]N, bidentate N[^]N and bridging N[^]N:N and N[^]N:O modes forming monomeric and supramolecular species including some with interesting photophysical properties.^{1,2,3} It is known that substitution of pyridines for quinolines red-shifts absorption and emission properties of derived metal complexes⁴ and many oligopyridine ligands have been investigated with one or more pyridines replaced with quinolines for this reason. However, the bisquinoline analogue of terpyridone, 2,6-bis(2-quinolyl)-4(1H)-pyridone **1**, is unknown. In light of the rich coordination chemistry and luminescence potentially available an investigation of this ligand was undertaken.

Results and discussion.

Synthesis of Ligand **1**

The synthesis of **1** was achieved by a variation on the Constable method for terpyridone,² reacting methyl quinaldate with acetone in the presence of sodium ^t-butoxide to give 1,5-bis(2-quinolyl)-pentane-1,3,5-trione following a



Scheme 1. Synthesis of 2,6-bis(2-quinolyl)-4(1H)-pyridone **1**. i) NaOtBu^t, THF 60°C 16h; ii) NH₄OAc, MeOH, 60°C, 16h.

double Claisen condensation. We find sodium ^tbutoxide a convenient and less hazardous replacement for sodium hydride in the synthesis of both **1** and 2,6-bis(2-pyridyl)-4(1H)-pyridone and it does not appear to lead to significant loss of yield in either case. 1,5-bis(2-quinolyl)-pentane-1,3,5-trione is recovered following aqueous work up as a fibrous yellow solid and is a difficult material to characterise as it is only sparingly soluble and appears to exist in solution as a gross mix of keto-enol tautomers and / or enol double bond isomers. However, condensation of the material recovered from the double Claisen condensation with ammonium acetate in methanol, following aqueous work up gave **1** in good yield (Scheme 1).

^a Department of Chemistry, Lancaster University, Bailrigg, Lancaster, LA1 4YB,.

^b School of Chemistry, Cardiff University, Main Building, Park Place, Cardiff, CF10 3AT. platts@cardiff.ac.uk

*email m.coogan@lancaster.ac.uk

Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/x0xx00000x

NMR Studies of bis-quinolylpyridone **1**.

As 2-terpyridone has been reported to exist in equilibrium between hydroxypyridine and pyridone forms⁵ an investigation of the tautomerism of **1** was undertaken by NMR. As with terpyridone, **1** predominantly exists in the keto (pyridone) form (C=O δ_c 182 ppm) in the less polar CDCl₃, but primarily in the enol (pyridinol) form in the more polar DMSO-d₆ (C-O δ_c 166.5 ppm).⁶

A closer examination showed some of the peaks to be unexpectedly broad, which made determining the quaternary carbon chemical shifts challenging, and completely prevented the observation of the potentially useful ¹⁵N chemical shifts via ¹H,¹⁵N-HMBC.⁷ Gradual dilution of a 176 mM DMSO-d₆ solution of **1** to 5 mM improved the linewidth (FWHM) of the OH from 68.0 to 17.5 Hz, but slightly *increased* that of the CH from 2.3 to 5.3 Hz, indicating that intramolecular processes are not the main cause of the line-broadening. Instead, titration of a 20 mM solution with small amounts of a 100 mM NEt₃/DMSO-d₆ stock solution was much more successful, with a final 1 mol. % (with respect to the ligand) NEt₃ reducing the linewidths to 0.73 and 0.49 Hz for the OH and CH, respectively. We attribute this improvement to the neutralisation of acidic impurities still remaining after the recrystallization, as evidenced by the appearance of the NEt₃ CH₂ as a quartet of doublets (*J* 7.3, 4.3 Hz) due to coupling with the methyl and NH⁺. Approximately 75% of the impurities were due to residual acetic acid, but 25% could not be directly accounted for. The presence of acid is likely to increase the rate of proton exchange between OH, NH and water and increase the rate of tautomerisation, both of which would serve to increase the linewidths. Recent work has shown that acetic acid solvates are readily accessible,¹⁰ so it is quite possible that small amounts of this crystalline form were simultaneously formed during the synthesis procedure.

4-pyridones are known to self-associate in solution,⁸ but the molecular weight calculated from diffusion experiments on 5–100 mM solutions in DMSO-d₆⁹ compared favourably with its true molecular weight, so it seems likely that this is not a significant factor. This is in keeping with observations that increasing steric bulk decreases the tendency towards self-association.^{8,9}

The greatly improved linewidth in DMSO-d₆ allowed for the identification of the minor keto tautomer by 2D NOESY and its quantification at 1.5 mol. %. The ¹H linewidths of the keto form were notably broader than that of the enol tautomer, but the *S/N* was too low to establish whether this was due to rotational isomerism or simply a relatively higher acidity of the NH over the OH. It was also possible to use ¹⁵N-HMBC to measure ¹⁵N chemical shifts of 279.4 ppm for the central pyridine ring, and 302.0 ppm for the quinoline nitrogen.

