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Ligand Redox Non-Innocence in $[\text{Co}^{\text{III}}(\text{TAML})]^{0/-}$ Complexes Affects Nitrene Formation

Nicolaas P. van Leest,[†] Martijn A. Tepaske,[†] Jean-Pierre H. Oudsen,[§] Bas Venderbosch,[§] Niels R. Rietdijk,[†] Maxime A. Siegler,[‡] Moniek Tromp,[§] Jarl Ivar van der Vlugt^{†,*} and Bas de Bruin^{†,*}

[†]Homogeneous, Supramolecular and Bio-Inspired Catalysis Group and [§]Sustainable Materials Characterization Group, van 't Hoff Institute for Molecular Sciences (HIMS), University of Amsterdam, Science Park 904, 1098XH Amsterdam, The Netherlands

[‡]Department of Chemistry, Johns Hopkins University, Baltimore, Maryland 21218, United States

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ABSTRACT: The redox non-innocence of the TAML scaffold in cobalt-TAML (Tetra-Amido Macrocyclic Ligand) complexes has been under debate since 2006. In this work we demonstrate with a variety of spectroscopic measurements that the TAML backbone in the anionic complex $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ is truly redox non-innocent, and that one-electron oxidation affords $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$. Multireference (CASSCF) calculations show that the electronic structure of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ is best described as an intermediate spin ($S = 1$) cobalt(III) center that is antiferromagnetically coupled to a ligand-centered radical, affording an overall doublet ($S = 1/2$) ground-state. Reaction of the cobalt(III)-TAML complexes with PhINNs as a nitrene precursor leads to TAML-centered oxidation, and produces nitrene radical complexes without oxidation of the metal ion. The ligand redox state (TAML^{red} or TAML^{sq}) determines whether mono- or bis-nitrene radical complexes are formed. Reaction of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ or $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ with PhINNs results in formation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns})]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns})_2]^-$, respectively. Herein, ligand-to-substrate single-electron transfer results in one-electron reduced Fischer-type nitrene radicals ($\text{N}^{\bullet}\text{Ns}^-$) that are intermediates in catalytic nitrene transfer to styrene. These nitrene radical species were characterized by EPR, XANES, and UV-Vis spectroscopy, high resolution mass spectrometry, magnetic moment measurements and supporting CASSCF calculations.

INTRODUCTION

The use of base metals and redox non-innocent (or redox-active) ligands in radical-type carbene, oxo and nitrene transfer reactions has evolved as a powerful tool for the direct functionalization of (unactivated) C–H bonds and olefins.¹ The functionalized products of these reactions are motifs in pharmaceuticals and agrochemicals, and are therefore highly valued.² N-group transfer reactivity is an efficient way to afford the direct synthesis of e.g. secondary amines and aziridines, of which the synthesis otherwise typically requires harsh reaction conditions or multiple steps.³ Generation of the essential catalytic metal-nitrene intermediates has been achieved with second and third row transition metals (Ru,⁴ Rh,⁵ Pd,⁶ Ag⁷ and Au⁸) as well as more abundant base metals (Mn,⁹ Fe,¹⁰ Co,¹¹ Ni¹² and Cu¹³).

Our group, in collaboration with the Zhang group, has studied the formation and reactivity of nitrene adducts of cobalt(II)-porphyrin complexes, which are competent catalysts for a range of (enantioselective) amination and aziridination reactions.^{11a,11d-j,14} The mono-nitrene species generated on cobalt upon reaction with an organic azide is most accurately described as a one-electron reduced Fischer-type nitrene radical.^{14b} This interesting electronic structure is the result of *metal-to-substrate* single-electron transfer (SET), wherein cobalt is oxidized from Co^{II} to Co^{III} , and the nitrene is reduced by one

electron to produce a nitrene radical ($\text{N}^{\bullet}\text{R}^-$) complex with single-electron population of the π symmetric Co–N antibonding orbital. Interestingly, reaction of cobalt(II)-porphyrins with iminodinanones (PhINNs, Ns = nosyl) led to formation of bis-nitrene radical species with two one-electron reduced Fischer-type nitrenes, wherein the second nitrene is reduced via *ligand-to-substrate* SET. Intrigued by these nitrene transfer catalysts, we became interested in the possibility of nitrene radical formation on square planar cobalt(III) platforms involving solely *ligand-to-substrate* single electron transfer,¹⁵ by studying systems containing redox-active ligands for which *metal-to-substrate* SET is difficult or even impossible.

When searching for suitable redox-active macrocyclic tetradentate ligand platforms that enforce a square planar coordination geometry around cobalt in an oxidation state higher than +II, we decided to investigate the Tetra-Amido Macrocyclic Ligand (TAML) platform designed by the group of Collins.¹⁶ The general structure of a TAML that met the aforementioned requirements is depicted in Scheme 1. Moreover, the potential redox non-innocence of TAML and related *o*-phenylenedicarboxamido complexes has been proposed in the literature, and for clarity we will follow the nomenclature as presented in Scheme 1 for the fully reduced tetra-anion (*red*), mono-oxidized tri-anionic ligand-centered radical (*sq*) and fully oxidized di-anion (*q*).^{16,17}

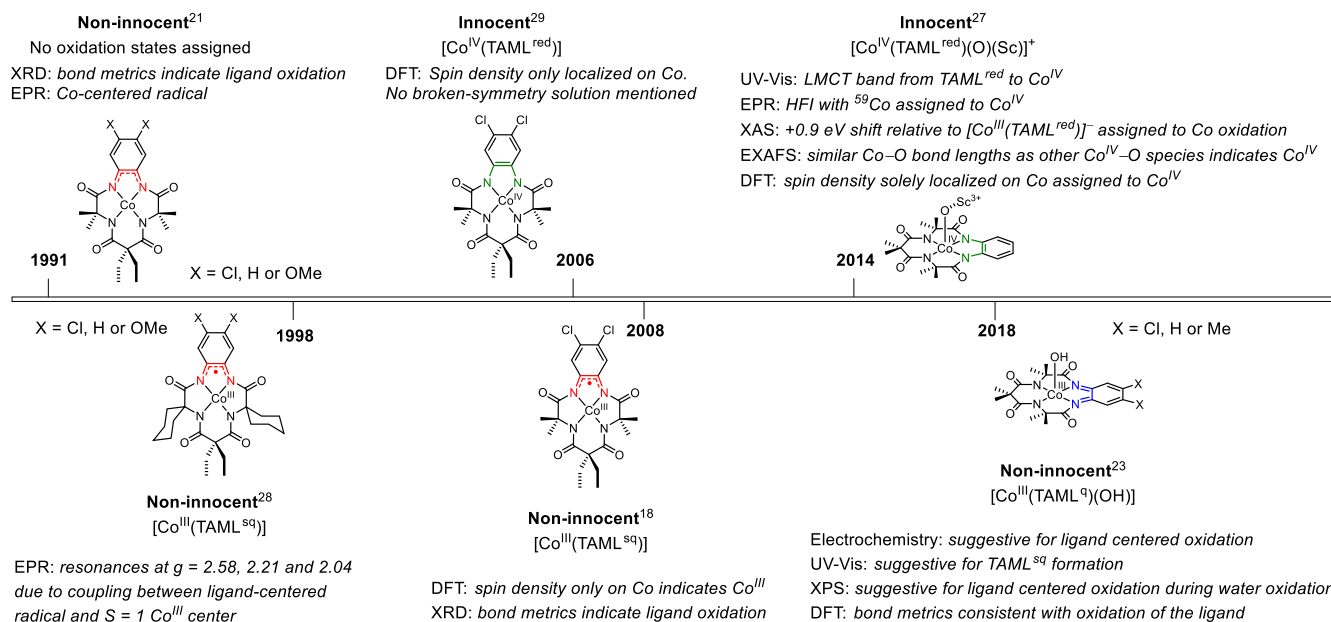
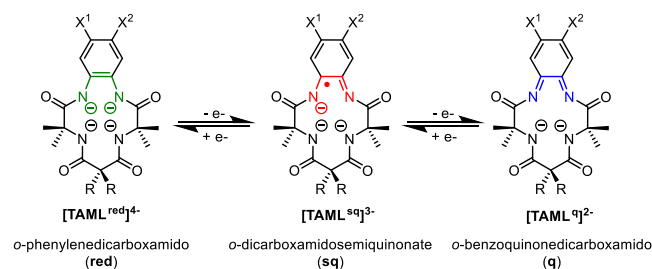


Figure 1. Interpretation of the ligand (non-)innocence in cobalt-TAML complexes in chronological order. HFI = hyperfine interaction.

