

Dialkyl Ether Formation at High-Valent Nickel

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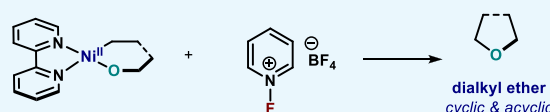
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ABSTRACT: In this article, we investigated the I_2 -promoted cyclic dialkyl ether formation from 6-membered oxanickelacycles originally reported by Hillhouse. A detailed mechanistic investigation based on spectroscopic and crystallographic analysis revealed that a putative reductive elimination to forge $C(sp^3)-OC(sp^3)$ using I_2 might not be operative. We isolated a paramagnetic bimetallic Ni^{III} intermediate featuring a unique $Ni_2(OR)_2$ (OR = alkoxide) diamond-like core complemented by a μ -iodo bridge between the two Ni centers, which remains stable at low temperatures, thus permitting its characterization by NMR, EPR, X-ray, and HRMS. At higher temperatures ($>-10^\circ C$), such bimetallic intermediate thermally decomposes to afford large amounts of elimination products together with iodoalkanol. Observation of the latter suggests that a $C(sp^3)-I$ bond reductive elimination occurs preferentially to any other challenging $C-O$ bond reductive elimination. Formation of cyclized THF rings is then believed to occur through cyclization of an alcohol/alkoxide to the recently forged $C(sp^3)-I$ bond. The results of this article indicate that the use of F^+ oxidants permits the challenging $C(sp^3)-OC(sp^3)$ bond formation at a high-valent nickel center to proceed in good yields while minimizing deleterious elimination reactions. Preliminary investigations suggest the involvement of a high-valent bimetallic Ni^{III} intermediate which rapidly extrudes the $C-O$ bond product at remarkably low temperatures. The new set of conditions permitted the elusive synthesis of diethyl ether through reductive elimination, a remarkable feature currently beyond the scope of Ni.

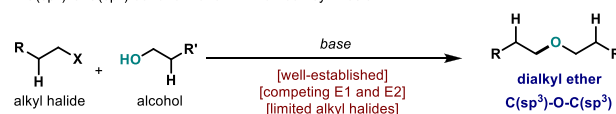


INTRODUCTION

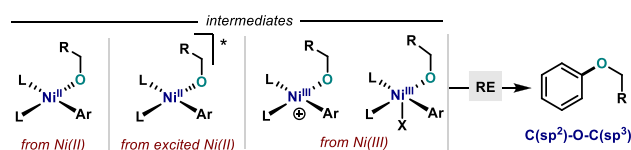
Dialkyl ethers constitute one of the most valuable functional groups, and their synthesis represents one of the oldest strategies to build chemical complexity. As a result, formation of $C-O$ bonds through the union of two organic fragments has prevailed one of the most powerful technologies, finding application across the chemical sciences: from covalent linkages and solid supports to crucial motifs in biologically active compounds.¹ From a synthetic point of view, formation of the $C-O$ bond has largely relied on the venerable Williamson ether synthesis,² which involves the union of an alcohol and an alkyl halide through a S_N2 reaction in the presence of a strong base (Scheme 1A). The high practicality and scalability of this transformation has placed it as a cornerstone reaction in both academic and industrial settings.³ Yet, the nucleophilic mechanism of the reaction is in turn its Achilles heel: the reaction efficiency is largely affected by the competitive alkoxide- or base-promoted E1 and E2 processes when secondary and tertiary alkyl halides are utilized. To circumvent these limitations, organic chemists have devoted their efforts in developing many strategies to produce highly coveted ethers—electrochemistry,⁴ organocatalysis,⁵ Lewis acid/base catalysis,⁶ among others. Nevertheless, one of the most promising alternatives in the literature to forge $C-O$ bonds relies on the mediation of transition metals. Exploiting their redox properties, transition metal catalysis has been demonstrated to be one of the pillars in the construction of these linkages.⁷ For example, Chan-Lam or Ullmann couplings

Scheme 1. (A) Williamson Ether Synthesis (Advantages and Pitfalls); (B) Existing Methods for $C(sp^2)-O$ Bond Formation Using Ni Catalysts; (C) $C(sp^3)-O-C(sp^3)$ Bond Formation from High-Valent Ni

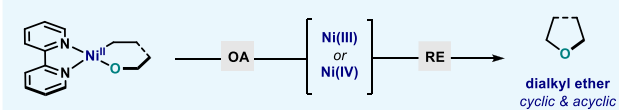
A. $C(sp^3)-O-C(sp^3)$ bond formation: Williamson synthesis



B. Reductive elimination of $C-O$ bonds from nickel complexes: a mechanistic overview



C. **This work:** Mechanistic analysis leads to $C(sp^3)-OC(sp^3)$ bond formation at a Ni center



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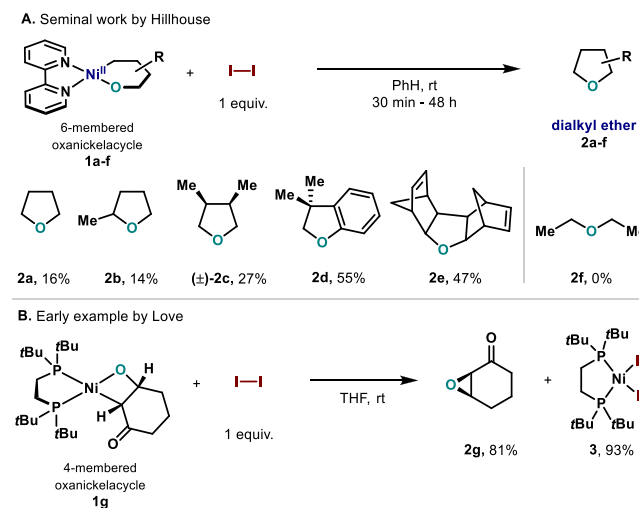


have experienced great success and found broad application for the synthesis of a myriad of highly relevant ethers.⁸ Despite the advantages associated with these methods, they have been largely dominated by linkages such as C(sp²)-O-C(sp²) and C(sp²)-O-C(sp³). In addition, Pd catalysis has also experienced tremendous development in this front, providing catalytic methods for C(sp²)-O bond formation.^{9–11} Yet, Pd complexes that permit construction of dialkylethers (C(sp³)-O-C(sp³)) through reductive elimination still present severe challenges.¹² A remarkable example of the formation of cyclic ethers is the Co-catalyzed radical cyclization of alkenols from Mukaiyama,¹³ which has found ample success in various synthetic endeavors.^{13c}

