Structural and photophysical properties of gold(I) phosphine-decorated 4,4'-diethynyl-2,2'-bipyridine ligands


Keywords: gold / heterocycles / alkyne / 2,2'-bipyridine / emission

Treatment of 4,4'-diethynyl-2,2'-bipyridine with R3PAuCl (R = Ph, 4-tolyl, Et, iPr) leads to the formation of a family of gold(I) phosphine decorated 4,4'-diethynyl-2,2'-bipyridine ligands. The solid state structures of the compounds are significantly affected by the change from aryl to alkyl substituted phosphines, while the progression from ethyl to isopropyl substituents leads to a subtle change in the packing that results in the propagation of two different polymeric chains motifs, both supported by close Au...Au contacts (3.1239(1) Å for R = Et, and 3.395(1) Å for R = iPr). In CH2Cl2 solution, each of compounds 1-4 is a dual emitter at room temperature. When the excitation wavelength is ≈238 nm, the emission spectra of 1 and 2 exhibit new bands at 288 and 570 nm at the expense of the original emissions. The photodegradation is not inconsistent with the formation of gold nanoclusters.

Introduction

The formation of Au–C σ-bonds by coupling terminal alkynes with Au(I) metal centres is well established.[1,2] Interest in gold(I)-containing compounds arises from their potential applications in advanced materials, largely because of their luminescent properties.[3–7] Among the first photoemissive gold(I) alkynyl complexes to be described in the literature were Ph3PAuC≡CPh and {PhC≡CAu}2(µ-dppe) (dppe = Ph2PCH2CH2PPh2). Che and coworkers[8] showed that these species possessed long-lived and emissive excited states in CH2Cl2 solution at ambient temperature, and upon going from solution to the solid state, a dramatic red shift in the emission maximum of {PhC≡CAu}2(µ-dppe) was observed. The presence of short (≈3.00-3.20 Å) Au...Au contacts in the solid state is thought to be fundamental to emissive behaviour.[3,9–11]

By combining phosphine and alkyne metal-binding domains at a gold(I) centre, one gains significant scope for the design of macromolecules:[12] rigid-rod P–Au–C– domains and a wide range of readily accessible mono-, bi- and polydentate phosphate ligands containing sp3 hybridized P atoms combine with a tendency for gold atoms to aggregate (so-called 'aurophilicity').[13–15,16] Of the 215 compounds in the CSD (CSD v 5.3 with May 2009 updates, Conquest v. 1.11) possessing R3PAuC≡C units, only 19[9,18–32] exhibit intermolecular aurophilic interactions resulting in one-dimensional Au...Au bonded polymers. Of particular relevance to our work is the trigold derivative ([Bu3PAuC≡CCH2OCH2CH2O)3CCH2Otpy (tpy = 2,2';6',2"-terpyridine) which forms one-dimensional chains with an oriented assembly in the solid state in which all the tpy units point in the same direction.[19] The incorporation into gold(I) alkyne-containing assemblies of additional metal-binding domains such as tpy,[19,30,33] bpy[30,32,34–37] (bpy = 2,2'-bipyridine) or 1,10-phenanthroline[38] allows the incorporation of redox- or photoactive metal centres for tuning (or switching if metal-binding is reversible) the physical properties of the material. For enhancing emission properties, molecular design considerations need to take into account the steric demands of the gold(I) substituents so that aurophilic interactions may be optimized. It should be noted, however, that even though close gold–gold contacts may appear to be accessible in the solid state, competition with hydrogen-bonding and other weak interactions may dominate in the packing.[33] In contrast, aurophilic and hydrogen-bonded interactions[39] or metal-binding[40] may act in a cooperative manner.

At a separation of ≈3 Å, the strength of an attractive Au...Au interaction is about 30 kJ mol–1,[41] and this is comparable with the dissociation enthalpy of a classical hydrogen bond.[15]

We report here a series of 2,2'-bipyridine ligands (Chart 1), decorated in the 4 and 4'-positions with phosphine gold(I) alkynyl units, and discuss the relationship between their solid state structures and their solid state and solution emissive properties.