The solubility of **1** in CDCl₃ was lower at 10 mM and gave similarly broad peaks, but again, the addition of NEt₃ greatly improved the linewidth, allowing quantification of the minor enol tautomer at 4.5 mol. %, in comparison to 0.9 mol. % for terpyridone at the same concentration. As expected, ¹⁵N-HSQC and ¹⁵N-HMBC experiments yielded substantially different

chemical shifts for the major keto tautomer: 127.3 ppm for the pyridone NH, and 297.2 ppm for the quinolyl N.

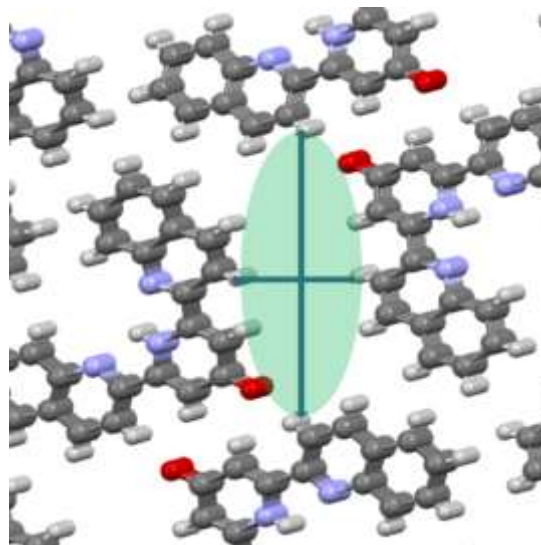


Figure 1. **1** in *I2/a* showing void channels running along *b*-axis.

Structure	Torsion Angles in Pyridone-form	Torsion Angles in Pyridinol-form
1 (<i>I2/a</i>)	2.6(2) and 2.6(2)	
1-a (P21/ <i>n</i>)	2.3(3) and 8.8(3)	21.29(17) and 20.84(17)
1-b (P-1)	1.58(16) and 2.74(16)	5.30(13) and 1.49(13)
	1.82(18) and 5.12(18)	23.35(15) and 13.8(3) (major) and 1.8(4) (minor)

Table 1: Torsion angles (degrees) measured by X-ray diffraction for N_{pyr}-C-C-N_{quin}

Structural Studies of bis-quinolylpyridone **1**.

Crystals of **1** demonstrate polymorphism, those grown from cooling a mixture of EtOH and H₂O give the monoclinic form in space group *I2/a*. In this structure the molecules are all in the pyridone form, with a C-O distance of 1.265(2) Å. This structure also contains continuous voids running along the *b*-axis occupied by solvent molecules. The other two polymorphs were formed by slow diffusion of Et₂O into concentrated solutions of **1** in either CDCl₃ or CD₃CN. From CD₃CN another monoclinic form in P21/*n* is obtained (**1-a**), this time with one molecule in pyridone form and another in the pyridinol form in the asymmetric unit. The C-O bond lengths are 1.264(2) and 1.343(3) Å and the two oxygen atoms are hydrogen-bonded as a pair along the *c*-axis. Finally, from CDCl₃, we have obtained the triclinic polymorph (**1-b**) that has an asymmetric unit containing 4 different molecules, in similar hydrogen-bond pairs. (Figures 1,2).

It is notable that in **1-b** one of molecules in the pyridinol form deviates significantly from planarity. Indeed, in both structures examination of the quinoline arms in relation to the central pyridine in these structures, one can observe (Table 1) that in general the pyridone molecules remain very close to planar,

whereas the pyridinol forms demonstrate larger deviations from planarity.

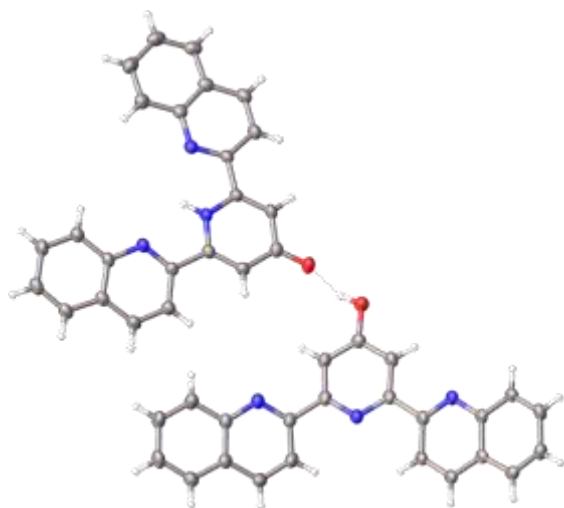


Figure 2: Solid-state molecular structure **1-a** demonstrating the pyridine-pyridyl tautomers.

