Chemical Science

www.rsc.org/chemicalscience



ISSN 2041-6539



EDGE ARTICLE Hayato Tsurugi, Kazushi Mashima *et al.* 2,2'-Bipyridyl formation from 2-arylpyridines through bimetallic diyttrium intermediate

Chemical Science

EDGE ARTICLE



Cite this: Chem. Sci., 2015, 6, 5394

2,2'-Bipyridyl formation from 2-arylpyridines through bimetallic diyttrium intermediate†

Yu Shibata, Haruki Nagae, Shiki Sumiya, Raphaël Rochat, Hayato Tsurugi* and Kazushi Mashima*

An alkylyttrium complex supported by an N,N'-bis(2,6-diisopropylphenyl)ethylenediamido ligand, (ArNCH₂CH₂NAr)Y(CH₂SiMe₃)(THF)₂ (**1**, Ar = 2,6-ⁱPr₂C₆H₃), activated an *ortho*-phenyl C–H bond of 2phenylpyridine (**2a**) to form a (2-pyridylphenyl)yttrium complex (**3a**) containing a five-membered metallacycle. Subsequently, a unique $C(sp^2)-C(sp^2)$ coupling of 2-phenylpyridine proceeded through a bimetallic yttrium intermediate, derived from an intramolecular shift of the yttrium center to an *ortho*position of the pyridine ring in **3a**, to yield a bimetallic yttrium complex (**4a**) bridged by two-electron reduced 6,6'-diphenyl-2,2'-bipyridyl. Aryl substituents at the *ortho*-position of the pyridine ring were key in order to destabilize the μ,κ^2 -(C,N)-pyridyldiyttrium intermediate prior to the C(sp²)–C(sp²) bond formation.

Received 2nd May 2015 Accepted 17th July 2015

DOI: 10.1039/c5sc01599e

www.rsc.org/chemicalscience

Introduction

Transition metal-catalyzed homo-coupling reactions of two arenes are an important category of C-C bond forming reactions to construct π -conjugated biaryl skeletons.^{1,2} For example, in the Ullmann coupling reaction, the activation of aryl C-X bonds of heteroarenes with a low-valent metal species, such as Cu powder or Ni(cod)₂, produces the corresponding biaryl compounds. Although these reductive homo-coupling reactions are frequently used, the formation of salt waste is inevitable, and thus there is a demand for a more atom-economical method to synthesize biaryl skeletons. The most direct protocol for $C(sp^2)-C(sp^2)$ bond formation is through C-H bond activation of aromatic compounds. To date, various transition metal complexes have been applied to such dehydrogenative biaryl coupling reactions.3 The most well-established mechanism for biaryl $C(sp^2)$ - $C(sp^2)$ bond formation mediated by a mononuclear species is reductive elimination of mononuclear di(aryl) metal species. However, some monoarylated metal species undergo biaryl $C(sp^2)$ - $C(sp^2)$ coupling reactions. In this reaction, two mechanisms are proposed to be involved, *i.e.* disproportionation to produce a di(aryl)metal and low-valent metal species,⁴ and an associative C-C bond formation mediated by two metal centers.^{5,6} In the associative mechanism, a bridged dimer species A is first formed through π -coordination of the aryl moiety to another metal center, followed by the formation of species **B**, which contains a 3-centered-2-electron bridging aryl moiety (Fig. 1). Subsequent C-C bond formation from species B produces the corresponding biaryl compound. A closely related reaction is the Glaser divne coupling reaction of terminal alkynes using a Cu catalyst, the mechanism of which involves a stepwise process through π -coordination of the C=C bond to a different metal center and a 3-centered-2-electron C(sp)-bridging dinuclear intermediate before the C-C bondforming step.⁷ Synthesis of the C=C π -coordination-bridged multimetallic species and mechanistic studies of the bimetallic aggregation-assisted C(sp)-C(sp) bond formation are feasible due to the strong coordinating ability of the alkyne moiety to the metal center; however, corresponding studies of arylmetal species and the mechanism of associative biaryl $C(sp^2)-C(sp^2)$ bond formation have not been established due to the weaker

View Article Online

View Journal | View Issue



Fig. 1 Aggregated monoarylated and monoalkynylated metal species. A: π -coordination-bridged bimetallic species. B: 3-centered-2-electron Ph-bridged bimetallic species. C: dianionic biphenyl-bridged bimetallic lanthanide complex. D: dianionic diyne-bridged bimetallic lanthanide complex.

