

Mesoionic Carbenes in Low- to High-Valent Vanadium Chemistry

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Cite This: *Inorg. Chem.* 2021, 60, 15421–15434

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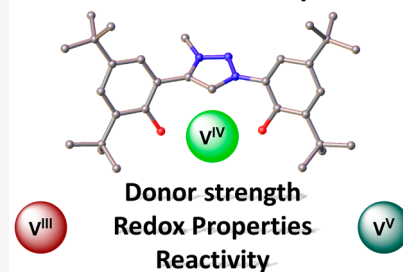
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ABSTRACT: We report the synthesis of vanadium(V) oxo complex **1** with a pincer-type dianionic mesoionic carbene (MIC) ligand **L**¹ and the general formula [VOCl(L¹)]. A comparison of the structural (SC-XRD), electronic (UV–vis), and electrochemical (cyclic voltammetry) properties of **1** with the benzimidazolinyldene congener **2** (general formula [VOCl(L²)]) shows that the MIC is a stronger donor also for early transition metals with low d-electron population. Since electrochemical studies revealed both complexes to be reversibly reduced, the stronger donor character of MICs was not only demonstrated for the vanadium(V) but also for the vanadium(IV) oxidation state by isolating the reduced vanadium(IV) complexes [Co(Cp*)₂][**1**] and [Co(Cp*)₂][**2**] ([Co(Cp*)₂] = decamethylcobaltocenium). The electronic structures of the compounds were investigated by computational methods. Complex **1** was found to be a moderate precursor for salt metathesis reactions, showing selective reactivity toward phenolates or secondary amides, but not toward primary amides and phosphides, thiophenols, or aryls/alkyls donors. Deoxygenation with electron-rich phosphines failed to give the desired vanadium(III) complex. However, treatment of the deprotonated ligand precursor with vanadium(III) trichloride resulted in the clean formation of the corresponding MIC vanadium(III) complex **6**, which undergoes a clean two-electron oxidation with organic azides yielding the corresponding imido complexes. The reaction with TMS-N₃ did not afford a nitrido complex, but instead the imido complex **10**. This study reveals that, contrary to popular belief, MICs are capable of supporting early transition-metal complexes in a variety of oxidation states, thus making them promising candidates for the activation of small molecules and redox catalysis.

MIC-Vanadium complexes



INTRODUCTION

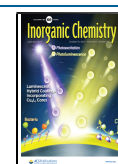
Almost two decades after the first report of an abnormal 5-imidazolinyldene carbene complex,¹ mesoionic carbenes have been developed into a distinguished ligand class.^{2,3} Among them, 1,2,3-triazole derived mesoionic carbenes, namely 1,2,3-triazolinyldenes,⁴ stand out by their modular synthesis via the copper-catalyzed [3 + 2] cycloaddition between azides and alkynes.^{5–7} After their initial reporting by Albrecht *et al.*,⁸ they quickly became prominent synthetic targets for (electro-) catalysis,^{9–23} supramolecular chemistry,^{24–27} magnetism,²⁸ and photochemistry due to their versatile synthesis and comparatively straightforward handling.^{29–36} Throughout these studies, a great effort has been made to decipher their electronic structure. Mesoionic carbenes are commonly believed to be strong σ -donor ligands paralleling heteroaryls; however, recent reports emphasize their π -accepting properties^{37–39} as demonstrated by the isolation of a reduced triazolinyldene ligand.⁴⁰ Nevertheless, most studies targeted hitherto late transition metals or main group elements, while early transition-metal complexes with mesoionic carbenes have been rarely explored.^{41–44} Arguably, this can be attributed to the relatively weak bond between N-heterocyclic carbenes and early transition metals.⁴⁵ However, this weak bond may be enhanced by harnessing anionic linkers. This strategy has allowed the isolation of a number of interesting metal

complexes^{46–49} of the early transition metals,^{50–71} the lanthanides,^{72–80} and the actinides.^{81–86}

Among the early transition metals, vanadium chemistry has witnessed a remarkable activity over the past 50 years and has been applied in heterogeneous and homogeneous catalysis,⁸⁷ small molecule activation,^{88–92} molecular magnetism,^{93–96} and spin qubits.^{97–100} Narrowing down the field to NHC vanadium complexes, since the first two reports of vanadium NHC complexes in 1994 by Roesky *et al.*¹⁰¹ and 2003 by Abernethy *et al.*,¹⁰² the utility of these complexes has mostly been explored in polymerization catalysis.^{103–107} Beyond this, only a few other applications of vanadium-NHC complexes have been examined, including small molecule activation^{108,109} and the neutralization of chemical warfare agents.¹¹⁰ Still, most vanadium NHC complexes refer to diamagnetic vanadium(V) complexes, while low-valent vanadium complexes have been rarely investigated.^{101,106,111–114}

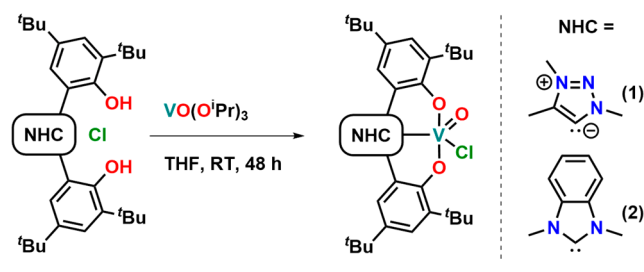
Received: July 9, 2021

Published: September 30, 2021



Inspired by Bellemin-Lapponnaz's ligand design, employing two anionic redox-active phenolate linkers,^{50–54,115–117} we have recently reported the first mesoionic carbene complexes of groups IV, V, and VI based on a 1,2,3-triazolinylidene scaffold.⁴⁴ Our initial report focused on niobium as a group V representative, and we thus expand this chemistry herein toward vanadium. We report the ligand's σ -donor strength by comparing the structural, spectroscopic, and electrochemical properties of the triazolinylidene complex **1** with its benzimidazolinylidene (benzNHC) congener **2**, proving the triazolinylidene ligand **L**¹ as the stronger donor. Furthermore, the salt metathesis reactivity of the new triazolinylidene complex **1** is presented, revealing moderate scope. While phenols and secondary amides give good and clean conversion, all other nucleophiles investigated gave no tractable reaction products. We furthermore report on the isolation of various vanadium complexes in the oxidation states +IV and +III, where the latter are potent precursors to vanadium(V) imido complexes.

Scheme 1. Synthesis of the Vanadium(V) NHC Complexes Following Protonolysis between the Corresponding Azolium Salts and VO(OⁱPr)₃



RESULTS AND DISCUSSION

Despite our previous finding that protonolysis between the triazolium salt [H₃L¹][Cl] and Ti(OⁱPr)₃Cl did not lead to quantitative deprotonation of the triazolium salt,⁴⁴ we decided to adopt this strategy using VO(OⁱPr)₃ as the vanadium source. To our delight, the reaction between the triazolium salt [H₃L¹][Cl] and VO(OⁱPr)₃, followed by the subsequent washing of the crude solids with hexane, afforded the MIC vanadium-oxo complex **1** as a dark green powder in yields of 85% (Scheme 1). The ¹H NMR spectrum of **1** in benzene confirmed the desired transformation due to the absence of the OH and triazolium-5H protons which revealed a C₁ symmetric species in solution. Unfortunately, due to the high quadrupolar moment of the ⁵¹V nucleus, we were not able to observe the characteristic ¹³C NMR resonance of the triazolinylidene carbon atom. Nevertheless, the absence of the triazolium-5C resonance at 131.9 ppm in the ¹³C NMR of **1** confirms the formation of a triazolinylidene complex of high-valent vanadium(V). Furthermore, a shift of the ⁵¹V resonance to –533 ppm (Figure S5) in the ⁵¹V NMR indicates the presence of a strong donor ligand. To set this value into context, we also synthesized the benzimidazolinylidene complex **2**, recently reported by LeRoux et al.^{118,119} The proton NMR of **2** shows a C_s symmetric species in solution where the absence of the OH and the benzimidazolium-2H protons indicates also the

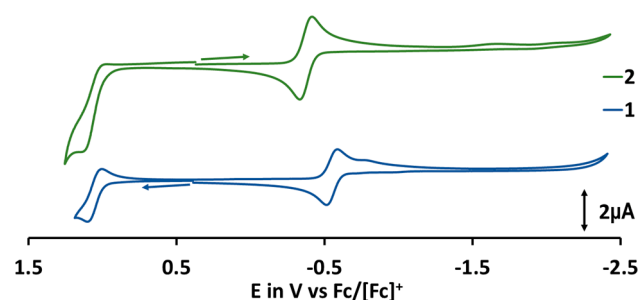


Figure 1. Cyclic voltammogram of **1** (blue) and **2** (green) in 0.1 M NBu₄PF₆ in CH₂Cl₂ at 298 K. Scan rate: 100 mV s⁻¹.