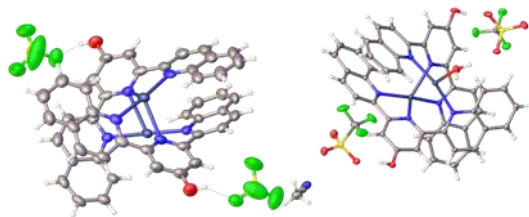


Figure 3: Solid-state molecular structures of **2** and **2-a** (respectively) demonstrating the bridging-helical arrangement of the ligand system.

Silver Complexation of bis-quinolylpyridone **1**.

As terpyridone is known to form a trimeric cage¹¹ upon reaction with rhenium pentacarbonyl halides followed by activation with silver salts, a similar approach was taken, treating **1** with equimolar $\text{Re}(\text{CO})_5\text{Br}$ followed by activation of the solid product with AgBF_4 . However, upon slow diffusion of diethyl ether into an acetonitrile solution of the product of this reaction, instead of a rhenium-based trimer, a dimeric silver species was isolated in which each Ag_2 unit is bound by two molecules of **1**. Closer examination revealed that the Ag_2L_2 unit showed an unusual connectivity and geometry with each of the terminal quinoline nitrogens of each ligand binding a different silver ion, while the central pyridone nitrogen of each ligand bridges between both silvers. The two molecules of **1** are themselves mutually entwined in a double helical arrangement around the central Ag-Ag axis (Figure 3). Although coordination to rhenium had been intended it is known that terpyridines only coordinate to the rhenium *fac*-tricarbonyl core in a bidentate mode and the derived complexes are labile⁴ whereas the isolated product involves

coordination all of the nitrogen donors and thus appears to be the thermodynamically preferred product of the reaction.

In order to demonstrate that the formation of the dimeric helical structure was not a feature of the unusual reaction conditions, ligand **1** was treated with AgBF_4 and AgOTf , and in each case crystals were obtained in which the double helical Ag_2L_2 structure was repeated (Figure 3). A number of other examples of Ag dimers featuring a single nitrogen bridging between the two centres are known¹² including a similar helical example derived from a chiral terpyridine in which the

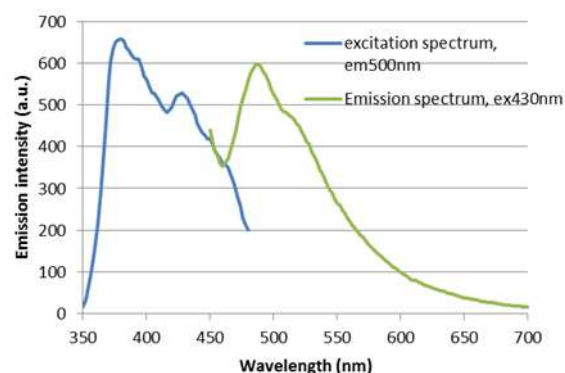


Figure 4: Excitation and emission spectra of ligand **1**.

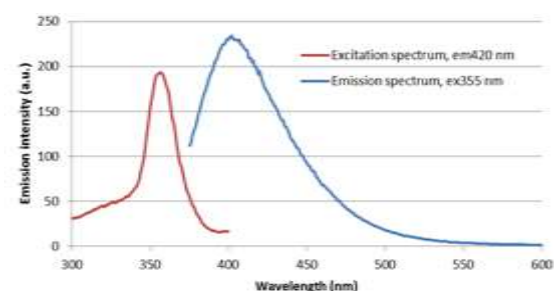


Figure 5: Excitation and emission spectra of complex **2**.

handedness of the helix is controlled by the ligand in solution, but not in the solid state.¹³

Photophysical Studies of bis-quinolylpyridone **1** and Silver Helicate **2**.

Formation of helicates from coordination of oligopyridines with a variety of metals is well precedented,^{1,13} but unexpectedly **2** was found to show intense luminescence, prompting an investigation of the photophysical properties of both **1** and **2**. Ligand **1** shows a remarkably low energy emission for a simple organic molecule, with λ_{max} emission 485 nm, with broad structured emission bands extending beyond 600 nm (Figure 4) giving visually yellow emission. In light of the unusually large Stokes shift the emission lifetime of **1** was investigated by Time Correlated Single Photon Counting (TCSPC) methods and found to have an intensity-weighted average lifetime of 1.4 ns, strongly indicating fluorescence from a singlet excited state. This suggests that the large Stokes shift derives from a geometrical arrangement in the singlet excited state, rather than a change of electronic configuration.

