

Cite this: *Chem. Sci.*, 2016, 7, 1996

A versatile method for the preparation of carbon–rhodium hybrid catalysts on graphene and carbon black†

Chin Min Wong,^{ab} D. Barney Walker,^{ab} Alexander H. Soeriyadi,^a J. Justin Gooding^a and Barbara A. Messerle^{*ab}

Strategies for combining the selectivity and efficiency of homogeneous organometallic catalysts with the versatility of heterogeneous catalysts are urgently needed. Herein a direct and modular methodology is presented that provides rapid access to well-defined carbon–rhodium hybrid catalysts. A pre-synthesized Rh(I) complex containing a carbene-triazole ligand was found to be stable for direct immobilization onto unactivated graphene, carbon black and glassy carbon electrodes. Characterization of the heterogeneous systems using X-ray photoelectron spectroscopy (XPS), thermogravimetric analysis (TGA), inductively coupled plasma-optical emission spectroscopy/mass spectrometry (ICP-OES/MS), Raman spectroscopy, scanning electron microscopy (SEM) and transmission electron microscopy (TEM) confirmed the well-defined nature of the hybrid catalysts. The hybrid catalysts show excellent activity, comparable to that of the homogeneous system for the hydrosilylation of diphenylacetylene, with turnover numbers ranging from 5000 to 48 000. These catalysts are the best reported to date for the hydrosilylation of diphenylacetylene. In common with conventional heterogeneous catalysts, high reusability, due to a lack of Rh metal leaching, was also observed for all carbon–rhodium complexes under investigation.

Received 6th October 2015
Accepted 7th December 2015

DOI: 10.1039/c5sc03787e

www.rsc.org/chemicalscience

Introduction

The assembly of highly efficient and selective homogeneous systems onto heterogeneous solid materials is a facile route to the development of optimized and recyclable hybrid systems with well-defined active sites.^{1–3} While a variety of catalysts covalently tethered to carbon based scaffolds (*e.g.* activated carbon, graphene/graphene oxide and carbon nanotubes) and other well known supports (*e.g.* aluminosilicates, organic resins and metal organic frameworks) have been reported,^{4–12} the fabrication of these hybrid systems is often complicated, requiring multiple reaction steps.

The development of hybrid catalysts consisting of well-defined organometallic complexes on solid supports can be established using both non-covalent interactions or covalent bonding. In terms of producing hybrid catalysts that are covalently bonded to the solid support, the synthesis route typically

occurs *via* at least two steps, where initially a precursor ligand is attached to the surface then – in a second and separate step – a transition metal centre is introduced. The first step is normally dictated by the nature of the functional groups on the solid surface. For example, reactive Si–OH moieties on silica surfaces can be effectively used to initiate ligand immobilization.^{13,14} However, certain supports require additional surface activation in order to achieve the functional groups that are needed for ligand modification. This is a common strategy for functionalization of graphene surfaces, whereby pre-oxidation of graphene to graphene oxide (GO) is often necessary prior to functionalization with an appropriate ligand motif.^{15–17} At times, reduction back to reduced GO is also performed after ligand immobilization and complexation with the metal precursor.^{18–20} This complicated multistep assembly to the hybrid system is not ideal for pristine materials, such as pure graphene, where it is desirable that its regular, conjugated surface is retained.^{21–24} The second step (complexation of metal with the immobilized ligand) in the stepwise assembly of hybrid catalysts could create an additional complication, as incomplete complexation of the immobilized ligand can occur. Furthermore, the common use of an excess of metal precursors increases the chance of nonspecific adsorption of metals onto the surface.

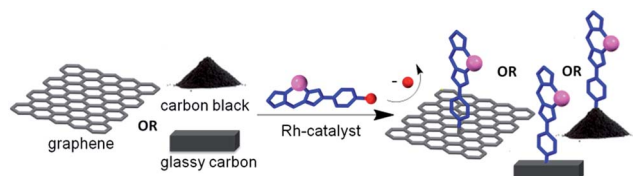
In designing new hybrid catalysts, it would be highly desirable to have a simple, general method for attaching optimized

^aDepartment of Chemistry, University of New South Wales, Sydney, NSW 2052, Australia

^bDepartment of Chemistry and Biomolecular Sciences, Macquarie University, Sydney, NSW 2109, Australia. E-mail: barbara.messerle@mq.edu.au

† Electronic supplementary information (ESI) available: Additional schemes, figures and tables, characterization of the complexes and hybrid complexes on carbon surfaces, XPS, TGA, ICP-MS/OES, IR, Raman, SEM, TEM and electrochemical data and calculations are included. Full catalysis and recycling tables are also included. See DOI: 10.1039/c5sc03787e





Scheme 1 Illustration of the direct attachment of catalyst to G, CB or GC without the need for oxidatively introduced functional groups.

metal complexes directly to a carbon support without the need for surface pre-activation or post-immobilization treatment with excess precious metal precursors. While direct functionalization of graphene is possible using aryl diazonium-based precursors,^{25–28} the harsh conditions required by this reaction are typically not compatible with finely tuned catalytically-active metal complexes. Herein we describe a modular protocol for the direct hybridization of a pre-formed Rh(I) catalyst with the surface of unactivated carbon materials (Scheme 1). We demonstrate the potential widespread utility of this method by functionalizing two different high surface area carbon allotropes, graphene (G) and carbon black (CB) and show that these are highly efficient catalysts. Furthermore, it is demonstrated that even low surface area glassy carbon (GC) electrodes can be readily transformed into functionalized, recyclable catalytic systems using this efficient methodology.

Results and discussion

Here, we introduce our novel homogeneous catalyst, $[2\text{Rh}(\text{CO})_2]\text{BPh}_4$ (Fig. 1) onto the carbon supports by directly immobilizing the pre-synthesized Rh(I) complex containing an aniline-functionalized bidentate carbene-triazole ligand, $[1\text{Rh}(\text{COD})]\text{BPh}_4$ (COD = cyclooctadiene) via the *in situ* generation of the diazonium moiety (Scheme 2). The aniline functionalized complex $[1\text{Rh}(\text{COD})]\text{BPh}_4$, in an acidic solution of nitromethane and sodium nitrite was added to a GC electrode, or stirred with CB or G, for 24 h. The carbon materials were then washed repeatedly, and suspended in a mixture of dichloromethane and pentane under an atmosphere of CO. The GC electrode was washed thoroughly and dried under a gentle flow of nitrogen to give $[\text{GC}-2\text{Rh}(\text{CO})_2]$. $[\text{CB}-2\text{Rh}(\text{CO})_2]$ or $[\text{G}-2\text{Rh}(\text{CO})_2]$ were isolated as black powders after copious washing and drying under vacuum at 25 °C overnight.

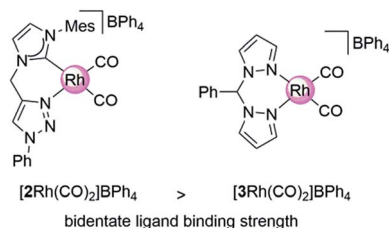
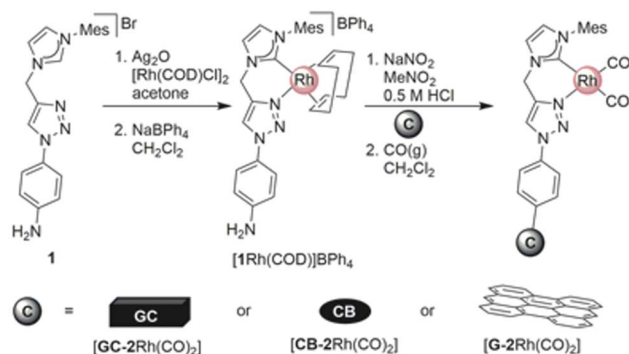


Fig. 1 Homogeneous Rh(I) complexes containing carbene-triazole (2) and bis-pyrazole (3) ligands.



Scheme 2 Synthesis of $[\text{GC}-2\text{Rh}(\text{CO})_2]$, $[\text{CB}-2\text{Rh}(\text{CO})_2]$ and $[\text{G}-2\text{Rh}(\text{CO})_2]$.