Scheme 1. General structure of the TAML scaffold and the potential redox non-innocence of the backbone. X¹ = Cl, H, NO₂, OMe. X² = Cl, H. R = Et, Me, F.¹⁶



Iron complexes of these TAML activators have found widespread use in oxidation chemistry, and TAML complexes with Cr, Mn, Fe, Co, Ni and Cu have been reported with many variations of the TAML scaffold.^{16,18} Interestingly, *ligand-centered oxidation* of an [Fe^V(TAML^{red})(NTs)]⁻ complex was shown to afford [Fe^V(TAML^{sq})(NTs)], which is a more active nitrene transfer species towards activated C–H bonds (bond dissociation energy between 75 and 80 kcal mol⁻¹) and thioanisole than the reduced analogue.¹⁹ A similar trend was observed for a manganese-imido complex, wherein [Mn^V(TAML^{red})(NMe₃)]⁻ (Me₃ = mesityl) proved to be unreactive, while the *metal-centered oxidized* complex [Mn^{VI}(TAML^{red})(NMe₃)] could be used for hydrogen atom transfer reactions and nitrene transfer to thioanisole.²⁰ Apparently the redox activity of the TAML ligand varies from complex to complex, depending on the metal and other ligands, and both metal- and ligand-centered redox processes can be used to influence nitrene-transfer reactivity.

Specific [Co^{III}(TAML^{red})]⁻ complexes²¹ have been used for electrochemical water oxidation^{22,23} and oxygen reduction,²⁴ cycloaddition of CO₂ to epoxides,²⁵ electrochemical sensing of H₂O₂,²⁶ oxo transfer to C–H bonds²⁷ and electron-transfer reactions.²⁸ However, to the best of our knowledge, no nitrene transfer reactions or stoichiometric reactions leading to formation of

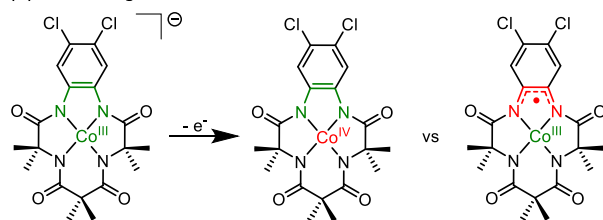
Co(TAML)-based imido- or nitrene-complexes have been reported to date. Moreover, contrary to chemistry with iron, the existence of TAML-centered redox processes in cobalt complexes is still under debate (Figure 1).

Collins *et al.*²¹ reported the synthesis and characterization of an anionic [Co^{III}(TAML^{red})]⁻ complex with a diamidophenyl backbone in 1991. The anionic parent complex was characterized as a triplet with an S = 1 Co center and a fully reduced o-phenylenedicarboxamido ligand. Oxidation of this complex afforded a neutral S = 1/2 system, for which crystallographic bond metrics indicated single-electron oxidation of the ligand and electron paramagnetic resonance (EPR) data hinted at a cobalt-centered radical (Figure 1). This data was interpreted in 1998 as corresponding to an S = 1 cobalt(III) center antiferromagnetically coupled to a ligand-centered radical ([Co^{III}(TAML^{sq})]⁻).²⁸ Ghosh *et al.*²⁹ reported an elaborate density functional theory (DFT) study on the ligand non-innocence of multiple variations of the TAML backbone, and suggested that the electronic structure of [Co^{III}(TAML^{sq})] is better described as [Co^{IV}(TAML^{red})] (Figure 1). Their assignment was based on the Mulliken spin density, which was solely localized on cobalt. Collins and coworkers¹⁸ critically re-interpreted these spin densities as being evidence of an S = 1 Co^{III} center. It should be noted that multireference post-Hartree-Fock methods were not accessible at the time, and possible broken-symmetry solutions were apparently not explored. As such, optional antiferromagnetic coupling between an S = 1 Co center and a ligand-centered radical could have remained hidden in the applied DFT calculations.

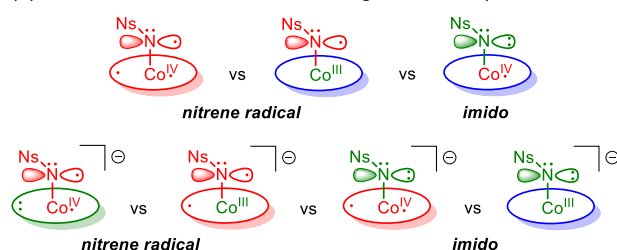
Innocent behavior of the TAML scaffold was claimed in an electrochemical study reported in 2014,³⁰ as well as in the characterization of a Lewis-acid stabilized oxo-complex [Co^{IV}(TAML^{red})(O)].²⁷ The TAML^{red} and Co^{IV} oxidation states in a Sc³⁺-bound [Co^{IV}(TAML^{red})(O)]²⁻ complex were based on UV-Vis, EPR, XAS (X-ray absorption spectroscopy) and EXAFS (extended X-ray absorption fine structure) studies, in combination with DFT calculated Mulliken spin densities.²⁷ On

the contrary, TAML-centered redox-activity in $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{OH})]$ was claimed in 2018 on the basis of UV-Vis, EPR and XPS (X-ray photoelectron spectroscopy) studies.²³

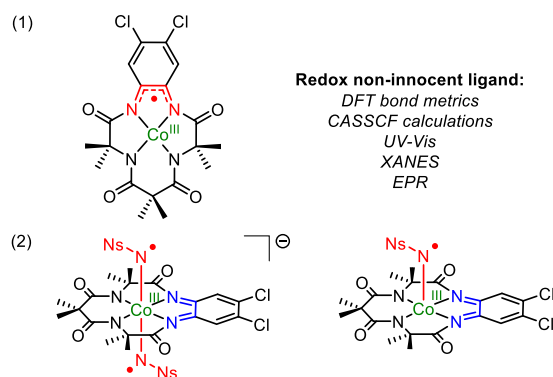
(A) Possible ligand redox non-innocence:



(B) Possible electronic structures of the targeted nitrene species:



(C) Main findings:



Ligand oxidation state influences nitrene formation:

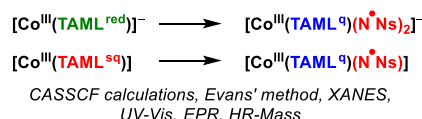


Figure 2. (A): Electronic structure questions regarding the redox non-innocence of the TAML scaffold. (B): Possible electronic structures of the targeted nitrene species. The ligand color-coding is as presented in Scheme 1. (C): Main findings with the assignment of the TAML scaffold being redox non-innocent in the coordination sphere of cobalt and its influence on nitrene (radical) formation.

