

## RESEARCH ARTICLE

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Cite this: *Inorg. Chem. Front.*, 2025, **12**, 3157

## Oxidation of phenols by the excited state of an osmium(vi) nitrido complex†

Yu-Zhong Lu,<sup>a</sup> Li-Xin Wang,<sup>a</sup> Rui-Yue Qi,<sup>a,b</sup> Jing Xiang,<sup>a</sup> Ji-Yan Liu<sup>\*a</sup> and Tai-Chu Lau<sup>\*a,c</sup>

The photoreaction of an osmium(vi) nitrido complex,  $[\text{Os}^{\text{VI}}(\text{N})(\text{L})(\text{CN})_3]^-$  (**OsN**), with various phenols has been investigated. Upon irradiation of **OsN** with visible light, the excited state (**OsN\***) is generated which reacts readily with a variety of phenols. **OsN\*** reacts with mono- and di-substituted phenols, including 2,6-dimethylphenol, 2,6-dichlorophenol and 4-methylphenol to afford the corresponding osmium(III) benzoquinone monoimine and osmium(IV) benzoquinone monoiminato complexes. On the other hand, in the reactions of **OsN\*** with bulky tri-substituted phenol such as 2,4,6-tri-*tert*-butylphenol, C–C bond cleavage occurred and  $[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{N}=\text{tBu}_2\text{Ph}_{(-2\text{H})}\text{O})]^-$  was formed as the major product. The electronic effects of various *para*-substituents (X) on the oxidation of phenols were investigated by the method of initial rates ( $R_x$ ). A Hammett plot of  $\log(R_x/R_{\text{H}})$  versus  $\sigma_p$  is linear with a  $\rho$  value of  $-0.54$ . A linear correlation of  $\log(R_x)$  with the oxidation potentials ( $E$ ) of phenols was also found with a slope of  $-0.80$ . On the other hand, no correlations were found between  $\log(R_x)$  and O–H bond dissociation energy (BDE), as well as the  $\text{p}K_{\text{a}}$  of phenols. The oxidation of phenols by **OsN\*** exhibits a negligible kinetic isotope effect (KIE),  $k(\text{C}_6\text{H}_5\text{OH})/k(\text{C}_6\text{D}_5\text{OD}) \sim 1$ . These results are consistent with a mechanism that involves an initial  $1\text{e}^-$  oxidation of the phenol followed by rapid proton transfer (ET-PT) to generate a phenoxy radical, this is followed by a N-rebound step to give the osmium products.

Received 7th December 2024,  
Accepted 26th February 2025

DOI: 10.1039/d4qi03144j

rsc.li/frontiers-inorganic

## Introduction

Transition metal nitrido ( $\text{M}\equiv\text{N}$ ) complexes are intermediates in  $\text{N}_2$  fixation and potentially useful reagents for the nitrogenation of organic substrates.<sup>1,2</sup> Compared to metal oxo ( $\text{M}=\text{O}$ ) species, there are relatively few nitrido complexes that are reactive towards common organic substrates. Examples include osmium(vi) nitrido complexes containing various N-based ligands such as 2,2'-bipyridine (bpy), 2,2':6',2''-terpyridine (tpy), and tris(1-pyrazolyl)methane (tpm); they are highly electrophilic and readily undergo N-atom transfer to a variety of organic substrates.<sup>3</sup> Moreover, Ru(vi) nitrido complexes bearing salen type ligands, such as  $[\text{Ru}^{\text{VI}}(\text{N})(\text{salchda})(\text{CH}_3\text{OH})]^+$  (**RuN**, *salchda* = *N,N'*-bis(salicylidene)-*o*-cyclohexyldiamine dianion),<sup>4</sup> and several iron(v/iv)<sup>5</sup> and Mn(v/vi)<sup>6</sup> nitrido

complexes are also highly reactive species that are capable of oxidizing various organic substrates.

The oxidation of phenols to quinones by various oxidants has been extensively studied, mainly because such reactions are relevant to many biological processes.<sup>7–13</sup> Phenols can be oxidized by various pathways (Fig. 1), the most common one is initial  $1\text{e}^-$  oxidation resulting in the formation phenoxy radicals, which is followed by rapid loss of the phenolic proton (consecutive electron transfer-proton transfer, ET-PT). Further rapid loss of  $1\text{e}^- + 1\text{H}^+$  results in the formation of quinones.<sup>14,15</sup> On the other hand, oxidation of phenols by some metal oxo species may involve an H-atom transfer mechanism (or concerted ET-PT).<sup>16–22</sup> Oxidation *via* an initial electrophilic attack on the aromatic ring of phenol has also been reported for a ruthenium(IV) species.<sup>23,24</sup>

Prior to this work, there was only one example on oxidation of phenols by a metal nitrido species. **RuN** readily reacts with phenols in the presence of pyridine to afford (*salchda*)ruthenium(II) *p*-benzoquinone imine complexes.<sup>25</sup> The reactions were proposed to proceed *via* an initial electrophilic attack by **RuN** at the aromatic ring of phenols.

In an attempt to design a highly active metal nitrido complex, we turned our attention to the excited state chemistry of  $\text{M}\equiv\text{N}$ . We reported recently the synthesis of a highly luminescent Os(vi) nitrido complex,  $[\text{Os}^{\text{VI}}(\text{N})(\text{L})(\text{CN})_3]^-$

<sup>a</sup>Key Laboratory of Optoelectronic Chemical Materials and Devices (Ministry of Education), School of Optoelectronic Materials and Technology, Jiangnan University, Wuhan, 430056, China. E-mail: xiangjing35991@sohu.com

<sup>b</sup>Hebei Key Laboratory of Heterocyclic Compounds, Handan University, Handan 056005, Hebei Province, China

<sup>c</sup>Department of Chemistry, City University of Hong Kong, Tat Chee Avenue, Kowloon Tong, Hong Kong, P. R. China. E-mail: bhtclau@cityu.edu.hk

† Electronic supplementary information (ESI) available. CCDC 2406139–2406142. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d4qi03144j>



## (a) Possible reaction pathways for phenol oxidation

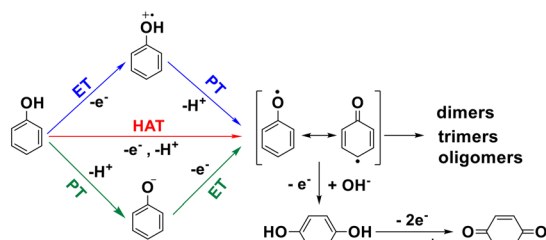
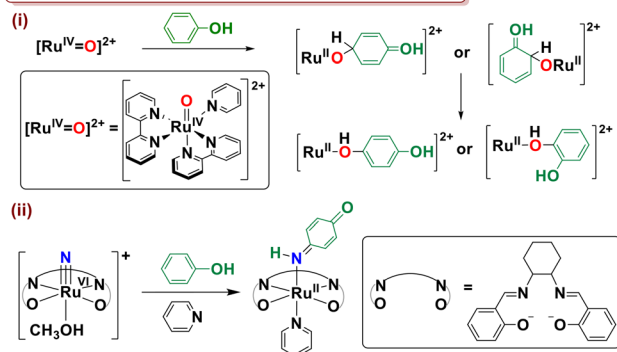
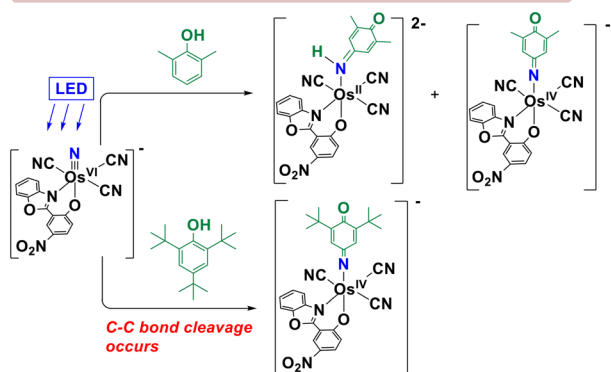
(b) Electrophilic attack by (i)  $\text{Ru}^{\text{IV}}\text{O}$  and (ii)  $\text{Ru}^{\text{VI}}\text{N}$ (c) This work: phenol oxidation by the excited state of  $\text{Os}^{\text{VI}}\text{N}$ 

Fig. 1 Oxidation pathways of phenols by oxidants.

