Hirshfeld Surface Analysis of anisylresorcinarene and Luminescence Studies of its Benzimidazoly Lappend Ru(II) Polypyridyl Complexes

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Abstract: The macrocyclic precursor, 2,8,14,20-tetraakis(4-methoxyphenyl)-4,6,10,12,16,18,22,24-octahydroxyalicyclic[4]resorcinarene (P1), has been synthesised and characterised using IR, NMR, ESI mass spectroscopy and single crystal X-ray diffraction analysis. P1 crystallizes in the monoclinic crystal system with P2_1/c space group. Its Hirshfeld surface analysis has been carried out. The other precursor, 2-(pyridin-2-yl)-1H-benzimidazole (P2) and complex precursors, dichlorobis(phenanthroline)ruthenium(II) hydrate (P3) and dichlorobis(2,2'-bipyridine)ruthenium(II) hydrate (P4) are prepared by the modification of reported procedures. The macrocyclic ligand, benzimidazolyl functionalized anisylresorcinarene (L), has been synthesised by the aminomethylation of P1 using P2 and characterised using IR and ESI mass spectrometry. Tetranuclearruthenium(II) phenanthroline and bipyridine complexes of L have been synthesized and characterized by IR, ESI-MS and electronic absorption spectroscopy. Upon excitation at the excitation maxima the complexes emit from the 2MLCT state at 600 and 624 nm, respectively. The emission band of complexes at 77 K in frozen CH_2CN is hypsochromically shifted to 594 and 616 nm, respectively with noticeable increase in the emission intensity and in quantum yields. The complexes show monoexponential decay profile in acetonitrile and have life times of 142 and 200 ns, respectively.

Keywords: Resorcinarene; Ru(II) complex; 2-(pyridin-2-yl)-1H-benzimidazole; Hirshfeld surface analysis

1. Introduction

The coherent design and synthesis of novel architectures are of current interest in the field of supramolecular chemistry [1]. The resorcinarenes, which are calixarenes formed from resorcinol building blocks linked by methylene groups [2], are utilized as supramolecular assemblies which coordinate through all three of their constituent parts: the lower rim, the phenolic groups, and the ortho position on the aromatic rings [3]. Mannich reaction is one of the main reactions employed in functionalization of resorcinarenes with polypyridylic ligands [4]. The rich photochemistry of polypyridylic ligands with almost all metals is related with π-back bonding. The luminescent properties of ruthenium(II) complexes of phenanthroline and bipyridine have been extensively studied due to their significant MLCT absorption in the visible region, their ability to undergo MLCT excitations, and photoreactivity of the MLCT excited states [5]. The current investigation involves the synthesis, characterization and luminescence studies of tetraneuclearruthenium (II) bis(phenanthroline) and bis (bipyridine) complexes of benzimidazolyl functionalized anisylcalixresorcinarene. The Hirshfeld surface analysis of anisylresorcinarene crystal is also carried out to understand the intermolecular interaction present in the molecule.

2. Experimental Section

2.1 Reagents

Anisaldehyde, cyanopyridine, phenanthroline, 2,2'-bipyridine, ruthenium(III) chloride trihydrate, resorcinol, o-phenylenediamine, o-phosphoric acid, sodium perchlorate monohydrate (Aldrich), ammonia, hydrochloric acid (35%), and lithium chloride (Fluka) were purchased and used as such.

2.2 Physical measurements

Single-crystal X-ray structure data collections were performed on a Bruker AXS Kappa Apex II CCD diffractometer with graphite monochromatic Mo (Kα) radiation (λ) 0.71073 Å. Infrared spectra were recorded on a Perkin-Elmer spectrometer RX-1 FT-IR Spectrometer in the range of 4000-400 cm⁻¹. The electrospray ionization mass spectra were recorded on a micromass Quattro II triple quadrupole mass spectrometer. ¹H NMR was recorded on a Bruker AVANCE III 500 MHz (AV 500) multinuclear NMR spectrometer at 25 °C. The electronic absorption spectra were recorded on a Shimadzu UV-2450 UV-Visible spectrophotometer controlled by the UV Probe version-2.33 software. Fluorescence spectra were recorded on a Fluorolog-3 FL3-221 spectrofluorometer. The emission lifetimes were carried out using nanosecond laser flash photolysis (Applied Photophysics, U.K.). Emission quantum yields (Φ) were calculated by integrating the area under the luminescence curves and by using equation 1 [6] where OD is optical density of the compound at the excitation wavelength and A is the area under the emission spectral curve. The standard used for the luminescence quantum yield measurements was [Ru(bpy)_3]Cl_2 (Φ = 0.04) [7] and corrected for the refractive index of the solvent.