Table 1. Oxidation and Reduction Potentials of the Vanadium(V)-NHC Complexes **1 and **2** Referenced vs Fc/[Fc]⁺ in 0.1 M NBu₄PF₆ Solutions in CH₂Cl₂ at 298 K**

complex	E ^{1/2} ox. (V)	ΔE (mV)	E ^{1/2} red. (V)	ΔE (mV)
1	1.05	80	–0.56	70
2	n.a.	n.o.	–0.37	90

formation of an NHC complex of vanadium(V). Similar to **1**, we could not observe the carbene carbon resonance in the ¹³C NMR spectrum of **2**. The ⁵¹V NMR signal of **2** is (compared to complex **1**) strongly shifted to lower fields resonating at –503 ppm (Figure S10), indicating a lower donor strength of the benzimidazolinylidene compared to the triazolinylidene ligand.¹²⁰ Unambiguous proof for the formation of the NHC and MIC complexes was obtained by single-crystal X-ray diffraction (SC-XRD) analysis. X-ray quality crystals of **1** and **2** could be grown by slow diffusion of pentane into a concentrated solution of the corresponding complexes in toluene or benzene, respectively (Figure 2). Complex **1** crystallized in the monoclinic space group P2₁/n as a toluene solvate, while complex **2** crystallized without additional solvent molecules in the asymmetric unit in the orthorhombic space group Pbc_a. The most striking difference in the molecular conformations of the complexes is that in the case of **2**, the benzannulated heterocycle is shifted out of plane compared to the triazolinylidene complex **1**, this pitch angle along the C1–V1 bond axis was found to be only 0.7(1)°. These structural parameters are well reproduced by density functional theory (DFT) calculations (Table S1) and are also discernible in the coordination environment around the vanadium center. While **1** adopts an almost perfect square pyramidal coordination environment in the solid state (τ₅ = 0.05), complex **2** is distorted with τ₅ = 0.20. The C1–V1 distances in **1** and **2** are 2.055(3) Å and 2.131(3) Å, suggesting a stronger metal carbene interaction in **1** compared to **2**. This agrees with the pronounced high-field shift of the ⁵¹V NMR resonances in **1** relative to **2**. The V1–O10 distances were determined to be 1.583(3) Å and 1.585(2) Å in **1** and **2**, showing only a minor influence of the NHC moiety toward the strength of the V=O bonds. However, the influence of the NHC unit on the V=O stretching frequencies in the IR is discernible with resonances at 986 cm⁻¹ (calculated: 1026 cm⁻¹) and 1000 cm⁻¹ (calculated: 1035 cm⁻¹) in **1** and **2**. These values are indicative for a weaker V=O multiple bond character in **1** in comparison to **2** due to arguably reduced π-donation from the terminal oxo ligand, and thus, corroborate stronger donor properties of the MIC ligand.

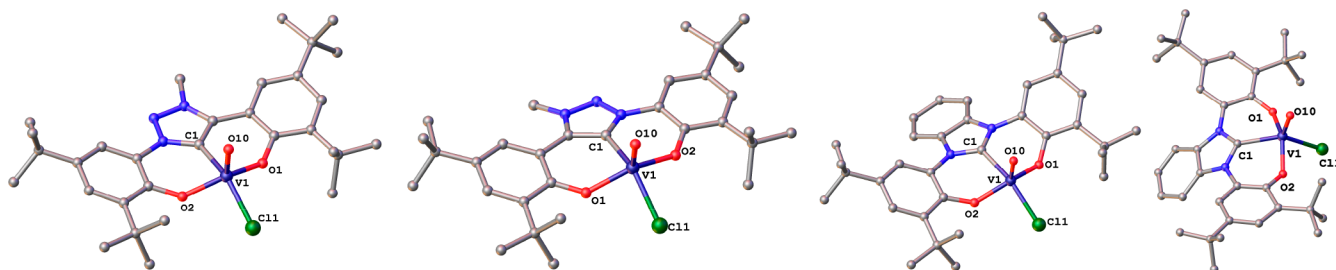
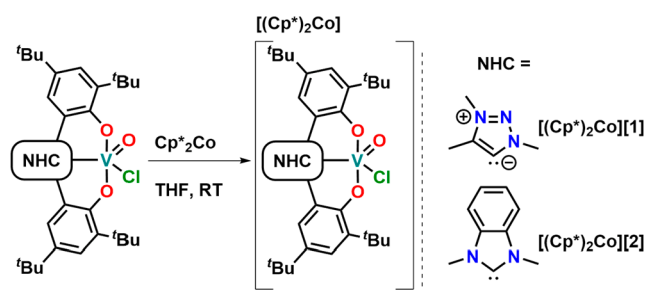


Figure 2. Molecular structures of **1**, $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$, **2**, and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$ (from left to right). Hydrogen atoms, solvent molecules, and counterions have been omitted for clarity.

Scheme 2. Reduction of Complexes **1** and **2** by Decamethylcobaltocene



To further probe the donor properties of the triazolin-versus the benzimidazolinyldene donor, we investigated the complexes by electrochemical methods. Cyclic voltammetry measured in dichloromethane revealed a reversible reduction corresponding to the V(V/IV) redox couple for both complexes (Figure 1). In agreement with the results from ^{51}V NMR and IR spectroscopy, the reduction potential (the reductions are vanadium centered, *vide infra*) for **1** appears 0.19 V cathodically shifted compared to the reduction potential of **2** (Table 1). This suggests a higher electron density at the vanadium center in **1**, which is in line with the higher σ -donor character of MIC relative to benzNHC ligands. Additionally, complex **1** showed one reversible (ligand-centered) oxidation, whereas for **2** it is at the edge of the solvent window and thus could not be evaluated (Figures S69–S72).

To reveal the site of reduction for the two vanadium complexes, we reduced the complexes with decamethylcobaltocene (Scheme 2). While a THF solution turned greyish upon reduction of **1**, the desired product precipitated as a bright green powder for **2**. Evans method in CD_2Cl_2 revealed a magnetic moment of 1.74 and 1.67 μ_B for $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$, which is in agreement with a vanadium(IV) redox state.¹²¹ Accordingly, the V=O stretches shift from 986 to 932 cm^{-1} (calculated: 1026 to 1007 cm^{-1}) and from 1000 to 968 cm^{-1} (calculated: 1035 to 1007 cm^{-1}) in $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$, respectively (see Figures S49–S52). X-ray quality crystals of both reduced complexes $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$ could be grown by the slow evaporation of dichloromethane out of a hexane/dichloromethane mixture (Figure 2).

While the general structural factors (e.g., coordination environment) resemble the same trends as the parent vanadium(V) complexes **1** and **2**, the vanadium donor atom distances increase slightly. For example, the vanadium carbene distances expand from 2.055(3) Å to 2.070(3) Å and from 2.131(3) Å to 2.153(7) Å during the reduction to

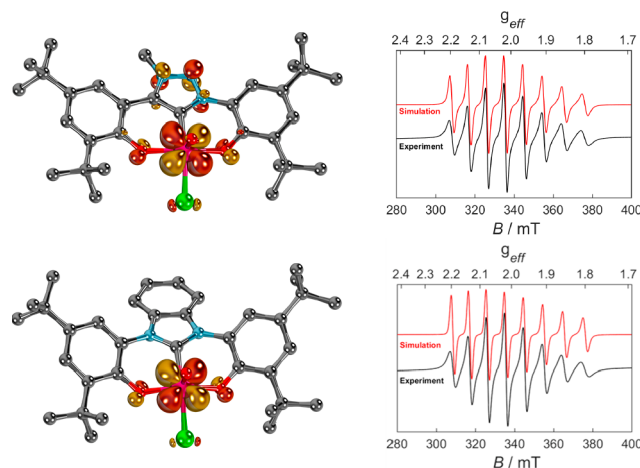
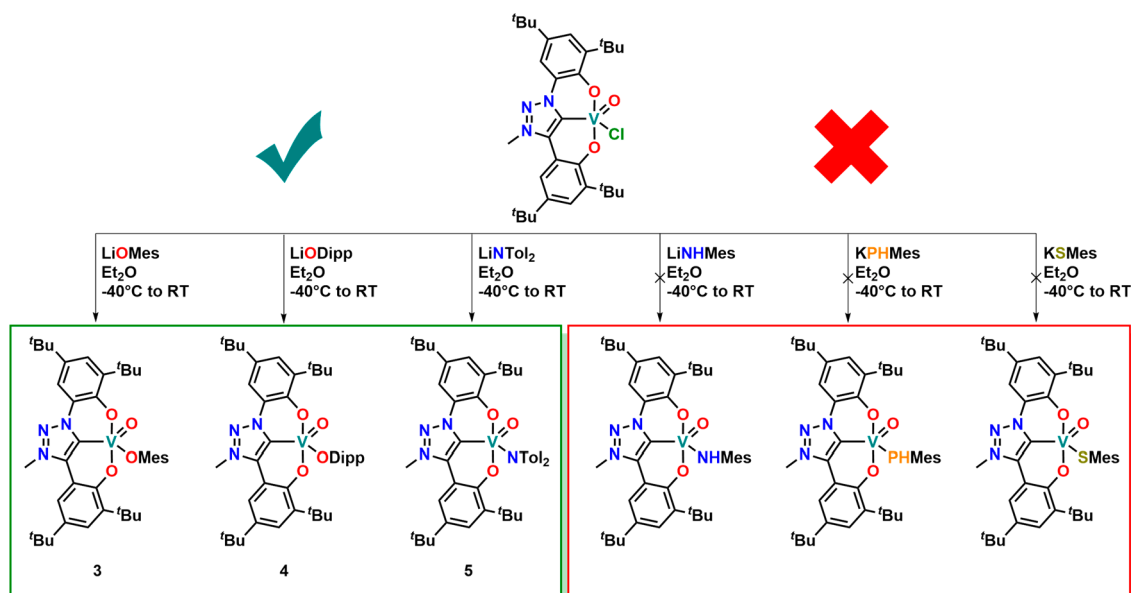