On application as a catalyst for hydroamination, our previously employed carbon supported bis-pyrazole system^{29,30} (Fig. 1) delivered remarkably high turnover numbers²⁹ but underwent some leaching of the Rh metal centre over time. The novel Rh(I) catalyst attached to the carbon surface here contains a carbene-triazole ligand system that has a high binding affinity for the Rh ion in order to create recyclable hybrid catalysts that exhibit minimal metal leaching.

While $[\text{GC}-2\text{Rh}(\text{CO})_2]$ was characterized using X-ray photoelectron spectroscopy (XPS), $[\text{CB}-2\text{Rh}(\text{CO})_2]$ and $[\text{G}-2\text{Rh}(\text{CO})_2]$ were characterized using XPS, thermogravimetric analysis (TGA), inductively coupled plasma-optical emission spectroscopy/mass spectrometry (ICP-OES/MS), Raman spectroscopy, scanning electron microscopy (SEM) and/or transmission electron microscopy (TEM).

In the XP spectra³¹ of $[\text{G}-2\text{Rh}(\text{CO})_2]$ (Fig. 2), the appearance of the N1s peak at *ca.* 402 eV³² and Rh3d_{5/2} peak at *ca.* 310 eV were observed. N1s and Rh3d peaks at similar binding energies were

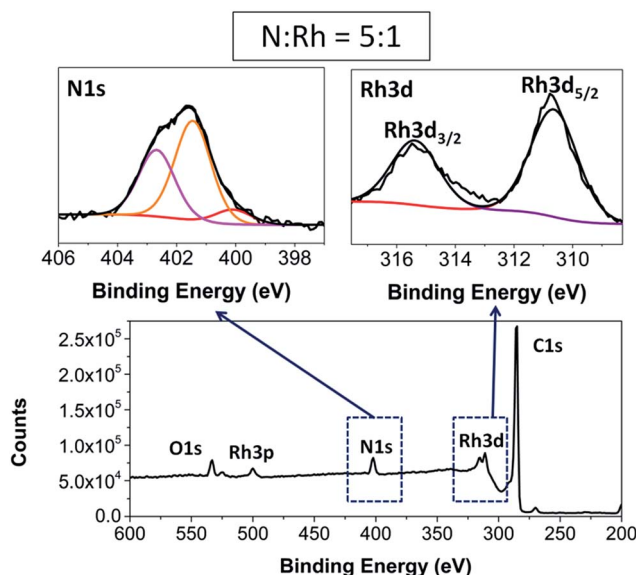


Fig. 2 XP survey scan of $[\text{G}-2\text{Rh}(\text{CO})_2]$ and selected narrow scan of N1s and Rh3d.

observed in the XP spectra of $[\text{GC-2Rh}(\text{CO})_2]$, $[\text{CB-2Rh}(\text{CO})_2]$ and the unattached complex $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ (see all XPS figures of hybrid complexes, unmodified carbon surfaces and homogeneous complex in the ESI, section SI-2†), consistent with analogous hybrid Rh(I) complexes on GC and CB reported previously.^{29,30} The atomic ratio of N : Rh for $[\text{GC-2Rh}(\text{CO})_2]$ and $[\text{G-2Rh}(\text{CO})_2]$ was found to be close to 5 : 1 as expected, which statistically means that only individual complexes are immobilized on the surfaces. In the case of $[\text{CB-2Rh}(\text{CO})_2]$, the ratio of N : Rh was found to be 3.8 : 1 suggesting that there may be a small amount of adventitious physisorbed Rh on the CB surface.

The thermal stability of $[\text{CB-2Rh}(\text{CO})_2]$ and $[\text{G-2Rh}(\text{CO})_2]$ was analyzed against that of unmodified CB and G using TGA under an atmosphere of nitrogen (Fig. 3). The thermogram of $[\text{CB-2Rh}(\text{CO})_2]$ displayed two weight loss processes at ca. 100–470 and 470–490 °C giving a total weight loss of 13% while the thermogram for $[\text{G-2Rh}(\text{CO})_2]$ displayed one weight loss process between ca. 350–650 °C with a total weight loss of 7% (see ESI, section SI-3†). Additionally, we observed that the thermogram of $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ only and a mixture of $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ and carbon black (see ESI, Fig. SI2†) showed a two weight loss trend from 100–380 °C due to the burn-off of the unattached metal complex similar to what was observed from the thermogram of $[\text{CB-2Rh}(\text{CO})_2]$.

The quantitative amount of Rh on CB and G obtained using ICP-OES was found to be 2.5 wt% for $[\text{CB-2Rh}(\text{CO})_2]$ and 0.84 wt% for $[\text{G-2Rh}(\text{CO})_2]$. This corresponds well to the estimated amount of Rh in $[\text{CB-2Rh}(\text{CO})_2]$ (2.6 wt%) and $[\text{G-2Rh}(\text{CO})_2]$ (1.3 wt%), calculated from the total weight loss obtained from TGA. For $[\text{GC-2Rh}(\text{CO})_2]$, cyclic voltammetry was used to determine the quantity of Rh complexes immobilized on the GC electrode to be $3.4 \times 10^{-10} \text{ mol cm}^{-2}$ or $1.2 \times 10^{-4} \text{ wt\% cm}^{-2}$ (see ESI, section SI-5†).

SEM and TEM images of $[\text{G-2Rh}(\text{CO})_2]$ showed that the integrity and morphology of the 2D structure is retained after treatment for immobilization of the Rh complex, and is similar to images of unmodified G (see ESI, section SI-8/9†). The Raman

spectra of $[\text{CB-2Rh}(\text{CO})_2]$ and $[\text{G-2Rh}(\text{CO})_2]$ showed a slight increase in the defects (D) band compared to the graphite (G) band following immobilization of the complex onto the surfaces (see ESI, section SI-7†), which indicates that both carbon–rhodium hybrids contained covalent linkages to the carbon black or graphene surface respectively, consistent with observations for similar covalent functionalization on carbon materials.^{33,34} An increase in the defects (D) band compared to the graphite (G) band of the carbon samples can be observed when comparing the peak intensity ratio of the D to G band of unmodified graphene, which was 0.79. After immobilization to form $[\text{G-2Rh}(\text{CO})_2]$, the peak intensity ratio of the D to G band increased to 0.82. Similarly, the peak intensity ratio of the D to G band for $[\text{CB-2Rh}(\text{CO})_2]$ was 1.0 which was higher when compared to the peak intensity ratio of D to G band of the unmodified carbon black of 0.92.

To summarize the surface characterization, we have demonstrated a straight-forward and effective method for the direct immobilization of Rh(I) complexes containing a mixed bidentate carbene-triazole ligand system onto carbon surfaces. The well-defined complexes were covalently anchored on GC, CB and G, showing that this method is versatile enough to be performed on various carbon surfaces with different surface areas and properties.

Catalysis

Following characterization of the hybrid catalysts, the efficiency of homogeneous complex $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ and hybrid complexes $[\text{GC-2Rh}(\text{CO})_2]$, $[\text{CB-2Rh}(\text{CO})_2]$ and $[\text{G-2Rh}(\text{CO})_2]$ were evaluated as catalysts for the hydrosilylation of diphenylacetylene (**4**) and triethylsilane at 50 °C and/or 25 °C using 0.02 mol% of Rh relative to the quantity of **4** (Scheme 3).³⁵ Transition metal-catalyzed hydrosilylation of alkynes allows the formation of alkenylsilanes in a straightforward, atom efficient and convenient method. Alkenylsilanes are versatile organosilicon compounds that can be used for further transformations in many organic synthesis reactions.