Other first-row transition metals (Cu, Fe, Ni)^{14,15} have also been demonstrated to excel as catalysts in various C–O coupling strategies.¹⁶ In particular, Ni-catalyzed transformations have gained tremendous momentum for their enormous capabilities in forging C–heteroatom bonds.¹⁷ A seminal work by Hartwig described the possibility to forge C(sp²)-O-C(sp³) bonds using Ni(COD)₂ and dppf as the optimal catalytic system.¹⁸ Encompassing Hartwig's precedent, Stradiotto described a general C(sp²)-O bond formation from L₂Ni^{II} complexes capitalizing on a newly designed set of phosphine-based ligands (L = CyPAd-DalPhos)¹⁹ (Scheme 1B). More recently, methods that replace the phosphine by simple diamine ligands have been reported, which rely on access to high-valent²⁰ or high-energy²¹ Ni complexes through light irradiation, which rapidly forge the C(sp²)-O bond. Mechanistic investigations on these latter approaches revealed that C(sp²)-O bond formation can proceed either via a Ni^{III} intermediate or via an excited Ni^{II} complex after energy transfer (Scheme 1B).²² In 2020, Ackermann and co-workers developed a nickel-catalyzed electrochemical C(sp²)-H alkoxylation, which proceeds through a Ni^{III} intermediate.²³ Recently, Nocera reported a Ni^I-catalyzed etherification protocol that mimics the reactivity of photoredox-catalyzed couplings without the use of a light source or photocatalyst.²⁴ Similarly to the metallaphotoredox protocols, it is believed that the high oxidation state of the Ni^{III} intermediate provides the necessary driving force to undergo C(sp²)-O bond linkage upon reductive elimination. Whereas realization of C(sp²)-O-C(sp³) through reductive elimination at a Ni center is well precedented and studied,²⁵ a fundamental mechanistic understanding of the analogous process to forge dialkyl ethers through C(sp³)-OC(sp³) reductive elimination still remains elusive with virtually no systematic studies on their feasibility (Scheme 1C).²⁶

In the 1990s, Hillhouse provided one of the first examples of C(sp³)-O-C(sp³) bond formation from well-defined oxanickelacycles (**1a–e**) bearing bipyridine ligands and using stoichiometric I₂ as oxidant (Scheme 2A).²⁷ In these seminal reports, involvement of high-valent nickel species such as Ni^{III}-I or Ni^{IV}-I complexes was suggested; yet, minimal evidence was provided, and such intermediates remained purely speculative. Interestingly, when I₂ was replaced by other oxidants such as O₂ or ferrocenium (Fc⁺) the C(sp³)-OC(sp³) bond formation was not observed.^{27d} In addition, higher yields were obtained for those complexes where β-hydride elimination is hampered by the limited conformations of the oxanickelacycles (Scheme 2A). As a result, formation of the C(sp³)-OC(sp³) bond was limited to cyclic products, as exemplified by the incapacity of **1f** to deliver the corresponding acyclic dialkyl ether.^{27b} On the basis of these early results, Love

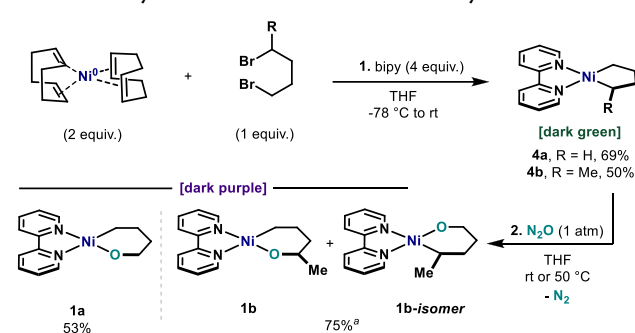
Scheme 2. I₂-Promoted C(sp³)-O-C(sp³) Bond Formation: (A) Hillhouse's Seminal Work with Bipyridine Oxanickelacycles; (B) Love's Example Using Strained Oxanickelacycle with Bidentate Phosphine



and co-workers capitalized on the I₂-promoted C–O bond formation and applied it to the oxanickelacyclobutane **1g** bearing a 1,2-bis(di-*tert*-butylphosphino)ethane (dtbpe) as the ligand (Scheme 2B).²⁸ The authors observed rapid and clean formation of the corresponding epoxide in good yield along with almost quantitative formation of the corresponding (dtbpe)Ni₂ (**3**). In this case, deleterious β-hydride elimination is not operative due to the presence of a ketone. To the best of our knowledge, these reports represent solitary examples present in the literature regarding formation of dialkyl ethers from well-defined organometallic species. Despite the powerful reactivity observed, no evidence of the intermediates involved has been reported. Yet, fundamental understanding of the key parameters that govern this particular transformation would provide tremendous insights for the design of future catalytic C(sp³)-O-C(sp³) ether syntheses. Herein, we report a comprehensive mechanistic study on the transformation originally described by Hillhouse: characterization of the reaction intermediates revealed formation of a robust and paramagnetic Ni^{III} dimer, which thermally decomposes to afford primarily elimination products. Additional mechanistic data suggests that direct C(sp³)-OC(sp³) reductive elimination from such Ni^{III} intermediate to forge simple THF rings is highly unlikely. On the contrary, experimental evidence supports an alternative mechanism based on a preferential C(sp³)-I reductive elimination followed by an intramolecular S_N2 reaction. Yet, all of these drawbacks were overridden by the replacement of I₂ by fluorine-containing oxidants which prevent not only competitive C–X reductive elimination but also deleterious elimination side reactions. In this manner, high yields of the cyclized tetrahydrofurans were obtained.