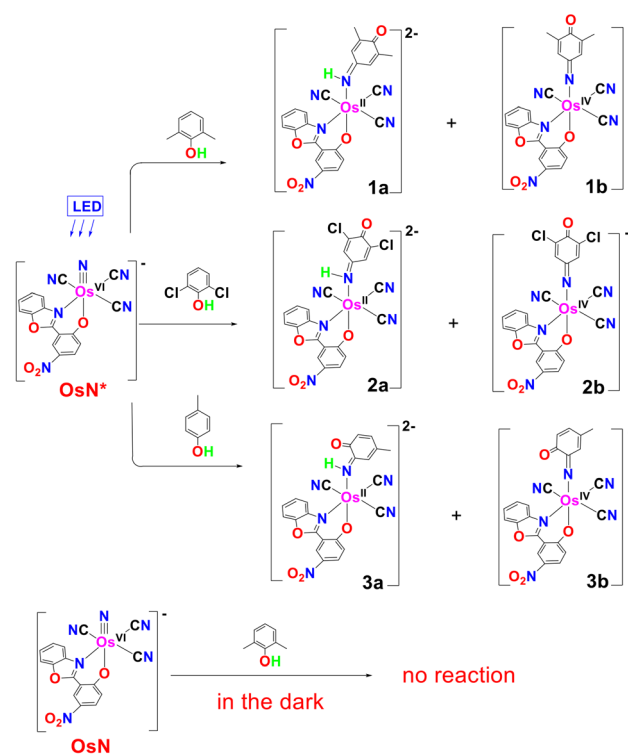
( $\text{OsN}$ ,  $\text{HL} = 2$ -(2-hydroxy-5-nitrophenyl)benzoxazole);<sup>26–33</sup>  $\text{OsN}$  is highly luminescent in both solid state and fluid solution (solid state:  $\lambda_{\text{em}} = 591$  nm,  $\phi = 11.7\%$ ,  $\tau = 1.90$   $\mu\text{s}$ ; in degassed  $\text{CH}_2\text{Cl}_2$ :  $\lambda_{\text{em}} = 594$  nm,  $\phi = 3.0\%$ ,  $\tau = 0.48$   $\mu\text{s}$ ). The CV of  $\text{OsN}$  shows an irreversible oxidation wave at  $E_{\text{pa}} = 1.88$  V and an irreversible reduction wave at  $E_{\text{pa}} = -0.99$  V vs. Saturated Calomel Electrode (SCE), which are tentatively assigned as the metal-centered  $\text{Os}^{\text{VII}/\text{VI}}$  and  $\text{Os}^{\text{VI}/\text{V}}$  process, respectively. The excited state ( $\text{OsN}^*$ ) of this complex is readily generated by visible light irradiation ( $\lambda > 460$  nm).  $\text{OsN}^*$  was found to be highly oxidizing and it reacts readily with various organic substrates, including alkanes, arenes, amines, alcohols, and dihydroxybenzenes.<sup>26–33</sup>

Herein, we reported the oxidation of phenols by  $\text{OsN}^*$ . These reactions have a number of unusual/novel features. In contrast to the oxidation of phenols by  $\text{RuN}$ , which occurs *via*

an electrophilic ring attack mechanism, oxidation by  $\text{OsN}^*$  occurs by an ET-PT followed by an N-rebound mechanism. Similar to oxidation by  $\text{RuN}$ , oxidation of various mono- and di-substituted phenols by  $\text{OsN}^*$  produced the corresponding osmium(II) benzoquinone imine complexes. However, because  $\text{OsN}^*$  is a powerful oxidant, further oxidation of the osmium(II) products by  $\text{OsN}^*$  occur to afford novel osmium(IV) iminato complexes as a second product. Moreover, oxidation of tri-substituted phenol such as 2,4,6- $\text{tBu}_3\text{C}_6\text{H}_2\text{OH}$  results in C–C bond cleavage of the substrate.

## Results and discussion

Upon irradiation with blue LED ( $\lambda > 460$  nm) for 24 h, the light-yellow  $\text{CH}_2\text{Cl}_2$  solution of  $\text{OsN}$  and 10 equiv. of 2,4,6- $\text{Me}_3\text{C}_6\text{H}_2\text{OH}$  rapidly turned dark red (Fig. 2). A mixture of the osmium(II) *p*-benzoquinone monoimine complex [ $\text{Os}^{\text{II}}(\text{L})(\text{CN})_3(\text{NH}=\text{Me}_2\text{Ph}_{(-2\text{H})}=\text{O})]^{2-}$  (**1a**) and osmium(IV) *p*-benzoquinone monoiminato complex [ $\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{N}=\text{Me}_2\text{Ph}_{(-2\text{H})}=\text{O})]^{2-}$  (**1b**), were isolated as  $\text{PPh}_4^+$  salts with 20% and 26% yields, respectively. Similarly, the photo-reaction of  $\text{OsN}$  with 2,6- $\text{Cl}_2\text{C}_6\text{H}_3\text{OH}$  afforded a mixture of ( $\text{PPh}_4$ )<sub>2</sub>[ $\text{Os}^{\text{II}}(\text{L})(\text{CN})_3(\text{NH}=\text{Cl}_2\text{Ph}_{(-2\text{H})}=\text{O})]^{2-}$  [( $\text{PPh}_4$ )<sub>2</sub>**2a**] and ( $\text{PPh}_4$ )<sub>2</sub>[ $\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{N}=\text{Cl}_2\text{Ph}_{(-2\text{H})}=\text{O})]^{2-}$  [( $\text{PPh}_4$ )<sub>2</sub>**2b**] with 32% and 25% yields, respectively. In addition, ( $\text{PPh}_4$ )<sub>2</sub>[ $\text{Os}^{\text{II}}(\text{L})(\text{CN})_3(\text{NH}_3)]^{2-}$  ( $\text{OsNH}_3$ )<sup>28</sup> was isolated with ~10% yield in reactions with these phenols. Controlled experiments showed that

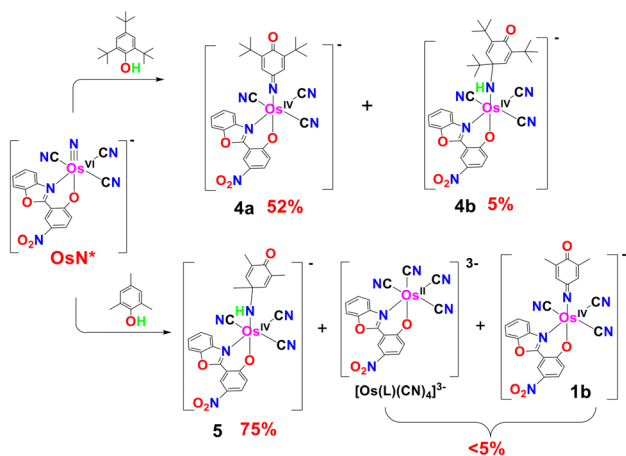
Fig. 2 Reactions of  $\text{OsN}$  with various phenols in the excited state and ground state.

no reaction between **OsN** and phenols was observed in the dark.