At 50 °C, $[\text{CB-2Rh}(\text{CO})_2]$ and $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ were both highly efficient catalysts, with the reaction reaching complete conversion of substrate **4** to product **5** in just 0.25 h (Fig. 4). While $[\text{G-2Rh}(\text{CO})_2]$ was slightly less efficient compared to $[\text{CB-2Rh}(\text{CO})_2]$ and $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$, the reaction still reached 56% conversion in 0.5 h and complete conversion in 1 h. The hydrosilylation reaction was repeated at 25 °C (to decrease the reaction rate to a more readily monitored rate) with $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ and $[\text{CB-2Rh}(\text{CO})_2]$ for better comparison. $[\text{CB-2Rh}(\text{CO})_2]$ and $[\text{2Rh}(\text{CO})_2]\text{BPh}_4$ each displayed very similar efficiency as catalysts for the reaction (Fig. 4). The reaction using

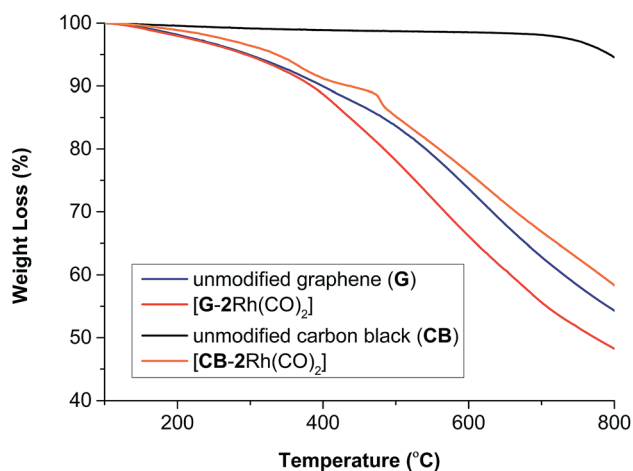
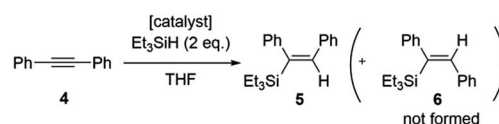


Fig. 3 TGA curves for unmodified CB and G, $[\text{G-2Rh}(\text{CO})_2]$ and $[\text{CB-2Rh}(\text{CO})_2]$ recorded under an atmosphere of nitrogen.



Scheme 3 Hydrosilylation reaction of diphenylacetylene **4** and triethylsilane.



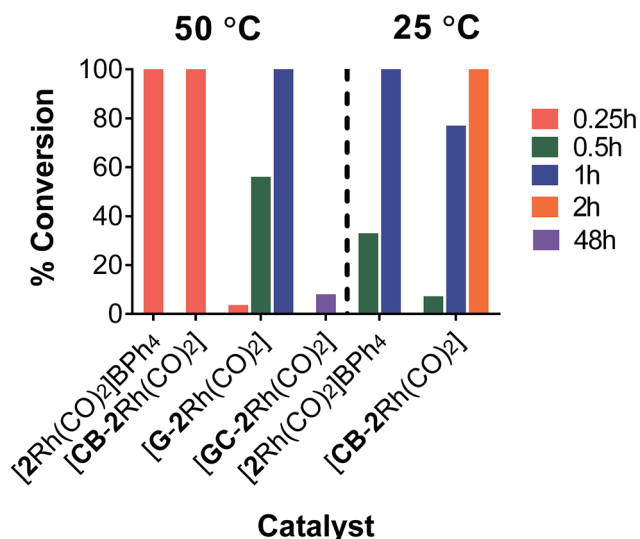


Fig. 4 Graph depicting the % conversion at times 0.25, 0.5, 1, 2 and 48 h for the hydrosilylation reaction using [2Rh(CO)₂]BPh₄, [CB-2Rh(CO)₂], [G-2Rh(CO)₂] and [GC-2Rh(CO)₂] at 50 °C or 25 °C.

[CB-2Rh(CO)₂] as catalyst was slightly slower, reaching 77% conversion to product 5 in 1 h compared to the complete conversion obtained using [2Rh(CO)₂]BPh₄ in the same time frame. This slightly lower efficiency of [CB-2Rh(CO)₂] could be due to the time required for diffusion of substrate to the carbon black support compared to the direct mixing of the two in the homogeneous reaction mixture. On completion, all reactions had turnover numbers (TON) close to 5000 (see ESI, Table S14[†]). These catalysts are the most efficient reported to date for the hydrosilylation of disubstituted alkyne, both homogeneous^{36–40} and heterogeneous.^{41–45}

On immobilizing the complex on a GC support ([GC-2Rh(CO)₂]), it was found that although the reaction was slow and showed only 8% conversion to product 5 in 48 h (Fig. 4), the TON for this reaction reached the very high value of 48 000 in that time. This correlates well with the fact that the GC electrode has a plate-like flat structure with a limited surface area (2.4 cm²) and hence holds only a small number of moles of immobilized catalyst relative to the substrate whereas there was a much higher mole ratio of catalyst to substrate immobilized on the high surface area CB and G. The high TON observed here for [GC-2Rh(CO)₂] is very similar to the turnover numbers we reported previously for GC supported catalysts.²⁹

In a series of control experiments at 50 °C, when using no catalyst or unmodified CB and G as the catalyst, the hydrosilylation reaction did not proceed even after 24 h (Table 1). In order to confirm that the carbon surfaces are covalently attached to the Rh(i) complexes, the homogeneous complex [2Rh(COD)]BPh₄, which lacks the aniline functional group that [1Rh(COD)]BPh₄ contains for the purpose of attachment, was mixed together with some unmodified G. The mixture was then treated using the same approach used to synthesize the hybrid catalysts, and the isolated G powder was tested for the hydrosilylation reaction. There was <0.08% conversion observed after 24 h, confirming that without the aniline functionalized group

Table 1 Control experiments for the hydrosilylation reaction

	Catalyst	Conv. ^a (%)	Time (h)
1	Unmodified CB only	0	24
2	Unmodified G only	0	24
3	G + [2Rh(COD)]BPh ₄ after synthesis treatment	0.08	24
4	No catalyst	0	24

^a Determined using ¹H NMR spectroscopy.

on the ligand, a covalent hybrid catalyst could not be assembled using the established synthesis.

Recyclability of hybrid catalysts

Initially, [GC-2Rh(CO)₂] was investigated for the hydrosilylation reaction over the course of three reaction cycles. After each run, the GC electrode was analyzed by XPS. The ratio of N : Rh on the surface of the GC remained constant at *ca.* 4.5 : 1 (see ESI, Table S15[†]), and this confirmed that the Rh complex remained on the GC surface even after multiple runs.

The low level of conversion to product 5 (<11%) obtained over an extended reaction time (48 h) when using [GC-2Rh(CO)₂] led to the investigation of the efficiency of hybrid catalysts on G and CB. In the recycling experiments for [CB-2Rh(CO)₂] (1.6 mol%) and [G-2Rh(CO)₂] (0.3 mol%), the hybrid catalysts were recycled over three 24 h cycles, and the filtrate and washings after each run were analyzed by ICP-MS to determine the Rh content. Both [G-2Rh(CO)₂] and [CB-2Rh(CO)₂] had excellent recycling ability (Fig. 5(a)), with complete conversion to product 5 after each run. The excellent recyclability was reinforced by ICP-MS analysis of the washings (Fig. 5(b)). When using [G-2Rh(CO)₂] as the catalyst, the ICP-MS showed hardly any leaching after each run, with less than 6% loss of Rh from the G surface after three runs. When using [CB-2Rh(CO)₂] as catalyst instead, an initial 12% loss of Rh from the CB surface was observed after the first cycle (Fig. 5(b)), consistent with the initial loss of the physisorbed Rh. This physisorbed Rh could not be removed prior to catalysis by normal washing procedures, and most likely requires the reaction conditions of the catalysis reaction to enable its' removal. After the second and third cycle, there was only a total of 3% loss of

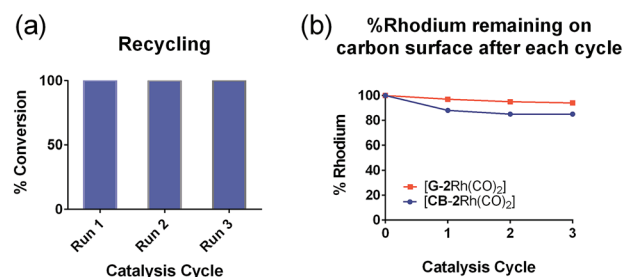


Fig. 5 Graphs depicting: (a) the % conversions of the hydrosilylation reaction for three reaction cycles when using [CB-2Rh(CO)₂] and [G-2Rh(CO)₂], which are shown on the same graph and (b) the % rhodium remaining on the carbon surfaces after each reaction cycle.