\[ \Phi = \frac{(OD_{std}A_{sample})^2}{(OD_{sample}A_{std})^2}\Phi_{std} \quad eq.1 \]

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Scheme 1: Synthesis of anisylresorcinarene (P1) and the ligand (L)

Scheme 2: Synthesis of tetranuclear ruthenium(II) complexes of L

2.3 Synthesis of precursor, anisylcalix[4]resorcinarene (P1)

To a solution of resorcinol (2.75 g, 25 mmol) in ethanol-water (7:3 v/v, 50 mL), was added hydrochloric acid (35%, 10 mL) stirred at 70 °C for 1 h. A solution of anisaldehyde (3.04 g, 25 mmol) in ethanol(50 mL) was added to the reaction mixture followed by water (30 mL). The resulting suspension was refluxed under stirring for 100 h and then cooled to room temperature. The solid product that separated out was filtered, washed with water, and dried to obtain a pale orange powder. The product formed was recrystallized in DMF and characterised.

Figure 1: 500 MHz $^1$H NMR spectrum of P1 in DMSO-$d_6$ at 25°C

Yield 4.85 g (85%), pale yellow powder, mp>300 °C (dec.). Analytical data calculated for C$_{56}$H$_{48}$O$_{12}$($M_r$ = 913): C, 73.21; H, 5.18. Found: C, 73.67; H, 5.30. IR data $\nu_{max}$/cm$^{-1}$ 3373 $\nu$(O–H), 2932 $\nu$(C–H) (aromatic), 2833 $\nu$(C–H)
(aliphatic), 1658, 1608 and 1510 δ(C–C) (aromatic), 1430 δ(C–H) (aliphatic), 1247 ν(C–O), 822 γ(C–H) (aromatic). 1H NMR data (500 MHz, DMSO-d6, 298 K) δ 5.44 (4H, s, CH2), 5.55 (4H, d), 6.09 (4H, s), 6.27 (8H, d), 6.50 (8H, m), 8.45 (8H, m). ESI MS: m/z 913 [M]+, 936 [M+Na]+.

The volume of the filtrate was reduced to 5 mL and 50 mL of sodium perchlorate solution was added with stirring whereupon a dark red solid separated out. The product was filtered, dried and recrystallized in hot acetonitrile.

**2.4 Synthesis of ligand (L)**

To a solution of P1 (1 g, 1.09 mmol) and 2-(pyridin-2-yl)-1H-benzimidazole (P2) (0.85 g, 4.38 mmol) in a solvent mixture of 2-methoxyethanol-ethanol (2:1 v/v, 60 mL), was added 37% aqueous solution of formaldehyde (0.12 ml, 4.38 mmol) under argon atmosphere and refluxed for 48 h. The resulting solution was flash evaporated to get red precipitate and resulting solution was flash evaporated to get red precipitate and recrystallized.

Yield 6.3 g (72 %) Red powder, Mp>300 °C (dec.). Analytical data calculated for C16H18N2O4 (M, = 274): C, 74.47; H, 4.86; N, 9.65. Found: C, 74.35; H, 4.68; N, 9.58. IR spectrum: ν(C–H) (aliphatic), 2926 ν(C–H) (aromatic), 2833 ν(C–H) (aromatic), 1605, 1510 ν(C=C) (aromatic), 1466 δ (CH2–N), 1246 ν(C–O), 745 ω(-CH).

ESI MS: m/z 1227 [M-(2P2+4OCH3+H)]+; 1257 [M-(2P2+3OCH3+H)]+; 1354 [M+(H)+2P2]+.


