Additions and intramolecular migrations of nucleophiles in cationic diruthenium μ-allenyl complexes

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Summary: the diruthenium μ-allenyl complex [Ru\textsubscript{2}(CO)(NCMe)(μ-CO){μ-η\textsuperscript{1}:η\textsuperscript{2}-C(H)=C=C(Me)(Ph)}(Cp)\textsubscript{2}][BF\textsubscript{4}], \textsuperscript{3b}, reacts with halide anions to yield the neutral derivatives [Ru\textsubscript{2}(CO)\textsubscript{2}(X){μ-η\textsuperscript{1}:η\textsuperscript{2}-C(H)=C=C(Me)(Ph)}(Cp)\textsubscript{2}] [X = Cl, \textsuperscript{4b}; X = Br, \textsuperscript{4c}; X = I, \textsuperscript{4d}]. Complex \textsuperscript{4b} undergoes isomerization to the unprecedented bridging vinyl-chlorocarbene species [Ru\textsubscript{2}(CO)(μ-CO){μ-η\textsuperscript{1}:η\textsuperscript{3}-C(Cl)C(H)=C=C(Me)(Ph)}(Cp)\textsubscript{2}], \textsuperscript{10}, upon filtration of a CH\textsubscript{2}Cl\textsubscript{2} solution through an alumina column.

Complex \textsuperscript{3b} reacts with an excess of NaBH\textsubscript{4} to give five products: the allene complex [Ru\textsubscript{2}(CO)\textsubscript{2}{μ-η\textsuperscript{2}:η\textsuperscript{2}-CH\textsubscript{2}=C=C(Me)(Ph)}(Cp)\textsubscript{2}], \textsuperscript{5}; the hydride species \textit{trans}-[Ru\textsubscript{2}(CO)\textsubscript{2}(μ-H){μ-η\textsuperscript{1}:η\textsuperscript{2}-CH=CH=C=C(Me)(Ph)}(Cp)\textsubscript{2}], \textsuperscript{6}, and \textit{cis}-[Ru\textsubscript{2}(CO)\textsubscript{2}(μ-H){μ-η\textsuperscript{1}:η\textsuperscript{2}-CH=CH=C=C(Me)(Ph)}(Cp)\textsubscript{2}], \textsuperscript{8}; the vinylalkylidene [Ru\textsubscript{2}(CO)(μ-CO){μ-η\textsuperscript{1}:η\textsuperscript{3}-C(H)C(H)=C=C(Me)(Ph)}(Cp)\textsubscript{2}], \textsuperscript{9}; and the cluster [Ru\textsubscript{3}(CO)\textsubscript{3}(μ-H)\textsubscript{3}(Cp)\textsubscript{3}], \textsuperscript{7}.

Studies on the thermal stabilities of \textsuperscript{5,6,8} and \textsuperscript{9} have suggested a plausible mechanism for the formation of these complexes and for the synthesis of \textsuperscript{10}.

Keywords: Diruthenium complexes; Allenyl; Bridging vinyl-chlorocarbene; Bridging hydride; Hydride migration

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Introduction

Allenyl ligands [-C\(\alpha\)(R)=C\(\beta\)=C\(\gamma\)(R')(R'')] coordinated in mono- [1], di- [1a, 2] and poly-nuclear complexes [1a, 3] have been the subject of much interest in recent years. In particular, di- and poly-nuclear species have attracted considerable attention: the unsaturated C\(_3\) chain readily undergoes coupling reactions with organic fragments [4], providing an easy route for the synthesis of new multisite-bound hydrocarbyl ligands [1a, 2b, 4, 5].

Allenyls display a variety of coordination modes. The most common in dinuclear complexes is the [\(\mu-\eta^1:\eta^2\)] fashion, with coordination through the C\(\alpha\) and C\(\beta\) carbons [1a, 6]; however, alternative modes of coordination have been reported [1a, 2d, 7].

Common methods of synthesis include nucleophilic substitution on propargyl halides [8] and C-C bond formation by reaction with diazocompounds [9]. A further strategy for the preparation of [\(\mu-\eta^1:\eta^2\)] allenyl complexes was recently reported [10], whereby the diruthenium complexes [Ru\(_2\)(CO)(\(\mu\)-CO){\(\mu-\eta^1:\eta^3\)}-C\(\alpha\)(H)C\(\beta\)C\(\gamma\)(OH)(Me)(R)C(O)](Cp)\(_2\) [R = Me, 1a; R = Ph, 1b], prepared from the cyclopentenone adduct [Ru\(_2\)(CO)(\(\mu\)-CO){\(\mu-\eta^1:\eta^3\)}-C(Ph)C(Ph)C(O)](Cp)\(_2\)] by alkyne exchange [11], are treated with HBF\(_4\), resulting in formation of the allenyl [Ru\(_2\)(CO)(\(\mu\)-CO){\(\mu-\eta^1:\eta^2\)}-C\(\alpha\)(H)=C\(\beta\)=C\(\gamma\)(Me)(R)](Cp)\(_2\)][BF\(_4\)] [R = Me, 2a; R = Ph, 2b] (scheme 1) [12].

Preliminary studies on the chemistry of 2 have shown that the allenyl ligand is reactive towards neutral organic molecules such as alkynes and diazocompounds [12]. Moreover, compounds 2a-b react readily with CH\(_3\)CN/Me\(_3\)NO to give the acetonitrile adducts [Ru\(_2\)(CO)(NCMe)(\(\mu\)-CO){\(\mu-\eta^1:\eta^2\)}-C(H)=C=C(Me)(R)](Cp)\(_2\)][BF\(_4\)] [R = Me, 3a; R = Ph, 3b], see scheme 1.