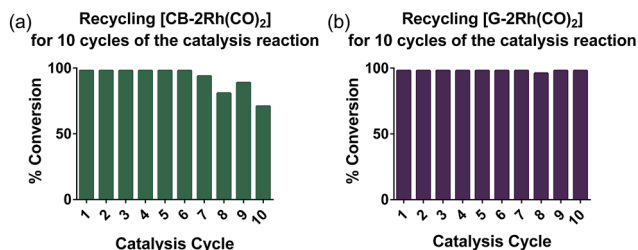


Fig. 6 Graphs depicting: (a) the % conversions of the hydrosilylation reaction for 10 reaction cycles when using [CB-2Rh(CO)₂] and (b) the % conversions of the hydrosilylation reaction for 10 reaction cycles when using [G-2Rh(CO)₂].

Rh observed. Again, such results confirm the stability of the immobilized Rh complex on the CB surface. These results also suggest that the use of the mixed bidentate carbene-triazole ligand system allows an enhanced capacity to recycle the Rh(I) complexes compared to our previously described hybrid system.^{29,30}

To further investigate the recyclability of the hybrid catalysts on G and CB, [CB-2Rh(CO)₂] and [G-2Rh(CO)₂] were reused over 10 cycles of the catalysis reaction. Each cycle was tested after 4 h at 25 °C (Fig. 6). [G-2Rh(CO)₂] was found to be an extremely recyclable heterogeneous catalyst and complete (>98%) conversions were obtained for all 10 cycles (Fig. 6(b)). When recycling [CB-2Rh(CO)₂], complete conversion (>98%) of the reaction was obtained for the first 7 cycles. Over the following 8–10 cycles, there was a slight 10–20% reduction in product formation (Fig. 6(a)), which suggests that [CB-2Rh(CO)₂] is slightly less recyclable compared to [G-2Rh(CO)₂]. The high recyclability of the hybrid catalysts shown here supports the benefit of immobilizing the highly efficient and well-defined homogeneous catalyst [2Rh(CO)₂]BPh₄ on the surface. [2Rh(CO)₂]BPh₄ is also shown to be a highly efficient homogeneous catalyst for the hydrosilylation of diphenylacetylene as it can catalyze up to 5 cycles of extra substrate addition before reaching a saturation point in the reaction mixture (see ESI, Fig. S22†). The development of the hybrid catalysts from the optimized homogeneous catalyst allows us to exploit the reusability of the homogeneous catalyst, providing us with highly recyclable hybrid catalyst that also allows for easy product separation.

Conclusions

In conclusion, during the investigations of a series of hybrid complexes on G, CB and GC as catalysts for the hydrosilylation of diphenylacetylene, it was found that [CB-2Rh(CO)₂] was the most efficient catalyst and [G-2Rh(CO)₂] was slightly less efficient than [CB-2Rh(CO)₂]. While [GC-2Rh(CO)₂] did not reach the same conversion levels over time, the very high TON achieved demonstrated excellent efficiency of this catalyst as well. The results also showed hybrid complex [CB-2Rh(CO)₂] was as efficient as the homogeneous complex [2Rh(CO)₂]BPh₄ in catalyzing this reaction. The catalysts investigated here could be utilized in very small amounts to give high TONs ranging from

5000 to 48 000. These catalysts are the most efficient of the homogeneous and heterogeneous catalysts reported to date for promoting the hydrosilylation of disubstituted alkyne. Finally, the recycling experiments also showed that the hybrid catalysts could be recycled for at least 10 times with hardly any loss of Rh from the surface. [G-2Rh(CO)₂] was found to be slightly more recyclable compared to [CB-2Rh(CO)₂]. All the results demonstrate the stability and reusability of the hydrid Rh(I) complexes containing the carbene-triazole mixed bidentate ligand system.

Experimental

General experimental and materials

Unless otherwise stated, all manipulations were carried out under inert atmosphere using standard Schlenk techniques or in a nitrogen or argon-filled Braun glovebox. All reagents were purchased from Aldrich Chemical Company Inc. or Alfa Aesar Inc. and used as received unless otherwise noted. Graphene⁴⁶ was contributed by Professor John Stride (UNSW). Glassy carbon electrodes (1 × 1 × 0.1 cm³) were purchased from Goodfellow. Carbon black (XC-72R) was supplied by Cabot Corporation. RhCl₃·xH₂O was purchased from Precious Metals Online PMO P/L. For the purposes of air sensitive manipulations and in the preparation of air sensitive metal complexes, tetrahydrofuran, dichloromethane, acetonitrile and pentane were dispensed from a PuraSolv solvent purification system. Methanol was distilled from magnesium methoxide under an atmosphere of nitrogen or argon. The bulk compressed gases argon (>99.999%) and carbon monoxide (>99.5%) were obtained from Air Liquide and used as received. Nitrogen gas for Schlenk line operation comes from in-house liquid nitrogen boil-off. 1-Azido-4-nitrobenzene,⁴⁷ [Rh(COD)Cl]₂,⁴⁸ 1-mesityl-3-(prop-2-yn-1-yl)-1H-imidazol-3-ium bromide⁴⁹ and 1-mesityl-3-((1-phenyl-1H-1,2,3-triazol-4-yl)methyl)-1H-imidazol-3-ium tetraphenylborate⁴⁹ were synthesized using literature procedures. ¹H and ¹³C{¹H} NMR spectra were recorded on Bruker DMX400, DMX500 and DMX600 spectrometers operating at 400, 500 and 600 MHz (¹H) respectively and 100, 125 and 150 MHz (¹³C) respectively. ¹H and ¹³C NMR chemical shifts were referenced internally to residual solvent resonances. Unless otherwise stated, spectra were recorded at 298 K and chemical shifts (δ), with uncertainties ±0.01 Hz for ¹H and ±0.05 Hz for ¹³C, are quoted in parts per million, ppm. Coupling constants (J) are quoted in Hz and have uncertainties of ±0.05 Hz for ¹H–¹H and ±0.5 Hz for ¹³C–¹⁰³Rh and ¹³C–¹¹B. Numbering and figures corresponding to NMR characterization data are located in the ESI, Table S1.† Deuterated solvents were purchased from Cambridge Stable Isotopes and used as received. Air sensitive NMR samples were prepared in an inert gas glovebox or by vacuum transfer of deuterated solvents into NMR tubes fitted with a Young's Teflon valve. For air sensitive NMR samples, dichloromethane-*d*₂ and chloroform-*d* were distilled over calcium hydride. Microanalyses were carried out at the Campbell Micro-analytical Laboratory, University of Otago, New Zealand or at the Research School of Chemistry, The Australian National University, Canberra, Australia. Mass spectra were acquired using a Thermo LTQ Orbitrap XL located in the



Bioanalytical Mass Spectrometry Facility (BMSF) in UNSW. *M* is defined as the molecular weight of the compound of interest or cationic fragment for cationic metal complexes.

