

Cite this: *Chem. Sci.*, 2019, 10, 7937

All publication charges for this article have been paid for by the Royal Society of Chemistry

Low-valent homobimetallic Rh complexes: influence of ligands on the structure and the intramolecular reactivity of Rh–H intermediates†

Pascal Jurt,^a Oleg G. Salnikov,^{bc} Thomas L. Gianetti,^{id *ad} Nikita V. Chukanov,^{bc} Matthew G. Baker,^a Grégoire Le Corre,^a Jaap E. Borger,^a Rene Verel,^{id a} Sébastien Gauthier,^e Olaf Fuhr,^{id g} Kirill V. Kovtunov,^{id bc} Alexey Fedorov,^{id af} Dieter Fenske,^g Igor V. Koptuyg,^{id bc} and Hansjörg Grützmacher,^{id *a}

Supporting two metal binding sites by a tailored polydentate trop-based (trop = 5*H*-dibenzo[*a,d*] cyclohepten-5-yl) ligand yields highly unsymmetric homobimetallic rhodium(I) complexes. Their reaction with hydrogen rapidly forms Rh hydrides that undergo an intramolecular semihydrogenation of two C≡C bonds of the trop ligand. This reaction is chemoselective and converts C≡C bonds to a bridging carbene and an olefinic ligand in the first and the second semihydrogenation steps, respectively. Stabilization by a bridging diphosphine ligand allows characterization of a Rh hydride species by advanced NMR techniques and may provide insight into possible elementary steps of H₂ activation by interfacial sites of heterogeneous Rh/C catalysts.

Received 2nd June 2019

Accepted 5th July 2019

DOI: 10.1039/c9sc02683e

rsc.li/chemical-science

Introduction

Bimetallic transition metal complexes have been investigated since the 1970s,¹ however despite numerous reports, only a few complexes are widely used in catalytic applications. The most prominent is a rhodium(II) acetate dimer and its derivatives exploited for carbene transfer reactions.² These and more recent bimetallic systems typically outperform monometallic alternatives in terms of activity and selectivity, functional group tolerance, and catalyst loading.^{3–5} The interest in homobimetallic catalysts has recently seen a renaissance.⁶ Selected catalytic applications include diazo-free cyclopropanation,⁷ small molecule activation,^{8,9} hydrogenation,¹⁰ hydroformylation¹¹ and C–C coupling reactions.¹² Notably, heterobimetallic complexes serve as models of the transmetalation step for the latter process.^{13–17} Most of these bimetallic species

fall into two groups: (i) heterobimetallic early-late transition metal complexes, or (ii) symmetric bimetallic complexes with a core consisting of two mid-to-late transition metals.¹⁸ Group (i) typically relies on ligands that combine in close proximity both soft and hard moieties, and feature a highly polar dative interaction from the late to the early transition metal.¹⁹ Group (ii) relies on symmetric bridging ligands, often leading to an apolar bimetallic interaction.²⁰ Complexes of the group (ii) have been of particular interest for the understanding of reactivity of small clusters.^{21–23}

Rhodium nanoparticles on support materials are widely used in heterogeneous catalysis and various industrial processes especially for hydrogenation–dehydrogenation reactions.²⁴ But the mode of interaction between the supported rhodium sites and H₂ is not fully understood.²⁵ A recent DFT study reported that hydrogen activation by small rhodium clusters on a carbon support features a bridging hydride species and a hydride ligand (Scheme 1a) that could be transferred to the carbon support.²⁶ However, such intermediates have not been observed experimentally to date. Low valent homobimetallic rhodium complexes serve as molecular models for supported Rh sites towards understanding the hydrogen activation on such materials.¹⁹ While representation of supported heterogeneous catalysts by a bimetallic model significantly reduces their complexity, this approach allows for a reliable identification of reaction products and is therefore insightful, despite the apparent oversimplification of the intrinsic complexity of heterogeneous catalysts. That said, examples of well-defined bimetallic Rh(I)–Rh(I) systems capable of activating H₂ are scarce.²⁷ Scheme 1b presents a rare dirhodium(I) complex that,

^aDepartment of Chemistry and Applied Biosciences, ETH Zürich, Switzerland. E-mail: hgruetzmacher@ethz.ch

^bInternational Tomography Center, SB RAS, Novosibirsk, Russia

^cNovosibirsk State University, Novosibirsk, Russia

^dDepartment of Chemistry and Biochemistry, University of Arizona, Tucson, USA. E-mail: tgianetti@email.arizona.edu

^eUniversité de Rennes, CNRS, ISCR-UMR 6226, F-35000 Rennes, France

^fDepartment of Mechanical and Process Engineering, ETH Zürich, Switzerland

^gInstitute of Nanotechnology, Karlsruhe Institute of Technology (KIT), Eggenstein-Leopoldshafen, Germany

† Electronic supplementary information (ESI) available: experimental spectra and modelling details. CCDC 1919533, 1919534, 1919542, 1919543 and 1919546. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9sc02683e



according to DFT calculations, activates dihydrogen leading to one bridging and one terminal hydride ligand, in a similar fashion as mentioned above for the supported rhodium centers. This mode of the hydrogen activation dissymmetrizes the dirhodium complex while simultaneously forming a stabilizing Rh–Rh bond.²⁸ Again, experimental evidence for such a dirhodium dihydride species is still lacking, although dissymmetric complexes with a dirhodium(i) core containing chloride ligands in place of the hydrides were reported.²⁹ A symmetric dirhodium dihydride complex, which forms an intermetallic bond, was characterized by NMR and IR (Scheme 1c).³⁰

In contrast to the exploitation of complexes with symmetric ligands, the advantages of dissymmetric ligands to control the reactivity of two adjacent Rh centers are underutilized.³¹ The dissymmetry of the electronic environment imposed by the ligand could not only enable otherwise inaccessible reactivity manifolds,^{19,32} but also allows modelling the environment and complexity of heterogeneous catalysts (metal-support interface, surface defects, *etc.*). In particular, a non-innocent ligand with multiple unsaturated C–C bonds could provide insights in the reactivity of Rh/C interfacial sites of metallic Rh nanoclusters or nanoparticles on carbon-based supports.

Previous work by the Grützmacher group showed that the bidentate concavely shaped tropPPh₂ (trop = 5*H*-dibenzo[*a,d*]cyclohepten-5-yl) ligand featuring both a σ -donor (Ph₂P group) and a π -accepting binding site (C=C_{trop}) enables a strong binding³³ to several transition metal centers including Pd,³⁴ Rh^{35,36} and Ir.^{36,37} In order to synthesize a homobimetallic low valent dirhodium complex, we developed a tailored ^{{(TMS)C \equiv C}₂tropPPh₂ ligand with a second binding site created by two alkyne moieties (*vide infra*). This framework supports a Rh₂(I) complex with labile triflate ligands *cis* to the polarized intermetallic Rh–Rh bond and demonstrates cooperativity of two metal sites in the hydrogen activation. However, resulting hydrides react intra-molecularly by adding hydrogen to the alkyne units of the ^{{(TMS)C \equiv C}₂tropPPh₂ ligand. We performed in}}

depth NMR studies, including the use of the parahydrogen-induced polarization (PHIP) technique,^{38–42} to elucidate the structure and transformations of these Rh hydride intermediates that convert the trop ligand into a carbene-like motif *via* semihydrogenation of the first C \equiv C bond. Remarkably, the semihydrogenation of the second C \equiv C bond proceeds with a different chemoselectivity forming a *cis*-olefinic ligand. Tuning the Rh coordination sphere by exchanging one triflate for a bis(diphenylphosphino)methane (dppm) ligand inhibits the semihydrogenation steps and allows to characterize the intermediate rhodium hydride species by NMR. Overall, reactivity of Rh₂ complexes in a carbon-rich ligand environment offers mechanistic insight on the net H₂ activation across a Rh–Rh bond and interaction of Rh–H species with a carbon support in Rh/C heterogeneous catalysts.

