Phosphorescent Tris-cyclometalated Pt(IV) Complexes with Mesoionic N-Heterocyclic Carbene and 2-Arylpyridine Ligands

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ABSTRACT: The synthesis, structure, photophysical properties, and electrochemistry of the first series of Pt(IV) tris-chelates bearing cyclometalated aryl-NHC ligands are reported. The complexes have the general formula [Pt(trz)(CAN)]+, combining two units of the cyclometalated, mesoionic aryl-NHC ligand 4-butyl-3-methyl-1-phenyl-1H-1,2,3-triazol-5-ylidine (trz) with a cyclometalated 2-arylpyridine [CAN = 2-(2,4-difluorophenyl)-pyridine (dfppy), 2-phenylpyridine (ppy), 2-(p-tolyl)pyridine (tpy), 2-(2-thienyl)pyridine (thpy), 2-(9,9-dimethylfluoren-2-yl)-pyridine (dfpy)], and presenting a mer arrangement or metalated aryls. They exhibit a significant photostability under UV irradiation and long-lived phosphorescence in the blue to yellow color range, arising from 3LC excited states involving the CAN ligands, with quantum yields of up to 0.34 in fluid solution and 0.77 in the rigid matrix at 298 K. The time-dependent density functional theory (TD-DFT) calculations reveal that nonemissive, deactivating excited states of ligand-to-metal charge-transfer (LMCT) character are pushed to high energies as a consequence of the strong heteroaromaticity of the carbene moieties, making the Pt(trz)3 subunit an essential structural component that enables efficient emissions from the chromophoric CAN ligands, with potential application for the development of different Pt(IV) emitters with tunable properties.

INTRODUCTION

The use of N-heterocyclic carbene ligands (NHCs) for the design of strongly luminescent transition-metal complexes has become widespread, mostly associated with the development of phosphors for organic light-emitting devices (OLEDs).1–8 Mesoionic NHCs have received special attention in this field because their exceptionally strong σ-donating ability makes them particularly well suited to induce large ligand-field splittings, raising the energies of dissociative, metal-centered (MC) excited states and reducing the nonradiative deactivation that takes place through the thermal population of such states.9–11 This effect has even been applied with remarkable success to extend the excited-state lifetimes of strongly deactivated first-row transition-metal complexes.9–11

Bidentate cyclometalated aryl-substituted NHC ligands (aryl-NHCs, CAN*) have been extraordinarily successful with the Ir(III)12–17 and Pt(II)18–26 ions as a replacement of cyclometalated 2-arylpyridines (CAN), enabling better photostabilities, wider color tunability, and higher emission efficiencies. These enhancements are brought about by the larger ligand-field splitting induced by the NHC moiety with respect to the pyridine and, consequently, the reduced thermal accessibility of MC states from the triplet, mixed ligand-centered/metal-to-ligand charge-transfer (3LC/MLCT) emissive state of Ir(III) and Pt(II) complexes. The most relevant tris-chelates bearing CAN* ligands are homoleptic Ir(III) complexes mer/fac-[Ir(CAN*)]12,16,17,21 and mixed-carbene variations,19 which can achieve blue phosphorescent emissions, thanks to the large π−π* gap of the ligands. Heteroleptic tris-chelates of the type [Ir(CAN*)(CAN)]2 have also been reported, in which the arylcarbene acts as a supporting, nonchromophoric ligand, whereas the emission is mostly determined by the CAN ligands,13,18,19,23–31 except for a few cases that incorporate CAN* ligands featuring low π−π* gaps.35–37 However, very few heteroleptic tris-chelates bearing two CAN* ligands are known, which include complexes [Ir(C*CAcC*AC)(CAN)]2 bearing a bis-aryl-NHC38 and [Ir(C*AcC*)(2NAN)] or [Ir(CAN*)2(NAN)]9, where NAN is a pyridylpyrazolate, pyridyldiazolate, pyridylbenzimidazolate,39–41 or bipyridyld.42 Such systems are interesting because the Ir(CAN*)2 subunit functions as a robust platform for the development of efficient emitters whose properties can be tuned by incorporating different chromophoric CAN or NAN ligands.

