

Efficient Electronic Communication of Two Ruthenium Centers through a Rigid Ditopic N-Heterocyclic Carbene Linker

Michael Nussbaum,^[a] Oliver Schuster,^{*,[a]} and Martin Albrecht^{*,[a, b]}

Abstract: A ditopic benzobis(carbene) ligand precursor was prepared that contained a chelating pyridyl moiety to ensure co-planarity of the carbene ligand and the coordination plane of a bound octahedral metal center. Bimetallic ruthenium complexes comprising this ditopic ligand [L_4Ru-C,N -bbi- C,N - RuL_4] were obtained by a transmetalation methodology (C,N -bbi- C,N =benzobis(N -pyridyl- N' -methyl-imidazolylidene)). The two metal centers are electronically decoupled when the ruthenium is in a pseudotetrahedral geometry imparted by a cymene spectator ligand ($L_4=[(cym)Cl]$). Ligand exchange of the Cl^- /cymene ligands for two bipyridine or four MeCN ligands induced a change of the coordination

geometry to octahedral. As a consequence, the ruthenium centers, separated through space by more than 10 Å, become electronically coupled, which is evidenced by two distinctly different metal-centered oxidation processes that are separated by 134 mV ($L_4=[(bpy)_2]$; bpy=2,2'-bipyridine) and 244 mV ($L_4=[(MeCN)_4]$), respectively. Hush analysis of the intervalence charge-transfer bands in the mixed-valent species indicates substantial valence delocalization in both complexes

(delocalization parameter $\Gamma=0.41$ and 0.37 in the bpy and MeCN complexes, respectively). Spectroelectrochemical measurements further indicated that the mixed-valent Ru^{II}/Ru^{III} species and the fully oxidized Ru^{III}/Ru^{III} complexes gradually decompose when bound to MeCN ligands, whereas the bpy spectators significantly enhance the stability. These results demonstrate the efficiency of carbenes and, in particular, of the bbi ligand scaffold for mediating electron transfer and for the fabrication of molecular redox switches. Moreover, the relevance of spectator ligands is emphasized for tailoring the degree of electronic communication through the benzobis(carbene) linker.

Keywords: ditopic ligands • metal–metal communication • mixed-valent state • N-heterocyclic carbene • ruthenium

Introduction

Molecular switches provide intriguing opportunities as active entities for the storage and processing of data.^[1] Metal complexes are particularly attractive units for such switching purposes, since they offer multiple read-out functions due to their potential activity in non-invasive events, such as reversible redox,^[2] spin transitions,^[3] and charge-transfer^[4] processes. The resulting electronic, magnetic, and optical signals constitute a diagnostic probe for the state of the switch.^[1–4] The information density can be considerably enhanced when metal centers are mutually coupled. For example, redox switching of a bimetallic species gives access to a stable mixed-valent state in addition to the fully ox-

idized and reduced states, if the metal centers are electronically coupled.^[5] Such mixed-valent states have distinct properties that depend on the degree of coupling.^[6] A valence-localized system will thus provide access to a molecular diode with the oxidized entity as acceptor and the reduced unit as donor site.^[7] Valence-delocalized systems will be excellent constituents of molecular wires^[8] with the possibility for tailoring activity through modification of the bridging ligand.

Intramolecular metal–metal coupling is critically dependent on the ability of the bridging ligand to function as an electronic relay.^[9] The two currently most popular classes of ligands that promote coupling in bimetallic systems are comprised of either ditopic bi- and oligopyridines,^[10] and of oligoacetylenes.^[11] While the latter type of ligands offers an excellent electronic link between the metal and the bridging ligand with a high degree of $d_M-\pi_L$ overlap, especially in metal–alkylidene and metal–alkynylidene oxidation states, the oligopyridine family dwells on the p -acidity of the chelating pyridyl ligand and its synthetic versatility for functionalization. In a simplistic view, N-heterocyclic carbenes (NHCs)^[12] represent a fusion of these two successful classes of ligands, featuring the synthetic versatility^[13] of pyridines with the (partial) p component of the $M-C_{NHC}$ bond^[14] reminiscent of acetylenes. Stimulated by these considerations, a variety of ditopic NHC ligands have been developed (e.g. **A–C**, Figure 1).^[15] However, even in complexes containing

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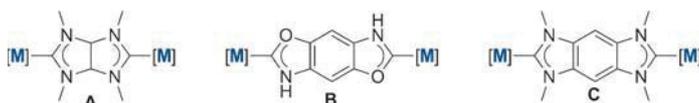


Figure 1. Relevant examples of ditopic NHC ligands.

the apparently highly conjugated benzobis(imidazolyliene) bridging ligand **C**, the metal–metal interaction was mediocre at best. Different explanations for the weak coupling of the metal centers in this molecular configuration have been put forward,^[16] including low $d_M-\pi_L$ overlap due to a mismatch of the ligand 2p orbital and the metal 4d orbitals,^[17] poor p electron delocalization between the arene and the NCN units in the bridging ligand,^[18] and an unfavorable orientation of the relevant d_M orbitals with respect to the π_L system that minimizes overlap.^[19] We have addressed this latter issue by incorporating rigid pyridyl donor groups into the NHC wingtips, which induce chelation and consequentially impart a coaxial arrangement of the metal and ligand orbitals, a concept that has been widely exploited in bipyridine chemistry.^[20] The constrained NHC coordination substantially enhances the intermetallic coupling between two ruthenium centers bridged by ligands of type **C**. The electronic Ru...Ru coupling is critically dependent on the nature of the spectator ligands and on the coordination geometry they impose. These results indicate that appropriately designed dicarbenes provide efficient bridging ligands for the fabrication of molecular switches, and they emphasize the relevance of the mutual orientation of ligand and metal orbitals.

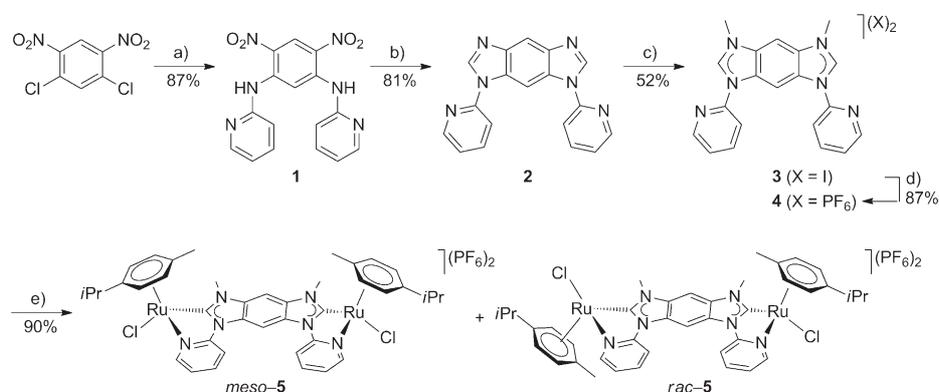
Results and Discussion

Synthesis of a ditopic pyridyl–carbene precursor and dimeric Ru₂ complexes: The synthetic approach towards chelating dicarbene ligands containing pyridyl donor groups was inspired by a methodology developed by Bielawski and co-workers for related monodentate ligand platforms (Scheme 1).^[21] Accordingly, amination of 2,4-dinitro-1,5-dichlorobenzene with 2-aminopyridine furnished the pyridyl-functionalized product **1**. Aminations performed in EtOH at 120 °C afforded **1** only after extended reaction times and furnished monosubstituted compounds as major products due presumably to the deactivating effect of the first pyridyl substituent. Yields and purity were substantially higher in a solvent-free reaction when using a large excess of 2-aminopyridine at 90 °C. Subsequent heterocycle formation was achieved

in a palladium-catalyzed one-pot reaction involving reduction of the nitro groups with formic acid followed by in situ cyclization with orthoformate, which afforded the pyridyl-substituted benzobis(imidazole) **2**. Cyclization induced a marked upfield shift of the aryl proton resonances from $\delta_H=9.38$ and 10.26 in **1** to 8.32 and 8.92 ppm in **2**. Moreover, a singlet at $\delta_H=8.62$ ppm was attributed to the imidazole proton ($\delta_C=142.2$ ppm). The reaction proceeds in high yields (> 80 %). The largest side product was identified by HPLC-MS (m/z 357 g mol^{-1}) as an ethoxy-adduct of **2**, possibly originating from incomplete EtOH elimination upon cyclization with HC(OEt)_3 . Formation of **2** was unambiguously confirmed by an X-ray crystal-structure determination (see Figure S1 in the Supporting Information), which revealed an essentially planar molecular structure for **2**. The central benzene ring is elongated along the $\text{CH}\cdots\text{CH}$ vector, a feature that was previously observed by Bielawski^[17] and us,^[18] albeit in a less-pronounced manner. Pairs of molecules of **2** are stacked along the crystallographic c -axis and give rise to supramolecular zig-zag layers in the a - b plane (see Figure S1 in the Supporting Information).