Given i) the contrasting descriptions of ligand and cobalt oxidation states in $[\text{Co}(\text{TAML})]$ complexes, ii) our interest in generating cobalt-nitrene radical intermediates via ligand-to-substrate SET and iii) the previous characterization of $[\text{Fe}(\text{TAML})(\text{imido})]$, $[\text{Mn}(\text{TAML})(\text{imido})]$ and $[\text{Co}(\text{TAML})-(\text{oxo})]$ complexes, we set out to answer the following research questions:

- (1) Is the ligand in $[\text{Co}(\text{TAML})]$ complexes redox non-innocent and can the different assignments in literature be reconciled? (See Figure 2A).
- (2) Can the $[\text{Co}(\text{TAML})]$ platform be used to generate (catalytically competent) cobalt-nitrene (radical) species, and what is the influence of the (ligand) oxidation state

on the (electronic) structure of the targeted nitrene (radical) species? (See Figure 2B).

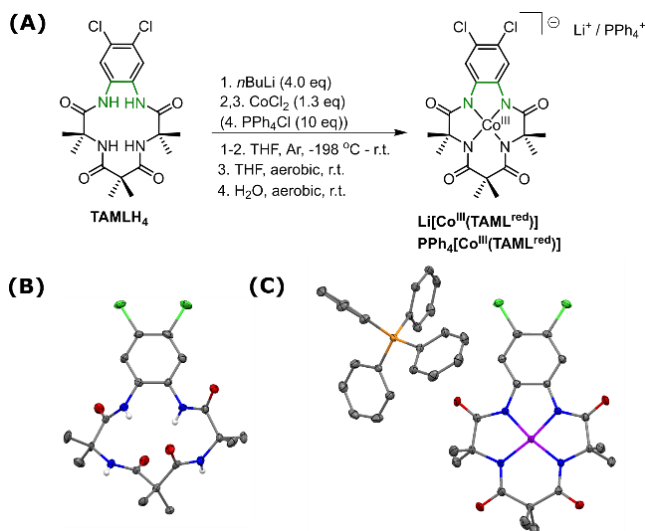
- (3) In case the TAML ligand platform is indeed redox-active, can we use this feature for ligand-to-substrate SET to produce nitrene radical species at square planar cobalt(III) species? (See Figure 2B).

The main findings of the investigations presented in this paper are summarized in Figure 2C.

RESULTS AND DISCUSSION

Ligand-Centered Oxidation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$. The parent $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ complex was obtained according to an adapted literature procedure.^{21,31} After a five-step synthetic procedure to obtain the ligand (**TAMLH₄**), coordination of Co^{II} to the fully deprotonated ligand (generated using *n*-BuLi) and aerobic oxidation afforded $\text{Li}[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$, or **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** after salt metathesis with **PPh₄Cl** (Scheme 2). Crystals suitable for single crystal X-ray diffraction (XRD) analysis of **TAMLH₄** and **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** were grown by vapor diffusion of pentane into concentrated THF solutions of the ligand or complex, respectively. The solid state structure of **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** displays a square planar geometry around cobalt and a non-coordinating THF molecule in the crystal lattice. As expected, and in accordance with literature,²¹ analysis of the crystallographic bond metrics (see Supporting Information) of the diamidophenyl ring in **TAMLH₄** and **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** supports the preservation of aromaticity upon coordination to cobalt, with the ligand being fully reduced ($(\text{TAML}^{\text{red}})^{4-}$) and the metal adopting the Co^{III} oxidation state. The effective magnetic moment of **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** , as determined via the Evans' method,³² indicated a triplet ($S = 1$) ground state ($\mu_{\text{eff}} = 2.94 \mu_{\text{B}}$). This is in accordance with literature and expected for an intermediate spin Co^{III} center with two parallel metal-centered unpaired electrons.²¹ The DFT optimized structure of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ in the triplet state at the BP86/def2-TZVP level of theory is consistent with these observations and the calculated bond metrics closely match the experimental bond lengths (see SI).

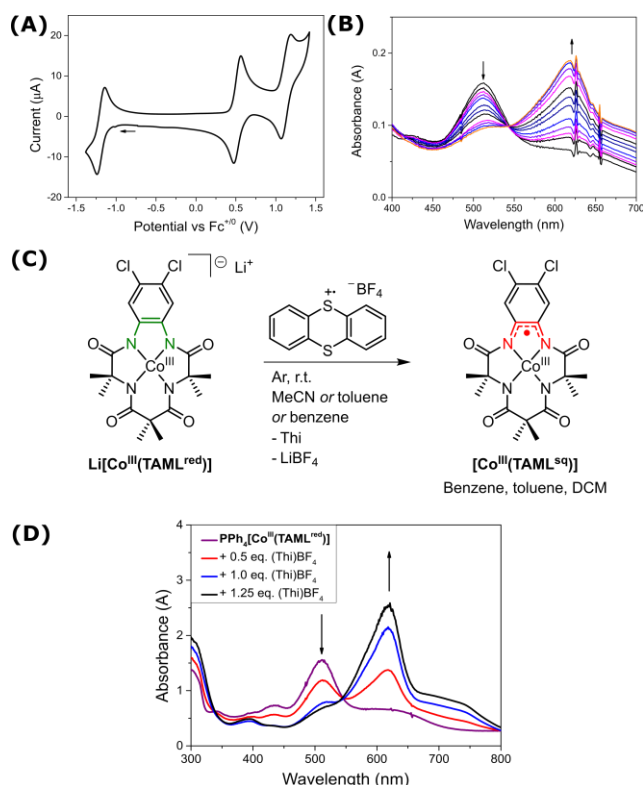
Scheme 2. (A): Formation of **Li $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** and **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** from **TAMLH₄**. Thermal displacement ellipsoid plots (50% probability level) of **TAMLH₄** (B) and **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$** (C). H-atoms (except for NH) and lattice solvent (THF for **PPh₄ $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$) removed for clarity.**



Electrochemical oxidation of **PPh₄[Co^{III}(TAML^{red})]** in CH₂Cl₂ (Scheme 3A) using cyclic voltammetry displays three fully reversible redox events at $E_{1/2} = -1.18$, $+0.53$ and $+1.13$ V vs. Fc^{+/0}, which are attributed to metal-centered reduction (Co^{III/II}) and two ligand-centered oxidations (TAML^{red/sq} and TAML^{sq/q}) respectively (*vide infra*).³³ UV-Vis spectroelectrochemical (UV-Vis-SEC) monitoring of the oxidation event at $+0.53$ V vs. Fc^{+/0} shows disappearance of **PPh₄[Co^{III}(TAML^{red})]** ($\lambda_{\text{max}} = 510$ nm) and concomitant appearance of the characteristic absorption band of **[Co^{III}(TAML^{sq})]** ($\lambda_{\text{max}} = 623$ nm) with an isosbestic point at 545 nm (Scheme 3B).^{21,23,34,35} For clarity we already assigned the electronic structure of **[Co^{III}(TAML^{sq})]** in the following descriptions. In the following sections we will further elaborate on the measurements and calculations leading to this assignment.

Chemical oxidation of TAML complexes with ceric ammonium nitrate ((NH₄)₂[Ce(NO₃)₆]) typically requires excess oxidant and large volumes of solvent to extract the product.²¹ For purple colored **PPh₄[Co^{III}(TAML^{red})]** and **Li[Co^{III}(TAML^{red})]**, oxidation with a stoichiometric amount of thianthrenium tetrafluoroborate ((Thi)BF₄) ($E^0_{1/2} = 0.86$ V vs. Fc^{+/0})³⁶ cleanly afforded the blue colored **[Co^{III}(TAML^{sq})]** complex (Scheme 3C). A UV-Vis titration gave identical data as the UV-Vis-SEC monitoring of the oxidation event at $+0.53$ V vs. Fc^{+/0} (Scheme 3D).