Department of Chemistry, Graduate School of Engineering Science, Osaka University, Toyonaka, Osaka 560-8531, Japan. E-mail: mashima@chem.es.osaka-u.ac.jp; tsurugi@chem.es.osaka-u.ac.jp

[†] Electronic supplementary information (ESI) available: Experimental details for the synthesis and characterization of Y complexes, ¹H NMR spectrum of the deuterium labelling experiment, and crystal data for **3e** (CCDC 1409167), **4a** (CCDC 1048580), **4b** (CCDC 1048581), **4c** (CCDC 1048582), **6a** (CCDC 1048583), **6c** (CCDC 1048584) and **6d** (CCDC 1048585). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c5sc01599e

 π -aromatic coordination to the metal center compared with C=C π -coordination.

Rare-earth metal complexes containing a mono(aryl)- or mono(alkynyl)metal moiety, generated by C-H bond activation of arenes and terminal alkynes using an alkylmetal species, also mediate $C(sp^2)$ - $C(sp^2)$ and C(sp)-C(sp) bond formation through the aggregation of two metal species. Because of the stability of the +3 oxidation state of the rare-earth metal center, C-C bond formation products, biaryls and diynes, have been trapped in their dianionic form to give bimetallic complexes such as C and **D** as reported by Fryzuk *et al.* (Fig. 1),^{5,8,9} even though access to low-valent rare-earth metal complexes has been reported by Evans et al.¹⁰ In our studies on the C-H bond activation of heteroaromatic compounds by rare-earth metal and early transition metal complexes,¹¹ we found that σ -bond metathesis and subsequent 2,2'-bipyridyl formation from 2-arylpyridines proceeded upon treatment of an alkyl complex of (ethylenediamido)yttrium (1) with 2-arylpyridine (2). During this transformation, the C-H bond adjacent to the nitrogen atom of the pyridine ring was selectively functionalized. Mononuclear (2-pyridylphenyl)yttrium complex 3 was detected and isolated as an intermediate in the formation of dianionic 2,2'-bipyridylbridged dinuclear yttrium complex 4 (Scheme 1). This is the first example of 2,2'-bipyridyl formation through bimetallic aggregation, even though dimerization of pyridine via C-H bond activation and insertion reactions has previously been reported by Teuben and Diaconescu, respectively.12 Catalytic 2,2'-bipyridyl formation via C-H bond activation has only been achieved using heterogeneous Pd/C and Ru cluster catalysts.13 In addition, steric and electronic tuning of the pyridine derivatives led to the isolation of dinuclear bis(μ,κ^2 -(C,N)-pyridyl)diyttrium, mononuclear κ^2 -(C,N)-pyridylyttrium, and 5-membered metallacycle complexes as possible intermediates in the C-C bond formation.

Results and discussion

We first treated an alkylyttrium complex (ArNCH₂CH₂NAr) $Y(CH_2SiMe_3)(THF)_2$ (1, Ar = 2,6-^{*i*}Pr₂C₆H₃)¹⁴ with 1 equiv. of 2-phenylpyridine (2a) in benzene at room temperature. The color of the solution changed immediately from pale yellow to orange and then to dark green, and green-colored crystals of 4a were precipitated (eqn (1)). The green crystals were sparingly soluble in aromatic and aliphatic solvents. The molecular structure of 4a was determined by X-ray diffraction studies, and its ORTEP drawing is shown in Fig. 2. During the reaction, 6,6'-diphenyl-



Scheme 1 Pyridylyttrium-mediated 2,2'-bipyridyl formation. $[Y] = (ArNCH_2CH_2NAr)Y.$

2,2'-bipyridyl was formed as a biaryl coupling product of 2phenylpyridine. Two (ethylenediamido)yttrium moieties are bridged by the two-electron reduced 6,6'-diphenyl-2,2'-bipyridyl ligand. The nitrogen atoms in the 2,2'-bipyridyl moiety are located on opposite sides, and the 2,2'-bipyridyl ligand coordinates to the two yttrium atoms in a μ – η^4 : η^4 -coodination mode. The bond length of Y1–N1 (2.344 Å) is shorter than the typical yttrium–nitrogen dative bond (*ca.* 2.5 Å),¹⁵ but longer than the Y1–N2 and Y1–N3 bonds (*ca.* 2.19 Å). The Y1–C2* bond (2.661 Å) is much longer than the yttrium–carbon covalent bond (*ca.* 2.45 Å),^{55,15c} The C1–C1* bond (1.396 Å) of the central 2,2'-bipyridyl moiety is similar in length to the analogous bond in two-electron reduced 2,2'-bipyridyl bound to two alkali metal centers with alternate planes (1.400 Å, rubidium).¹⁶