The UV-vis absorption spectrum (see ESI) of complex **2** showed an unexpectedly low energy band appearing as a shoulder on the more intense higher energy absorptions with an apparent maximum around 350 nm. Upon excitation at 350 nm, **2** unexpectedly showed a broad emission band centred around 440 nm and extending to much lower energy (Figure 5). Further investigation revealed an excitation maximum at 355 nm and a series of emission bands of varying intensity contributing to the low energy emission profile. Unusually, it appears that upon formation of complex **2** the emission profile of ligand **1** suffers a *blue*-shift, presumably as

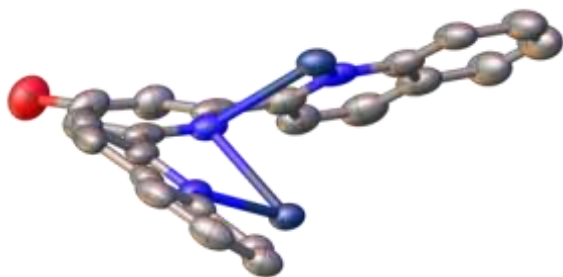


Figure 6. Simplified solid-state structure of **2** with anions and hydrogen atoms omitted, highlighting the torsion angles between pyridine and quinoline moieties.

π - π^* in nature (Figure 7, predicted spectra and orbital and compositions are reported in ESI).

Large Stokes shift and tailing emission bands are often indicative of *d*-metal involvement in photophysics, so the photophysics of **2** were also examined with theoretical methods. TD-DFT prediction of the absorption spectrum is in broad agreement with experiment, with lowest energy band predicted to lie at 330 nm, a transition dominated by HOMO-LUMO excitation. As shown in Figure 8, the former is largely metal *d* in character, with some contribution from N lone pairs, while the latter (see Figure 9) is a π^* ligand-based

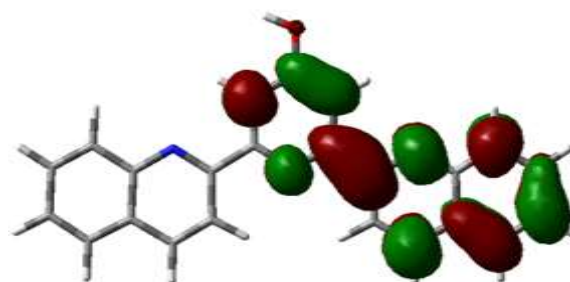
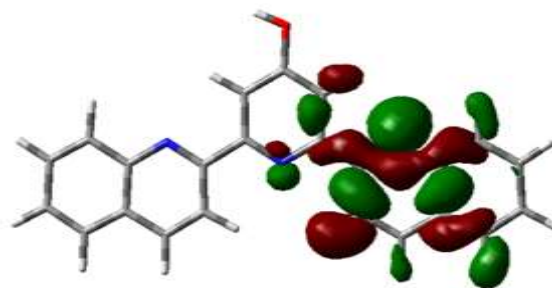


Figure 7. HOMO-4 (top) and LUMO (bottom) of **1** (pyridinol form, ZINDO).

orbital, such that excitation is assigned to be MLCT. Singlet emission, following relaxation of the first excited state, is predicted to come at 497 nm (TD-DFT) or 443 nm (ZINDO), the latter again in better agreement with experiment. Once again, this transition involves HOMO-LUMO combination, but at the optimal geometry of the excited state, the character of these orbitals changes. Now, both are ligand based, HOMO localised mainly on quinolines and LUMO on pyridines, with small contributions from Ag, such that the photophysics of **2** seems to be best characterised as ILCT emission resulting from MLCT/ILCT absorption.

In light of the theoretical finding that a singlet excited state gave the best fit for the observed emission bands, regardless of the large Stokes shift we re-examined the photophysical properties of complex **2** using time resolved methods. TCSPC

Structure	Torsion angle (degrees) for N _{pyr} -C-C-N _{quin}
2	31.5(7); 30.9(7); 37.8(6); 25.9(8) and 30.2(7); 30.8(7); 30.8(7); 31.2(8)
2-a	Chelate: 22.1(3); 27.5(3); Bridge: 34.5(3); 42.6(3)

Table 2: Torsion angles in silver complexes measured by X-ray diffraction

a result of the loss of planarity leading to reduced conjugation and thus higher energy photophysics (average C-N-N-C torsion angle quinoline – pyridine = 33°, overall planar offset quinoline – quinoline = 66°, see figure 6, Table 2). However, the emission profile observed is highly unusual and in no way characteristic of a single isolated quinoline ring coordinated to a metal which is not photophysically active, with a Stokes shift of around 100 nm and structured emission tailing to > 550.

Theoretical Studies of bis-quinolylpyridone **1** and Silver Helicate **2**.