Scheme 3. (A): Cyclic voltammogram of PPh₄[Co^{III}(TAML^{red})] in DCM (see SI for details). (B): UV-Vis-SEC oxidation of PPh₄[Co^{III}(TAML^{red})] in DCM (see SI for details). (C): Oxidation of [Co^{III}(TAML^{red})]⁻ to [Co^{III}(TAML^{sq})] with (Thi)BF₄. (D): UV-Vis titration of PPh₄[Co^{III}(TAML^{red})] in DCM (0.15 mM) with increasing amounts of (Thi)BF₄.



characteristic for a net $S = 1/2$ system with unpaired electron density on cobalt ($g_{\text{iso}} = 2.22$) (Figure 3A). EPR measurements at 10 K in a toluene glass showed a rhombic signal with $g_x = 2.03$, $g_y = 2.16$, $g_z = 2.54$ and partially unresolved cobalt hyperfine interactions (HFIs) (Figure 3B). The inclusion of ⁵⁹Co ($I = 7/2$ nucleus) HFIs ($A^{\text{Co}}_{\text{x}} = 5.0$ MHz, $A^{\text{Co}}_{\text{y}} = 50.0$ MHz, $A^{\text{Co}}_{\text{z}} = 20.0$ MHz) is necessary for accurate simulation of the spectrum. The DFT calculated cobalt HFIs are overestimated (B3LYP/def2-TZVP: $g_x = 2.04$, $g_y = 2.25$, $g_z = 2.26$, $A^{\text{Co}}_{\text{x}} = 166.3$ MHz, $A^{\text{Co}}_{\text{y}} = 199.8$ MHz, $A^{\text{Co}}_{\text{z}} = 641.3$ MHz), which we attribute to the erroneous description of multireference systems with DFT methods (*vide infra*). Interestingly, the isotropic X-band EPR spectrum measured in MeCN (Figure 3C) revealed an eight-line pattern at $g_{\text{iso}} = 2.00$, attributed to hyperfine coupling with cobalt ($A^{\text{Co}}_{\text{iso}} = 36.0$ MHz) in **[Co^{III}(TAML^{sq})(MeCN)]**, which is in excellent agreement with the DFT calculated parameters (B3LYP/def2-TZVP: $g_{\text{iso}} = 2.00$, $A^{\text{Co}}_{\text{iso}} = 34.2$ MHz). Notably, this species has a single-reference doublet electronic structure with the unpaired electron residing in a cobalt-ligand π^* orbital (strongly delocalized over cobalt and the ligand; see SI).

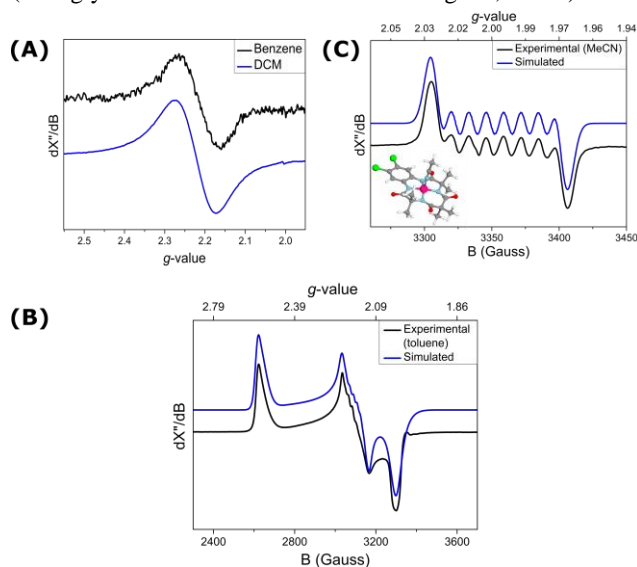


Figure 3. (A): X-band EPR spectrum of **[Co^{III}(TAML^{sq})]** in benzene at r.t. (black line: microwave freq. 9.390167 GHz, mod. amp. 4 G, power 2.518 mW) and CH₂Cl₂ (blue line: microwave freq. 9.3966 GHz, mod. amp. 5 G, power 2.000 mW) with $g_{\text{iso}} = 2.22$. (B): experimental (black) and simulated (blue) X-band EPR spectrum of **[Co^{III}(TAML^{sq})]** in toluene at 10 K. Microwave freq. 9.365984 GHz, mod. amp. 4 G, power 2.000 mW. Simulation parameters: $g_x = 2.03$, $g_y = 2.16$, $g_z = 2.54$, $A^{\text{Co}}_{\text{x}} = 5.0$ MHz, $A^{\text{Co}}_{\text{y}} = 50.0$ MHz, $A^{\text{Co}}_{\text{z}} = 20.0$ MHz, linear A-strain -0.018 (z direction), quadratic A strain -18 (x direction) and -2 (y direction). (C): Experimental (black) and simulated (blue) X-band EPR spectrum of **[Co^{III}(TAML^{sq})(MeCN)]** at r.t. in MeCN and DFT (BP86/def2-TZVP/disp3) optimized structure. Microwave freq. 9.3886 GHz, mod. amp. 3 G, power 0.7962 mW. Simulated (calculated; B3LYP/def2-TZVP) parameters: $g_{\text{iso}} = 2.00$ (2.00) and $A^{\text{Co}}_{\text{iso}} = 36.0$ (34.2) MHz.

The effective magnetic moment of **[Co^{III}(TAML^{sq})]** ($\mu_{\text{eff}} = 1.88 \mu_{\text{B}}$, Evans' method) was found to be consistent with an overall net doublet ($S = 1/2$) ground state. Room temperature (r.t.) X-band EPR studies in CH₂Cl₂ or toluene reveal a signal

The cobalt oxidation state of the four-coordinate complexes was further investigated using Co K-edge X-ray near edge structure (XANES) analysis. The Co K-edge XANES spectra of **PPh₄[Co^{III}(TAML^{red})]** and **[Co^{III}(TAML^{sq})]** in toluene are compared in Figure 4. The edge position (determined at the half-step height) was 7721 eV for both complexes. Both spectra are identical, which is in line with the same oxidation state (+III) and similar coordination geometry of cobalt in the two

complexes. The +III oxidation state of cobalt was already found in $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ (according to XRD-derived bond metric analysis, *vide supra*) and the observed edge position is equal to a related $[\text{Co}^{\text{III}}(\text{TAML})]^-$ complex.²⁷ The shoulder at approximately 7715 eV in the Co K-edge XANES spectra is typical for square planar Co complexes, including square planar Co-porphyrin complexes and a related cobalt-TAML complex.^{14b,27} The main edge feature arises primarily from $1s \rightarrow 4p$ electron transitions, whereas the feature at 7715 eV is commonly assigned to $1s \rightarrow 4p_z$ and ligand to metal charge transfer (LMCT) shakedown transitions.³⁷

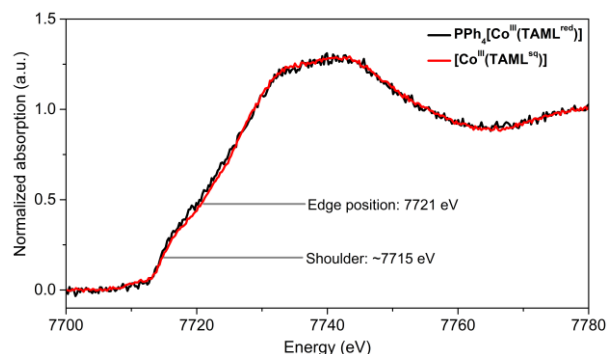


Figure 4. Co-K edge XANES analysis of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ (black) and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ (red) in toluene.