In addition to 2-phenylpyridine, 2-arylpyridines with methyl and methoxy groups at the *para*-position of the phenyl group were applicable to the 2,2'-bipyridyl formation. Complex **4b** was isolated in quantitative yield after treatment of **1** with 1 equiv. of 2-(4-methylphenyl)pyridine (**2b**) at room temperature for 48 h. When 2-(4-methoxyphenyl)pyridine (**2c**) was used as the substrate, heating the reaction mixture at 50 °C for 68 h led to the formation of green crystals of **4c** in 36% yield. The low isolated yield of **4c** was attributed to decomposition during the C–C bond forming process. Because complexes **4b** and **4c** had



Fig. 2 Molecular structure of complex 4a with 30% thermal ellipsoids. All hydrogen atoms and isopropyl groups are omitted for clarity. Selected bond lengths (Å) and angles (°): Y1–N1, 2.344(5); Y1–C1, 2.746(5); Y1–C1*, 2.857(5); Y1–C2*, 2.661(5); N1–C1, 1.424(6); C1–C2, 1.476(7); C1–C1*, 1.396(10); Y1–N2, 2.194(4); Y1–N3, 2.192(5); N1–Y1–C2*, 70.68(16); N2–Y1–N3, 80.78(17). Dihedral angle between N1–Y1–C2* and N1–C1–C1*–C2* planes, 109.6.

poor solubility in aliphatic and aromatic solvents and low stability in coordinating solvents, characterization of **4b** and **4c** was based only on X-ray diffraction studies and combustion analyses.¹⁷

We next conducted a deuterium labelling experiment. The addition of C_6D_5 derivative $2a \cdot d_5$ to complex 1 in C_6H_6 resulted in the formation of the same green crystals together with a mixture of SiMe₄ and SiMe₄- d_1 , the former indicating direct C-H bond activation at the *ortho*-position of the pyridine ring by the alkylyttrium moiety, and the latter indicating C-D bond activation of the *ortho*- C_6D_5 position by Y–CH₂SiMe₃ followed by an intramolecular shift of the yttrium atom to the *ortho*-position of the pyridyl before the C–C bond forming process (*vide infra*). These processes are consistent with subsequent oxidative quenching of the crystalline compound by CCl₄ to give a mixture of d_{8^-} , d_{9^-} , and d_{10} -6,6'-diphenyl-2,2'-bipyridyl, as evidenced by the intensity (69%-H) of the singlet signal corresponding to the *ortho*-position of the phenyl ring at δ_H 8.18 (Scheme 2).

In addition, when **1** was reacted with **2a** at room temperature in THF for 9 days to allow complete C–H bond activation, 5membered metallacyclic complex **3a** was isolated in 98% yield. Complex **3a** was stable and no further coupling reaction was detected in THF, but dissolution of **3a** in C_6D_6 afforded the bimetallic compound **4a** quantitatively. This clearly indicated that the 5-membered metallacyclic complex **3a** is a metastable species in benzene that could lead to a subsequent intramolecular shift of the yttrium center to the *ortho*-position of the pyridine ring, followed by C–C bond formation to afford **4a** (Scheme 3).

When 2-arylpyridines 2c and 2d were treated with 1 in C_6D_6 at room temperature, C–H bond activation of the aryl ring proceeded to form five-membered metallacyclic complexes 3cand 3d (Scheme 4). On heating the C_6D_6 solution of 3c at $50 \,^{\circ}C$, green microcrystals were precipitated from the reaction mixture, as determined by eqn (1). In contrast, 3d was stable in C_6D_6 at $50 \,^{\circ}C$. We thus presumed that the C–C bond formation was affected by electron-donating/-withdrawing substituents *meta* to the metallated carbon; THF coordination to yttrium for 3a and 3b was weaker than that for 3c and 3d in benzene, leading to easy dissociation of THF from yttrium for 3a and 3band C–C bond formation to form 4a and 4b at room temperature.

CCl₄

D/H

D/H

D/H = 31/69

Scheme 2 Deuterium labelling experiment.

2a-d₅ Ď

C₆H₆, rt

SiMe₄/SiMe₄-d₁



Scheme 3 Stepwise metallacycle and C-C bond formation



Scheme 4 Effect of substituents on the aryl ring of 2-arylpyridine on the C–C bond formation step.

