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Reactivity of azido terpyridine Pd(II) and Pt(II) complexes towards 4,4,4-trifluoro-2-butynoic acid: structural insight into the triazolato coordination mode†

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The mono- and binuclear azido terpyridine square-planar complexes of ionic formulas, [Pd₂(N₃)₂L]²⁺ and [Pt(N₃)L]⁺ (L = 1,4-bis(2,2':6',2''-terpyridin-4'yl)benzene), underwent the catalyst-free [3 + 2] cycloaddition coupling with 4,4,4-trifluoro-2-butynoic acid at ambient temperature affording the corresponding triazolato complexes. A mixture of triazolato isomers was generated by these inorganic click reactions. An increase in the solubility of the compounds was achieved by replacing the azido ligand with a triazolato ligand. By calculating the vibrational modes and comparing the total electronic and zero-point energy values, the local minimum structures of the complexes and the nature of the predominant triazolato isomer were verified. The theoretical work was complemented with natural bond analysis to get an insight into the natural charge and electronic arrangement of the metal ion, the hybridization of M–L bonds and strength of M–N bonds.

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1. Introduction

One of the key challenges in the field of bioinorganic chemistry is the chemo-selective functionalization of biomolecules with biologically active coordination and organometallic compounds. Actually, some biomolecules can be modified *via* 1,3-dipolar cycloaddition in a targeted manner. Azide complexes have thus far been subjected to click coupling with electrophiles such as alkynes and isothiocyanate to produce five-membered rings.^{1,2} Initially, [Pd(N₃)₂(PPh₃)₂] reacted with trifluoro-acetonitrile, under catalyst-free conditions, affording a tetrazolate complex.³ In their study of the correlation between the reaction kinetics and the nature of the substituent R on the nitrile RCN, Beck and Fehlhammer showed that the rate of the reaction is improved as the withdrawing nature of R group increased.⁴

Later, Paul's research team showed the benefits of using a square-planar complex of the general formula [MN₃L]⁺ (L is a tridentate ligand) in the cycloaddition reactions.⁵ Later on,

several investigations on the reactivity of metal azides of Mo,⁶ W,⁶ Mn,⁷ Ru,⁸ Pd,^{9,10} Pt,^{9,11} Re,^{12–14} and Rh^{15,16}, ions towards some electrophiles were reported. Despite the terminal attachment of the azido ligand to the metal center, the discovered triazolato crystal structures proved unequivocally that the kinetically formed N1 triazolato isomer favored isomerization to the N2 counterpart. The bulkiness of the electrophile, the type of the ancillary ligand, the geometry (square-planar *vs.* octahedral) of the metal complexes, the synthesis and storage conditions, as well as other factors, all play a role in the N2 triazolato isomerization.¹

Kim and coworkers discovered that the click reactions of azido square-planar complexes with isothiocyanates proceeded site-specifically yielding tetrazole-thiolato bound complexes.^{17,18} Pd(II) thiolato–tetrazolate complexes, with *N,N,N*-tridentate ligands, have recently shown higher antifungal activity (MIC = 0.31 μM) against *Cryptococcus neoformans* and *Candida albicans* than their comparable chlorido complexes (MIC = 0.95–1.90 μM) and the reference medication Fluconazole (MIC = 0.41–26.1 μM).¹⁰

Recently, it was examined how four different alkynes reacted to two square-planar Pd(II) and Pt(II) complexes functionalized with the *N,N,S*-tridentate ligand.¹⁹ With the exception of two reactions involving Pt(II) azide and di-alkyl acetylenedicarboxylate, which resulted in a non-equivalent mixture of N1 and N2-isomers, the kinetically generated N1-triazolato isomers tended to isomerize to N2-analogues. Early in the experiment, according to the ¹⁹F NMR kinetics profiles, three triazolato isomers were formed. By the time the experiment was over, all but the dominant N2-triazolato bound isomer remained.¹⁹ Next,

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the reactions of azido Pd(II) and Pt(II) complexes of 2,6-bis(1-ethylbenzimidazol-2'-yl)pyridine with dimethyl acetylenedicarboxylate and 4,4,4-trifluoro-2-butyric acid ethyl ester produced a mixture of the two (or three) anticipated triazolite isomers.¹¹ NMR spectroscopy was used to examine the stability of the triazolite complexes, the mechanism of the [3 + 2] cycloaddition coupling and the nature of the intermediate. The isomer ratio of the triazolite-bound Pt(II) complexes appeared to be controlled by the nature of the solvent. The first example of the catalyst-free 1,3-dipolar cycloaddition reactions of (*trans*-, *trans*-), diazido dihydroxy dipyridine Pt(IV) complex with some electrophiles was reported by Farrer and Sadler.²⁰

In this contribution, we report the reactivity of azido Pd(II) and Pt(II) complexes of 1,4-bis(2,2':6',2''-terpyridin-4'yl)benzene (Scheme 1) towards 4,4,4-trifluoro-2-butyric acid at the ambient temperature under the catalyst-free conditions. Using density functional theory calculations, together with spectral tools, it was possible to learn more about the molecular structures, vibrational properties, and the electronic properties of the anticipated triazolite isomers. In order to learn more about the nature, and strength of the hybridization of the bonds in the coordination sphere surrounding the d⁸ metal ion (Pd(II) or Pt(II)), Weinhold and coworkers' natural bond orbital (NBO) analysis²¹ was carried out for the triazolite isomers.

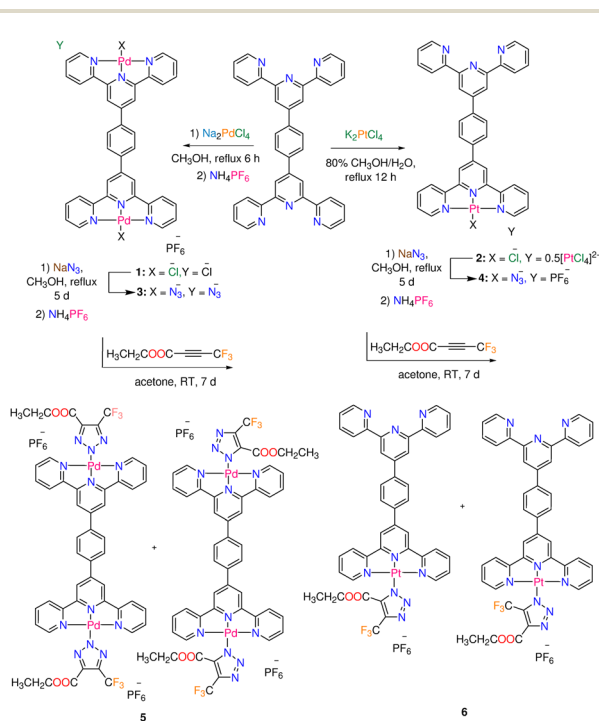
2. Results and discussion

2.1. Synthesis and characterization of complexes

Terephthalaldehyde and eight equivalents of 2-acetylpyridine were reacted in a mixture of ammonia and potassium *tert*-