The benzobis(imidazole) was selectively quaternized with excess MeI to yield the benzobis(imidazolium) salt **3**. The substantial downfield shift of the NCHN proton (from $\delta_H=8.62$ to 10.84 ppm) and to a lesser extent of the aryl protons ($\delta=$ ca. 0.8 ppm) in the ¹H NMR spectrum supports alkylation of the imidazole unit. Moreover, the NMR resonances of the pyridyl fragment are essentially unaltered upon alkylation, which indicates that imidazole methylation is selective. Anion exchange with aqueous $(\text{NH}_4)\text{PF}_6$ gave **4** and was carried out to avoid issues due to potential halide scrambling in the subsequent (trans)metalation step.

Complexation of the ditopic carbene precursor was accomplished according to a classical metalation–transmetalation protocol by using Ag_2O and $[\text{RuCl}_2(\eta^6\text{-}p\text{-cymene})]_2$.^[22] While the transruthenation afforded pure products when performed in CH_2Cl_2 , reaction with Ag_2O in this solvent was exceedingly slow, presumably due to the limited solubility of the dicationic salt **4**. Therefore, formation of the carbene



Scheme 1. Synthesis of complexes **5**: a) 2-NH₂-pyridine, 90 °C, 3 d; b) HCOONa, HCOOH, Pd/C, HC(OEt)₃, 100 °C, 36 h; c) MeI, MeCN, microwave irradiation, 110 °C, 1 h; d) (NH₄)PF₆, H₂O; e) Ag₂O, MeOH, 24 h, then $[\text{RuCl}_2(p\text{-cymene})]_2$, CH₂Cl₂, 3 d.

silver intermediate was performed in MeOH and the solvent was then changed to CH₂Cl₂ for the transmetalation, without any purification or characterization of the probably polymeric silver complex.^[23] Complex **5** is an air-stable orange solid. While HPLC-MS analysis suggests a pure compound, NMR spectroscopic analysis shows two distinct sets of signals that were attributed to the *rac* and *meso* diastereoisomers of **5** as a consequence of the chirality of the complex at each ruthenium center.^[15,24] Solubility tests revealed that one of the two isomers is less soluble in MeCN and was thus successfully separated by repetitive precipitation and filtration. This isomer is stable in solution over several days, which demonstrates that racemization at ruthenium is very slow. Hence, 2D NMR spectroscopic measurements allowed all resonances to be assigned to one or the other isomer. However, we were unable to confidently assign the *meso* and *rac* designations based on spectroscopic data, and crystallization attempts have failed to date.

Successful double metalation was indicated by the pertinent cymene/benzobis(carbene) integral ratios and by the carbenic resonance at $\delta_{\text{C}}=202.8$ and 203.3 ppm, corresponding to more than $\delta=50$ ppm downfield shift compared to the ligand precursor (Table 1). As expected for a chiral com-

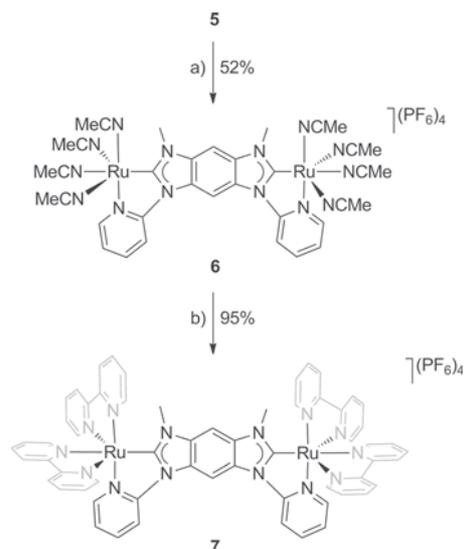
Table 1. Selected NMR spectroscopic shifts of complexes **5–7** and the ligand precursor **4**.^[a]

Compound	δ_{C} NCN	δ_{H} H _{Ar}	δ_{C} C _{Ar} -H	δ_{H} H _{Ar'}	δ_{C} C _{Ar'}
4	146.7	9.65	103.6	9.08	99.7
5	203.3	8.56	97.8	8.20	96.7
5'	202.8	7.84	95.9	7.69	95.9
6	207.6	8.58	95.9	8.11	94.8
7	211.1	8.92	96.2	7.69	93.7

[a] All data in CDCl₃ except **4** (in [D₆]DMSO); Ar denotes the position β to the pyridyl-substituted nitrogen, Ar' the position β to the methyl-substituted nitrogen atom; **5** and **5'** are the two diastereoisomers of **5** (*meso* and *rac*).

plex, all four aromatic cymene protons are magnetically inequivalent, and the *i*Pr group appears as two distinct high-field doublets for each isomer. Chelation of the pyridyl group is supported by the marked deshielding of the C6-bound pyridyl protons ($\delta = >0.4$ ppm, see Figure S2 in the Supporting Information). Moreover, the resonances of the protons of the central arene ring experience a diagnostic upfield shift upon ruthenation and appear at $\delta_{\text{H}}=7.84$ and 7.69 ppm in one isomer, and at $\delta_{\text{H}}=8.20$ and 8.56 ppm in the other one (cf. $\delta_{\text{H}}=9.08$ and 9.65 ppm in the ligand precursor **4**). These shift differences suggest a strong electronic coupling of the metal center with the central arene moiety.

Substitution of the ancillary ligands was achieved according to established procedures.^[25] Thus, reaction of complex **5** with AgPF₆ in MeCN at reflux temperature gave the solvento complex **6** (Scheme 2). The reaction progress is conveniently monitored optically as the bright-orange solution turns colorless upon displacement of the Cl⁻ and cymene ligands. The NMR spectra of complex **6** reveals only one set of signals and thus supports the notion of diastereomeric



Scheme 2. Synthesis of complexes **6** and **7**: a) AgPF₆, MeCN, reflux, 20 h; b) 2,2'-bipyridine, MeNO₂, reflux, 20 h.

mixtures in **5**. Evidently, the ruthenium centers are achiral in **6**. The carbene carbon atom is further deshielded (from $\delta_{\text{C}}=203$ to 207.6 ppm), and also the central arene protons undergo a slight downfield shift, $\delta_{\text{H}}=8.11$ and 8.58 ppm (see Figure S2 in the Supporting Information). Three different types of MeCN ligands are resolved, with one set ($\delta_{\text{H}}=2.09$ ppm) twice as intense as the other two sets ($\delta_{\text{H}}=2.60$ and 1.96 ppm, respectively). These features support an octahedral coordination geometry of the ruthenium centers with two MeCN in a mutual *trans* position and hence symmetry related. Based on the stronger donor properties of NHCs versus pyridine ligands, the most deshielded resonance was tentatively attributed to the MeCN ligand *trans* to the pyridyl donor and the resonance at highest field to the MeCN *trans* to the carbene.

Crystals of **6** suitable for an X-ray diffraction analysis were obtained by slow diffusion of Et₂O into a MeCN solution of the complex. The molecular structure (Figure 2, Table 2) confirms the octahedral geometry of the two ruthenium centers, with only little angular distortion. The bite angle of the *C,N*-bidentate ligand is 78.5(4)°, which is in line with related structures featuring pyridyl-substituted NHC ligands.^[26] As a consequence of this acute bite angle, the overall molecular shape of **6** is curved.^[27] The Ru–N bonds for the MeCN ligands *trans* to the carbenes (2.115(6) and 2.113(6) Å) are significantly longer than the remaining Ru–N distances (2.02–2.04 Å, Table 1). This elongation is in agreement with a substantially stronger *trans* influence of the carbene ligand, which is often correlated with its strong σ -donor ability. The metal–metal separation in **6** is 10.537(2) Å and hence some 0.2 Å shorter than the Ru...Ru distance in a bimetallic complex with an analogous ditopic ligand that lacks chelating wingtip groups.^[18] This difference is almost fully accounted for by the shorter Ru–C bond upon chelation, namely, 1.95(1) Å in **6** versus 2.065(4) Å in the monodentate analogue.