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When compared with other d⁶ metal ions, the photophysical properties of Pt(IV) complexes have received much less attention. In previous studies, we have shown that cyclometalated Pt(IV) complexes with 2-arylpyridine ligands may exhibit very efficient and long-lived phosphorescence, 43−48 which makes them potentially useful as luminescence-based sensors, photosensitizers, or photocatalysts. Their emissive excited states are essentially ¹⁵C in character, with very little metal-to-ligand charge-transfer (MLCT) excited states, may become thermally accessible from the emissive state. Such states have dissociative character because dσ* orbitals are strongly antibonding, providing a pathway for nonradiative deactivation or photochemical reactivity. 49 Therefore, an indispensable requirement for Pt(IV) complexes to reach high emission efficiencies is the presence of suitable strong σ-donor ligands, which induce a large ligand-field splitting and raise the energy of MLCT states.

Recently, we reported Pt(IV) complexes of the types [PtCl₂(CAN*)₂(CAN)] and [PtCl₂(CAN*)₂(CAN+C)], where CAN* is a cyclometalated, mesoionic aryl-NHC ligand of the 1,2,3-triazolylidene subclass, which constituted the first examples of Pt(IV) emitters bearing a carbene ligand. 51,52 Derivatives of the type [PtCl₂(CAN*)₂(CAN)] exhibited strong ³LC emissions involving the CAN ligand, with significantly higher quantum efficiencies with respect to the homologous C₂-symmetrical [PtCl₂(CAN+C)] complexes as a consequence of the electronic effects of the carbene moiety. However, their synthesis presented problems associated with the difficult cyclometalation of the aryl-NHC ligand, resulting in relatively low yields. Herein, we present a straightforward methodology involving two consecutive cyclometalations of aryl-NHC ligands that has allowed the synthesis of complexes of the type [Pt(CAN*)₂(CAN)]. These species are the first Pt(IV) tris-chelates bearing cyclometalated aryl-NHC ligands and show intense phosphorescent emissions that can be modulated through the variation of the CAN ligand, demonstrating the usefulness of the Pt(CAN*)₂ subunit as a platform for the design of efficient emitters.

### RESULTS AND DISCUSSION

**Synthesis and Characterization.** The synthetic route to the target tris-cyclometalated Pt(IV) complexes is shown in Scheme 1. The reaction of the dimeric platinum precursor ((Pr₃N)_2[Pt₂Cl₆]) with 4 molar equiv of the in situ-prepared silver carbene intermediate “Ag(trzH)” (trzH = 4-butyl-3-methyl-1-phenyl-1H-1,2,3-triazol-5-ylidene) in 1,2-dichloroethane at 80 °C led to the selective formation of trans-[PtCl₂(trzH)] (I), which could be isolated in 85% yield. The trans geometry of I was established from an X-ray diffraction analysis (see below). The reason for the exclusive formation of this isomer is probably that, upon coordination of the first trzH ligand, the Pt−C₁ bond trans to the carbene carbon becomes highly labile and is rapidly abstracted by the silver ion, resulting in the coordination of a second trzH ligand in this position. Consistent with this, the attempts to obtain a complex with only one trzH ligand by employing a 1:2 molar ratio (dimeric platinum precursor to carbene) were unsuccessful, resulting always in the formation of I. Similar results have been previously observed for the reactions of K₂[PtCl₄] with other silver carbene. 53−57

The ¹H and ¹³C NMR spectra of complex I show two sets of resonances for the trzH ligand in very similar proportions (Figure S1), pointing to the presence of two different conformational isomers that interconvert very slowly at room temperature as a consequence of restricted rotation about the Pt−C bond. This possibility was confirmed by a variable-temperature NMR study in DMSO-d₆, which showed that the different pairs of aromatic and aliphatic signals coalesce in the range 35−60 °C (Figure S2). Using the Eyring equation, 58 a free energy of activation of ΔG° = 15.0 kcal/mol was calculated for this process at the coalescence temperature of the NCH₂ protons (60 °C). Several examples of restricted rotation of NHC ligands about the metal−C bond have been previously reported. 53,59−61

The crystal structure of I is shown in Figure 1. The crystal analyzed corresponded to the conformer with an antiparallel orientation of the phenyl and butyl substituents of the trzH ligands. The Pt−C₁ bonds lie along a crystallographic 2-fold axis and therefore the coordination around the metal is strictly planar. The mean plane of the triazolylidene ring is rotated by 68.85° with respect to the coordination plane. The Pt−C₁ bond distance of 2.027(2) Å is typical of Pt(II) complexes with mutually trans NHC ligands. 53−55,62−64

Treatment of a CH₂Cl₂ solution of I with PhICl₂ led to the oxidation to Pt(IV) and the electrophilic metatation of the pendant aryl group of one of the trzH ligands, resulting in the formation of the monocyclometalated species [PtCl₃(trz)₂]⁻.