![Chart 1. Structures of compounds 1-4 with the numbering scheme used for NMR spectroscopic assignments; for 1 and 2, phenyl ring is labelled B.](image-url)
Results and Discussion

Synthesis and solution characterization

Compounds 1-4 (Chart 1) were prepared by the reaction of 4,4'-diethynyl-2,2'-bipyridine\(^{42}\) with the appropriate R\(_3\)PAuCl in a mixture of toluene, CH\(_2\)Cl\(_2\) and disopropylamine in the presence of Cul.\(^{43}\) The reactions were monitored by spot thin layer chromatography and under ambient conditions, the reactions required 12-16 hours to reach completion. However, upon heating in THF or CH\(_2\)Cl\(_2\) at 50°C in a microwave reactor, each reaction was complete within 30 minutes.

The base peak in the electrospray mass spectrum of 1 appeared at m/z 1121.8 and corresponded to the [M + H]\(^+\) ion. Very low intensity peaks at m/z 721.7 (5%) and 1579.4 (5%) were assigned to [Au(Ph\(_3\)P)\(_2\)]\(^+\) and [M + AuPPh\(_3\)]\(^+\)\(^{44}\). In contrast, the base peak in the ESI mass spectrum of 2 corresponded to [Au{(4-Tol)\(_3\)P}2\(^+\) (m/z = 805.2), and the next most intense peak to [M + AuP(4-Tol)\(_3\)]\(^+\) (m/z = 1705.0, 59%). The [M + H]\(^+\) ion was observed only as a low intensity peak (m/z = 1205.1, 28%). This same pattern of peaks was also observed for 3 and 4, but for these compounds, the base peak corresponded to [M + AuP(4-Tol)\(_3\)]\(^+\) (m/z = 1147.1) or [M + AuP(4-Tol)\(_3\)]\(^+\) (m/z = 1273.2), and the [M + H]\(^+\) peak was observed with relative abundances of 10 and 3%, respectively.

The \(^1\)H and \(^13\)C NMR spectra of the compounds were in accord with the symmetrical structure shown in Chart 1. However, it was not possible to resolve signals in any of the \(^13\)C NMR spectra for the alkyne carbon atoms. The \(^31\)P NMR absorption and emission data are discussed later in the paper.