The chemistry of the derivatives 3 offers promise in that acetonitrile often acts as a labile ligand in organometallic chemistry. For dinuclear species acetonitrile removal is a crucial step in allowing coordination, at one of the two metal centres, of either anionic nucleophiles such as hydride, halides and pseudo-halides [13], or neutral organic fragments [14]. In a number of cases this substitution is the precursor of an intramolecular coupling reaction involving the species introduced and the pre-existing bridging hydrocarbyl ligand; these reactions, particularly those leading to carbon-carbon bond formation, have attracted interest because they may act as models for heterogeneously catalyzed processes occurring on metal surfaces [14a,c].
It has been previously demonstrated that complex 3a undergoes nucleophilic substitution under mild conditions by chloride ion, towards which 2a-b are completely unreactive, affording the neutral allenyl derivative \([\text{Ru}_2(\text{CO})_2(\text{Cl})\{\mu-\eta^1:\eta^2-\text{C(H)=C=CMe}_2\}](\text{Cp})_2]\), 4a (scheme 1), the structure of which has been determined by X-ray diffraction [12]. We report herein our studies on the reactivity of 3b towards hydride and halide ions, aimed at exploring the possibility of coupling reactions between the allenyl ligand and species introduced to the metal centre by nitrile removal.

**Scheme 1**

![Scheme 1](image-url)
Results and discussion

1. Reactivity of acetonitrile derivatives of diruthenium allenyl complexes with NaBH₄

The acetonitrile compound $[\text{Ru}_2(\text{CO})(\text{NCMe})(\mu-\text{CO})\{\mu-\eta^1:\eta^2-\text{C(H)}=\text{C}(\text{Me})(\text{Ph})\}(\text{Cp})_2][\text{BF}_4]$, 3b, is obtained rapidly by treating a dichloromethane solution of 2b with one equivalent of Me₃NO dissolved in acetonitrile (scheme 2).

![Scheme 2](image)

Complex 3b was characterized by IR spectroscopy and used in situ for subsequent reactions with nucleophiles. The IR spectrum of 3b (in CH₂Cl₂ solution) shows two bands at 2005 and 1870 cm⁻¹, corresponding to the presence of one terminal and one semi-bridging carbonyl ligands respectively. Complex 3b probably exists as mixtures of interconverting isomers, due to $\sigma$, $\pi$ and cis/trans fluxionality [10, 12], allowing the NCMe ligand to be bound at either ruthenium atom.

The reaction of 3b, in THF solution, with NaBH₄ was investigated. IR spectroscopy revealed, after one hour, the complete disappearance of the starting material, and subsequent removal of the solvent gave a yellow-brown residue. The IR spectrum of this (in CH₂Cl₂ solution) exhibited two absorptions at 1981 cm⁻¹ (s) and 1919 cm⁻¹ (vs). The residue was chromatographed on an alumina column maintained at –35 °C, leading to the separation of five products (see scheme 3): $[\text{Ru}_2(\text{CO})_2\{\mu-\eta^2:\eta^2-\text{CH}_2=\text{C}(\text{Me})(\text{Ph})\}(\text{Cp})_2]$, 5; trans-$[\text{Ru}_2(\text{CO})_2(\mu-\text{H})\{\mu-\eta^1:\eta^2-\text{CH}=\text{C}(\text{Me})(\text{Ph})\}(\text{Cp})_2]$, 6; $[\text{Ru}_2(\text{CO})_3(\mu-\text{H})(\text{Cp})_3]$, 7; cis-$[\text{Ru}_2(\text{CO})_2(\mu-\text{H})\{\mu-\eta^1:\eta^2-\text{CH}=\text{C}(\text{Me})(\text{Ph})\}(\text{Cp})_2]$, 8; and $[\text{Ru}_2(\text{CO})(\mu-\text{CO})\{\mu-\eta^1:\eta^3-\text{C(H)}\text{C(H)}=\text{C}(\text{Me})(\text{Ph})\}(\text{Cp})_2]$, 9. Complex 7, the formation of which appears to be the
consequence of some fragmentation (yield 7 %), is a known species [15], and was identified by IR spectroscopy and mass spectrometry, in addition to its characteristic deep blue colour. The new compounds 5, 6, 8 and 9 have been fully characterized by IR and NMR spectroscopy, elemental analysis and mass spectrometry.

Scheme 3

Complex 5 is the formal result of nitrile displacement and hydride attack on the $C_\alpha$ carbon of the allenyl unit of 3b. The complex consists of a [Ru$_2$(Cp)$_2$(CO)$_2$] core, with the two carbonyls occupying terminal sites at each metallic centre, and an allene ligand which is
coordinated to the ruthenium atoms through its \( \pi \) bonds. Dinuclear compounds containing the \( \mu-\eta^2:\eta^2 \) allene moiety have been described [16], some containing the \([M_2(Cp)_2(CO)_2]\) skeleton (\( M = Ru [17], Fe [18] \)). Furthermore, the same \( \eta^2:\eta^2 \) allene ligand has also been reported to be coordinated in polynuclear species [19].