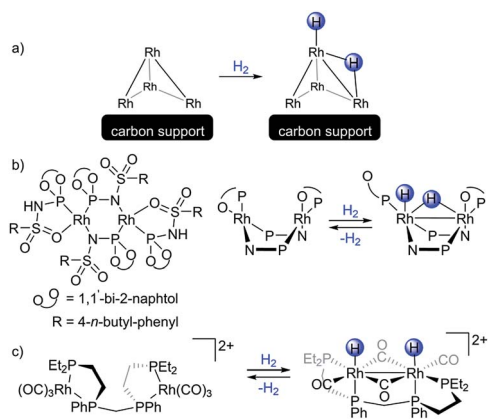
Results and discussion

Synthesis and characterization of 10,11-di-(trimethylsilyl) acetylene-5*H*-dibenzo[*a,d*]cyclohepten-5-diphenylphosphine and its Rh₂ complexes

The tropketone **1**⁴³ was converted to the polydentate trop ligand **5** in four steps with an overall yield of 34%, utilizing a Sonogashira protocol and conventional functional group transformation reactions (Fig. 1, top panel). Mixing **5** with one equivalent of [(C₂H₄)₂RhCl]₂ leads to the chloro bridged dimer **6** having two adjacent rhodium centers per trop ligand (83% yield). The abstraction of the chloride ligands in **6** with silver triflate gives the monomeric homobimetallic complex **7** in 82% yield. According to single crystal X-ray diffraction of **7**, two triflate ligands bridge between the two rhodium centers. The addition of one equivalent of diphenylphosphinomethane (dppm) displaces one triflate ligand to form the dppm adduct **8** (83% yield, Fig. 1, bottom panel).

Complexes **6–8** were characterized by single crystal X-ray diffraction methods. All complexes possess a distorted square planar geometry around Rh1 and a nearly ideal square planar environment around Rh2 ($\tau_{\text{Rh1}} = 0.29, 0.46$ and 0.37 , and $\tau_{\text{Rh2}} = 0.06, 0.01$ and 0.03 for **6**, **7** and **8** respectively).⁴⁴ The Rh–Rh contact in monomeric **7** is 2.6297(2) Å, that is 0.21 Å shorter than in dimeric **6** (2.8464(3) Å, Table 1). This shortening of the Rh–Rh distances is accompanied by an elongation of the P–Rh bond from 2.1829(3) Å in the chloro bridged dimer **6** to 2.2115(5) Å in the monomeric triflate complex **7**, indicating that the Rh–Rh interaction in **7** is stronger. However, the bond lengths of the coordinated C–C multiple bonds as well as the respective C–Rh distances do not differ much between **6** and **7** (Table 1). Dppm adduct **8** has a Rh–Rh bond of 2.7691(7) Å, an intermediate value between those of **6** and **7**. Additional significant changes are observed in the alkene/alkyne bonds *trans* to P2 and P3, which are elongated in **8** (Rh1-ct(C5–C6) 1.981(7) Å and Rh2-ct(C1–C2) 2.232(7) Å in **8** compared to Rh1-ct(C5–C6) 1.913(2) Å and Rh2-ct(C1–C2) 2.062(2) Å in **7**), explained by the higher *trans* influence of the phosphine ligand.⁴⁵

The ¹³C NMR olefinic chemical shifts for the chloro bridged dimer **6** and the monomeric triflate complex **7** are similar ($\delta^{13}\text{C}$



Scheme 1 (a) Hydrogen activation as calculated by DFT for small rhodium clusters on a carbon support. (b) A bimetallic Rh hydrogenation catalyst (left) and its proposed hydrogen activation mode, according to DFT (right). (c) A rare dirhodium(II) dihydride characterized by NMR.