1-Mesityl-3-((1-(4-nitrophenyl)-1*H*-1,2,3-triazol-4-yl)methyl)-1*H*-imidazol-3-ium bromide, nitro functionalized ligand precursor to ligand 1

1-Mesityl-3-(prop-2-yn-1-yl)-1*H*-imidazol-3-ium bromide (1020 mg, 3.33 mmol) was reacted with 1-azido-4-nitrobenzene (547 mg, 3.33 mmol) in a deoxygenated solution of isopropanol : water (2 : 1, 50 mL). After stirring for 5 minutes, sodium ascorbate (132 mg, 0.666 mmol, 20 mol%) followed by copper sulfate pentahydrate (41.3 mg, 0.165 mmol, 5 mol%) was added to the orange mixture. The reaction mixture was stirred for 24 hours to give an orange-red precipitate in an orange solution. The precipitate was collected *via* filtration and washed thoroughly with a saturated solution of Na₂EDTA until the filtrate turned from green to colourless. The precipitate was then dried under vacuum to give the ligand as a red solid. The ligand can be recrystallized from dichloromethane : pentane as red crystals (712 mg, 46%). ¹H NMR (400 MHz, CDCl₃): δ 10.31 (br s, 1H, Im-H2), 9.63 (s, 1H, Tz-H5'), 8.38 (d, ³*J* = 9.3 Hz, 2H, *m*-CH of ArNO₂), 8.13 (d, ³*J* = 9.3 Hz, 2H, *o*-CH of ArNO₂), 8.08 (t, ³*J* = 1.5 Hz, 1H, Im-H5), 7.14 (t, ³*J* = 1.5 Hz, 1H, Im-H4), 6.98 (s, 2H, *m*-CH of Mes), 6.22 (s, 2H, CH₂), 2.32 (s, 3H, *p*-CH₃ of Mes), 2.04 (s, 6H, *o*-CH₃ of Mes) ppm; ¹³C{¹H} NMR (100 MHz, CDCl₃): 147.5 (C_q of PhNO₂), 142.0 (Tz-C4'), 141.7 (*p*-CCH₃ of Mes), 140.9 (*ipso*-C to NH₂ of PhNH₂), 138.1 (Im-C2), 134.2 (*o*-CCH₃ of Mes), 130.6 (*ipso*-C of Mes), 130.0 (*m*-CH of Mes), 125.6 (*m*-CH of ArNO₂), 124.9 (Tz-C5'), 123.4 (Im-C5), 123.3 (Im-C4), 120.9 (*o*-CH of ArNO₂), 44.5 (CH₂), 21.2 (*p*-CH₃ of Mes), 17.8 (*o*-CH₃ of Mes) ppm; HRMS (ESI⁺, MeOH): *m/z* (%): calculated for [C₂₁H₂₃N₆O₂]⁺ = [M]⁺ = 389.1721; found [M]⁺ = 389.1718 (100%) amu; elemental analysis: found C, 51.85; H, 4.68; N, 17.21; calculated for C₂₁H₂₃BrN₆O₂·0.25CH₂Cl₂: C, 52.03; H, 4.42; N, 17.13%.

3-((1-(4-Aminophenyl)-1*H*-1,2,3-triazol-4-yl)methyl)-1-mesityl-1*H*-imidazol-3-ium bromide, (1)

The nitro functionalized ligand precursor above (605 mg, 1.29 mmol) was dissolved in methanol (30 mL). Pd/C (10% w/w, 54.4 mg, 0.0511 mmol, 4 mol%) was then added to the orange solution. After stirring for 5 minutes, hydrazine hydrate (1.3 mL, 26.8 mmol) was added slowly to the reaction mixture. The red-black mixture was heated at reflux under argon overnight. The mixture was filtered through Celite® which was washed thoroughly with excess methanol. The filtrate was dried with anhydrous magnesium sulfate, filtered and the solvent was evaporated to give the crude product as a yellow oil. The crude product was recrystallized from a dichloromethane : pentane mixture to yield ligand 1 as a fluffy pale yellow solid (526 mg, 93%). ¹H NMR (400 MHz, CDCl₃): δ 10.4 (br s, 1H, Im-H2), 8.93 (s, 1H, Tz-H5'), 8.03 (apparent t, ³*J* = 1.5 Hz, 1H, Im-H5), 7.50 (d, ³*J* = 8.8 Hz, 2H, *o*-CH of ArNH₂), 7.10 (apparent t, ³*J* = 1.5 Hz, 1H, Im-H4), 7.00 (s, 2H, *m*-CH of Mes), 6.74 (d, ³*J* = 8.5 Hz, 2H, *m*-CH of ArNH₂), 6.23 (s, 2H, CH₂), 3.93 (s, 2H, NH₂), 2.34 (s, 3H,

p-CH₃ of Mes), 2.05 (s, 6H, *o*-CH₃ of Mes) ppm; ¹³C{¹H} NMR (100 MHz, CDCl₃): δ 147.5 (C_q of ArNH₂), 141.7 (*p*-CCH₃ of Mes), 140.7 (Tz-C4'), 137.9 (Im-C2), 134.3 (*o*-CCH₃ of Mes), 130.7 (*ipso*-C of Mes), 130.04 (*m*-CH of Mes), 128.2 (*ipso*-C to NH₂ of ArNH₂), 123.8 (Tz-C5'), 123.3 (Im-C5), 122.94 (Im-C4), 122.3 (*o*-CH of ArNH₂), 115.4 (*m*-CH of ArNH₂), 44.7 (CH₂), 21.2 (*p*-CH₃ of Mes), 17.8 (*o*-CH₃ of Mes) ppm; HRMS (ESI⁺, MeOH): *m/z* (%): calculated for [C₂₁H₂₃N₆]⁺ = [M]⁺ = 359.1979; found [M]⁺ = 359.1974 (100%) amu; elemental analysis: found C, 57.70; H, 5.55; N, 19.03; calculated for C₂₁H₂₃BrN₆: C, 57.41; H, 5.28; N, 19.13%.

[1Rh(COD)]BPh₄

Ligand 1 (189 mg, 0.429 mmol) was dissolved partially as a suspension in acetone (25 mL). Ag₂O (87.6 mg, 0.378 mmol) was added to the suspension prior to heating the reaction mixture at reflux under argon for one hour. After cooling the reaction mixture to room temperature, [Rh(COD)Cl]₂ (106 mg, 0.214 mmol) was added. The reaction mixture was heated again at reflux for another one hour and then stirred at room temperature overnight under argon. The mixture was filtered through Celite® which was washed thoroughly with excess acetone. The solvent was removed *in vacuo* and the crude product was dissolved in dichloromethane (15 mL). NaBPh₄ (147 mg, 0.430 mmol) was added and white solid precipitates immediately. The reaction mixture was stirred vigorously at room temperature for 45 minutes to ensure complete reaction. The mixture was filtered through Celite® which was washed thoroughly with dichloromethane. The solution was concentrated *in vacuo* to ca. 2 mL which was added slowly dropwise to a vigorously stirred solution of pentane (30 mL) in an ice bath. Yellow precipitate forms immediately. After five minutes of stirring, the solvent was filtered and the precipitate dried under vacuum to give [1Rh(COD)]BPh₄ as a bright yellow solid (323 mg, 85%). ¹H NMR (400 MHz, CD₂Cl₂): δ 7.40 (m, 8H, *o*-CH of BPh₄), 7.24 (d, ³*J*_{H-H} = 8.9 Hz, 2H, *o*-CH of ArNH₂), 7.04 (m, 2H, *m*-CH of Mes overlapped with Tz-H5'), 7.04 (s, 1H, Tz-H5' overlapped with *m*-CH of Mes), 7.00 (t, ³*J*_{H-H} = 7.5 Hz, 8H, *m*-CH of BPh₄), 6.89 (d, ³*J*_{H4-H5} = 1.9 Hz, 1H, Im-H5), 6.85 (t, ³*J*_{H-H} = 7.1 Hz, 3H, *p*-CH of BPh₄), 6.75 (d, ³*J*_{H4-H5} = 1.9 Hz, 1H, Im-H4), 6.72 (d, ³*J*_{H-H} = 8.9 Hz, 2H, *m*-CH of ArNH₂), 4.92 (m, 2H, CH of COD), 4.76 (s, 2H, CH₂), 4.02 (s, 2H, NH₂), 3.64 (m, 2H, CH of COD), 2.37 (s, 3H, *p*-CH₃ of Mes), 2.25 (m, 2H, CH₂ of COD), 2.10 (s, 6H, *o*-CH₃ of Mes), 1.97 (m, 6H, CH₂ of COD) ppm; ¹³C{¹H} NMR (100 MHz, CD₂Cl₂): δ 176.2 (d, ¹*J*_{Rh-C} = 51.2 Hz, Im-C2), 164.4 (q, ¹*J*_{B-C} = 49.3 Hz, *o*-CH of BPh₄), 149.0 (C_q of ArNH₂), 140.1 (*p*-CCH₃ of Mes), 140.0 (C_q of Tz), 136.3 (br s, *o*-CH of BPh₄), 135.9 (*o*-CCH₃ of Mes), 135.5 (*ipso*-C of Mes), 129.5 (*m*-CH of Mes), 127.1 (*ipso*-C to NH₂ of ArNH₂), 126.2 (q, ⁴*J*_{B-H} = 2.6 Hz, *m*-CH of BPh₄), 123.1 (Im-C4), 122.6 (*o*-CH of ArNH₂), 122.5 (Im-C5), 122.3 (*p*-CH of BPh₄), 121.2 (Tz-C5'), 115.3 (*m*-CH of ArNH₂), 96.7 (d, ²*J*_{Rh-C} = 7.9 Hz, 2C, CH of COD), 79.1 (d, ²*J*_{Rh-C} = 12.4 Hz, 2C, CH of COD), 45.1 (CH₂), 32.6 (2C, CH₂ of COD), 29.2 (2C, CH₂ of COD), 21.2 (*p*-CH₃ of Mes), 18.6 (*o*-CH₃ of Mes) ppm; HRMS (ESI⁺, MeOH): *m/z* (%): calculated for [C₂₉H₃₄N₆Rh]⁺ = [M]⁺ = 569.1900; found [M]⁺ = 569.1893 (100%); calculated for [M - C₈H₁₁Rh]⁺ = 359.1979; found [M - C₈H₁₁Rh]⁺ = 359.1977