The spectroscopic data for 5 are characteristic of its diiron and diruthenium analogues. The IR spectrum (in \( CH_2Cl_2 \) solution) exhibits two carbonyl bands, at 1933 and 1915 cm\(^{-1} \). The \(^1\)H NMR spectrum shows the presence in solution of two isomers in comparable amounts. In analogous compounds the only configuration exhibited by the Cp ligands is trans, and it is likely that this is also the case here. For example, the complex \([Ru_2(CO)_2{\mu-\eta^2:\eta^2-\text{C}_a(H)(\text{CO}_2\text{Et})=\text{C}_\beta=\text{C}_\gamma(H)(\text{CO}_2\text{Me})]}(\text{Cp})_2] \) has been shown to exist in two trans forms, both characterized by X ray diffraction [17]. The two forms differed in the orientations of the substituents at \( C_\gamma \), with one isomer having the \( \text{CO}_2\text{Me} \) group and the \( C_\alpha-H \) in axial and equatorial sites, respectively, with regard to the Ru-Ru bond axis, while in the other the \( \text{CO}_2\text{Me} \) and the \( C_\alpha-H \) lie in equatorial and axial positions, respectively. In both isomers, the \( \text{CO}_2\text{Et} \) function is in an axial position. On this basis, it is likely that the two isomers of 5 have the structures 5a and 5b, in which the orientations of the \( C_\gamma \) substituents (Ph and Me) are different. The \(^1\)H NMR data suggest that in 5a Ph is axial and Me is equatorial, whereas in 5b the opposite configuration is adopted. Indeed, in the \(^1\)H NMR spectrum of 5a the methyl group (\( \delta 1.82 \) ppm) is shielded compared with that in 5b (\( \delta 2.17 \) ppm), while the opposite trend is seen for the Cp resonances [5a: \( \delta 5.04, 4.87 \) ppm; 5b \( \delta 4.65, 4.35 \) ppm]. These features indicate that the Me group is close to the Ru-Ru axis in 5a while in 5b it is oriented away from it (scheme 4).

**Scheme 4**
The Cα-H protons of 5 resonate as doublets at ca. 3.6 and 2.7 ppm, with a coupling constant of ca. 4 Hz. In the 13C NMR spectrum, the Cα resonance is shifted significantly upfield (e.g. for 5b at 29.7 ppm) with respect to complex 2b [12], as a consequence of the loss of alkylidene character. The Cβ and Cγ resonances occur at ca. 190 and 60 ppm.

Complexes 6 and 8, the latter being the highest yield product (36 %), derive from 3b by nitrile displacement with hydride ion, followed by rearrangement without involving the allenyl ligand.

The IR spectrum of 6 exhibits two carbonyl bands, at 1972 and 1918 cm−1, with the second more intense than the first. Conversely, the IR spectrum of 8 shows bands at 1984 and 1921 cm−1, with the latter being less intense than the former.

The NMR spectra of 6 and 8 reveal the absence of fluxionality and contain single sets of resonances. In the 1H NMR spectra, the bridging hydrides are at typically high field chemical shifts, at −13.73 ppm (6) and −14.20 ppm (8). The salient 1H and 13C NMR features of 6 and 8 do not differ significantly from the data available for the parent 2b. Thus, the Cα-H proton resonates as a quartet (due to coupling with Cγ-Me) at 8.80 ppm (6) and 9.12 ppm (8). Moreover, the Cα, Cβ and Cγ carbons are observed at ca. 115, 165 and 110 ppm respectively. The CO carbons resonate at values typical of terminal carbonyl ligands (e.g. for 6 at 202.4 and 203.1 ppm).

Complexes 6 and 8 are isomers, which could arise from either cis-trans arrangements of the Cp ligands or from axial and equatorial orientations of the Cγ substituents. The former is the most likely on the basis of the IR and NMR data. Cis and trans isomers of complexes containing the [M2(Cp)2(CO)2] unit (M = Fe, Ru) in several cases have distinctive frequencies for their higher carbonyl stretching vibration, the latter being at higher values in cis isomers than in the trans forms [13c-e, 20]. On this basis, a cis configuration is attributable to complex 8 and a trans configuration to complex 6. This attribution is supported by comparing the 1H NMR resonances of the Cp ligands of 6 and 8 with those of the cis and trans isomers of the parent complex 2b [12], and by the order in which compounds 6 and 8 are eluted from the chromatographic column. Thus 6, which is expected to be less polar than 8, is eluted with a less polar solvent. Moreover, the Cγ-Me resonances for 6 and 8, in both the 1H and 13C NMR spectra, are very similar (e.g. in the 1H NMR spectra: for 6, δ 2.25 ppm; for 8, δ 2.24 ppm), while the Cα resonances in the 13C NMR spectra differ significantly (for 6: δ = 113.5 ppm; for
8: \( \delta = 118.1 \) ppm). The opposite trends should be apparent if compounds 6 and 8 have different orientations of the methyl and phenyl groups at \( C_\gamma \) and the same arrangement of the Cp ligands. In 6 and 8 the \( C_\gamma \) substituents are expected to adopt the same orientations found in the precursor 2b, where the hindered phenyl group points away from the Cp rings, lying in an axial position.

The structures shown in scheme 3, with bridging rather than terminal hydride, are as expected for compounds 6,8. Hydrides are usually found acting as bridging ligands in diruthenium complexes [14a, 21], and examples of terminal coordination are rare [22]. IR spectroscopy is strongly diagnostic in determining which coordination is adopted by hydride in a dinuclear complex, since the band due to terminal Ru-H stretching is observable at ca. 1960 cm\(^{-1}\) [22a,c]. No such band is observed in the IR spectra of 6,8.