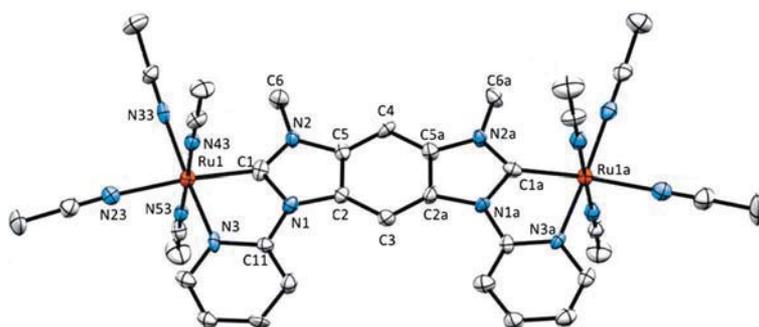


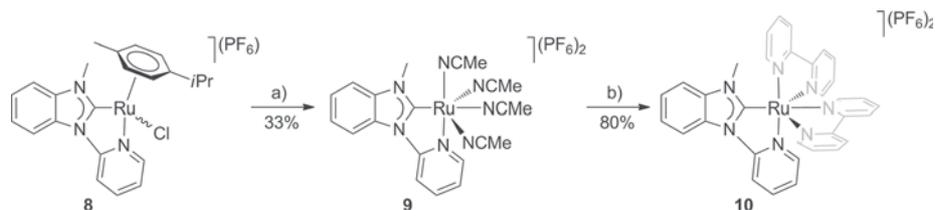
Figure 2. ORTEP view of complex **6** (50% probability ellipsoids; Hydrogen atoms, counter-ions and co-crystallized MeCN molecules omitted for clarity).

Table 2. Selected bond lengths [Å] and angles [°] for complexes **6** and **10**.

	6 ^[a]		10
Ru1–C1	1.955(6)	1.945(6)	1.979(4)
Ru1–N3	2.053(6)	2.059(6)	2.068(5)
Ru1–N23	2.115(6)	2.113(6)	2.126(3)
Ru1–N33	2.042(6)	2.038(6)	2.046(6)
Ru1–N43	2.034(6)	2.025(6)	2.074(5)
Ru1–N53	2.026(6)	2.022(6)	2.062(6)
C1–N1	1.388(9)	1.393(9)	1.393(8)
C1–N2	1.346(9)	1.380(8)	1.350(7)
N1–C2	1.346(9)	1.413(8)	1.416(5)
N2–C5	1.395(8)	1.368(9)	1.413(6)
N1–C1–N2	105.2(6)	105.0(6)	104.2(5)
C1–N1–C11	116.2(6)	116.7(6)	117.3(5)
C1–Ru1–N3	78.4(3)	78.5(3)	77.7(2)
C1...C1a	6.690(8)	6.690(8)	–
Ru...Ru	10.537(2)	10.537(2)	–

[a] Right column pertains to east end of the complex (atom labels in Figure 2).

Subsequent ligand exchange with bpy was straightforward in MeNO₂ and yielded bright orange complex **7** in an excellent yield (90%). Introduction of bpy directly starting from the cymene complex **5** in the presence of AgPF₆ was less successful and afforded a mixture of compounds containing both bipyridine and MeCN ligands bound to the ruthenium center. The NMR spectroscopic data of complex **7** indicates the presence of two diastereoisomers; however, the congestion of the aromatic region due to the presence of two sets of five inequivalent pyridyl fragments did not allow proton and carbon resonances to be fully resolved and unambiguously assigned. Most diagnostic are the shielding of the pyridyl proton in α -position as well as a shift of the carbene resonance to higher field ($\delta_C = 211.1$ ppm). In addition, the chemical shift difference between the two central arene protons increases upon bpy coordination. The two resonances appear in complex **7** at $\delta_H = 7.69$ and 8.92 ppm and are thus



Scheme 3. Synthesis of complexes **9** and **10**: a) AgPF₆, MeCN, reflux, 20 h; b) 2,2'-bpy, MeNO₂, reflux, 20 h.

some $\delta = 1.2$ ppm apart, whereas $\delta < 0.5$ ppm in **6** (see Figure S2 in the Supporting Information). Again, the sensitivity of these central protons towards peripheral ligand exchange reactions suggests that electronic coupling of the metal center with these protons and hence with the center of the molecule is effective.

Synthesis of monocarbene complexes:

For comparative purposes, analogous monometallic complexes were prepared by a route similar to the synthesis of the bimetallic complexes **5–7**. Thus, stirring of complex **8**^[25e] at reflux temperature in MeCN in the presence of AgPF₆ gave the solvento complex **9** and subsequent addition of bpy yielded complex **10** as a monometallic analogue of **7** (Scheme 3). Similar spectroscopic trends as deduced for the bimetallic complexes were observed. For example, the resonance of the carbene nucleus gradually shifts upfield upon introducing MeCN and subsequently bpy spectator ligands ($\delta_C = 198.2, 204.0,$ and 207.9 ppm for **8, 9,** and **10**, respectively). Moreover, four coordinated MeCN molecules were spectroscopically identified in **9** in the 1:2:1 ratio as discussed for the bimetallic complex **6** (see above).

Crystals of **9** featured severe disorder in co-crystallized solvent molecules and PF₆⁻ anions, which prevented refinement to an acceptable convergence. Hence, discussion of structural data of the cationic section is limited (see Figure S3 in the Supporting Information). Crystals of **10** (Figure 3) were of better quality. The molecular structure

confirms the expected connectivity pattern. The Ru–C (1.979(4) Å) and Ru–N_{pyridine} bonds (2.068(5) Å) are slightly longer than in **6**, and consequentially, the bite-angle is more acute, 77.7(2)°. As in **6** and **9**, the carbene exerts the strongest *trans* influence and the Ru–N23 bond *trans* to the NHC moiety is significantly longer (2.126(3) Å) relative to the other Ru–N_{pyridine} bonds (2.046(6)–2.074(5) Å).

Electrochemical analyses: Cyclovoltammetry (CV) measurements of complex **5** in CH₂Cl₂ show a single and reversible oxidation/reduction process with $E_{1/2}$ (**5**²⁺/**5**) = +1.606 V versus SCE (see Figure S4 in the Supporting Information,

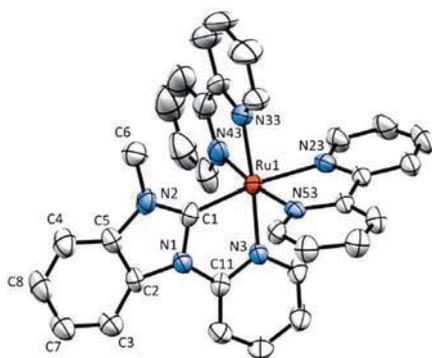


Figure 3. ORTEP view of **10** (50% probability ellipsoids; hydrogen atoms and non-coordinating PF_6^- anions omitted for clarity).