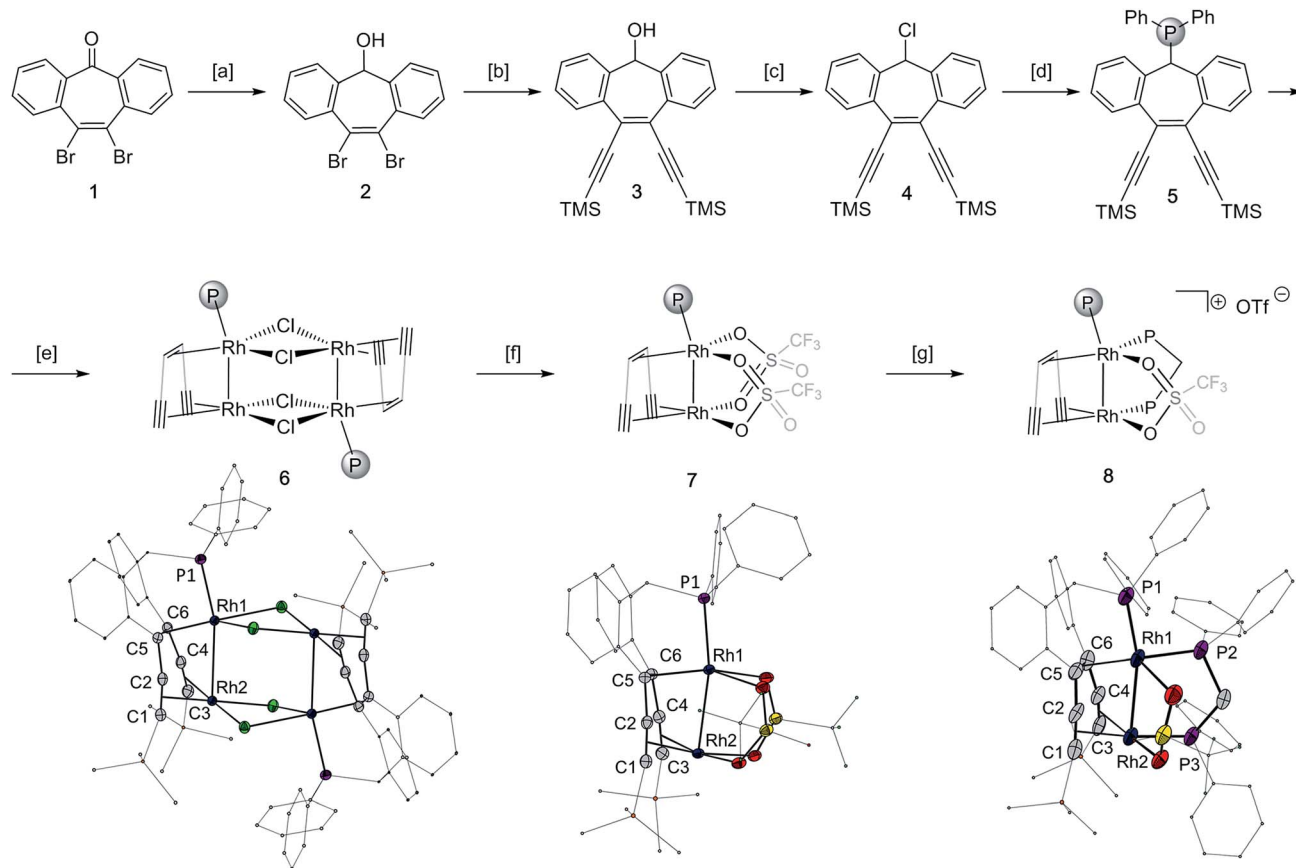


Fig. 1 [a] NaBH_4 , MeOH, 98%. [b] $\text{TMS-C}\equiv\text{CH}$, 5 mol% $[\text{Pd}(\text{PPh}_3)_4]$, CuI, toluene, Et_3N , 76%. [c] SOCl_2 , DCM, 0°C , 80%. [d] LiPPh_2 , toluene, 63%. [e] $[(\text{C}_2\text{H}_4)\text{RhCl}]_2$, benzene, 82%. [f] AgOTf , DCM, 83%. [g] $\text{Ph}_2\text{PCH}_2\text{PPh}_2$, DCM/hexane, 80%. The representation of ligand 5 is simplified in structures 6–8 for clarity. The crystal structures of bimetallic complexes 6–8 are given below the respective chemical structures and have the solvent molecules and non-coordinated triflate anions omitted for clarity.

Table 1 Selected bond lengths (Å) and angles ($^\circ$) for complexes 6 to 8. ct: center of coordinated multiple bond

Bond length or angle	6	7	8
Rh1–Rh2	2.8464(3)	2.6297(2)	2.7691(7)
Rh1–P1	2.1829(3)	2.2115(5)	2.2385(17)
C1–C2	1.242(3)	1.246(3)	1.235(10)
C3–C4	1.247(4)	1.248(3)	1.230(9)
C5–C6	1.499(3)	1.495(3)	1.475(10)
ct(C1–C2)–Rh2	2.077(2)	2.062(2)	2.232(7)
ct(C3–C4)–Rh2	2.054(3)	2.057(2)	2.051(6)
ct(C5–C6)–Rh1	1.962(2)	1.913(2)	1.981(7)
P1–Rh1–Rh2	162.708(19)	168.488(16)	164.92(5)
Rh1–Rh2–ct(C1–C2)	78.83(7)	87.48(6)	80.82(19)
Rh1–Rh2–ct(C3–C4)	81.01(7)	83.07(6)	80.93(19)
ct(C1–C2)–Rh2–ct(C3–C4)	89.67(10)	82.19(6)	85.9(2)
P1–Rh1–ct(C5–C6)	91.43(7)	90.56(8)	91.4(2)

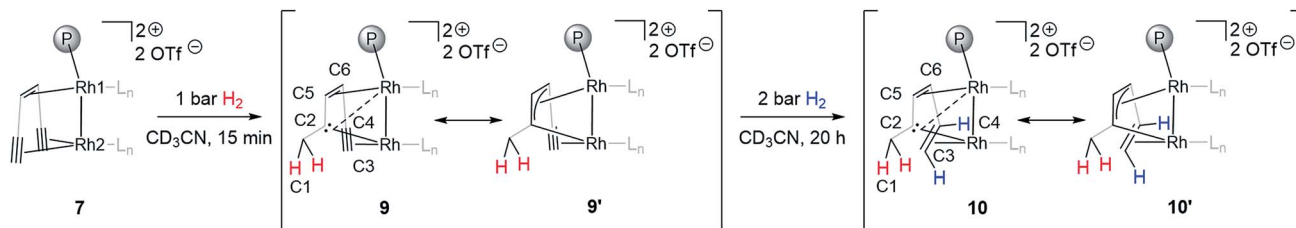
= 37.9 vs. 37.0 ppm for 6 and 7, respectively). However, a strong shielding is observed for both triple bonds in the triflate complex ($\delta^{13}\text{C} = 99.4$ to 86.6 ppm for $\text{TMS-C}\equiv\text{C}$ and $\delta^{13}\text{C} = 74.0$ to 65.3 for $\text{TMS-C}\equiv\text{C}$ for 6 and 7, respectively) suggesting that Rh2 site in 7 is more electron-rich than in 6. Likely, the Rh–Rh bond can be best described as a dative bond where electron

donation from Rh2 into the antibonding orbital of the Rh1–P bond occurs, similar to the bonding in early-late bimetallic transition metal complexes.¹⁹ For complex 8, this is supported by calculations, as the HOMO–2 and HOMO–3 orbitals show a clear overlap between the two metal centers, with a larger orbital contribution of Rh2 (Fig. S1†).

Reactivity of $[\text{Rh}_2\{\text{(TMS)C}\equiv\text{C}\}_2\text{tropPPh}_2]$ complexes with H_2

Adding hydrogen (1–2 bar) to the headspace of a J. Young NMR tube with monomeric triflate complex 7 dissolved in CD_3CN and following the reaction progress by NMR allows observing a stepwise hydrogenation of the two triple bonds of the supporting $\{\text{(TMS)C}\equiv\text{C}\}_2\text{tropPPh}_2$ ligand. The first alkyne semihydrogenation step proceeds quantitatively within 15 min and, strikingly, yields carbene species 9 (Scheme 2). Such chemoselective hydrogenation is uncommon and was only previously observed for alkynes bound to d^8 metal centers with a $\text{Pt}(\text{II})-(\mu\text{-H})_3\text{-Pt}(\text{II})$ core.⁴⁶ The semihydrogenation of the remaining triple bond leads to a coordinated alkene 10 (Scheme 2) and proceeds with a slower rate requiring ca. 20 h at 2 bar of H_2 for quantitative conversion. The carbene complex 9 and the carbene alkene complex 10 display characteristic signals in ^{13}C NMR spectra assigned to the bridging carbene at $\delta^{13}\text{C} = 171.1$ and





Scheme 2 Stepwise semihydrogenation of **7** in CD_3CN at room temperature followed by *in situ* NMR spectroscopy. L_n is CD_3CN . The representation of ligand **5** (Scheme 1) was simplified for clarity.