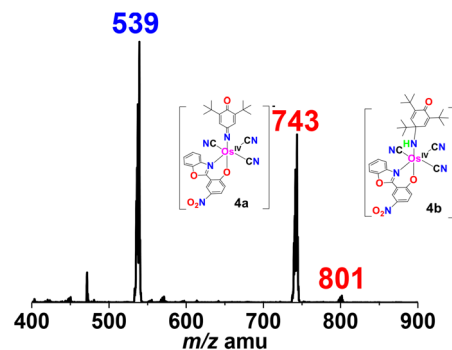
On the other hand, in the photoreaction with 4-methylphenol, **OsN** undergoes electrophilic attack at the *ortho* position to afford an  $[\text{PPh}_4]_2[\text{Os}^{\text{II}}(\text{L})(\text{CN})_3(o\text{-NH}=\text{MePh}_{(-2\text{H})}=\text{O})]$  [**(PPh<sub>4</sub>)<sub>2</sub>3a**] and  $[\text{PPh}_4][\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(o\text{-N}=\text{MePh}_{(-2\text{H})}=\text{O})]$  [**(PPh<sub>4</sub>)<sub>3</sub>b**], with yields of 37% and 32%, respectively.

The photoreactions of **OsN** with 2,4,6-tri-*tert*-butylphenol and 2,4,6-tri-methylphenol were also investigated (Fig. 3); for these substrates with three alkyl substituents, direct electrophilic attack on the aromatic ring by **OsN**\* may be inhibited. Reaction of **OsN**\* with 2,4,6-*t*-Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH afforded  $[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(p\text{-N}=\text{tBu}_2\text{Ph}_{(-2\text{H})}=\text{O})]^-$  (**4a**), isolated as  $\text{PPh}_4^+$  salt with ~52% yield. The structure of **4a** (see characterization below) reveals formal C–C bond cleavage of 2,4,6-*t*-Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH; GC/MS and GC show that the 2-methylpropene was formed with ~50% yield. <sup>1</sup>H NMR spectrum of (**PPh<sub>4</sub>**)**4a** shows that the remaining two *t*-Bu groups are symmetry-related, consistent with the *para* attack of 2,4,6-*t*-Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH by **OsN**\*. ESI/MS (–ve mode) of the photoreaction solution of **OsN** and 2,4,6-*t*-Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH shows a predominant peak at *m/z* 743 (Fig. 4), attributed to  $[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(p\text{-N}=\text{tBu}_2\text{Ph}_{(-2\text{H})}=\text{O})]^-$  (**4a**). There is also a very minor peak at *m/z* 801, (estimated to be 5%), which is tentatively assigned to  $[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{NH}-\text{tBu}_3\text{Ph}_{(-2\text{H})}=\text{O})]^-$  (**4b**), arising from an electrophilic attack of **OsN** at the *para* position of the phenol.

In the case of 2,4,6-Me<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH with less bulky methyl substituents, electrophilic ring attack by **OsN**\* occurred predominantly at the *para* position, and the amido complex,  $[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{NH}(\text{Me}_3\text{Ph}_{(-2\text{H})}=\text{O}))]^-$  (**5**), was isolated as  $\text{PPh}_4^+$  salt with ~75% yield. The ESI/MS (–ve mode) of the product solution shows a predominant peak at *m/z* = 675 due to complex **5**. In addition, there are two minor peaks at *m/z* 659.1 and 275.6, which are tentatively assigned to **1b** and  $[\text{Os}^{\text{II}}(\text{L})(\text{CN})_4]^{3-}$ , respectively. These two species should arise from C–C bond cleavage of 2,4,6-Me<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH; however, the total yields of these two products are estimated to be <5%.



**Fig. 3** Photoreaction of **OsN** with 2,4,6-*t*-Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH and 2,4,6-Me<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH.



**Fig. 4** ESI/MS of photoreaction of **OsN** with 10 equiv. 2,4,6-*t*-Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH for 4 h showing a predominant peak at *m/z* 743 and a small peak at *m/z* 801.

All products have been characterized by IR, UV/vis, CV, ESI/MS, and <sup>1</sup>H NMR (Fig. S1–S12†). All the Os(II) and Os(IV) compounds are diamagnetic, as evidenced by the sharp resonances found in the normal range in their <sup>1</sup>H NMR spectra. The diamagnetism of these compounds is consistent with the low spin d<sup>6</sup> and d<sup>4</sup> electronic configurations for Os(II) and Os(IV), respectively. The IR spectra of (**PPh<sub>4</sub>**)**2a** and (**PPh<sub>4</sub>**)**2a** show  $\nu(\text{C}\equiv\text{N})$  stretches at 2096, 2070 cm<sup>–1</sup> and 2108, 2082 cm<sup>–1</sup>, and  $\nu(\text{C}=\text{O})$  stretches at 1609 cm<sup>–1</sup> and 1610 cm<sup>–1</sup>, respectively. The IR spectra of (**PPh<sub>4</sub>**)**1b** and (**PPh<sub>4</sub>**)**2b** show  $\nu(\text{C}\equiv\text{N})$  stretches at 2145, 2134 cm<sup>–1</sup> and 2153, 2142 cm<sup>–1</sup>, and  $\nu(\text{C}=\text{O})$  stretches at 1629 cm<sup>–1</sup> and 1639 cm<sup>–1</sup>, respectively which are at higher wavenumbers as compared with (**PPh<sub>4</sub>**)**2a** and (**PPh<sub>4</sub>**)**2a**. Similar  $\nu(\text{C}\equiv\text{N})$  and  $\nu(\text{C}=\text{O})$  stretches are also found in (**PPh<sub>4</sub>**)**2a** and (**PPh<sub>4</sub>**)**3b**.

As shown in Fig. S6,† the UV/vis spectra of these compounds show ligand-centered  $\pi\text{--}\pi^*$  transitions in the UV regions. In the osmium(II) products (**PPh<sub>4</sub>**)**2a**, (**PPh<sub>4</sub>**)**2a**, and (**PPh<sub>4</sub>**)**2a**, there are strong absorption bands at 500–650 nm with molar extinction coefficients ( $\epsilon$ ) of the order of 10<sup>4</sup> M<sup>–1</sup> cm<sup>–1</sup>, which are assigned to Os(II) to *p*-benzoquinone monoiimine charge transfer (MLCT) transitions. Notably, there is an intense absorption band with  $\epsilon > 6 \times 10^4$  M<sup>–1</sup> cm<sup>–1</sup> in (**PPh<sub>4</sub>**)**2a**. In the osmium(IV) products (**PPh<sub>4</sub>**)**1b**, (**PPh<sub>4</sub>**)**2b** and (**PPh<sub>4</sub>**)**3b**, the broad absorption bands at ~450 nm are probably due to LMCT transitions.

The cyclic voltammograms (CVs) of the osmium complexes (**PPh<sub>4</sub>**)**2a**, (**PPh<sub>4</sub>**)**2a**, and (**PPh<sub>4</sub>**)**2a** in CH<sub>3</sub>CN exhibit a reversible/quasi-reversible Os<sup>III/II</sup> couples at –0.67 V, –0.38 V and –0.78 V vs. Fc<sup>+/0</sup>, respectively (Fig. 5). There is also an irreversible wave at  $E_{\text{pc}} = -1.46$  V, –1.35 V and –1.72 V, respectively which are tentatively assigned to the reduction of the benzoquinone imine ligands. The oxidation waves at  $E_{1/2} = 1.25$  V,  $E_{\text{pc}} = 1.46$  V and 1.23 V, respectively, are assigned to Os<sup>IV/III</sup>. The CVs of the osmium(IV) products **1b**, **2b** and **3b** exhibit irreversible Os<sup>IV/III</sup> waves at  $E_{\text{p}} = -0.65$  V, –0.39 V and –0.66 V, respectively (Fig. 6); while the waves at  $E_{\text{p}} = -1.61$  V, –1.35 V and –1.49 V, respectively are tentatively assigned to the reduction of benzoquinone imine ligands.

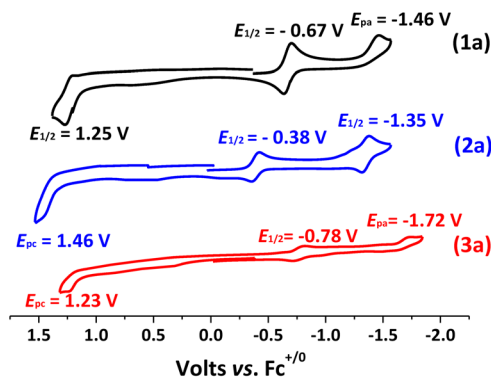


Fig. 5 CVs for **1a**, **2a**, and **3a** in MeCN solutions containing 0.1 M  $[t\text{Bu}_4\text{N}](\text{PF}_6)$  with a scan rate =  $0.1 \text{ V s}^{-1}$ .