RESULTS AND DISCUSSION

Initially, we considered that oxanickelacycles **1a** and **1b** provided an excellent platform to investigate an oxidative C(sp³)-OC(sp³) bond formation (Scheme 3). Both complexes represent challenging substrates to undergo intramolecular C–O bond formation (Scheme 2A) due to the dynamic behavior of the alkoxide anion as a result of the fluxionality of the alkyl backbone. Subsequently, diverse

Scheme 3. Synthetic Route to Oxanickelacyclohexanes³¹

^aThis yield comprises a mixture of 85:15 of **1b** and **1b-isomer**.

conformations of the CH and CH₂ groups can be adopted which could lead to unproductive β -hydride elimination or simple E2. With these potential drawbacks in mind, we set out to synthesize nickelacyclopentanes **4a** and **4b** as described in the literature:^{27,29} the corresponding 1,4-alkyldibromides (1.0 equiv) were reacted with an excess of $\text{Ni}(\text{COD})_2$ (2.0 equiv) and bipyridine (bipy, 4.0 equiv) at $-78\text{ }^\circ\text{C}$ in THF. After warming up to $25\text{ }^\circ\text{C}$, the mixture is filtered and the dark-green complexes **4a** and **4b** are obtained in 69% and 50% yield, respectively (step 1, Scheme 3). Subsequently, **4a** and **4b** were subjected to O-atom insertion using N_2O following Hillhouse's procedure.²⁷ After exposing **4a** and **4b** to N_2O atmosphere (1 atm) in THF, oxanickelacyclohexane complexes **1a** and **1b** were obtained as deep purple solids in 53% and 75% yield, respectively (step 2, Scheme 3). It is important to mention that to access high-purity **1a** and **1b**, further filtration is required through Avicel³⁰ in order to remove some colloidal nickel particles.³¹ Complexes **1a** and **1b** exhibit remarkable stability in the solid state and can be stored in the freezer of the glovebox. Yet, solutions of **1a** and **1b** slowly degrade, probably through decoordination of the alkoxide ligand, which complicates purification. Indeed, purities that ranged from 85% to 93% could be achieved for **1a** and **1b** after a series of crystallizations and filtrations. Such Ni^{II} complexes are square planar and easily characterized by NMR in the diamagnetic region.³¹ Whereas **1a** offers inherent symmetry, **1b** is not symmetric and different products can arise in the O-atom insertion step (step 2, Scheme 3). Compound **1b** has both a 1° and 2° carbons, and the selectivity for the O insertion was found to be 85:15, favoring isomer **1b** over **1b-isomer**.³² However, considering that the reductive elimination of both isomers would lead to the same THF product **2b**, no further separation was attempted. For clarity, only structure **1b** will be used in the following schemes.

With these complexes in hand, cyclic voltammetry studies were performed in order to gain insight into their redox properties. As shown in Figure 1, the cyclic voltammogram of complex **1b** in CD_3CN revealed two oxidative waves at -0.70 and $+0.31$ V against Fc/Fc^+ . Interestingly, the first oxidative wave is not reversible, whereas the second wave is quasi-reversible. On the basis of other similar CVs for well-defined cyclic (bipy) Ni^{II} (alkyl)(aryl) and (terpy) Ni^{II} (C_4F_8) complexes,³³ we tentatively assigned the redox potentials to the corresponding single-electron oxidation $\text{Ni}^{\text{II}}/\text{Ni}^{\text{III}}$ and $\text{Ni}^{\text{III}}/\text{Ni}^{\text{IV}}$ couples, respectively. It is worth pointing out that the low values for both processes manifest the facility of **1b** to access high-valent Ni intermediates.³⁴

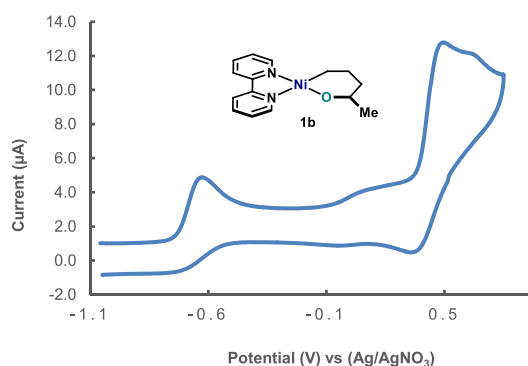
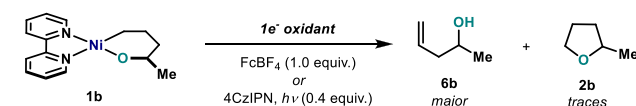


Figure 1. Cyclic voltammogram of **1b** (1.0 mM) in CD_3CN , recorded versus Ag/AgNO_3 electrode, using $n\text{-Bu}_4\text{NPF}_6$ (0.2 M) as electrolyte, under argon, with a scan rate of $100\text{ mV}\cdot\text{s}^{-1}$. Potentials are then converted to the Fc/Fc^+ couple.³¹

On the basis of the oxidation potential obtained for **1b** ($E_{1/2}(\text{Ni}^{\text{II}}/\text{Ni}^{\text{III}}) = -0.70$ V, $E_{1/2}(\text{Ni}^{\text{III}}/\text{Ni}^{\text{IV}}) = +0.31$ V vs Fc/Fc^+ in CD_3CN), it is reasonable to propose that oxidation of this complex to the corresponding Ni^{III} should be feasible using Fc^+ or O_2 .^{27d} However, Hillhouse already noticed that Fc^+ and O_2 did not lead to any C–O bond formation in good yield. Indeed, when **1b** was oxidized with FcBF_4 , mainly the elimination product was obtained (**6b**) with only trace amounts of C(sp³)–OC(sp³) bond formation (Scheme 4).