166.3 ppm and the methylene carbon at $\delta^{13}\text{C} = 35.5$ and 36.7 ppm, respectively. The methylene group formed is identified by the two diastereotopic proton signals in the ^1H NMR spectrum at $\delta^1\text{H} = 2.38, 3.18$ and 1.99, 3.03 ppm for **9** and **10**, respectively, showing a geminal $^2J_{\text{HH}}$ coupling of 12.1 and 13.1 Hz. 2D NMR experiments (COSY, HSQC and HMBC) support the structural assignment of complexes **9** and **10**; this data is presented in the ESI (Fig. S2–S5[†]). The bridging nature of the carbene ligand is further characterized by two rather different J_{CRh} coupling constants for **9** ($^1J_{\text{CRh}} = 33.9, 11.1$ Hz) and for **10** ($^1J_{\text{CRh}} = 35.7, 11.2$ Hz), which suggests that the Rh2 center has a closer contact to the carbene carbon than Rh1 (Scheme 2). Results of a single crystal X-ray analysis of **10** are consistent with the proposed structure, although the quality of data for **10** is rather poor (Fig. S6[†]). The observed $^1J_{\text{CRh}}$ coupling constants of 33.9 and 35.7 Hz are similar to the earlier reported values for dinuclear rhodium(i) complexes with bridging carbenes which likewise show $^1J_{\text{CRh}}$ in the range of 30–32 Hz.⁶ The assignment of *cis*-semihydrogenation in **10** is supported by two ^1H NMR signals at $\delta^1\text{H} = 2.58$ and 5.07 ppm coupled to each other with $^3J_{\text{HH}} = 11.3$ Hz.

The two olefinic carbons of the central trop double bond in **6–8** (C5 and C6 in Fig. 1) are strongly shielded, most likely due to the anisotropic effects of the neighboring alkyne group (the chemical shifts range from $\delta^{13}\text{C} = 37.0$ –48.7 ppm). Upon hydrogenation, these carbons show a remarkable difference in the chemical shift (**9**: $\delta^{13}\text{C} = 43.3$ and 98.8 ppm, **10**: $\delta^{13}\text{C} = 62.9$

and 100.0 ppm for C6 and C5 respectively, see Scheme 2). The chemical shift change in C6 is consistent with hydrogenation of the alkyne to alkene, lowering its anisotropic effect. However, the strong deshielding of C5 in both complexes indicates an overlap between the C2 carbene p orbital and the trop double bond, leading to an allyl-like bonding around Rh1.⁴⁷ This inference is further confirmed by the J_{CRh} coupling constants, which are much smaller for the C5 carbon than for the C6 carbon (Table 2, numbering in the trop ligand is according to Scheme 2). A similar bonding motif was reported with a dinickel(i) core in the solid state.⁴⁸ To summarize, complexes **9** and **10** contain a carbene carbon in conjugation with the central double bond of the trop ligand and this allyl-type ensemble is bridging to the Rh(i)–Rh(i) fragment. The ligand sphere of Rh2 in **9** and **10** in solution is likely stabilized by CD_3CN .

Furthermore, a characteristic shielding is observed in the ^{31}P NMR spectra when comparing the trop phosphorus signal at $\delta^{31}\text{P} = 104.6$ ppm in the monomeric triflate complex **7** with the respective signals at $\delta^{31}\text{P} = 59.7$ and 67.0 ppm in the carbene complex **9** and the carbene alkene complex **10**. This change is accompanied by a shift to a smaller $^1J_{\text{Rh1P}}$ coupling constant in **9** and **10** ($^1J_{\text{Rh1P}} = 127.2$ and 136.0 Hz) as compared to that in **7** ($^1J_{\text{Rh1P}} = 185.6$ Hz), indicating a higher trans influence of the Rh2 center in **9** and **10** compared to **7**. This suggests weakening of the P–Rh bond that is offset by strengthening of the Rh–Rh bond, attributed to the interaction with the bridging carbene ligand.⁴⁹ The latter could also be viewed as a distorted

Table 2 Selected NMR data of complexes **7**, **9**, and **10**

Nucleus	Position	7		9		10	
		δ (ppm)	J (Hz)	δ (ppm)	J (Hz)	δ (ppm)	J (Hz)
^{13}C	C1	75.4	$J_{\text{CRh}} = 11.9$	35.5		36.7	
	C2	97.0	$J_{\text{CRh}} = 7.7$	171.1	$J_{\text{CRh}} = 33.9, 11.1$	166.3	$J_{\text{CRh}} = 35.7, 11.2$
	C3	75.4	$J_{\text{CRh}} = 11.9$	76.9	$J_{\text{CRh}} = 11.6$	57.3	$J_{\text{CRh}} = 15.6$
	C4	97.0	$J_{\text{CRh}} = 7.7$	101.7	$J_{\text{CRh}} = 8.1$	81.5	$J_{\text{CRh}} = 12.4, 2.7$
	C5	38.1	$J_{\text{CRh}} = 14.9$	98.8	$J_{\text{CRh}} = 7.6$	100.0	$J_{\text{CRh}} = 7.1$
	C6	38.1	$J_{\text{CRh}} = 14.9$	43.3	$J_{\text{CRh}} = 11.0$	62.9	$J_{\text{CRh}} = 10.8$
^1H	C1H ₂			2.38, 3.18	$^2J_{\text{HH}} = 12.1$	1.99, 3.03	$^2J_{\text{HH}} = 13.1$
^{31}P		104.6	$^1J_{\text{PRh1}} = 185.6$	59.7	$^1J_{\text{PRh1}} = 127.2$	67.0	$^1J_{\text{PRh1}} = 136.0$
			$^2J_{\text{PRh2}} = 4.6$		$^2J_{\text{PRh2}} = 18.4$		$^2J_{\text{PRh2}} = 7.1$
^{103}Rh	P–Rh1–Rh2	–6852	$^1J_{\text{Rh1P}} = 186$	–7184	$^1J_{\text{Rh1P}} = 126$	–7163	$^1J_{\text{Rh1P}} = 136$
	P–Rh1–Rh2	— ^a		–6733		–7066	

^a No proton showed a significant coupling to Rh2 in the ^1H – ^{103}Rh HMQC spectrum.