In addition to the isolation of stable five-membered metallacyclic complex 3d, the reaction of 1 with benzo[h]quinolone atroom temperature gave <math>3e in quantitative yield (eqn (2)). Complex 3e was isolated as microcrystals suitable for X-ray diffraction studies. Although the quality of the crystallographic data from the X-ray diffraction studies was insufficient, we determined the overall structure of 3e, in which a C-H bond of benzo[h]quinolone was activated to form a five-membered metallacycle as shown in the ESI (Fig. S1†). Complex 3e was not converted to the C-C bond formation product analogous to 4ac, probably due to the low flexibility of the benzo[h]quinolonescaffold.



In sharp contrast to the reaction of **1** with 2-arylpyridines **2a**-**2c**, treatment of **1** with 1 equiv. of pyridine (**5a**) or 4- or 3substituted pyridines (**5b-d**) afforded bis(μ , κ^2 -(C,N)-pyridyl) diyttrium complexes **6a-d** as poorly soluble yellow microcrystals (Scheme 5). The ORTEP drawing of **6a** is shown in Fig. 3. The μ , κ^2 -(C,N)-pyridyl ligand is positioned at the bridging part of the bimetallic structure. The bond lengths of Y–N1 (2.328(4) Å) and Y*–C1 (2.560(5) Å) are longer than those found for mononuclear κ (C,N)-pyridylyttrium complexes.^{17,18} Two yttrium atoms, two bridging carbons, and the two nitrogen atoms of the bridging

This article is licensed under a Creative Commons Attribution 3.0 Unported Licence.

THF

SiMea

 $(Ar = 2, 6^{-i} Pr_2 C_6 H_3)$

THE



Scheme 5 Reactions of alkylyttrium complex 1 with pyridine derivatives.

pyridines are located in the same plane. The ¹H NMR spectrum of **6a** displays four resonances corresponding to the bridging pyridine ring at $\delta_{\rm H}$ 9.18 (3-py), 8.81 (6-py), 7.46 (4-py), and 6.80 (5-py). A significant downfield shift of the resonance at the 3-py position might be due to the proximity of the C-H bond to the metal fragment. When 3,5-dimethylpyridine (**5e**) was used as the substrate, mononuclear yttrium complex **7e** was isolated in 97% yield. In the ¹³C NMR spectrum, a doublet signal was observed for the carbon atom attached to the yttrium center at $\delta_{\rm C}$ 219.6 (¹ $J_{\rm Y-C} = 35.2$ Hz), which is in the typical range for mononuclear arylyttrium complexes.¹⁸ Even after heating solutions of complexes **6a–d** and **7e**, which contained a 2-pyridylyttrium moiety in the molecular structure, C–C coupling products were not detected in the reaction mixture; decomposition of the complexes was observed, and

no single species was isolated from the reaction mixture. 2-Trimethylsilylpyridine (**5f**) was also reacted with yttrium complex **1** to form (dimethylpyridylsilyl)methylyttrium complex **8f** *via* $C(sp^3)$ -H bond activation. In this case, an intramolecular shift of the yttrium center to form 2-pyridylyttrium species or 6,6'-bis(trimethylsilyl)-2,2'-bipyridyl formation was not observed.

Based on our findings for the alkylyttrium-mediated C-H bond activation and C-C coupling reaction, we propose a mechanism for 2,2'-bipyridyl formation as shown in Scheme 6. First, alkylyttrium complex 1 cleaves a C-H bond at the orthoposition on the phenyl ring of 2a to produce five-membered metallacycle complex 3a. Complex 3a is isomerized to threemembered metallacycle intermediate E. Initial formation of the five-membered metallacycle prior to formation of the three-membered metallacycle, as the major pathway, was confirmed by the deuterium labelling experiment as shown in Scheme 2, where mono-deuterated $SiMe_4$ - d_1 was generated and one H atom was incorporated into the phenyl ring. Direct formation of the intermediate E from complex 1 was plausible as a minor pathway, and this was confirmed by the detection of SiMe₄ in the deuterium labelling experiment. Isomerization between three- and five-membered metallacycles was similarly reported by Diaconescu et al. for rare-earth metal complexes. Although the isomerization trend is opposite to the report by Diaconescu et al., they mentioned that the pyridyl carbanion is more stable $(2.8 \text{ kcal mol}^{-1})$ than the phenyl carbanion for the phenylpyridyl anion. We presume that the relative stabilities of the three- and five-membered metallacycles are significantly affected by the attached metal fragment.19 The effect of substituents of the 2-arylpyridines on the C-C bond formation as shown in Scheme 4 indicates that the dissociation of the coordinating THF from the yttrium is key for further isomerization. Although the three-membered metallacycle intermediate E dimerized as a doubly μ,κ^2 -(C,N)-bridged dinuclear structure, similar to divttrium complexes 6a-d, introduction of aryl groups at the ortho-position of the pyridine ring might destabilize the μ,κ -(C,N)-bridging mode of the pyridine moiety to afford 4a through 3-centered-2-electron aryl-bridged intermediate F.



