Phosphorus Analogues of [Ni(bpy)2]: Synthesis and Application in CarboxytextendashHalogen Bond Activation

Leitl, J.; Coburger, P.; Scott, D.J.; Ziegler, C.G.P.; Hierlmeier, G.; Wolf, R.; van Leest, N.P.; Bruin, B. de; Hörner, G.; Müller, C.

DOI
10.1021/acs.inorgchem.0c01115

Publication date
2020

Document Version
Final published version

Published in
Inorganic Chemistry

License
Article 25fa Dutch Copyright Act

Citation for published version (APA):
Phosphorus Analogues of [Ni(bpy)_2]: Synthesis and Application in Carbon–Halogen Bond Activation

J. Leitl, P. Coburger, D. J. Scott, C. G. P. Ziegler, G. Hiermeier, R. Wolf,* N. P. van Leest, B. de Bruin,* G. Hörner, and C. Müller*

ABSTRACT: The neutral, homoleptic pyridylphosphininenickel(0) complex [Ni(2-Py-4,6-Ph_2-PC_5H_2)_2] (1) has been obtained by reaction of the formal Ni(0) source [(IPr)Ni(H_C=CHSiMe_3)_2] with 2 equiv of 2-(2'-pyridyl)-4,6-diphenylphosphinine (L). Compound 1 can be oxidized both electrochemically and through the use of ferrocenium salts, to afford the corresponding Ni(I) complexes [1][BF_4], [1][THF]PF_6, and [1][BArF_4]. The structures of these salts reveal an interesting dependence on the nature of the anion. While [1][BF_4] and [1][THF]PF_6 show trigonal-bipyramidal coordination of Ni in the solid state, [1][BArF_4] exists as a dinuclear Ni(I) complex and possesses a bridging phosphinine moiety in a rare μ_2 mode. Reactions of 1 with halobenzenes highlight the noninnocent behavior of the aromatic phosphinine ligand, leading to the formation of oxidized Ni complexes but not to classical oxidative addition products. The reaction of 1 with bromobenzene affords the Δ^5 phosphinine 2 and the bipyramidal Ni(I) complex [1][Br], whereas a more unconventional oxidation product 3 is formed from the reaction of 1 and iodobenzene.

INTRODUCTION

Nickel(0) complexes have become ubiquitous in homogenous catalysis and are used for a wide range of processes, including alkene and alkyne oligomerization reactions, as well as Kumada-type cross couplings. The complex bis(2,2'-bipyridine)nickel(0) (A; Figure 1) is an archetypal example of a Ni(0) species with versatile catalytic properties and has found diverse applications in various (electro)chemical reactions and catalysis.

Replacing the pyridyl moieties in A with valence isoelectronic phosphinine (also referred to as phosphabenzenes) units can have a significant impact on both the electronic structure and reactivity of the corresponding coordination compound. Considering the versatile applications of A and bipyridine complexes, as well as the fact that monophosphininenickel(0) complexes are well-investigated, it is quite surprising that phosphorus-containing analogues of A are extremely rare. In fact, the known examples appear to be limited to just a single species described by Le Floch and co-workers, who reported that reactions of 2,2':(4,5-dimethyl)biphosphinine with nickel(0) sources, such as Ni(cod), (cod = 1,5-cyclooctadiene), afford the homoleptic bis(biphosphinine) complex B (Figure 1). However, the reactivity of this complex—and almost all related chelating phosphininenickel complexes—remains unexplored.

We have recently become interested in the coordination chemistry of the donor-functionalized phosphinine 2-(2'-pyridyl)-4,6-diphenylphosphinine (L), a “hybrid” P,N ligand that can be considered to be a cross between previously explored biphosphinines and the ubiquitous bipyridines. L possesses two electronically distinct binding sites—a “soft” P and a “hard” N donor site—which can have a significant impact on its coordination properties and reactivity. Although a large variety of 4d and 5d transition-metal complexes of L have been prepared, syntheses of 3d metal complexes remain scarce. Nevertheless, we have recently demonstrated that the combination of a late 3d metal with ligand L can lead to versatile coordination chemistry and, even more importantly, to useful reactivity, as exemplified by the facile cleavage of one C=O bond in CO_2 by the Fe complex C (Figure 1).

EXPERIMENTAL DETAILS

All experiments were performed under an atmosphere of dry argon using standard glovebox and Schlenk-line techniques. Tetrahydrofur-

Received: April 16, 2020
Published: July 2, 2020
an (THF), toluene, diethyl ether, and n-hexane were purified, dried, and degassed using an Mibrn SPS800 solvent purification system. Benzene was distilled and degassed with Schlenk techniques. Deuterated THF and benzene were purchased from Sigma-Aldrich and used as received. Bromobenzene and iodobenzene were purchased from Sigma-Aldrich and used as received. Deuterated THF and benzene were purchased from Sigma-Aldrich. 

- **Figure 1.** Previously reported bidentate, chelated phosphinine-containing transition-metal complexes (A–C) and the new nickel pyridylphosphinine complexes ([Ni(2-Py-4,6-Ph2-PC5H2)2](1), [1]Br, and 3) described herein. 18c6 = [18]crown-6.