A solution of the ligand L (0.17 g, 0.10 mmol) and [Ru(phen)2Cl2] 2H2O (0.21 g, 0.4 mmol) or [Ru(bpy)2Cl2] 2H2O (0.20 g, 0.4 mmol) in DMF (50 mL) was refluxed with stirring under an argon atmosphere for 12 h. The resultant dark suspension was cooled and filtered.
3. Results and Discussion

3.1 Synthesis and Characterisation of organic precursors, ligand and complexes.

The acid catalyzed condensation of resorcinol (1 equiv) with 4-methoxybenzaldehyde (1 equiv) upon refluxing in ethanol-water mixture gives P1(Scheme 1). It is characterized by CHN analysis, IR, NMR, ESI mass spectroscopy and single crystal x-ray diffraction analysis. The absence of aldehyde (C=O) stretching vibration and presence of intermolecular hydrogen bonded hydroxyl (-OH) stretching in FT-IR spectrum of P1 confirms the condensation reaction. The 'H NMR spectrum shows resonance signals for aromatic, hydroxyl, methoxy and methine bridge protons (Figure 1). The mass spectrum of P1 shows a peak at m/z 913 due to molecular ion species, [M]+ (C_{34}H_{34}O_{12}H) (Figure 2). The compound 2-(pyridin-2-yl)-1H-benzimidazole (P2) was prepared by the condensation of equimolar amount of 2-cyanopyridine and o-phenylenediamine in o-phosphoric acid by using the modified procedure reported by Addison [8]. The benzimidazolyl pyridine functionalized anisylcalix[4]resorcinarene (L) was synthesized by the reaction of anisylcalix[4]resorcinarene (P1) with 2-(pyridin-2-yl)-1H-benzimidazole (P2) and formaldehyde in solvent mixture of ethanol and 2-methoxyethanol (Scheme 1). The ligand L is characterized by CHN analysis, IR, and mass spectrometry. In FT-IR spectrum of L, the band at 1466 cm⁻¹ is due to scissoring of-CH₂ connected to tertiary amine, confirming the formation of CH₂bridge in the ligands. The ESI-TOF mass spectrum of L showed a peak at m/z 1473 corresponding to the fragment [M-(2C₆H₄OCH₃+3H₂O)]⁺ formed by the removal of two methoxyphenyl groups and three water molecules from the molecular ion. The peak at m/z 1354 is due to the fragment [(M+H)-2P2]⁺ formed by the loss of two benzimidazolyl pyridine groups followed by the addition of one hydrogen atom to the molecular ion (Figure 3). The precursor complexes, [Ru(phen)Cl₂]·2H₂O (P3) and [Ru(bpy)Cl₂]·2H₂O (P4) were synthesized by the reaction of RuCl₃·3H₂O (1 equiv) and phenanthroline bipyridine (2 equiv) in refluxing DMF in the presence of lithium chloride as reported by Sullivan et al [9].

The reaction of ruthenium precursor complexes, P3 or P4, with the ligand L (1 equiv) in N,N-dimethylformamide, as per the Scheme 2, formed the stable complexes of I and 2, respectively. The tetranuclear complexes were characterized by CHN analysis, IR, ESI MS, and electronic absorption spectroscopy. The IR spectral bands at 1097 and 1088 cm⁻¹ confirm the presence of perchlorate counter ion in complexes 1 and 2, respectively. The ESI-TOF mass spectra of the complexes give the peaks corresponding to the fragment ions of the complexes (Figure 4 and 5).

3.2 Crystal structure of P1

The ORTEP representation of P1 with all atoms presented as 50% probability ellipsoid is given in Figure 6. The compound P1 crystallizes in monoclinic crystal system with P₂₁̅C space group. The cell parameters are a = 17.4950(5) Å, b = 11.6056(3) Å, c = 19.7364(6) Å, α = γ = 90°, β = 74.923(3)°, V = 3630.4(3) Å³ and Z = 2. The molecule possesses a cis-trans-trans (ctt) configuration and a rigid chair-like conformation. The asymmetric unit is comprised of half of the anisylresorcinarene molecule which can be completed using the two fold rotation axis (axis passing through the centre of the molecule). The solvent molecules are found to be highly disordered throughout the crystal lattice and so they are removed using SQUEEZE program. The resorcinol hydroxyl groups are found to involve in the intermolecular H-bonding with the -OCH₂ group of the nearby molecule that the C-H···O interactions between the two molecules also stabilize the crystal lattice (Figure 7).