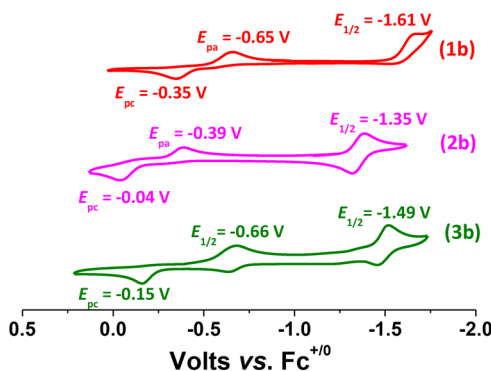


Fig. 6 CVs for **1b**, **2b**, and **3b** in MeCN solutions containing 0.1 M  $[t\text{Bu}_4\text{N}](\text{PF}_6)$  with a scan rate =  $0.1 \text{ V s}^{-1}$ .

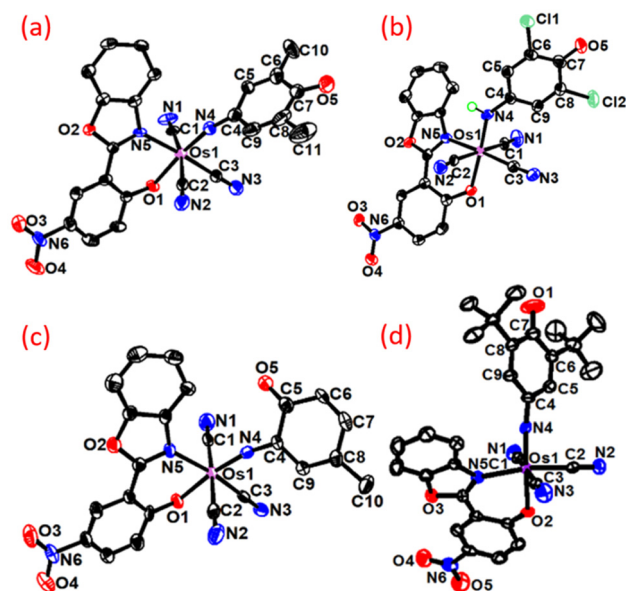


Fig. 7 The structures of (a) **1b**, (b) **2a**, (c) **3b** and (d) **4a**.

Table 1 Selected bond parameters ( $\text{\AA}$ ,  $^\circ$ ) for **1b**, **2a**, **3b** and **4a**

	<b>1b</b>	<b>2a</b>	<b>3b</b>	<b>4a</b>
Os1–C1	2.074 (5)	2.053 (4)	2.082 (5)	2.081(5)
Os1–C2	2.077 (5)	2.070 (4)	2.081 (5)	2.029(4)
Os1–C3	2.025 (5)	2.009 (4)	2.034 (6)	2.066(5)
Os1–N4	1.781 (4)	1.927 (3)	1.768 (4)	1.779(4)
Os1–N5	2.117 (4)	2.129 (3)	2.118 (4)	2.115(3)
Os1–O1	2.010 (3)	2.066 (3)	2.028 (3)	—
Os1–O2	—	—	—	2.022(3)
Os1–N4–C4	167.2 (4)	136.6 (3)	167.9 (5)	163.8(3)

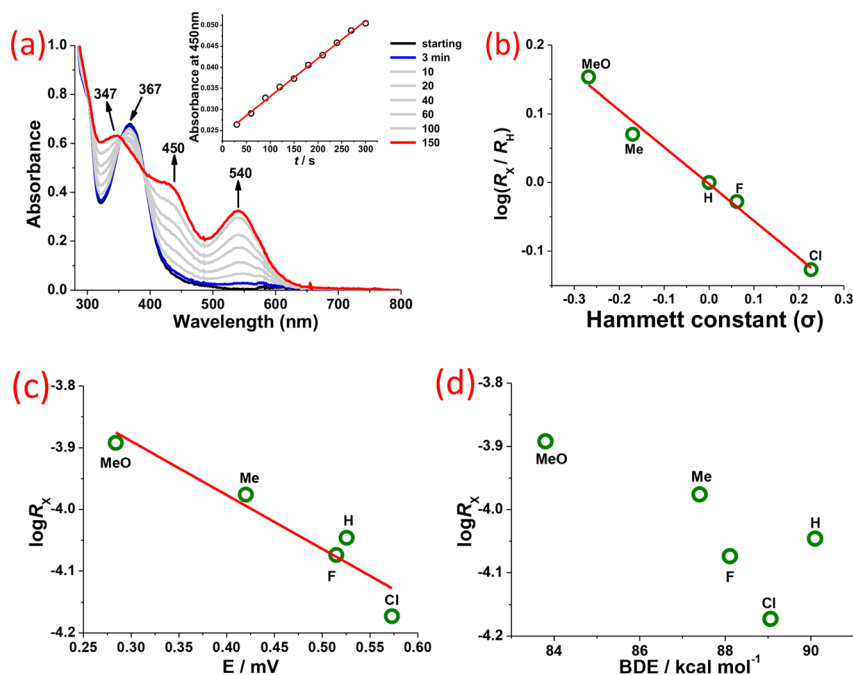


Fig. 8 (a) UV/vis spectral changes for the photoreaction of **OsN** ( $2.5 \times 10^{-5} \text{ M}$ ) with phenol ( $1.8 \times 10^{-3} \text{ M}$ ) in  $\text{C}_2\text{H}_4\text{Cl}_2$ . Inset shows the time-trace absorbance at 450 nm. (b) Hammett plot for the photoreaction of **OsN** with 4-substituted phenols in  $\text{C}_2\text{H}_4\text{Cl}_2$ . Slope =  $-0.54$ . Intercept =  $-0.00217$ . (c) Plot of  $\log(R_x)$  vs.  $E^\circ$ . (d) Plot of  $\log(R_x)$  vs. BDE.





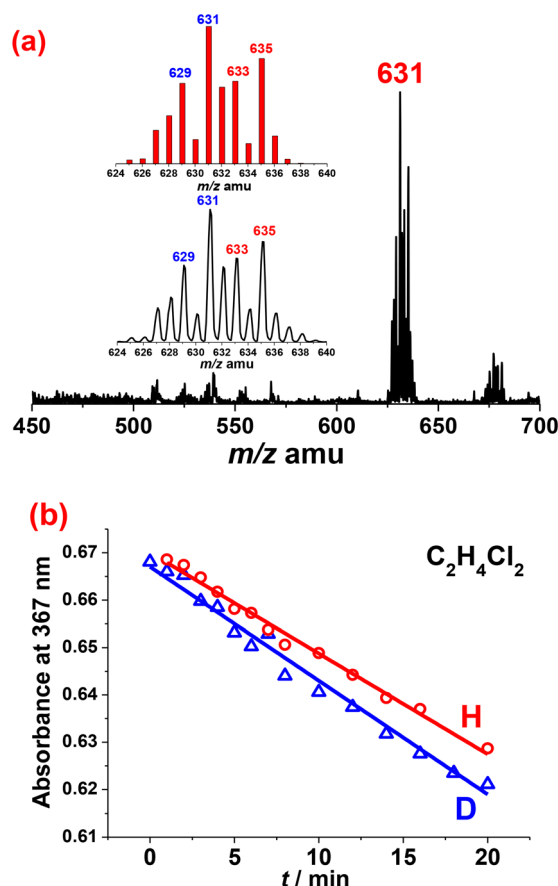
The molecular structures of **(PPh<sub>4</sub>)<sub>1</sub>b**, **(PPh<sub>4</sub>)<sub>2</sub>2a**, **(PPh<sub>4</sub>)<sub>3</sub>3b**, and **(PPh<sub>4</sub>)<sub>4</sub>4a** have been determined by X-ray crystallography (Fig. 7). Selected bond parameters are summarized in Table 1. The *mer*-configuration of **OsN** is retained in these complexes. There are two PPh<sub>4</sub><sup>+</sup> in **2a** and only one PPh<sub>4</sub><sup>+</sup> in **1b** and **3b**. The Os–N4 bond length in **2a** is 1.927(3) Å, which is significantly shorter than related Os–N bond lengths in [Os<sup>III</sup>{N(H)C(NH<sub>2</sub>)}(L<sup>1</sup>)(CN)<sub>3</sub>]<sup>–</sup> and [Os<sup>III</sup>{N(H)CN}(L<sup>1</sup>)(CN)<sub>3</sub>]<sup>2–</sup> (HL<sup>1</sup> = 2-(2-hydroxyphenyl)benzoxazole),<sup>32,33</sup> indicating the presence of strong  $\pi$  back-bonding interaction between Os<sup>II</sup> and the benzoquinone monoimine ligand. On the other hand, the Os–N4 bond lengths are 1.781(4) and 1.768(4) Å, respectively for **1b** and **3b**, which are much shorter than that in **2a**, indicating Os–N double bond character in these Os(IV) complexes. This is also evidenced by the more linear Os1–N4–C4 bond angles of 167.2(4)° and 167.9(5)° than that in **2a** (136.6(3)°). The bond parameters of **3b** and **4a** are essentially identical to those in **1b**, as the metal centers have the same oxidation state.