- **Synthesis.** Synthesis of 1. Compound 1 was prepared in an Mbrn argon glovebox. 1 is sensitive toward moisture and air. It is soluble and stable in benzene, THF, and toluene. A solution of L (2.5 equiv, 125 mg, 0.385 mmol) in benzene (2 mL) was added dropwise to a solution of D (1 equiv, 100 mg, 0.154 mmol) in benzene at room temperature. An immediate color change from yellow to deep purple was observed. The reaction mixture was stirred for 16 h at room temperature. Volatiles were completely removed, and the deep-purple residue was washed with n-hexane (3 x 2 mL) in order to remove IPr. The remaining residue was extracted with benzene (4 x 2 mL) and layered with n-hexane (12 mL). 1 was isolated as deep-purple crystals after decanting the mother liquor, washing with n-hexane (3 x 2 mL), and drying under vacuum. Crystals suitable for single-crystal XRD were grown from the slow diffusion of n-hexane into a benzene solution of 1. Yield: 40 mg, 37%. 

- **X-band electron paramagnetic resonance (EPR) spectra** were recorded on Bruker Avance 400 spectrometers referenced externally to 85% H3PO4(aq). The assignment of 1H and 13C NMR signals was confirmed by two-dimensional (COSY, HSQC, HMBC) experiments. 

- **Magnetic susceptibilities** in solution were determined by the Evans NMR method in THF-d8 at room temperature. UV–vis spectra were recorded using a Varian Cary 50 spectrometer. Elemental analyses were determined by the analytical department of the University of Regensburg. Cyclic voltammograms were recorded with a CH Instruments electrochemical analyzer. Gas chromatography with a flame ionization detector (GC-FID): Shimadzu GC2010plus. Carrier gas: H2; Column: Restek Rxi (30 m x 0.25 mm x 0.25 µm) Carrier gas: H2; Standard heating procedure: 50 °C (2 min), 25 °C min⁻¹ → 280 °C (5 min). Gas chromatography-mass spectrometry (GC-MS) HP6890 system with a 7683B injector and an Agilent 7820A system. Column: HP-5, 19091J-413 (30 m x 30.25 mm x 0.25 µm). Carrier gas: N2. Calibration of substrates and products with internal standard n-pentadecane and analytically pure samples. The experimental X-band electron paramagnetic resonance (EPR) spectra were recorded on a Bruker EMX spectrometer (Bruker BioSpin Rheinstetten) equipped with a He temperature-controlled cryostat system (Oxford Instruments). The g values were calculated with the ORCA software package at the B3LYP/def2-TZVP level of theory. The spectrum was analyzed and simulated using the WSEP program of Prof. Frank Neese. 

- **Single-crystal X-ray diffraction (XRD)** data were recorded on an Agilent Technologies SuperNova diffractometer with Cu Kα radiation (λ = 1.54184 Å). Either semiempirical multisite absorption corrections or analytical ones were applied to the data. The structures were solved with SHELXTL and least-squares refinements on F² were carried out with SHELXL. The H atoms were located in idealized positions and refined isotropically with a riding model. 

- **NMR spectra** were recorded on Bruker Avance 400 spectrometers (1H, 13C) and Bruker Avance 500 spectrometers (1H, 31P) referenced externally to 85% H3PO4(aq). The assignment of 1H and 13C NMR signals was confirmed by two-dimensional (COSY, HSQC, HMBC) experiments. The remaining residue was extracted with benzene (4 x 2 mL) and layered with n-hexane (12 mL). 1 was isolated as deep-purple crystals after decanting the mother liquor, washing with n-hexane (3 x 2 mL), and drying under vacuum. Crystals suitable for single-crystal XRD were grown from the slow diffusion of n-hexane into a benzene solution of 1. Yield: 40 mg, 37%. 

- **Elemental analysis.** Calculated for C43H30N2P Ni (Mw = 709.39 g·mol⁻¹): C, 74.50; H, 4.55; N, 3.95. Found: C, 74.98; H, 4.70; N, 3.77. UV–vis [THF; λₘ𝑎ₓ (nm)] (ε (cm⁻³ mol⁻¹ g⁻¹)) = 290 (98632), 520 (29740), 816 (20814). 1H NMR (400.13 MHz, 300 K, THF-d8): δ 7.05–7.11 (m, 6H, H of C₂=Ph), 7.18–7.21 (m, 2H, H of NC₅H₄), 7.33–7.37 (m, 2H, H of C₂=Ph), 7.41–7.45 (m, 4H, H of C₂=Ph), 7.79–8.17 (m, 4H, H of C₂=Ph), 7.85–8.66 (m, 2H, H of NC₅H₄). 13C NMR (100.61 MHz, 300 K, THF-d8): δ 139.0 (d, 4H, H of C₂=Ph), 138.4 (d, 4H, H of C₂=Ph), 130.4 (d, 4H, H of C₂=Ph), 129.0 (d, 4H, H of C₂=Ph), 128.6 (d, 4H, H of C₂=Ph), 127.4 (t, 4H, H of C₂=Ph), 127.4 (t, 4H, H of C₂=Ph), 124.8 (m, C of C₂=Ph), 143.9 (m, C of C₂=Ph), 145.1 (m, C of C₂=Ph), 146.6 (m, C of C₂=Ph), 153.1 (m, C of C₂=Ph). 