Table 3). This redox potential is about 0.5 V higher than that of neutral $[\text{RuCl}_2(\text{cym})(\text{NHC})]$ complexes^[18] and thus demonstrates the lower electron density in the cationic chelates, which is also reflected in the NMR deshielding of the

Table 3. Electrochemical data for complexes **5–10**.^[a]

Complex	solvent	$E_{1/2}/\text{V}$	$\Delta E/\text{mV}^{[b]}$	$\Delta E_{1/2}/\text{mV}$	$K_c^{[c]}$
5 / Ru_2	CH_2Cl_2	+1.606	151	–	–
6 / Ru_2	MeCN	+1.644, +1.888	n.d.	244	$10^{4.13}$
7 / Ru_2	CH_2Cl_2	+1.434	122	–	–
7 / Ru_2	MeNO_2	+1.283, +1.417	95, 95	134	$10^{2.26}$
8 / Ru	CH_2Cl_2	+1.583	116	–	–
9 / Ru	MeCN	+1.619	104	–	–
10 / Ru	CH_2Cl_2	+1.446	103	–	–
10 / Ru	MeNO_2	+1.287	110	–	–

[a] 0.1 M $[\text{NBu}_4][\text{PF}_6]$ as the supporting electrolyte, sweep rate: 50 mV s^{-1} , potentials referenced to internal Fc^+/Fc ($E_{1/2} = 0.41 \text{ V}$). [b] $\Delta E_p = E_{pa} - E_{pc}$. [c] $K_c = \exp(\Delta E_{1/2}/0.059)$.

ligand protons (see above). No further oxidation occurred up to +1.9 V. The cathodic current of the oxidation was considerably larger than the anodic current. Such behavior has previously been attributed to the adsorption of the oxidized analyte to the electrode surface.^[28] In contrast, the corresponding monometallic complex **8** shows no such adsorption effect, but a similar $E_{1/2}$ ($\mathbf{8}^+/\mathbf{8}$) = +1.583 V. Differential pulse voltammetry (DPV) measurements did not show any substantial broadening of the signal recorded for **5** relative to that of the monometallic analogue **8**, which indicated that the two metal centers in **5** are essentially decoupled (Class I according to Robin and Day).^[29] Measurements in MeCN or MeNO_2 were inconclusive and showed either an irreversible oxidation (MeCN), or no redox-process at all (MeNO_2). Displacement of the chloride ligand may account for the different behavior of **5** and **8** in these solvents as compared to CH_2Cl_2 .

Complex **6** was measured in MeCN because its solubility in CH_2Cl_2 is low and NMR spectroscopic experiments showed decomposition in MeNO_2 . A reversible redox process was observed at $E_{1/2}$ ($\mathbf{6}^{2+}/\mathbf{6}$) = +1.574 V, comparable to the monometallic model complex **9** ($E_{1/2}$ ($\mathbf{9}^+/\mathbf{9}$) = +1.619 V;

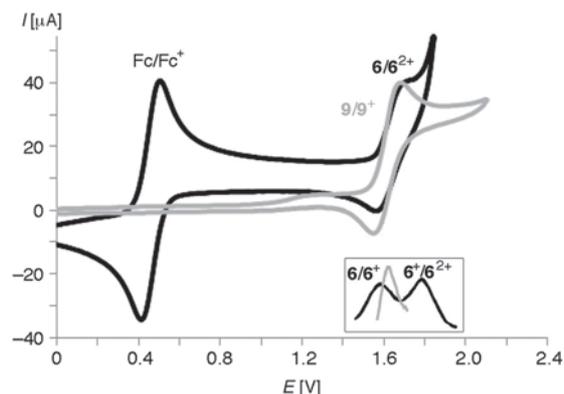


Figure 4. CV plots and relevant DPV sections (inset) of complexes **6** and **9** (1 mM in dry MeCN with 0.1 M $[\text{NBu}_4][\text{PF}_6]$ as the supporting electrolyte, 50 mV s^{-1} scan rate, Fc^+/Fc used as internal standard with $E_{1/2}$ (Fc^+/Fc) = 0.41 V vs. SCE).

Figure 4). No further oxidation occurred up to +2.1 V. Compound **6** is unstable at potentials higher than +1.8 V and decomposed during electrochemical cycles, due presumably to the poor stabilization of the ruthenium(III) center by MeCN ligands. However, DPV measurements in an optically transparent thin-layer electrochemical (OTTLE) cell^[30] revealed two separate processes of similar intensity which were attributed to a stepwise oxidation of the two ruthenium(II) centers with $E_{1/2}$ ($\mathbf{6}^+/\mathbf{6}$) = +1.644 and $E_{1/2}$ ($\mathbf{6}^{2+}/\mathbf{6}^+$) = +1.888 V. These data suggest that a mixed-valence $\text{Ru}^{\text{II}}/\text{Ru}^{\text{III}}$ state is accessible. Based on the 244 mV potential difference between the two oxidation processes and the ensuing comproportionation constant $K_c = 10^{4.13}$,^[31] complex **6** corresponds to a Class II/III borderline system according to the Robin and Day classification.^[29,32] These results demonstrate for the first time that efficient electronic communication of two metal centers through a rigid carbene linker is achievable. This intermetallic coupling confirms that conjugation through the benzobis(carbene) linker is effective, thus corroborating the NMR sensitivity of the central arene protons towards peripheral functionalization (cf. NMR discussion above).

Qualitatively similar data were recorded for the bpy ligated bimetallic complex **7**. CV and DPV measurements in MeNO_2 show two distinct redox processes at $E_{1/2}$ ($\mathbf{7}^+/\mathbf{7}$) = +1.283 and $E_{1/2}$ ($\mathbf{7}^{2+}/\mathbf{7}^+$) = +1.417 V (Figure 5). The lower oxidation potential compared to **6** underlines the stronger donor properties of bpy versus MeCN.^[33] The stability of the oxidized species is much improved and the oxidations are fully reversible. The peak potential separation $\Delta E_{1/2} = 134 \text{ mV}$ is smaller than for the acetonitrile complex **6**, yielding a comproportionation constant $K_c = 10^{2.26}$ (Class II system in the Robin and Day classification). The monometallic analogue **10** undergoes one oxidation with $E_{1/2}$ ($\mathbf{13}^+/\mathbf{13}$) = +1.287 V, identical to the first oxidation of **7**. When measured in CH_2Cl_2 , complex **7** showed only one single redox process in CV and DPV ($E_{1/2}$ ($\mathbf{7}^{2+}/\mathbf{7}$) = +1.434 V). The asymmetric shape of the DPV signal and the large

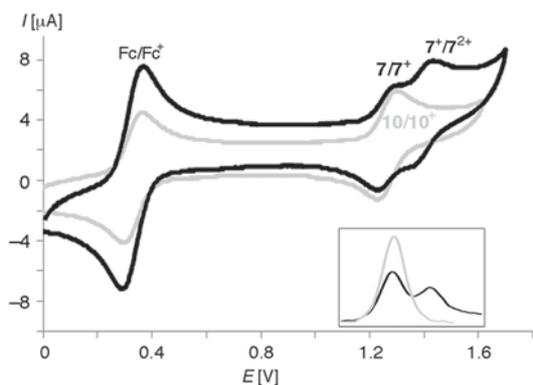


Figure 5. CV plots and relevant DPV sections (inset) of complexes **7** and **10** (1 mm in dry MeNO₂ with 0.1 M [NBu₄][PF₆] as the supporting electrolyte, 50 mV s⁻¹ scan rate, Fc⁺/Fc used as internal standard with $E_{1/2}$ (Fc/Fc⁺) = 0.35 V vs. SCE).

anodic peak current hint to fast decomposition in this solvent (see Figure S5 in the Supporting Information).

Spectroelectrochemistry and Hush analysis: The mixed-valent species derived from complexes **6** and **7** were further investigated by spectroelectrochemical methods by using an OTTL cell.^[30,34] Complex **6** shows a strongly potential-dependent absorption behavior (Figure 6). In the absence of

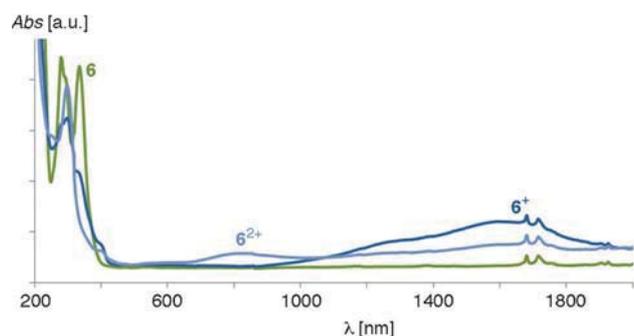


Figure 6. Absorption spectra (MeCN solutions) of complexes **6** at 0.0 V (green), **6**⁺ at +1.37 V (dark blue), and **6**²⁺ at +1.6 V (light blue).

an applied potential, the complex features a strong absorption band at 333 nm, tentatively assigned to a metal-to-ligand charge-transfer band. At +1.37 V, the spectrum shows a markedly reduced intensity of the UV absorption and a broad absorption in the NIR region (λ_{\max} = 1593 nm, ϵ = 13 100 M⁻¹ cm⁻¹) that is diagnostic for an intervalence charge-transfer (IVCT) and thus supports the formation of a mixed-valent Ru^{III}/Ru^{II} species **6**⁺. Further increase of the potential to +1.6 V reduces the IVCT intensity. The fully oxidized Ru^{III}/Ru^{III} species **6**²⁺ is characterized by a shallow absorption maximum at λ_{\max} = 838 nm and a transparent absorption window until nearly 300 nm. Complex **7** features qualitatively identical properties, including a MLCT band at λ_{\max} = 411 nm, a broad IVCT band centered at 1652 nm (ϵ = 2780 M⁻¹ cm⁻¹) upon applying a +1.23 V potential, and

a weak absorption at 733 nm for the fully oxidized species **7**²⁺ at +1.5 V (see Figure S6 in the Supporting Information).