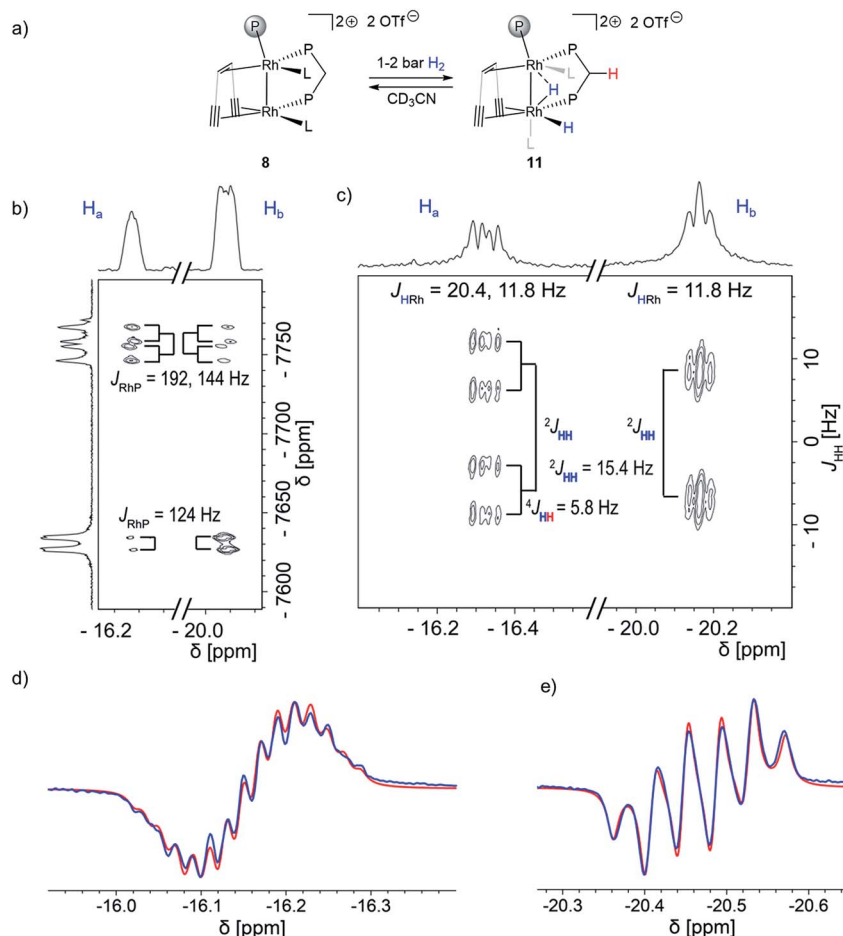
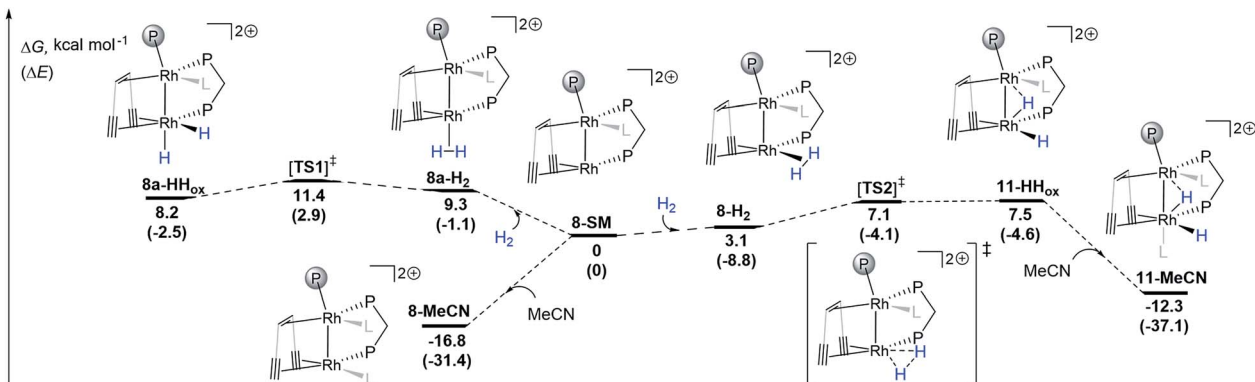


Fig. 2 Proposed activation of hydrogen by complex **8** (L is one acetonitrile ligand) and relevant spectroscopic data: (a) reaction of **8** with hydrogen, as proposed on the basis of NMR experiments, (b) J-resolved $^1\text{H}\{^{31}\text{P}\}$ spectrum, (c) $^1\text{H}-^{103}\text{Rh}$ HMQC spectrum, (d) PHIP ^1H spectrum (blue) with simulation (red), of hydride H_a , (e) PHIP $^1\text{H}\{^{31}\text{P}\}$ spectrum (blue) with simulation (red) of hydride H_b . The representation of ligand **5** was simplified in structures **8**, **11** and **12** for clarity.

dirhodacyclopropane.⁵⁰ Analysis of the ^{103}Rh NMR data reveals that Rh1 is significantly shifted upfield when comparing **7** with **9** and **10** ($\delta^{103}\text{Rh} = -6852, -7184, \text{ and } -7163$ ppm for **7**, **9** and **10**, respectively) further supporting a more electron rich Rh core. Comparing **9** and **10** using the ^{103}Rh NMR shift of Rh2 suggests

that the Rh2 site in **10** is more electron rich than in **9** ($\delta^{103}\text{Rh} = -6733, -7066$ ppm for **9** and **10**, respectively),⁵¹ which is consistent with a change in the ligand sphere from an alkyne to a weaker π accepting alkene. These results indicate that the bimetallic core becomes more electron rich with each hydrogenation step.



Scheme 3 DFT calculations (Gaussian09, $\omega\text{B97X-D/def2-SVP}$) for the hydrogen activation pathways from **8-SM** to **8a-HH_{ox}** and **11-HH_{ox}**. L is one acetonitrile ligand. The transition states were confirmed to connect the two respective energy minima by IRC calculations (see ESI†).



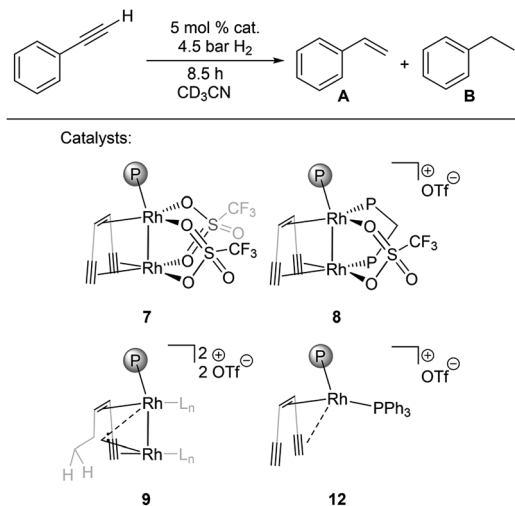


Fig. 3 Catalytic semihydrogenation of phenylacetylene by bimetallic catalysts 7–9 and the benchmark monometallic catalyst 12 (L_n is CD₃CN, ((TMS)C≡C)₂tropPPh₂ ligand is simplified for clarity).

Table 3 Results of the catalytic semihydrogenation of phenylacetylene. Conversion and selectivities are reported after 8.5 hours of the reaction

Entry	Catalyst	Conv. (%)	Sel. A (%)	Sel. B (%)	k _{obs} (10 ⁻² h ⁻¹)
1	7	39	90	10	5.9
2	8	79	97	3	23
3	9	39	97	3	6.7
4	12	16	>99	—	0.95

We have also performed the semihydrogenation of the ligand in a stepwise manner, where the carbene complex **9** was formed first under H₂, followed by deuteration to the carbene alkene complex **10** under a D₂ atmosphere. In this case, deuterium is only incorporated at the double bond (blue hydrogen atoms in Scheme 2) indicating irreversible ligand hydrogenation. This observation is consistent with the results of an experiment where **10** was formed *in situ* under an atmosphere of H₂ and subsequently placed under an atmosphere of D₂. In this case no deuterium incorporation is observed within 14 h, indicating that both hydrogenation steps are irreversible (Fig. S7 and S8†).

To gain further insight on how the H₂ molecule is activated at the Rh₂ fragment, *parahydrogen* (*p*-H₂)^{52,53} was used instead of H₂ for the hydrogenation of monomeric triflate complex **7**, resulting in no detectable hyperpolarized Rh hydride intermediates. Interestingly, the only PHIP hyperpolarized signals observed in the ¹H NMR spectra are assigned to the methylene CH₂ protons of carbene complex **9** (Fig. S10 and S11†), which establishes a pairwise hydrogen addition to the triple bond of the trop ligand when forming the bridging carbene complex **9**.