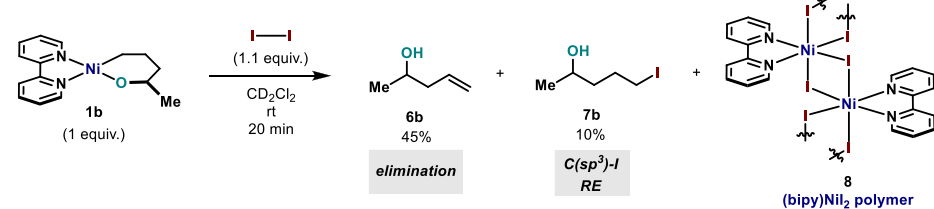
Scheme 4. Attempts to C–O Bond Formation after One-Electron Oxidation of **1b** To Access High-Valent Ni^{III} 

Other single-electron oxidants were tested, such as photocatalyst 4CzIPN (2,4,5,6-tetra(9*H*-carbazol-9-yl)-isophthalonitrile (**5**),³⁵ which would also lead to cationic Ni^{III} intermediates. Yet, no desired THF ring **2b** could be detected in the crude mixture. These results suggest that a C(sp³)–OC(sp³) reductive elimination from cationic Ni^{III} intermediates is unlikely (vide infra).

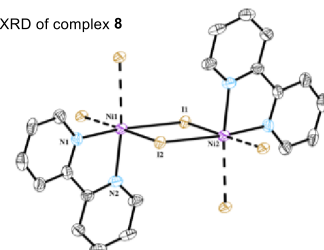
To gain further insight into the C–O bond-forming event, the reaction reported by Hillhouse and co-workers was repeated: complex **1b** (1.0 equiv) was reacted with I_2 (1.1 equiv) in CD_2Cl_2 at $25\text{ }^\circ\text{C}$.³⁶ To our surprise, upon addition of I_2 to a solution of **1b**, the color of the reaction mixture quickly changed from deep purple to orange. After 20 min of stirring at $25\text{ }^\circ\text{C}$, brown solids precipitated from the solution. At this point, the solids were filtered off and the filtrate was analyzed by ^1H NMR and GC-MS. Surprisingly, the desired cyclic ether **2b** was not observed (Scheme 5A). On the other hand, significant amounts of pent-4-en-2-ol (**6b**, 45%) were obtained, presumably through side-elimination reactions. More importantly, peaks relative to 5-iodopentan-2-ol (**7b**, 10%) were detected and successfully assigned.³¹ The presence of **7b** was further confirmed by GC-MS. After solubilizing the solids, analysis by ^1H NMR revealed a set of broad paramagnetic peaks, suggesting the presence of ligated bipy–Ni species. HRMS analysis of the solid indicated that a possible structure could be (bipy) NiI_2 ,²⁸ although dimeric compounds could not be ruled out.³⁷ Slow crystallization in CD_2Cl_2 at $-20\text{ }^\circ\text{C}$ led to crystals suitable for X-ray analysis, which unequivocally

Scheme 5. (A) Hillhouse I_2 -Promoted $C(sp^3)$ – O – $C(sp^3)$ Bond Formation Reaction (Analysis of the Fate of the Organic and Inorganic Compounds); (B) X-ray Structure of Complex 8;³¹ (C) Ni^{III} Bimetallic Intermediate 9a–b; (D) 1H NMR of Paramagnetic Complex 9a^{31b}

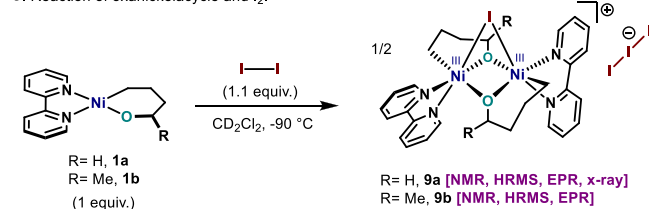
A. Analysis of the organic and inorganic species in the I_2 mediated $C(sp^3)$ – O bond formation from oxanickelacycles



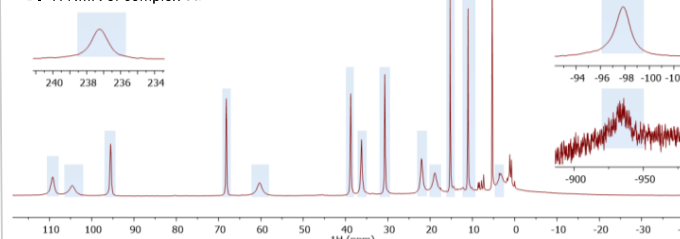
B. XRD of complex 8



C. Reaction of oxanickelacycle and I_2 .



D. 1H NMR of complex 9a



^aDisordered iodine atoms are omitted for clarity. Selected distances (Angstroms): Ni1–I1 = Ni2–I2 = 2.78; Ni1–I2 = Ni2–I1 = 2.83; Ni1–Ni1 = Ni1–Ni2 = 2.07. Selected angles (degrees): I1–Ni–I2 = 89.2; Ni1–I1–Ni2 = 94.2. ^bInsets correspond to peaks at 237, -98 , and -930 ppm.

confirmed formation of a polymeric (bipy)NiI₂ (**8**) consisting of octahedral Ni complexes linked together by μ -iodo bridges (Scheme 5B).

Considering the redox potential of iodine ($E_{1/2}(I_2/I_2^{\bullet-}) = +0.21$ V vs Fc^+/Fc),³⁸ it is reasonable to assume formation of a high-valent nickel Ni^{III} – I complex upon mixing **1a** and **1b** with I_2 . However, similarly to the case of Fc^+ , formation of Ni^{IV} is highly unlikely. To investigate whether Ni^{III} – I intermediates are formed in the reaction system, we decided to monitor the reaction by 1H NMR at low temperature. Upon adding a cold solution of I_2 (1.1 equiv) to a solution of **1a** in a J-Young tube at $-90^\circ C$, a rapid color change was observed from deep purple to deep orange (Scheme 5C). Complete conversion of **1a** to a new set of well-defined signals was observed. The 1H NMR spectrum at $-90^\circ C$ exhibits peaks ranging from -930 to $+237$ ppm, pointing to a paramagnetic complex (Scheme 5D).³¹ Upon storing a concentrated solution at $-35^\circ C$, good-quality crystals formed. X-ray analysis unambiguously determined that the intermediate consisted of a symmetric cationic Ni^{III} bimetallic complex (**9a**). This complex represents a unique and unprecedented structure of a dinuclear Ni^{III} with several structural and electronic interesting features (Figure 2). First, **9a** contains two Ni^{III} atoms in an octahedral arrangement with a large Ni–Ni distance of 2.84 Å, thus suggesting no metal–metal interaction.³⁹ In addition, **9a** features a rather unique μ -iodo bridge unifying both Ni atoms with a highly strained Ni1–I1–Ni2 angle of 58.07° . As a result, the I is bridging both Ni^{III} centers through an elongated Ni–I bond (Ni1–I1 = Ni2–I1, 2.92 Å). It is important to mention that Ni^{III} dimers with one bridging halogen atom have been recently proposed as intermediates in $C(sp^2)$ – X ($X = Br, Cl, I$) bond formation.^{39a,40} However, the most striking feature of **9a** is the diamond-like core formed by the Ni atoms and the alkoxide ligands. Two μ_2 -bridging alkoxide anions join the two Ni centers in a highly symmetric environment (Ni1–O1 = Ni1–O2 = Ni2–O1 = Ni2–O2, 2.00 Å) with angles of Ni1–O1–Ni2 = 92.72° and O1–Ni1–O2 = 79.57° . The Ni–O distances for **9a** are in the range of other bis(μ_2 -oxide)-