In agreement with previous studies,^{27,29} DFT calculations with various GGA and hybrid functionals (BP86, B3LYP, PBE and OPBE; see SI for details) gave unsatisfactory results for the Co(TAML)-type complexes under investigation. An illustrative example of the problem encountered with DFT is found in the challenging description of the net-doublet ground state of the $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ complex. Distinguishing between a genuine Co^{IV} complex and a multireference electronic structure solution involving antiferromagnetic coupling between an $S = 1$ Co^{III} center and a TAML ligand-centered radical (as indicated by the B3LYP broken symmetry DFT solution) is very difficult, if not impossible, when relying only on single reference computational methods (such as DFT).³⁸ We therefore decided to turn to multireference N-Electron Valence State Perturbation Theory (NEVPT2)-corrected Complete Active Space Self Consistent Field (CASSCF) calculations for a proper description of the electronic structures of the Co(TAML)-type complexes described in this paper.³⁹

CASSCF calculations were initiated on the anionic $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ complex by inclusion of all cobalt d -orbitals and those ligand π -orbitals (L_π) that could have an interaction with cobalt. In the final CASSCF(14,13) calculation, all initial orbitals were preserved in the active space, except for the d_{xy} orbital, which is uncorrelated (occupancy of 2.00).⁴⁰ A selection of the most relevant active orbitals with their occupancies (in parentheses) is given in Figure 5A. Löwdin population analysis of the electronic configuration of the d -shell gave $(d_{xy})^{2.00}(d_{z^2})^{1.99}(d_{yz})^{1.02}(d_{xz})^{1.02}$, consistent with the assigned +III oxidation state of cobalt. Notably, the L_π orbital at $-0.268 E_h$ has a weak bonding interaction with the d_{xz} orbital and is fully filled (occupancy 1.91), consistent with the fully reduced oxidation state of the ligand.

CASSCF(13,12) calculations on the neutral $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ complex included a similar active space as for the parent anionic complex, and revealed substantial multi-reference character. The uncorrelated d_{z^2} and d_{xy} orbitals (occupancy 2.00) were not preserved in the active space.⁴⁰ The reduced charge on

the complex causes increased stabilization of the cobalt d -orbitals compared to the parent anionic complex, which increases overlap between the d_{xz} and L_π orbitals (Figure 5B). Due to this stabilization, the bonding and antibonding combinations of the d_{xz} and L_π orbitals are comprised of substantial contributions from both d_{xz} and L_π . As a result of three orbitals being close in energy, significant population of the L_π - d_{xz} antibonding combination (occupancy 0.64) from the $d_{xz}+L_\pi$ bonding combination (occupancy 1.38) occurs, while the d_{yz} orbital is singly occupied (1.07). The net-doublet ground state of the neutral $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ complex is thus best described as an $S = 1$ Co^{III} center that is antiferromagnetically coupled to an $S = 1/2$ TAML-centered radical, leading to a net-doublet system with a $(d_{xy})^{2.00}(d_{z^2})^{2.00}(d_{xz}+L_\pi)^{1.38}(d_{yz})^{1.07}(L_\pi-d_{xz})^{0.64}$ electronic structure, in agreement with the early interpretation of Collins.²⁸ Excitation energies derived from the CASSCF(13,12) calculation revealed that the absorption band observed at $\lambda_{\text{max}} = 623 \text{ nm}$ (Scheme 3B and D) is indeed characteristic for the ligand-centered radical. The corresponding calculated excitation (at 625 nm) is comprised of ligand-centered $L_\pi \rightarrow L_\pi$ - d_{xz} and metal-to-ligand ($d_{xz}+L_\pi \rightarrow L_\pi$ - d_{xz} and $d_{yz} \rightarrow L_\pi$ - d_{xz}) charge-transfer processes, with the ligand-centered radical orbital being the acceptor in all cases.

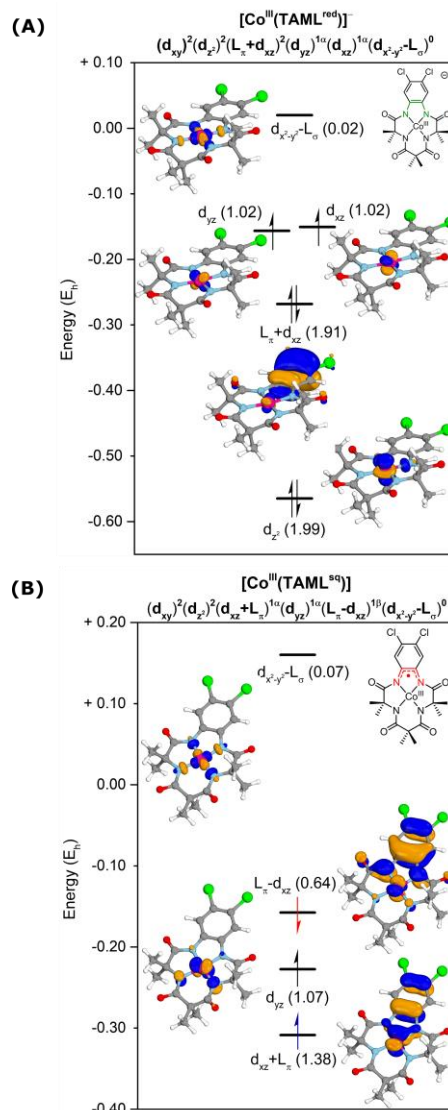


Figure 5. Relevant active orbitals and occupancies (in parenthesis) of NEVPT2-corrected CASSCF(14,13) on $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ (A) and CASSCF(13,12) on $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ (B).

The combined data from magnetic moment measurements, EPR, UV-Vis and XANES spectroscopy and NEVPT2-CASSCF calculations reveal that oxidation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ is ligand-centered, giving rise to the formation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$, wherein cobalt retains its +III oxidation state.

Synthesis and Characterization of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ via Ligand-to-Substrate SET. With a proper understanding of their electronic structure, confirming that both complexes are square planar cobalt(III) species featuring a redox-active ligand, but in two different ligand oxidation states, we next set out to investigate nitrene formation at the anionic $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ and neutral $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ complexes. We were particularly interested in exploring the influence of the ligand oxidation state on the structure and overall composition of the targeted nitrene adducts.

Addition of one equivalent of the nitrene precursor PhINNs to $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ in CH_2Cl_2 at r.t. led to a mixture of starting material, mono-nitrene adduct $[\text{Co}(\text{TAML})(\text{NNs})]^-$ and trace amounts of bis-nitrene adduct $[\text{Co}(\text{TAML})(\text{NNs})_2]^-$, as revealed by negative mode electrospray ionization high resolution mass spectrometry (ESI-HRMS⁻) analysis. Upon addition of ten equivalents PhINNs to $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ in CH_2Cl_2 or toluene at r.t., quantitative formation of bis-nitrene species $[\text{Co}(\text{TAML})(\text{NNs})_2]^-$ was achieved, based on ESI-HRMS⁻ and UV-Vis analysis (Scheme 4 and Figure 6A,B).⁴¹ While bis-nitrene formation was readily achieved for the anionic complex upon addition of an excess of PhINNs, addition of ten equivalents of PhINNs to the neutral complex $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ in CH_2Cl_2 or toluene at r.t. led to quantitative formation of only the mono-nitrene species $[\text{Co}(\text{TAML})(\text{NNs})]$, as shown by ESI-HRMS⁻ and UV-Vis analysis (Scheme 4 and Figure 6C, D). Also the addition of the alternative nitrene source PhINTs (ten equivalents) to $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ in CH_2Cl_2 or toluene at room temperature led to formation of mono-nitrene complex $[\text{Co}(\text{TAML})(\text{NTs})]$, according to ESI-HRMS⁻ data. The effective magnetic moments

of $[\text{Co}(\text{TAML})(\text{NNs})_2]^-$ ($2.75 \mu_B$) and $[\text{Co}(\text{TAML})(\text{NNs})]$ ($1.53 \mu_B$) are consistent with formation of (net) triplet ($S = 1$) and doublet ($S = 1/2$) systems, respectively. For clarity, we already included the assigned oxidation states of the ligand and cobalt for the anionic bis-nitrene ($[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$) and neutral mono-nitrene ($[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$) in Scheme 4 and Figure 6 and the following text. In the next sections we will further elaborate on the measurements and calculations leading to these assignments.