Next, we added 1–2 bar of H₂ to the solution of dppm adduct **8** in CD₃CN in a J. Young NMR tube and observed no formation of the bridging carbene-like moiety. Instead, a mixture of **8** and a Rh dihydride complex **11** (Fig. 2a) is formed that is stable for

several days at room temperature. Two characteristic hydride signals are observed at approximately δ¹H = −16.2 and −20.5 ppm. Dihydride **11** was further characterized by low temperature NMR experiments as well as using the PHIP technique. The removal of the H₂ (or the D₂) atmosphere from a J. Young NMR tube containing a mixture of the dppm adduct **8** and the dihydride **11** cleanly reforms **8**, indicating a fully reversible hydrogen activation (Fig. S12 and S13†). This is further supported by EXSY spectroscopy, variable temperature NMR and a partially negative line-shape (PNL) of the *ortho*-hydrogen peak in PHIP NMR (Fig. S14–S17†).^{38,54,55} Since PNL does also occur in PHIP experiments with **7**, the same mode of hydrogen activation is likely occurring with both species, **7** and **8**. Upon formation of *cis*-dihydride **11**, the Rh1 NMR resonance δ¹⁰³Rh1 = −7758 ppm is significantly shifted to lower frequencies. This low-frequency shift exceeds even the ones observed for **9** (δ¹⁰³Rh1 = −7184 ppm) and **10** (δ¹⁰³Rh1 = −7163 ppm). The Rh2 nucleus (δ¹⁰³Rh = −7630 ppm) shows likewise a strong shift to lower frequencies relative to the ¹⁰³Rh2 nuclei in **9** and **10** (δ¹⁰³Rh2 = −6733 and −7066 ppm, respectively), which is in agreement with the influence of strongly σ donating hydrides on both metal centers (Fig. 2b). ¹H{³¹P} J-resolved 2D NMR spectroscopy reveals the J_{HH} and J_{HRh} coupling constants (Fig. 2c). The two hydrides remain coupled in complex **11**, revealed by the ²J_{HH} = 15.4 Hz splitting in the indirect dimension. The hydride at δ¹H = −16.2 ppm shows an additional ⁴J_{HH} = 5.8 Hz coupling to another proton, assigned by COSY to one of the methylene protons of the dppm ligand (red in Fig. 2a). The NMR characterization of **11** was also completed by ¹H, ¹³C, ¹⁹F, ²⁹Si and ³¹P spectra (Fig. S21–S27†). Altogether, this data confirmed that dppm adduct **8** activates hydrogen reversibly and is in equilibrium with the dihydride species **11**. No semihydrogenation of the triple bonds is observed in this case.

Recording the J-resolved spectrum with ³¹P decoupling allows extracting the J_{HRh} coupling constants (Fig. 2c). While the hydride at δ¹H = −16.3 ppm appears as a doublet of doublets (J_{HRh} = 20.4, 11.8 Hz), the hydride at δ¹H = −20.2 ppm appears as a pseudo-triplet (J_{HRh} = 11.8 Hz). The observed ²J_{HH} coupling of 15.4 Hz is larger than typically found in traditional *cis* hydrides formed *via* oxidative addition (²J_{HH} = 7.2–9.5 Hz),⁵⁶ suggesting another geometry since higher coupling constants indicate larger angles between the substituents. Overall, these results are consistent with a bridging geometry for the dihydride **11** (Fig. 2a). Assignment of H_a *cis* to the intermetallic bond is based on the observation of a remarkable long-range ⁴J_{HH} coupling of H_a to a CH₂ proton (⁴J_{HH} = 5.8 Hz, highlighted red in Fig. 2a and b). Unfortunately, recording a J-resolved spectrum with ¹⁰³Rh decoupling was not successful due to the large difference in the chemical shift, and it was not possible to eliminate the J_{HRh} couplings from both metal centers at once. Therefore, the J_{HP} coupling constant could not be accessed with this approach. However, the ¹H–³¹P coupling constants could be extracted by fitting the observed PHIP signals (Fig. 2d and e). The PHIP spectrum in Fig. 2e was recorded with a ³¹P decoupling, which selectively eliminates only ³¹P couplings arising from the dppm moiety. This allows distinguishing coupling constants between the hydrides and dppm/trop phosphorous



centers. In addition, PHIP experiments allowed to determine the sign of the $^2J_{\text{HH}}$ coupling constant between the two hydrides which is negative $^2J_{\text{HH}} = -15.4$ Hz.

The proton H_a shows an $^2J_{\text{HP}}$ coupling to the dppm ^{31}P center of $^2J_{\text{HP}} = 13$ and 6 Hz, indicating a *cis* $^2J_{\text{HP}}$ coupling.⁵⁶ The second hydrogen atom H_b is likely close to the *cis* position of the trop phosphine center, as suggested by the couplings of $^2J_{\text{HP}} = 24$ and 18 Hz. The coupling to the second dppm phosphorous center is substantially smaller ($J_{\text{HP}} = 4$ Hz). However, the large J_{HH} coupling constant as well as the similar coupling constants to both Rh centers (resulting in the pseudo-triplet in Fig. 2b) are consistent with an interaction with the second Rh center. This assignment allows reporting a *cis* coupling constant of $^2J_{\text{HRh}} = 11.8$ Hz. This indicates that the close Rh–Rh contact is preserved after the addition of dihydrogen. Having assigned H_a as the terminal hydride, while H_b interacts with the two Rh centers unsymmetrically, the Rh–Rh core can be described either as a Rh(II)–Rh(II) center or a Rh(I) \rightarrow Rh(III) polarized bond, due to the bridging nature of H_b . Both descriptions imply a close Rh–Rh contact. We tested if this hydride reactivity could also be observed with other ligands and subjected tricyclohexylphosphine, diphenylphosphine oxide and triazabicyclodecene ligands to conditions of the *in situ* PHIP experiments demonstrating that only the dppm ligand leads to the hydride species. However, all PHIP experiments with the added ligand except triazabicyclodecene showed the hyperpolarized signals of methylene CH_2 protons with very similar chemical shifts and coupling constants as observed for **9**. In addition, a partially negative lineshape signal for the *ortho*hydrogen was observed for all ligands, suggesting that hydrogen activation is reversible (Fig. S31–S33[†]).