### Substituent effects

The photoreactions of **OsN** with various 4-substituted phenols (4-X-C<sub>6</sub>H<sub>4</sub>OH; X = MeO, Me, H, F, Cl) were also investigated by UV/vis spectroscopy (Fig. S10†). The initial rates ( $R_x$ ) of these reactions were obtained at 450 nm (Fig. 8a).  $R_x$  was found to increase with increasing electron donating properties of the substituents: MeO > Me > H > F > Cl. A linear correlation was obtained in the Hammett plot of  $\log(R_x/R_H)$  versus the substituent constant ( $\sigma$ ), with a  $\rho$  value of  $-0.54 \pm 0.01$  (Fig. 8b), indicating that the phenol center is more positive in the transition state.  $\log(R_x)$  also correlates well with the oxidation potentials ( $E^{\text{ox}}$ ) of these phenols with a slope of  $-0.80$  (Fig. 8c).<sup>34</sup> The results indicate that the reaction may proceed *via* an initial rate-limiting 1e<sup>–</sup> transfer (ET) from phenols to **OsN**<sup>\*</sup>. No clear correlation between  $\log(R_x)$  and pK<sub>a</sub> of phenols, suggesting that proton transfer (PT) is not involved in the rate-limiting step. There is also no clear relationship between  $\log(R_x)$  and the O–H bond dissociation energies (BDEs) of the phenols (Fig. 8d),<sup>35</sup> which does not support a hydrogen atom transfer (HAT) mechanism for the oxidation of phenols by **OsN**<sup>\*</sup>.

### Kinetic isotope effects (KIE)

The KIE for the reaction of **OsN**<sup>\*</sup> with phenol was determined by ESI/MS, using an equimolar mixture of phenol (C<sub>6</sub>H<sub>5</sub>OH) and d<sup>6</sup>-phenol (C<sub>6</sub>D<sub>5</sub>OD) as substrate. As shown in Fig. 9a, the KIE value was estimated to be  $\sim(1 \pm 0.05)$  from the ratios of the most intense peaks for two Os(IV) benzoquinone monoiminato products, ( $m/z = 631$  and  $635$  for [Os<sup>IV</sup>(L)(CN)<sub>3</sub>(N=C<sub>6</sub>H<sub>4</sub>=O)]<sup>–</sup> and [Os<sup>IV</sup>(L)(CN)<sub>3</sub>(N=C<sub>6</sub>D<sub>4</sub>=O)]<sup>–</sup> respectively), assuming that the spraying and ionization efficiencies of the two ions are similar. The isotopic distribution pattern obtained is also in agreement with the simulated one. The KIE is also determined by UV/vis spectroscopy. As shown in Fig. S11,† the UV/vis spectral changes were obtained from the photoreaction of **OsN** with phenol (C<sub>6</sub>H<sub>5</sub>OH) and d<sup>6</sup>-phenol (C<sub>6</sub>D<sub>5</sub>OD) under the same conditions. The reactions were followed by the change in absorbance at 367 nm, which gives a KIE of  $R_H/R_D = 0.97 \pm$



**Fig. 9** (a) ESI/MS for the photoreaction of **OsN** with equimolar of phenol and d<sup>6</sup>-phenol showing a predominant peak around  $m/z$  631. Inserts show the expanded isotopic distribution of peak  $m/z$  631, which agrees with the simulated isotopic distribution of  $m/z$  631 and 635 with a mole ratio of 0.505 : 0.495 (KIE  $\sim 1$ ). (b) Time trace for absorbance at 367 nm of **OsN** ( $2.5 \times 10^{-5}$  M) with 100 equiv. of phenol (red line) and d<sup>6</sup>-phenol (blue line) in C<sub>2</sub>H<sub>4</sub>Cl<sub>2</sub>. The linear fitting gives initial rates of  $R_H = (-2.13 \pm 0.06) \times 10^{-3}$ ,  $r^2 = 0.98$  and  $R_D = (-2.40 \pm 0.10) \times 10^{-3}$ ,  $r^2 = 0.99$ ;  $R_H : R_D = 0.97$ .

0.01 (Fig. 9b). Based on the above results, the photoreaction of **OsN** with phenol exhibits negligible KIE, indicating that C–H bond cleavage is not involved in the rate-limiting step, in line with the conclusion obtained from investigation of substituent effects.

### Proposed mechanism

Based on the above experiments, the proposed mechanism for the photoreaction of **OsN** with phenols is illustrated in Fig. 10 (using 2,6-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>OH as an example). The initial step is rate-limiting 1e<sup>–</sup> oxidation of 2,6-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>OH (ET), followed by rapid proton transfer (PT) to give the **Os<sup>V</sup>NH** and phenoxy radical, which is supported by a linear correlation between  $\log(R_x)$  and  $E$  of the phenols with a slope of  $-0.80$ . The proposed mechanism is also supported by the linear Hammett correlation with  $\rho$  value of  $-0.54$  and a negligible KIE effect ( $k(\text{C}_6\text{H}_5\text{OH})/k(\text{C}_6\text{D}_5\text{OD}) \sim 1.03$ ). Moreover, the initial rate ( $R$ ) for phenolate is 60 times faster than that of phenol, further vali-



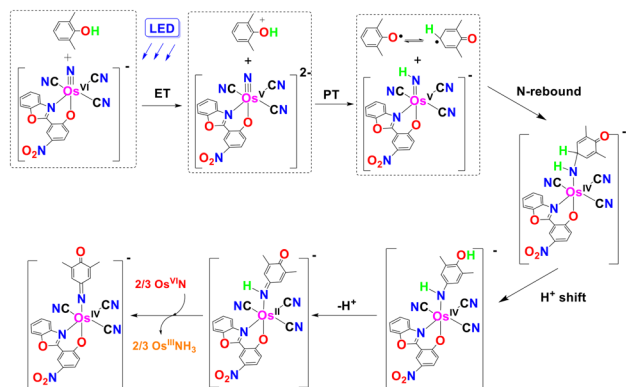


Fig. 10 The proposed reaction mechanism for the reaction of  $\text{OsN}^*$  with 2,6- $\text{Me}_2\text{C}_6\text{H}_3\text{OH}$ .