Scheme 4. Synthesis of bis-nitrene (radical) complex $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ and mono-nitrene (radical) complex $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NR})]$ (R = Ns, Ts) from $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$, respectively.

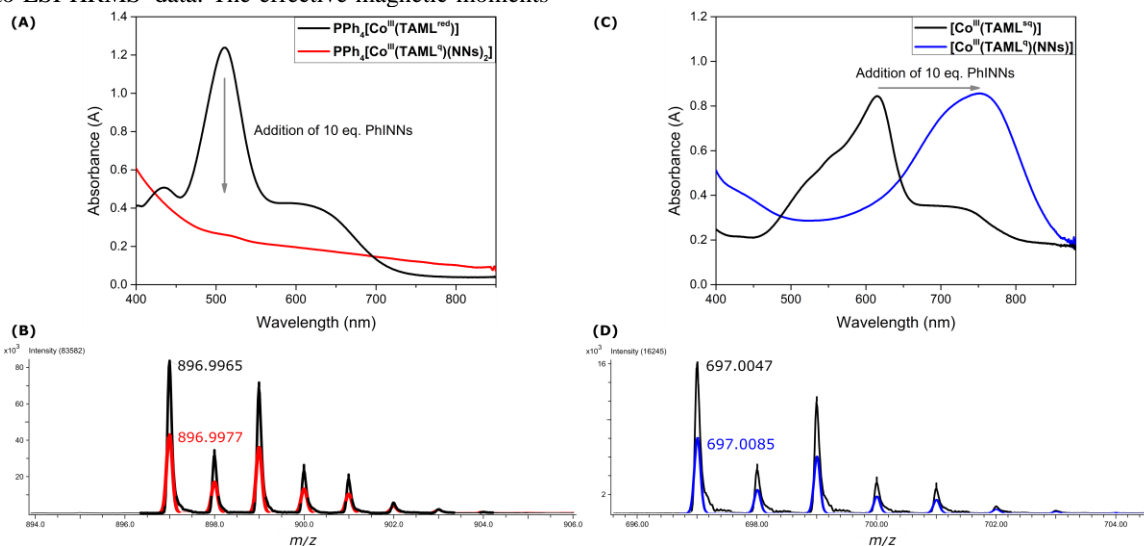
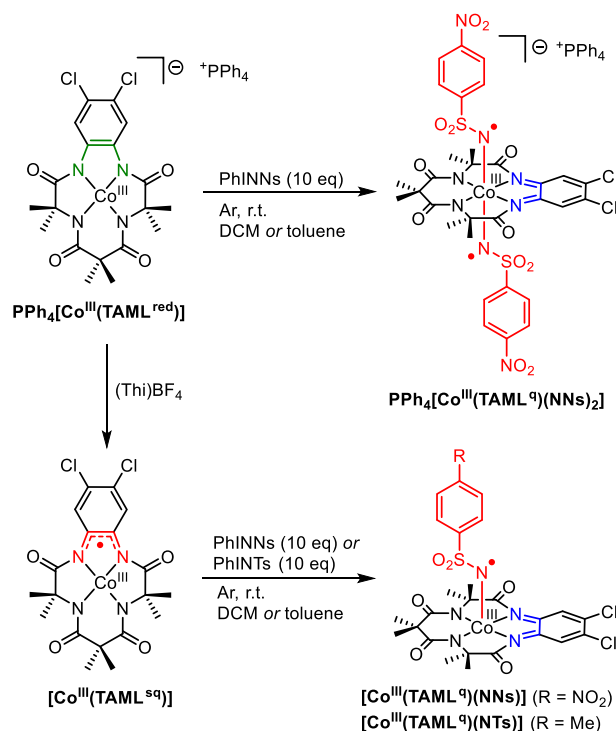


Figure 6. (A): UV-Vis spectrum of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ (red) upon reaction of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ (150 μM in CH_2Cl_2 , black) with 10 equivalents of PhINNs. (B) ESI-HRMS⁻ spectrum (black) and simulated spectrum (red) for $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$. (C): UV-Vis spectrum of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ (blue), formed by addition of 10 equivalents of PhINNs to $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ (78 μM in CH_2Cl_2 , black). (D) ESI-HRMS⁻ spectrum (black) and simulated spectrum (blue) for $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$.

As can be expected for an integer spin system, the anionic bis-nitrene complex $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$ is X-band EPR silent, both at r.t. and at 10 K. The neutral mono-nitrene complex $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ displays an isotropic EPR signal (Figure 7A) at $g_{\text{iso}} = 2.091$ at r.t., showing well-resolved ^{59}Co ($A_{\text{Co}}^{\text{Co}} = 89.5$ MHz) and poorly resolved (but necessary for accurate simulation) ^{14}N ($A_{\text{N}}^{\text{N}} = 18.9$ MHz) HFI. The anisotropic low temperature (20 K) EPR spectrum of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ recorded in a toluene glass displays a slightly rhombic signal with a small g -anisotropy and multiple hyperfine coupling interactions, consistent with a net-doublet ground state (Figure 7B).⁴² The r.t. EPR spectrum of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NTs})]$ proved to be similar to that of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ (see SI).

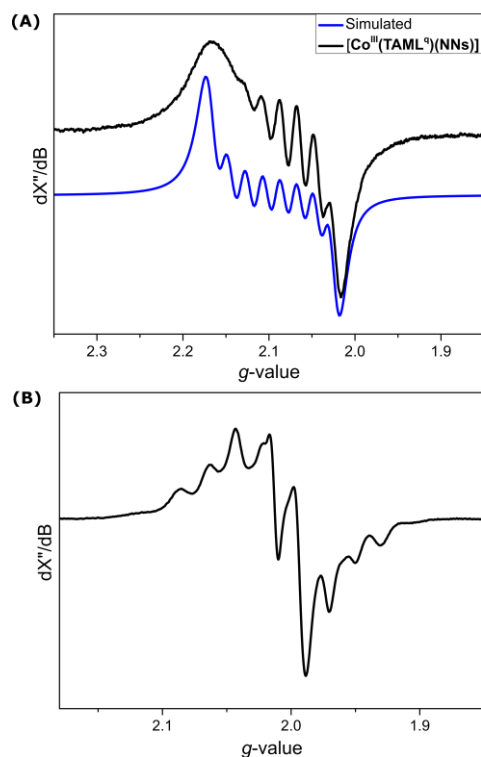


Figure 7. (A): Experimental (black) and simulated (blue) X-band EPR spectrum of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ at r.t. in toluene. Microwave freq. 9.3716 GHz, mod. amp. 2.000 G, power 6.325 mW. Simulated parameters: $g_{\text{iso}} = 2.091$, $A_{\text{Co}}^{\text{Co}} = 89.5$ (34.2) MHz and $A_{\text{N}}^{\text{N}} = 18.9$ MHz. (B): X-band EPR spectrum of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ in toluene glass at 20 K. Microwave freq. 9.376 GHz, mod. amp. 2.000 G, power 6.325 mW.