- **X-ray crystallography.** The crystal structures were solved with SHELXTL and least-squares refinements on F² were carried out with SHELXL. The H atoms were located in idealized positions and refined isotropically with a riding model. The solvent of the green reaction mixture was completely removed, and the orange product was characterized by sublimation under high vacuum (50 °C; ca. 10⁻³ mbar). The remaining brown residue was extracted with THF (2 mL) and layered with n-hexane (4 mL). After crystallization for 2 days at ~35 °C, 1 was isolated as a brown powder. Crystals suitable for single-crystal XRD were grown from the slow diffusion of n-hexane.
into a toluene solution of [1]BF₆. Yield: 43 mg, 77%. Elem. anal. Calcd for C₃₀H₁₈N₅PF₆ (Mₙ = 796.20 g mol⁻¹): C, 66.38; H, 4.05; N, 3.52. Found: C, 65.17; H, 4.26; N, 3.35. UV–vis [THF; \( \lambda_{\text{max}} / \text{nm} \) (\( \epsilon_{\text{max}} / \text{L mol}^{-1} \cdot \text{cm}^{-1} \))]: 284 (38665), 338 (sh, 15633), 466 (8545). \( \mu \) (THF-d₈): 2.0(1) \( \mu_{\text{B}} \).

**Synthesis of [1]THF**. Compound [1]THF was prepared in an MBraun argon glovebox. [1]THF was sensitive toward moisture and air. [1]THF is soluble and stable in benzene, THF, and toluene. 1 (50 mg, 0.07 mmol) was dissolved in THF (1 mL), and FeC₆F₆ (23 mg, 0.07 mmol) was added at room temperature. After stirring for 1 day at \(-35^\circ\text{C}\), [1]THF/FeC₆F₆ could be isolated as a deep-brown powder. Crystals suitable for single-crystal XRD were grown from the slow diffusion of n-hexane into a THF solution of [1]THF/FeC₆F₆. Yield: 86 mg, 78%. Elem. anal. Calcd for C₁₅H₁₀N₃P·FeC₆F₆ (Mₙ = 926.47 g mol⁻¹): C, 66.96; H, 4.09; N, 3.55. Found: C, 64.83; H, 4.75; N, 3.21. UV–vis [THF; \( \lambda_{\text{max}} / \text{nm} \) (\( \epsilon_{\text{max}} / \text{L mol}^{-1} \cdot \text{cm}^{-1} \))]: 279 (55432), 330 (22957), 454 (26207), 512 (sh), 605 (sh). \( \mu \) (THF-d₈): 1.79(1) \( \mu_{\text{B}} \).

**Synthesis of [2]PF₆**. Compound [2]PF₆ was prepared in an MBraun argon glovebox. [2]PF₆ is sensitive toward moisture and air. [2]PF₆ is soluble and stable in benzene, THF, and toluene. 1 (50 mg, 0.07 mmol) was dissolved in toluene (1 mL), and toluene (3 mL) and drying under vacuum. 1 (50 mg, 0.07 mmol) was added to a diethyl ether solution of [2]PF₆ (74 mg, 0.07 mmol) were dissolved in THF (2 mL) at room temperature. An immediate color change from deep purple to brown was observed. After stirring overnight, the solution turned green, and the solvent of the reaction mixture was completely removed. Ferrocene was removed by washing with n-hexane. Bromobenzene (7.5 g, 0.07 mmol) was added to a toluene (1 mL), and this reaction mixture was completely removed. Ferrocene was removed by sublimation under high vacuum (50 °C, ca. 10⁻² mbar). The remaining brown residue was extracted with THF (2 mL) and layered with n-hexane (6 mL). After crystallization for 1 day at \(-35^\circ\text{C}\), [2]PF₆ could be isolated as a deep-brown powder. Crystals suitable for single-crystal XRD were grown from the slow diffusion of n-hexane into a THF solution of [2]PF₆/FeC₆F₆. Yield: 43 mg, 86%. Elem. anal. Calcd for C₁₅H₁₀N₃P·FeC₆F₆ (Mₙ = 796.20 g mol⁻¹): C, 66.96; H, 4.09; N, 3.55. Found: C, 64.83; H, 4.75; N, 3.21. UV–vis [THF; \( \lambda_{\text{max}} / \text{nm} \) (\( \epsilon_{\text{max}} / \text{L mol}^{-1} \cdot \text{cm}^{-1} \))]: 279 (55432), 330 (22957), 454 (26207), 512 (sh), 605 (sh). \( \mu \) (THF-d₈): 1.79(1) \( \mu_{\text{B}} \).

**Synthesis of [3]BF₄**. Compound [3]BF₄ was prepared in an MBraun argon glovebox. [3]BF₄ is soluble in benzene, THF, and toluene. 1 (50 mg, 0.07 mmol) was dissolved in THF (2 mL) and drying under vacuum, 1 (50 mg, 0.07 mmol) was added to a toluene (1 mL), and this reaction mixture was heated to 60 °C overnight. The deep-purple solution turned deep red upon heating. The mixture was cooled to room temperature and layered with n-hexane (8 mL). After crystallization at room temperature overnight, a deep-green solid was isolated by decanting the fluorescent pink solution. After washing with n-hexane (2 × 2 mL) and drying under vacuum, 3 was isolated as a deep-green powder. Crystals suitable for single-crystal XRD were grown from the slow diffusion of n-hexane into a toluene solution of [3]BF₄. Yield: 38 mg (64%). Elem. anal. Calcd for C₁₅H₁₀N₃P·FeC₆F₆ (Mₙ = 843.96 g mol⁻¹): C, 53.80; H, 3.85; N, 1.66. Found: C, 53.99; H, 3.58; N, 1.98. UV–vis [THF; \( \lambda_{\text{max}} / \text{nm} \) (\( \epsilon_{\text{max}} / \text{L mol}^{-1} \cdot \text{cm}^{-1} \))]: 285 (24370), 450 (4788), 614 (2600). \( \mu \) (THF-d₈): 3.2(1) \( \mu_{\text{B}} \). The decanted fluorescent pink solution was evaporated to dryness, and the remaining pink residue was extracted with n-hexane (3 × 2 mL). Slow evaporation at room temperature gave pure 2 as a pink powder. Yield: 10 mg, 30%.