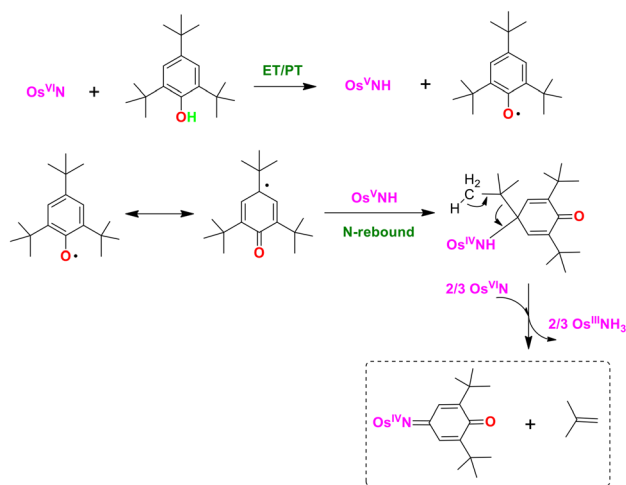


Fig. 11 The proposed mechanism for the C–C bond cleavage of 2,4,6- $\text{t-Bu}_3\text{C}_6\text{H}_2\text{OH}$  by  $\text{OsN}^*$ .

dating the initial ET mechanism (Fig. S12†). This is followed by tautomerism and N-rebound to give an  $\text{Os}^{\text{IV}}$  amido species. An internal  $2e^-$  redox results in the  $\text{Os}^{\text{II}}$  hydroquinone monoimine product. The other product,  $\text{Os}^{\text{IV}}$  hydroquinone monoiminato complex, was formed by further  $2e^-$  oxidation by  $\text{OsN}^*$  ( $2/3\text{OsN}^* + \mathbf{1a} \rightarrow \mathbf{1b} + 2/3\text{OsNH}_3$ ). This last step is supported by the isolation of  $\text{OsNH}_3$  in 10% yield.

For the photoreaction of  $\text{OsN}$  and 2,4,6- $\text{t-Bu}_3\text{C}_6\text{H}_2\text{OH}$ , the first four steps are similar (Fig. 11), *i.e.*  $1e^-$  oxidation of 2,4,6- $\text{t-Bu}_3\text{C}_6\text{H}_2\text{OH}$  (ET), followed by rapid proton transfer (PT) to give the  $\text{Os}^{\text{V}}\text{NH}$  and phenoxyl radical; followed by tautomerism and N-rebound to give an  $\text{Os}^{\text{IV}}$  amido species. Further  $2e^-$  oxidation of this  $\text{Os}^{\text{IV}}$  amido species by  $\text{OsN}^*$  leads to C–C bond cleavage with the formation of **4a**, 2-methylpropene, and  $\text{OsNH}_3$ .

## Conclusion

In conclusion, we have demonstrated novel reactivity of the excited state of an osmium(vi) nitrido complex towards

phenols. The photoreactions of  $\text{OsN}$  with the parent phenol, as well as mono- and di-substituted phenols afforded  $\text{Os}^{\text{II}}$  *p*-benzoquinone monoimine and osmium(IV) *p*-benzoquinone monoiminato complexes. On the other hand, oxidation of bulky tri-substituted phenol such as 2,4,6- $\text{t-Bu}_3\text{C}_6\text{H}_2\text{OH}$  resulted in C–C bond cleavage of the substrate. Mechanistic studies indicate that the photoreactions proceed *via* an initial  $1e^-$  oxidation followed by a rapid proton transfer (ET-PT) to generate phenoxyl radicals, this is followed by a N-rebound step to give the osmium products. We believe that our work is a significant contribution to  $\text{M}\equiv\text{N}$  excited state as well as phenol oxidation chemistry.

## Experimental

$(\text{PPh}_4)_2[\text{Os}^{\text{II}}(\text{L})(\text{CN})_3(\text{NH}=\text{Me}_2\text{PhOH}_{(-2\text{H})})]$  [( $\text{PPh}_4$ )<sub>2</sub>**1a**] and  $(\text{PPh}_4)[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{N}=\text{Me}_2\text{PhOH}_{(-2\text{H})})]$  [( $\text{PPh}_4$ )**1b**]

10 tubes each containing  $\text{OsN}$  (5 mg, 5.7  $\mu\text{mol}$ ) and 2,6- $\text{Me}_2\text{C}_6\text{H}_3\text{OH}$  (73 mg, 0.6 mmol) in  $\text{CH}_2\text{Cl}_2$  under argon were irradiated with blue LED light for 24 h, whereby the light-yellow solutions turned dark red. The solutions were combined, and the solvent was removed under reduced pressure. The residue was washed with diethyl ether (100 ml) to remove the unreacted 2,6- $\text{Me}_2\text{C}_6\text{H}_3\text{OH}$ . The residue was dissolved in a minimum amount of  $\text{CH}_2\text{Cl}_2$  and then loaded onto a silica gel column. The first yellow band  $(\text{PPh}_4)[\text{Os}^{\text{IV}}(\text{L})(\text{CN})_3(\text{N}=\text{Me}_2\text{Ph}_{(-2\text{H})}=\text{O})]$  [( $\text{PPh}_4$ )**1b**] was eluted by  $\text{CH}_2\text{Cl}_2$ /acetone (v : v, 4 : 1). Yellow needle crystals were obtained from slow diffusion of diethyl ether into a MeCN solution of ( $\text{PPh}_4$ )**1b**. The second blue band was eluted by  $\text{CH}_2\text{Cl}_2$ /acetone/MeOH (v : v : v, 8 : 2 : 1). The solvent was removed under reduced pressure. The solid obtained was dissolved in  $\text{H}_2\text{O}$  (10 ml) and excess  $\text{PPh}_4\text{Cl}$  (30 mg) was added to give the blue precipitate. The crude product was further purified by slow diffusion of diethyl ether into a  $\text{CH}_2\text{Cl}_2$  solution of ( $\text{PPh}_4$ )<sub>2</sub>**1a**. Yield for ( $\text{PPh}_4$ )<sub>2</sub>**1a**: 15 mg, 20% (based on the  $\text{OsN}$  consumed). Selected IR (KBr disc,  $\text{cm}^{-1}$ ):  $\nu(\text{N-H})$  3248;  $\nu(\text{C}\equiv\text{N})$  2096 and 2070;  $\nu(\text{C}=\text{O})$  1645;  $\nu(\text{C}=\text{N})$  1609;  $\nu(\text{N}=\text{O})$  1303.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ ):  $\delta$  8.92 (s, 1H, Ar-H), 7.92–7.78 (m, 8H, Ar-H), 7.77–7.64 (m, 18H, Ar-H), 7.62–7.48 (m, 18H, Ar-H), 7.38 (d,  $J = 8.0$  Hz, 1H, Ar-H), 7.29 (m, 1H, Ar-H), 7.19 (t,  $J = 8.0$  Hz, 1H, Ar-H), 6.67 (s, 1H, Ar-H), 1.69 (s, 6H). ESI/MS (–ve mode) in MeOH: at  $m/z$  329.7 ( $[\text{M}]^{2-}$ ); Anal. Calcd for  $\text{C}_{72}\text{H}_{56}\text{N}_6\text{O}_5\text{P}_2\text{Os}$ : C, 64.66; H, 4.22; N, 6.28. Found: C, 64.80; H, 4.25; N, 6.32%. UV/Vis ( $\text{CH}_2\text{Cl}_2$ ):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [ $\text{mol}^{-1} \text{dm}^3 \text{cm}^{-1}$ ]): 269 (22 472), 276 (21 630), 287 (16 020), 344 (14 050), 377sh (11 870), 432 (11 270), 541 (17 340), 586 (17 670). Yield for ( $\text{PPh}_4$ )**1b**: 15 mg, 26% (based on the  $\text{OsN}$  consumed). Selected IR (KBr disc,  $\text{cm}^{-1}$ ):  $\nu(\text{C}\equiv\text{N})$  2145 and 2135;  $\nu(\text{C}=\text{O})$  1630;  $\nu(\text{C}=\text{N})$  1611.  $^1\text{H}$  NMR (400 MHz,  $\text{CDCl}_3$ ):  $\delta$  9.04 (s, 1H, Ar-H), 8.19 (s, 2H, Ar-H), 8.07 (d,  $J = 9.5$  Hz, 1H, Ar-H), 7.96–7.88 (m, 4H, Ar-H), 7.84–7.74 (m, 9H, Ar-H), 7.72 (d,  $J = 7.4$  Hz, 1H, Ar-H), 7.69–7.61 (m, 8H, Ar-H), 7.53 (t,  $J = 8.3$  Hz, 2H, Ar-H), 6.92 (d,  $J = 9.2$  Hz, 1H, Ar-H), 3.15 (s, 6H,  $-\text{CH}_3$ ). ESI-MS (–ve mode) in MeOH:  $m/z$  659 ( $\text{M}^-$ ); Anal. Calcd for  $\text{C}_{48}\text{H}_{35}\text{N}_6\text{O}_5\text{OsP}$ : C, 57.82; H, 3.54; N, 8.43. Found: C, 57.93; H, 3.60; N, 8.31%. UV/Vis ( $\text{CH}_2\text{Cl}_2$ ):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [ $\text{mol}^{-1}$



$\text{dm}^3 \text{cm}^{-1}$ ): 269 (23 710), 277 (25 070), 288 (23 480), 330 (22 610), 432 (33 300).

