Effects of ligands on the migratory insertion of alkenes into rhodium–oxygen bonds†‡

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Migratory insertions of olefins into metal–oxygen bonds are elementary steps of important catalytic processes, but well characterised complexes that undergo this reaction are rare, and little information on the effects of ancillary ligands on such reactions has been gained. We report a series of alkoxo alkene complexes of rhodium(I) that contain a range of bidentate ligands and that undergo insertion of the alkene. Our results show that complexes containing less electron-donating ancillary ligands react faster than their counterparts containing more electron-donating ancillary ligands, and that complexes possessing ligands with larger bite angles react faster than those with smaller bite angles. External added ligands had several effects on the reactions, including an inhibition of olefin isomerisation in the product and acceleration of the displacement of the product from complexes of ancillary ligands with small bite angles. Complementary computational studies help elucidate the details of these insertion processes.

Introduction

Migratory insertions are a fundamental process of organometallic chemistry that occur in a wide range of commonly practiced catalytic reactions. The most common classes of olefin insertions involve reactions of metal-hydride, -alkyl, or -aryl complexes, and these reactions are part of many catalytic processes, including hydrogenation, olefin polymerisation, and arene alkylation and vinylation. The migratory insertion of alkenes into metal–heteroatom bonds is much less common than insertions into metal–hydrogen and metal–carbon bonds, and information on the influence of the properties of the metal–ligand fragments on the rates of these reactions is limited. The pathways by which C–N bonds with alkenes form are emerging, and studies have begun to reveal the factors that dictate when this bond forms by insertion into a metal–nitrogen bond and when it forms by external attack on a bound alkene. These studies also have begun to reveal the factors that control the rates of the insertion processes. However, related information on the insertions of alkenes into metal–oxygen bonds is lacking.

The insertions of alkenes into metal–oxygen bonds has been demonstrated to occur during several important catalytic reactions, including the classic Wacker oxidation under conditions of low chloride concentration, but examples of discrete complexes that undergo insertions of olefins into metal–alkoxide bonds are rare and have occurred in most cases with activated alkenes. For example, tetrafluoroethylene inserts into

Scheme 1 Insertion of alkenes into metal–alkoxide bonds.
the platinum(n)–methoxy bond of Pt(DPPE)(OMe)(Me) (Scheme 1A), acrylonitrile inserts into the platinum(n)–phenoxo bond in Pt(PEt3)2(OPh)(H), and tetracyanoethene inserts into the Pt(II)–O bond of a platinaoxetane. Yet, the only direct observation of the insertion of an unactivated alkene into a transition-metal–alkoxy bond is the reaction of PEt3-ligated rhodium(II) alkene complexes that insert coordinated alkenes (Scheme 1B). No discrete complexes of any metal containing other types of ancillary ligands have been reported that undergo this elementary reaction.

We report a series of rhodium(i) alkoxo alkene complexes that contain a range of bidentate phosphine ligands and that undergo insertion of the alkene to form cycloisomerised product. With these compounds, we have been able to reveal several effects of the ancillary diphosphine ligand and external ligand on the rate of the insertion of alkenes into the rhodium alkoxides. First, the steric properties of the ancillary ligand strongly affect the rate of the insertion, as well as the mechanism of the reaction after insertion; second, the electronic properties of the ligand affect the rate of insertion such that more electron-poor ligands lead to faster reactions than do more electron-rich ligands; third, free ligand (PPh3), which was added to bind the resulting Rh(i) hydride to form a stable product, appears to promote dissociation of the insertion-metal bond and thereby limit hydride-induced isomerisation of the complex of the terminal alkene to that of the internal alkene.

In contrast, product dissociation occurs without prior association of PPh3 when the steric congestion around the metal center is low. In contrast, product dissociation occurs without prior association of PPh3 when the steric congestion around the metal is higher.

**Results and discussion**

**Synthesis and insertion reactions of rhodium alkoxo olefin complexes**

A series of alkoxorhodium complexes shown in Fig. 1 was generated at low temperature. Addition of alcohol 2 to rhodium(i) silylamido diphosphine complexes 1a–m (see ESI† for the synthesis and characterisation of 1a–m) at −78 °C quantitatively generated Rh(i) alkene alkoxo complexes 3a–m. Complexes 3a–m were characterised in situ by 1H and 31P{1H} NMR spectroscopy at −40 °C. This set of complexes contains a series of diphosphine ligands with varying backbones connecting the phosphorus atoms and substituents on the phosphorus atoms that alter their structural and electronic properties.