butoxide to produce 1,4-bis(2,2':6',2''-terpyridin-4'yl)benzene (L) in good yield (77%) and purity.²² Square-planar complexes **1** and **2** (Scheme 1), with ionic formulas of [Pd₂Cl₂L]⁺ and [PtClL]⁺, respectively, were obtained by reactions of terpyridine ligand L and either Na₂PdCl₄ (in methanol) or K₂PtCl₄ (in methanol/water). Under comparable experimental conditions, it appears that the reaction between the Pd(II) precursor and L afforded a binuclear complex **1**, whereas the same reaction with Pt(II) precursor gave a mononuclear complex **2**. The IR spectra of the free ligand and complexes (**1** and **2**) are given in the ESI (Fig. S1–S3†). Complexes **1** and **2** are poorly soluble in DMF, however they breakdown in DMSO, which could be likely attributed to the displacement of chlorido ligands by solvent molecules. These metal-based compounds were therefore subjected to solid state NMR {¹³C, ¹⁵N, ¹⁹F, and ³¹P} analysis (Fig. S4–S8†). In ¹⁹F NMR spectrum of **1** (Fig. S6†), the PF₆⁻ signal of the counter ion is detected at δ = -69.4 ppm. The ¹⁵N NMR spectrum of **1** is characterized by two strong signals at δ = (-154.5, -159.1) ppm as well as three weak signals at δ = (-76.0, -82.7, -85.9) ppm.²³ The presence of two metal centers in **1** may complicate the ¹⁵N NMR spectrum. The ¹⁵N NMR spectrum of **2** (Fig. S8†) displays four signals at δ = -67.5, -71.1, -80.6, and -174.3 ppm. The Δ¹⁵N_{coord} parameter, the difference between the chemical shifts of free nitrogen atom and M–N bond, gives an idea about the coordination of the heteroatom.²³ Compared to **1**, the Δ¹⁵N_{coord} of the signal at δ = -174.3 ppm of **2** is relatively high, which is typical for both Pt central atom and the nitrogen *trans*- to Cl.^{23,24} The atmospheric solids analysis probe mass spectrometry (ASAP-MS) spectrum of **1** (in the positive mode, Fig. S9†) shows two distinct fragments at *m/z* 968.9147 (calcd 968.9150) and *m/z* 683.0772 (calcd 683.0768) equivalent to {[Pd₂Cl₂L]PF₆}⁺ and {[PdClL]}⁺, respectively. The ASAP-MS spectrum of **2** (Fig. S10†) displays a unique fragment at *m/z* 770.1393 (calcd 770.1393) corresponding to {[PtCl]}⁺.

As shown in Scheme 1, azido Pd(II) and Pt(II) complexes (**3** and **4**) were obtained by ligand exchange and gathered in pure form. The distinctive vibrational spectral feature in spectra of **3** and **4** is the prominent ν(N₃) band at 2037 and 2022 cm⁻¹, respectively (Fig. S11 and 12†). The solubility of **3** and **4** in DMF is poor, while both complexes decompose in DMSO. Thus, these metal-azide complexes were analysed with solid-state NMR {¹³C, ¹⁵N, ¹⁹F, and ³¹P} (Fig. S13–S19†). In ¹⁹F and ³¹P NMR spectra of **3** (Fig. S15 and S16†), the PF₆⁻ signals are detected at δ = -69.3 and -142.9 ppm, respectively. The ¹⁵N NMR spectrum of **3** (Fig. S14†) shows a set of six signals at δ = (-75.9, -84.6, -86.6), and (-158.6, -160.4, -164.8) ppm corresponding to the terminal and central pyridine rings as well as three signals at δ = -122.0, -207.8 and -309.6 ppm equivalent to the coordinated azido ligand.^{25,26} The most clear ¹⁵N signals in the NMR spectrum of **4** (Fig. S18†) are observed at δ = -68.2, -71.6, and -306.4 ppm. The ¹⁹F signal of PF₆⁻ is seen at δ = -69.4 ppm in the spectrum of **4**. Two distinct fragments at *m/z* 982.9961 (calcd 982.9957) and 688.1186 (calcd 688.1184) matching to {[Pd₂(N₃)₂L]·PF₆}⁺, and {[Pd(N₃)L]}⁺ are visible in the ASAP-MS spectrum of **3** (Fig. S20†). There is a distinct peak (Fig. S21†) at *m/z* 778.1813 (calcd 778.1799) in the mass spectrum of **4**, which corresponds to {[Pt(N₃)L]}⁺. The reactivity of azido complexes (**3**



Scheme 1 Synthesis of Pd(II) and Pt(II) complexes **1**–**6** functionalized with terpyridine derivative L (main species are shown in the cases of **5** and **6**).



and 4) towards $\text{CF}_3\text{-C}\equiv\text{C-COOCH}_2\text{CH}_3$ was examined at the ambient temperature. The inorganic click reactions (iClick) were carried out by suspending the azido complexes in an acetone solution of the additional alkyne, as illustrated in Scheme 1. To precipitate the complexes in their purest form, a saturated solution of ammonium hexafluorophosphate was added to the triazolite reaction mixtures. The formation of triazolite complexes 5 and 6 is clearly determined by the disappearance of the strong $\nu(\text{N}_3)$ band, and the growth of the distinctive (C=O) band of the triazolite moiety at around 1727 cm^{-1} (Fig. S22 and S23†). Unlike complexes 1–4, the triazolite derivatives 5 and 6 have good solubility in DMSO. The positively charged mass fragments (Fig. S24†) at m/z 854.1431 (calcd 854.1426) and m/z 1315.0461 (calcd 1315.0440) in the spectrum of 5 are assigned to $\{[\text{Pd}_2(\text{triazolato})_2\text{L}]\cdot\text{PF}_6\}^+$, and $\{[\text{Pd}(\text{triazolato})\text{L}]\}^+$ in that order. The prominent peak at m/z 1110.2287 (calcd 1110.1639) designated to $\{\text{Na}[\text{Pt}(\text{triazolato})\text{L}]\cdot\text{PF}_6\}^+$ is what distinguishes the ESI MS spectrum of 6 (Fig. S25†).