The stability of the oxidized species was evaluated by time-dependent monitoring of the IVCT absorbance and the band around 800 nm, respectively. Thus exposing complex **6** to a potential of +1.46 V for prolonged periods of time induces a gradual decay of the absorbance at 1590 nm to about 50% within 1 h (see Figure S7 in the Supporting Information). At a higher potential (+1.6 V), the decay is even faster and the absorbance at 820 nm is depleted within few minutes ($t_{1/2}$ = ca. 2 min), which suggests a low stability of **6**⁺ and even a lower one for **6**²⁺. Moreover, in a CV measurement of **6** between +1.0 and +2.0 V, the IVCT absorption intensity in the anodic scan is reduced to about 20% of the absorption during the cathodic scan (Figure 7a).^[35] These measurements thus corroborate the poor stability of oxidized **6**⁺ and **6**²⁺ as deduced from initial CV measurements (see above).

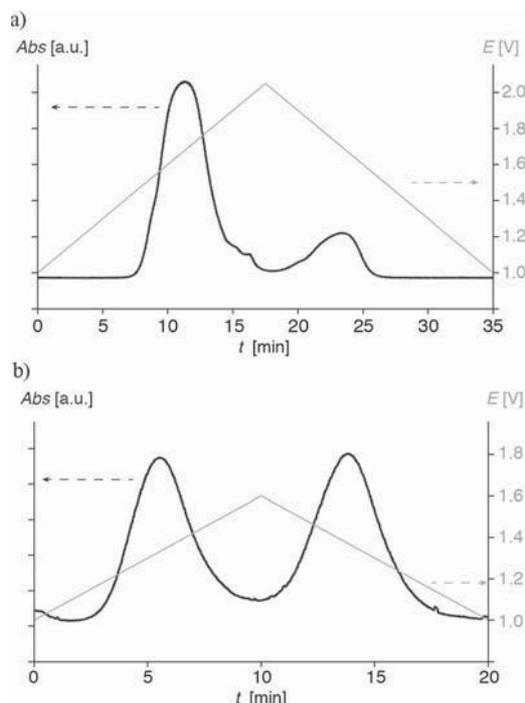


Figure 7. a) Absorbance change at 1590 nm upon a redox cycle between 1.0–2.0 V of **6** (1 mm in MeCN, 1 mV s⁻¹ scan rate); b) Absorbance change at 1730 nm upon a redox cycle between 1.0–1.6 V of **7** (1 mm in MeNO₂, 1 mV s⁻¹ scan rate).

In contrast, the bpy spectator ligands greatly enhance the stability of the oxidized forms of **7** and hence improve the reversibility substantially. For example, the decay of the IVCT absorbance band of the mixed-valent Ru^{II}/Ru^{III} species **7**⁺ is only moderate (~5% decrease within 100 min), probably due to disproportionation. The fully oxidized Ru^{III}/Ru^{III} complex **7**²⁺ did not show any signs of degradation (see Figure S8 in the Supporting Information). Moreover,

the IVCT absorbance of the species at 1730 nm is equally intense in anodic and cathodic scans (Figure 7b). These results underline the relevance of a judicious choice of spectator ligands for both stability and electronic coupling.^[36]

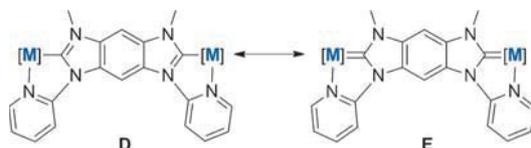
Analysis of the IVCT bands by Marcus' theory^[37] and by methods developed by Hush^[5b,d] allows the degree of coupling of the metal centers to be further detailed (Table 4). The IVCT bands for **6**⁺ and **7**⁺ is considerably asymmetric

Table 4. Spectroscopic analysis of mixed-valence complexes **6**⁺ and **7**⁺.^[a]

	Solvent	n_{\max}	ϵ_{\max}	$\Delta n_{1/2}$		asymmetry ^[b]	H_{ab} ^[c]	Γ
				(obs)	(calcd)			
6 ⁺	MeCN	6277	13078	2412	3808	1.87	729 cm ⁻¹ (0.0904 eV)	0.37
7 ⁺	MeNO ₂	6042	2783	2219	3736	2.06	377 cm ⁻¹ (0.0467 eV)	0.41

[a] n in cm⁻¹, ϵ in M⁻¹cm⁻¹. [b] Asymmetry of the IVCT band determined as a ratio $\Delta n_{1/2}(\text{high})/\Delta n_{1/2}(\text{low})$. [c] Ru...Ru distance $R = 10.537 \text{ \AA}$ from X-ray crystal structure of **6**.

(see Figure S9 in the Supporting Information), with the bandwidth at half height on the high energy side, $\Delta\nu_{1/2}(\text{high})$, about twice as large as that at the low energy side, $\Delta\nu_{1/2}(\text{low})$. The observed bandwidth at half height $\Delta\nu_{1/2}(\text{obs}) = 2412 \text{ cm}^{-1}$ for **6**⁺ and 2219 cm^{-1} for **7**⁺. These values are significantly smaller than the values calculated from the Hush approximation for a valence-trapped class II system ($\Delta\nu_{1/2}(\text{calcd})$ 3808 and 3736 cm^{-1} for **6**⁺ and **7**⁺, respectively).^[5d,38] These deviations suggest a substantial degree of valence delocalization. The electronic coupling parameter H_{ab} was calculated to be $729 (0.090 \text{ eV})$ and $377 \text{ cm}^{-1} (0.047 \text{ eV})$, respectively.^[5] The different H_{ab} values determined for **6**⁺ and **7**⁺ may potentially hint to a strong solvent dependence, which is characteristic for a class II system.^[11b] However, since all further electrochemical and spectroscopic analyses point to borderline class II/III systems, we attribute this difference to a better $d_{\text{M}}-\pi_{\text{L}}$ overlap in complex **6** containing weakly donating ancillary MeCN ligands. Weaker bonding of ancillary ligands strengthens the M-C_{NHC} bond and hence increases the electronic coupling of the metal centers, thus increasing the relevance of resonance structures **E** (Scheme 4). Accord-



Scheme 4. Resonance structures **A** and **B** emphasizing the electronic separation and coupling, respectively, of two coordinated metal centers.

ingly, the valence delocalization in **6**⁺ is more pronounced than in complex **7**⁺ containing two bpy spectator ligands. In agreement with the classification of **6**⁺ and **7**⁺ as borderline valence-trapped/delocalized mixed-valent systems, the delocalization parameter Γ is very close to 0.5.

Conclusion

By introducing rigidly planar chelating pyridyl wingtip groups, ditopic benzobis(carbene) ligands become effective linkers for imparting electronic coupling between two coordinated ruthenium centers. Intermetallic communication is critically triggered by appropriate alignment of the metal 4d and the ligand p orbitals as a key principle to overcome the hitherto poor coupling in dicarbene species. Small overlap and hence insignificant Ru...Ru coupling results from coordination of the chelate to pseudotetrahedral metal coordination geometries as in the cymene complex **5**, and also emanates from monodentate carbene coordination, since the carbene is typically rotated out of the metal coordination plane by 30–45°. In contrast, chelation to octahedral metal centers as in **6** and **7** induces sufficiently large $d_{\text{M}}-\pi_{\text{L}}$ overlap for imparting a high degree of intermetallic electronic communication, reminiscent of the poor versus pronounced π bonding established in pyridine and bipyridine complexes, respectively. These data lend further support to the relevance of π contributions to the metal-NHC bond.

Appropriate substitution of the ancillary ligands enhances the efficiency of the electronic communication and thus provides an opportunity for tailoring the electronic properties of the redox switch. Thus, while spectroelectrochemical analyses indicate that both MeCN and bpy spectator ligands provide borderline Class II/III systems, valence delocalization is more pronounced in complex **6** with peripheral MeCN ligands than in complex **7** containing bpy spectator ligands. Besides affecting the stability of the different oxidation states and the reversibility of the redox processes, spectator ligands thus also influence the degree of communication. This relevance needs to be taken into account when designing a next generation of molecular switches based on the bis(carbene) scaffold.