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**Scheme 1. Synthetic Route to the Target Tris-cyclometalated Pt(IV) Complexes**

![Scheme 1](https://doi.org/10.1021/acs.inorgchem.2c02039)
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Figure 1. Structure of complex 1 (thermal ellipsoids at 50% probability). Hydrogen atoms are omitted. Selected bond distances (Å) and angles (°): Pt–C1, 2.027(2); Pt–C11, 2.3253(8); Pt–C12, 2.3132(7); C1–Pt–C11, 90.35(6); C1–Pt–C12, 89.65(6); and Cl1–Pt–C12, 180.0.

Figure 2. Structures of complexes 2 (a) and the cations of 4b (b) and 4d (c) (thermal ellipsoids at 50% probability). Hydrogen atoms are omitted. Selected bond distances (Å) and angles (°): 2: Pt–C1, 2.046(2); Pt–C9, 2.032(2); Pt–C14, 2.089(2); Pt–C11, 2.4509(5); Pt–C12, 2.3190(5); Pt–C13, 2.3395(5); C1–Pt–C9, 80.95(8); and C1–Pt–C14, 173.90(8). 4b: Pt–C1, 2.037(3); Pt–C9, 2.071(3); Pt–C14, 2.043(4); Pt–C22, 2.065(4); Pt–N7, 2.094(3); Pt–C31, 2.087(3); C1–Pt–C9, 80.22(14); C14–Pt–C22, 79.79(14); and N7–Pt–C31, 79.69(13). 4d: Pt–C1, 2.034(2); Pt–C9, 2.0414(19); Pt–C14, 2.0393(19); Pt–C22, 2.0837(19); Pt–N40, 2.1164(17); Pt–C31, 2.0790(19); C1–Pt–C9, 80.06(8); C14–Pt–C22, 80.05(8); and N–Pt–C31, 79.46(7).

(trzH)] (2), which was isolated in 96% yield. The 1H NMR spectrum of 2 showed an aromatic resonance flanked by 195Pt satellites at δ = 6.99 ppm (JHH = 37 Hz), arising from the proton ortho to the metalated carbon of a phenyl ring. The crystal structure (Figure 2) corroborated the presence of a cyclometalated trz and a coordinated trzH and revealed that the carbene moieties remain mutually trans, resulting in a mer disposition of metalated carbons or chloride ligands.

The metalation of the pendant phenyl group of the remaining trzH ligand in 2 was achieved by heating at 120 °C in 1,2-dichlorobenzene in the presence of a base (Na2CO3), which resulted in the formation of the bis-cyclometalated complex [PtCl2(trz)2] (3, 78% yield). Its 1H NMR spectrum shows a single set of resonances arising from equivalent cyclometalated trz ligands. In addition, the proton ortho to the metalated carbon of the phenyl ring is strongly shielded (δ = 6.49 ppm, JHH = 52 Hz), indicating that it is affected by the diamagnetic current of an orthogonal aromatic ring. These data demonstrate a C2-symmetrical configuration with mutually cis chloride ligands.

The introduction of different cyclometalated 2-arylpyridines (C\(\text{A}\text{N}\)) as the third chelating ligand was accomplished by reacting 3 with 2.4 equiv of AgOTf (OTf\(^{-}\) = trifluoromethanesulfonate) and an excess of the 2-arylpyridine in 1,2-dichlorobenzene at 120 °C, which afforded complexes \([\text{Pt(trz)}_2(\text{C}\text{A}\text{N})]\)OTf in 72–80% yields \([\text{C}\text{A}\text{N}] = \text{dppy (4a)}\), \text{ppy (4b)}\), \text{tpy (4c)}\), \text{thpy (4d)}\), \text{flpy (4e)}]. The 1H NMR spectra of these complexes show three distinctive aromatic resonances flanked by 195Pt satellites arising from the protons ortho to the metalated aryls, which are significantly shielded because they are directed toward orthogonal aromatic rings (δ = 7.04–6.29 ppm).