(78%) amu; elemental analysis: found C, 71.62; H, 6.09; N, 9.44; calculated for $C_{53}H_{54}BN_6Rh$: C, 71.62; H, 6.12; N, 9.46%.

[2Rh(COD)]BPh₄

Sodium ethoxide (14.7 mg, 0.225 mmol) was added to a stirred suspension of [Rh(COD)Cl]₂ (53.5 mg, 0.109 mmol) in ethanol (10 mL). The orange-yellow mixture was stirred at room temperature for 10 minutes. A solution of 1-mesityl-3-((1-phenyl-1*H*-1,2,3-triazol-4-yl)methyl)-1*H*-imidazol-3-ium tetraphenylborate ligand⁴ (144 mg, 0.217 mmol) in tetrahydrofuran (20 mL) was added slowly dropwise into the stirred suspension. The reaction mixture was stirred at room temperature for three hours. The solvent was removed *in vacuo* to give the crude product as a bright yellow solid. The crude product was dissolved in dichloromethane and filtered through Celite® which was washed thoroughly with excess dichloromethane. The solvent was concentrated to *ca.* 2 mL. Pentane (25 mL) was then added slowly and a bright yellow solid precipitates. The solvent was filtered off and the solid was dried under vacuum to yield [2Rh(COD)]BPh₄ as a bright yellow solid (126 mg, 64%). ¹H NMR (400 MHz, CDCl₃): δ 7.55 (m, 8H, *o*-CH of BPh₄), 7.48 (m, 3H, *o*- and *p*-CH of Ph), 7.38 (m, 2H, *m*-CH of Ph), 7.02 (s, 2H, *m*-CH of Mes overlapped with *m*-CH of BPh₄), 7.00 (t, ³*J*_{H-H} = 7.2 Hz, 8H, *m*-CH of BPh₄ overlapped with *m*-CH of Mes), 6.84 (t, ³*J*_{H-H} = 7.3 Hz, 4H, *p*-CH of BPh₄), 6.76 (d, ³*J*_{H4-H5} = 1.8 Hz, 1H, Im-H5), 6.65 (d, ³*J*_{H4-H5} = 1.8 Hz, 1H, Im-H4), 6.58 (s, 1H, Tz-H5'), 4.89 (m, 2H, CH of COD), 4.44 (s, 2H, CH₂), 3.63 (m, 2H, CH of COD), 2.38 (s, 3H, *o*-CH₃ of Mes), 2.25 (m, 2H, CH₂ of COD), 2.09 (s, 6H, *p*-CH₃ of Mes), 1.96 (m, 6H, CH₂ of COD) ppm; ¹³C{¹H} NMR (100 MHz, CDCl₃): δ 175.3 (d, ¹*J*_{Rh-C} = 52.7 Hz, Im-C2), 164.4 (q, ¹*J*_{B-C} = 49.6 Hz, *ipso*-C of BPh₄), 140.1 (C_q of Tz), 139.8 (*p*-CCH₃ of Mes), 136.3 (br s, *o*-CH of BPh₄), 135.8 (*o*-CCH₃ of Mes), 135.6 (*ipso*-C of Ph), 135.2 (*ipso*-C of Mes), 130.1 (*p*-CH of Ph), 130.0 (*o*-CH of Ph), 129.4 (*m*-CH of Mes), 126.0 (q, ⁴*J*_{B-C} = 2.8 Hz, *m*-CH of BPh₄), 122.8 (Im-C5), 122.6 (Im-C4), 122.1 (*p*-CH of BPh₄), 121.8 (Tz-C5'), 120.8 (*m*-CH of Ph), 96.3 (d, ²*J*_{Rh-C} = 7.8 Hz, 2C, CH of COD), 78.9 (d, ²*J*_{Rh-C} = 12.5 Hz, 2C, CH of COD), 44.6 (CH₂), 32.4 (2C, CH₂ of COD), 29.0 (2C, CH₂ of COD), 21.2 (*p*-CH₃ of Mes), 18.6 (*o*-CH₃ of Mes) ppm; HRMS (ESI⁺, MeOH): *m/z* (%): calculated for [C₂₉H₃₃N₅Rh]⁺ = [M]⁺ = 554.1786, found [M]⁺ = 554.1776 (100%) amu; elemental analysis: found C, 69.76; H, 6.19 and N, 7.80; calculated for C₅₃H₅₃BN₅Rh·0.5CH₂Cl₂: C, 70.13; H, 5.94 and N, 7.64%.

[2Rh(CO)₂]BPh₄

[2Rh(COD)]BPh₄ (117 mg, 0.133 mmol) was dissolved in dichloromethane (15 mL). The yellow solution was degassed *via* three cycles of freeze-pump-thaw. The reaction mixture was placed under an atmosphere of carbon monoxide gas in a balloon. The solution turned pale yellow in colour after stirring at room temperature for one hour. Pentane (45 mL) was added and yellow solid precipitates in solution with vigorous stirring. The solvent was filtered and the solid was washed with pentane (2 × 2 mL). The precipitate was dried *in vacuo* to yield [2Rh(CO)₂]BPh₄ as a pale yellow solid (75.6 mg, 70%). ¹H NMR (500 MHz, CD₂Cl₂): δ 7.59 (m, 3H, *o*- and *p*-CH of Ph), 7.53 (m,