Warming the alkoxo alkene complexes 3a–m led to the formation of furan products 4a–c, isomerised alkoxo alkene complexes 5a–m, and rhodium hydride species (Scheme 2). Methylenebis(trifluoromethyl)phenyl FF (1a) is the direct product from insertion of the alkene moiety of 3a–m into the rhodium–alkoxy bond and subsequent β-hydride elimination. Dihydrofurans 4b–c are likely formed by reaction of rhodium hydride byproducts with the alkene moiety of 4a. Isomerisation of the double bond in 3a–m gave alkoxo alkene complexes 5a–m with an internal alkene bound to rhodium. These complexes did not undergo alkene insertion. The combination of alkoxo alkene complex 3f and an equivalent of [Rh(DPPP)(PPh3)H] yielded a mixture of products including 52% of the isomeric alkoxo alkene complex 5f. This result indicates that rhodium hydride byproducts induce isomerisation of the complex of the terminal alkene to that of the internal alkene.

To occupy open coordination sites on the rhodium hydride species and thereby limit hydride-induced isomerisation...
reactions, reactions were run in the presence of free phosphine ligand. Triphenylphosphine was selected as the external ligand because it prevented isomerisation without reacting with alkoxo alkene complexes 3a–m (see Fig. S97†). With 1–20 equivalents of PPh3 added to the reaction of DPPP-ligated complex 3f, the amount of isomerised furans was reduced, and the amount of methylene furan 4a depended on the amount of added PPh3. Reactions conducted with higher concentrations of PPh3 formed less isomerised product than those conducted with lower concentrations of PPh3. The yield of 4a was also higher from reactions with a more strongly binding added ligand, such as PEt3 or P(OPh)3, instead of PPh3 (see Table S3†).

Broadly speaking, reactions of complexes containing more electron-poor diphosphines gave higher yields of furan products (96% and 88% respectively). Those of complexes containing more electron-rich diphosphines gave lower yields of furan products (88% and 80% respectively). The electronic properties of an ancillary LX-type ligand from cyclo-alkene complex 3f were indistinguishable from each other (3f: 4.1 ± 0.1 × 10−4 s−1; t1/2 = 30 min; 3f: kobs = 1.0 ± 0.1 × 10−3 s−1), indicating that the reactions of alkoxo alkene complexes 3b and 3f are zero order in added PPh3, respectively, when more than 1 equivalent of PPh3 is added.

Having confirmed that PPh3 is not interfering with the migratory insertion process, we analysed the effects of the ancillary ligands in 3a–m on this reaction by comparing the rate constants for reaction of a series of complexes containing different ligands. The effect of the electronic properties of the ancillary ligands was revealed by comparing the rate constants for complexes 3a–c, which contain three DPPF derivatives as ligands. The kinetic data (3a containing p-OMe ligand: kobs = 2.8 × 10−4 s−1; 3b containing p-H ligand: kobs = 4.2 × 10−4 s−1; 3c containing p-CF3 ligand: kobs = 5.6 × 10−4 s−1) indicate that the less electron-rich alkoxo alkene complexes react faster than more electron-rich alkoxo alkene complexes.

Two studies describing the effect of electronic properties of an ancillary ligand on the rate of insertion into a metal–amido complex have been previously reported.11,12 Wolfe reported the intramolecular insertion of alkynes into bisphosphine-ligated palladium(II)–amido bonds.13 The electronic effect of the ancillary ligand on this process was examined for complexes of the same three DPPF derivatives as in the study presented here. Complexes containing less electron-rich ancillary ligands underwent migratory insertion of the alkene into the metal–amido bond faster than complexes containing more electron-rich ligands. The similarity of the system studied by Wolfe and the system reported here allows a direct comparison of the electronic effects of these phosphines on the two insertion processes. The p-CF3 containing amido complex reacted more than five times faster than the analogous p-OMe containing complex, while the p-CF3 containing alkoxo complex reacted only two times faster than the analogous p-OMe containing complex. These olefin insertions into metal–alkoxo bonds in the current work are less sensitive to the electronic properties of the ligand than are the prior migratory insertions into metal–amido bonds.