Our results provide unambiguous evidence that N-heterocyclic carbene-based ligands are suitable linkers for the fabrication of molecular redox switches. They constitute synthetically versatile alternatives to the more popular oligopyridine- and alkyne-derived linkers, thus offering to novel synthons for application in organometallic polymer chemistry, in molecular electronics, and potentially also in computing.

Experimental Section

General comments: Starting materials were commercially available and used as received. Solvents were purified by using a Thermovac alumina/catalyst column system. The synthesis of complexes **8** and **10** was described previously.^[25e] Where indicated, reactions were conducted in a Biotage Initiator Microwave Synthesizer. ¹H and ¹³C(¹H) NMR spectra were recorded with Bruker spectrometers at RT unless stated otherwise.

Chemical shifts (δ) are given in ppm relative to signals of residual protio solvent (coupling constants J in Hz). Signals were assigned with the aid of two-dimensional cross-coupling experiments. Elemental analyses were performed by the Microanalytical Laboratory of the ETH Zürich (Switzerland).

Electrochemistry: Electrochemical studies were carried out by using an EG&G Princeton Applied Research Potentiostat, Model 273A, employing a gas-tight three electrode cell under an argon atmosphere. A glassy carbon disk with 3.41 mm² surface area was used as the working electrode and was polished before each measurement. The reference electrode was Ag/AgCl and the counter electrode was a Pt wire. In all experiments, Bu₄NPF₆ (0.1 M in dry solvent) was used as the supporting electrolyte with analyte concentrations of approximately 1 mM. Measurements were performed at different scan rates (50–250 mV s⁻¹). All redox potentials were referenced to the ferrocenium/ferrocene couple (Fc⁺/Fc) as internal standard with $E_{1/2}$ (Fc⁺/Fc) versus SCE = +0.46 V in CH₂Cl₂, +0.40 V in MeCN, +0.35 V in MeNO₂.^[39]

Spectroelectrochemical studies were performed by using an EG&G Princeton Applied Research Potentiostat, Model 273A and a Perkin-Elmer UV/Vis-NIR Spectrometer, Model Lambda 900, employing an optically transparent thin-layer electrochemical (OTTLE) cell.^[30] In all experiments, Bu₄NPF₆ (0.1 M in MeCN or MeNO₂) was used as supporting electrolyte with analyte concentrations of approximately 1 mM. For Hush analysis,^[5] data were first treated to remove solvent artifacts by subtracting the absorbance at ground potential, assuming that there is no relevant absorbance in the NIR region but background and solvent. The baseline was corrected and absorbance values were smoothed over 9 data points. The IVCT band of **6**⁺ (ν_{\max} = 6277 cm⁻¹) contains a shoulder towards high energy. Assuming that the shoulder arises from a minor decomposition product, it was removed by curve fitting. Transformation from absorbance to molar extinction ϵ was calculated via Beer-Lambert law (1 mM concentrations, path length d = 0.2 mm for the OTTLE cell). Integrals were calculated numerically and yielded induced dipole moment μ_{ge} = 7.31 D for **6**⁺ and 3.16 D for **7**⁺ according to Equation (1):

$$\mu_{\text{ge}} = 0.09584 \left(\int \epsilon(\nu) \nu d\nu / \nu_{\max} \right)^{1/2} \quad (1)$$

The electronic coupling was calculated according to Equation (2):

$$H_{\text{ab}} = \mu_{\text{ge}} \times \nu_{\max} / (e \times R) \quad (2)$$

with R taken from the Ru...Ru distance in **6** as determined by X-ray crystallography (10.537 Å) and is the elementary charge. The delocalization parameter Γ was calculated according to Equation (3):

$$\Gamma = 1 - (\nu_{1/2(\text{obs})} / \nu_{1/2(\text{calcd})}) \quad (3)$$

with $\nu_{1/2(\text{calcd})} = (2310 \times \nu_{\max})^{1/2}$.

Synthesis of 1: 1,5-Dichloro-2,4-dinitrobenzene (2.76 g, 11.6 mmol) and 2-aminopyridine (24.0 g, 255 mmol) were stirred neat at 90 °C for 3 days. The mixture was poured into MeCN (50 mL) and water was added until a yellow precipitate formed. This precipitate was isolated by filtration and dried in vacuo to yield **1** as an analytically pure yellow solid (3.59 g, 87%). ¹H NMR (360 MHz CDCl₃): δ = 10.64 (brs, 2H; NH), 10.26 (s, 1H; H_{Ar}), 9.38 (s, 1H; H_{Ar}), 8.44 (d, ³J_{HH} = 4.5 Hz, 2H; H_{py}), 7.75 (t, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 7.17 (d, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 7.09 ppm (dd, ³J_{HH} = 4.5, ³J_{HH} = 8.0 Hz, 2H; H_{py}); ¹³C{¹H} NMR (75 MHz, CDCl₃): δ = 152.5 (C_{py}-C), 148.1 (C_{py}-H), 143.4 (C_{Ar}-NO₂), 138.2 (C_{py}-H), 127.9 (C_{Ar}-H), 126.6 (C_{Ar}-NH), 119.3 (C_{py}-H), 114.9 (C_{py}-H), 103.4 ppm (C_{Ar}-H); elemental analysis calcd (%) for C₁₆H₁₂N₆O₄ (352.31): C 54.55, H 3.43, N 23.84; found C 54.71, H 3.52, N 23.88.

Synthesis of 2: Compound **1** (800 mg, 2.3 mmol) was dissolved in a mixture of HC(OEt)₃ (26 mL) and HCOOH (85%, 2.2 mL, 58 mmol) under argon atmosphere. Sodium formate (3.56 g, 52 mmol) and Pd/C 10% (0.6 mg, 0.52 mmol) were added and the reaction was stirred at 100 °C for 36 h. After cooling, the reaction mixture was filtered and the filtrate treated with aqueous HCl (6%) and stirred for 10 min. The yellow solu-

tion was treated with aqueous NaOH (10%) until basic and allowed to stand for 1 h. The resulting solid was collected by filtration, washed with H₂O, and dried to yield crude **2** (567 mg, 81%). Recrystallization from CH₂Cl₂/pentane mixture afforded an analytically pure sample. ¹H NMR (300 MHz CDCl₃): δ = 8.92 (s, 1H; H_{Ar}), 8.66 (dd, ³J_{HH} = 4.8, ⁴J_{HH} = 1.5 Hz, 2H; H_{py}), 8.62 (s, 2H; NCHN), 8.32 (s, 1H; H_{Ar}), 7.95 (td, ³J_{HH} = 8.1, ⁴J_{HH} = 1.5 Hz, 2H; H_{py}), 7.65 (d, ³J_{HH} = 8.1 Hz, 2H; H_{py}), 7.33 ppm (dd, ³J_{HH} = 8.1, ³J_{HH} = 4.9 Hz, 2H; H_{py}); ¹³C{¹H} NMR (75 MHz, CDCl₃): δ = 150.3 (C_{py}-C), 149.5 (C_{py}-H), 142.2 (NCHN), 142.2 (C_{Ar}-N), 139.0 (C_{py}-H), 130.6 (C_{Ar}-N), 121.6 (C_{py}-H), 114.2 (C_{py}-H), 111.0 (C_{Ar}-H), 96.2 ppm (C_{Ar}-H); elemental analysis calcd for C₁₈H₁₂N₆ (312.33) × 1/8 CH₂Cl₂: C 67.41, H 3.82, N 26.02; found: C 67.78, H 4.10, N 25.61.