Structural determinations

Crystals of 2\{1\} Et\(_2\)O and 2 Et\(_2\)O suitable for single X-ray diffraction were grown by slow diffusion of Et\(_2\)O into a CH\(_2\)Cl\(_2\) solution of 1, or a toluene/CH\(_2\)Cl\(_2\) solution of 2. The structures of the compounds are shown in Figures 1 and 2, respectively. Whereas 2 is centrosymmetric, I contains two independent gold atoms. The bpy unit in both derivatives adopts the expected transoid conformation, with the angle between the least squares planes of the two pyridine rings being 8.8(3)\(^\circ\) in 1 and constrained by symmetry to 0\(^\circ\) in 2. The environment around the P atoms is unexceptional, with P–C bond distances in the range 1.790(5)–1.825(6), and 1.812(1)–1.816(1) Å, respectively. There is only a small deviation from linearity along the P–Au–C\(_{bpy}\) units, with the smallest angle subtended at any of the gold and alkyne carbon atoms being 174.6(2)\(^\circ\) at Au2 in 1, and 176.83(5)\(^\circ\) at Au1 in 2. In both 2\{1\} Et\(_2\)O and 2 Et\(_2\)O, the diethyl ether molecules are disordered. The Et\(_2\)O molecule was modelled over two positions in each compound; in 2 Et\(_2\)O, the O atom was common to both positions. The centrosymmetric pair of molecules of 1 in the unit cell engage in edge-to-face interactions between phenyl rings. Figure 3 highlights these interactions and also shows how the alkyne carbon atoms C32 and C33 form close contacts to a CH unit of a phenyl ring on an adjacent molecule. The C31C32 alkyne unit also interacts with a phenyl ring of another molecule of 1 (C32...H361\(^{iii}\) = 2.88, C32...C36\(^{iii}\) = 3.435(9) Å, symmetry code iii = 1 + x, y, z). In 2 Et\(_2\)O, similar close contacts are observed between molecules of 2, but this time involving the methyl group of one tolyl substituent (C7...H212\(^{ii}\) = 2.85 Å, C7...C21\(^{ii}\) = 3.764(3) Å, symmetry code ii = –1 + x, y, z). Packing motifs based on C–H...C\(_{alkyne}\) contacts are well established in the solid state structures of alkenes with aromatic substituents,\(^{44-49}\) and C–H...C\(_{alkyne}\) interactions are ubiquitous among compounds containing R\(_3\)PAuC=C units (CSD v 5.3 with May 2009 updates, Conquest v. 1.11),\(^{17}\) involving both aromatic and aliphatic C–H units. There are no significant intermolecular Au...Au contacts in either 1 or 2, the closest separations being Au2...Au2\(^{i}\) = 4.763(1) Å in 1 (symmetry code i = 2 – x, y, 1 – z), and Au1...Au1\(^{iii}\) = 5.303(1) Å in 2 (symmetry code iii = 1 – x, 1 – y, –z). In the light of the structure of compound 4 (see below), the lack of Au...Au interactions cannot simply be attributed to the steric demands of the triarylphosphine units (Tolman cone angles for PPh\(_3\) and P(4-Tol)=145°)\(^{30}\).

Fig. 1 Molecular structure of compound 1 in 2\{1\} Et\(_2\)O (ellipsoids plotted at the 50% probability level). Selected bond parameters: Au1–P1 = 2.263(1), Au1–C32 = 2.008(6), Au2–P2 = 2.071(1), Au2–C12 = 2.015(7), C11–C12 = 1.160(9), C31–C32 = 1.184(8) Å; P1–Au1–C32 = 174.6(2), P2–Au2–C12 = 177.7(2), C3–C11–C12 = 179.4(7), Au2–C12–C11 = 179.6(6), C8–C31–C32 = 176.4(6), C31–C32–Au1 = 175.7(5)°.

Fig. 2 Molecular structure of compound 2 (ellipsoids plotted at the 50% probability level) in 2 Et\(_2\)O. Selected bond distances and angles: Au1–P1 = 2.2802(3), Au1–C7 = 1.997(1), C6–C7 = 1.213(2) Å; P1–Au1–C7 = 176.83(5), C5–C6–C7 = 177.1(2), C6–C7–Au1 = 179.8(2)°. Symmetry code i = –x, –y, 1–z.
Single crystals of 3 and 4 were grown by slow diffusion of Et₂O into a CH₂Cl₂ solution of each compound. Figures 3 and 4 depict the molecular structures and give selected bond parameters. In the triethylphosphine derivative 3, the gold atoms are in independent (but chemically similar) environments, while the triisopropylphosphine derivative 4 is centrosymmetric. In 4, one of the isopropyl groups is disordered and has been modelled over two sites of fractional occupancies 0.64 and 0.36. The backbone of each of 3 and 4 is significantly more bowed than those in 1 and 2, with the smallest angles subtended at gold or alkyne carbon atoms being 166.7(2)° at C14 in 3, and 172.8(3)° at Au1 in 4. The bipy unit is constrained by symmetry to planarity in 4, and in 3, the angle between the least squares planes of the two pyridine rings is 11.3(1)°. Figure 6 illustrates that there are clear similarities in the overall packing of 3 and 4, with weak interactions between alky1 C–H groups and the bpy domains of adjacent molecules, as well as C–H...alkyne contacts (Table 1).