The crystal structures of complexes 4b and 4d were solved by X-ray diffraction analyses (Figure 2) and are completely analogous. The carbene moieties are mutually trans, and the coordinated nitrogen and metalated carbon of the C\(\text{A}\text{N}\) ligand are trans to each of the metalated aryls of the trz ligands. Therefore, they retain the disposition of trz ligands found in their precursor, resulting in a mer configuration of metalated aryl rings. In the case of 4d, the positions of the thiophene and pyridine rings were found disordered, with one of the orientations presenting a much higher occupancy factor than the other (ca. 81:19). Hence, the Pt–C9 or Pt–C22 bonds in 4d can be considered as predominantly trans to the pyridyl or thiyenyl rings, respectively (Figure 2), the latter being significantly longer because of the stronger trans influence of the metalated carbon. However, in 4b the positions of the pyridyl and phenyl rings of the ppy ligand were not distinguished by the refinement model, resulting in very similar Pt–C9 and Pt–C22 bond distances.

Photophysical Properties. The electronic absorption spectra of 4a–e in CHCl\(_2\) solution (Table 1, Figure 3) show structured absorptions in the 250–400 nm range that can be ascribed to primarily \(1\text{LC}\) transitions within the ligands. The shapes and energies of the observed bands are very similar to those of complexes \([\text{PtMe(\text{C}(\text{C}\text{A}\text{N}))}]^{4+}\) and \([\text{PtCl}_2(\text{trz})(\text{C}\text{A}\text{N})]\)\(^{2+}\) with the respective 2-arylpyridine ligands, implying...
that the spectra are dominated by CAN-centered absorptions, whereas those involving the trz ligands must be obscured. The lowest-energy absorption maximum shifts from 321 to 370 nm along the sequence 4a \rightarrow 4e, as the expected energies of the lowest π−π* transition of the CAN ligand decrease. As observed for [PtMe(Cl)(flpy)]_2, complex 4e presents a significantly higher molar absorptivity (25900 M\(^{-1}\) cm\(^{-1}\)) with respect to the rest of the derivatives (8100−12100 M\(^{-1}\) cm\(^{-1}\)). Intense absorptions are advantageous for applications such as photocatalysis or bioimaging.

Before examining their luminescence, the photostability of 4a−e was checked by irradiating their solutions in CD\(_2\)CN in quartz NMR tubes with UV light (310 nm) for 6 h at room temperature. Only in the cases of 4a−c were traces of decomposition products observed in the \(^1\)H NMR spectra (ca. 2% of the initial concentration; Figures S15−S19). This behavior is noteworthy because certain tris-chelate Pt(IV) complexes with a mer configuration of metalated aryl groups, mer-[Pt(CAN)]\(_3^2\) (CAN = dfppy, ppy, tpy), isomerize to the fac complexes under irradiation with UV light as a consequence of the population of LMCT excited states.\(^{43,45}\) Instead, complexes 4a−e produce significant luminescent emissions, which were characterized from deaerated CH\(_2\)Cl\(_2\) solutions and poly(methyl methacrylate) (PMMA) matrices (2 wt %) at 298 K and frozen butyronitrile (PrCN) glasses at 77 K. The emission data at 298 K are summarized in Table 2, and the emission spectra in CH\(_2\)Cl\(_2\) solution are shown in Figure 4.