Synthesis of 3: Compound **2** (567 mg, 1.8 mmol) was suspended in a mixture of MeI (2.0 mL, 32 mmol) and MeCN (0.1 mL). The reaction mixture was sealed in a pressure tube and heated to 110 °C for 48 h. After cooling, the reaction mixture was filtered and the residue was washed with a minimum amount of MeCN and Et₂O and dried in vacuo, yielding **3** as an analytically pure and slightly hygroscopic off-white powder (569 mg, 52%). ¹H NMR (400 MHz, [D₆]DMSO): δ = 10.84 (s, 2H; NCHN), 9.65 (s, 1H; H_{Ar}), 9.13 (s, 1H; H_{Ar}), 8.85 (d, ³J_{HH} = 4.5 Hz, 2H; H_{py}), 8.36 (t, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 7.65 (d, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 7.33 (dd, 2H, ³J_{HH} = 8.0, ³J_{HH} = 4.5 Hz, 2H; H_{py}), 4.33 ppm (s, 6H; N-CH₃); ¹³C{¹H} NMR (100 MHz, [D₆]DMSO): δ = 149.4 (C_{py}-C), 147.5 (C_{py}-H), 146.6 (NCHN), 140.8 (C_{py}-H), 131.4 (C_{Ar}-N), 128.5 (C_{Ar}-N), 125.2 (C_{py}-H), 116.8 (C_{py}-H), 103.6 (C_{Ar}-H), 99.7 (C_{Ar}-H), 34.5 ppm (N-CH₃); elemental analysis calcd for C₂₀H₁₈N₆ (596.21) × 1/2 H₂O: C 39.69, H 3.16, N 13.89; found: C 39.57, H 3.26, N 13.82.

Synthesis of 4: Compound **3** (569 mg, 1.0 mmol) was dissolved in H₂O (80 mL) and a solution of (NH₄)PF₆ (3.23 g, 20 mmol) in H₂O (10 mL) was added under vigorous stirring. The formed precipitate was collected by filtration and dried in vacuo to yield **4** as an analytically pure white powder (502 mg, 87%). ¹H NMR (360 MHz, [D₆]DMSO): δ = 10.80 (s, 2H; NCHN), 9.65 (s, 1H; H_{Ar}), 9.08 (s, 1H; H_{Ar}), 8.85 (d, ³J_{HH} = 4.1 Hz, 2H; H_{py}), 8.35 (t, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 8.10 (d, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 7.77 (dd, ³J_{HH} = 8.0, ³J_{HH} = 4.1 Hz, 2H; H_{py}), 4.32 ppm (s, 6H; N-CH₃); ¹³C{¹H} NMR (90 MHz, [D₆]DMSO): δ = 149.7 (C_{py}-C), 147.6 (C_{py}-H), 146.7 (NCHN), 140.9 (C_{py}-H), 131.5 (C_{Ar}-N), 128.6 (C_{Ar}-N), 125.4 (C_{py}-H), 116.8 (C_{py}-H), 103.5 (C_{Ar}-H), 99.7 (C_{Ar}-H), 34.5 ppm (N-CH₃); elemental analysis calcd (%) for C₂₀H₁₈F₁₂N₆P₂ (632.33): C 37.99, H 2.87, N 13.29; found: C 38.12, H 3.06, N 13.24.

Synthesis of complex 5: Ag₂O (81 mg, 0.35 mmol) was added to a suspension of benzobisimidazolium salt **4** (100 mg, 0.16 mmol) in MeOH (10 mL). The reaction mixture was stirred in the dark under argon at RT for 24 h. All volatiles were evaporated and the residue was suspended in dry CH₂Cl₂ (10 mL). After addition of [RuCl₂(*p*-cymene)]₂ (106.4 mg, 0.17 mmol), the suspension was stirred at RT for 72 h protected from light. The mixture was then centrifuged and the supernatant was decanted and filtered through a short pad of Al₂O₃. After evaporation and drying in vacuo, complex **5** was obtained as an orange solid in a 1:1 mixture of diastereomers (167 mg, 90%). ¹H NMR (400 MHz CDCl₃), *isomer A*: δ = 9.27 (d, ³J_{HH} = 5.2 Hz, 2H; H_{py}), 7.90–7.96 (m, 4H; H_{py}), 7.84 (s, 1H; H_{Ar}), 7.69 (s, 1H; H_{Ar}), 7.49–7.56 (m, 2H; H_{py}), 6.37 (d, ³J_{HH} = 6.3 Hz, 2H; H_{cym}), 6.23 (d, ³J_{HH} = 6.3 Hz, 2H; H_{cym}), 6.14 (d, ³J_{HH} = 6.5 Hz, 2H; H_{cym}), 5.78 (d, ³J_{HH} = 6.5 Hz, 2H; H_{cym}), 3.98 (s, 6H; N-CH₃), 2.38 (septet, ³J_{HH} = 6.6 Hz, 2H; CHMe₂), 2.28 (s, 6H; cym-CH₃), 0.92 (d, ³J_{HH} = 6.6 Hz, 6H; CH-CH₃), 0.83 (d, ³J_{HH} = 6.6 Hz, 6H; CH-CH₃); *isomer B*: δ = 9.30 (d, ³J_{HH} = 5.6 Hz, 2H; H_{py}), 8.56 (s, 1H; H_{Ar}), 8.45 (d, ³J_{HH} = 8.3 Hz, 2H; H_{py}), 8.33 (t, ³J_{HH} = 8.0 Hz, 2H; H_{py}), 8.20 (s, 1H; H_{Ar}), 7.56 (t, ³J_{HH} = 6.3 Hz, 2H; H_{py}), 6.31 (d, ³J_{HH} = 6.3 Hz, 2H; H_{cym}), 6.28 (d, ³J_{HH} = 6.3 Hz, 2H; H_{cym}), 6.05 (d, ³J_{HH} = 6.5 Hz, 2H; H_{cym}), 5.72 (d, ³J_{HH} = 6.5 Hz, 2H; H_{cym}), 4.43 (s, 6H; N-CH₃), 2.46 (septet, ³J_{HH} = 6.6 Hz, 2H; CHMe₂), 2.22 (s, 6H; cym-CH₃), 0.92 (d, ³J_{HH} = 6.6 Hz, 6H; CH-CH₃), 0.83 ppm (d, ³J_{HH} = 6.6 Hz, 6H; CH-CH₃); ¹³C{¹H} NMR (100 MHz CDCl₃), *isomer A*: 202.8 (C-Ru), 154.4 (C_{py}-C), 153.0 (C_{py}-H), 143.4 (C_{py}-H), 134.5 (C_{Ar}-N), 127.8 (C_{Ar}-N), 125.0 (C_{py}-H), 115.1 (C_{py}-H), 111.7 (C_{cym}-C), 108.4 (C_{cym}-C), 96.7 (C_{Ar}-H), 95.9 (C_{Ar}-H), 94.3 (C_{cym}-H), 92.9 (C_{cym}-H), 89.5 (C_{cym}-H), 85.2 (C_{cym}-H), 37.5 (N-CH₃), 32.1 (CHMe₂), 22.7 (CH-CH₃), 22.5 (C-CH₃), 19.6 (cym-CH₃);

isomer **B**: $\delta = 203.3$ (C–Ru), 157.4 (C_{py} –C), 153.0 (C_{py} –H), 142.9 (C_{py} –H), 135.7 (C_{Ar} –N), 128.8 (C_{Ar} –N), 124.3 (C_{py} –H), 114.9 (C_{py} –H), 111.2 (C_{cym} –C), 108.4 (C_{cym} –C), 97.8 (C_{Ar} –H), 95.9 (C_{Ar} –H), 94.3 (C_{cym} –H), 94.0 (C_{cym} –H), 89.3 (C_{cym} –H), 85.4 (C_{cym} –H), 37.4 (N–CH₃), 32.1 (CHMe₂), 22.6 (C–CH₃), 22.5 (C–CH₃), 19.4 ppm (cym–CH₃); elemental analysis calcd (%) for C₄₀H₄₂Cl₂F₁₂N₆P₂Ru₂ (1171.79) × CH₂Cl₂: C 39.18, H 3.69, N 6.69; found C 38.98, H 3.90, N 6.97.