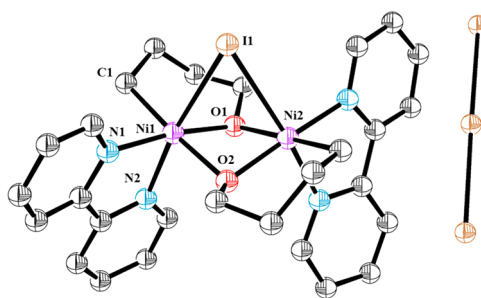


Figure 2. X-ray structure of compound **9a**.³¹ Hydrogen atoms and disordered iodide atoms in I_3^- counterion are omitted for clarity. Selected distances (Angstroms): Ni1–Ni2 = 2.84; Ni1–I1 = Ni2–I1 = 2.92; Ni1–O1 = Ni1–O2 = Ni2–O1 = Ni2–O2 = 2.00; Ni1–Ni1 = 1.99; Ni1–Ni2 = 2.05; Ni1–C1 = 2.013. Selected angles (degrees): Ni1–I1–Ni2 = 58.07° ; Ni1–O1–Ni2 = 92.72° ; O1–Ni1–O2 = 79.57° .

bridging Ni^{III} complexes known in the literature.⁴¹ The complex is complemented by the bipyridine ligands with similar Ni–N distances for both N (Ni1–N1, 1.99 Å; Ni1–N2, 2.05 Å). Finally, the remaining position of the octahedron is occupied by the alkyl residue with Ni– $C(sp^3)$ distances resembling those reported for other Ni^{III} – $C(sp^3)$ bonds (Ni1–C1, 2.013 Å).⁴² A parallel behavior was observed when complex **1b** was reacted with I_2 . However, attempts to obtain crystals of the Ni^{III} intermediate were unsuccessful. Bimetallic complexes **9a** and **9b** were further characterized by HRMS both in ESI^+ and in ESI^- modes.³¹ Finally, due to the paramagnetic nature of complexes **9a** and **9b**, further structural and electronic characterization was attempted by electron paramagnetic resonance (EPR).

The EPR spectra of complexes **9a** and **9b** were recorded at the X-band, 30 K, and are depicted in Figure 3. As expected, the EPR spectra of **9a** and **9b** are very similar but show subtle differences in line splitting and intensity ratio between the different spectral features. The multiple line splitting suggests an electron–electron spin–spin interaction between the two Ni^{III} centers. This means that the dimer structure found in the

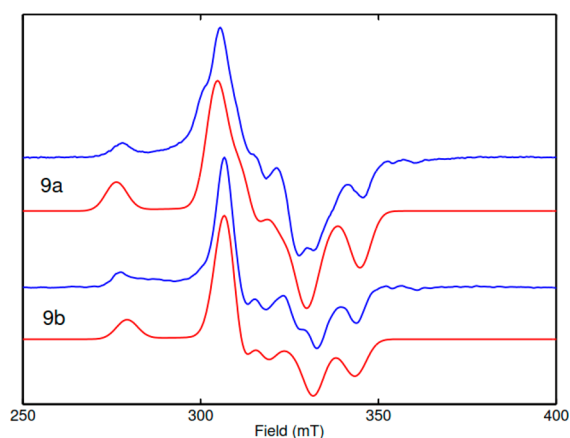


Figure 3. X-band EPR (9.623 GHz) of complexes **9a** and **9b** recorded at 30 K (blue traces).³¹ Experimental parameters: 1 mW, 100 kHz, 7.5 G field modulation. Red traces represent the Easyspin⁴³ (esfit) simulation with the following parameters: $g(9a) = (2.081, 2.155, 2.279)$; $g(9b) = (2.084, 2.144, 2.287)$. Dipolar interaction $D(9b) = 517$ MHz; $D(9a) = 550$ MHz. J coupling < 50 MHz.

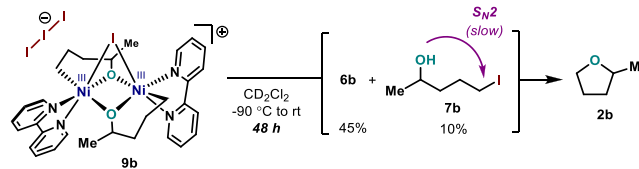
solid state (Figure 2) is retained in solution for both complexes. Making use of the symmetry properties of the dimer complex,³¹ we were able to simulate the EPR spectra as shown in Figure 3. The g -matrix principal values obtained for **9a** (2.081, 2.155, 2.279) and **9b** (2.084, 2.144, 2.287) are similar to what has been observed for a similar N,N -ligand-coordinated Ni^{III} monomer complex (2.03, 2.14, 2.20).⁴⁰ The magnetic interaction between the two Ni^{III} centers is dominated by the dipolar contribution found to be (0.9, 1.1, -2)*550 MHz for **9a** and (0.9, 1.1, -2)*517 MHz for **9b**. The J coupling between the two Ni^{III} centers is very small (50 MHz), and its effect on the EPR is only visible as a small splitting at the center of the spectrum. As confirmed by DFT analysis,³¹ the two Ni^{III} centers effectively behave as isolated $S = 1/2$ systems. This is in agreement with NMR analysis, estimating the magnetic susceptibility of the dimer complex using the Evans method to be $S = 1/2$ for each Ni^{III} center.³¹