The Co K-edge XANES spectra for $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ are shown in Figure 8. As was observed for the parent complexes $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$, the edge position for both cobalt-nitrene complexes is detected at 7721 eV, suggesting that the cobalt centers in all four complexes have the same overall +III oxidation state. Interestingly, the intense shoulder absorption at 7715 eV observed in the spectra of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ (corresponding to $1s \rightarrow 4p + \text{LMCT}$ shakedown transitions characteristic for square planar cobalt complexes), is no longer visible in the nitrene adducts, thus suggesting that both complexes undergo changes in coordination number and/or geometry. Solely based on Co K-edge XANES data, no differentiation between a

five- or a six-coordinated cobalt complex can be made. This was also observed in related $\text{Co}^{\text{III}}(\text{porphyrin})$ -mono- and bis-nitrene complexes that displayed an octahedral coordination environment, with an axial co-ligand (NsNH_2 , NsNH^- , H_2O , or solvent) present in case of the mono-nitrene species.^{14b} Moreover, an additional low intensity pre-edge feature at 7711 eV is observed clearly for $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ (insert in Figure 8). The pre-edge feature in the XANES spectrum of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ is not well-resolved due to moderate data quality caused by low solubility of the complex. These pre-edge features arise from $1s \rightarrow 3d$ transitions, and in centrosymmetric (i.e. square planar and octahedral) complexes these transitions are weak due to quadrupole transitions.^{14b} However, symmetry breaking enables $3d-4p$ -hybridization of metal atomic orbitals, causing the pre-edge to gain intensity due to dipole-allowed transitions. It thus seems that $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ bears an unidentified sixth coordinating co-ligand (octahedral coordination geometry), but is not fully centrosymmetric. However, similar low-intensity pre-edge features have been observed in a five-coordinate cobalt-TAML complex²⁷ and therefore a square pyramidal coordination around cobalt cannot be fully excluded for $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$.

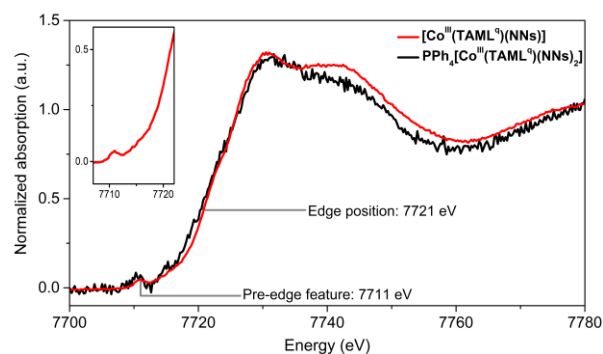


Figure 8. Co-K edge XANES analysis of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ (black) and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ (red) in toluene. Insert: zoom of the pre-edge feature for $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})(\text{NNs})]$.

Consistent with the abovementioned experimental results, DFT calculations (BP86, def2-TZVP, disp3, m4 grid) indicate that formation of the neutral mono-nitrene complex $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ ($S = 1/2$; $\Delta G_{298\text{K}}^{\circ} = -20.3$ kcal mol⁻¹) from $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ ($S = 1/2$; reference point) is energetically more favorable than formation of the neutral bis-nitrene adduct $[\text{Co}(\text{TAML})(\text{NNs})_2]$ ($S = 1/2$; $\Delta G_{298\text{K}}^{\circ} = -14.5$ kcal mol⁻¹). However, the corresponding formation energies of the anionic mono- and bis-nitrene complexes $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})(\text{NNs})]^-$ ($S = 1$; $\Delta G_{298\text{K}}^{\circ} = -27.9$ kcal mol⁻¹) and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$ ($S = 1$; $\Delta G_{298\text{K}}^{\circ} = -29.9$ kcal mol⁻¹) from $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ ($S = 1$; reference point) are nearly equal (see SI).

NEVPT2-corrected CASSCF calculations were performed to accurately describe the electronic structure of the nitrene species. All cobalt d -orbitals, ligand L_{π} and nitrene localized p -orbitals were included in the active spaces. CASSCF(14,13) calculations on $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]$ showed that the d_{xy} orbital is not preserved in the active space (occupancy 2.00)⁴⁰ and that the d_{z^2} orbital forms bonding (nitrene- N^1 and N^2 localized,

occupancy 1.94) and antibonding (mostly d_{xz} localized, occupancy 0.07) combinations with the nitrene N_{pz} orbitals. The d_{yz} and d_{xz} orbitals are both filled (occupancies 1.97 and 1.95, respectively) and the L_{π} - d_{xz} (occupancy 0.10) is virtually empty. Given that the L_{π} orbital was doubly-filled in $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ (*vide supra*), this implies that formation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$ from $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ is associated with two-electron oxidation of the ligand. Interestingly, both nitrene nitrogen atoms bear a single unpaired electron in their N_{py}/N_{px} orbitals (both occupancies 1.00). The electronic structure is thus best described as $(d_{xy})^{2.00}(d_{yz})^{1.97}(d_{xz})^{1.95}(N_{pz}^1 + N_{pz}^2 + d_{xz})^{1.94}(N_{px}^2)^{1.00}(N_{py}^1)^{1.00}$, consistent with a Co^{III} center, fully oxidized TAML backbone (TAML^{q}) and two one-electron reduced Fischer-type nitrene radical substrates ($\text{N}^{\bullet}\text{Ns}^-$).⁴³ Moreover, the cobalt(III) center has undergone a spin transition from intermediate spin in $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ to low spin in $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns}^-)_2]^-$ upon formation of the bis-nitrene radical species. As a result, the net total spin state does not change in the process, and remains a triplet spin state ($S = 1$). The most relevant active orbitals and their occupation numbers are shown in Figure 9A. In addition, excitation energies derived from the CASSCF(14,13) calculation revealed that no intense absorption bands are expected in the 400-850 nm region (see SI), consistent with the experimental spectrum depicted in Figure 6A.

The complex bears some resemblance to the previously reported cobalt-porphyrin bis-nitrene ($[\text{Co}^{\text{III}}(\text{TPP}^{\bullet})(\text{N}^{\bullet}\text{Ns}^-)_2]$)¹⁴ and ruthenium-porphyrin bis-imido ($[\text{Ru}^{\text{VI}}(\text{TPP})(\text{NTs})_2]$)⁴⁴ complexes (TPP = tetraphenylporphyrin). The ruthenium bis-imido complex is formed exclusively via metal-centered oxidation processes. However, while in the cobalt-porphyrin complex double nitrene-radical formation is the result of combined *metal-to-substrate* and (porphyrin) *ligand-to-substrate* SET processes, formation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns}^-)_2]^-$ is an *entirely (double) ligand-to-substrate* single-electron transfer process.

In a very similar fashion, CASSCF(13,12) calculations on $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$ reveal π ($d_{yz}+N_{py}$) and σ ($d_{xz}+N_{pz}$) bonding interactions between cobalt and the nitrene, with occupations of 1.93 and 1.86 electrons respectively.⁴⁵ The d_{xz} orbital is filled (occupancy 1.91) and the formerly half-filled L_{π} - d_{xz} orbital is now unoccupied (occupancy 0.12), indicating single-electron oxidation of the ligand (i.e. from TAML^{sq} to TAML^{q}). The single unpaired electron of the complex is mainly localized on the nitrene moiety (N_{py} - d_{yz} , occupancy 1.06), again consistent with $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns}^-)]$ being a Fischer-type nitrene radical complex with a net π -bond order between cobalt and the nitrene of ~ 0.5 .⁴³ As for the anionic bis-nitrene complex, the neutral mono-nitrene complex is generated via *ligand-to-substrate* SET. Once again the cobalt(III) ion does not change its oxidation state in the process, but it does undergo a spin flip from intermediate spin in $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ to low spin in $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns}^-)]$. The most relevant active orbitals and their occupations are shown in Figure 9B. Notably, neither $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns}^-)_2]^-$ nor $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{N}^{\bullet}\text{Ns}^-)]$ show significant multi-reference character.