Our group reported an additional migratory insertion of an alkene into a metal–amido bond in which the effect of the electronic properties of an ancillary LX-type ligand from cyclo-metalation of an aryl dialkyl phosphate at the ortho carbon of the aryl group was investigated.11 A substituent at the para position of the aryl ring was varied, and this single change in substituent from −CF3 to −OMe led to nearly an order of magnitude difference in the rate constant for the elementary process. The electronic effect of these phosphines on the two insertion processes. The p-CF3 containing amido complex reacted more than five times faster than the analogous p-OMe containing complex, while the p-CF3 containing alkoxo complex reacted only two times faster than the analogous p-OMe containing complex. These olefin insertions into metal–alkoxo bonds in the current work are less sensitive to the electronic properties of the ligand than are the prior migratory insertions into metal–amido bonds.

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step of insertion. While this reaction system is significantly different from the one reported here, it implies, together with the work of Wolfe,\textsuperscript{10} that the migratory insertion of alkynes into metal–alkoxy bonds are less sensitive to changes in the electronic properties of the ancillary ligands than the insertion of alkynes into metal–amido bonds. Together, these data begin to provide a general trend for migratory insertions into late metal–heteroatom bonds.\textsuperscript{10,12,29}

Complexes containing diphosphines with biphenyl linkages between the two phosphorus atoms (3h–i) reacted faster than the structurally related DPPB-ligated complex 3g, which has an alkyl ligand containing the same number of carbon atoms between the phosphorus atoms of the ancillary ligand as are present in the biaxial linkages of biaxial bisphosphines. The rates of reaction of the complexes of biphenyl diphosphines were even too fast to measure conveniently at 35 °C; thus, the rate constants for reaction of these two complexes were obtained at 0 °C (3h: \(k_{\text{obs}} = 4.5 \times 10^{-4} \text{ s}^{-1}\); 3i: \(k_{\text{obs}} = 2.1 \times 10^{-4} \text{ s}^{-1}\)). These high rates of reaction can be explained by the lower electron donation of the biaxial backbones in the ancillary ligands of complexes 3h–i than of the ferrocenyl or aliphatic backbones in complexes 3a–g.

The effect of the ligand bite angle was illustrated by comparing the rate constants for the structurally related complexes 3j–m (3j: \(3.2 \times 10^{-4} \text{ s}^{-1}\); 3k: \(2.2 \times 10^{-4} \text{ s}^{-1}\); 3l: \(7.2 \times 10^{-4} \text{ s}^{-1}\); 3m: \(18 \times 10^{-4} \text{ s}^{-1}\)). The complexes containing the ligands with the largest bite angles in this set, sixanthos- ligated 3l and xanthos-ligated 3m, reacted faster than homoxantphos-ligated 3l and DPEphos-ligated complexes 3j and 3k. Similarly, the observed rate constant for DPPB-ligated complex 3g (\(14 \times 10^{-4} \text{ s}^{-1}\)) was higher than the rate constant for DPPP-ligated complex 3f (\(8.4 \times 10^{-4} \text{ s}^{-1}\)). These results show that an increase in the size of the bite angle accelerates the insertion of alkynes into metal–alkoxy bonds. Analogous results were previously described for the insertion of alkynes into palladium(n)-amido,\textsuperscript{30,31} palladium(n)-methyl,\textsuperscript{42} and Group V metal–hydrido bonds.\textsuperscript{41} A larger bite angle likely increases the steric congestion of the alkene complexes prior to insertion and therefore makes insertion more feasible.

**Effect of free phospine on the reaction rates**

The relative rates for the reactions of DETPP-ligated complex 3d and DiPrPP-ligated complex 3e initially appeared to contradict the steric effect deduced from analysing the effect of bite angle on the reactions of complexes 3j–m. In the presence of 20 equivalents of PPh\(_3\), the less sterically hindered complex 3d reacted faster (\(k_{\text{obs}} = 7.5 \times 10^{-4} \text{ s}^{-1}\) at 35 °C) than the more sterically hindered complex 3e (\(k_{\text{obs}} = 2.2 \times 10^{-4} \text{ s}^{-1}\) at 35 °C). As discussed later in this paper, PPh\(_3\) appears to displace the product of insertion and \(\beta\)-hydrogen elimination from the hydrido alkene complex INT-B when the bite angle is small. Because this displacement reaction would be associative (Scheme 3), coordination of PPh\(_3\) to INT-B containing DETPP would be faster than coordination to the more sterically con- gested complex containing DiPrPP, causing the reaction of DETPP-ligated 3d to be faster than reaction of DiPrPP-ligated 3e when conducted in the presence of free PPh\(_3\). This proposal also implies that displacement of the product, rather than migratory insertion, is rate-limiting for the more hindered complex 3e in the presence of PPh\(_3\).