Figure 3. Compounds $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ (top left) and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$ (bottom left) feature unpaired electrons in the 3d(yz) orbitals (QROs). X-band EPR spectra of $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ (top right) and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$ (bottom right) of 5 mM solutions in CH_2Cl_2 at 300 K; black traces show the experimentally observed spectra and red traces the corresponding simulations. Hydrogen atoms have been omitted for clarity.

$[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$, respectively. Similarly, the phenolate distances elongate by almost 0.1 Å (compare Table S3). This is in line with the larger ionic radius of a vanadium(IV) compared to a vanadium(V) ion, which suggests the reduction is vanadium centered. The center of the redox processes was further corroborated by EPR spectroscopy (Figure 3, right). Both complexes, $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$ show a characteristic eight-line spectrum, which is consistent with a single unpaired electron located at the metal (^{51}V , 99.75% natural abundance, $I = 7/2$). Values of $g_{\text{iso}} = 1.9715$ and $g_{\text{iso}} = 1.9666$ and $a = [274.1915, 268.8744, 258.6040 \text{ MHz}]$ and $a = [256.8, 267.7, 261.9 \text{ MHz}]$ for $[\text{Co}(\text{Cp}^*)_2][\mathbf{1}]$ and $[\text{Co}(\text{Cp}^*)_2][\mathbf{2}]$, respectively, are comparable with other vanadium(IV) complexes, for example, a four-coordinate vanadium alkylidene.¹²¹ The electronic structure was corroborated by scalar relativistic DFT calculations at the ZORA-PBE-D3BJ/def2-TZVPP//ZORA-PBE-D3BJ/def2-SVP level of theory,^{122–132} which indicate a vanadium centered SOMO (quasi-restricted orbital QRO, Figure 3, left) with only small orbital overlap with the supporting ligand.

Further evidence for the stronger donor character of the mesoionic carbene ligand L^1 compared to the benzimidazolinyldene L^2 can be extracted from UV–vis spectroscopy. When changing from the MIC complex **1** to the benzNHC complex **2**, the charge-transfer (CT) band located at 387 nm for **1** shifts

Scheme 3. Salt Metathesis Reactivity of Complex 1 toward Various Chalcogen and Pnictogen-Based Nucleophiles as Well as Aryl Anion Donors



to 420 nm for **2**. Based on time-dependent DFT calculations (Figures S74–S89), we assign this band as essentially ligand-to-ligand charge transfer (LLCT) from the phenolate moieties to the NHC bridge, with only minor contribution from the metal. As the benzNHC is a stronger acceptor ligand compared to the MIC, this band is red-shifted in **2** compared to **1**.¹³³ The broad bands located between 500 and 800 nm in **1** and **2** are assigned to the respective ligand-to-metal charge transfers (LMCTs). This assignment is in line with their disappearance in the reduced complexes [Co(Cp*)₂][**1**] and [Co(Cp*)₂][**2**] (see Figure 4). Overall, the results from NMR, electrochemistry, and UV–vis absorption spectroscopy support the notion of MICs being stronger donors than benzNHCs in vanadium complexes.

To further explore the chemical potential of the triazolinylidene complex **1**, we turned our interest toward salt metathesis replacing the remaining chloride ligand (Scheme 3). Mixing the parent lithium salts with **1** in

–40 °C cold diethyl ether, followed by recrystallization from hexane, gave the pure mesitolate (**3**), 2,6-diisopropylphenolate (**4**), and 4,4'-ditolylamide (**5**) complexes in good to moderate yields. To our surprise, the reaction with primary amides (LiNHMes), thiophenolates (KSMeS), and phosphanides (KPHMes) failed to give well-defined products under the above-described conditions. Similarly, anionic alkyl or aryl donors, unrelated to their source (lithium or Grignard reagents), resulted in complicated reaction mixtures from which no defined reaction products could be isolated. The NMR spectroscopic and structural analyses (Figure 5) of the

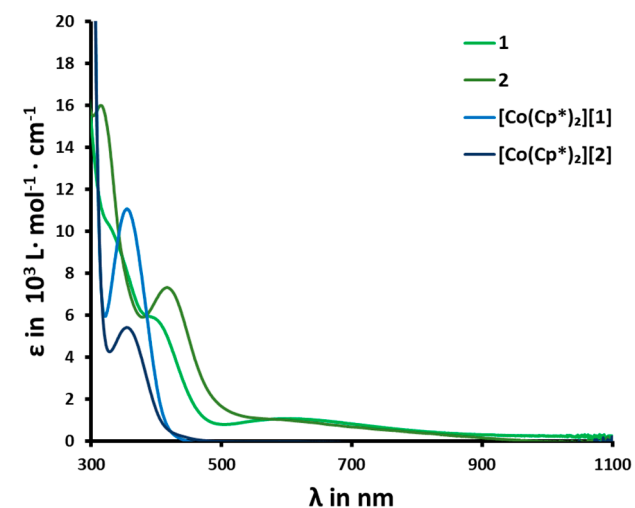


Figure 4. Stacked UV–vis spectra of the complexes **1**, **2**, [Co(Cp*)₂][**1**], and [Co(Cp*)₂][**2**] in CH₂Cl₂ at 298 K.

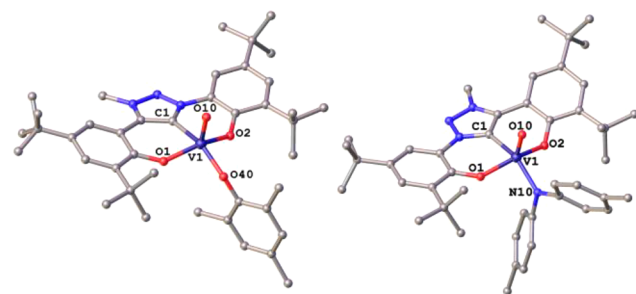
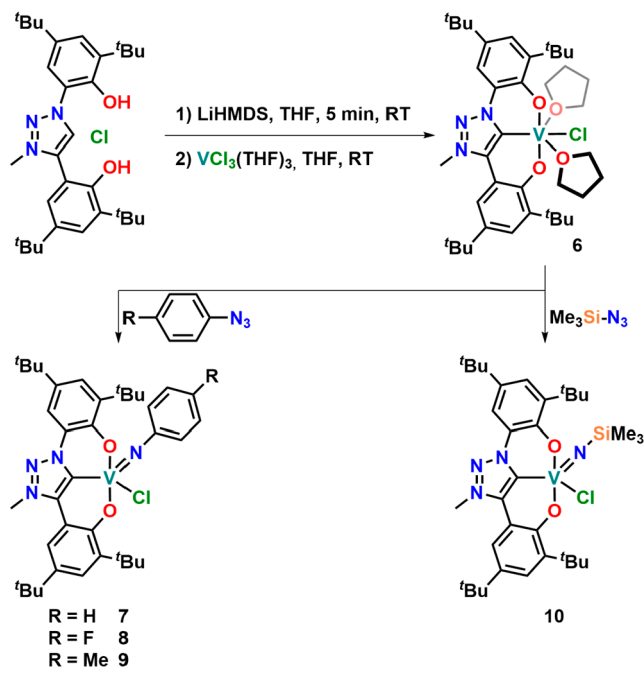


Figure 5. Molecular structures of **3** (left) and **5** (right). Hydrogen atoms and lattice solvent molecules have been omitted for clarity.

complexes **3**–**5** resemble the expected characteristics. For further information we refer to Figures S11–S25 and Tables S2 and S3 in the Supporting Information. Notably, we found that the coligand had a strong influence on the UV–vis spectroscopic features of the complexes. While the halide complex **1** was deep green ($\lambda_{\text{max}} = 582 \text{ nm}$; $\epsilon = 1000 \text{ L mol}^{-1} \text{ cm}^{-1}$), the phenolate complexes **3** and **4** are a deep purple color ($\lambda_{\text{max}} = 534 \text{ nm}$; $\epsilon = 4400 \text{ L mol}^{-1} \text{ cm}^{-1}$ for complex **4**), and the amide complex **5** is dark teal ($\lambda_{\text{max}} = 660 \text{ nm}$; $\epsilon = 7200 \text{ L mol}^{-1} \text{ cm}^{-1}$, compare also Figure S68).