The literature claims that the N1 or N2 atoms of the triazolite ligand, synthesized from symmetrical alkyne, can bind M^{n+} ion. Despite the azido ligand being terminally attached to Pd(II) or Pt(II) ions, X-ray crystallographic examination of numerous triazolite complexes with M^{n+} ions other than d^8 ions revealed conclusive evidence for the isomerization of the kinetically generated N1 (or N3) isomer into the N2 analogue.^{12,27,28} A combination of two (or three) isomers can occur when the conversion of N1 (or N3) to N2 is incomplete.^{29,30} The geometry of the reacting azide complex (square-planar, piano-stool or octahedral), the bulkiness of the electron-poor alkynes, the experimental conditions (coordinating solvents, reactants ratio, temperature, etc.), as well as the nature of the ancillary ligands all have an impact on the final triazolite coordination mode.¹ In our situation, the reaction of 3 (or 4) with the asymmetric alkyne is anticipated to result in the formation of three triazolite bound modes (N1, N2 and N3) (Fig. 1). The triazolite isomers bearing $-\text{COOCH}_2\text{CH}_3$ and $-\text{CF}_3$ groups close to the ligand framework, respectively, are referred to as N1 and N3, while the name N2 denotes the coordination of the central triazolite N to the metal centre (Fig. 1). The NMR spectra of 5 are shown in Fig. S26–S30.† The ^1H NMR spectrum of 5 (Fig. S26†) is broad with the primary feature being two ethyl ester signals at 4.39 and 1.41 ppm as well as few aromatic signals in the range of 7.81–8.74 ppm. Two minor ^1H NMR signals for the ethyl ester group are seen clearly at around 4.12

and 1.08 ppm supporting the presence of another isomer. The ^{13}C NMR spectrum of 5 (Fig. S27†) shows the characteristic signals for $\text{CF}_3\text{-C}$ at $\delta = 138.7$ ppm (q, $^2J_{\text{C,F}} = 39.4$ Hz), and 121.7 ppm (q, $^1J_{\text{C,F}} = 267.0$ Hz) as well as three unique signals for C=O, $-\text{CH}_2$ and $-\text{CH}_3$ functional groups at 159.2, 60.9 and 13.7 ppm, respectively. The ^{19}F and ^{31}P signals of PF_6^- are seen at $\delta = -70.2$ ppm (d, $^1J_{\text{PF}} = 711$ Hz) and $\delta = -144.2$ (sept, $^1J_{\text{PF}} = 711$ Hz) in the spectra of 5, in that order. As indicated by the few tiny signals observed in the ^{19}F NMR spectrum of 5 (Fig. S28†), the presence of two metal centres in 5 undoubtedly increases the number of possible triazolite isomers. According to ^{19}F NMR spectroscopy (Fig. S28†), the cycloaddition reaction of 5 with 4,4,4-trifluoro-2-butynoic acid ethyl ester gave mainly a mixture of N1-, and N2-isomers, with the $-\text{CF}_3$ group displaying two singlet signals of different intensities at $\delta = -59.03$, and -58.95 ppm, respectively. If the triazolite N3-isomer is present in the reaction mixture of 5, its ^{19}F NMR signal should be at least 4 ppm apart from those of N1 and N2.¹¹ The up-field change in the $-\text{CF}_3$ ^{19}F -signal from $\delta = -52$ ppm in the starting coupling agent to higher chemical-shift values in 5 could suggest triazolite formation.¹² Complex 5 has low solubility in most organic solvents except DMSO, therefore we couldn't dissolve it in non-coordinating solvents. We discovered in prior contributions to the iClick reaction that performing the NMR measurements in DMSO increases the conversion of N1- into N2-isomer¹¹ and consequently we allocated the highest intensity ^{19}F NMR signal at $\delta = -58.95$ ppm to the N2-isomer. According to ^1H and ^{19}F NMR studies, the reaction of 4 with the $-\text{CF}_3$ alkyne produced the three expected triazolite isomers (Fig. 1). The presence of isomers and the prospect of rapid isomerization during the NMR experiments make determining its ^1H NMR spectrum highly challenging. The crucial spectrum characteristics for the progress of the inorganic click reaction and the formation of the triazolite molecule, on the other hand, were discovered. This includes the appearance of ethyl ester signals in the ^1H NMR spectrum, as well as the presence of ^{19}F NMR signals with greater chemical-shift values than the original CF_3 alkyne signal. We detected the presence of PF_6^- counterion by monitoring the ^{19}F and ^{31}P NMR signals, as we had done before with the chlorido and azido complexes.

2.2. Density functional theory calculations

The analytical and spectroscopic evidence in the preceding section demonstrated the mono- and binuclear nature of the isolated triazolite Pt(II) and Pd(II) complexes. In 5 and 6, tridentate terpyridine derivative and monodentate triazolite ligand are what make up the square-planar arrangement that surrounds the metal core. Utilizing the Becke 3-parameter (exchange) Lee–Yang–Parr functional^{31,32} and the LANL2DZ basis set,^{33,34} ground-state geometry optimization and the vibrational analysis of all the isomers were performed in order to get insight into the triazolite binding mode of the formed isomers and their stabilities. The harmonic frequency analysis was used to verify that the obtained geometries were local minima. Except for the N2 isomer of the Pt(II) complex, we could not discover any imaginary modes in any of the optimized

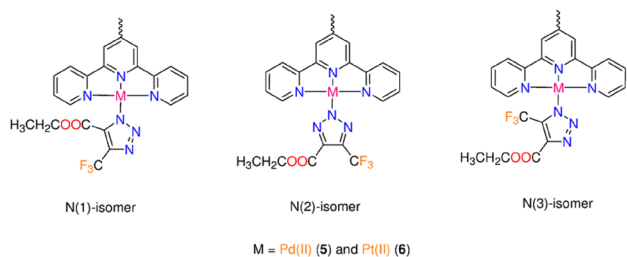


Fig. 1 The three triazolite bound isomers formed from reaction of 3 (or 4) with 4,4,4-trifluoro-2-butynoic acid ethyl ester.



geometries. Next, the total electronic and zero-point energy values were compared (Table S1†). Comparatively, it is important to note that the N2-isomer of the Pd(II) complex is more stable than N1 and N3 isomers by 3.5 and 19.7 kJ mol⁻¹, respectively, and that the N1 is therefore about 16.2 kJ mol⁻¹ more stable than the N3 isomer. In terms of the presence of the N1 and N2 isomers, these conclusions agree with the experimental NMR data. The calculated bond lengths and bond angles around the d⁸ metal ions are presented in Tables S2 and Table S3.† The variation in the bonds and angles of the three isomers were compared to each other. The triazolate rings in the N1 and N3 isomers of **5** are *trans* to one another and rotated off the plane by 57.0° and 55.0°, respectively, according to the dihedral angle of N73–N69–Pd–N35. The triazolate moiety in N2 isomer is nearly in the plane of the terpyridine ligand system at equal distances from terpyridine framework. The two Pd(II) moieties of the binuclear system in the three isomers are nearly

perpendicular to one another and have the same bond length, as illustrated in Fig. 2. As opposed to the N1 and N3 isomers, the two N–N bonds of the triazolate ring are nearly equivalent in the N2 isomer. The distance to the triazolato ligand is 2.027, 2.028 and 2.032 Å in the three isomers of **5**, respectively.