Having identified and characterized **9**, we set out to explore its reactivity. Upon slowly warming solutions of **9a** and **9b** in CD₂Cl₂ from -90 to 25 °C, several interesting observations were made. First, the chemical shifts of complexes **9a** and **9b** are highly dependent on the temperature, which further confirms the paramagnetic nature of **9**.³¹ Moreover, **9a** and **9b** have a remarkable stability across a wide range of temperatures, from -90 to -10 °C. Beyond -10 °C, rapid evolution of **9a** and **9b** to terminal alkenes **6a** and **6b** and iodoalcohols **7a** and **7b** is observed. While traces of THF could be detected in the case of **9a**, no detectable amount of the C(sp³)-OC(sp³) bond formation product **2b** was observed for **9b**.⁴⁴ This last observation indicates that the C(sp³)-I reductive elimination is kinetically more favorable to any other C-O bond-forming event at Ni. Whereas such C(sp³)-I bond formation proceeds via a reductive elimination from Ni^{III}-I or direct attack of the I counterion to the Ni-C(sp³) bond in a S_N2 fashion is currently unknown.^{42,45} However, a similar system was recently reported by Diao, suggesting that C(sp³)-I bond formation could proceed through monomeric square pyramidal Ni^{III} complexes.⁴⁰ Hence, it is plausible to think that after dissociation of **9a** and **9b**, a similar process could be operative.

The absence of **2b** upon warming **9b** to 25 °C together with rapid consumption of oxanickelacycle **1b** at -90 °C toward

preferential formation of **6b** and **7b** led us to consider that the C(sp³)-OC(sp³) bond formation pathway accounting for the ca. 10–14% yield of **2b** may arise from a slow intramolecular cyclization of the iodoalkoxide/iodoalcohol in a S_N2 fashion.⁴⁶ Indeed, after 48 h of reaction time in the NMR tube without stirring, the iodoalcohol formed initially slowly evolves to form **2b** (Scheme 6).³¹ It is well established that 5-iodoalkoxides can

Scheme 6. Thermal Decomposition of Complex **9b** and Hypothetical S_N2 Reaction from **7b**



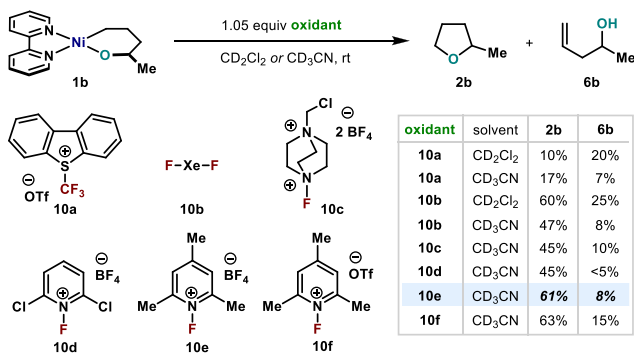
undergo intramolecular 5-*exo-tet* cyclization to afford cyclic ethers.⁴⁷ This experimental evidence supports the lack of reactivity when attempting formation of open-chain ethers such as Et₂O (**2f**) due to the much slower rates for intermolecular S_N2 reactions.⁴⁸

With these results in hand, we addressed such a defying and elusive reductive elimination. It was clear that other oxidants that enable access to high-valent Ni species should be scrutinized.⁴⁹ When I₂ was replaced by Umemoto's reagent (*S*-(trifluoromethyl)dibenzothiophenium triflate, TDTT, **10a**),^{33a} a low yield of **2b** was observed (10%). A reduced amount of side product **6b** was obtained when CD₃CN was used instead. During monitoring studies at variable temperatures, HCF₃ (boiling point = -82.1 °C) was detected. Formation of fluoroform suggests the involvement of CF₃-Ni-H intermediates and points to alkenol **6b** being formed through β-hydride elimination pathways. In addition to alkenol **6b** and HCF₃, other byproducts containing C(sp³)-CF₃ were also identified by ¹⁹F NMR, which was consistent with formation of high-valent Ni intermediates.⁵⁰ Despite the low yields, to the best of our knowledge, this challenging C(sp³)-CF₃ bond formation is unprecedented at a Ni center.⁵¹ At this point, it was quite evident that competitive C(sp³)-X reductive eliminations (X = I, CF₃) should be suppressed if the challenging C(sp³)-O-C(sp³) is to be achieved. Hence, we speculated that the presence of F ligands in the coordination sphere of a high-valent Ni intermediate would dramatically reduce the observed side reactions due to the high kinetic barrier to forge C(sp³)-F bonds.⁵² When **1b** was mixed with 1.05 equiv of XeF₂ in CD₂Cl₂ or CD₃CN, immediate reaction took place and the desired ether **2b** was observed as the major product in 60% or 47% yield, respectively.^{50a,52f,53} Interestingly, formation of **6b** remained minor in CD₂Cl₂ and could be largely suppressed in CD₃CN (8%). It is important to mention that products derived from a putative C(sp³)-F reductive elimination were only observed in trace amounts.³¹ In this line, when XeF₂ was replaced by SelectFluor (**10c**) in CD₃CN a similar outcome was obtained with a 45% yield of **2b** along with a minimal amount of **6b** (10%). We then investigated several commercially available substituted 1-fluoro-pyridinium salts (**10d-f**) as they have increased solubility in CH₃CN.⁵⁴ Using **10d**, 45% **2b** and <5% **6b** were obtained. Gratifyingly, when using NFTPb (**10e**, *N*-fluoro-2,4,6-trimethylpyridinium tetrafluoroborate) and NFTPt (**10f**, *N*-fluoro-2,4,6-trimethylpyridinium triflate) an

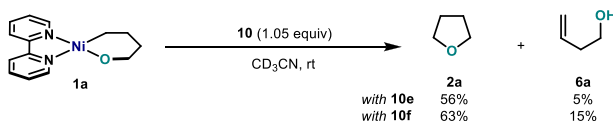
increase in the yield of **2b** was observed (61% and 63%, respectively). Notice that when using BF_4^- as counterion, formation of **6b** could be minimized to a residual 8%. As expected, **2a** was also obtained in a satisfactory 56% and 63% yield when using **10e** and **10f**, respectively (Scheme 7B).