Compound 9 (and its analogue 11, see later) belongs to a large class of diruthenium and diiron complexes which have been reported in the literature. Such complexes consist of a \( \text{cis-}[\text{M}_2(\text{Cp})_2(\text{CO})(\mu-\text{CO})] \) core (\( M = \text{Ru, Fe} \)) and a bridging \( \mu-\eta^1:\eta^3 \)-vinylalkylidene ligand (scheme 5, structure I), which has been alternatively described as \( \mu-\eta^1:\eta^3 \)-allylidene (scheme 5, structure II) or 1,2-dimetalla-cyclo-pent-3-ene (scheme 5, structure III) [23].

\[ \text{Scheme 5} \]

The spectroscopic characteristics of 9 are consistent with those of the analogous species. The IR spectrum (in CH\(_2\)Cl\(_2\) solution) shows two bands at 1951 and 1774 cm\(^{-1}\), accounting for terminal and bridging carbynyls, respectively. In the \(^1\)H NMR spectrum, which has a single set of resonances, the \( C_\alpha-H \) proton resonates as a doublet (\( J 7.2 \) Hz), due to coupling with \( C_\beta-H \), at the typically low field (\( \delta 10.75 \) ppm) expected for a proton bound to a bridging
alkylidene carbon; Cβ-H is seen at δ 4.71 ppm. In the 13C NMR spectrum, the resonances of the Cα, Cβ and Cγ carbons are found at δ 155.4, 82.7 and 71.1 ppm respectively. These values are consistent with the alkylidene character of Cα and with the olefin-bound-to-metal nature of Cβ and Cγ.

2. Studies on the thermal stabilities of the products of the reaction of 3b with NaBH₄

The IR spectrum of the mixture formed in the reaction of 3b with NaBH₄ (see above) indicates that the mixture contains mainly compounds 6 and 8, and suggests that the other products isolated by chromatography might derive from 6 and 8 through subsequent intramolecular rearrangements.

In order to investigate this possibility, solutions of complexes 5, 6, 8 and 9 were heated and monitored via IR spectroscopy. These studies have suggested a plausible mechanism for the formation of 6, 5 and 9, shown in scheme 6. Moreover, it has been possible to deduce a scale of thermodynamic stability, with the vinyl-alkylidene species 9 as the most stable product.

This result is in accordance with previous findings, which indicate that µ-allenyl complexes undergo nucleophilic additions preferentially at the central carbon (Cβ) of the C₃ chain [24], although examples of additions at Cα have also been reported [5a, 25].

Compound 8 is the main fraction collected by chromatography, in spite of the fact it is not stable in boiling n-heptane solution. This abundance is probably due to the following kinetic reasons: i) complex 8 results directly from 3b by nitrile substitution with hydride; ii) the configuration of the Cp ligands (cis) is that presumably favoured in the parent cationic allenyl complexes [12]. In other words, the synthesis of 8 consists of simple substitution at the metal centres and would not require a rearrangement of the Cp rings.

Complex 8 undergoes clean and complete cis-to-trans isomerisation after 30 minutes in heptane at reflux temperature, to give 6. Cis-trans isomerizations are common in dinuclear complexes of ruthenium and iron. Adams and Cotton proposed a mechanism for the cis-trans isomerization of [Fe₂(μ-CO)₂(CO)₂(Cp)₂] which is now a fundamental example of the dynamic behaviour of metal carbonyl compounds [26]. Moreover, cis-trans interconversion has been investigated in [Ru₂(μ-CO)₂(CO)₂(Cp)₂] [27] and related dinuclear complexes [21e, 28].

Compound 6 transforms cleanly, in boiling heptane over 7 hours, into the µ-η²:η² allene species 5b. It is noteworthy that complex 5b is obtained in the absence of 5a under these
conditions, contrary to what was found after chromatography at low temperature. It may be concluded that 5b is thermodynamically more stable than 5a. Formation of 5a in significant amounts at low temperature is probably because it has the same arrangement of the Cγ substituents as the allenyl precursor (see scheme 4). The rearrangement which leads from 6 to 5b consists of hydride migration from the metal site to the Cα carbon of the allenyl moiety. It is probable that, both on the alumina column and in boiling heptane, compounds 5a,b derive from 6 (and not from 8) since this route does not require any modification of the arrangement of the Cp ligands.

Finally, if an n-heptane solution of 5b is heated for 42 hours, formation of 9 occurs, accompanied by some decomposition, as a result of hydrogen migration from Cα to Cβ, together with trans-to-cis isomerisation. Compound 9 is stable in boiling heptane solution upon further heating (7 hours).

Scheme 6
Table 1
Relevant $^1$H NMR resonances for the compounds involved in the reactions shown in scheme 6.

<table>
<thead>
<tr>
<th>Complex</th>
<th>8</th>
<th>6</th>
<th>5b</th>
<th>9</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\mu$-H</td>
<td>−14.20</td>
<td>−13.73</td>
<td></td>
<td></td>
</tr>
<tr>
<td>$C_\alpha$-H</td>
<td>9.12</td>
<td>8.80</td>
<td>3.72, 2.85</td>
<td>10.75</td>
</tr>
<tr>
<td>$C_\beta$-H</td>
<td></td>
<td></td>
<td></td>
<td>4.71</td>
</tr>
</tbody>
</table>
3. Reactivity of acetonitrile derivatives of diruthenium allenyl complexes with halides

The possibility of introducing simple anions at the metal sites of the diruthenium species 3, and observing subsequent migration to the allenyl ligand, led us to investigate the reactivity of 3b with halides. The reactions of 3b with Cl\(^-\) (from LiCl), Br\(^-\) (from LiBr) and I\(^-\) (from KI) occur rapidly in the presence of a large excess of the salt in THF solution, to afford the neutral products [Ru\(_2\)(CO)\(_2\)(X){µ-\(\eta^1:\eta^2\)-C\(_\alpha\)(H)=C\(_\beta\)=C\(_\gamma\)(Me)(Ph)}(Cp)\(_2\)] [X = Cl, 4b; X = Br, 4c; X = I, 4d] (scheme 7).