To test this hypothesis, the rates for the reactions of complexes 3d–e were measured in the absence of PPh\(_3\) to eliminate effects of the free ligand on product displacement. The reaction rates were too slow to be measured at 35 °C for both complexes, so the reactions were conducted at 70 °C. At this temperature, the more sterically congested complex 3e (\(k_{\text{obs}} = 8.8 \times 10^{-4} \text{ s}^{-1}\) at 70 °C) reacted significantly faster than its less sterically congested counterpart 3d (\(3.0 \times 10^{-4} \text{ s}^{-1}\) at 70 °C). This result is consistent with increased steric bulk on the ancillary ligands leading to faster rates of insertion and supports the hypothesis that increased steric hindrance of the ancillary ligand increases the rate of the insertion step, but retards the association of PPh\(_3\), thereby leading to slower reaction of 3e than of 3d in the presence of PPh\(_3\).

The large accelerating effect of PPh\(_3\) that was observed for the reactions of 3d–e was further investigated by conducting the reactions of other complexes in the presence and absence of PPh\(_3\). Similar rate accelerating effects were observed for complexes 3f–i, but the reaction was zero order in PPh\(_3\) in each case when more than 1 equivalent of PPh\(_3\) was added. In contrast, the rate constants for the reactions of complexes 3b and 3m were the same in the presence and absence of added PPh\(_3\). Thus, the reaction rates of rhodium diphosphine alkoxy alkene complexes containing ancillary ligands with natural bite angles greater than 99° were unaffected by added PPh\(_3\), whereas the reaction rates of rhodium diphosphine alkoxy alkene complexes containing ancillary ligands with bite angles smaller than 99° were faster in the presence of PPh\(_3\) than in the absence of this ligand, but zero order in PPh\(_3\) in the presence of 1–20 equivalents of PPh\(_3\).

We propose that 4a dissociates from INT-B without prior association of PPh\(_3\) for complexes containing ancillary ligands with bite angles >99° as in 3b and 3m (Scheme 3). In complexes containing ancillary ligands with bite angles <99°, the steric congestion around the metal centre is smaller. Therefore, direct dissociation from INT-B is less favoured, and coordination of free PPh\(_3\) (INT-C) leads to displacement of methylene-tetrahydrofuran 4a. Although the added PPh\(_3\) does not affect the rate of the reactions of the complexes containing ligands with bite angles greater than 99°, it does affect the products of the reaction. Binding of the external ligand to the resulting
rhodium hydride species inhibits isomerisation of 4a \((\text{vide supra})\).

To gain additional information on the role of \(\text{PPh}_3\), the rate constant for reaction of 3f was measured in the presence of 0.1 to 1 equivalent of \(\text{PPh}_3\). In each reaction, the initial reaction rate was the same, but the rate of conversion sharply decreased (more than that of an exponential decay) during the reaction, and at the time at which the rate decreased was shorter for reactions with lower initial concentrations \(\text{PPh}_3\) (see Table S1 and Fig. S2\(^\dagger\)). These data indicate that \(\text{PPh}_3\) is consumed in the reaction and once no free \(\text{PPh}_3\) is present, the reaction rate is smaller, as described above for reactions conducted in the absence of \(\text{PPh}_3\).

**Computational studies on the insertion process**

Computational studies were conducted with the Gaussian16 suite of programs\(^\dagger\) to analyse the feasibility of the proposed mechanism. The pathway for the reaction of DPPP-ligated complex 3f was calculated as a model for complexes containing ancillary ligands with bite angles smaller than 99° (Fig. 2). Geometries and energies were calculated at the PBE0-D3(BJ) level of theory\(^{41-48}\) using a mixed def2 basis set\(^{49,50}\) (see ESI\(\dagger\) for details), including SMD solvent correction for toluene\(^\dagger\) and thermal correction to 308 K. Additional single point calculations were performed for selected structures at the M06-D3/SDD/6-311+G(d,p) level of theory\(^{43-47}\) including the same thermal and solvent corrections. This combination of functional and basis set has been previously employed to study a rhodium(i)-catalysed reaction including a migratory insertion step.\(^{58}\)