Scheme 7. (A) Screening of Oxidants for the Oxidatively Induced $\text{C}(\text{sp}^3)\text{--OC}(\text{sp}^3)$ Bond Formation;^a (B) Application to the Synthesis of THF

A. Screening of fluorinated oxidants in the reductive elimination^a



B. Synthesis of THF via reductive elimination^a

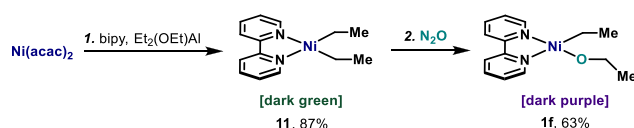


^aReaction conditions: oxanickelacycle **1b** (1 equiv), oxidant **10** (1.05 equiv) in CD_3CN or CD_2Cl_2 at 25 °C.

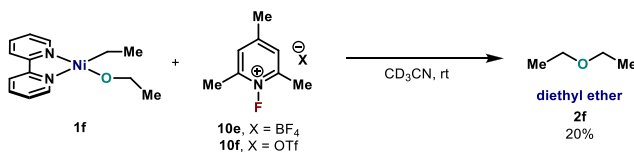
Having identified a set of conditions that enable formation of cyclic $\text{C}(\text{sp}^3)\text{--O--C}(\text{sp}^3)$ bonds at high-valent Ni centers, we speculated that a similar pathway should be valid for formation of acyclic ethers. In order to study this possibility, an acyclic precursor was synthesized through a two-step sequence (Scheme 8A). Following the protocol from Ikeda,⁵⁵ we initially synthesized (bipy)NiEt₂ (**11**) by reacting bipy, Ni(acac)₂, and (EtO)AlEt₂ in Et₂O for 50 h at 25 °C. After isolation via

Scheme 8. (A) Synthesis of (bipy)NiEt₂ **11** and (bipy)Ni(Et)(OEt) **1f**;^a (B) Oxidatively Induced Synthesis of Diethyl Ether from Nickel Complex **1f**^b

A. Synthesis of the alkyl-alkoxide Ni^{III} precursor



B. Unprecedented oxidatively-induced acyclic ether bond formation



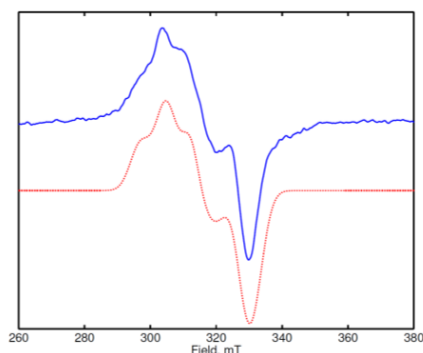
^aReaction conditions: (step 1) Ni(acac)₂ (1 equiv), bipy (1 equiv) Et₂AlOEt (3 equiv) in Et₂O, from -50 to 25 °C, 50 h, **11** 87% isolated yield; (step 2) **11** (1 equiv), N₂O (1 atm) in THF at 25 °C, **1f** 63% isolated yield. ^bReaction conditions: oxanickelacycle **1f** (1 equiv), **10e** and **10f** (1.05 equiv) in CD_3CN at 25 °C, 1 min. Yields determined by ¹H NMR using mesitylene as internal standard.

filtration (87%), the deep green solid **11** is subjected to N₂O atmosphere at 25 °C, allowing smooth O insertion into the Ni–C(sp³) bond,^{27a,b} leading to (bipy)Ni(Et)(OEt) (**1f**) in 63% yield. At this point, **1f** was subjected to our optimized oxidation conditions using **10f** in CD_3CN at 25 °C (Scheme 8B). Gratifyingly, rapid formation of Et₂O (**2f**) was observed in 20% yield at 25 °C. A similar result was obtained using **10e**. Formation of **2f** from **1f** represents a unique example of $\text{C}(\text{sp}^3)\text{--OC}(\text{sp}^3)$ bond formation and contrasts with the results obtained from Hillhouse, where $\text{C}(\text{sp}^3)\text{--OC}(\text{sp}^3)$ bond formation from acyclic Ni^{II} complexes could not be achieved (see Scheme 2A). This unprecedented result for unbiased, acyclic substrates combined with the use of simple pyridinium salt **10e** and **10f** as a mild oxidant provides an interesting proof-of-concept for the development of new strategies based on Ni and may open the door to new avenues for catalytic dialkyl ether syntheses in the future.