Likewise, the N1 isomer of the Pt(II) complex is more stable than the N3 analogue by about 4.35 kJ mol⁻¹. Like the N2-isomer of **5**, the triazolate moiety of the N2 isomer in **6** is almost in the plane of the terpyridine ligand system with similar distances from the terpyridine framework. The distance to the triazolato ligand is 2.038, 2.033 and 2.042 Å in the three isomers of **6**, respectively, that are longer than the equivalent ones in the Pd(II) isomers. The triazolate ring in the N1 and N3 isomers of **6** is rotated from the plane by 55.0° and 52.0° as defined by the dihedral angles of N69–N67–Pt–N35. When compared to those found in the N1 and N3 isomers, the distances between Pt(II) ion and nitrogen atoms of the terpyridine framework in the N2 isomer are shorter (Fig. 3).

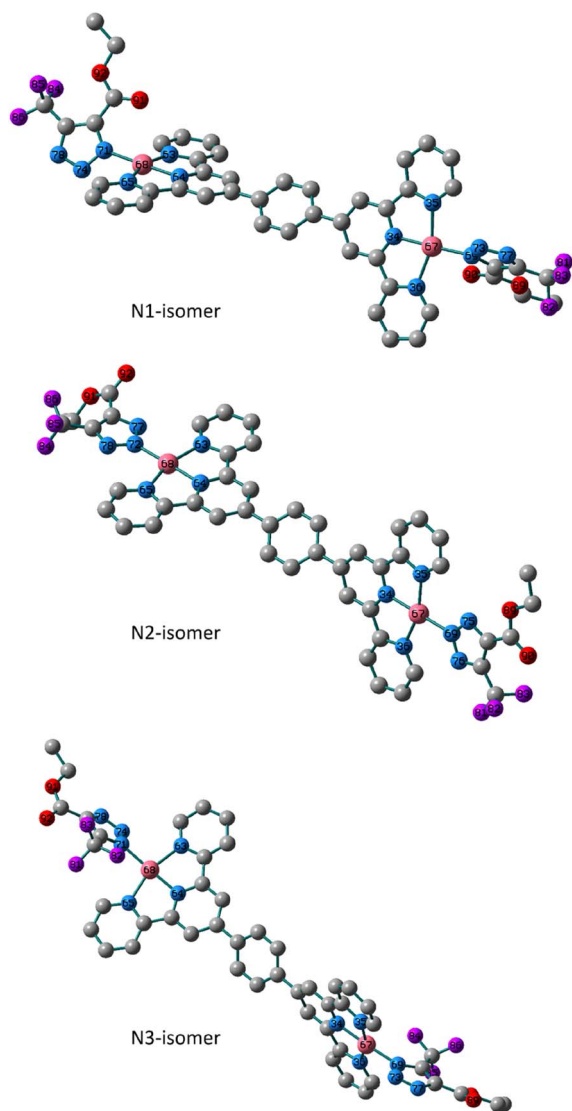


Fig. 2 Local minimum structures of the optimized triazolate binuclear Pd(II) isomers of **5** (N1, N2 and N3) calculated at B3LYP/LANL2DZ level of theory. (Hydrogen-atoms were omitted for the clarity).

2.3. Natural bond orbital analyses

It was feasible to determine the natural charge, type of bonding, type of hybridization, and strength of the M–N bonds in all the three triazolate isomers of compounds **5** and **6** using the natural bond orbital (NBO) analysis of Weinhold and co-workers²¹ and the second order perturbation theory analysis. In addition, the NBO technique has been widely employed in the evaluation of

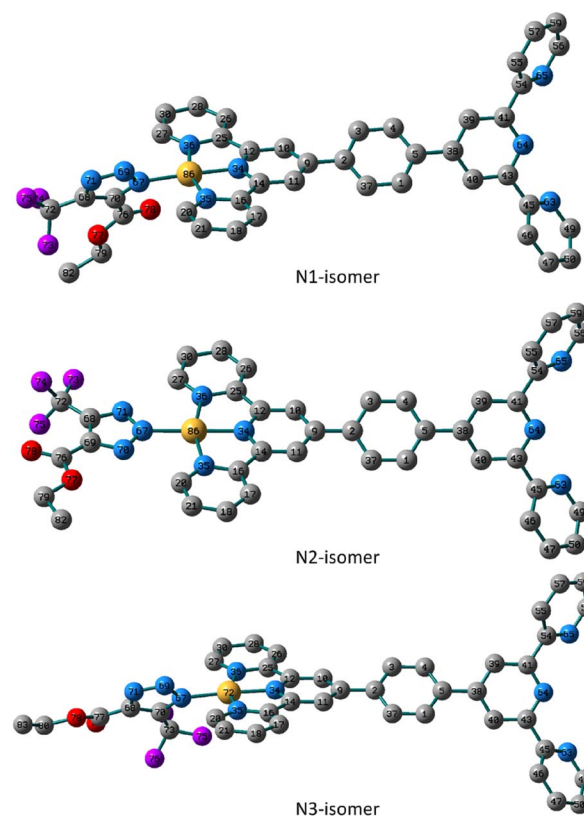


Fig. 3 Local minimum structures of the optimized triazolate binuclear Pt(II) isomers of **6** (N1, N2 and N3) calculated at B3LYP/LANL2DZ level of theory. (Hydrogen-atoms were omitted for the clarity).