The data at 77 K and the complete series of excitation and emission spectra are included in the Supporting Information. Vibronically structured emissions are observed in all cases, characterized by large Stokes Shifts and lifetimes in the hundreds of microseconds range, which demonstrate a \(^3\)LC emissive state. Given that emission energies decrease in the same order as the lowest-energy absorption, the involved ligand is clearly the cyclometalated 2-arylpyridine, and therefore the trz ligands play a supporting role. In the case of the flpy derivative 4e, a secondary emission band at a higher energy is assigned as fluorescence on the basis of its very short lifetime (<0.2 ns). This band represents a very small fraction of the emitted photons, with a quantum yield of \(\Phi_p \approx 0.005\) in both CH\(_2\)Cl\(_2\) and PMMA. We have previously reported dual fluorescent/phosphorescent emissions from Pt(IV) complexes bearing flpy or other CAN ligands with extended π systems,\(^46,65\) which are due to a relatively less efficient intersystem crossing to the triplet manifold as a consequence of a lower metal orbital contribution to the involved excited states and the reduced spin−orbit coupling effects induced by the metal. Excitation spectra monitored at the phosphorescent emission band correlate with the corresponding absorption profiles in all cases. The excitation spectrum of 4e monitored...
at the fluorescence band coincides with the one monitored at the phosphorescence band in the lower-energy region but shows some differences at higher energies that we tentatively attribute to relatively inefficient internal conversion between higher-lying \( ^1\text{LC(trz)} \) states and the lowest \( ^1\text{LC(ppy)} \) state (see Figure S21 for details). Compared with fac-\([\text{Pt(CAN)}_2]^{+} , [\text{PtMe(Cl)(CAN)}] , [\text{PtCl}_2(\text{trz})(\text{CAN})]^{+} \) the phosphorescent emissions of 4a–e are somewhat blue-shifted, probably as a consequence of the stronger \( \pi \)-acceptor character of the trz ligands relative to 2-arylpyridines,\(^{66} \) leading to a lower energy of metal \( dz \) orbitals and hence a lower MLCT contribution to the emissive excited state.

Quantum yields vary in the range 0.10–0.34 in \( \text{CH}_2\text{Cl}_2 \) and 0.39–0.77 in PMMA matrix and reach the highest values for the derivatives bearing a ppy-based ligand (4a–c). The only previously reported luminescent Pt(IV) tris-chelates with a mer arrangement of metalated aryl rings contain at least one CAN ligand of a relatively low energy for the \( \pi-\pi^* \) transition, namely, mer-\([\text{Pt(ppy)}]^{165} \) and the heteroleptic derivatives mer-\([\text{Pt(ppy)}_2(\text{flpy})]^{165} \) and mer-\([\text{Pt(CAN)}_2(\text{C'AN})^+ \) with CAN = dipp, ppy, C'AN = thpy, 1-phenylisooquinoline (piq),\(^{165} \) and their quantum yields were in the range from 0.03 (for mer-\([\text{Pt(ppy)}_2(\text{piq})]^{+} \) to 0.08 (for mer-\([\text{Pt(ppy)}]^{+} \)).

The radiative and nonradiative rate constants (\( k_r \) and \( k_{nr} \), respectively) for the phosphorescent emissions were calculated assuming that the triplet emissive state is formed with unit efficiency. This assumption introduces a negligible error in the case of 4e because the fluorescence emission has a very low quantum yield.\(^{65} \) The \( k_r \) values are similar to those of complexes fac-\([\text{Pt(CAN)}_2]^{+} \) and \([\text{PtCl}_2(\text{trz})(\text{CAN})]^{+} \) with the same CAN ligands. The lower quantum yields of 4d and 4e are mainly attributable to their lower radiative rates, which are typically found for Pt(IV) complexes bearing thpy\(^{45,46,52} \) and flppy\(^{65} \) ligands and can be explained by a relatively poor metal-ligand orbital overlap, leading to decreased MLCT contributions to the emissive state. The \( k_{nr} \) values are drastically reduced in PMMA matrix in all cases, resulting in significantly higher quantum yields. This indicates that nonradiative deactivation in \( \text{CH}_2\text{Cl}_2 \) solution occurs mainly through molecular motion and collisions with solvent molecules.

**Electrochemistry.** The cyclic voltammograms of complexes 4a–e were registered in MeCN solution and are shown in Figure 5. The potentials of the observed redox processes and estimations of highest occupied/lowest unoccupied molecular orbital (HOMO/LUMO) energies are listed in Table 3. A single irreversible oxidation wave is observable within the accessible potential window for all complexes except 4e, which produces two irreversible waves. The anodic peak potentials decrease in the sequence 4a \( \rightarrow \) 4e, corresponding to increasing HOMO energies as the CAN ligand becomes more electron donating. Therefore, the HOMO is essentially a \( \pi \) orbital of the CAN ligand in all cases, which agrees with the density functional theory (DFT) calculations on 4c (see below). The estimated HOMO energies are similar to those of other Pt(IV) complexes with the respective CAN ligand in a similar coordination environment, i.e., homoleptic mer-\([\text{Pt(CAN)}]^{+} \) or heteroleptic mer-\([\text{Pt(CAN)}_2(\text{C'AN})^+ \) complexes,\(^{66} \) whereas the fac isomers [Pt(CAN)(trz)]\(^{+} \) and the bis-cyclometalated complexes \([\text{PtCl}_2(\text{CAN})(\text{trz})]^{+} \) show lower values.