Scheme 7

Compounds 4b-d were purified by quick filtration through alumina and fully characterized by IR and NMR spectroscopy, elemental analysis and mass spectrometry. The molecular structures of 4b-d are likely to resemble that determined in the solid state for [Ru\(_2\)(CO)\(_2\)(Cl){µ-\(\eta^1:\eta^2\)-C\(_\alpha\)(H)=C\(_\beta\)=C\(_\gamma\)(Me)}(Cp)\(_2\)], 4a; this latter complex, which has a trans geometry for the Cp rings, contains two carbonyls, one terminally bonded to one ruthenium atom, and one semi-bridging. The X ray structure of 4a revealed that the allenyl ligand bridges the two ruthenium atoms in an identical fashion to that found for 2b [12], with a \(\sigma\) bond to one metal and a \(\pi\) bond to the other. Moreover, the chloride ligand is bound to the ruthenium which is \(\sigma\)-bound to the allenyl.

The mono-halide species 4b-d show very similar spectroscopic patterns, analogous to that of 4a. The IR spectra (in CH\(_2\)Cl\(_2\) solution) exhibit two bands at ca. 1985 and 1890 cm\(^{-1}\), corresponding to one terminal and one semibridging carbonyl, respectively. The \(^1\)H NMR
spectra of the compounds, recorded at room temperature, show the absence of fluxionality. Moreover, the NMR spectra of 4b contain two sets of signals, in \textit{ca.} 4:1 ratio. It is proposed that the two sets are due to \textit{cis} and \textit{trans} isomers, in agreement with what was reported for 4a, although it is difficult to deduce which set of resonances is which. The salient $^1$H and $^{13}$C NMR features of 4b-d do not significantly differ from the data available for 4a, 2b, 6 and 8. Thus, a resonance at \textit{ca.} 10 ppm is seen for the $C_\alpha$-$H$ proton, while the carbons of the allenyl chain resonate, for example in the case of 4d, at $\delta$ 120.7 ($C_\alpha$), 160.4 ($C_\beta$) and 113.7 ($C_\gamma$) ppm. Moreover, the resonances of the carbonyl ligands are found at \textit{ca.} 202 and 213 ppm, in accord with the terminal and semi-bridging coordination modes, respectively, suggested by IR data.

The substitution of one nitrile ligand by halides has been previously described for dinuclear complexes containing the unit $[\text{M}_2\text{Cp}_2(\text{CO})_2]$, $\text{M} = \text{Fe, Ru}$ \cite{13b-e}.

The reaction which leads to 4c is accompanied by fragmentation, since small amounts (12 \%) of the known monomer $[\text{Ru(Cp})(\text{CO})_2(\text{Br})]$ \cite{29} were isolated upon chromatography.

More interestingly, we have observed that compound 4b, in dichloromethane solution, is not stable upon slow filtration through the alumina used for chromatographic separations. Thus, passing a CH$_2$Cl$_2$ solution of 4b through an alumina column (see experimental) results in clean formation of an orange product which has been identified as $[\text{Ru}_2(\text{CO})(\mu-\text{CO})\{\mu-\eta^1:\eta^3-\text{C}_\alpha(\text{Cl})\text{C}_\beta(\text{H})=\text{C}_1(\text{Me})(\text{Ph})\}(\text{Cp})_2]$, 10, an isomer of 4b (scheme 8).

\textbf{Scheme 8}

\begin{center}
\includegraphics[width=0.8\textwidth]{scheme8.png}
\end{center}

Evidence for terminal and symmetrically bridging carbonyls in 10 is provided by IR absorptions at 1985 and 1832 cm$^{-1}$ respectively. The $^1$H NMR spectrum displays resonances for the phenyl protons (7.80-7.05 ppm), the olefinic $C_\beta$-$H$ (5.59 ppm), the Cp rings (5.26, 4.57 ppm), and the $C_\gamma$-$\text{Me}$ group (1.48 ppm), respectively. In particular, the absence of a low-field
signal, attributable to an alkyldiene proton, suggests the modification of the allenyl moiety present in the starting compound 4b. Unfortunately, complex 10 is not sufficiently soluble to record a satisfactory $^{13}$C NMR spectrum, or to grow crystals suitable for X-ray analysis. However, elemental analysis for C, H and Cl, and the mass spectrum confirmed the identity of 10.

According to the sequence shown in scheme 6, the bridging vinyl-chlorocarbene ligand of complex 10 might be the result of chloride migration from the metal to the C$_\alpha$ carbon of the allenyl ligand, followed by hydrogen migration from C$_\alpha$ to C$_\beta$. Nonetheless, attempts to convert 4b into 10 upon heating in solution were unsuccessful: complex 4b is unaltered after 15 hours in boiling toluene. This experimental evidence suggests that the possibility that complex 10 forms by a reaction of 4b with CH$_2$Cl$_2$ on alumina, rather than by intramolecular rearrangements of 4b, should not be excluded. Somewhat surprisingly, the transformation depicted in scheme 8 is limited to compound 4b, since the analogous species 4a,4c-d are stable upon passage through alumina. Terminal halocarbene ligands are well known species [30], however only a few cases of bridging halocarbenes have been reported, and they usually involve the CF$_2$ unit bridging between two metal centres [31]. Compound 10 contains the first example of a bridging vinyl-chlorocarbene ligand.

**Conclusions**

The substitution of a carbonyl ligand by an acetonitrile molecule in cationic diruthenium allenyl complexes allows the introduction of hydride and halide ions to the metal sites under mild conditions.

The resulting products undergo successive rearrangements which involve the bridging allenyl ligand. These can be promoted by: i) thermal treatment, ii) filtration through alumina, and iii) both. Hydride migration from the metal site to the C$_\alpha$ carbon of the allenyl moiety provides a $\mu$-$\eta^2$-$\eta^2$ allene adduct. The latter can be isolated and characterized, but it is further convertible into a more stable vinyl-alkylidene derivative, by hydrogen migration from C$_\alpha$ to C$_\beta$.