Table 1. C–H...N non-classical hydrogen bonds and C–H...Calkyne interactions in 3 and 4.

<table>
<thead>
<tr>
<th></th>
<th>D–H...A</th>
<th>H...A / Å</th>
<th>D...A / Å</th>
<th>D–H–A°</th>
<th>Symmetry codes a–d</th>
</tr>
</thead>
<tbody>
<tr>
<td>3</td>
<td>C19–H191...N2¹</td>
<td>2.66</td>
<td>3.617(4)</td>
<td>171</td>
<td>1 – x, 1/2 + y, 1/2 – z</td>
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<tr>
<td></td>
<td>C25–H252...N1²</td>
<td>2.79</td>
<td>3.717(4)</td>
<td>159</td>
<td>x, y, –1 + z</td>
</tr>
<tr>
<td></td>
<td>C23–H231...C14⁴</td>
<td>2.88</td>
<td>3.659(4)</td>
<td>144</td>
<td>1/2 + x, 1/2 + y, 1/z</td>
</tr>
<tr>
<td>4</td>
<td>C11–H11A...N1³</td>
<td>2.88</td>
<td>3.84(1)</td>
<td>161</td>
<td>1/2 + x, 1/2 + y, 1 – z</td>
</tr>
<tr>
<td></td>
<td>C10–H10B...C6'[a]</td>
<td>2.95</td>
<td>3.90(3)</td>
<td>163</td>
<td>x – y, 1/2 – z</td>
</tr>
<tr>
<td></td>
<td>C10–H10B...C7'[a]</td>
<td>2.94</td>
<td>3.78(3)</td>
<td>143</td>
<td>1/2 – x, 1/2 – y, 1/2 + z</td>
</tr>
</tbody>
</table>

[a] H10B is major occupancy site for disordered H atom; the site for minor occupancy H10D also permits a comparable C–H...C interaction.

In contrast to the situation in 2{1}Et₂O and 2Et₂O, the gold atoms in 3 and 4 approach closely. In 3, the Au1...Au2 separation is 3.1239(1) Å (Figure 6), and in 4, Au1...Au2 is 3.395(1) Å (symmetry code ii = 1 – x, y, 1/2 – z). These close contacts lead to the assembly of polymeric chains. However, careful inspection of Figure 6 (in which the Au...Au contacts in each chain are shown as black lines) shows that the directions in which the infinite chains are propagated are significantly different. This is quantified in the noticeable difference between the Calkyne–Au...Au–Calkyne dihedral angles in 3 and 4. In 4, C7–Au1...Au2³–C7″ is 101.5° (symmetry code ii = 1 – x, y, 1/2 – z), whereas in 3, the dihedral angle C12–Au1...Au2²–C14¹ is –127.8° (symmetry code i = 1 + x, 3/2 – y, 1/2 + z). We propose that the primary intermolecular interactions controlling the overall assembly are weak hydrogen bonds and C–H...π and van der Waals interactions. In each solid state structure, each gold atom has two neighbouring gold atoms, one at a separation of <3.4 Å and a second at just over twice this distance (7.389 Å in 3 and 8.654 Å in 4). The difference in the Tolman cone angle of the phosphine (PET₃, 132° and PPr₃, 160°),[50] appears to be sufficient to tune the packing so that the dominant aurophilic interactions are between spatially distinct pairs of gold atoms resulting in the different chain assemblies shown in Figure 7.
Absorption and emission properties