**(PPh<sub>4</sub>)<sub>2</sub>[Os<sup>II</sup>(L)(CN)<sub>3</sub>(NH=Cl<sub>2</sub>PhOH<sub>(-2H)</sub>)] [(PPh<sub>4</sub>)<sub>2</sub>2a] and (PPh<sub>4</sub>)<sub>2</sub>[Os<sup>IV</sup>(L)(CN)<sub>3</sub>(N=Cl<sub>2</sub>PhOH<sub>(-2H)</sub>)] [(PPh<sub>4</sub>)<sub>2</sub>2b]**

The synthesis and isolation of (PPh<sub>4</sub>)<sub>2</sub>2a and (PPh<sub>4</sub>)<sub>2</sub>2b are similar to that of (PPh<sub>4</sub>)<sub>2</sub>1a and (PPh<sub>4</sub>)<sub>2</sub>1b except that 2,6-Cl<sub>2</sub>C<sub>6</sub>H<sub>3</sub>OH was used instead of 2,6-dimethylphenol. Yield for (PPh<sub>4</sub>)<sub>2</sub>2a: (25 mg, 32%, based on the OsN consumed). Selected IR (KBr disc,  $\text{cm}^{-1}$ ):  $\nu(\text{N-H})$  3235;  $\nu(\text{C}\equiv\text{N})$  2108 and 2082;  $\nu(\text{C=O})$  1638;  $\nu(\text{C=N})$  1609;  $\nu(\text{N=O})$  1307; <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  8.96 (d,  $J$  = 2.9 Hz, 1H), 7.90–7.82 (m, 8H), 7.80 (d,  $J$  = 2.9 Hz, 1H), 7.77–7.68 (m, 16 H), 7.65–7.56 (m, 16H), 7.55–7.48 (m, 2H), 7.37–7.29 (m, 3H), 7.23 (s, 1H); 6.78 (d,  $J$  = 9.4 Hz, 1H). ESI-MS (–ve mode) in MeOH:  $m/z$  350.2 [M]<sup>2–</sup>. Anal. Calcd for C<sub>70</sub>H<sub>50</sub>Cl<sub>2</sub>N<sub>6</sub>O<sub>5</sub>OsP<sub>2</sub>: C, 61.00; H, 3.66; N, 6.10. Found: C, 61.10; H, 3.71; N, 6.18%. UV/Vis (CH<sub>2</sub>Cl<sub>2</sub>):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [mol<sup>–1</sup> dm<sup>3</sup> cm<sup>–1</sup>]): 262sh (18 090), 269 (19 810), 276 (18 740), 285sh (12 760), 338 (12 110), 363sh (9360), 417sh (7070), 593 (60 850). Yield for (PPh<sub>4</sub>)<sub>2</sub>2b: 15 mg, 25% (based on the OsN consumed). Selected IR (KBr disc,  $\text{cm}^{-1}$ ):  $\nu(\text{C}\equiv\text{N})$  2153 and 2141;  $\nu(\text{C=O})$  1639;  $\nu(\text{C=N})$  1614. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  9.08 (s, 1H, Ar–H), 8.50 (s, 2H, Ar–H), 8.21 (d,  $J$  = 8.9 Hz, 1H, Ar–H), 7.97–7.89 (m, 4H, Ar–H), 7.84–7.75 (m, 9H, Ar–H), 7.72–7.58 (m, 11H, Ar–H), 7.13 (d,  $J$  = 9.2 Hz, 1H, Ar–H). ESI-MS (–ve mode) in MeOH:  $m/z$  699 [M]<sup>–</sup>; Anal. Calcd for C<sub>46</sub>H<sub>29</sub>Cl<sub>2</sub>N<sub>6</sub>O<sub>5</sub>OsP<sub>2</sub>: C, 53.23; H, 2.82; N, 8.10. Found: C, 53.29; H, 2.76; N, 8.21%. UV/Vis (CH<sub>2</sub>Cl<sub>2</sub>):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [mol<sup>–1</sup> dm<sup>3</sup> cm<sup>–1</sup>]): 270 (40 050), 277 (44 140), 288sh (42 330), 331sh (37 880), 447 (60 490).

**(PPh<sub>4</sub>)<sub>2</sub>[Os<sup>II</sup>(L)(CN)<sub>3</sub>(NH=MePhOH<sub>(-2H)</sub>)] [(PPh<sub>4</sub>)<sub>2</sub>3a] and (PPh<sub>4</sub>)<sub>2</sub>[Os<sup>IV</sup>(L)(CN)<sub>3</sub>(N=MePhOH<sub>(-2H)</sub>)] [(PPh<sub>4</sub>)<sub>2</sub>3b]**

The synthesis and isolation of (PPh<sub>4</sub>)<sub>2</sub>3a and (PPh<sub>4</sub>)<sub>2</sub>3b are similar to that of (PPh<sub>4</sub>)<sub>2</sub>1a and (PPh<sub>4</sub>)<sub>2</sub>1b except that 4-MeC<sub>6</sub>H<sub>4</sub>OH was used instead of 2,6-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>OH. Yield for (PPh<sub>4</sub>)<sub>2</sub>3a, 28 mg, 37%. Selected IR (KBr disc,  $\text{cm}^{-1}$ ) for (PPh<sub>4</sub>)<sub>2</sub>3a:  $\nu(\text{N-H})$  3253;  $\nu(\text{C}\equiv\text{N})$  2125 and 2086;  $\nu(\text{C=O})$  1645;  $\nu(\text{C=N})$  1612. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  9.03 (d,  $J$  = 2.9 Hz, 1H, Ar–H), 8.01 (s, 1H), 7.99 (s, 1H), 7.80–7.86 (m, 8H), 7.77–7.68 (m, 16 H), 7.65–7.56 (m, 16H), 7.18–7.10 (t,  $J$  = 8.0 Hz, 1H), 7.08 (d,  $J$  = 10.2 Hz, 1H), 6.72 (m, 2H), 6.38 (m, 2H), 5.32 (s, 1H) 1.27 (s, 3H), 3.50 (s, 3H, Me). ESI-MS (–ve mode) in MeOH:  $m/z$  323.3 [M]<sup>2–</sup>. Anal. Calcd for C<sub>71</sub>H<sub>54</sub>N<sub>6</sub>O<sub>5</sub>OsP<sub>2</sub>: C, 64.44; H, 4.11; N, 6.35. Found: C, 64.10; H, 4.21; N, 6.18%. UV/Vis (CH<sub>2</sub>Cl<sub>2</sub>):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [mol<sup>–1</sup> dm<sup>3</sup> cm<sup>–1</sup>]): 262sh (18 090), 269 (19 810), 276 (18 740), 285sh (12 760), 338 (12 110), 363sh (9360), 417sh (7070), 593 (60 850). Yield for (PPh<sub>4</sub>)<sub>2</sub>3b, 18 mg, 32%. Selected IR (KBr disc,  $\text{cm}^{-1}$ ):  $\nu(\text{C}\equiv\text{N})$  2139 and 2130;  $\nu(\text{C=O})$  1649;  $\nu(\text{C=N})$  1614. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  9.07 (d,  $J$  = 2.8 Hz, 1H), 8.65 (dd,  $J$  = 6.3, 2.8 Hz, 1H) 8.13 (dd,  $J$  = 9.3, 2.9 Hz, 1H), 7.90–7.82 (m, 4H), 7.77–7.84 (m, 8H), 7.65–7.73 (m, 10H), 7.60–7.53 (m, 2H), 7.21 (dd,  $J$  = 9.8, 2.2 Hz, 1H), 6.99 (d,  $J$  = 9.3 Hz, 1H); 5.77 (d,  $J$  = 9.8 Hz, 1H) 3.58 (s, 3H). ESI-MS (–ve mode) in MeOH:  $m/z$  645 [M]<sup>–</sup>. Anal. Calcd for C<sub>47</sub>H<sub>33</sub>N<sub>6</sub>O<sub>5</sub>OsP<sub>2</sub>: C, 57.43; H, 3.38; N, 8.55. Found: C, 57.10;