intra- and intermolecular contacts, and it provides a suitable basis for studying charge transfer or conjugative interactions in molecular systems. NBO analysis was carried out to determine the contribution of atomic orbitals (AOs) to NBO σ and π hybrid orbitals for bonded atom pairs. There are three NBO hybrid orbitals defined: bonding orbital (BD), lone pair (LP), and core (CR). The ideal Lewis structure is built from Lewis σ -type (donor) NBOs and non-Lewis σ -type (acceptor) NBOs. Covalency in molecules is defined by filled NBOs, whereas anti-bonds stand for empty valence-shell capacity and span regions of the atomic valence space that are suitably unsaturated by covalent electrons. Weak valence anti-bond occupancies indicate intricate deviation from an idealised localised Lewis picture, *i.e.*, real “delocalization effects.” The electronic configuration of the metal centre (Pd or Pt) of the triazolite isomers (N1, N2 and N3), its 4d (or 5d) electronic distribution and its natural atomic charge are presented in Table S4.† According to the NBO analysis, there appears to be no substantial difference between the electronic structures of the triazolite isomers of **5**. For example, the electronic configurations of the Pd ions are essentially identical, and the natural charges of the metal ions have not changed significantly, with all values falling within the range of 0.68069–0.69165. The difference between the two natural charges in the N3 isomer is insignificant (10^{-4} e), but it is 0.0023e in the N2 isomer. The computed natural charge on Pd atom is much lower than the formal charge (+2) due to electron density donation from the terpyridine and triazolite ligands. The population of 4d(Pd) of 9.33 does not match to the oxidation state Pd(II), which is consistent with ligand-to-d(Pd) electron transfer. On the other hand, the electronic structures of the two metal ions in the same molecule do not differ much. For example, the electronic arrangements of Pd ions in N2-isomer of **5** are nearly the same, $[\text{Kr}]5s^{0.32}4d^{8.80}5p^{0.20}5d^{0.01}$. The calculated natural charges of Pd ions are 0.68069e and 0.68299e. The occupancies of 4d orbitals are as follows: $d_{xy}^{1.97206}$ $d_{xz}^{1.94229}$ $d_{yz}^{1.85229}$ $d_{x^2-y^2}^{1.1085}$ $d_{z^2}^{1.92903}$ and $d_{xy}^{1.95252}$ $d_{xz}^{1.96486}$ $d_{yz}^{1.774671}$ $d_{x^2-y^2}^{1.47997}$ $d_{z^2}^{1.65949}$. In comparison, the difference in the natural charges between the two metal centres in the N2 isomer appears to be attributable mainly to a change in the electron occupancies between $d_{x^2-y^2}$ and d_{z^2} orbitals.

Hyperconjugation is described as a stabilising effect caused by the overlap of one occupied and another electron deficient orbital. The hyper-conjugative interaction energy between the donor–acceptor orbitals was calculated using the second-order perturbation technique of the Fock Matrix in NBO basis. The stronger the electronic interactions between the donors such as ligand coordination sites and acceptors *e.g.*, metal ions, the higher the value of second order interaction energy (E^2). The higher the inclination of electron donors to donate to electron acceptors, and the larger the level of conjugation of the entire system. These interactions can be identified by observing a rise in the electron density of the anti-bonding orbital, which weakens the respective bonds. According to NBO analysis, the four bonds between Pd and the N-atoms can be described as donation of electron density from a lone pair (LP) orbital on each nitrogen atom to palladium molecular orbitals. The E^2 values of LP(1)N34 \rightarrow , LP(1)N35 \rightarrow , LP(1)N36 \rightarrow , and LP(1)N69

\rightarrow RY*(1)Pd are 0.34, 0.88, 0.88, and 0.83 kcal mol $^{-1}$, respectively. The corresponding E^2 values of the other metal centre in the same molecule, LP(1)N63 \rightarrow , LP(1)N64 \rightarrow , LP(1)N65 \rightarrow , and LP(1)N72 \rightarrow RY*(1)Pd are 0.91, 0.38, 0.91 and 0.90 kcal mol $^{-1}$, respectively. It can be seen that the interactions between the terminal nitrogen atoms of the terpyridine ligand and Pd ion are equal, and they are stronger than those of the central terpyridine nitrogen and triazolite nitrogen atom. There is a minor discrepancy in the E^2 values of the M–N bonds between the two metal centres.

Like the Pd(II) triazolite complexes, the electronic configuration of the Pt(II) ion in the three isomers ($[\text{Xe}]6s^{0.47}5d^{8.59}6p^{0.20}6d^{0.01}$) are the same (Table S4†). Thus, 67.99 core electrons, 9.26 valence electrons (on 6s, 5d, and 6p atomic orbitals), and 0.01 Rydberg electrons (on 6d orbital) give the 77.25 electrons. The natural charges of the Pt ion have not changed significantly among the three isomers, 0.71638e (N1), 0.73683e (N2) and 0.70854e (N3). The calculated natural charge on Pt atom is substantially lower than the formal charge (+2) due to electron density donation from the surrounding ligands. The population of 5d(Pt) of 9.27 does not match to the oxidation state Pt(II), which is consistent with ligand-to-d(Pt) electron transfer. In the N2 isomer, the σ (Pt–N34) bond is formed from a $sp^{2.85}$ hybrid on terpyridine N34 atom (which is the mixture of 26.00% s, and 74.00% p atomic orbitals) and $sp^{0.40}d^{5.87}$ hybrid on platinum atom (which is the mixture of 13.77% s, 5.46% p and 80.77% d atomic orbitals). NBO analysis indicated that the σ (Pt–N34) bond is strongly polarized towards the nitrogen atom with about 75.86% of electron density concentrated on the nitrogen atom. In the most stable triazolite isomer, the N1 isomer, the four bonds formed by Pt and N-atoms are defined as electron density donations from a lone pair (LP) orbital on each nitrogen atom to Pt molecular orbitals. The E^2 values of LP(1)N34 \rightarrow , LP(1)N35 \rightarrow , LP(1)N36 \rightarrow , and LP(1)N67 \rightarrow RY*(4)Pt are 0.15, 0.48, 0.39, and 0.73 kcal mol $^{-1}$, respectively. It can be seen that the interaction between the triazolite N atom and Pt ion is stronger than those of the terpyridine nitrogen atoms.

2.4. Time-dependent density functional theory calculations

To get an insight into the nature of electronic transitions anticipated to be seen in the electronic spectra of the three triazolite isomers of complexes **5** and **6**, time-dependent density functional theory (TDDFT) calculations were performed in the singlet state using a Becke 3-parameter (exchange) Lee–Yang–Parr functional^{31,32} with LANL2DZ basis set^{33,34} and PCM solvation model (DMSO). GAUSSSUM was used to simulate the electronic spectra. A Gaussian convolution with a full-width at half-maximum (FWHM) of 3000 was used to incorporate each excited state. The three triazolite isomers of **5** have virtually identical electronic absorption spectra (Fig. S31†), having two main absorption bands at around 307 and 370 nm. The band at 370 nm shifts slightly when we move from N1, to N3, and finally to N2. The lowest energy band at 307 in all the three isomers is due to HOMO \rightarrow LUMO. As seen in Fig. 4, HOMO is primarily formed of π (triazolite), whereas LUMO is primarily composed of the terpyridine π -framework with a tiny



at 305 nm (HOMO−9 → LUMO/HOMO−3 → LUMO+1) is unique as it comprised of π - π^* within the triazolate moiety as well as LLCT as that observed in the case of the other isomers. Therefore, all the three isomers of **5** demonstrated two kinds of transitions, LLCT and MLCT.