The first reduction wave is observed at very similar potentials for all complexes and is irreversible, except for 4e, which shows a quasi-reversible wave (Figure S24). Additional reversible processes are observed at more negative potentials, corresponding to the reduction and subsequent reoxidation of species arising from the first reduction. The very similar LUMO energies indicate that this orbital is the same for all complexes and is not affected by the CAN ligand. On the basis of DFT calculations on complex 4c, the LUMO is composed of the combined lowest \( \pi^* \) orbitals of the trz ligands.

**Computational Study.** For a more precise understanding of the properties of complexes 4, DFT and time-dependent DFT (TD-DFT) calculations have been carried out for the tpy derivative 4c. Complete details are presented in the Supporting Information, including fragment contributions to the frontier orbitals (Table S2). Figure 6 shows an orbital energy diagram, including selected isosurfaces. The HOMO is mainly composed of the highest \( \pi \) orbital of the tpy ligand with some \( dz^* \) orbital contribution from the metal (ca. 3%), whereas the LUMO is made of the lowest \( \pi^* \) orbital of the trz ligands and is similarly distributed over them. The LUMO+1 is the lowest \( \pi^* \) orbital of the tpy ligand. The lowest molecular orbital with \( dz^* \) character is LUMO+4.

The TD-DFT results predict essentially LC transitions within the tpy ligand as the most intense, lowest-energy singlet excitations (Table S3), in agreement with the above interpretation of the experimental absorption spectrum. Analogous transitions involving the trz ligand are predicted at higher energies and have lower oscillator strengths. Ligand-to-ligand charge-transfer (LLCT) transitions between the tpy and trz ligands are predicted to occur at low energies, but their oscillator strengths are extremely low and therefore they cannot be identified in the experimental spectrum. The first three triplet excitations correspond to LC transitions within each of the cyclometalated ligands (Table S4), the lowest one involving the tpy, which is consistent with the assignment of the emissive state. The lowest triplet LMCT excitation involving an electronic promotion to LUMO+4 is 1.26 eV above the emissive state (\( ^3\text{T}_{15} \), Table S4; \( \Delta E = 0.64 \) or 0.78 eV for
Table 3. Electrochemical Dataa and HOMO/LUMO Energy Estimationsb for Complexes 4a−e

| complex | $E'_{pa}$c | $E'_{pc}d$ | $E_{1/2}e$ | $E_{HOMO}$ | $E_{LUMO}$ | $\Delta E_{HOMO-LUMO}$ |
|---------|------------|------------|------------|------------|------------|------------------------|
| 4a      | 2.11       | −1.92, −2.17 | −2.04, −2.46 | −6.63      | −2.90      | 3.73                   |
| 4b      | 1.89       | −1.99, −2.23 | −2.08, −2.35, −2.52 | −6.46      | −2.82      | 3.64                   |
| 4c      | 1.79       | −1.99       | −2.16, −2.34, −2.55 | −6.37      | −2.81      | 3.56                   |
| 4d      | 1.68       | −1.92, −2.03, −2.29 | −2.48 | −6.28      | −2.86      | 3.42                   |
| 4e      | 1.58, 1.94 | −1.88, −2.07, −2.27, −2.39 | −6.18      | −2.88      | 3.30                   |

*a In V vs SCE, registered in a 0.1 M solution of (Bu₄N)PF₆ in dry MeCN at 100 mV s⁻¹. b In eV. c Irreversible anodic peak potentials. d Irreversible cathodic peak potentials. e For the reversible waves.