Figure 6: ORTEP representation of P1 (CCDC # 1566460) with atoms represented as 40% probability thermal ellipsoids.

Figure 7: Two dimensional arrangement of P3 crystal with short contacts.

3.3 Hirshfeld surface analysis of P1

Intermolecular interactions play a vital role in the crystal packing of molecules in supramolecular chemistry. The quantitative measurements of these interactions are possible by Hirshfeld surface (HS) analysis of the crystal [10]. The HSs of P1 in the Figure 8 show surfaces that have been mapped with dnorm shape index and curvedness. The intense red spots in the dnorm surface near to resorcinol ring are due to H-bond interaction present in the molecule and the light red spots are due to C-H and CH-π interactions.

Figure 8: Hirshfeld surfaces of P1 (dnorm shape index and curvedness).
The shape index map on the HSs shows a complementary red hollows and blue bumps where two molecular HSs touch one another. Curvedness shows relatively large regions of green with r.m.s. curvature near unity, which is separated by dark blue ‘edges’ with large r.m.s. curvature. Occasional highlights of yellow and red indicate hydrogen bonds on the surface. The quantitative measures like volume (V_{H1}), area (S_0), globularity (G) and asphericity (Ω) were also computed using HS analysis. The term, globularity (G) is found to be <1 which indicates that the molecular surface is more structured, not a sphere. The asphericity, the measure of anisotropy [10] is found to be 0.11 for P1.

### 3.4 Fingerprint plots of P1

Two dimensional summaries of the intermolecular interactions in the crystal system are provided by a fingerprint plot of d_{ij} versus d_{lc}, Figure 9 displays the two dimensional fingerprint plots for the P1 molecule which involve features like spike of various length and thickness corresponding to all the intermolecular interaction present within the crystal structure. The wing like peripheral spikes represent C-H contacts in fingerprint plot whereas hydrogen bonding are present as longer and thinner spikes and the non-directional H···H contacts are characterized by broader spikes.

**Figure 9: Fingerprint regions of P1 (all interactions-100%, H-H -72.9%, O-H-19.9%, C-H 7.3%)**

### 3.5 Luminescence properties of complexes at 298 and 77 K.

The tetranuclear ruthenium(II) complexes exhibit electronic absorption bands at 263, 317 and 332 nm for 1, and 289, 317 and 333 nm for 2 assignable to the ligand centered π-π* transitions and broad absorption bands at 445 and 457 nm for 1 and 2, respectively, characteristic of the 1MLCT transition. The emission spectra of the complex were recorded in CH3CN at 298 K and 77 K. Upon excitation onto their excitation maxima, 1 and 2 complexes exhibit emission bands at 600 and 624 nm, respectively due to 1MLCT emission at 298 K. The complexes show blue shift in emission maxima in comparison with the parent compound [([bpy]3Ru(P4)])3+ [11].

**Figure 10: Electronic absorption spectra of the complexes 1 and 2 in CH3CN at 298K.**

The emission band of complexes 1 and 2 at 77 K in frozen CH3CN is hypsochromically shifted to 594 and 616 nm, respectively with noticeable increase in the emission intensity due to rigidochromic effect which is characteristic of typical MLCT emitters [12]. The quantum yields of 1 and 2 have increased from 0.003 to 0.04; and 0.013 to 0.23, respectively, when the temperature is decreased from 298 K to 77 K. The complexes show monoexponential decay profile in acetonitrile and have life time of 142 and 200 ns for 1 and 2, respectively. The absorption and emission spectra are depicted in Figures 10 and 11, respectively.

**Figure 11: Emission spectra of complexes 1 and 2 in CH3CN at 298K and 77K.**

### 4. Conclusion

A new benzimidazolyl functionalized anisylcalix[4]resorcinarene and its tetranuclearRu(II) polypyridyl complexes have been prepared. The study demonstrates the versatility of calixresorcinarenes in the assembly of polynuclear ruthenium(II) complexes. The methodology developed in the present work can be exploited to synthesize a series of polypyridine functionalized calixresorcinarenes. The tetranuclearRu(II) complex is stable in solution and solid state under ambient conditions and is soluble in organic solvents. The ligand offers a wide scope in constructing a plethora of complexes by inserting different metal cations to vary and fine-tune the electronic properties in such polynuclear assemblies.

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### References


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