The dominant band in the absorption spectrum of a CH$_2$Cl$_2$ solution of 4,4'-diethynyl-2,2'-bipyridine is at 233 nm, with lower intensity, overlapping bands at 298 and 310 nm, the latter being red-shifted from the absorption at 280 nm in 2,2'-bipyridine (Figure 8). These absorptions arise from alkyne and bpy $\pi^* \leftarrow \pi$ and $\pi^* \leftarrow n$ transitions. Figure 8 also shows the absorption spectra of the free phosphines Ph$_3$P and (4-Tol)$_3$P, both of which shows absorption maxima at 264 nm. On going from 4,4'-diethynyl-2,2'-bipyridine to compounds 1-4, the highest energy band in each spectrum lies between 235 and 239 nm and is presumably alkyne-centred (Figures 8 and 9). The more intense absorptions around 260 nm for compounds 1 and 2 compared to 3 and 4 are attributed to the presence of the aryl substituents in the former. Above ≈260 nm, there is a common pattern of absorptions across the series of derivatives (Figure 9), and by comparison with related systems, we propose that these bands arise from bpy/alkyne $\pi^* \leftarrow \pi$ transitions with involvement from Au orbitals.

In CH$_2$Cl$_2$ solution, each of compounds 1-4 is a dual emitter at room temperature (Figure 10). The general shape of the emission spectra is similar to that reported by Vicente et al. for the triphenylphosphine gold(I) derivative of 5-ethyl-2,2'-bipyridine. Excitation of compound 1 (in CH$_2$Cl$_2$) at 239 nm leads to emission bands at 344 and 438 nm, and a well-defined shoulder appears at 466 nm when the compound is excited at 268 or 285 nm, i.e. in the region of the bpy/alkyne-centred absorptions. Similar behaviour is observed for 2-4. For each of derivatives 1-4, the lower energy emission is the more intense of the two, the reverse of that observed for 5-
shows that the energy and intensity of the emissions are virtually independent of the phosphine substituent. The excitation spectrum of a CH₂Cl₂ solution of 1 showed that each of the emissions at 344, 438 and 466 nm originates from the broad absorptions between 250 and 320 nm.

Fig. 10 Emission spectra of compounds 1-4: the key for the curves is the same as in Figure 9. Concentration of each CH₂Cl₂ solution is 1.2 x 10⁻³ mol dm⁻³ (* = first harmonic).

When sequential solution (CH₂Cl₂) emission spectra of compounds 1 and 2 were recorded over a period of about 30 minutes (~20-25 spectra), we observed a decay of the bands shown in Figure 10 and the growth of emission bands at 288 and 570 nm (Figure 11) when the excitation wavelength was 238 nm. No such changes were observed when the compounds were excited at higher wavelengths. Absorption spectra recorded immediately after each series of emission spectra revealed new absorptions at 323 and 366 nm when λ_ex had been 238 nm, and no change in the absorption spectra when λ_ex = 253 (for compound 2), 269 or 286 nm (for compounds 1 and 2). We have been unable to determine the nature of the photodegradation product, but tentatively propose that it arises from cleavage of the alkyne–gold bond since the absorption band at ≈238 nm is most likely to be alkyne-centred. The trialkylphosphine derivatives 3 and 4 behave similarly, although the emissions from the degradation products are very much less intense than from compounds 1 and 2. The observation of the emission at 570 nm upon photodegradation is not inconsistent with the formation of small gold nanoparticles.

Fig. 11 Emission spectrum of a CH₂Cl₂ solution of compound 2 as a function of time (λ_ex = 238 nm; * = first harmonic.).

Conclusions

We have shown that bis(gold(I) phosphine)-decorated 4,4'-diethynyl-2,2'-bipyridine ligands can be readily made by reaction of 4,4'-diethynyl-2,2'-bipyridine with R₃PAuCl (R = Ph, 4-tolyl, Et, iPr). The solid state structures of the four compounds (1-4) show a dependence upon the presence of aryl or alkyl substituted phosphine, and the change from ethyl to isopropyl substituents leads to a subtle alteration in the packing that results in the propagation of two different polymeric chains motifs, both supported by close Au...Au contacts (3.1239(1) Å for R = Et, and 3.395(1) Å for R = iPr). In CH₂Cl₂ solution, each of compounds 1-4 is a dual emitter at room temperature. When the excitation wavelength is ≈238 nm, the emission spectra of the compounds decay over a period of about 30 minutes, and for each of 1 and 2, new bands at 288 and 570 nm become the dominant features of the emission spectrum. The photodegradation is not inconsistent with the formation of gold nanoparticles.