**CONCLUSIONS**

A method to obtain the C₂-symmetrical, bis-cyclometalated species [PtCl₂(trz)₂] (3) has been developed, involving an oxidation of the Pt(II) bis-carbene complex, trans-[PtCl₂(trzH)₂] (1), and the successive electrophilic metalations of the pendant phenyl groups of the trzH ligands. Complex 3 is an excellent precursor for the preparation of cationic tris-cyclometalated complexes [Pt(trz)₃(CAN)]OTf (4) via chloride abstraction with AgOTf in the presence of 2-arylpyridine ligands. These are the first reported Pt(IV) tris-chelates bearing cyclometalated aryl-NHC ligands. Despite presenting a mer configuration of metalated aryls, they are significantly photostable under irradiation with UV light, in sharp contrast with most of the homologous mer-[Pt(CAN)₃]-OTf complexes with the same CAN ligands, which undergo photoisomerization reactions. In addition, they show significant phosphorescent emissions in different media, arising from ³LC states involving the CAN ligands. The computational results show that the energies of deactivating ³LMCT states are high enough not to have an adverse effect on the emissions via thermal population from the emissive state, which is consistent with the observed photostabilities and can be explained by the strong σ-donation from the NHC moieties. Hence, the Pt(trz)₂ subunit is demonstrated as a suitable platform upon which phosphorescent Pt(IV) tris-chelates with tunable emission energies can be built by incorporating different cyclometalated 2-arylpyridines as chromophoric ligands, opening the way to the development of new classes of emissive Pt(IV) complexes by employing other chromophoric bidentate ligands.

**EXPERIMENTAL SECTION**

**General Considerations and Instrumentation.** Preparations were carried out under atmospheric conditions, except for those that required silver reagents, which were conducted in the dark under an N₂ atmosphere. Synthesis grade solvents were employed in all cases. The triazolium iodide salt, [Pr₂N][PtCl₆]₆8 and PhICl₄ were synthesized according to reported procedures. All other reagents were obtained from commercial sources. Elemental analyses were carried out with a LECO CHNS-932 microanalyzer. Electrospray ionization high-resolution mass spectra (ESI-HRMS) were recorded on an
Agilent 6220 Accurate-Mass time-of-flight (TOF) LC/MS. NMR spectra were registered on a 600 MHz Bruker Avance spectrometer at 298 K. The variable-temperature NMR spectra of I were registered on a 300 MHz Bruker Avance spectrometer. Chemical shifts (δ) were referenced to residual signals of nondeuterated solvent and are given in ppm downfield from tetramethylsilane. Abbreviations: s, singlet; d, doublet; t, triplet; q, quartet; p, pentet; sext, sextet; and m, multiplet.

**Preparation of trans-[PtCl₂(trz)₂]** (1). The triazolium salt (250 mg, 0.73 mmol) and Ag₂O (110 mg, 0.47 mmol) were mixed in 1,2-dichloroethane (15 mL), and the resulting suspension was stirred at 50 °C for 14 h and then filtered through Celite. (NTP₂)[PtCl₆] (178 mg, 0.38 mmol) was added to the filtrate, and the mixture was stirred for 14 h at 80 °C and filtered through Celite. The filtrate was evaporated to dryness, and the residue was triturated with MeOH (3 × 4 mL) to give a white solid, which was recrystallized from CH₂Cl₂/ Et₂O and vacuum-dried to give 1. Yield: 217 mg (85%).

**Data for [Pt(trz)₂(dpppy)]OTf (4a)**: Yield: 61 mg (73%).

**General Procedure for the Preparation of [Pt(trz)₂(CN)OTf]** (4). A Carius tube was charged with complex 3 (60 mg, 0.09 mmol), AgOTf (54 mg, 0.21 mmol), the NAC ligand (0.45 mmol), and 1,2-dichlorobenzene (2 mL), and the mixture was stirred at 120 °C for 1 h. After cooling down to room temperature, CH₂Cl₂ (10 mL) was added, and the mixture was filtered through Celite. An excess of NaN₃ was then added, and the suspension was stirred at 110 °C and filtered through Celite. Partial evaporation of the filtrate and addition to Et₂O (10 mL) led to the precipitation of a white solid, which was collected by filtration and vacuum-dried to give the corresponding complex 4.