**RESULTS AND DISCUSSION**

Encouraged by our previous results with Fe, we were motivated to investigate the coordination compounds of the pyridylphosphinine ligand L with other late 3d metals. Given the paucity of prior Ni complexes containing chelating phosphinine ligands—and, in particular, the lack of corresponding reactivity studies—we were specifically motivated to study the reactivity of L toward suitable low-valent Ni sources and decided to start with the formal Ni(0) source D. While the 1:1 reaction of D and L resulted in a mixture of several inseparable products (Figure S11), the reaction of D with 2.5 equiv of L led to the selective formation of a single phosphorus-containing species (1), characterized by a singlet in the \( ^{31}\text{P} [\text{H}] \) NMR spectrum at \( \delta = 179.4 \text{ ppm} \) (Scheme 1; for the spectrum, see Figure S5). Isolation of 1 was achieved by the removal of volatiles (including H₂C=CH₂Et) under vacuum, the removal of IPr by washing with n-hexane, and the layering of a purple solution of 1 in benzene with n-hexane at room temperature for 2 days, to afford a pure, crystalline material in 35% yield.

Compound 1 was fully characterized by NMR and UV–vis spectroscopy, elemental analysis, and cyclic voltammetry (CV; see the Supporting Information, SI). The collected data are all consistent with the formulation of 1 as the 1:2 homoleptic chelate complex [Ni(L₂)] (Scheme 1), which is also in line with the reaction stoichiometry employed.
Scheme 1. Synthesis of 1 Using D as a Ni(0) Precursor

\[
\begin{align*}
\text{Dipp} & \quad \text{Me}_2\text{Si} \quad \text{SiMe}_3 \quad \text{Ph} \\
\rightarrow & \quad 2.5 \text{ L} \\
\rightarrow & \quad 1 \\
\end{align*}
\]

Figure 2. Molecular structure of 1 in the solid state (left). Ellipsoids are drawn at the 40% probability level; H atoms are omitted for clarity; four phenyl groups are displayed in wireframe for clarity. Selected bond lengths [Å] and bond angles [deg]: Ni1–N1 2.1355(15), Ni1–P1 2.091(2), Ni1–P2 2.092(2), P1–Ni1–P2 128.15(8), N1–Ni1–N2 110.4(2), P1–Ni1–N2 124.67(18), N1–Ni1–P2 129.03(17), P1–Ni1–N1 84.05(17), P2–Ni1–N2 84.28(17), C1–P1–C5 102.0(3), C1′–P2–C5′ 102.2(3), C1′–P2–C5′ 102.0(3). Cyclic voltammograms of 1 (right; from -400 to +100 mV). As expected homoleptic chelate complex, where the Ni center is anticipated, the solid-state molecular structure shows the planar and tetrahedral geometries, respectively. The P–Ni–N bite angles are 84.05(17)° for P1–Ni1–N1 and 84.28(17)° for P2–Ni1–N2. The C1–P1–C5 angle of 1 is approximately 102°, close to the one in free phosphinines (ca. 100°). This is in line with earlier observations that electron-rich metal fragments cause only a marginal opening of the C1–P1–C5 angle in the coordinated phosphinine. In contrast, this effect is much more pronounced in coordination compounds, in which the phosphinine ligand is bound to more electron-poor metal centers with reduced π-back-donation ability. Those complexes are usually characterized by a high reactivity of the P=C double bond toward nucleophilic attack.

Complex 1 is the first example of a Ni complex containing a pyridylphosphinine ligand. The molecular structure of 1 is reminiscent of ubiquitous 2,2′-bipyridinenickel(0) complexes, which are known for their versatile redox properties and corresponding utilization in the electrochemical reduction of aryl halides, for example.6–11,54 Thus, in order to determine the redox properties of 1, cyclic voltammograms were recorded in THF using [nBu4N][PF6] as a supporting electrolyte. Several irreversible and quasi-reversible oxidation and reduction processes are apparent (Figure 2). However, perhaps the most interesting process is the quasi-reversible oxidation observed at $E_{1/2} = -0.6$ V versus Fc/Fc+. Because this feature is well-behaved on the CV time scale, it was anticipated that 1 might also be oxidized in a preparative manner using a suitable chemical oxidant. Thus, 1 was treated with several different ferrocenium salts FcX [Fc = ferrocenium; X = BF4, PF6, and BArF4, where $Ar^f = 3,5-(CF3)2C6H3] in order to obtain the corresponding cationic Ni complexes [1]BF4, [1(THF)]PF6, and [12](BArF4)2 (Scheme 2), which could all be characterized crystallographically.