The reaction of the complex containing DPPP was chosen for computational studies because this reaction was one influenced by the presence of \(\text{PPh}_3\). The activation free energy for the migratory insertion of the alkene moiety in 3f into the rhodium–alkoxo bond was calculated to be 20.5 kcal mol\(^{-1}\) (TS1). The free energy of the resulting rhodium alkyl complex INT-A is predicted to be 7.7 kcal mol\(^{-1}\) higher than that of 3f. These values correspond well to those reported for the migratory insertion of ethylene and propene into rhodium hydrido complexes.\(^{28}\) The free energy barrier for \(\beta\)-hydride elimination from INT-A is only 2.1 kcal mol\(^{-1}\) (TS2), and the free energy of the product (INT-B) is identical to that of INT-A. \(\beta\)-Hydride elimination, therefore, is predicted to be rapid and reversible.

If \(\text{PPh}_3\) is present in the current mixture, it can readily bind to the rhodium centre of INT-B to form INT-C (Path 3). The dissociation of methylenetetrahydrofuran 4a from INT-C is predicted to be exergonic by 9.0 kcal mol\(^{-1}\). Thus, all calculations predict that migratory insertion would be expected to be the rate limiting step in this reaction, and this prediction is consistent with our kinetic data on the effect of added ligand on the rate of the overall process.

In the absence of \(\text{PPh}_3\), Path 3 cannot occur. Direct dissociation of 4a from INT-B leads to the combination of rhodium hydride and alkene products that are computed, together, to lie 28.3 kcal mol\(^{-1}\) uphill of the starting complex (Path 1). This high energy makes isomerisation processes possible by reinsertion of the rhodium hydride moiety into the double bond of the alkene ligand (TS3, Path 2). Indeed, 4a formed in only 13% yield when the reaction of 3f was conducted without added \(\text{PPh}_3\), and internal alkene 4b and the complex 5f containing the isomerised alkoxo olefin were formed as the major products in 42% each (see ESI\(\dagger\) for details).

The energy values obtained from a second single point calculation with the M06 functional were indistinguishable from the results that were calculated with the PBE0 functional. Most important, the energy of the transition state leading to the isomerised rhodium alkyl species TS3 is predicted to be nearly identical to the energy of the transition state TS1 for the initial migratory insertion.

**Characterisation of rhodium alkyl complexes generated in the absence of \(\text{PPh}_3\)**

The reaction of DPPP-ligated complex 3f was monitored by \(^{31}\text{P}\) \({}_1\text{H}\) NMR spectroscopy in the absence of \(\text{PPh}_3\) at 35 °C. Complexes corresponding to two sets of doublets of doublets at 48.4 and 16.3 ppm, and 50.4 and 12.5 ppm accumulated. The coupling constants for these two species were nearly identical. Therefore, we assume that these complexes are either diastereomers or constitutional isomers in which the rhodium centre is bound to different carbon atoms on the product methylenetetrahydrofuran scaffold. Similar \(^{31}\text{P}\) \({}_1\text{H}\) NMR signals were observed during the reaction of the BINAP-ligated complex 3h (see ESI\(\dagger\)). In this case, the \(^{31}\text{P}\) \({}_1\text{H}\) NMR spectrum contained four sets of peaks, presumably corresponding to two sets of diastereomers resulting from the axial chirality of the BINAP ligand. For both ligands, the complexes accumulated over time when the reaction was monitored at 35 °C, but decayed after the reaction was run to completion at 70 °C. Thus, the observed complexes are potential intermediates to the final product 4a or to the side products 4b-c resulting from isomerisation observed in the absence of \(\text{PPh}_3\).
Information about the identity of these complexes was deduced from a series of NMR experiments. $^{31}$P($^1$H) DOSY NMR spectroscopy showed that the diffusion constants of the complexes are almost the same as the diffusion constants of the alkoxo alkene complexes 3f and 5f, suggesting that they are a similar size and likely monomeric. No hydrides of appropriate intensity were detected by $^1$H NMR spectroscopy at time points at which the complexes had accumulated, and no coupling between the small hydride peaks and the $^3$P($^1$H) NMR signals was detected by $^3$P($^1$H)-$^2$H-HMBC NMR spectroscopy. Thus, the observed complexes are not hydrido alkene complexes like INT-A.