The electronic absorption spectra (Fig. S32†) of the triazolate isomers of **6** are characterised by two major bands at roughly 296 and 400 nm, with the higher energy band having a greater oscillator strength value than the lower energy band. The band at around 400 nm is due to HOMO → LUMO. As shown in Fig. 5, HOMO orbital of N1 isomer is contained upon π -system of the phenyl ring of the terpyridine ligand, and LUMO is of d(Pt). Thus, the band at 400 nm is assigned to LMCT. For N2 and N3-isomers, the transitions at 408 and 403 nm may be assigned to LLCT/MLCT initiating from π (uncoordinated terminal pyridyl rings) and terminating at the electron-deficient π -system of the coordinated pyridyl rings/d(Pt). Changing the coordination mode of the triazolate ring from N1 to N2 or N3 appears to have a significant impact on the overall electronic structure of this class of Pt(II) complexes. The band at 293 nm in N1 isomer may be assigned to LLCT from π (phenyl) → π^* (coordinated terminal pyridyl ring). The band at around 295 nm in both N2 and N3-isomers may be attributed to LLCT from the electron donor free-terminal pyridyl rings to π -system of the terpyridine ring. Overall, it seems that the electronic structure of the N2 isomer is more similar to that of N3.

3. Conclusion

To evaluate the opportunity of functionalizing biomolecules with $[\text{Pd}_2(\text{Cl})_2\text{L}]^{2+}$ and $[\text{Pt}(\text{Cl})\text{L}]^+$ (L = 1,4-bis(2,2':6',2''-terpyridin-4'yl)benzene) through the inorganic click reaction, the chlorido ligand(s) was exchanged with N_3^- group, which was then coupled with electron poor alkyne (4,4,4-trifluoro-2-butyne acid) under the catalyst-free conditions, yielding triazolate analogues. Under similar experimental conditions, it appears that the reaction between the Pd(II) precursor and terpyridine ligand afforded a binuclear complex, whereas the same reaction with Pt(II) precursor gave a mononuclear complex. The chloro and azido complexes are insoluble or break down in most organic solvents, however, the corresponding triazolate complexes are soluble in the same solvents. Therefore, the solubility of this class of compounds could be improved by exchanging the chlorido ligand for a triazolate moiety. Solid-state NMR and ASAP-MS were applied to elucidate the structures of insoluble complexes.

The reactivity of the azido complexes towards 4,4,4-trifluoro-2-butyne acid as well as the nature of the isolated triazolate isomers were performed by NMR spectroscopy particularly ^{19}F NMR. In our situation, the reactions of the azido complexes with the asymmetric alkyne is anticipated to result in the formation of three triazolate bound modes (N1, N2 and N3). According to the ^{19}F NMR spectroscopy, the cycloaddition reaction of the azido Pd(II) complex gave mainly a mixture of N1-, and N2-isomers. Parallely, the total electronic and zero-point energy values of the three isomers of the Pd(II) binuclear complex were compared. The calculations showed that N2-isomer is more stable than N1

and N3 analogues as well as the N1 isomer is more stable than N3 that agrees with the experimental data. In the example of the Pt(II) complex, the comparison of total electronic and zero-point energy values suggested that the most prevalent species of this complex should be either N1 or N3. Time-dependent density functional calculations showed that there is no significant difference in the electronic structures of the three isomers and the observed transitions might be assigned to LLCT or MLCT. According to the natural bond orbital analysis, there appears to be no substantial difference, in terms of natural charge and the populations of the d-subshell, between electronic structures of the three triazolate isomers. It can see that the interactions between the terminal nitrogen atoms of the terpyridine ligand and Pd ion are equal, according to the values of second order interaction energy ($E^{(2)}$), and they are stronger than those of the central terpyridine nitrogen and triazolate nitrogen atom. On the contrary, it can see that the interaction between the triazolate N atom and Pt ion is stronger than those of the terpyridine nitrogen atoms. Overall, this contribution sheds light on how to bio-conjugate square-planar complexes using the inorganic click reaction, how to improve the solubility of extremely insoluble intriguing complexes, and the nature of the generated species in relation to the nature of the metal ions.

4. Experimental section

4.1. Materials and instruments

On oven-dried glassware, the experiments were conducted in anhydrous solvents. Ammonium hexafluorophosphate, sodium azide, sodium tetrachloropalladate, and potassium tetrachloroplatinate(II), were all purchased from commercial sources and utilised exactly as they were given. 1,4-Bis(2,2':6',2''-terpyridin-4'yl)benzene,²² and 4,4,4-trifluoro-2-butyne acid ethyl ester,³⁵ were synthesized by following the published schemes. Metal azides can degrade quickly and unexpectedly. Scratching metal azides can result in explosive reactions. Only small-scale synthesis should be carried out, and heating of solid compounds must be avoided at all costs. As a result, handling and purification must be done with the utmost care. The CHN elemental compositions were recorded using either the HEKtech EA 3000 elemental analyzer or the Elementar Vario MICRO cube CHN analyzer. On a Thermo Fisher Exactive Plus Orbitrap mass, electrospray ionisation mass spectra were collected at a resolution of $R = 70,000$ and a solvent flow rate of 50 L min. To collect vibrational spectra in the solid state, a Nicolet 380 FT-IR spectrometer with a smart iFTR adapter was employed. ^1H , ^{13}C , ^{19}F and ^{31}P NMR spectra were recorded with Bruker-Avance 500 (^1H , 500.13 MHz; $^{13}\text{C}\{^1\text{H}\}$, 125.77 MHz; ^{19}F , 470.59 MHz; ^{31}P , 202.46 MHz) or Bruker-Avance 400 (^1H , 400.40 MHz; $^{13}\text{C}\{^1\text{H}\}$, 100.68 MHz; ^{31}P , 162.08 MHz) spectrometers. Using two-dimensional NMR techniques, $\{^1\text{H}, ^1\text{H}\}$ COSY and $\{^1\text{H}, ^{13}\text{C}\}$ HSQC, NMR signal assignments were carried out.