Synthesis of complex 6: Complex **5** (140 mg, 0.12 mmol) was dissolved in MeCN (10 mL) and AgPF₆ (121 mg, 0.48 mmol) was added. The mixture was heated to reflux and stirred for 20 h. After cooling, the reaction mixture was poured into H₂O (10 mL). The formed precipitate was filtered, washed with Et₂O (3 × 5 mL) and dried to yield analytically pure complex **6** as an off-white powder (91 g, 52%). X-ray quality crystals were grown by slow diffusion of Et₂O into a MeCN solution of **6** at –20 °C. ¹H NMR (400 MHz, CDCl₃): $\delta = 9.05$ (d, ³J_{HH} = 5.7 Hz, 2H; H_{py}), 8.58 (s, 1H; H_{Ar}), 8.48 (d, ³J_{HH} = 8.2 Hz, 2H; H_{py}), 8.33 (t, ³J_{HH} = 8.2 Hz, 2H; H_{py}), 8.11 (s, 1H; H_{Ar}), 7.56 (dd, ³J_{HH} = 8.2, ³J_{HH} = 5.7 Hz, 2H; H_{py}), 4.41 (s, 6H; N–CH₃), 2.60 (s, 6H; NCCH₃), 2.09 (s, 12H; NCCH₃), 1.96 ppm (s, 6H; NCCH₃); ¹³C{¹H} NMR (100 MHz, CDCl₃): $\delta = 207.6$ (C–Ru), 156.4 (C_{py} –C), 154.7 (C_{py} –H), 142.2 (C_{py} –H), 136.0 (C_{Ar} –N), 129.9 (C_{Ar} –N), 127.9 (NCMe), 126.2 (NCMe), 125.4 (NCMe), 123.4 (C_{py} –H), 114.3 (C_{py} –H), 95.9 (C_{Ar} –H), 94.8 (C_{Ar} –H), 35.9 (N–CH₃), 4.9 (NC–CH₃), 4.4 (NC–CH₃), 1.1 ppm (NC–CH₃); elemental analysis calcd (%) for C₃₆H₄₀F₂₄N₁₄P₄Ru₂ (1452.02): C 29.80, H 2.78, N 13.52; found C 29.67, H 2.89, N 13.30.

Synthesis of complex 7: A pressure tube was charged with complex **6** (60 mg, 41 mmol), 2,2-bipyridine (60 mg, 0.38 mmol), and MeNO₂ (4 mL) and stirred at 110 °C for 20 h. After cooling, the orange reaction mixture was poured into Et₂O (20 mL) and the formed precipitate was filtered off, washed with Et₂O, and dried to give complex **7** as an orange solid (86 mg, 95%). Recrystallization from CH₂Cl₂/pentane gave an analytically pure sample and crystals suitable for an X-ray diffraction analysis. NMR spectroscopy revealed a 1:1 mixture of two diastereomers. ¹H NMR (500 MHz, CDCl₃): $\delta = 8.92$ (s, 2H; H_{Ar}), 8.64 (d, 4H; H_{bpy}), 8.63 (d, 4H; H_{bpy}), 8.62 (d, 2H; H_{py}), 8.61 (d, 2H; H_{bpy}), 8.59 (d, 2H; H_{py}), 8.56 (d, 2H; H_{bpy}), 8.55 (d, 2H; H_{bpy}), 8.53 (d, 2H; H_{bpy}), 8.26 (t, 4H; H_{bpy}), 8.24 (d, 2H; H_{py}), 8.21 (t, 4H; H_{bpy}), 8.20 (d, 2H; H_{py}), 8.15 (t, 2H; H_{bpy}), 8.12 (t, 2H; H_{bpy}), 8.10 (t, 2H; H_{bpy}), 8.09 (t, 2H; H_{bpy}), 8.09 (t, 2H; H_{py}), 8.06 (t, 2H; H_{py}), 8.00 (d, 2H; H_{bpy}), 7.98 (d, 2H; H_{bpy}), 7.86 (t, 4H; H_{bpy}), 7.85 (d, 2H; H_{bpy}), 7.80 (t, 4H; H_{bpy}), 7.79 (d, 2H; H_{bpy}), 7.69 (s, 2H; H_{Ar}), 7.58 (dd, 4H; H_{bpy}), 7.45 (dd, 2H; H_{bpy}), 7.44 (dd, 2H; H_{bpy}), 7.42 (dd, 2H; H_{bpy}), 7.41 (dd, 2H; H_{bpy}), 7.40 (dd, 2H; H_{py}), 7.31 (dd, 4H; H_{bpy}), 7.30 (dd, 2H; H_{py}), 3.50 ppm (s, 12H; N–CH₃); ¹³C{¹H} NMR (125 MHz, CDCl₃): $\delta = 211.1$ (C–Ru), 158.3–156.2 (C_{py}^2), 156.7–150.3 (C_{py}^3), 141.5–139.2 (C_{py}^4), 136.7 (C_{Ar} –N), 130.3 (C_{Ar} –N), 130.3–124.1 (C_{py}^5), 125.9–114.4 (C_{py}^6), 96.2 (C_{Ar} –H), 93.7 (C_{Ar} –H), 34.0 ppm (N–CH₃); elemental analysis calcd (%) for C₆₀H₄₈F₂₄N₁₄P₄Ru₂ (1747.11): C 41.25, H 2.77, N 11.22; found C 40.96, H 3.21, N 11.01.

Synthesis of complex 9: Complex **8** (50 mg, 80 mmol) was dissolved in MeCN (8 mL) and AgPF₆ (45 mg, 0.18 mmol) was added. The reaction mixture was stirred at reflux temperature for 20 h. After cooling, all volatiles were evaporated and the residue was dissolved in a minimum amount of H₂O and extracted with CH₂Cl₂ (3 × 5 mL). The combined organic layers were washed with H₂O (5 mL) and dried over Mg₂SO₄. The solution was poured into pentane (30 mL) and the formed precipitate was filtered and washed with pentane. After drying in vacuo complex **9** was obtained as an off-white solid (45 mg, 33%). X-ray quality crystals were grown by slow diffusion of Et₂O into a MeCN solution of complex **9** at –20 °C. Elemental analysis calcd (%) for C₂₁H₂₃F₁₂N₇P₂Ru (764.45): C 32.99, H 3.03, N 12.83; found: C 33.23, H 2.91, N 13.04; ¹H NMR (300 MHz, CDCl₃): $\delta = 8.97$ (dd, ³J_{HH} = 5.8, ⁴J_{HH} = 0.9 Hz, 1H; H_{py}), 8.27–8.13 (m, 3H; H_{Ar} + 2H_{py}), 7.81–7.73 (m, 1H; H_{Ar}), 7.64–7.51 (m, 2H; H_{Ar}), 7.47 (ddd, ³J_{HH} = 7.2, ³J_{HH} = 5.8, ³J_{HH} = 1.3 Hz, 1H; H_{py}), 4.26 (s, 3H; N–CH₃), 2.56 (s, 3H; NC–CH₃), 2.07 (s, 6H; NC–CH₃), 1.96 ppm (s, 3H; NC–CH₃); ¹³C{¹H} NMR (100 MHz, CDCl₃): $\delta = 204.0$ (C–Ru), 156.9 (C_{py} –C), 154.4 (C_{py} –H), 141.9 (C_{py} –H), 138.2 (C_{Ar} –N), 132.7 (C_{Ar} –N), 127.7 (NCMe), 126.1 (C_{Ar} –H), 123.0 (C_{Ar} –H), 125.1 (NCMe), 123.0 (C_{py} –H), 118.7 (NCMe), 113.7 (C_{py} –H), 112.9 (C_{Ar} –H), 112.3 (C_{Ar} –H), 35.3

(N–CH₃), 4.8 (NC–CH₃), 4.4 (NC–CH₃), 1.4 ppm (NC–CH₃); elemental analysis calcd (%) for C₂₁H₂₃F₁₂N₇P₂Ru (764.45): C 32.99, H 3.03, N 12.83; found: C 33.23, H 2.91, N 13.04.

Crystal structure determinations: Suitable single crystals were mounted on a StoeMark II-Imaging Plate Diffractometer System (Stoe & Cie, 2002) equipped with a graphite-monochromator. Data collection was performed by using MoK α radiation ($\lambda = 0.71073$ Å) with a nominal crystal to detector distance of 135 mm. All structures were solved by direct methods by using SHELXS-97 and refined by full-matrix least-squares on F^2 for all data using SHELXL-97.^[40] The hydrogen atoms were included in calculated positions and treated as riding atoms using SHELXL-97 default parameters. Anisotropic thermal displacement parameters were used for all non-hydrogen atoms. Four independent molecules and eight disordered PF₆[–] anions are present in the unit-cell of **9**. Attempts to resolve the disorder of the PF₆[–] ions and solvent molecules by using the SQUEEZE routine^[41] were unsuccessful due to the low quality of the data. Further crystallographic details are compiled in the Supporting Information. CCDC-numbers 949564 (**2**), 949565 (**6**), and 949566 (**10**) contain the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.

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