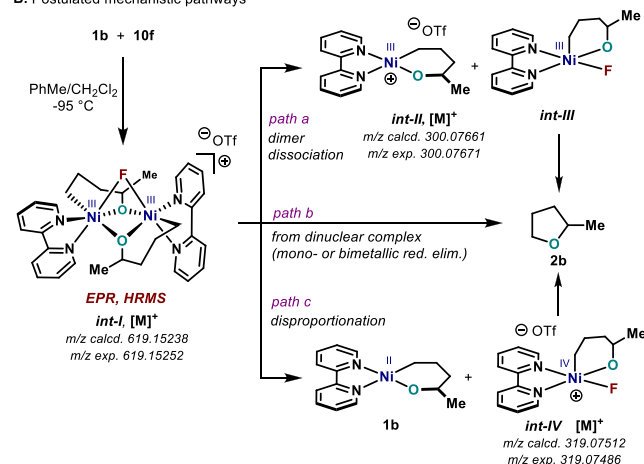
Successful formation of $\text{C}(\text{sp}^3)\text{--O--C}(\text{sp}^3)$ bonds using N-fluoropyridinium reagents posed the question of what is the exact nature of the high-valent Ni species involved in the process. On the basis of the CV results for **2b** (Figure 1) together with the oxidation results using Fc⁺, O₂, and 4CzIPN photocatalyst (Scheme 4),²⁷ it is evident that access to Ni^{III} species is facile;⁴⁹ yet, cationic Ni^{III} are not capable of C–O bond formation due to fast elimination side reactions. It is important to point out that owing to the extremely fast reaction rates for C–O bond formation, mechanistic investigations on this particular system pose an experimental challenge. Indeed, **1a**, **1b**, **10e**, and **10f** react at -90 °C in <1 min, and no Ni intermediates could be detected spectroscopically (¹H and ¹⁹F NMR). Attempts to stabilize high-valent Ni species using tripodal ligands (tris(pyrazolyl)borate⁵⁶ or tris(pyridyl)methane)^{33a,49} resulted in degradation or failed to incorporate O to the Ni^{II}–C(sp³) bond. To our delight however, a signal could be detected by EPR from reaction of **1b** with **10f**. Such species were very short lived, but a sufficient amount could be trapped after rapid (1 s) mixing at -95 °C (melting point of PhMe) and subsequent freezing in liquid N₂ (Scheme 9A). The EPR spectrum showed multiple splitting features consistent with a Ni^{III} dimer species (*int-I*, Scheme 9B). The width of the spectrum, however, is reduced with respect to that of **9a** and **9b**, suggesting a reduction in the dipolar interaction between the two Ni^{III} centers. Indeed, spectral fits resulted in a smaller value (299 vs 517 and 550 MHz), whereas the *g*-tensor only differed slightly.³¹ It is important to point out that although the fitting parameters for *int-I* are probably not unique due to the few spectral features and large number of free parameters, the current fits would be consistent with a Ni^{III} dimer with symmetry properties resembling **9a**.⁵⁷ Additional information about the putative intermediates was provided by mass spectrometry analysis. When a low-temperature mixture of **1b** and **10f** was analyzed by mass spectrometry, a *m/z* corresponding to *int-I* could be detected. In addition, *m/z* consistent with structures such as *int-II* (or **1b**) and *int-IV* were also identified.⁵⁸ Although the nature of the exact species prior to reductive elimination still remains elusive, several possibilities are envisaged. On one hand, dissociation of *int-I* would afford *int-II* and *int-III* (path a).⁵⁹ An alternative pathway would involve a disproportionation of *int-I* into starting oxanickelacycle **1b** and *int-IV* (path c).⁶⁰ Reductive elimination could then occur from either *int-III* or *int-IV*. The high degree of elimination obtained in Scheme 4 when using Fc⁺ or 4CzIPN would argue against path

Scheme 9. (A) EPR of *int-I* at 20 K;^a (B) Mass Spectrometry Results and Postulated Mechanistic Pathways

A. EPR of the mixture of **1b** + **10f** in PhMe/CH₂Cl₂



B. Postulated mechanistic pathways



^aExperimental conditions: Power = 2.0 mW, modulation (100 kHz) amplitude 7.5 G. Dotted red trace represents the Easyspin⁴³ (esfit) simulation with parameters $g = (2.103, 2.200, 2.227)$. Dipolar interaction $D(\text{int-I}) = 299$ MHz. J coupling = 100 MHz.

a as a major contributor. Moreover, another possible pathway could involve direct reductive elimination from the dinuclear complex *int-I* (path b), which has been postulated for certain C–heteroatom and C–C bond-forming events.^{40,61} In an attempt to discern between mechanistic pathways, we carried out the oxidation of **1b** with only 0.5 equiv of **10f**. In this case, one-half of the yield of **2b** observed in Scheme 7A was obtained (35–40%) without trace amounts of **6b**.³¹ The absence of elimination byproducts suggests that cationic Ni^{III} species might not be present and adds additional evidence about path a not being operative. However, the variety of pathways by which the C–O bond could be formed manifests the need for further mechanistic investigations to fully elucidate the nature of the intermediates involved.

CONCLUSIONS

In conclusion, we studied the seminal I₂-promoted C–O bond formation reported by Hillhouse toward formation of cyclic ethers (**2a** and **2b**). A detailed mechanistic investigation revealed formation of **9a** and **9b**, an unprecedented Ni^{III} bimetallic structure as a cationic intermediate, which was fully characterized by NMR, EPR, HRMS, and X-ray in the case of **9a**. These paramagnetic complexes feature a unique Ni₂(OR)₂ (OR = alkoxide) diamond-like core complemented

by a μ -iodo bridge between the two Ni centers. The anionic counterion of the complexes consists of the linear I₃[−], which remains in the outer sphere of the robust bimetallic cation. Thermal decomposition of **9** beyond −10 °C leads primarily to elimination products (**6**). In addition, substantial amounts of iodoalkanols (**7**) were detected through preferential C(sp³)–I reductive elimination. The low yields obtained for **2a** and **2b** are postulated to arise from an intramolecular S_N2 reaction from **7** over long periods of time. This manifests that the original mechanistic picture for direct C(sp³)–OC(sp³) bond formation through reductive elimination is extremely challenging for simple THF rings. Cyclic voltammetry studies as well as a survey of oxidants identified the use of fluoropyridinium reagents **10e** and **10f** as excellent candidates to afford good yields of **2** while minimizing formation of elimination byproducts (**6**). In addition, this new set of conditions was successfully applied in the elusive synthesis of acyclic diethyl ether (**2f**) from a well-defined Ni^{II} complex. Preliminary mechanistic studies revealed that upon oxidation of **1** with **10**, a highly reactive intermediate could be detected in solution by EPR and HRMS, which is consistent with a Ni^{III} dimeric structure (*int-I*). Efforts to fully characterize the high-valent species involved after oxidation and prior to reductive elimination are currently under investigation in our laboratory. We believe the findings reported here open the door to new avenues for Ni catalysis and could aid practitioners in the field to unravel novel metal-catalyzed ether synthesis.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.0c07381>.

Experimental procedures and analytical data (¹H, ¹⁹F, and ¹³C NMR, EPR, X-ray, HRMS, CV) (PDF)

CIF file for compound **8**, also available free of charge from the www.ccdc.cam.ac.uk under CCDC number 2014828 (CIF)

CIF file for compound **9a**, also available free of charge from the www.ccdc.cam.ac.uk under CCDC number 2014830 (CIF)

CIF file for compound **S1**, also available free of charge from the www.ccdc.cam.ac.uk under CCDC number 2014829 (CIF)

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Notes

The authors declare no competing financial interest.

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