DFT calculations ($\omega\text{B97X-D/def2-SVP}$) support the proposed reversible hydrogen activation of the monomeric triflate complex **8** leading to the dihydride **11** (Scheme 3). From the computed **8-SM** intermediate, two H_2 adducts can be formed: with the H_2 molecule coordinated *cis* (**8-H₂**) or *trans* (**8a-H₂**) to Rh1 (Scheme 3). Both complexes can undergo oxidative addition steps *via* activated complexes which are located at energetically low lying transition states ($[\text{TS1}]^\ddagger$: $\Delta G = 11.4$ kcal mol⁻¹; $[\text{TS2}]^\ddagger$: $\Delta G = 7.1$ kcal mol⁻¹), leading to dihydrides **8a-HH_{ox}** and **11-HH_{ox}**, respectively. With the exception of **11-MeCN**, all intermediates are relatively close in energy and are expected to be accessible from **8-SM** at room temperature. Since **11-HH_{ox}** has an open coordination site *trans* to Rh1, binding of one acetonitrile ligand occurs giving a more stable species **11-MeCN** ($\Delta G = -19.8$ kcal mol⁻¹) with a distorted octahedral coordination environment around Rh2, as expected for Rh(III) d⁶ complexes. The energy of the coordinatively saturated **11-MeCN** is only slightly higher than that of **8-MeCN** ($\Delta G = -12.3$ and -16.8 kcal mol⁻¹, respectively). These two complexes are therefore expected to slowly interconvert and be observable in solution, which is indeed supported by NMR spectroscopy. We note that $\Delta\Delta G$ values from DFT calculations and variable temperature NMR measurements obtained from a Van't Hoff plot (Fig. S16[†]) are in reasonable agreement ($\Delta\Delta G = 4.5$ and 1.1 kcal mol⁻¹, respectively). It is of note that since the geometrical reorganization in going from $[\text{TS2}]^\ddagger$ to **11-HH_{ox}** is

only minimal, the barrier for this step is very low and we attribute the small positive energy difference between **11-HH_{ox}** and $[\text{TS2}]^\ddagger$ to the numerical inaccuracy of the applied DFT method. In addition, we note that the bimetallic hydrogen activation pathway involving a four-membered M_2H_2 transition state was not considered because this reaction is symmetry forbidden.⁵⁷

Interestingly, a bridging hydride resembling that in **11** has been proposed and calculated in both rhodium-based homogeneous bimetallic²⁸ and heterogeneous systems.²⁶ This observation further underlines the utility of low valent homobimetallic complexes with ligands containing alkene and alkyne binding sites in understanding elementary steps in heterogeneous catalysts which are deposited on carbon support materials.

Bimetallic dirhodium complexes **7–9** and the monometallic catalyst **12** were then tested as catalysts for the semi-hydrogenation of phenylacetylene (5% catalyst loading, 25 °C, Fig. 3, for details see ESI[†]). No activation period was observed (Fig. S41[†]). The dppm adduct **8** showed highest performance with a selectivity for styrene of 96% at 78% conversion and k_{obs} around 3–5 times higher than those for monomeric triflate complex **7** and carbene complex **9** (Table 3, entries 1–3). The similar rates k_{obs} observed with **7** and **9** indicate that the *in situ* hydrogenation of **7** to **9** takes place under the catalytic conditions and leads to the same active species, consistent with the presence of the hyperpolarized CH_2 signals characteristic for **9** in these catalytic reactions according to *in situ* PHIP NMR (Fig. S42[†]). Importantly, PHIP results also show that the dihydride species **11** form from **8** under catalytic conditions (Fig. S43[†]).

Remarkably, the monometallic catalyst **12** converts phenylacetylene significantly slower than all tested bimetallic complexes, with a rate of $k_{\text{obs}} = 9.5 \times 10^{-3}$ h⁻¹ (Table 3, entry 4). We speculate that the second metal serves as an electron reservoir which helps to avoid the formation of an inert trop-Rh(III) d⁶ species⁵⁸ formed by oxidative addition of dihydrogen. PHIP experiment performed under catalytic conditions with the ddpm adduct **8** indicates a pairwise hydrogen transfer to phenylacetylene forming styrene, *i.e.* hydrogen is activated by **8** to give dihydride **11**, followed by the transfer of the two hydrides to the same substrate molecule (Fig. S43[†]).

Conclusions

The $\{\text{TMS}\text{C}\equiv\text{C}\}_2\text{tropPPh}_2$ molecule with one phosphane, one alkene, and two alkyne donor sites was designed to allow the synthesis of a new family of low-valent homobimetallic Rh(I)–Rh(I) complexes. The reactivity of these complexes provides insight into possible metal–metal cooperation in hydrogen activation reactions. The unique ligand environment of the Rh2 site in $\{\text{TMS}\text{C}\equiv\text{C}\}_2\text{tropPPh}_2$ complexes makes this ligand an interesting model for the interface between metal clusters and particles and a carbon support material as found in Rh/C heterogeneous catalysts. Two distinct modes of intramolecular hydrogenation of the ligand were identified leading to a bridging carbene or an olefin ligand. When stabilized by a dppm ligand, an intermediate hydride species could be



characterized by advanced NMR methods which show that this species has a structure with a bridging and a terminal hydride ligand.

Conflicts of interest

The authors declare no conflicts of interest.

Acknowledgements

We gratefully acknowledge financial support of Swiss National Science Foundation. I. V. K. acknowledges the grant from the Russian Science Foundation (19-13-00172) for the support of hydrogenation experiments with parahydrogen. K. V. K thanks SB RAS integrated research program (# 0333-2018-0006/II.1.13) for parahydrogen activation studies and Ministry of Science and Higher Education of the RF (AAAA-A16-116121510087-5) for access to NMR equipment. A. F. thanks the Holcim Stiftung for the Habilitation fellowship.