In light of the unexpectedly large Stokes shift of ligand **1** and the blue-shifted photophysics of its complex, **2**, theoretical calculations were used to probe the structure and photophysics of **1** and **2** in more detail. DFT in simulated CH₃CN indicates that pyridone tautomer is 6 kJ mol⁻¹ lower in energy than the pyridine form. Both forms are predicted to have similar absorption (strong bands at 296 nm for pyridone, 283 nm for pyridine) and singlet emission (376 nm for pyridone, 388 nm for pyridine) spectra, where all bands are π - π^* in nature. Given the poor agreement between experiment and theory for emission, we also tested the performance of the semi-empirical ZINDO approach: this predicts singlet emission at 364 nm for pyridone and 496 nm for pyridine. The latter is in striking agreement with the main peak in Fig. 3. ZINDO emission for the **1** pyridinol form at 496 nm is dominated by transition HOMO-4 \leftarrow LUMO, at the excited state geometry: as shown by orbital plots this is essentially

measurements in both solution and solid state gave identical results indicating an intensity-weighted average lifetime of 2.5 ns, strongly suggestive of singlet emission, regardless of the involvement of a second row transition metal. The close match between lifetime in the solid and solution states suggest that the emissive species is the same in solution and solid states, i.e. that the structure of the helicate observed crystallographically is maintained in solution. As it is important to establish that the complex maintains the dimeric structure in solution, and mass spectrometry did not identify an intact $[\text{Ag}_2\text{L}_2]$ ion (showing only a variety of fragments the largest of which matches an $[\text{AgL}_2]$ ion) 1H NMR DOSY experiments were undertaken to assess the solution speciation. As the complex is insoluble in solvents for which reliable DOSY molecular mass determination is established internal standards were used (see ESI) and indicate that the molecular mass of the solution species closely matches that of a mass standard chosen to be close to that of the helicate and far greater than a standard approximately the mass of a monomeric complex (See ESI).

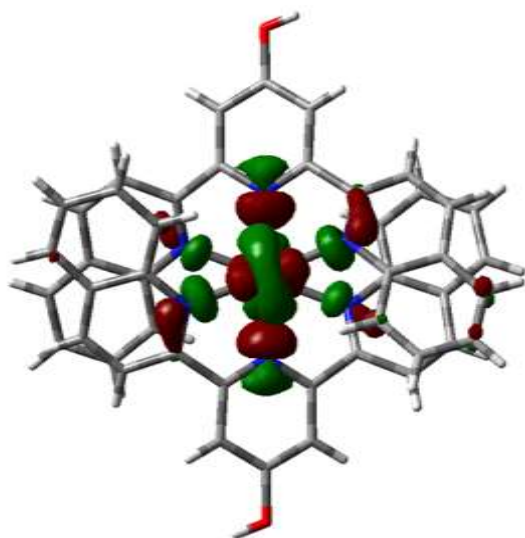


Figure 8. Ground state HOMO of **2** (ZINDO).

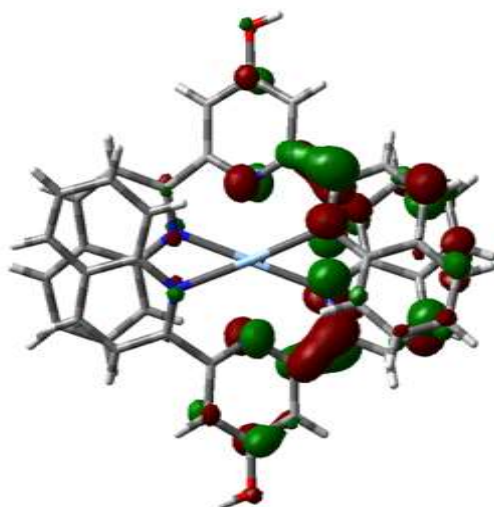


Figure 9. Ground state LUMO of **2** (ZINDO).

Further evidence is provided by the considerable ^{15}N chemical shift changes on coordination of nitrogen to silver. The helicate **2** gave ^{15}N shifts of 247.8 for the pyridone and 268.0 for the quinoyl group, or -31.6 and -34.0 ppm compared to the free ligand in DMSO-d_6 . These coordination shifts are entirely typical for Ag-N complexes.⁷ While the solubility of **1** in MeCN-d_3 was too low to measure ^{15}N shifts in the same solvent, the coordination shift is more than an order of magnitude larger than any known solvent-induced shift changes.