m-CH of Ph), 7.44 (m, 8H, *o*-CH of BPh₄), 7.06 (s, 2H, *m*-CH of Mes), 6.98 (t, ³*J*_{H-H} = 7.5 Hz, 8H, *m*-CH of BPh₄ overlapped with Im-H4), 6.97 (d, ³*J*_{H4-H5} = 1.8 Hz, 1H, Im-H4 overlapped with *m*-CH of BPh₄), 6.93 (d, ³*J*_{H4-H5} = 1.8 Hz, 1H, Im-H5), 6.82 (t, ³*J*_{H-H} = 7.2 Hz, 4H, *p*-CH of BPh₄), 6.70 (s, 1H, Tz-H5'), 4.41 (s, 2H, CH₂), 2.38 (s, 3H, *p*-CH₃), 2.06 (s, 6H, *o*-CH₃) ppm; ¹³C{¹H} NMR (125 MHz, CD₂Cl₂): δ 184.7 (d, ¹*J*_{Rh-C} = 55.9 Hz, CO), 184.6 (d, ¹*J*_{Rh-C} = 70.6 Hz, CO), 172.1 (d, ¹*J*_{Rh-C} = 46.9 Hz, Im-C2), 164.5 (q, ¹*J*_{B-C} = 49.3 Hz, *ipso*-C of BPh₄), 141.2 (*p*-CCH₃ of Mes), 140.8 (C_q of Tz), 136.2 (br s, *o*-CH of BPh₄), 136.1 (*o*-CCH₃), 135.7 (*ipso*-C of Ph), 135.2 (*ipso*-C of Mes), 131.1 (*p*-CH of Ph), 130.5 (*o*-CH of Ph), 129.9 (*m*-CH of Mes), 126.4 (q, ⁴*J*_{B-C} = 2.6 Hz, *m*-CH of BPh₄), 124.1 (Im-C5), 123.4 (Im-C4), 122.5 (*p*-CH of BPh₄), 122.3 (Tz-C5'), 121.4 (*m*-CH of Ph), 44.6 (CH₂), 21.3 (*p*-CH₃ of Mes), 18.4 (*o*-CH₃ of Mes) ppm; HRMS (ESI⁺, MeOH): *m/z* (%): calculated for [C₂₃H₂₁N₅O₂Rh]⁺ = [M]⁺ = 502.0750; found [M]⁺ = 502.0703 (100%) amu; FTIR (solid) ν CO = 2083, 2019 (s) cm⁻¹; elemental analysis: found C, 66.42; H, 4.95; N, 8.49; calculated for C₄₇H₄₁BN₅O₂Rh·0.5CH₂Cl₂: C, 66.03; H, 4.90; N, 8.52%.

Immobilization of [CB-2Rh(CO)₂] and [G-2Rh(CO)₂]

The complex [1Rh(COD)]BPh₄ (24.8 mg, 0.0279 mmol) was dissolved in a degassed solution of nitromethane (10 mL). The mixture was cooled in an ice bath for 15 minutes before concentrated hydrochloric acid (0.5 mL, 0.5 M) was added dropwise into the stirred yellow solution while keeping the mixture between 0–5 °C. Sodium nitrite (1.90 mg, 0.0279 mmol) was then added and the mixture was stirred vigorously for another 5 minutes. After taking the mixture out of the ice bath, carbon black or graphene (50 mg) was added while the mixture was stirred vigorously to ensure that the carbon powders was diffused in solution properly. The mixture was allowed to gradually warm up to room temperature. The mixture was stirred under nitrogen at room temperature overnight. The modified carbon powder was separated from the reaction media *via* centrifugation (4000 rpm) for 5 minutes. It was consecutively washed and stirred vigorously with a methanol : MilliQ water (50 : 50, 20 mL) and methanol (5 × 10 mL) followed by centrifugation after each washing. The washed powder was dried under vacuum in a dessicator for 24 hours. The modified carbon powders was then dispersed with vigorous stirring in a dichloromethane : pentane solution (1 : 1, 20 mL). The mixture was degassed *via* three cycles of freeze-pump-thaw and then placed under an atmosphere of carbon monoxide in a balloon (1 atm). The mixture was left to react for one hour before the modified powder was separated from the reaction solvent *via* centrifugation. The modified powder was washed and stirred vigorously with pentane (3 × 10 mL) and methanol (3 × 10 mL) followed by centrifugation after each washing. The washed powder was then dried under vacuum in the dessicator for 24 hours to give either [CB-2Rh(CO)₂] (37.2 mg, 72%) and [G-2Rh(CO)₂] (32.0 mg, 68%).

Immobilization of [GC-2Rh(CO)₂]

The glassy carbon electrode was polished sequentially in alumina slurries (distilled water suspension) of 1.0, 0.3 and 0.05 μm on micropolishing cloth (Buehler, IL, USA) and rinsed



with distilled water between each step. It was then washed thoroughly with ethanol and soaked for 30 min in dichloromethane. The complex $[1\text{Rh}(\text{COD})]\text{BPh}_4$ (24.8 mg, 0.0279 mmol) was dissolved in a degassed solution of nitromethane (10 mL). The mixture was cooled in an ice bath for 5 minutes before concentrated hydrochloric acid (0.5 mL, 0.5 M) was added dropwise into the stirred yellow solution while keeping the mixture between 0–5 °C. Sodium nitrite (1.90 mg, 0.0279 mmol) was then added followed by the glassy carbon (GC) electrode. The mixture was left to sit in the ice bath for a further hour before allowing the mixture to gradually warm up to room temperature and remained at room temperature overnight. The modified GC electrodes was separated from the reaction media and washed with MilliQ water (5 mL). It was then consecutively washed with a methanol : MilliQ water (50 : 50, 20 mL) and methanol (5 × 10 mL). The washed modified GC electrode was dried under a gentle flow of nitrogen. The modified GC electrode was then dispersed in a dichloromethane : pentane solution (1 : 1, 20 mL). The mixture was degassed *via* three cycles of freeze–pump–thaw and then carbon monoxide was slowly bubbled into the mixture for 15 minutes. The modified GC electrode was washed with pentane (3 × 10 mL) and then methanol (3 × 10 mL). The washed powder was then dried under a gentle flow of nitrogen to give with $[\text{GC-2Rh}(\text{CO})_2]$.

Procedure for hydrosilylation reaction using the homogeneous catalyst, glassy carbon (GC), carbon black (CB) and graphene (G) hybrid catalysts

The reactions were performed in glass v-vials fitted with a magnetic stirring bar and a Teflon screw-caps. For the glassy carbon hybrid catalyst, the electrode (1 × 1 × 0.1 cm), diphenylacetylene (0.1 M), triethylsilane (0.2 M) and tetrahydrofuran (2 mL) were weighed into the glass vial in the glovebox. For the homogenous catalyst (0.3 mg), carbon black (0.8 mg) or graphene (0.5 mg) hybrid catalysts, the catalyst (0.02 mol% Rh relative to amount of diphenylacetylene) and diphenylacetylene (3.3 M) were weighed in air into the vial. The vials was then brought into the glovebox and triethylsilane (6.6 M) and tetrahydrofuran (0.06–0.6 mL) was added. The reactions vials were stirred and heated in an oil bath at 50 °C for 0.25–24 hours. At set time intervals until completion of the reaction, the reaction vial was removed from the oil bath and immediately cooled in an ice bath to room temperature.

For the reaction vial containing the glassy carbon hybrid catalyst, the reaction mixture was removed from the vial and solvent evaporated. The product was dissolved in CDCl_3 for NMR analysis. The reaction vial containing the hybrid catalysts on carbon black and graphene was centrifuged and the supernatant was collected into a round bottom flask. The solvent was evaporated and CDCl_3 was added to dissolve the products for NMR analysis. In all cases, the product conversions were calculated by integration of the chosen product peak in the ^1H NMR spectrum against the integration of the chosen starting material (diphenylacetylene) peak.

During the recycling experiments, the glassy carbon electrode catalyst was removed from the reaction mixture, washed

with methanol thoroughly (6 mL) and the electrode was dried under a gentle flow of nitrogen for 1 minute and reused subsequently. The carbon–rhodium catalysts on carbon black and graphene was washed with methanol (3 × 2 mL) and centrifuged to remove the solvent used in between each cycle. The carbon–rhodium catalyst was then dried under vacuum and reused in the next cycle. All supernatant and washings were collected and analyzed by ICP-MS.

Acknowledgements

Financial support from the University of New South Wales and the Australian Research Council are gratefully acknowledged. This research was supported under the Australian Research Council's Discovery Projects funding scheme (project numbers DP130101838 and DP150103065). We are grateful to Prof. John Stride for providing the graphene and Cabot Corporation for the donation of the carbon black (Vulcan XC-72R). We thank Dr Karen Privat of the Mark Wainwright Analytical Centre (UNSW) for collecting the SEM images and Mr Fei Han for collecting the TEM images. CMW thanks the Australian Government for the award of an International Postgraduate Research Scholarship (IPRS).