Experimental Section

General: ¹H and ¹³C NMR spectra were recorded at 295 K on Bruker Avance DRX-500 or DPX-400 MHz spectrometers; for ¹H and ¹³C, chemical shifts are relative to residual solvent peaks (TMS δ 0 ppm) and for ¹³C P are with respect to external reference 85% aqueous H₃PO₄. A Shimadzu FTIR-8400S spectrophotometer was used to record IR spectra (solid samples on a Golden Gate diamond ATR accessory). Electrospray ionization (ESI) mass spectra were recorded using Finnigan MAT LCQ or Bruker esquire 3000 Plus instruments. Electronic absorption and emission spectra were recorded on a Varian-Cary 5000 spectrophotometer and Shimadzu RF-5301 PC spectrophotometer, respectively. Microwave reactions were performed in a Biotage Initiator 8 reactor. Solvents were distilled before use (water content monitored by Karl-Fischer titration), and all reactions were carried out under N₂.

R₃PAuCl (R = Et, ‘Pr, Ph, 4-Tol) (4-Tol = 4-tolyl), were prepared from HAuCl₄·3H₂O (Sigma-Aldrich) by a published procedure. For Ph₃PAuCl and (4-Tol)₃PAuCl, the reactions were carried out at -5°C; remaining syntheses were carried out at -20°C. The R₃PAuCl products were purified by washing with hot hexane.

Compounds 1-4 were prepared by either method 1 and 2, with yields being similar for either route for a given phosphine.

Method 1. R₃PAuCl (R = Et, ‘Pr, Ph, 4-Tol), 4,4'-diethynyl-2,2'-bipyridine and CuI were dissolved in a mixture of CH₂Cl₂ (or THF) (6 mL) and toluene (2 mL). Diisopropylamine (predistilled, 2 mL) was added to make the solution basic. The reaction mixture was stirred at room temperature in the dark (12-16 h), after which it was filtered and the solvent removed from the filtrate in vacuo. The crude material was purified by preparative plate chromatography in the dark (Al₂O₃, CH₂Cl₂).

Method 2. R₃PAuCl (R = Et, ‘Pr, Ph, 4-Tol), 4,4'-diethynyl-2,2'-bipyridine and CuI were added to argon-degassed THF (or CH₂Cl₂) (8 mL). Diisopropylamine (predistilled, 2 mL) was added to make the solution basic. The reaction mixture was placed in a vial in a microwave reactor (50°C, 30 min), after which it was filtered and the solvent removed from the filtrate. The crude material was purified by preparative plate chromatography in the dark (Al₂O₃, CH₂Cl₂).

Compound 1. R₃PAuCl (126 mg, 255 μmol, 4,4'-diethynyl-2,2'-bipyridine and CuI were dissolved in a mixture of CH₂Cl₂ (or THF) (6 mL) and toluene (2 mL). Diisopropylamine (predistilled, 2 mL) was added to make the solution basic. The reaction mixture was stirred at room temperature in the dark (12-16 h), after which it was filtered and the solvent removed from the filtrate. The crude material was purified by preparative plate chromatography in the dark (Al₂O₃, CH₂Cl₂).