Table 1. Geometry Indices ($\tau_4$ and $\tau_5$)55,56 and Magnetic Moments ($\mu_{eff}$) THF-d8 300 K of Reported Ni Complexes

<table>
<thead>
<tr>
<th>compound</th>
<th>$\tau_4$</th>
<th>$\tau_5$</th>
<th>$\mu_{eff}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.80</td>
<td>0.53</td>
<td>2.0(1)</td>
</tr>
<tr>
<td>[1]BF4</td>
<td></td>
<td>0.71</td>
<td>1.7(1)</td>
</tr>
<tr>
<td>[1]Br</td>
<td></td>
<td>0.79</td>
<td>2.1(1)</td>
</tr>
<tr>
<td>3</td>
<td>0.95</td>
<td>3.2(1)</td>
<td></td>
</tr>
</tbody>
</table>

The solid-state molecular structures of [1]BF4 and [1(THF)]PF6 reveal a common structural motif: trigonal-bipyramidal Ni(I) complexes coordinated by two ligands L and direct equatorial coordination of an additional ligand (Figure 3). In [1]BF4, the tetrafluoroborate anion shows an interaction of one F atom with the metal center [Ni1–F1 distance: 2.2729(9) Å]. The value of the geometry index for [1]BF4 ($\tau_5 = 0.53$) is between those expected for square-pyramidal ($\tau_5 = 0$) and trigonal-bipyramidal ($\tau_5 = 1$; Table 1) complexes. For [1(THF)]PF6, the fifth coordination site at Ni is occupied by a THF solvent molecule [Ni1–O1 2.1355(15) Å], rather than by the counteranion, which can be attributed to the reduced donor strength of the PF6− anion relative to BF4− (Figure 3). A similar coordination motif was observed following the
The magnetic moments of \([\text{I}]\text{BF}_4\) and \([\text{I}(\text{THF})]\text{PF}_6\) were determined by the Evans NMR method and are consistent with \(S = \frac{1}{2}\) Ni(I) centers (Table 1). EPR measurements of \([\text{I}]\text{BF}_4\) and \([\text{I}(\text{THF})]\text{PF}_6\) were recorded in toluene glasses at 20 and 40 K, respectively, and also confirmed the presence of a single unpaired electron at each Ni center (for full details, see the SI). The EPR signal of \([\text{I}]\text{BF}_4\) appears almost isotropic because of the broad and overlapping signals of the \(g\) tensor. However, it is better described as a rhombic system, which is indicated by the asymmetric shape of the signal. The observed \(g\) tensor with the simulated principal components \(g_{11} = 2.195\), \(g_{22} = 2.127\), and \(g_{33} = 2.060\) is consistent with a Ni-centered radical in a system with small \(g\) anisotropy (for the spectrum, see Figure S20). The EPR spectrum of \([\text{I}(\text{THF})]\text{PF}_6\) displays similarly overlapping signals of the \(g\) tensor and can again be described as a rhombic system with small \(g\) anisotropy. Poorly resolved hyperfine interactions were observed on (at least) one \(g\) tensor, and satisfactory simulation was achieved after the inclusion of hyperfine coupling (135 and 150 MHz) to \(^{31}\text{P}\) (\(I = \frac{1}{2}\)) on two \(g\) tensors. The observed \(g\) tensor (simulated \(g\) tensor: \(g_{11} = 2.200\), \(g_{22} = 2.129\), and \(g_{33} = 2.035\)) is again consistent with a Ni-centered radical (for the spectrum, see Figure S21). These results are consistent with the density functional theory (DFT) calculations carried out on \([\text{I}]\text{BF}_4\), which show a large amount of spin density on Ni (0.85) and reproduce the experimental \(g\) tensor quite well [TPSSH/IGLO-III+CP(PPP); see the SI].

The reaction of \(1\) with FcBaB\(_4\) leads to a product with a notably different structure, at least in the solid state. In this case, the product \([\text{I}_2]\text{BaB}_4\) is formed, which crystallizes as a dicatonic dinuclear Ni complex fragment with two Ni(I) centers bridged by one P atom of one phosphinine ligand L. The charge of the dication is balanced by two BArF\(_4\) anions, which show no close contacts with either Ni center (Figure 3). Compound \([\text{I}_2]\text{BaB}_4\) is a rare example of a complex in which the phosphinine ligand shows a \(\mu-P\) bridging coordination mode.\(^5\) The corresponding bipyridyl \(\mu-C\) bridging mode is even scarcer (despite the fact that these complexes were studied far more extensively) because of the reduced electronic versatility of the lighter heterocycle.\(^6\) The Ni–N and Ni–P distances in \([\text{I}_2]\text{BaB}_4\) are similar to those in \([\text{I}]\text{BF}_4\) and \([\text{I}(\text{THF})]\text{PF}_6\) except for the bridging Ni–P three distance, which is slightly elongated \(2.4071(11)\) Å vs mean distance 2.246 Å, as expected for a bridging interaction. The bridging P3 atom is tetrahedrally coordinated between the two Ni(I) centers with a geometry index of \(r_3 = 0.86\).

Also, in contrast to \([\text{I}]\text{BF}_4\) and \([\text{I}(\text{THF})]\text{PF}_6\) the EPR signal of \([\text{I}_2]\text{BaB}_4\) in a toluene glass at 30 K is well resolved and shows a rhombic system with small \(g\) anisotropy and hyperfine coupling interactions with two \(^{31}\text{P}\) nuclei along three \(g\) tensors (\(A_{11}^{P} = A_{11}^{P} = 205\) MHz, \(A_{22}^{P} = A_{22}^{P} = 170\) MHz, and \(A_{33}^{P} = A_{33}^{P} = 175\) MHz). The spectrum again indicates that a Ni-centered radical is present (simulated \(g\) tensors: \(g_{11} = 2.2170\), \(g_{22} = 2.1450\), and \(g_{33} = 2.0195\); Figure S22). The found \(g\) values (close to 2) and well-defined
hyperfine coupling interactions are indicative for an isolated S = 1/2 species. Thus, the EPR data might indicate that [1_u]([BArF]_4)_2 dissociates in solution into two monocations [Ni(I)L_2]^+. This is also consistent with DFT calculations on the monocation, which yield a rhombic g tensor (g = 2.123, g = 2.116, and g = 2.015) that is in good agreement with the experimental results.