By further probing the versatility of complex **1**, we examined oxo-exchange reactions, with emphasis on generating vanadium imido complexes. To reach this goal, we applied

Scheme 4. Synthesis of a Vanadium(III) Triazolinylidene Complex and Its Subsequent Conversion into Aryl and Alkyl Imido Complexes



isocyanates as the imido source, liberating carbon dioxide to provide the driving force for this process. Although this strategy was successful for a plethora of vanadium imido complexes,^{134,135} in the present case, even at elevated temperatures (60, 80, or 120 °C), no conversion could be observed. In one experiment (120 °C) with 3,5-bis-(trifluoromethyl)phenyl isocyanate, we observed its cyclotrimerization to yield the corresponding isocyanuric amide. It has been reported that electropositive metals, that is, Lewis acids, catalyze this process.^{136–140} However, NHCs and NHOs are also known to be potent catalysts for this transformation.^{141,142} As the presence of free carbenes at elevated temperatures (e.g., due to minimal thermal decomposition) cannot be fully ruled out, the catalytically active species remains unclear.

Since the direct generation of imido complexes from the oxo complexes had failed, we sought other strategies. Another versatile access to metal imido complexes is the direct reduction of organic azides by low-valent metal complexes. Thus, we initially aimed to generate a low-valent vanadium(III) complex by deoxygenating the parent vanadium(V) complex **1** with triethylphosphine. Although

this strategy met with success for the deoxygenation of molybdenum(VI) benzimidazolinylidene complexes,^{143,144} no useful reaction products could be isolated in the present case. Similarly, switching to other phosphines such as triphenylphosphine or trimethylphosphine turned out to be unproductive. We consequently turned our focus to installing the triazolinylidene ligand **L**¹ directly on vanadium(III).¹¹³ Although our attempts to isolate the fully deprotonated and free triazolinylidene $\text{Li}_2[\text{L}^1]$ have failed so far, deprotonation of $[\text{H}_3\text{L}^1][\text{Cl}]$ with LiHMDS (HMDS = hexamethyldisilazide) at room temperature, followed by the immediate addition of the deprotonated triazolinylidene to $\text{VCl}_3(\text{THF})_3$ resulted in the formation of a brown suspension. After extraction with toluene and washing of the crude solids with hexane, we isolated the desired vanadium(III) complex **6** as an orange powder in 46% yield (Scheme 4). The complex is remarkably sensitive toward air and moisture and decomposes in the glovebox at room temperature within several days even in the solid state. However, it turned out to be stable for a couple of weeks at -40 °C. The formation of the desired vanadium(III) complex was initially evident by the strong paramagnetic nature of its ¹H NMR, revealing an effective magnetic moment of $2.71 \mu_{\text{B}}$, which is in line with the presence of a d^2 electron configuration and is comparable to previously reported vanadium(III) complexes.¹⁰⁹ Despite numerous attempts and due to the high sensitivity of the complex, only low-quality crystals of the complex could be obtained from concentrated diethyl ether solutions at -40 °C. In any case, the connectivity of the molecule was unambiguously determined, confirming the vanadium(III) oxidation state (Figure 6, left). In addition to the triazolinylidene and the halide ligands, complex **6** was found to further hold two tetrahydrofuran donors, creating an octahedral coordination environment around the vanadium center. As expected for an $S = 1$ spin system, the EPR spectrum at room temperature did not reveal any observable signals (Figure S90).^{121,145} Computational investigation revealed both SOMOs to be vanadium centered (Figure 7) with only small orbital overlap with the supporting ligand. Accordingly, and in agreement with the experiment, these calculations indicate an adiabatic singlet–triplet gap of $\Delta E = -47 \text{ kJ mol}^{-1}$ in favor of the triplet state. Note that the prediction of two SOMOs is consistent with two irreversible waves of **6** observed in the cyclic voltammogram at 0.09 and 1.03 V vs Fc/Fc^+ in acetonitrile (Figure S73).

Having the vanadium(III) complex **6** in hand, we turned back our interest to the generation of vanadium(V) imido complexes following an azide reduction strategy. As envisioned, complex **6** reacts smoothly with organic azides such as phenyl, 4-fluorophenyl, or 4-methylphenyl azide to

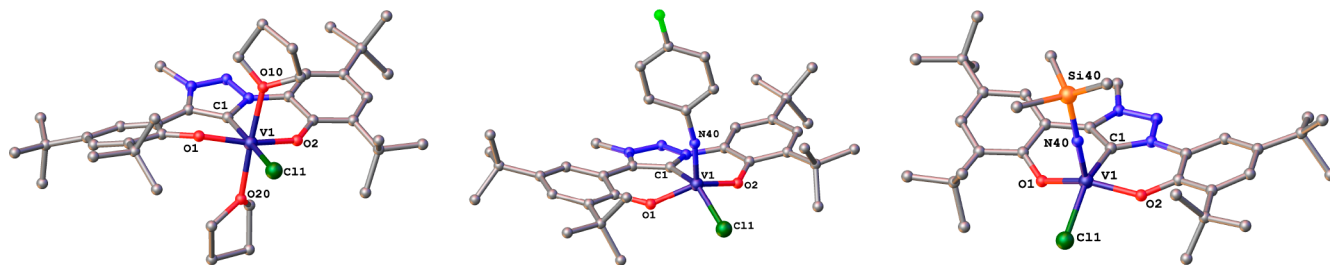


Figure 6. Molecular structures of the vanadium(III) complex **6** and the imido complexes **8** and **10** (from left to right). Hydrogen atoms and lattice solvent molecules have been omitted for clarity.

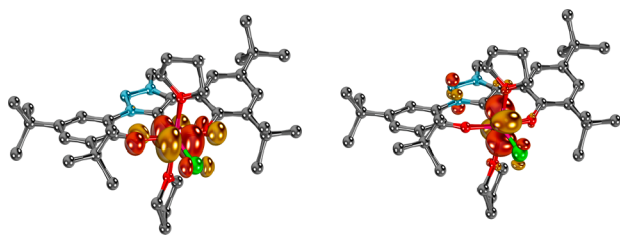


Figure 7. Compound **6** features unpaired electrons in the $3d(xz)$ and $3d(xy)$ orbitals (QROs). Hydrogen atoms have been omitted for clarity.

form the corresponding vanadium(V) imido complexes **7**, **8**, and **9**. Upon addition of the azides to benzene solutions of complex **6** and heating the samples to $60\text{ }^{\circ}\text{C}$, gas evolution was observed, and the reddish-brown solution gradually changed to deep green. The reaction could be followed by ^1H NMR spectroscopy, revealing the clean conversion to the desired complexes, which can be isolated by evaporation of benzene and washing the green powders with hexane in yields of 46–100% (Scheme 4). While the proton resonances of the complexes show the expected characteristics, the ^{13}C carbene resonance could not be observed again due to the large quadrupole moment of the ^{51}V nucleus. Similar problems occurred during our attempts to observe the imido nitrogen atom using ^1H – ^{15}N HMBC NMR experiments. However, in the ^{51}V NMR, the complexes show resonances at -398.0 , -396.8 , and -378.2 ppm for **7**, **8**, and **9** respectively (Figures S32, S38, and S43). To unambiguously prove the structural identity, X-ray quality crystals of complex **8** were grown from concentrated diethyl ether solution (Figure 6, middle). The complex crystallizes in the monoclinic space group $P2_1/n$ with one molecule in the asymmetric unit. The vanadium center is penta-coordinate by the ligand, the imido nitrogen atom, and a chlorido ligand in a slightly distorted square pyramidal coordination environment ($\tau_5 = 0.10$). The V1–N40 distance was found to be $1.644(2)$ Å and is comparable to previously reported vanadium(V) imido complexes. The metal carbene distance V1–C1 was found to be $2.048(2)$ Å, which compares well to complex **1** and can be found in the Supporting Information in Tables S2 and S3.