Interestingly, ligand-to-substrate SET combined with a metal-based spin flip effectively leads to a shift of the spin density from the metal to the nitrene nitrogen(s) in both the neutral

mono-nitrene and the anionic bis-nitrene complexes, without oxidation of cobalt and without changing the net total spin state of the complex. The redox events clearly occur on the TAML backbone (electron donor) and the nitrene (electron acceptor), wherein the former undergoes one-electron or two-electron oxidation to accommodate one or two nitrene radical(s) on the Co^{III} -center.

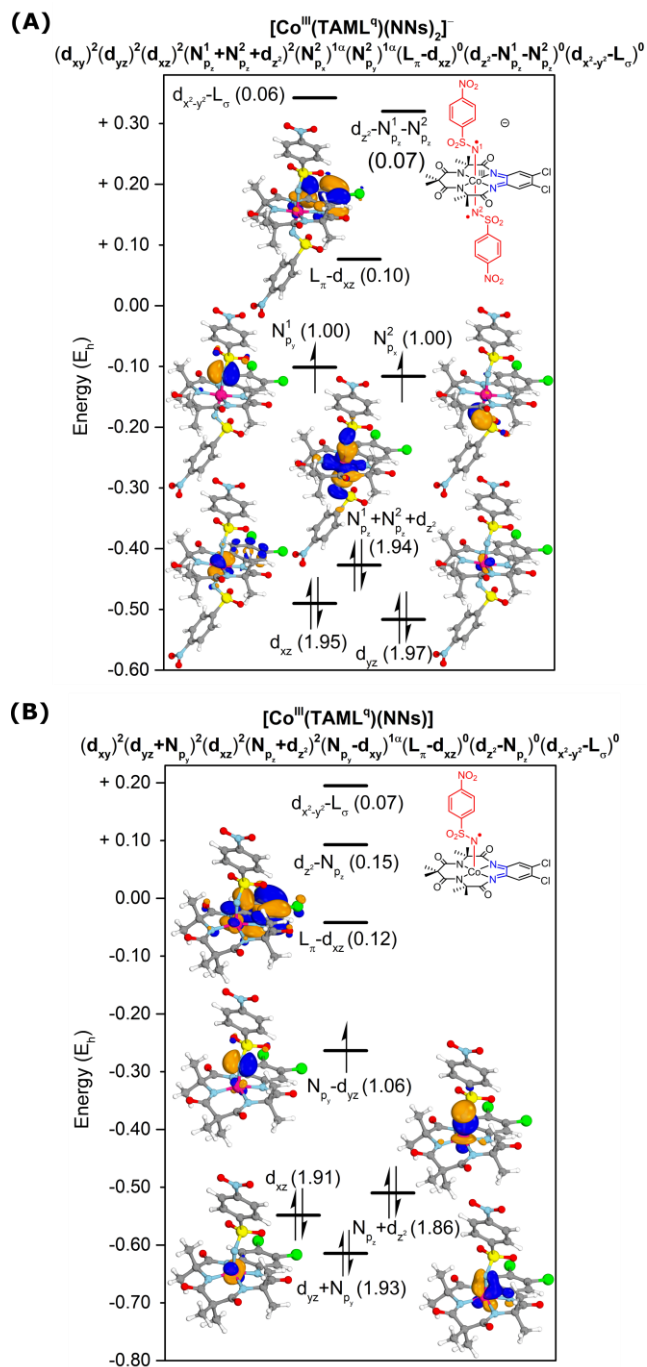
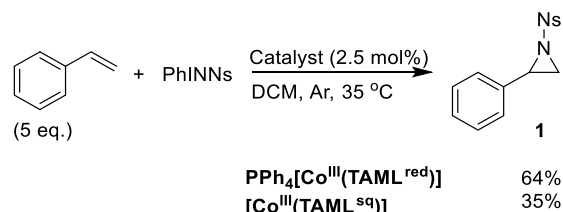


Figure 9. Most relevant active orbitals and occupancies (in parenthesis) of NEVPT2-corrected CASSCF(14,13) ($[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$, top) and CASSCF(13,12) ($[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$, bottom) calculations.

Intrigued by the influence of the ligand oxidation state on the structure of the nitrene species, the mono- and bis-nitrene

species were probed for catalytic nitrene transfer reactivity in the benchmark aziridination of styrene (Scheme 5).^{1a,3} A remarkable difference in yield of aziridine product **1** was observed when using $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ (64%) or $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ (35%) as the catalyst in nitrene transfer reactions from PhINNs to styrene, suggesting that the anionic bis-nitrene and neutral mono-nitrene exhibit markedly different activity and/or stability properties. A thorough investigation on the applicability and mechanisms of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ as aziridination catalysts is the subject of current investigations, that will be reported in due time. At this point it is worth mentioning that for cobalt-TAML complexes the *reduced (anionic)* $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ species are apparently more effective nitrene-transfer catalysts than the corresponding *oxidized (neutral)* $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ species, while for iron- and manganese TAML complexes the reverse was observed.^{19,20}

Scheme 5. Catalytic aziridination of styrene to afford **1 using $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$.**



CONCLUSIONS

In this work we have conclusively shown that the ligand in Co(TAML) complexes is redox-active. Oxidation of $[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]^-$ using $(\text{Thi})\text{BF}_4$ cleanly affords $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ via ligand-centered oxidation, with the electronic structure being best described as an intermediate spin ($S = 1$) cobalt(III) center that is antiferromagnetically coupled to a ligand-centered radical ($S = 1/2$).

Interestingly, cobalt-nitrene adducts of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ and $\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})$ can be cleanly generated from PhINNs via ligand-to-substrate single-electron transfer to afford $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})_2]^-$ and $[\text{Co}^{\text{III}}(\text{TAML}^{\text{q}})(\text{NNs})]$, respectively. CASSCF calculations revealed that both nitrene complexes are best described as one-electron reduced Fischer-type nitrene radicals. The formation of a bis-nitrene adduct of $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ is attributed to the availability of two electrons within the reduced TAML framework for double ligand-to-substrate SET, whereas only one electron can be used for ligand-to-substrate SET on $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$, which therefore affords the mono-nitrene adduct. Intriguingly, in both cases the combination of ligand-to-substrate SET and a spin flip from intermediate spin ($S = 1$) to

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low spin ($S = 0$) at the cobalt(III) center effectively results in a shift of the spin density from the metal to the nitrene moieties, without oxidation of cobalt and without changing the net total spin state of the complex.

Preliminary catalytic styrene aziridination reactions using $\text{PPh}_4[\text{Co}^{\text{III}}(\text{TAML}^{\text{red}})]$ or $[\text{Co}^{\text{III}}(\text{TAML}^{\text{sq}})]$ as the catalyst reveal remarkable differences in activity/stability between the two systems. More elaborate studies on the underlying mechanisms, synthetic applicability and differences between the two complexes in nitrene transfer catalysis will be reported in the near future.

ASSOCIATED CONTENT

Supporting Information. Experimental details, synthetic procedures, relevant NMR, EPR, HRMS, XRD, UV-Vis, electrochemical and XANES data, geometries (xyz coordinates) of stationary points (DFT) and description of the CASSCF calculations. This material is available free of charge via the Internet at <http://pubs.acs.org>.

AUTHOR INFORMATION

Corresponding Author

*Dr. Ir. J. I. van der Vlugt. j.i.vandervlugt@uva.nl

*Prof. Dr. B. de Bruin. b.debruin@uva.nl

Notes

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ABBREVIATIONS

CASSCF, Complete Active Space Self Consistent Field; DCM, dichloromethane; DFT, density functional theory; EPR, electron paramagnetic resonance; HFI, hyperfine interaction; Ns, nosyl; r.t., room temperature; TAML, tetra-amido macrocyclic ligand; XANES, X-ray near edge structure.

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