**Preparation of [PtCl₂(trz)₂]** (2). To a solution of 1 (198 mg, 0.28 mmol) in CH₂Cl₂ (25 mL) was added PhCl (94 mg, 0.84 mmol), and the mixture was stirred at room temperature for 1 h. Partial evaporation of the resulting solution under reduced pressure (2 mL) and addition of Et₂O (10 mL) led to the precipitation of a white solid, which was collected by filtration and vacuum-dried to give 2. Yield: 200 mg (96%).

**General Procedure for the Preparation of [Pt(trz)₂]** (3). A Carius tube was charged with complex 2 (75 mg, 0.10 mmol), Na₂CO₃ (54 mg, 0.51 mmol), and 1,2-dichlorobenzene (3 mL), and the mixture was stirred at 120 °C for 14 h. After cooling down to room temperature, Et₂O (10 mL) was added, and the precipitate was collected by filtration. The product was extracted with CH₂Cl₂ (5 × 5 mL). Partial evaporation of the combined extracts under reduced pressure (2 mL) and addition of Et₂O (10 mL) led to the precipitation of a white solid, which was collected by filtration and vacuum-dried to give 3. Yield: 56 mg (78%).

**Data for [Pt(trz)₂]** (3a): Yield: 61 mg (76%).

**Data for [Pt(trz)₂]** (3b): Yield: 217 mg (85%).
Data for [Pt(trz)(tpy)]OTf (4c). Yield: 65 mg (80%). 1H NMR (600 MHz, CDCl3): δ = 8.04 (d, Jsub = 8.2 Hz, 1H, CH), 7.97–7.91 (m, 2H, CH), 7.80 (d, Jsub = 8.0 Hz, 1H, CH), 7.78 (dd with satellites, Jsub = 8.0, 1.4 Hz, Jsat = 8–11 Hz, 1H, CH), 7.29 (td, Jsub = 7.7, 1.1 Hz, 1H, CH), 7.25 (td, Jsub = 7.7, 1.0 Hz, 1H, CH), 7.15 (td, Jsub = 7.4, 1.0 Hz, 1H, CH), 7.07–7.00 (m, 3H, CH3), 6.69 (dd with satellites, Jsub = 7.2, 1.3 Hz, Jsat = 25 Hz, 1H, CH), 6.60 (dd with satellites, Jsub = 7.8, 1.2 Hz, Jsat = 50 Hz, 1H, CH), 6.54 (s with satellites, Jsub = 27 Hz, 1H, CH), 4.10 (s, 3H, NCH3), 4.09 (s, 3H, NCH3), 2.12 (s, 3H, CH3), 2.06–1.95 (m, 2H, CH2), 1.86–1.79 (m, 1H, CH2), 1.78–1.72 (m, 1H, CH2), 1.20–1.11 (m, 2H, CH2), 0.73 (s, Jsub = 7.2 Hz, 3H, CH3). 13C NMR (150 MHz, CDCl3): δ = 167.1 (PtC), 155.1 (C), 151.3 (C), 151.1 (PtC = 532 Hz, C), 150.2 (CH), 146.3 (C), 145.0 (C), 144.6 (C), 144.2 (C), 141.4 (C), 143.4 (C), 142.9 (Jsub = 37 Hz, C), 142.6 (C), 141.8 (C), 140.3 (CH), 138.7 (C), 135.9 (CH), 133.0 (Jsub = 22 Hz, CH), 131.0 (Jsub = 28 Hz, CH), 128.6 (CH), 127.5 (CH), 126.5 (CH), 126.3 (Jsub = 790 Hz, C), 126.0 (Jsub = 42 Hz, CH), 125.7 (CH), 123.6 (CH), 123.3 (CH), 121.1 (CH), 120.7 (CH), 120.2 (Jsub = 20 Hz, CH), 116.5 (Jsub = 30 Hz, CH), 116.4 (Jsub = 18 Hz, 1H, CH), 47.1 (C), 37.1 (NCH3), 37.0 (NCH3), 32.4 (CH2), 32.3 (CH2), 27.7 (CH2), 27.6 (CH3), 24.3 (CH2), 24.2 (CH2), 23.2 (CH3), 23.1 (CH3), 14.0 (CH3), 13.8 (CH3). HRMS (ESI+) m/z: [M+] Calcd for C44H42F5N10PtS: 893.3617; Found: 893.3632.

Analytical and computational methods and data (PDF)

Access Codes
CCDC 2177317–2177320 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

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Notes
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