Notes and references

- 1 *Heterogenized Homogeneous Catalysts for Fine Chemicals Production*, ed. P. Barbaro and F. Liguori, Springer, Dordrecht, The Netherlands, 2010.
- 2 D. J. Cole-Hamilton, *Science*, 2003, **299**, 1702–1706.
- 3 C. W. Jones, *Top. Catal.*, 2010, **53**, 942–952.
- 4 A. Schaetz, M. Zeltner and W. J. Stark, *ACS Catal.*, 2012, **2**, 1267–1284.
- 5 C. Freire, C. Pereira and S. Rebelo, *Catalysis*, 2012, **24**, 116–203.
- 6 D. T. Genna, A. G. Wong-Foy, A. J. Matzger and M. S. Sanford, *J. Am. Chem. Soc.*, 2013, **135**, 10586–10589.
- 7 M. Yoon, R. Srirambalaji and K. Kim, *Chem. Rev.*, 2012, **112**, 1196–1231.
- 8 D. Gajan and C. Coperet, *New J. Chem.*, 2011, **35**, 2403–2408.
- 9 U. Diaz, D. Brunel and A. Corma, *Chem. Soc. Rev.*, 2013, **42**, 4083–4097.
- 10 Y. Zhang and S. N. Riduan, *Chem. Soc. Rev.*, 2012, **41**, 2083–2094.
- 11 *Carbon Materials For Catalysis*, ed. P. Serp and J. L. Figueiredo, 2009.
- 12 E. Auer, A. Freund, J. Pietsch and T. Tacke, *Appl. Catal.*, 1998, **173**, 259–271.
- 13 F. Rascon, R. Wischert and C. Coperet, *Chem. Sci.*, 2011, **2**, 1449–1456.
- 14 M. C. Kung, M. V. Riofski, M. N. Missaghi and H. H. Kung, *Chem. Commun.*, 2014, **50**, 3262–3276.
- 15 S. Kumar, P. Kumar and S. L. Jain, *J. Mater. Chem. A*, 2014, **2**, 18861–18866.
- 16 Q. Zhao, Y. Zhu, Z. Sun, Y. Li, G. Zhang, F. Zhang and X. Fan, *J. Mater. Chem. A*, 2015, **3**, 2609–2616.



- 17 H. Su, Z. Li, Q. Huo, J. Guan and Q. Kan, *RSC Adv.*, 2014, **4**, 9990–9996.
- 18 Y.-T. Xi, P.-J. Wei, R.-C. Wang and J.-G. Liu, *Chem. Commun.*, 2015, **51**, 7455–7458.
- 19 M. Blanco, P. Alvarez, C. Blanco, M. V. Jimenez, J. Fernandez-Tornos, J. J. Perez-Torrente, L. A. Oro and R. Menendez, *Carbon*, 2015, **83**, 21–31.
- 20 N. Shang, S. Gao, C. Feng, H. Zhang, C. Wang and Z. Wang, *RSC Adv.*, 2013, **3**, 21863–21868.
- 21 J. K. Wassei and R. B. Kaner, *Acc. Chem. Res.*, 2013, **46**, 2244–2253.
- 22 D. K. James and J. M. Tour, *Acc. Chem. Res.*, 2013, **46**, 2307–2318.
- 23 B. F. Machado and P. Serp, *Catal. Sci. Technol.*, 2012, **2**, 54–75.
- 24 C. N. R. Rao, A. K. Sood, K. S. Subrahmanyam and A. Govindaraj, *Angew. Chem., Int. Ed.*, 2009, **48**, 7752–7777.
- 25 G. L. C. Paulus, Q. H. Wang and M. S. Strano, *Acc. Chem. Res.*, 2013, **46**, 160–170.
- 26 P. Huang, L. Jing, H. Zhu and X. Gao, *Acc. Chem. Res.*, 2013, **46**, 43–52.
- 27 C. Peng, Y. Xiong, Z. Liu, F. Zhang, E. Ou, J. Qian, Y. Xiong and W. Xu, *Appl. Surf. Sci.*, 2013, **280**, 914–919.
- 28 E. Bekyarova, M. E. Itkis, P. Ramesh, C. Berger, M. Sprinkle, W. A. de Heer and R. C. Haddon, *J. Am. Chem. Soc.*, 2009, **131**, 1336–1337.
- 29 A. A. Tregubov, K. Q. Vuong, E. Luais, J. J. Gooding and B. A. Messerle, *J. Am. Chem. Soc.*, 2013, **135**, 16429–16437.
- 30 A. A. Tregubov, D. B. Walker, K. Q. Vuong, J. J. Gooding and B. A. Messerle, *Dalton Trans.*, 2015, **44**, 7917–7926.
- 31 The nature of the counter-ions for the positively charged immobilized complexes could not be established unambiguously by XPS, which was consistent with previous reports: D. R. Laws, J. Sheats, A. L. Rheingold and W. E. Geiger, *Langmuir*, 2010, **26**, 15010–15021.
- 32 We also observed in all hybrid cases, a very minor peak within the N1s peak at about 399–400 eV indicative of the azo bond (N=N) formation due to some incomplete diazonium reaction and this is consistent with previous reports: A. L. Gui, E. Luais, J. R. Peterson and J. J. Gooding, *ACS Appl. Mater. Interfaces*, 2013, **5**, 4827–4835.
- 33 J. M. Englert, C. Dotzer, G. Yang, M. Schmid, C. Papp, J. M. Gottfried, H.-P. Steinruck, E. Spiecker, F. Hauke and A. Hirsch, *Nat. Chem.*, 2011, **3**, 279–286.
- 34 A. C. Ferrari and J. Robertson, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2000, **61**, 14095–14107.
- 35 In all cases, the *E* isomer (5) of the alkene product was formed in >99 : 1 ratio to the *Z* isomer (6).
- 36 J. Berding, J. A. van Paridon, V. H. S. van Rixel and E. Bouwman, *Eur. J. Inorg. Chem.*, 2011, 2450–2458.
- 37 M. Blug, X.-F. Le Goff, N. Mezailles and P. Le Floch, *Organometallics*, 2009, **28**, 2360–2362.
- 38 L. Yong, K. Kirleis and H. Butenschoen, *Adv. Synth. Catal.*, 2006, **348**, 833–836.
- 39 M. R. Chaulagain, G. M. Mahandru and J. Montgomery, *Tetrahedron*, 2006, **62**, 7560–7566.
- 40 N. Imlinger, K. Wurst and M. R. Buchmeiser, *J. Organomet. Chem.*, 2005, **690**, 4433–4440.
- 41 W. Guo, R. Pleixats, A. Shafir and T. Parella, *Adv. Synth. Catal.*, 2015, **357**, 89–99.
- 42 M. Planellas, W. Guo, F. Alonso, M. Yus, A. Shafir, R. Pleixats and T. Parella, *Adv. Synth. Catal.*, 2014, **356**, 179–188.
- 43 R. Cano, M. Yus and D. J. Ramon, *ACS Catal.*, 2012, **2**, 1070–1078.
- 44 F. Alonso, R. Buitrago, Y. Moglie, J. Ruiz-Martinez, A. Sepulveda-Escribano and M. Yus, *J. Organomet. Chem.*, 2010, **696**, 368–372.
- 45 M. Chauhan, B. J. Hauck, L. P. Keller and P. Boudjouk, *J. Organomet. Chem.*, 2002, **645**, 1–13.
- 46 M. Choucair, P. Thordarson and J. A. Stride, *Nat. Nanotechnol.*, 2009, **4**, 30–33.
- 47 S. W. Kwok, J. R. Fotsing, R. J. Fraser, V. O. Rodionov and V. V. Fokin, *Org. Lett.*, 2010, **12**, 4217–4219.
- 48 G. Giordano and R. H. Crabtree, *Inorg. Synth.*, 1990, **28**, 88–90.
- 49 K. Q. Vuong, M. G. Timerbulatova, M. B. Peterson, M. Bhadbhade and B. A. Messerle, *Dalton Trans.*, 2013, **42**, 14298–14308.