When the EPR measurement of [1_u]([BArF]_4)_2 is performed in a 2-methyltetrahydrofuran glass at 20 K instead of a toluene glass, a well-resolved rhombic system with small g anisotropy is again observed (Figure 4). However, in this case, additionally resolved hyperfine couplings to two equivalent ^31P (I = 1/2), two equivalent ^14N (I = 1), and two equivalent ^1H (I = 1/2) nuclei are also observed. Calculations indicate that the proton hyperfine couplings are caused by the o-H atoms of the pyridyl groups (see the SI for details). The simulated g tensor (g = 2.21460, g = 2.1460, and g = 2.0212) is similar to that obtained in toluene and so is again consistent with a Ni-centered radical. The presence of monocationic [Ni(I)] centers is additionally confirmed by the magnetic moment of 1.7(1) μ_B (in THF-d_8 at 300 K) per Ni atom measured by the Evans NMR method.

In addition to simple outer-sphere oxidation, we were also interested in investigating more complex reactions of 1, particularly given the complete lack of analogous studies for other nickel phosphinine complexes. Ni(0) complexes, in general, are very versatile and find various applications in electrochemistry and catalysis. For example, Ni(0) species play important roles in cross-coupling reactions (e.g., Kumada reaction) because of their redox properties and ability to undergo oxidative additions with substrates such as aryl halides. Thus, we were motivated to investigate the reactivity of 1 with the simple model cross-coupling substrates bromobenzene and iodobenzene (Scheme 3). When 1 equiv of bromobenzene was added to 1 in benzene, no change was observed at room temperature, either by eye or by ^31P{1H} NMR spectroscopy. However, when the temperature was increased to 60 °C, a color change from deep purple to fluorescent red could be observed. According to GC-FID analysis, a maximum of ca. 70% bromobenzene was consumed (despite complete consumption of 1, as indicated by ^31P{1H} NMR spectroscopy) alongside the formation of a new species which shows a single ^31P{1H} resonance arising as a sharp singlet at δ = 4.7 ppm. This signal splits into a complex multiplet in the ^31P NMR spectrum. In order to characterize the corresponding species 2, the solvent of the fluorescent red reaction mixture was completely evaporated, and the remaining residue was extracted into n-hexane.

Crystals of 2 were obtained by slow evaporation of the resulting fluorescent orange n-hexane solution. Single-crystal XRD revealed its identity as a 1,1-diphenyl-λ^3-phosphinine (Figure 5), which accounts for the observed fluorescent properties. This structure formally arises from the addition of two phenyl groups to the P center of the

---

**Scheme 3. Reaction of 1 with Haloarenes PhX (X = Br, I) at 60 °C**

(i) Reaction in benzene, 2 (15% isolated yield) and [1]Br (45%). (ii) Reaction in toluene, 2 (30%) and 3 (64%). Possible contributions to the electronic structure of 3, including Ni d^8 (3-I), Ni d^9 (3-II), and Ni d^8 (3-III) centers.
phosphinine ligand L. As reported for other \( \lambda^3 \)-phosphinines, the P\(-\)C single bonds (mean distance 1.763 Å vs sum of covalent radii 1.86 Å), while the C\(-\)C bonds are in the range of C\(-\)C double bonds (mean distance 1.395 Å vs sum of covalent radii 1.34 Å).77 The almost planar phosphorus heterocycle is characterized by the small fold angle of 1.50(8)°, and the phenyl substituents at the P atom show bond lengths consistent with P\(-\)C single bonds (mean distance 1.817 Å).77 Similar to the only other \( \lambda^3 \)-pyridylphosphinine reported in the literature, the pyridyl group is essentially coplanar with the phosphinine ring \( [N1\text{--}C6\text{--}C5\text{--}C4 = 174.9(1)^\circ] \), while the phenyl groups in 4 and 6 positions of the heterocycle are not in-plane with the central hexagon. This is in accordance with our expectations for the steric demand of the N lone pair, which is smaller than a CH group of a phenyl moiety.76,78

The \( n \)-hexane-insoluble material remaining after separation of 2 could be dissolved in THF to give a red NMR-silent solution. Single crystals suitable for XRD were obtained after layering with \( n \)-hexane. The molecular structure revealed the formation of a trigonal-bipyramidal Ni(I) complex, \([1]Br\), containing two L and one additional bromide ligand (Figure 5), in a structure very similar to those observed for \([1]Br\) and \([1'(THF)]PF_6\). The \( \sigma \) bonds between the Ni center and the donor atoms of L (mean distances: Ni\(-\)N 2.105 Å and Ni\(-\)P 2.208 Å) are elongated compared to 1, most likely caused by the higher coordination number of the Ni atom. In comparison with the cationic complexes \([1]BF_4\) and \([1'(THF)]PF_6\), \([1]Br\) displays similar Ni\(-\)P distances but shortened Ni\(-\)N distances. The Ni\(-\)Br1 bond length of 2.4889(4) Å is in the range of covalent Ni\(-\)Br bonds (sum of covalent radii 2.430 Å),77 while the geometry index of \( \tau_b = 0.79 \) is consistent with a distorted trigonal-bipyramidal structure (Table 1). The magnetic moment of \([1]Br\) \( [2.1(1) \mu_B \text{ in THF-}d_8 \text{ at 300 K}] \) was determined by the Evans NMR method and is consistent with an \( S = 3/2 \) Ni(I) center (Table 1). Additionally, EPR measurements in a toluene glass at 20 K indicate the presence of a Ni(I) species with a metal-centered radical (Figure S24; simulated g tensor: \( g_{xx} = 2.190, g_{yy} = 2.125, \) and \( g_{zz} = 2.060)\), similar to \([1]BF_4\) and \([1'(THF)]PF_6\). The calculated spin density on Ni (0.82) as well as the calculated g tensor are in good agreement with the experimental data (Table S5).