In sight of the swift reduction of organic azides, we next turned our interest toward the use of “inorganic” azides, which we envisioned to form vanadium nitrido complexes. These complexes are relevant intermediates in the context of dinitrogen activation and valorization. Indeed, reacting complex **6** with 1 equiv of TMS- N_3 resulted in the clean conversion to diamagnetic **10** at $60\text{ }^{\circ}\text{C}$ (Scheme 4). After workup, the ^1H NMR spectrum of the new complex shows a single resonance at -0.31 ppm integrating nine protons, which is indicative of the remainder of the TMS group on the nitrogen atom and thus the formation of a TMS-imido complex. Indeed, Mendiola and co-workers recently reported that the cleavage of a TMS group from a TMS-imido ligand to form the corresponding nitride complex is not trivial also for other group 5 metals, for example, tantalum(V).¹⁴⁶ The ^{51}V -NMR shows a single resonance at -473.5 ppm (Figure S48) which is high-field shifted compared to the aryl-imido complexes **7**–**9**. However, this high-field shift can be attributed to the stronger donor character of alkyl/TMS imido versus and

aryl imido ligands. Unfortunately, due to the quadrupolar moment of vanadium, the detection of both the imido nitrogen atom as well as of the silicon atom using ^{29}Si NMR spectroscopy failed. Nevertheless, X-ray quality crystals of complex **10** could be grown from slow evaporation of a concentrated diethyl ether solution at room temperature (Figure 6, right). The structural analysis confirmed the formation of the terminal imido instead of the desired nitrido complex. The structural properties of complex **10** resemble those of complex **8** and are given in the Supporting Information in Tables S2 and S3. In contrast to previous reports,¹⁴⁷ applying sodium azide as a nitrogen/azide source resulted in complicated (paramagnetic) mixtures, from which no defined products could be isolated.

CONCLUSIONS

We have extended the use of mesoionic carbenes with an early transition metal, vanadium. Using combined spectroscopic, electrochemical, and computational methods, we have shown that mesoionic carbenes are stronger donors than classical NHCs in early transition-metal chemistry as well. The high-valent oxo-vanadium(V) complexes are of moderate use for salt metathesis, reacting cleanly only with phenolates and secondary amides. Remarkably, the mesoionic carbene ligand supports vanadium in three oxidation states (III/IV/V). This is a rare report of a structurally characterized low-valent vanadium(III) complex supported by an NHC ligand^{112–114,148} and the first of a mesoionic carbene stabilizing a low-valent early transition metal. Complex **6** is a powerful two-electron reductant and forms the corresponding high-valent vanadium(V) imido complexes with azides. In recent years, NHC-imido vanadium complexes have attracted a large interest as polymerization catalysts as well as in nitrene transfer reactions.¹⁴⁹ Following the concept of extreme π -loading effects¹⁵⁰ and given the numerous examples of superior activity of MIC based catalysts over their NHC congeners,¹⁵¹ we believe that our findings will create further interest in the use of mesoionic carbenes in early transition-metal-mediated reactions. Additionally, the redox-active nature of the phenolate tethers will also be of large interest in other catalytic reactions as well as small molecule activation.

EXPERIMENTAL SECTION

General Remarks. If not otherwise mentioned, all transformations were carried out in an argon-filled glovebox under inert conditions. Solvents were dried by an MBraun SPS system and stored over activated molecular sieves (3 Å) for at least 1 day. C_6D_6 was dried over sodium/benzophenone and CDCl_3 and CD_2Cl_2 over calcium hydride, followed by vacuum transfer and three freeze–pump–thaw cycles. The proligand $[\text{H}_3\text{L}^1][\text{Cl}]$ was synthesized following a literature known procedure.⁴⁴ LiOMes, $\text{LiN}(\text{Tol})_2$, and LiNHMe were obtained by deprotonating the corresponding phenol or aniline in pentane using *n*-BuLi and filtering off the products. In a similar way, KSMes and KPHMes were obtained by deprotonating the corresponding thiophenol and primary phosphine using KHMDS in toluene.¹⁵² 4-Methylphenyl azide¹⁵³ and 4-fluorophenyl azide¹⁵⁴ were synthesized following previously reported methods using *tert*-butyl nitrite and trimethylsilyl azide in acetonitrile. Decamethylcobaltocene, triethylphosphine, trimethylsilyl azide, sodium azide, and $\text{VO}(\text{O}^i\text{Pr})_3$ were used as received by commercial suppliers. NMR spectra were collected at ambient temperature on a Bruker AV-300, Ascent 400, AV-500, or an Ascent 700 spectrometer. ^1H and ^{13}C NMR chemical shifts (δ) are reported in ppm and were calibrated to residual solvent peaks. ^{51}V NMR chemical shifts have been calibrated to VOCl_3 in CDCl_3 as an external standard. It needs to be mentioned

at this point that ^{51}V NMR is extremely sensitive and minor impurities (0.5% <) can still be observed, even though the remaining characterization data appear to be clean. This explains the minor impurities observed in the ^{51}V NMR spectra of the complexes **1**, **4**, **5**, **7**, **8**, **9** and **10**. Elemental analysis was performed using an Elementar vario microcube instrument. IR spectra were collected using a Bruker Alpha IR spectrometer. Cyclic voltammetry was recorded using a BioLogic potentiostat and a three-electrode array (working electrode: glassy carbon, counter electrode: platinum, reference electrode: silver). EPR spectra were recorded from 5 mM solutions at 300 K using a Bruker Magnetech 5000 EPR spectrometer (microwave frequency, 9.46 GHz; microwave power, 5 mW; modulation amplitude, 0.5 mT). CW spectra were processed using MATLAB and EasySpin software package (see Supporting Information for details).¹⁵⁵

Synthetic Procedures. General Procedure for the Synthesis of 1 and 2. The synthesis of the complexes was adapted from the literature.⁵⁴ If not otherwise stated, the corresponding azolium salt (1 equiv, 1 mmol) was mixed with $[\text{VO}(\text{O}^i\text{Pr})]$ (1 or 1.2 equiv) in THF (30 mL) and stirred at room temperature for 2 days. The solvent was evaporated, and the resulting solids were suspended in hexane (20 mL) and stirred for 1 h at room temperature. The resulting suspension was filtered, and the solids were washed with minimal amounts of pentane (10 mL) and dried on the frit inside the glovebox to give the desired vanadium(V) complexes in yields of 78% and higher.

$[\text{VO}(\text{L}^1)]$ (**1**). From $[\text{H}_3\text{L}^1][\text{Cl}]$ (1 equiv., 1 mmol, 528 mg) and $\text{VO}(\text{O}^i\text{Pr})_3$ (1 equiv., 1 mmol, 244 mg). Yield: 85% (0.85 mmol, 503 mg). ^1H NMR (C_6D_6 , 298 K, 700 MHz, in ppm): δ = 8.06 (d, J = 2.4 Hz, 1H, Aryl-H), 7.75 (d, J = 2.4 Hz, 1H, Aryl-H), 7.73 (d, J = 2.4 Hz, 1H, Aryl-H), 7.10 (d, J = 2.4 Hz, 1H, Aryl-H). 3.17 (s, 3H, $\text{N}-\text{CH}_3$), 1.87 (s, 18H, $\text{C}(\text{CH}_3)_3$), 1.38 (s, 9H, $\text{C}(\text{CH}_3)_3$). 1.35 (s, 9H, $\text{C}(\text{CH}_3)_3$); $^{13}\text{C}\{^1\text{H}\}$ NMR (C_6D_6 , 298 K, 176 MHz, in ppm): δ = 162.5 (Aryl-C-OH), 1.55 (Aryl-C-OH), 143.7 (Aryl-C), 142.8 (Aryl-C), 141.6 (Aryl-C), 140.7 (Aryl-C), 140.1 (Aryl-C), 127.1 (Aryl-CH), 126.3 (Aryl-CH), 124.3 (Aryl-C), 118.6 (Aryl-CH), 113.6 (Aryl-CH), 113.2 (Aryl-C), 39.7 (N- CH_3), 36.9 ($\text{C}(\text{CH}_3)_3$), 36.7 ($\text{C}(\text{CH}_3)_3$), 35.4 ($\text{C}(\text{CH}_3)_3$), 35.2 ($\text{C}(\text{CH}_3)_3$), 32.2 ($\text{C}(\text{CH}_3)_3$), 32.1 ($\text{C}(\text{CH}_3)_3$), 30.8 ($\text{C}(\text{CH}_3)_3$), 30.7 ($\text{C}(\text{CH}_3)_3$); ^{51}V NMR (C_6D_6 , 298 K, 184 MHz, in ppm): δ = -533 (s). Elemental analysis (%) calcd for $\text{C}_{31}\text{H}_{43}\text{N}_3\text{O}_3\text{VCl}$: C, 62.89; H, 7.32; N, 7.10; found C, 62.53; H, 7.12; N, 6.86.