H, 3.71; N, 8.18%. UV/Vis (CH<sub>2</sub>Cl<sub>2</sub>):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [mol<sup>–1</sup> dm<sup>3</sup> cm<sup>–1</sup>]): 262sh (18 090), 269 (19 810), 276 (18 740), 285sh (12 760), 338 (12 110), 363sh (9360), 417sh (7070), 593 (60 850).

**(PPh<sub>4</sub>)<sub>2</sub>[Os<sup>IV</sup>(L)(CN)<sub>3</sub>(N=<sup>t</sup>Bu<sub>2</sub>Ph<sub>(-2H)</sub>=O)] [(PPh<sub>4</sub>)<sub>2</sub>4a] and (PPh<sub>4</sub>)<sub>2</sub>[Os<sup>IV</sup>(L)(CN)<sub>3</sub>(NH=<sup>t</sup>Bu<sub>3</sub>PhOH<sub>(-2H)</sub>)] [(PPh<sub>4</sub>)<sub>2</sub>4b]**

The synthetic route of (PPh<sub>4</sub>)<sub>2</sub>4a and (PPh<sub>4</sub>)<sub>2</sub>4b is similar to that of (PPh<sub>4</sub>)<sub>2</sub>1a and (PPh<sub>4</sub>)<sub>2</sub>1b, except that the 2,4,6-<sup>t</sup>Bu<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH is used instead of 2,6-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>OH. The first yellow band (PPh<sub>4</sub>)<sub>2</sub>4a was eluted by CH<sub>2</sub>Cl<sub>2</sub>/acetone (v:v, 10:1). Yellow needle crystals were obtained from the slow diffusion of diethyl ether into an acetone solution of (PPh<sub>4</sub>)<sub>2</sub>4a. The solvent was removed under reduced pressure.

Yield for (PPh<sub>4</sub>)<sub>2</sub>4a: 32 mg, 52%. Selected IR (KBr disc,  $\text{cm}^{-1}$ ):  $\nu(\text{C}\equiv\text{N})$  2142 and 2128,  $\nu(\text{C=O})$  1638;  $\nu(\text{C=N})$  1615;  $\nu(\text{N=O})$  1307. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  9.03 (d,  $J$  = 2.8 Hz, 1H, Ar–H), 8.12 (s, 2H, Ph–H), 8.05 (dd,  $J$  = 9.3, 2.9 Hz, 1H, Ar–H), 7.92 (dd,  $J$  = 8.0, 6.0 Hz, 4H, PPh<sub>4</sub>–H), 7.83–7.76 (m, 9H, Ar–H and PPh<sub>4</sub>–H), 7.73 (d,  $J$  = 7.2 Hz, 1H, Ar–H), 7.68–7.62 (m, 8H, PPh<sub>4</sub>–H), 7.54 (dtd,  $J$  = 13.9, 7.8, 6.2 Hz, 2H, Ar–H), 6.89 (d,  $J$  = 9.4 Hz, 1H, Ar–H), 1.28 (s, 18H, CH<sub>3</sub>). ESI-MS (–ve mode) in MeOH:  $m/z$  743 [M]<sup>2–</sup>. Anal. Calcd for C<sub>54</sub>H<sub>47</sub>N<sub>6</sub>O<sub>5</sub>OsP<sub>2</sub>: C, 59.99; H, 4.38; N, 7.77. Found: C, 60.10; H, 4.31; N, 7.78%. UV/Vis (CH<sub>2</sub>Cl<sub>2</sub>):  $\lambda_{\text{max}}$  [nm] ( $\epsilon$  [mol<sup>–1</sup> dm<sup>3</sup> cm<sup>–1</sup>]): 230 (27 380), 277 (15 620), 288 (14 640), 330 (13 030), 428 (20 930).

**(PPh<sub>4</sub>)<sub>2</sub>[Os<sup>IV</sup>(L)(CN)<sub>3</sub>(NH-Me<sub>3</sub>PhOH<sub>(-H)</sub>)] [(PPh<sub>4</sub>)<sub>2</sub>5]**

The synthetic route of (PPh<sub>4</sub>)<sub>2</sub>5 is similar to that of (PPh<sub>4</sub>)<sub>2</sub>1a and (PPh<sub>4</sub>)<sub>2</sub>1b, except that the 2,4,6-Me<sub>3</sub>C<sub>6</sub>H<sub>2</sub>OH is used instead of 2,6-Me<sub>2</sub>C<sub>6</sub>H<sub>3</sub>OH. (PPh<sub>4</sub>)<sub>2</sub>5 was purified by silica gel CH<sub>2</sub>Cl<sub>2</sub>/acetone (5:3) as the eluent. Yield: 43 mg, 75% (based on the OsN consumed). UV/Vis (CH<sub>2</sub>Cl<sub>2</sub>) for (PPh<sub>4</sub>)<sub>2</sub>5:  $\lambda_{\text{max}}$  [nm] ( $\epsilon/\text{M}^{-1} \text{cm}^{-1}$ ): 233 (42 430), 270 (18 030), 277 (19 430), 293 (20 360), 353 (20 460), 457 (6420). Selected IR (KBr disc,  $\text{cm}^{-1}$ ) for 5:  $\nu(\text{C}\equiv\text{N})$  2135 and 2127;  $\nu(\text{N-H})$  3247;  $\nu(\text{C=N})$  1610. ESI-MS (–ve mode) in MeOH:  $m/z$  675 [M]<sup>–</sup>. Anal. Calcd for C<sub>49</sub>H<sub>39</sub>N<sub>6</sub>O<sub>5</sub>OsP<sub>2</sub>: C, 58.09; H, 3.88; N, 8.30. Found: C, 58.10; H, 3.81; N, 8.28%. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  8.93 (d,  $J$  = 2.8 Hz, 1H, Ar–H), 7.96 (d,  $J$  = 6.6 Hz, 1H, Ar–H), 7.89 (t,  $J$  = 7.2 Hz, 4H), 7.75 (td,  $J$  = 7.6, 3.7 Hz, 8H), 7.65–7.57 (m, 10H, Ar–H and PPh<sub>4</sub>–H), 7.43 (t,  $J$  = 7.1 Hz, 1H, Ar–H), 7.20 (t,  $J$  = 8.4 Hz, 1H, Ar–H), 6.78 (s, 2H, Ph–H), 6.57 (d,  $J$  = 9.4 Hz, 1H, Ar–H), 2.23 (s, 3H, CH<sub>3</sub>), 1.27 (s, 6H, CH<sub>3</sub>).

## Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

## Data availability

The authors will supply the relevant data in response to reasonable requests.



## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was supported by the National Natural Science Foundation of China (22371092), the Excellent Discipline Cultivation Project by JHUN (2023XKZ038), the Project of Hebei Key Laboratory of Heterocyclic Compounds (no. KF202402), and the Graduate Scientific Research Foundation of Jiangnan University (KYCXJJ202426).

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