Notes and references

- H. Vahrenkamp, *Angew. Chem., Int. Ed.*, 1978, **17**, 379–392.
- M. P. Doyle, *Chem. Rev.*, 1986, **86**, 919–939.
- H. M. L. Davies and D. Morton, *Chem. Soc. Rev.*, 2011, **40**, 1857–1869.
- H. M. L. Davies and J. R. Manning, *Nature*, 2008, **451**, 417.
- J. Adams and D. M. Spero, *Tetrahedron*, 1991, **47**, 1765–1808.
- J. F. Berry and C. M. Thomas, *Dalton Trans.*, 2017, **46**, 5472–5473.
- Y.-Y. Zhou and C. Uyeda, *Angew. Chem., Int. Ed.*, 2016, **55**, 3171–3175.
- J. Ye, R. C. Cammarota, J. Xie, M. V. Vollmer, D. G. Truhlar, C. J. Cramer, C. C. Lu and L. Gagliardi, *ACS Catal.*, 2018, **8**, 4955–4968.
- R. B. Siedschlag, V. Bernales, K. D. Vogiatzis, N. Planas, L. J. Clouston, E. Bill, L. Gagliardi and C. C. Lu, *J. Am. Chem. Soc.*, 2015, **137**, 4638–4641.
- M. K. Karunananda and N. P. Mankad, *ACS Catal.*, 2017, **7**, 6110–6119.
- M. E. Broussard, B. Juma, S. G. Train, W.-J. Peng, S. A. Laneman and G. G. Stanley, *Science*, 1993, **260**, 1784–1788.
- D. R. Pye and N. P. Mankad, *Chem. Sci.*, 2017, **8**, 1705–1718.
- M.-E. Moret, in *Higher Oxidation State Organopalladium and Platinum Chemistry*, ed. A. J. Canty, Springer Berlin Heidelberg, Berlin, Heidelberg, 2011, pp. 157–184.
- R. J. Oeschger, D. H. Ringger and P. Chen, *Organometallics*, 2015, **34**, 3888–3892.
- D. Serra, M.-E. Moret and P. Chen, *J. Am. Chem. Soc.*, 2011, **133**, 8914–8926.
- R. J. Oeschger and P. Chen, *J. Am. Chem. Soc.*, 2017, **139**, 1069–1072.
- R. J. Oeschger and P. Chen, *Organometallics*, 2017, **36**, 1465–1468.
- J. P. Krogman and C. M. Thomas, *Chem. Commun.*, 2014, **50**, 5115–5127.
- R. C. Cammarota, L. J. Clouston and C. C. Lu, *Coord. Chem. Rev.*, 2017, **334**, 100–111.
- D. C. Powers and T. Ritter, *Acc. Chem. Res.*, 2012, **45**, 840–850.
- P. M. Maitlis, H. C. Long, R. Quyoum, M. L. Turner and Z.-Q. Wang, *Chem. Commun.*, 1996, 1–8.
- P. M. Maitlis, *J. Organomet. Chem.*, 2004, **689**, 4366–4374.
- P. M. Maitlis and V. Zanotti, *Catal. Lett.*, 2008, **122**, 80–83.
- C. H. Bartholomew and R. J. Farrauto, *Hydrogenation and Dehydrogenation of Organic Compounds*, John Wiley and Sons, Hoboken, NJ, 2nd edn, 2005.
- P. Serna and B. C. Gates, *Acc. Chem. Res.*, 2014, **47**, 2612–2620.
- A. Dutta and P. Mondal, *J. Phys. Chem. C*, 2018, **122**, 16925–16939.
- A. S. Weller and J. S. McIndoe, *Eur. J. Inorg. Chem.*, 2007, 4411–4423.
- F. W. Patureau, S. de Boer, M. Kuil, J. Meeuwissen, P.-A. R. Breuil, M. A. Siegler, A. L. Spek, A. J. Sandee, B. de Bruin and J. N. H. Reek, *J. Am. Chem. Soc.*, 2009, **131**, 6683–6685.
- A. J. Esswein, A. S. Veige and D. G. Nocera, *J. Am. Chem. Soc.*, 2005, **127**, 16641–16651.
- R. C. Matthews, D. K. Howell, W.-J. Peng, S. G. Train, W. D. Treleaven and G. G. Stanley, *Angew. Chem., Int. Ed. Engl.*, 1996, **35**, 2253–2256.
- J. F. Berry and C. C. Lu, *Inorg. Chem.*, 2017, **56**, 7577–7581.
- I. G. Powers and C. Uyeda, *ACS Catal.*, 2017, **7**, 936–958.
- J. Thomaier, S. Boulmaâz, H. Schönberg, H. Rüegger, A. Currao, H. Grützmacher, H. Hillebrecht and H. Pritzkow, *New J. Chem.*, 1998, **22**, 947–958.
- L. Bettucci, C. Bianchini, W. Oberhauser, M. Vogt and H. Grützmacher, *Dalton Trans.*, 2010, **39**, 6509–6517.
- H. Schönberg, S. Boulmaâz, M. Wörle, L. Liesum, A. Schweiger and H. Grützmacher, *Angew. Chem., Int. Ed.*, 1998, **37**, 1423–1426.
- F. Breher, H. Rüegger, M. Mlakar, M. Rudolph, S. Deblon, H. Schönberg, S. Boulmaâz, J. Thomaier and H. Grützmacher, *Chem.-Eur. J.*, 2004, **10**, 641–653.
- S. Boulmaâz, S. Loss, H. Schönberg, S. Deblon, M. Wörle, R. Nesper, H. Grützmacher and M. Mlakar, *Chem. Commun.*, 1998, 2623–2624.
- A. S. Kiryutin, G. Sauer, A. V. Yurkovskaya, H.-H. Limbach, K. L. Ivanov and G. Buntkowsky, *J. Phys. Chem. C*, 2017, **121**, 9879–9888.
- R. A. Green, R. W. Adams, S. B. Duckett, R. E. Mewis, D. C. Williamson and G. G. R. Green, *Prog. Nucl. Magn. Reson. Spectrosc.*, 2012, **67**, 1–48.
- T. C. Eisenschmid, R. U. Kirss, P. P. Deutsch, S. I. Hommeltoft, R. Eisenberg, J. Bargon, R. G. Lawler and A. L. Balch, *J. Am. Chem. Soc.*, 1987, **109**, 8089–8091.
- C. R. Bowers and D. P. Weitekamp, *J. Am. Chem. Soc.*, 1987, **109**, 5541–5542.
- S. B. Duckett and N. J. Wood, *Coord. Chem. Rev.*, 2008, **252**, 2278–2291.
- B. Le Bourdonnec and R. E. Dolle, *US Pat.*, 20080275131, 2008.



- 44 A. W. Addison, T. N. Rao, J. Reedijk, J. van Rijn and G. C. Verschoor, *J. Chem. Soc., Dalton Trans.*, 1984, 1349–1356.
- 45 P. K. Sajith and C. H. Suresh, *Dalton Trans.*, 2010, **39**, 815–822.
- 46 G. Banditelli and A. L. Bandini, *Organometallics*, 2006, **25**, 1578–1582.
- 47 W. A. Herrmann, *Angew. Chem., Int. Ed.*, 1978, **17**, 800–812.
- 48 C. A. Laskowski and G. L. Hillhouse, *Chem. Sci.*, 2011, **2**, 321–325.
- 49 A. Pidcock, R. E. Richards and L. M. Venanzi, *J. Chem. Soc. A*, 1966, 1707–1710.
- 50 W. A. Herrmann, J. Plank, D. Riedel, M. L. Ziegler, K. Weidenhammer, E. Guggolz and B. Balbach, *J. Am. Chem. Soc.*, 1981, **103**, 63–75.
- 51 P. B. Graham, M. D. Rausch, K. Taeschler and W. Von Philipsborn, *Organometallics*, 1991, **10**, 3049–3052.
- 52 O. G. Salnikov, H.-J. Liu, A. Fedorov, D. B. Burueva, K. V. Kovtunov, C. Copéret and I. V. Koptug, *Chem. Sci.*, 2017, **8**, 2426–2430.
- 53 K. V. Kovtunov, D. A. Barskiy, R. V. Shchepin, A. M. Coffey, K. W. Waddell, I. V. Koptug and E. Y. Chekmenev, *Anal. Chem.*, 2014, **86**, 6192–6196.
- 54 V. V. Zhivonitko, V.-V. Telkki, K. Chernichenko, T. Repo, M. Leskelä, V. Sumerin and I. V. Koptug, *J. Am. Chem. Soc.*, 2014, **136**, 598–601.
- 55 D. A. Barskiy, K. V. Kovtunov, I. V. Koptug, P. He, K. A. Groome, Q. A. Best, F. Shi, B. M. Goodson, R. V. Shchepin, A. M. Coffey, K. W. Waddell and E. Y. Chekmenev, *J. Am. Chem. Soc.*, 2014, **136**, 3322–3325.
- 56 A. Koch and J. Bargon, *Inorg. Chem.*, 2001, **40**, 533–539.
- 57 G. Trinquier and R. Hoffmann, *Organometallics*, 1984, **3**, 370–380.
- 58 F. F. Puschmann, H. Grützmacher and B. d. Bruin, *J. Am. Chem. Soc.*, 2010, **132**, 73–75.