To test whether the observed species are rhodium alkyl complexes, we added acid to the solution containing them. After generation of these species from complex 3f, a large excess of concentrated hydrochloric acid was added. The complexes were fully consumed, and 5-methyl-2,2-diphenyltetrahydrofuran$^{36}$ (6) was formed, albeit in 9% yield, as determined by $^1$H NMR spectroscopy using TMB as internal standard (Scheme 4). Alkenes 7 and 8 were identified as side products of this reaction in a combined yield of 36% among further unidentified side products. The formation of 7 and 8 can be explained by decomposition of complexes 3f and 5f in the presence of strong acid, respectively. The observation of tetrahydrofuran 6 as a product of this reaction suggests that protodemetalation of a rhodium–alkyl complex occurred. Rhodium–alkyl complexes are common intermediates but rarely observed and characterised in catalytic reactions.$^{44}$ We attempted to determine the position at which the protodemetalation occurred by adding deuterated acid instead of protic acid, but the $^2$H NMR spectrum of the crude reaction mixture was too complex to determine the site of deuteration of the organic product. GC/HRMS analysis of the crude reaction mixture showed a product with the expected mass of 6d, which further confirms the hypothesis that the observed species are rhodium alkyl complexes that undergo demetalation in the presence of strong acid.

Further information about the connectivity between rhodium and the organic ligand in the observed complexes was instead deduced from a series of NMR experiments. No coupling to rhodium was observed in the $^{13}$C($^1$H) NMR spectrum when the terminal carbon of the double bond in alcohol 2 was labelled with $^{13}$C.$^{45}$ This result indicates that the observed complex is not INT-A. In addition, monitoring the reaction of the alkoxo alkene complex from a variant of alcohol 2 containing deuterated phenyl groups by $^2$H NMR spectroscopy indicated the absence of rhodium–aryl interactions. Based on these experiments and the observation of two similar species in the reaction, it is likely that the complexes are the diastereomeric mixture of rhodium alkyl complexes in which the rhodium is bound to the 4-position of the methylenetetrahydrofuran scaffold (see Scheme 4 for the structure and the ESI‡ for a more detailed discussion).$^{42}$ This proposal that the observed complex contains rhodium bound to the 4-position of the tetrahydrofuryl unit is consistent with isomerisation processes dominating in the absence of added ligand when the bite angle of the ancillary ligand is small.

Conclusions

The studies described here reveal the effects of bound ancillary and free external ligands on the first set of well-characterised complexes containing varied ancillary ligands that insert alkene into rhodium–alkoxy bonds. Our results demonstrate that the insertion process is faster for complexes containing ancillary ligands that are more electron-poor and for which the bite angle is larger than for complexes with ancillary ligands that are more electron rich and for which the bite angle is smaller. Varying the electronic properties of the ancillary ligand appears to have a smaller effect on the rate of reaction than varying the steric properties of the ancillary ligand. The observed trends are similar to previously reported trends for the insertion of olefins into late transition metal–amido and –alkyl bonds.$^{10,12,19,40}$ Together, these results provide a comprehensive picture of the influences of ancillary ligands on migratory insertion processes of olefins.

Added ligand was found to affect the reaction in several ways. First, this external ligand affected the product distribution by limiting isomerisation processes caused by rhodium hydride byproducts. Second, the rates of reactions of complexes with bite angles smaller than 99° increased significantly in the presence of PPh$_3$, but the rates of reactions of complexes containing ligands with bite angles greater than 99° were unaffected by the presence of PPh$_3$. Computational results indicate that this difference in the effect of added PPh$_3$ is because PPh$_3$ changes the mechanism of product release from dissociative, in the absence of added ligand, to associative, in the presence of added ligand. The higher barrier for dissociation of the organic product than reinsertion of the rhodium hydride into the alkene bond enables isomerisation processes to dominate during reactions conducted without added PPh$_3$. A rhodium complex that is likely an alkyl intermediate in this isomerisation process accumulated during reactions without added PPh$_3$. We expect these findings to accelerate the design of new reactions involving migratory insertions of alkynes.

Conflicts of interest

There are no conflicts to declare.

![Scheme 4](image-url)
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Notes and references

62. A side product that is likely formed through a related metal alkyl complex was reported by Hay and Wolfe for cyclisation reactions resulting in arylated tetrahydrofurans (see ref. 23).