Likewise, an unprecedented example of a bridging vinyl-chlorocarbene species might be the result of chloride migration from the ruthenium centre to the C$_\alpha$ carbon of the allenyl ligand, occurring simultaneously with hydrogen migration from C$_\alpha$ to C$_\beta$. 
Experimental Section

General Data. All reactions and general manipulations were carried out under a nitrogen atmosphere, using solvents dried by distillation from appropriate drying agents. Chromatographic separations were carried out on columns of deactivated alumina (6% w/w water), at –35 °C. All reactions were monitored by IR spectroscopy. IR spectra were recorded on a Perkin-Elmer 1710 FT-IR spectrometer, using calcium fluoride cells of 1 mm path length. EI and FAB mass spectra were recorded on a Fisons Autospec instrument. Elemental analyses were performed by the Microanalytical laboratory of the School of Chemistry, University of Bristol. All NMR measurements were performed on JEOL GX 400 spectrometer, operating in Fourier transform mode. The chemical shifts for $^1$H and $^{13}$C were referenced to internal TMS. All the reagents were commercial products of the highest purity available and used as received, except for $[\text{Ru}_2(\text{CO})_2(\mu-\text{CO})_2(\mu-\eta^2-\text{C(H)}=\text{C}=\text{C(Me)(Ph)})_2][\text{BF}_4]$, 2b, which was prepared according to the published method [12].

Synthesis of $[\text{Ru}_2(\text{CO})(\text{NCMe})(\mu-\text{CO})_2(\mu-\eta^1:\eta^2-\text{C(H)}=\text{C}=\text{C(Me)(Ph)})_2][\text{BF}_4]$, 3b. Complex $[\text{Ru}_2(\text{CO})_2(\mu-\text{CO})_2(\mu-\eta^1:\eta^2-\text{C(H)}=\text{C}=\text{C(Me)(Ph)})_2][\text{BF}_4]$, 2b (200 mg, 0.316 mmol), was dissolved in CH$_2$Cl$_2$ (20 mL) and treated with a solution of Me$_3$NO in MeCN (3.3 mL, 0.33 mmol). The mixture was stirred for 15 minutes. The solvent was then removed under reduced pressure, giving a brown residue. The IR spectrum of the residue, dissolved in CH$_2$Cl$_2$, indicated clean formation of 3b [ν(CO) 2005 (vs), 1870 (s) cm$^{-1}$].

Reaction of $[\text{Ru}_2(\text{CO})(\text{NCMe})(\mu-\text{CO})(\mu-\eta^1:\eta^2-\text{C(H)}=\text{C}=\text{C(Me)(Ph)})_2][\text{BF}_4]$, 3b, with NaBH$_4$. Complex $[\text{Ru}_2(\text{CO})(\mu-\text{CO})(\text{NCMe})(\mu-\eta^1:\eta^2-\text{C(H)}=\text{C}=\text{C(Me)(Ph)})_2][\text{BF}_4]$, 3b, freshly prepared by reacting $[\text{Ru}_2(\text{CO})_2(\mu-\text{CO})_2(\mu-\eta^1:\eta^2-\text{C(H)}=\text{C}=\text{C(Me)(Ph)})_2][\text{BF}_4]$, 2b (400 mg 0.633 mmol), with Me$_3$NO in MeCN solution, was dissolved in THF (30 mL) and treated with NaBH$_4$ (240 mg, 6.34 mmol). The mixture was stirred for 1 h, then the solvent was removed under reduced pressure. The resulting residue was introduced to an alumina column (15 x 2.5 cm) and chromatographed at –35 °C. The following products were collected by using varying amounts of n-hexane and CH$_2$Cl$_2$ as eluent and isolated as powders upon removal of the solvent under vacuum.

$[\text{Ru}_2(\text{CO})_2(\mu-\eta^2-\text{C}_a\text{H}_2=\text{C}_b(\text{Me})(\text{Ph}))_2][\text{BF}_4]$, 5a-b (eluent: hexane/CH$_2$Cl$_2$ 95:5; colour: pale yellow). Yield: 30 mg, 9.1 %. Anal. Calcd. for C$_{22}$H$_{20}$O$_2$Ru$_2$: C, 50.96; H, 3.89.
Found: C, 50.80; H, 4.00. FAB-MS: 520 m/z. IR (CH₂Cl₂) ν(CO) 1933 (m, sh), 1915 (s) cm⁻¹.

¹H NMR (CD₂Cl₂, 298 K) 5a: δ 7.72-6.63 (5 H, Ph); 5.04, 4.87 (s, 10 H, Cp); 3.52, 2.60 (d, 2J_HH = 4.0 Hz, 2 H, C₆H₂); 1.82 (s, 3 H, Me); 5b: δ 7.72-6.63 (5 H, Ph); 4.65, 4.35 (s, 10 H, Cp); 3.72, 2.85 (d, 2J_HH = 4.3 Hz, 2 H, C₆H₂); 2.17 (s, 3 H, Me). 5a/5b ratio 6:5. ¹³C{¹H} NMR (CD₂Cl₂, 298 K) 5a: δ 210.1, 206.5 (CO); 189.9 (C₆β); 149.7 (ipso-Ph); 128.5, 127.7, 126.4 (Ph); 88.0, 86.1 (C₆); 62.1 (C₆γ); 28.8 (Me); 25.0 (C₆δ); 5b: δ 208.0, 207.1 (CO); 187.8 (C₆β); 150.7 (ipso-Ph); 127.7, 125.4, 124.5 (Ph); 87.3, 84.0 (C₆); 56.9 (C₆γ); 31.3 (Me); 29.7 (C₆δ).

trans-[Ru₂(CO)₃(μ-H){μ-η¹:η²-C₆H=Ĉ₆=C₆(Me)(Ph)}(Cp)]₂, 6 (eluent: hexane/CH₂Cl₂ 90:10; colour: yellow). Yield: 50 mg, 15 %. Anal. Calcd. for C₂₂H₂₀O₂Ru₂: C, 50.96; H, 3.89.