Compound 2. (4-Tol)₃PAuCl (105 mg, 196 μmol, 4,4'-diethynyl-2,2'-bipyridine and CuI (68.1 μmol) and CuI (0.2 mg, 1 μmol) were added to argon-degassed THF (or CH₂Cl₂) (8 mL). Diisopropylamine (predistilled, 2 mL) was added to make the solution basic. The reaction mixture was placed in a vial in a microwave reactor (50°C, 30 min), after which it was filtered and the solvent removed from the filtrate. The crude material was purified by preparative plate chromatography in the dark (Al₂O₃, CH₂Cl₂).
NMR (126 MHz, CDCl3): δ ppm = 156.1 (C3'), 149.1 (C4'), 142.2 (C5'), 134.4 (d, Jc = 14 Hz, C6'), 134.2 (C7'), 131.0 (d, Jc = 12 Hz, C8'), 126.9 (d, Jc = 59 Hz, C9'), 126.5 (C10'), 124.4 (C11'), 21.7 (C12') (C not observed); 13C NMR (162 MHz, CDCl3): δ ppm = 40.6 (s); UV-Vis (CH2Cl2): λmax (α) = 239 (105000), 253 sh (88000), 269 (69000), 286 (71000), 303 (28000), 315 nm (20000 dm3 mol–1 cm–1); emission (CH2Cl2, T = 298 K): λmax = 285 nm; Φα = 338, 486 nm; ESI-MS (CH2Cl2): m/z = 805.2 [Au4(i-Pr3P)+] (calc. 805.3). 4H elemental analysis calecd (%) for C31H36Au2N2P2 = 55.56, H 4.66, N 2.16. 4H elemental analysis calecd (%) for C31H36Au2N2P2: Et2O/H2O C 55.56, H 4.66, N 2.16, found C 55.67, H 4.54, N 2.12.

Compound 3. Et3PAuCl (68.7 mg, 196 µmol), 4,4′-diethyl-2,2′-bipyridine (20 mg, 98 µmol) and Cal (0.2 mg, 1 µmol). 3 was isolated as white solid (25.6 mg, 30.8 µmol, 31.4%). 1H NMR (500 MHz, CDCl3): δ ppm = 8.50 (dd, J = 5.0, 0.8 Hz, 2H, H3'), 8.33 (dd, J = 14.0, 2.8 Hz; 2H, H3'), 7.27 (dd, J = 5.0, 1.6 Hz, 2H, H3'), 1.81 (dq, J9a = 9.6, J9b = 7.7 Hz, 2H, H3'). (12) 13C NMR (126 MHz, CDCl3): δ ppm = 151.6 (C1'), 149.1 (C2'), 134.2 (C3'), 126.9 (C4'), 126.6 (C5'), 9.1 (C6') (C not observed); CP-305000, 303 (23000), 316 nm (21000 dm3 mol–1 cm–1); emission (CH2Cl2, T = 298 K): λmax = 285 nm; Φα = 338, 486 nm; ESI-MS (CH2Cl2): m/z = 805.2 [Au4(i-Pr3P)+] (calc. 805.3). 4H elemental analysis calecd (%) for C31H36Au2N2P2: Et2O/H2O C 55.56, H 4.66, N 2.16, found C 55.67, H 4.54, N 2.12.

References


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Supporting Information

CCDC 739607 (21) Et2O, 739604 (2 Et2O), 739605 (3) and 739606 (4) contain the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.


Entry for the Table of Contents

Layout 1:

Bis(gold(I) phosphine)-decorated 4,4'-diethynyl-2,2'-bipyridine ligands can be made by reaction of 4,4'-diethynyl-2,2'-bipyridine with R₃PAuCl (R = Ph, 4-tolyl, Et, iPr). The solid state structures of these derivatives show a dependence upon the presence of aryl or alkyl substituted phosphine; the change from ethyl to isopropyl substituents leads to packing differences that result in the propagation of two different polymeric chains motifs, both supported by close Au...Au contacts (3.1239(1) Å for R = Et, and 3.395(1) Å for R = iPr). In CH₂Cl₂ solution at room temperature, each compound is a dual emitter. When the excitation wavelength is ≈238 nm, the emission spectra decay over a period of ≈30 minutes, and for the phenyl and 4-tolyl derivatives, new bands at 288 and 570 nm become dominant emissions.

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