Given the unusual structure of the Ag_2N_6 core in **2**, we also examined the bonding and stabilisation of this complex using theoretical methods. Atoms-in-Molecules (AIM) analysis located a bond critical point (bcp) corresponding to an Ag-Ag bond ($\rho_{\text{pcb}} = 0.024$ au). Each Ag also has two strong Ag-N ($\rho_{\text{pcb}} = 0.079$ au) bonds to quinoline N and two weaker Ag-N ($\rho_{\text{pcb}} = 0.036$ au) to pyridine N, supporting the assignment of the latter as bridging interactions. In addition, stacking interactions between quinolines are found (8 bcp's with ρ_{pcb} between 0.005 and 0.007 au). Decomposing **2** into constituent parts sheds more light on the stability of this complex: breaking **2** into two 'AgN₃' species in simulated CH_3CN has $\Delta G = +123$ kJ mol⁻¹, i.e. **2** is strongly stabilised due to Ag-Ag and Ag-N bonding. In contrast, stacking does not contribute significantly to overall stabilisation: removing Ag⁺ species from **2** and to estimate ΔG for $\text{L}_2 \rightarrow 2\text{L}$ results in -23 kJ mol⁻¹.

Conclusions

2,6-bis(2-quinoyl)-4(1H)-pyridone **1**, exists in solution as a mixture of tautomers with the pyridone tautomer favoured by less polar solvents and exists in the solid state in a variety of polymorphs including one which shows solvent-filled channels. Solutions of **1** show intense room temperature luminescence with an unusually large Stokes shift and low energy emission which is assigned to a π - π^* transition of the hydroxypyridine tautomer. Coordination of **1** to Ag(I) gives binuclear double helicates which themselves show luminescence, which is, unusually, blue-shifted compared to the ligand. The emission from the binuclear helicates originates from singlet ILCT emission resulting from MLCT/ILCT absorption. Future studies will explore the coordination chemistry of this new ligand with other metal ions. In light of the fluorescence of the compounds reported herein it is possible that more soluble analogues may have applications in, *inter alia*, fluorescent cell imaging.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank Dr. David Rochester, Lancaster University, for Mass Spectrometry.

Experimental

All starting materials, reagents and solvents were purchased from commercial suppliers and used as supplied unless otherwise stated. ^1H , ^{13}C , and ^{15}N NMR were recorded at 400, 101 and 41 MHz respectively on a Bruker Avance III 400 equipped with a broadband observe probe (BBO) and referenced to the deuterium lock shift unless otherwise reported. Chemical shifts are reported in ppm, and coupling constants in Hz. IR spectra were recorded on an Agilent Technologies Cary 630 FTIR Spectrometer as thin films or solids and are reported in wavenumbers (cm^{-1}). UV vis spectra were recorded on an Agilent Technologies Cary 60. Steady state emission and excitation spectra were recorded on an Agilent Technologies Cary Eclipse. Time-resolved spectra were recorded on a PicoQuant FluoTime 300 exciting with an LDH-P-C-375 and decays analysed with the program FluoFit.

1,5-bis(2-quinolyl)-pentane-1,3,5-trione.

To a flame dried round bottom flask equipped with a stirrer bar and nitrogen bubbler was added methyl quinaldate (2.00 g, 10.7 mmol) and sodium t-butoxide (2.50 g, 26.0 mmol) and THF (30 ml). To the resulting slurry was added acetone (375 μL , 295 mg, 5.10 mmol) and the resulting thick red suspension was heated at 60 °C for 16 h. To the cooled mixture was added NH_4Cl (sat. aq. 100 ml), the mixture stirred vigorously for 10 minutes and the allowed to stand for 1 h. The solid orange product was collected by filtration, dried under vacuum. The crude product was recrystallised from toluene, separated by vacuum filtration, washed with diethyl ether and dried in air to give the title compound as an orange fibrous solid (888 mg, 2.41 mmol, 47%). A second crop was obtained upon evaporation of the mother liquor over 72 h in air, identical to the material above (372 mg, 1.01 mmol, 19%). Total yield of recrystallised material, 66%. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 8.32 (1H, d, $J=8.3\text{Hz}$) (H1,1'), 8.21 (1H, d, $J=5.0\text{Hz}$) (H5,5'), 8.19 (1H, d, $J=5.0\text{Hz}$) (H4,4'), 7.89 (1H, d, $J=8.0\text{Hz}$) (H6,6'), 7.79 (1H, t, $J=13.8\text{Hz}$) (H2,2'), 7.64 (1H, t, $J=13.7\text{Hz}$) (H3,3'), 7.08 (0.7H, s) (H7'), 2.31 (2H, s) (H7). IR ν max: 1723 (C=O), 1604, 1498, 1451, 1243 cm^{-1} . Due to low solubility this material was used without further characterisation.

2,6-Bis(2-quinolyl)-4(1H)-pyridone 1.