Found: C, 50.79; H, 3.95. FAB-MS: 520 m/z. IR (CH₂Cl₂) ν(CO) 1972 (m), 1918 (s) cm⁻¹. ¹H NMR (CD₂Cl₂, 298 K) δ 8.80 (q, 5J_HH = 2.5 Hz, 1 H, C₆H); 8.43-6.98 (5 H, Ph); 5.16, 4.76 (s, 10 H, Cp); 2.25 (d, 2J_HH = 2.5 Hz, 3 H, Me); -13.73 (s, 1 H, μ-H). ¹³C{¹H} NMR (CD₂Cl₂, 298 K) δ 203.1, 202.4 (CO); 162.7 (C₆β); 140.1 (ipso-Ph); 128.4, 126.4, 125.0 (Ph); 113.5 (C₆α); 110.9 (C₆γ); 85.2, 85.0 (Cp); 22.2 (Me).

[Ru₃(CO)₅(μ-H)₃(Cp)₃], 7 (eluent: hexane/CH₂Cl₂ 90:10; colour: blue). Yield: 39 mg, 7.0 % [15].

cis-[Ru₂(CO)₃(μ-H){μ-η¹:η²-C₆H=Ĉ₆=C₆(Me)(Ph)}(Cp)]₂, 8 (eluent: hexane/CH₂Cl₂ 80:20; colour: yellow). Yield: 117 mg, 36 %. Anal. Calcd. for C₂₂H₂₀O₂Ru₂: C, 50.96; H, 3.89.

Found: C, 51.10; H, 3.77. FAB-MS: 520 m/z. IR (CH₂Cl₂) ν(CO) 1984 (vs), 1921 (s) cm⁻¹. ¹H NMR (CD₂Cl₂, 298 K) δ 9.12 (q, 5J_HH = 2.5 Hz, 1 H, C₆H); 7.81-6.57 (5 H, Ph); 5.14, 5.06 (s, 10 H, Cp); 2.24 (d, 2J_HH = 2.5 Hz, 3 H, Me); -14.20 (s, 1 H, μ-H). ¹³C{¹H} NMR (CD₂Cl₂, 298 K) δ 202.6, 201.7 (CO); 162.7 (C₆β); 141.1 (ipso-Ph); 128.2, 126.5, 125.0 (Ph); 118.1 (C₆α); 112.3 (C₆γ); 84.7, 84.1 (Cp); 22.3 (Me).


Found: C, 51.14; H, 3.79. FAB-MS: 520 m/z. IR (CH₂Cl₂) ν(CO) 1951 (vs), 1774 (s) cm⁻¹. ¹H NMR (CD₂Cl₂, 298 K) δ 10.75 (d, 3J_HH = 7.2 Hz, 1 H, C₆H); 7.58-6.50 (5 H, Ph); 5.08, 4.88 (s, 10 H, Cp); 4.71 (d, 3J_HH = 7.2 Hz, 1 H, C₆H); 1.59 (s, 3 H, Me). ¹³C{¹H} NMR (CD₂Cl₂, 298 K) δ 239.4 (μ-CO); 200.5 (CO); 155.4 (C₆α); 146.8 (ipso-Ph); 128.0, 127.7, 125.6 (Ph); 88.6, 85.7 (Cp); 82.7 (C₆β); 71.1 (C₆γ); 38.9 (Me).
Studies on the thermal stabilities of 5, 6, 8 and 9. Complex 8 (150 mg, 0.289 mmol) was dissolved in n-heptane (15 mL) and the solution was heated at reflux temperature for 30 minutes. The solution was then cooled to room temperature and the solvent was removed under vacuum, giving a yellow powder. IR and NMR spectroscopic data indicated complete conversion into complex 6. The latter was purified by filtration on alumina. Yield: 130 mg, 87%. The residue was then dissolved in n-heptane (15 mL), and heated at reflux temperature for 7 hours. The solution was then cooled and powdery 5b was cleanly obtained upon removal of the solvent. The identity of 5b was established by IR and 1H NMR spectroscopy. Yield: 120 mg, 80%. The yellow complex was dissolved in n-heptane (15 mL) and heated at reflux. IR spectroscopy revealed progressive formation of 9 and consumption of 5b, which was completed after 42 hours. The final mixture was chromatographed on an alumina column and a yellow band was collected by using a mixture of CH2Cl2 and n-hexane (1:1 v:v) as eluent. Removal of the solvent gave a yellow powder, which was identified as complex 9 by IR and 1H NMR spectroscopy. Yield: 79 mg, 53%. Finally, a solution of complex 9 was boiled in n-heptane (10 mL) for 7 hours, but no variation was detected via IR.