Fig. 3 Molecular structure of complex **6a** with 30% thermal ellipsoids. All hydrogen atoms are omitted for clarity. Selected bond lengths (Å): Y1–N1, 2.328(4); Y1–C1, 2.681(5); Y1–C1*, 2.560(5); Y1–N2, 2.216(4); Y1–N3, 2.219(4); Y1–O1, 2.422(3).



Scheme 6 Plausible mechanism for the reductive dimerization of 2-phenylpyridine.

Conclusions

We demonstrated that yttrium-mediated 2,2'-bipyridyl formation proceeded through a bimetallic pyridylyttrium intermediate. Introduction of aryl substituents at the *ortho*-position of the pyridine ring destabilized the μ,κ^2 -(C,N)-bridged intermediate to accelerate associative bimetallic C(sp²)-C(sp²) bond formation. Further application of such bimetallic-mediated coupling reactions with not only rare-earth metal complexes but also early transition metal complexes is ongoing in our laboratory.

Acknowledgements

Y. S. acknowledges the financial support from the JSPS Postdoctoral Fellowship. H. N. expresses his special thanks for the financial support provided by the JSPS Research Fellowships for Young Scientists. H. T. acknowledges financial support from a Grant-in-Aid for Young Scientists (A) of the Ministry of Education, Culture, Sports, Science, and Technology, Japan. This work was supported by the Core Research for Evolutional Science and Technology (CREST) Program of the Japan Science and Technology Agency.

Notes and references

- 1 (*a*) T. D. Nelson and R. D. Crouch, *Org. React.*, 2004, **63**, 265; (*b*) Y. Yamamoto, Copper-mediated aryl-aryl bond formation leading to biaryls. A century after the Ullmann breakthrough, in *Copper-Mediated Cross-Coupling Reactions*, ed. G. Evano and N. Blanchard, John Wiley & Sons, Inc.: Hoboken, NJ, USA, 2014, p. 335.
- 2 (a) G. Bringmann, R. Walter and R. Weirich, Angew. Chem., Int. Ed. Engl., 1990, 29, 977; (b) J. Hassan, M. Sevignon, C. Gozzi, E. Schulz and M. Lemaire, Chem. Rev., 2002, 102, 1359.
- 3 For recent reviews of dehydrogenative coupling reactions, see: (a) C. S. Yeung and V. M. Dong, Chem. Rev., 2011, 111, 1215; (b) C.-L. Sun, B.-J. Li and Z.-J. Shi, Chem. Rev., 2011, 111, 1293; (c) C. Liu, H. Zhang, W. Shi and A. Lei, Chem. Rev., 2011, 111, 1780; (d) S. H. Cho, J. Y. Kim, J. Kwak and S. Chang, Chem. Soc. Rev., 2011, 40, 5068; (e) J. A. Ashenhurst, Chem. Soc. Rev., 2010, 39, 540; (f) L. Ackermann, R. Vicente and A. R. Kapdi, Angew. Chem., Int. Ed., 2009, 48, 9792. For recent reports of 2,2'-bipyridine formation, dehydrogenative see; (g)T. Kawashima, T. Takao and H. Suzuki, J. Am. Chem. Soc., 2007, **129**, 11006; (h) T. Takao, T. Kawashima, H. Kanda, R. Okamura and H. Suzuki, Organometallics, 2012, 31, 4817.
- 4 (*a*) A. Nakamura and S. Otsuka, *Tetrahedron Lett.*, 1974, **15**, 463; (*b*) T. T. Tsou and J. K. Kochi, *J. Am. Chem. Soc.*, 1979, **101**, 7547.
- 5 For reports of biaryl formation from phenyllanthanide complexes, see: (a) M. D. Fryzuk, J. B. Love and S. J. Rettig, *J. Am. Chem. Soc.*, 1997, 119, 9071; (b) M. D. Fryzuk, L. Jafarpour, F. M. Kerton, J. B. Love, B. O. Patrick and S. J. Rettig, *Organometallics*, 2001, 20, 1387.