To a stirred suspension of 1,5-bis(2-quinolyl)-pentane-1,3,5-trione (888 mg, 24.1 mmol) in methanol (25 ml) at 60 °C was added ammonium acetate (1.50 g, 19.4 mmol) and the stirred mixture heated at 60 °C for 66 h. After cooling to room temperature, the reaction mixture was treated with water (100 ml) and a yellow gel-like substance separated by prolonged vacuum filtration. The product was recrystallised from an ethanol acetone mixture to give the title compound as a white solid (595 mg, 1.77 mmol, 74%). ^1H NMR (400 MHz, CD_3OD): δ 8.51 (1H d, $J=8.1\text{Hz}$, H8,8') 8.28 (1H, d, $J=8.2\text{Hz}$, H4,4') 8.20 (1H, d, $J=8.2\text{Hz}$, H3,3') 8.03 (1H, d, $J=8.2\text{Hz}$, H5,5') 7.91 (1H, m, H6,6') 7.72 (1H, m, H7,7') 7.33 (1H, s, pyridone H3,3') $^{13}\text{C}\{^1\text{H}\}$ NMR (400 MHz, $\text{DMSO}-d_6$): δ 166.6, 156.7, 155.6, 147.6, 137.5, 130.5, 129.7, 128.6, 128.4, 127.6, 119.3, 109.8. IR ν max: 3271, 3055, 1634 (C=O), 1564 1502, 1429, 1258 cm^{-1} . m/z (ESI, HRMS 350.1277, $\text{C}_{23}\text{H}_{15}\text{N}_3\text{O}$ predicts

350.1288. $\text{C}_{23}\text{H}_{15}\text{N}_3\text{O}\cdot 2\text{H}_2\text{O}$ requires C 71.7 H 4.97 N 10.9 found C 71.7 H 4.67 N 11.1 %

Disilver(I)-bis(2,6-bis(2-quinolyl)-4(1H)-pyridonyl) tetrafluoroborate 2.

To a round bottom flask was added ligand **1** (67.0 mg, 0.191 mmol) and silver (I) tetrafluoroborate (40 mg, 0.206 mmol) and acetonitrile (20 ml) and the mixture heated at reflux for 5 minutes during which time the reactants passed into solution. The mixture was then concentrated under vacuum to 2 ml and treated with diethyl ether (2 ml) giving a white precipitate of the title compound (68 mg, 0.128 mmol, 67%). ^1H NMR (400 MHz, CD_3CN): δ 8.07 (1H, d, $J=8.1\text{Hz}$, H8,8') 7.80 (1H, d, $J=7.7\text{Hz}$, H5,5') 7.63 (1H, s, pyridine H3,3') 7.58 (3H, m, H3,3',4,4',7,7') 7.49 (1H, m, H6,6'). $^{13}\text{C}\{^1\text{H}\}$ NMR (400 MHz, CD_3CN , saturated solution, some peaks not observed): δ 206.6, 138.8, 131.2, 129.4, 128.4, 128.2, 128.1, 112.6. m/z (ESI, HRMS) observed 805.1483 $\text{C}_{46}\text{H}_{30}\text{N}_6\text{O}_2\text{Ag}$ (L_2Ag) requires 805.1476. $\text{C}_{46}\text{H}_{30}\text{N}_6\text{O}_2\text{Ag}_2\text{B}_2\text{F}_8\cdot \text{H}_2\text{O}$ requires C 50.0 H 2.92 N 7.60 found C 49.8 H 2.71 N 7.99 %

Theoretical Calculations

Geometry of **1** and **2** were fully optimised without any symmetry constraint using the M06-2X functional¹⁴ and a basis set consisting of 6-31G(d) on light atoms¹⁵ and Stuttgart-Dresden basis/ECP on Ag.¹⁶ TD-DFT calculations were performed using the range-separated CAM-B3LYP method,¹⁷ with the same basis set. All such calculations used Gaussian09.¹⁸ ZINDO/S¹⁹ calculations on DFT optimised geometries were carried out using the ORCA package.²⁰

Notes and references

For X ray data in CIF format see CCDC deposition numbers 1572595-1572599.

- See e.g. A. M. W. Cargill Thompson, *Coord. Chem. Rev.*, 1997, 160, 1; E. C. Constable, *Chem. Soc. Rev.*, 2007, 36, 246-253; H. Hofmeier and U. S. Schubert, *Chem. Soc. Rev.*, 2004, 33, 373-399; U. S. Schubert and C. Eschbaumer, *Angew. Chem. Int. Ed. Eng.*, 2002, 41, 2892; A. Wild, A. Winter, F. Schlutter and U. S. Schubert.
- E. C. Constable and M. D. Ward, *J. Chem. Soc., Dalton Trans.*, 1990, 1405.
- M. A. Hayes, C. Meckel, E. Schatz, M. D. Ward, *J. Chem. Soc. Dalton Trans.* 1992, 703.
- S. Hostachy, C. Policar and N. Delsuc, *Coord. Chem. Rev.*, 2017, 351, 172; L. Wei, J. W. Babich, W. Ouellette and J. Zubieta, *Inorg. Chem.*, 2006, 45, 3057; R. H. Platel, M. P. Coogan and J. A. Platts, *Dalton Trans.*, 2015, 44, 1170.
- E. Murguly, T. B. Norsten and N. Branda, *J. Chem. Soc., Perkin Trans. 2*, 1999, 2789-2794.
- S. Wang, J. Chen, L. Li, *Int. Res. J. Pure Appl. Chem.*, 2016, 11, 1-6.
- L. Pazderski, *Ann. Rep. NMR Spectrosc.*, 2013, 80, 33-179.
- P. Beak, J. B. Covington, S. G. Smith, J. M. White, J. M. Zeigler, *J. Org. Chem.*, 1980, 45, 1354-1362.
- R. Evans, Z. Deng, A. K. Rogerson, A. S. McLachlan, J. J. Richards, M. Nilsson, G. A. Morris, *Angew. Chem. Int. Ed.*, 2013, 52, 3199-3202.
- P. Florio, C. J. Coghlan, C.-P. Lin, K. Saito, E. M. Campo, W. R. Jackson, M. T. W. Hearn, *Aust. J. Chem.*, 2014, 67, 651-656.