Synthesis of [Ru2(CO)2(X){µ-η¹:η²-Ca(H)=Cβ=Cγ(Me)(Ph)}(Cp)] [X = Cl, 4b; X = Br, 4c; X = I, 4d]. To a THF solution (20 mL) of complex [Ru2(CO)(NCMe)(µ-CO){µ-η¹:η²-C(H)=C=CH(Me)(Ph)}(Cp)] [BF4], 3b, freshly prepared from [Ru2(CO)2(µ-CO){µ-η¹:η²-C(H)=C=CH(Me)(Ph)}(Cp)] [BF4], 2b (200 mg 0.316 mmol), LiCl (260 mg, 6.13 mmol) was added. The resulting mixture was stirred for 1 h, then it was introduced to a short alumina column (3 x 2.5 cm). Quick filtration with a mixture of THF and methanol (95:5 v:v) afforded an orange solution of 4b, which gave an ochre yellow powder upon removal of the solvent under reduced pressure. Yield: 133 mg, 76%. Anal. Calcd. for C22H19ClO2Ru2: C, 47.78; H, 3.46; Cl, 6.41. Found: C, 47.65; H, 3.23; Cl, 6.19. EI-MS: 554 m/z. IR (CH2Cl2) ν(CO) 1989 (vs), 1887 cm⁻¹. 1H NMR (CDCl3, 298 K) Major isomer: δ 9.80 (q, 5JHH = 1.9 Hz, 1 H, CαH); 7.77-7.24 (5 H, Ph); 5.30, 4.78 (s, 10 H, Cp); 2.28 (d, 5JHH = 1.9 Hz, 3 H, Me); Minor isomer: δ 9.87 (q, 5JHH = 1.9 Hz, 1 H, CαH); 7.77-7.24 (5 H, Ph); 5.21, 4.84 (s, 10 H, Cp); 1.92 (d, 5JHH = 1.9 Hz, 3 H, Me). Isomer ratio 4:1. 13C{1H} NMR (CDCl3, 298 K) δ 213.0 (s-CO); 201.0 (CO); 158.4 (Cβ); 138.6 (ipso-Ph); 130.5 (Cα); 128.3, 126.2 (Ph); 114.6 (Cγ); 88.3, 86.1 (Cp); 22.6 (Me).
Compounds 4c-d were prepared by the same procedure described for 4b, by reacting 3b with the appropriate halide salt (LiBr, KI).

4c (yield: 85 %; colour: orange). Anal. Calcd. for C_{22}H_{19}BrO_2Ru_2: C, 44.23; H, 3.21; Br, 13.37. Found: C, 44.35; H, 3.15; Br, 13.12. FAB-MS: 598 m/z. IR (CH_2Cl_2) ν(CO) 1986 (vs), 1890 (s) cm\(^{-1}\). \(^1\)H NMR (d\(_8\)-toluene, 298 K) δ 9.80 (q, 5 J\(_{HH}\) = 2.0 Hz, 1 H, C\(_α\)H); 7.70-7.19 (5 H, Ph); 4.80, 4.60 (s, 10 H, Cp); 2.02 (d, 5 J\(_{HH}\) = 2.0 Hz, 3 H, Me).

13C{\(^1\)H} NMR (d\(_8\)-toluene, 298 K) δ 210.5 (s-CO); 201.9 (CO); 160.1 (C\(_β\)); 138.9 (ipso-Ph); 129.0 (C\(_α\)); 128.1, 127.9, 126.0 (Ph); 113.9 (C\(_γ\)); 88.1, 86.4 (Cp); 29.0 (Me).

4d (yield: 89 %; colour: orange). Anal. Calcd. for C_{22}H_{19}IO_2Ru_2: C, 41.00; H, 2.97; I, 19.69. Found: C, 41.09; H, 3.06; I, 19.84. FAB-MS: 646 m/z. IR (CH_2Cl_2) ν(CO) 1984 (vs), 1891 (s) cm\(^{-1}\). \(^1\)H NMR (d\(_8\)-toluene, 298 K) δ 9.82 (q, 5 J\(_{HH}\) = 2.2 Hz, 1 H, C\(_α\)H); 7.54-6.63 (5 H, Ph); 4.47, 4.28 (s, 10 H, Cp); 1.97 (d, 5 J\(_{HH}\) = 2.2 Hz, 3 H, Me).

13C{\(^1\)H} NMR (d\(_8\)-toluene, 298 K) δ 209.2 (s-CO); 202.9 (CO); 160.4 (C\(_β\)); 140.0 (ipso-Ph); 128.6, 126.5, 126.1 (Ph); 120.7 (C\(_α\)); 113.7 (C\(_γ\)); 88.2, 86.9 (Cp); 30.4 (Me).

**Synthesis of [Ru_2(CO)(μ-CO){μ-η\(^1\):η\(^3\)-C\(_α\)(Cl)C\(_β\)=C(Me)(Ph)}(Cp)_2]**, 10. Complex [Ru_2(CO)_2(Cl){μ-η\(^1\):η\(^3\)-C(H)=C(C(Me)(Ph))}(Cp)_2], 4b (120 mg, 0.217 mmol), was dissolved in dichloromethane (15 mL) and introduced to an alumina column (15 x 2.5 cm). An orange band corresponding to 10 was collected by using neat CH_2Cl_2 as eluent. Compound 10 was obtained as an air stable orange powder upon removal of the solvent under reduced pressure. Yield: 85 mg, 71 %. Anal. Calcd. for C_{22}H_{10}ClO_2Ru_2: C, 47.78; H, 3.46; Cl, 6.41. Found: C, 47.62; H, 3.29; Cl, 6.30. EI-MS: 554 [M\(^+\)], 526 [M\(^+\) – CO], 519 [M\(^+\) – Cl] m/z. IR (CH_2Cl_2) ν(CO) 1985 (vs), 1832 (s) cm\(^{-1}\). \(^1\)H NMR (CD_2Cl_2, 298 K) δ 7.80-7.05 (5 H, Ph); 5.59 (s, 1 H, C\(_β\)H); 5.26, 4.57 (s, 10 H, Cp); 1.48 (s, 3 H, Me).

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References


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