Although Ni-mediated cross-coupling reactions usually involve Ni(0) and Ni(II) species,79 Ni(I) complexes analogous to \([1]Br\) have also been suggested as viable intermediates and may also show pronounced catalytic activity.76,78 Some other nickel(I) halide complexes have also been isolated from the reaction of Ni(0) complexes with PhX.85,86

The reaction of 1 was also investigated toward 1 equiv of iodobenzene under analogous conditions (toluene, 60 °C) but was found to lead to a significantly different outcome. In this case, full consumption (>99%) of iodobenzene took place according to GC-FID analysis. Compound 2 could again be isolated after extraction with \( n \)-hexane. However, the remaining residue showed a deep-green color rather than deep red, as had been observed using PhBr. Crystals were obtained from a toluene solution of this residue, after layering with \( n \)-hexane. Single-crystal XRD revealed a new product 3, which shows a significantly different structure in the solid state than \([1]Br\) (Figure 5). Most notably, 3 has clearly lost one of its phosphinine ligands L. Moreover, the Ni center—which is tetrahedrally coordinated (\( \tau_b = 0.95; \) Table 1)—is bound to two iodide ligands, as well as by a formally anionic 1-Ph-\( \lambda^3 \)-phosphinine ligand, which is derived by the addition of an extra phenyl substituent to ligand L. The Ni\(-\)P distance \( [2.2509(6) \text{ Å}] \) is in the range of Ni\(-\)P single bonds (sum of covalent radii 2.21 Å), while the Ni\(-\)N1 \( [1.9941(19) \text{ Å}] \) bond length is elongated in a manner similar to in 1. The Ni\(-\)N1 distances (mean value 2.514 Å) are in the range of typical Ni\(-\)I bonds (sum of covalent radii 2.43 Å). We anticipate that 3 is formed by oxidative addition of the carbon\(-\)halogen bond to the Ni center and subsequent transfer of the aryl fragment to the ligand. A similar reaction of a (TPP)Ni(C_6H_5)(acac) complex (TPP = 2,4,6-triphenyl-phosphinine; acac = acetylacetone) has been proposed by Lehmkuhl et al.87

Unlike \([1]Br\), the electronic structure of 3 cannot be assigned easily. The measured value of the magnetic moment
[3.2(1) in THF-$d_4$ at 300 K] lies between those expected for two and three unpaired electrons (Table 1). The magnetic susceptibility of a solid sample of 3 was measured with a SQUID magnetometer in the temperature range 100 K < $T$ < 400 K. The results are shown in Figure 6. These data reveal that the magnetic moment continuously increases from $\mu_{\text{eff}} = 2$ at 105 K to 4 at 400 K. These data suggest that a thermally activated spin transition occurs from an $S = 1/2$ ground state into an $S = 3/2$ excited state. The latter dominates at the upper limit of the accessible temperature window. In addition, it is noteworthy that the measured effective moment significantly exceeds the expected spin-only value for an $S = 3/2$ system at high temperature ($T > 350$ K). This observation might be explained by unquenched orbital contributions to the magnetic moment of the $S = 3/2$ species, which is a common phenomenon for complexes with a tetrahedral $d^3$ configuration at the metal atom.87

The EPR spectrum of 3 recorded at 20 K in a toluene glass shows overlapping and broad signals for the $g$ tensors. The simulated $g$ tensor ($g_{11} = 2.390, g_{22} = 2.242,$ and $g_{33} = 2.120$) is consistent with a Ni-centered radical in a rhombic $S = 1/2$ system (Figure 7).

In the case of 3, the experimental data could not be reproduced in good agreement using standard DFT (for details, see the SI). Furthermore, a rather large spin density was calculated at the Ni atom (1.23), hinting at a broken-symmetry solution. Indeed, a broken-symmetry treatment at the TPSS0-D3BJ/def2-TZVP level shows an interesting electronic structure. Here, an intermediate-spin Ni center ($S = 1, d^3$) is antiferromagnetically coupled to a phosphacyclohexadienyl radical, resulting in an overall $S = 1/2$ species (3-II; for details, see the SI). A small energy separation between the broken-symmetry doublet and quartet states was calculated by DFT ($\Delta E_{\text{doublet-quartet}} = 1.1$ kcal-mol$^{-1}$ at the TPSS0-D3BJ/def2-TZVP level of theory). However, dedicated multi-reference calculations, namely, CASSCF-NEVPT2 with an active space of 13 electrons in 12 orbitals, reveal a much more complicated electronic structure of the $S = 1/2$ ground state of 3. These calculations show that all three mesomeric structures (3-I, 3-II, and 3-III), shown in Scheme 3 (vide supra), contribute significantly to the overall electronic structure of 3. Thus, 3 exhibits significant multireference character, which makes an assignment of the oxidation state of the Ni atom somewhat ambiguous. However, both broken-symmetry DFT and CASSCF clearly show the redox-active behavior of the anionic 1-phenylphosphacyclohexadienyl ligand in 3. Furthermore, the CASSCF-NEVPT2 estimation of the $g$ tensor is in qualitative agreement with the experiment and reproduces the observed significant anisotropy of the $g$ tensor ($g_\parallel = 2.562, g_\perp = 2.227, g_e = 2.173$).