4.2. Synthesis

4.2.1. Synthesis of chloro complexes. 1: Methanol (40 mL) was added to a round-bottom flask that had already been



charged with sodium tetrachloropalladate (1.07 mmol; 315 mg) and 1,4-bis(2,2':6',2''-terpyridin-4'yl)benzene (0.50 mmol; 270 mg). After that, the reaction mixture was heated to reflux for 6 h. Yellow precipitate was filtered out, cleaned with methanol, chloroform, and diethyl ether, and then dried under vacuum. The product was suspended in methanol/water (75% v/v) solution mixture of ammonium hexafluorophosphate. Stirring was done overnight. Greenish-yellow precipitate was collected by filtration, washed with water, diethyl ether, and dried *in vacuo*. Complex 1 decomposes in DMSO- d_6 and had no solubility in the rest of the organic solvents. Based on elemental analysis, it appears that the counterion exchange is incomplete. Yield: 79% (397 mg, 0.39 mmol). IR (ATR): $\nu = 3113$ (w, CH), 1606 (s, CC/CN), 1479, 1399, 1247, 832 (s), 786, 741 cm^{-1} . ^{13}C -NMR (solid-state, 100.68 MHz): $\delta = 159.3, 156.4, 153.9, 152.3, 151.0, 147.6, 146.3, 142.2, 141.5, 140.0, 138.5, 136.6, 133.6, 130.5, 128.0, 123.6, 122.9, 119.9, 118.0, 115.1, 114.5$ ppm. ^{15}N NMR (solid-state, 40.56 MHz): $\delta = -159.1, -154.5, -85.9, -82.7, -76.0$ ppm. ^{19}F -NMR (solid-state, 396.5 MHz): $\delta = -69.4$ (br, PF_6^-) ppm. ASAP (ESI $^+$): 968.9147 $\{[\text{Pd}_2\text{Cl}_2\text{L}]\text{PF}_6\}^+$, 683.0772 $\{[\text{PdClL}]\}^+$. $[\text{Pd}_2\text{Cl}_2\text{L}]\text{Cl}\cdot\text{PF}_6\cdot\text{H}_2\text{O}$ ($\text{C}_{36}\text{H}_{24}\text{Cl}_3\text{F}_6\text{N}_6\text{PPd}_2\cdot\text{H}_2\text{O}$), C, 42.27; H, 2.56; N, 8.22; found C 42.67, H 2.74, N 8.30.

2: A 5 mL aqueous solution of K_2PtCl_4 (1.0 mmol; 415 mg) was poured to a round-bottomed flask that was already charged with terpyridine derivative (0.5 mmol; 270 mg) and 50 mL of methanol. For 12 h, the reaction mixture was heated to reflux, at which point brown precipitate formed. Filtration was done. The product was cleaned with water and diethyl ether, and then dried under vacuum. Yield: 66% (312 mg, 0.42 mmol). IR (ATR): $\nu = 3515$ (br, OH_2), 3060 (w, CH), 1584 (m, CC/CN), 1471, 1394, 1256, 1120, 896, 787, 739 cm^{-1} . Complex 2 decomposes also in DMSO- d_6 and did not dissolve in the remaining organic solvents. ^{13}C -NMR (solid-state, 100.68 MHz): $\delta = 155.3, 154.7, 152.7, 149.5, 145.1, 144.4, 143.9, 142.7, 140.6, 135.4, 134.3, 131.0, 129.1, 127.8, 126.3, 125.7, 124.9, 123.7, 122.6, 120.8, 119.3$ ppm. ^{15}N NMR (solid-state, 40.56 MHz): $\delta = -177.0, -174.3, -80.8, -70.9, -67.3$ ppm. ASAP (ESI $^+$): 770.1393 $\{[\text{M}-0.5\text{PtCl}_4]\}^+$. $\text{C}_{36}\text{H}_{24}\text{ClN}_6\text{Pt}\cdot 0.5\text{PtCl}_4\cdot 4\text{H}_2\text{O}$: C, 44.32; H, 2.89; N, 8.61, found C, 44.20; H, 2.95; N, 8.60.

4.2.2. Synthesis of azido complexes. 3: Sodium azide (0.015 mol; 1 g) was added to a round-bottomed flask that had been charged with complex 1 (0.35 mmol; 350 mg) and 100 mL of a 60% (v/v) methanol–water mixture. For 5 days, the reaction mixture was heated at 50 °C. Then, ammonium hexafluorophosphate (6.13 mmol; 1 g) was added to the reaction mixture. Stirring was continuing for 6 h. A yellow precipitate was filtrated off, rinsed with water and diethyl ether, and then dried for a few days under vacuum. Complex 3 is insoluble in most organic solvents and breaks down in DMSO- d_6 . Yield: 73% (289 mg, 0.25 mmol). IR (ATR): $\nu = 3113$ (w, CH), 2037 (s, N_3), 1605 (s, CC/CN), 1478, 1399, 1292, 1249, 832, 784, 742 cm^{-1} . ^{13}C -NMR (solid-state, 100.68 MHz): $\delta = 159.1, 157.1, 155.0, 154.0, 152.4, 150.7, 147.7, 144.0, 141.3, 139.8, 138.2, 136.8, 135.4, 133.8, 130.5, 128.0, 126.4, 123.7, 119.8, 118.4, 115.3, 114.3$ ppm. ^{15}N NMR (solid-state, 40.56 MHz): $\delta = -309.6, -207.8, -164.7, -160.4, -158.6, -122.0, -86.6, -84.6, -75.9$ ppm. ^{31}P -NMR (solid-state, 161.94 MHz): $\delta = -142.9$

(sept, $^1J_{\text{PF}} = 693$ Hz, PF_6^-) ppm. ^{19}F -NMR (solid-state, 396.5 MHz): $\delta = -69.3$ (br, PF_6^-) ppm. ASAP (ESI $^+$): 982.9969 $\{[\text{Pd}_2(\text{N}_3)_2\text{L}]\cdot\text{PF}_6\}^+$, 688.1186 $\{[\text{Pd}(\text{N}_3\text{L})]\}^+$. $\text{C}_{36}\text{H}_{24}\text{F}_{12}\text{N}_{12}\text{P}_2\text{Pd}_2\cdot 6\text{H}_2\text{O}$: C 35.00, H 2.94, N 13.60, found C 34.63, H 2.57, N 12.75. The elemental analysis data for synthetic coordination and organometallic complexes are frequently complicated by the sensitivity of M–C, M–N, ... *etc.* bonds to oxidation.³⁶ Additionally, the elemental analyses of compounds containing metal azide and triazolate occasionally exceed the allowable error. As a result, it is normal for the reported inaccuracy (>0.4%) in the elemental analysis of several azide and triazolate compounds.^{7,37}