$[\text{VO}(\text{L}^2)]$ (**2**). From $[\text{H}_3\text{L}^2][\text{Cl}]$ (1 equiv., 1 mmol, 563 mg) and $\text{VO}(\text{O}^i\text{Pr})_3$ (1.2 equiv., 1.2 mmol, 293 mg). Yield: 91% (0.91 mmol, 571 mg). ^1H NMR (C_6D_6 , 298 K, 700 MHz, in ppm): δ = 7.81 (m, 2H Aryl-H), 7.67 (d, J = 2.4 Hz, 2H, Aryl-H), 7.65 (d, J = 2.4 Hz, 2H, Aryl-H), 7.00 (m, 2H, Aryl-H), 1.87 (s, 18H, $\text{C}(\text{CH}_3)_3$), 1.36 (s, 18H, $\text{C}(\text{CH}_3)_3$); $^{13}\text{C}\{^1\text{H}\}$ NMR (C_6D_6 , 298 K, 176 MHz, in ppm): δ = 155.4 (Aryl-C-OH), 143.3 (Aryl-C), 139.9 (Aryl-C), 133.2 (Aryl-C), 128.6 (Aryl-C), 125.6 (Aryl-CH), 123.6 (Aryl-C), 123.4 (Aryl-CH), 115.4 (Aryl-CH), 114.4 (Aryl-CH), 36.2 ($\text{C}(\text{CH}_3)_3$), 34.9 ($\text{C}(\text{CH}_3)_3$), 31.7 ($\text{C}(\text{CH}_3)_3$), 30.3 ($\text{C}(\text{CH}_3)_3$); ^{51}V NMR (C_6D_6 , 298 K, 184 MHz, in ppm): δ = -503 (s). Elemental analysis (%) calcd for $\text{C}_{35}\text{H}_{44}\text{N}_2\text{O}_3\text{VCl}$: C, 67.03; H, 7.07; N, 4.47; found C, 66.94; H, 7.12; N, 4.21.

General Procedure for $[\text{Co}(\text{Cp}^*)_2][1]$ and $[\text{Co}(\text{Cp}^*)_2][2]$. The parent vanadium(V) complexes **1** or **2** (1 equiv.) were dissolved in THF and stirred for 10 min at room temperature. A solution of decamethylcobaltocene (1 equiv.) in THF was added, and the reaction mixtures were stirred for 5 h at room temperature.

$[\text{Co}(\text{Cp}^*)_2][\text{VO}(\text{L}^1)]$ ($[\text{Co}(\text{Cp}^*)_2][1]$). From complex **1** (1 equiv., 0.25 mmol, 148 mg) and $\text{Co}(\text{Cp}^*)_2$ (1 equiv., 0.25 mmol, 83 mg). After 5 h the reaction was filtered, and the solvent was evaporated. The greenish-gray residue was dissolved in CH_2Cl_2 and filtered again and concentrated to 1 mL. Hexane was added until the solution became turbid. One drop of dichloromethane was added to redissolve all solids, and the mixture was left to stand in an openly capped vial inside the glovebox for 2 days to let the dichloromethane evaporate. This formed large green blocks of $[\text{Co}(\text{Cp}^*)_2][1]$. Yield: 76% (0.19 mmol, 175.1 mg). Elemental analysis (%) calcd for

$\text{C}_{51}\text{H}_{74}\text{N}_3\text{O}_3\text{VCoCl}$: C, 66.48; H, 7.99; N, 4.56; found C, 66.75; H, 7.66; N, 4.28, μ_{eff} = 1.74 μB .

$[\text{Co}(\text{Cp}^*)_2][\text{VO}(\text{L}^2)]$ ($[\text{Co}(\text{Cp}^*)_2][2]$). From complex **2** (1 equiv., 0.25 mmol, 157 mg) and $\text{Co}(\text{Cp}^*)_2$ (1 equiv., 0.25 mmol, 83 mg). After 5 h, the green suspension was filtered, and the green solids were washed with another 5 mL of THF and 5 mL of pentane. The green solids were then dissolved in 5 mL of dichloromethane, and the solution was concentrated to 1 mL. Hexane was added until the solution became turbid. One drop of dichloromethane was added to redissolve all solids, and the mixture was left to stand in an openly capped vial for 2 days inside the glovebox to let the dichloromethane evaporate. This formed large green blocks of $[\text{Co}(\text{Cp}^*)_2][1]$. Yield: 69% (0.172 mmol, 165 mg). Elemental analysis (%) calcd for $\text{C}_{55}\text{H}_{74}\text{N}_2\text{O}_3\text{VCoCl}$: C, 69.06; H, 7.80; N, 2.93; found C, 68.37; H, 7.54; N, 2.82, μ_{eff} = 1.68 μB .

General Procedure for the Salt Metathesis Reactions. In a 20 mL scintillation vial, vanadium complex **1** was dissolved in 5 mL Et_2O and cooled to -40°C . In a separate vial, the corresponding lithium salt (phenolate or amide) was dissolved/suspended in 2 mL Et_2O and cooled to -40°C as well. This solution was then added dropwise at -40°C to the solution of the vanadium complex and slowly warmed to room temperature overnight while stirring. The deeply colored solutions are filtered to remove any lithium chloride formed during the reaction, and the solvent was evaporated under high vacuum. The colored residues were then dissolved in hexane, filtered again, and concentrated to approximately 0.5 mL. Storing these solutions at -40°C resulted in the formation of the corresponding complexes in moderate to good yields overnight.

$[\text{VO}(\text{OMes})(\text{L}^1)]$ (**3**). Following the general procedure, LiOMes (1 equiv., 0.1 mmol, 14 mg) was added to **1** (1 equiv., 0.1 mmol, 59 mg). Dark purple solid. Yield: 98% (68 mg, 0.098 mmol). ^1H NMR (C_6D_6 , 298 K, 400 MHz, in ppm) δ 8.17 (d, J = 2.4 Hz, 1H, Aryl-H), 7.73 (d, J = 2.4 Hz, 1H, Aryl-H), 7.69 (d, J = 2.5 Hz, 1H, Aryl-H), 7.13 (d, J = 2.3 Hz, 1H, Aryl-H), 6.92 (s, 1H, Mesityl-H), 6.71 (s, 1H, Mesityl-H), 3.07 (s, 3H, N- CH_3), 3.02 (s, br, 3H, C- CH_3), 2.32 (s, br, 3H, C- CH_3), 2.20 (s, 3H, C- CH_3), 1.69 (s, 9H, C- $(\text{CH}_3)_3$), 1.62 (s, 9H, C- $(\text{CH}_3)_3$), 1.41 (s, 9H, C- $(\text{CH}_3)_3$), 1.38 (s, 9H, C- $(\text{CH}_3)_3$). ^{13}C NMR (C_6D_6 , 298 K, 101 MHz, in ppm) δ 162.02 (Aryl-C-OH), 155.16 (Aryl-C-OH), 141.59 (Aryl-C), 141.39 (Aryl-C), 140.49 (Aryl-C), 139.86 (Aryl-C), 132.23 (Aryl-C), 129.51 (Aryl-C), 128.99 (Aryl-CH), 128.87 (Aryl-CH), 126.18 (Aryl-CH), 125.53 (Aryl-CH), 125.27 (Aryl-CH), 122.96 (Aryl-CH), 118.52 (Aryl-CH), 113.82 (Aryl-CH), 113.68 (Aryl-CH), 38.82 (N- CH_3), 36.21 ($\text{C}(\text{CH}_3)_3$), 36.11 ($\text{C}(\text{CH}_3)_3$), 34.81 ($\text{C}(\text{CH}_3)_3$), 34.66 ($\text{C}(\text{CH}_3)_3$), 31.86 ($\text{C}(\text{CH}_3)_3$), 31.77 ($\text{C}(\text{CH}_3)_3$), 30.04 ($\text{C}(\text{CH}_3)_3$), 29.90 ($\text{C}(\text{CH}_3)_3$), 21.13 (C- CH_3), 18.49 (C- CH_3), 17.99 (C- CH_3). ^{51}V NMR (C_6D_6 , 298 K, 79 MHz, in ppm) δ -566.66. Elemental analysis (%) calcd for $\text{C}_{40}\text{H}_{54}\text{N}_3\text{O}_4\text{V}_1$: C, 69.44; H, 7.87; N, 6.07; found C, 69.81; H, 8.16; N, 5.91.