- 11 M. P. Coogan, V. Fernandez-Moreira, B. M. Kariuki, S. J. Pope and F. L. Thorp-Greenwood, *Angew. Chem. Int. Ed. Engl.* 2009, **48**, 4965-8.
- 12 I. Bassanetti, M. Mattarozzi, M. Delferro, T. J. Marks, L. Marchio; *Eur. J. Inorg. Chem.* 2016, 2626, doi:10.1002/ejic.201501236; M. P. Carranza, B. R. Manzano, F. A. Jalon, A. M. Rodriguez, L. Santos, M. Moreno; *Inorg. Chem.* 2010, **49**, 3828, doi:10.1021/ic902563d; A. Cingolani, Effendy, F. Marchetti, C. Pettinari, R. Pettinari, B. W. Skelton, A. H. White; *Inorg. Chem.* 2004, **43**, 4387, doi:10.1021/ic0497376; F. Hintermaier, S. Mihan, M. Gerdan, V. Schunemann, A. Trautwein, W. Beck; *Chem. Ber.* 1996, **129**, 571, doi:10.1002/cber.19961290515]
- 13 G. Baum, E. C. Constable, D. Fenske, C. E. Housecroft, T. Kulke; *Chem. Commun.* 1998, 2659; F-L. Hu, H-F. Wang, D. Guo, H. Zhang, J-P. Lang and J. E. Beve, *Chem. Commun.*, 2016, **52**, 7990-7993; Zhen-Hong Wei, H-X. Li, W-H. Zhang, Z-G. Ren, Y. Zhang, J-P. Lang and B. F. Abrahams, *Inorg. Chem.*, 2008, **47**, 10461-10468.
- 14 Y. Zhao and D. G. Truhlar, *Theor. Chem. Acc.*, 2008, **120**, 215.
- 15 R. Ditchfield, W. J. Hehre, and J. A. Pople, *J. Chem. Phys.*, 1971, **54**, 724; P. C. Hariharan and J. A. Pople, *Theor. Chem. Acc.*, 1973, **28**, 213.
- 16 D. Andrae, U. Haeussermann, M. Dolg, H. Stoll, and H. Preuss, *Theor. Chem. Acc.*, 1990, **77**, 123.
- 17 T. Yanai, D. Tew, and N. Handy, *Chem. Phys. Lett.*, 2004, **393**, 51.
- 18 Gaussian 09, Revision C.01, M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, G. Scalmani, V. Barone, B. Mennucci, G. A. Petersson, H. Nakatsuji, M. Caricato, X. Li, H. P. Hratchian, A. F. Izmaylov, J. Bloino, G. Zheng, J. L. Sonnenberg, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, T. Vreven, J. A. Montgomery, Jr., J. E. Peralta, F. Ogliaro, M. Bearpark, J. J. Heyd, E. Brothers, K. N. Kudin, V. N. Staroverov, T. Keith, R. Kobayashi, J. Normand, K. Raghavachari, A. Rendell, J. C. Burant, S. S. Iyengar, J. Tomasi, M. Cossi, N. Rega, J. M. Millam, M. Klene, J. E. Knox, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. W. Ochterski, R. L. Martin, K. Morokuma, V. G. Zakrzewski, G. A. Voth, P. Salvador, J. J. Dannenberg, S. Dapprich, A. D. Daniels, O. Farkas, J. B. Foresman, J. V. Ortiz, J. Cioslowski, and D. J. Fox, Gaussian, Inc., Wallingford CT, 2010.
- 19 F. Neese, *Wiley interdisciplinary Reviews - Computational Molecular Science*, 2012, **2**, 73.
- 20 M. C. Zerner, G. H. Lowe, R. F. Kirchner, and U. T. Mueller-Westerhoff, *J. Am. Chem. Soc.*, 1980, **102**, 589.