4: A round-bottomed flask was filled with 200 mg of compound 2 (0.21 mmol) and 100 mL of a 60% (v/v) acetone–water solution before being charged with sodium azide (0.015 mol; 1 g). For 5 days, the reaction mixture was heated at 50 °C. Then, ammonium hexafluorophosphate (6.13 mmol; 1 g) was added to the reaction mixture. Stirring was continuing for 6 h. A brown precipitate was filtrated off, rinsed with water and diethyl ether, and then dried for a few days under vacuum. Complex 4 is insoluble in most organic solvents and breaks down in DMSO- d_6 . Yield: 70% (138 mg, 0.15 mmol). IR (ATR): $\nu = 3062$ (w, CH), 2022 (s, N_3), 1581 (m, CC/CN), 1468, 1390, 1259, 837, 786, 739 cm^{-1} . ^{13}C -NMR (solid-state, 100.68 MHz): $\delta = 155.2, 154.6, 152.8, 149.6, 145.2, 144.5, 142.6, 140.7, 136.0, 135.2, 134.1, 130.9, 129.0, 127.9, 126.4, 125.5, 123.7, 122.6, 121.3, 120.7, 119.3$ ppm. ^{15}N NMR (solid-state, 40.56 MHz): $\delta = -68.2, -71.6, -306.4$ ppm. ^{19}F -NMR (solid-state, 396.5 MHz): $\delta = -69.4$ (br, PF_6^-) ppm. ASAP (ESI $^+$): 777.1800 $\{[\text{PtN}_3\text{L}]\}^+$, 541.2138 $\{\text{L} + \text{H}\}^+$. $[\text{PtN}_3\text{L}]\cdot\text{Y}\cdot\text{H}_2\text{O}$ ($\text{Y}=\text{N}_3$ or PF_6^-) ($(\text{C}_{36}\text{H}_{24}\text{N}_9\text{Pt})_2\cdot\text{N}_3\cdot\text{PF}_6\cdot\text{H}_2\text{O}$): C, 49.12; H, 2.86; N, 16.71; found C 49.17, H 3.04, N 15.98.

4.2.3. Synthesis of triazolate complexes. 5: Complex 3 (0.09 mmol; 100 mg) was suspended in anhydrous acetone in a small round-bottomed flask (20 mL). The reaction mixture was then given 0.1 mmol (17 mg) of 4,4,4-trifluoro-2-butynoic acid ethyl ester. The reaction mixture was stirred until the solution became clear, which took 7 days. Then, ammonium hexafluorophosphate (6.13 mmol; 1 g) was added to the reaction mixture. Stirring was continuing for 6 h. A greenish-yellow precipitate was filtrated off, rinsed with water and diethyl ether, and then dried for a few days under vacuum. Yield: 62% (81 mg, 0.05 mmol). IR (ATR): $\nu = 3085$ (w, CH), 1727 (s, C=O), 1606 (m, CC/CN), 1440, 1312, 1248, 1135, 1028, 830, 784, 748, 724 cm^{-1} . ^1H NMR (d_6 -DMSO, 500.13 MHz) (broad spectrum): $\delta = 8.74, 8.63, 8.47, 8.34, 8.17, 7.81$, (aromatic signals), 4.39 (CH_2), 1.41 (CH_3) ppm. ^{13}C -NMR (d_6 -DMSO, 125.77 MHz): $\delta = 159.2$ (C=O), 156.4, 154.5, 152.4, 142.8, 138.7 (q, $^2J_{\text{C,F}} = 39.4$ Hz, C– CF_3), 137.6, 136.5, 129.1, 128.6, 127.8, 121.7 (q, $^1J_{\text{C,F}} = 267.0$ Hz, – CF_3), 121.4, 60.9 (– CH_2), 13.7 (– CH_3) ppm. ^{19}F NMR (d_6 -DMSO, 490.59 MHz): $\delta = -70.2$ (d, $^1J_{\text{PF}} = 711$ Hz, PF_6), –58.95 (s, CF_3 , major species, I) and –59.04 (s, CF_3 , minor species, II) ppm. ^{31}P NMR (d_6 -DMSO, 162.08 MHz): $\delta = -144.2$ (sept, $^1J_{\text{PF}} = 711$ Hz, PF_6) ppm. MS (ESI $^+$, DMSO/acetone): $m/z = 1315.0461$ $\{[\text{Pd}_2(\text{triazolato})_2\text{L}]\cdot\text{PF}_6\}^+$, 854.1431 $\{[\text{Pd}(\text{triazolato})\text{L}]\}^+$. $\text{C}_{48}\text{H}_{34}\text{F}_{18}\text{N}_{12}\text{O}_4\text{P}_2\text{Pd}_2\cdot 4\text{H}_2\text{O}$: C 37.64, H 2.76, N 10.97, found C 37.71, H 2.37, N 9.85.



6: To a small round flask charged with complex **4** (0.1 mmol; 100 mg) and acetone (10 mL), 4,4,4-trifluoro-2-butynoic acid ethyl ester (0.1 mmol; 17 mg) was added. The reaction mixture was stirred until the solution became clear, which took 7 d. Filtration was done and then one mL of a saturated solution of NH_4PF_6 was added. Stirring was continuing for 6 h. A brown precipitate was filtrated off, rinsed with water and diethyl ether, and then dried for a few days under vacuum. Yield: 53 (63 mg, 0.05 mmol). IR (ATR): $\nu = 3325, 2992$ (w, CH), 1728 (s, C=O), 1606 (s, CC), $1436, 1310, 1139, 1061, 829, 786$ cm^{-1} . ^{31}P NMR (d_6 -DMSO, 162.08 MHz): $\delta = -144.2$ (sept, $^1J_{\text{PF}_6} = 712$ Hz, PF_6) ppm. ^{19}F NMR (d_6 -DMSO, 490.59 MHz): $\delta = -70.1$ (sept, $^1J_{\text{PF}_6} = 699$ Hz, PF_6), $-59.89, -60.13$ and -60.26 ppm. MS (ESI⁺, DMSO/acetone): m/z 1110.2287 $\{\text{Na}[\text{Pt}(\text{triazolato})\text{L}]\cdot\text{PF}_6\}^+$. $\text{C}_{42}\text{H}_{29}\text{F}_3\text{N}_9\text{O}_2\text{Pt}\cdot 3\text{H}_2\text{O}$: C 44.14, H 3.09, N 11.03, found C 43.85, H 3.14, N 10.79.

4.3. Density functional theory calculations

The B3LYP^{31,32}/LANL2DZ^{33,34} approach was used to fully optimize the geometries of the triazolote isomers of **5** and **6** in the ground state. To get an insight into the local minimum structures of the triazolote isomers were verified by computing the vibrational modes at the same level of theory. Natural bond orbital analyses and TDDFT were performed using the previous mentioned DFT approach. All the calculations were carried out using Gaussian03.³⁸ Visualization of the optimized structures and frontier molecular orbitals were done with Gaussview03.³⁹

Data availability

Data is in ESI[†] file.

Author contributions

Ahmed Mansour: conceptualization, supervision, investigation, validation, resources, formal analysis, software, writing – original draft, writing – review & editing, resources. Krzysztof Radacki: methodology, investigation, resources. Gamal A. E. Mostafa, writing – review & editing, resources. Essam A. Ali, writing-review & editing, resources. Ola Shehab: investigation, writing – review & editing.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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