$[\text{VO}(\text{ODipp})(\text{L}^1)]$ (**4**). Following the general procedure, LiODipp (1 equiv., 0.05 mmol, 10 mg) was added to **1** (1 equiv., 0.05 mmol, 30 mg). Dark purple solid. Yield: 96% (35 mg, 0.048 mmol). ^1H NMR (C_6D_6 , 298 K, 700 MHz, in ppm) δ 8.26 (d, 1H, J = 2.3 Hz, Aryl-H), 7.84 (d, 1H, J = 2.0 Hz, Aryl-H), 7.79 (d, 1H, J = 2.4 Hz, Aryl-H), 7.40 (dd, 1H, J = 7.6 Hz, J = 1.6 Hz, Aryl-H), 7.22 (d, 1H, J = 2.3 Hz, Aryl-H), 7.15 (dd, 1H, J = 7.8 Hz, J = 2.0 Hz, Aryl-H), 7.00 (t, 1H, J = 7.6 Hz, Aryl-H), 4.95 (m, 1H, $\text{CH}(\text{CH}_3)_2$), 3.55 (m, 1H, $\text{CH}(\text{CH}_3)_2$), 3.16 (s, 3H, NCH₃), 1.83 (d, 6H, J = 6.8 Hz, CHCH_3), 1.82 (s, 9H, CCH_3), 1.57 (d, 6H, J = 6.9 Hz, CHCH_3), 1.72 (s, 9H, CCH_3), 1.48 (s, 9H, CCH_3), 1.47 (s, 9H, CCH_3), 1.22 (d, 6H, J = 7.0 Hz, CHCH_3), 1.22 (d, 6H, J = 6.8 Hz, CHCH_3). ^{13}C NMR (C_6D_6 , 175 MHz, in ppm): δ 167.27 (C-OH), 162.52 (C-OH), 155.68, 142.07, 141.80, 141.67, 140.99, 140.38, 140.11, 139.21, 126.63, 126.00, 125.53, 124.02, 123.53, 123.46, 118.88, 114.14, 114.07, 39.20 (NCH₃), 36.56 (CCH₃), 36.51 (CCH₃), 35.12 (CCH₃), 34.99 (CCH₃), 32.16 (CCH₃), 32.05 (CCH₃), 30.59 (CCH₃), 30.38 (-CCH₃), 28.58 (CHCH₃), 28.55 (CHCH₃), 24.57 (CHCH₃), 24.40 (CHCH₃), 24.06 (CHCH₃), 23.37 (CHCH₃). Elemental analysis (%) calcd for $\text{C}_{43}\text{H}_{60}\text{N}_3\text{O}_4\text{V}_1$: C, 70.34; H, 8.24; N, 5.73; found C, 69.04; H, 8.21; N, 5.68. The low carbon value results from

potential carbide formation, which is a common problem for early transition metals.

[VO(NTol)₂(L¹)] (5). Following the general procedure, LiN(Tol)₂ (1 equiv., 0.05 mmol, 10 mg) was added to **1** (1 equiv., 0.05 mmol, 30 mg). Dark blue solid. Yield: 56% (27 mg, 0.028 mmol). ¹H NMR (C₆D₆, 298 K, 700 MHz, in ppm): δ 8.19 (d, J = 2.5 Hz, 1H, Aryl-H), 8.14 (d, J = 8.3 Hz, 2H, Aryl-H), 7.17 (s, 1H, Aryl-H), 7.16 (d, J = 8.3 Hz, 2H, Aryl-H), 7.13 (d, J = 8.6 Hz, 2H, Aryl-H), 7.11 (d, J = 2.3 Hz, 1H, Aryl-H), 6.83–6.80 (m, 2H, Aryl-H), 3.07 (s, 3H, NCH₃), 2.23 (s, 3H, CH₃), 1.88 (s, 3H, CH₃), 1.70 (s, 18H, CCH₃), 1.41 (s, 9H, CCH₃), 1.39 (s, 9H, CCH₃). ¹³C NMR (C₆D₆, 298 K, 175 MHz, in ppm): δ 162.48 (C-OH), 155.51 (C-OH), 154.80, 151.36, 142.82, 141.70, 140.29, 140.11, 139.00, 136.01, 129.61, 129.48, 128.42, 125.88, 125.86, 125.15, 122.82, 121.44, 118.80, 113.96, 38.65 (NCH₃), 36.38 (CCH₃), 36.25 (CCH₃), 34.70 (CCH₃), 34.54 (CCH₃), 31.90 (CCH₃), 31.81 (CCH₃), 30.59 (CCH₃), 30.50 (CCH₃), 20.89 (CH₃), 20.66 (CH₃). Elemental analysis (%) calcd for C₄₅H₅₇N₄O₃V: C, 71.79; H, 7.63; N, 7.44; found C, 71.44; H, 7.28; N, 7.16.

[VCl(L¹)(THF)₂] (6). In a 20 mL scintillation vial, triazolium proligand [H₃L¹][Cl] (1 equiv., 1.50 mmol, 792 mg) was dissolved in 5 mL THF. To the stirring, bright yellow solution, a THF solution of lithium hexamethyldisilazide (3.3 equiv., 4.95 mmol, 828 mg) was added dropwise over a period of 10 min. After 30 min at ambient temperature, a suspension of VCl₃(THF)₃ (1 equiv., 1.50 mmol, 560 mg) was added at once. The mixture turned dark red/brown after the addition. After 15 h the mixture was filtered, the solvent was removed by evaporation. The dark brown residue was redissolved in toluene, and precipitated lithium chloride was filtered off. Toluene was removed under reduced pressure, and the residue was washed several times with a small amount hexane to give **6** as an orange-brown powder (496 mg, 46%). μ_{eff} (Evans method, C₆D₆) = 2.71 μ_B. Elemental analysis (%) calcd for C₃₅H₅₁N₃O₃V₁LiCl: C, 60.87; H, 7.44; N, 6.08; found C, 61.00; H, 7.13; N 6.04.

General Procedure for Imido Complexes. In a 10 mL J. Young Schlenk flask, vanadium(III) complex **6** (1.0 equiv., 0.10 mmol, 72 mg) was dissolved in 5 mL of benzene. The corresponding azide was added to the solution, and the mixture was heated to 60 °C. After 24 h, the solvent was lyophilized. Residues were washed several times with hexane to afford clean product. X-ray quality crystals were grown from concentrated diethyl ether solution at ambient temperature.

[V(N-Ph)Cl(L¹)] (7). From azidobenzene (1.2 equiv., 0.12 mmol, 14 mg). Dark green solid. Yield: 100% (67 mg, 0.10 mmol). ¹H NMR (C₆D₆, 298 K, 400 MHz, in ppm) δ 8.18 (d, J = 2.4 Hz, 1H, Aryl-H), 7.80 (d, J = 2.3 Hz, 1H, Aryl-H), 7.78 (d, J = 2.4 Hz, 1H, Aryl-H), 7.18 (s, 1H, Aryl-H), 6.68–6.61 (m, 2H, Phenyl-H), 6.38 (dd, J = 8.5, 7.0 Hz, 2H, Phenyl-H), 6.31–6.25 (m, 1H, Phenyl-H), 3.16 (s, 3H, NCH₃), 1.99 (s, 9H, C(CH₃)₃), 1.97 (s, 9H, C(CH₃)₃), 1.38 (s, 9H, C(CH₃)₃), 1.35 (s, 9H, C(CH₃)₃). ¹³C NMR (C₆D₆, 298 K, 101 MHz, in ppm) δ 142.22 (Aryl-C), 141.35 (Aryl-C), 140.13 (Aryl-C), 127.39 (Aryl-CH), 127.23 (Aryl-CH), 126.03 (Aryl-CH), 125.37 (Aryl-CH), 124.83 (Aryl-CH), 123.74 (Aryl-CH), 118.24 (Aryl-CH), 113.32 (Aryl-CH), 112.53 (Aryl-CH), 38.69 (N(CH₃)), 36.56 (C(CH₃)₃), 36.44 (C(CH₃)₃), 34.87 (C(CH₃)₃), 34.71 (C(CH₃)₃), 31.82 (C(CH₃)₃), 31.74 (C(CH₃)₃), 30.26 (C(CH₃)₃), 30.20 (C(CH₃)₃). ⁵¹V NMR (C₆D₆, 298 K, 79 MHz, in ppm) δ -397.98. Elemental analysis (%) calcd for C₃₇H₄₈N₄O₂V₁Cl₁·C₆H₆: C, 69.30; H, 7.30; N, 7.52; found C, 69.18; H, 7.51; N 6.94.