**CONCLUSION**

In summary, we have presented the synthesis and crystallographic characterization of the homoleptic 2-(2'-pyridyl)-4,6-phosphininenickel(0) complex 1. First reactivity studies on the phosphinine-based Ni complex 1 show that it can be conveniently oxidized using ferrocenium salts FeC$.X$. Oxidation with FeBF$_4$ and FeC$_2$P$_6$ leads to the formation of Ni(1) complexes [1]BF$_4$ and [1(THF)]PF$_6$, which possess similar trigonal-bipyramidal structural motifs. Oxidation of 1 with FeC(BA$^-$)$_2$, on the other hand, affords a dicationic dinuclear Ni(1) complex [1$_2$](BA$^-$)$_2$, containing a bridging phosphinine ligand in the rare $\mu_2$-coordination mode. The reaction of 1 with bromobenzene leads to formation of the $\lambda_2$-phosphinine 2 and the trigonal-bipyramidal Ni(1) complex [1]Br. In contrast, the reaction of 1 with iodobenzene results in the formation of 2 and the tetrahedral Ni complex 3, which is based on a formally anionic 1-Ph-A$^-$-phosphinine. Quantum-chemical calculations on 3 highlight the redox-active behavior of the phosphinine moiety, which prohibits a clear assignment of an oxidation state to the Ni center. The carbon–halogen bond splitting by a transition-metal complex and transfer of the aryl fragment to the ligand highlight the chemical noninnocence of phosphinine ligands in transition-metal complexes. Investigations on the mechanism of formation of [1]Br and 3 and further reactivity studies on 1 are currently underway in our laboratories.

**ASSOCIATED CONTENT**

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.inorgchem.0c01115.
Full synthetic details, NMR, UV–vis, and EPR spectra, and crystallographic refinement and computational details (PDF)

Accession Codes
CCDC 2007973 and 1988434–1988439 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: +44 1223 336033.

Author Information

Corresponding Authors
R. Wolf — Institute of Inorganic Chemistry, Universität Regensburg, 93040 Regensburg, Germany; Email: robert.wolf@ur.de

B. de Bruin — van’t Hoff Institute for Molecular Sciences, University of Amsterdam, 1098 XH Amsterdam, The Netherlands; orcid.org/0000-0002-3482-7669; Email: b.debruin@uva.nl

C. Müller — Institute of Chemistry and Biochemistry, Freie Universität Berlin, 14195 Berlin, Germany; Email: c.mueller@fu-berlin.de

Authors
J. Leitl — Institute of Inorganic Chemistry, Universität Regensburg, 93040 Regensburg, Germany
P. Coburger — Institute of Inorganic Chemistry, Universität Regensburg, 93040 Regensburg, Germany
D. J. Scott — Institute of Inorganic Chemistry, Universität Regensburg, 93040 Regensburg, Germany
C. G. P. Ziegler — Institute of Inorganic Chemistry, Universität Regensburg, 93040 Regensburg, Germany
G. Hiermeier — Institute of Inorganic Chemistry, Universität Regensburg, 93040 Regensburg, Germany
N. P. van Leest — van’t Hoff Institute for Molecular Sciences, University of Amsterdam, 1098 XH Amsterdam, The Netherlands
G. Hörner — Department of Chemistry, Inorganic Chemistry IV, Universität Bayreuth, 95440 Bayreuth, Germany; orcid.org/0000-0002-3883-2879

Complete contact information is available at: https://pubs.acs.org/10.1021/acs.inorgchem.0c01115

Notes
The authors declare no competing financial interest.

Acknowledgments
We thank Prof. Birgit Weber (Universität Bayreuth, Bayreuth, Germany) for access to a SQUID magnetometer and helpful discussions. Funding by the Deutsche Forschungsgemeinschaft (Grants WO1496/9-1 and MU1657/5-1) and Alexander von Humboldt Foundation (postdoctoral fellowship to D.J.S.) is gratefully acknowledged.

References


(29) See ref 28 for oxidative reactions of Ni(biphosphinine) complexes with C3Cl6.


(47) Schmid, B.; Venanzi, L. M.; Gerfin, T.; Gramlich, V.; Mathey, F. Synthesis and Spectroscopic Properties of the Complexes [M2(DIEN)2(NIPHOS)2][SF6]2 (M = Ir, DIEN = 1,5-Cyclo-
octadiene (COD); M = Rh, DIEN = Norbornadiene (NBD); NIPHOS = 2-(2′-Pyridyl)-4,5-Dimethylphosphinine. X-Ray Crystal Structure of [Ir(COD)(NIPHOS)][[SbF_6]_2]. Inorg. Chem. 1992, 31 (24), 5117–5122.


(65) Mao, Y.; Lim, K. M. H.; Li, Y.; Ganguly, R.; Mathey, F. The Original Coordination Chemistry of 2-Phosphophenol with Copper(I) and Gold(I) Halides. Organometalics 2013, 32 (12), 3562–3565.