[V(N-4-F-phenyl)Cl(L¹)] (8). From 1-azido-4-fluorobenzene (1.1 equiv., 0.11 mmol, 15 mg). Dark green solid. Yield: 83% (57 mg, 0.083 mmol). ¹H NMR (C₆D₆, 298 K, 400 MHz, in ppm) δ 8.18 (d, J = 2.4 Hz, 1H, Aryl-H), 7.80 (d, J = 2.3 Hz, 1H, Aryl-H), 7.78 (d, J = 2.5 Hz, 1H, Aryl-H), 7.18 (d, J = 2.4 Hz, 1H, Aryl-H), 6.46 (dd, J = 9.0, 5.1 Hz, 2H, Aryl-H), 5.94 (t, J = 8.8 Hz, 2H, Aryl-H), 3.15 (s, 3H, NCH₃), 1.98 (s, 9H, C(CH₃)₃), 1.97 (s, 9H, C(CH₃)₃), 1.37 (s, 9H, C(CH₃)₃), 1.34 (s, 9H, C(CH₃)₃). ¹³C NMR (C₆D₆, 298 K, 101 MHz, in ppm) δ 142.36 (Aryl-C), 141.47 (Aryl-C), 126.09 (Aryl-CH), 125.44 (Aryl-CH), 123.70 (Aryl-CH), 118.22 (Aryl-CH), 115.06 (Aryl-CH), 114.83 (Aryl-CH), 113.31 (Aryl-CH), 38.65 (NCH₃), 36.56 (C(CH₃)₃), 36.43 (C(CH₃)₃), 31.81 (C-

(CH₃)₃), 31.73 (C(CH₃)₃), 30.23 (C(CH₃)₃), 30.17 (C(CH₃)₃). ¹⁹F NMR (C₆D₆, 298 K, 376 MHz, in ppm) δ -109.57. ⁵¹V NMR (C₆D₆, 298 K, 79 MHz, in ppm) δ -396.83. Elemental analysis (%) calcd for C₃₇H₄₇N₄O₂F₁V₁Cl₁·0.5 C₆H₆: C, 66.34; H, 6.96; N, 7.74; found C, 63.15; H, 6.80; N 7.39. The low carbon value results from potential carbide formation, which is a common problem for early transition metals.

[V(N-4-Me-phenyl)Cl(L¹)] (9). From 1-azido-4-methylbenzene (1.1 equiv., 0.11 mmol, 15 mg). Dark green solid. Yield: 46% (31 mg, 0.046 mmol). ¹H NMR (C₆D₆, 298 K, 400 MHz, in ppm) δ 8.18 (d, J = 2.4 Hz, 1H, Aryl-H), 7.80 (d, J = 2.3 Hz, 1H, Aryl-H), 7.78 (d, J = 2.5 Hz, 1H, Aryl-H), 7.18 (d, J = 2.3 Hz, 1H, Aryl-H), 6.59 (d, J = 8.4 Hz, 2H, Aryl-H), 6.19–6.16 (m, 2H, Aryl-H), 3.16 (s, 3H, NCH₃), 2.00 (s, 9H, C(CH₃)₃), 1.99 (s, 9H, C(CH₃)₃), 1.62 (s, 3H, CCH₃), 1.38 (s, 9H, C(CH₃)₃), 1.36 (s, 9H, C(CH₃)₃). ¹³C NMR (C₆D₆, 298 K, 101 MHz, in ppm) δ 142.07 (Aryl-C), 141.18 (Aryl-C), 137.54 (Aryl-C), 128.62 (Aryl-CH), 125.97 (Aryl-CH), 125.32 (Aryl-CH), 124.93 (Aryl-CH), 123.79 (Aryl-CH), 118.26 (Aryl-CH), 113.34 (Aryl-CH), 112.58 (Aryl-CH), 38.62 (NCH₃), 36.58 (C(CH₃)₃), 36.46 (C(CH₃)₃), 31.84 (C(CH₃)₃), 31.76 (C(CH₃)₃), 30.27 (C(CH₃)₃), 30.21 (C(CH₃)₃), 20.88 (CCH₃). ⁵¹V NMR (C₆D₆, 298 K, 79 MHz, in ppm) δ -378.16. Due to the high sensitivity of the complex, no satisfactory elementary analysis could be obtained.

[V(N-TMS)Cl(L¹)] (10). From azidotrimethylsilane (1.1 equiv., 0.11 mmol, 13 mg). Dark greyish-green solid. Yield: 83% (55 mg, 0.083 mmol). ¹H NMR (C₆D₆, 298 K, 400 MHz, in ppm) δ 8.20 (d, J = 2.3 Hz, 1H, Aryl-H), 7.77 (d, J = 2.3 Hz, 1H, Aryl-H), 7.74 (d, J = 2.4 Hz, 1H, Aryl-H), 7.24 (d, J = 2.3 Hz, 1H, Aryl-H), 3.27 (s, 3H, NCH₃), 1.96 (s, 9H, C(CH₃)₃), 1.95 (s, 9H, C(CH₃)₃), 1.37 (s, 9H, C(CH₃)₃), 1.34 (s, 9H, C(CH₃)₃), -0.31 (s, 9H, Si(CH₃)₃). ¹³C NMR (C₆D₆, 298 K, 101 MHz, in ppm) δ 141.73 (Aryl-C), 140.82 (Aryl-C), 126.09 (Aryl-CH), 125.37 (Aryl-CH), 117.93 (Aryl-CH), 113.11 (Aryl-CH), 112.55 (Aryl-CH), 38.77 (NCH₃), 36.56 (C(CH₃)₃), 36.44 (C(CH₃)₃), 34.81 (C(CH₃)₃), 34.64 (C(CH₃)₃), 31.79 (C(CH₃)₃), 31.71 (C(CH₃)₃), 30.33 (C(CH₃)₃), 30.25 (C(CH₃)₃), 0.02 (Si(CH₃)₃). ⁵¹V NMR (C₆D₆, 298 K, 79 MHz, in ppm) δ -473.46. Elemental analysis (%) calcd for C₃₄H₅₂N₄O₂Si₁V₁Cl₁: C, 61.57; H, 7.90; N, 8.45; found C, 59.05; H, 8.00; N 8.27. The low carbon value results from potential carbide formation, which is a common problem for early transition metals.

X-ray Crystallography. Single crystals for X-ray diffraction experiments were performed at the analytical facility of the University of Paderborn or at the University of Innsbruck. All crystals were kept at 120(2) K or 153(2) K throughout data collection. Data collection was performed using either the ApexIII software package on a Bruker D8 Venture (Paderborn) or on a Bruker D8 Quest instrument (Innsbruck). Data refinement and reduction were performed using the Bruker ApexIII suite 2021. All structures were solved with SHELXT¹⁵⁶ and refined using the OLEX 2 software package.¹⁵⁷ Strongly disordered solvent molecules were removed using the SQUEEZE operation.¹⁵⁸ All nonhydrogen atoms were refined anisotropically, and hydrogen atoms were included at the geometrically calculated positions and refined using a riding model. For further crystallographic details, see Tables S2 and S3 in the Supporting Information.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.inorgchem.1c02087>.

NMR spectra, IR spectra, UV–vis spectra, crystallographic details, cyclic voltammograms and computational details (PDF)

Accession Codes

CCDC 2079467, 2079465, 2079472, 2079471, 2079474, 2079469, 2079473 and 2080686 contain the supplementary crystallographic data for this paper. These data can be obtained

free of charge via www.ccdc.cam.ac.uk/data_request/cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: + 441223 336033.

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Author Contributions

The project was designed and created by S.H. All experiments involving the synthesis of metal complexes were carried out by F.R.N. and S.L. The proligand $[\text{H}_3\text{L}^1][\text{Cl}]$ was synthesized by M.B. and F.R.N. EPR spectra were recorded and simulated by D.L., K.R.F., and D.J.L. X-ray diffraction analysis was performed by S.H. and K.W. ^{51}V NMRs were recorded by H.K. Computations were performed by D.M. The manuscript was written by F.R.N., D.M., and S.H. and proofread by all authors.

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

We are grateful to the Daimler and Benz Foundation, the Fonds der Chemischen Industrie, the Northrhine-Westphalian Academy of Science, Humanities, and the Arts as well as the University of Paderborn and the University of Innsbruck for funding of this work. Fabian A. Watt and Prof. Dr. Jan Paradies are kindly acknowledged for helpful discussions.

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