- 6 For reports of biaryl formation from arylmetal complexes of late transition metals, see: (a) W. V. Konze, B. L. Scott and G. J. Kubas, J. Am. Chem. Soc., 2002, 124, 12550; (b) T. V. Ramakrishna and P. R. Sharp, Organometallics, 2004, 23, 3079; (c) M. W. Wallasch, D. Weismann, C. Riehn, S. Ambrus, G. Wolmershauser, A. Lagutschenkov, G. Niedner-Schatteburg and H. Sitzmann, Organometallics, 2010, 29, 806.
- 7 (a) N. Mizuno, K. Kamata, Y. Nakagawa, T. Oishi and K. Yamaguchi, *Catal. Today*, 2010, 157, 359; (b) G. Zhang, H. Yi, G. Zhang, Y. Deng, R. Bai, H. Zhang, J. T. Miller, A. J. Kropf, E. E. Bunel and A. Lei, *J. Am. Chem. Soc.*, 2014, 136, 924; (c) R. Bai, G. Zhang, H. Yi, Z. Huang, X. Qi, C. Liu, J. T. Miller, J. Kropf, E. E. Bunnel, Y. Lan and A. Lei, *J. Am. Chem. Soc.*, 2014, 136, 16760.
- 8 For related dianionic diyne-ligated rare-earth metalcomplexes, see: (*a*) W. J. Evans, R. A. Keyer and J. W. Ziller, *Organometallics*, 1990, **9**, 2628; (*b*) H. J. Heeres, J. Nijhoff, J. H. Teuben and R. D. Rogers, *Organometallics*, 1993, **12**, 2609; (*c*) T. Dube, J. Guan, S. Gambarotta and G. P. A. Yap, *Chem.-Eur. J*, 2001, 7, 374.
- 9 For related dianionic biaryl-ligated rare-earth metal complexes, see: (a) W. Huang, F. Dulong, T. Wu, S. I. Khan, J. T. Miller, T. Cantat and P. L. Diaconescu, *Nat. Commun.*, 2013, 4, 1448; (b) W. Huang and P. L. Diaconescu, *Eur. J. Inorg. Chem.*, 2013, 4090; (c) W. Huang, P. M. Abukhalil, S. I. Khan and P. L. Diaconescu, *Chem. Commun.*, 2014, 50, 5221.
- 10 (a) M. R. MacDonald, J. W. Ziller and W. J. Evans, J. Am. Chem. Soc., 2011, 133, 15914; (b) M. R. MacDonald, J. E. Bates, M. E. Fieser, J. W. Ziller, F. Furche and W. J. Evans, J. Am. Chem. Soc., 2012, 134, 8420; (c) M. R. MacDonald, J. E. Bates, J. W. Ziller, F. Furche and W. J. Evans, J. Am. Chem. Soc., 2013, 135, 9857; (d) C. M. Kotyk, M. R. MacDonald, J. W. Ziller and W. J. Evans, Organometallics, 2015, 34, 2287; (e) G. Meyer, Angew. Chem., Int. Ed., 2014, 53, 3550.
- 11 (a) H. Tsurugi, K. Yamamoto and K. Mashima, *J. Am. Chem.* Soc., 2011, 133, 732; (b) H. Kaneko, H. Nagae, H. Tsurugi and K. Mashima, *J. Am. Chem. Soc.*, 2011, 133, 19626.
- 12 For reports of related 2,2'-bipyridyl formation through an insertion-dimerization pathway, see: (a) B.-J. Deelman, W. M. Stevels, J. H. Teuben, M. T. Lakin and A. L. Spek, Organometallics, 1994, 13, 3881; (b) H. S. Soo, P. L. Diaconescu and C. Cummins, Organometallics, 2004, 23, 498; (c) C. T. Carver and P. L. Diaconescu, J. Am. Chem. Soc., 2008, 130, 7558; (d) C. T. Carver, D. Benitez, K. L. Miller, B. N. Williams, E. Tkatchouk, W. A. Goddard III and P. L. Diaconescu, J. Am. Chem. Soc., 2009, 131, 10269; (e) C. T. Carver, B. N. Williams, K. R. Ogilby and P. L. Diaconescu, Organometallics, 2010, 29, 835; (f) S. Jie and P. L. Diaconescu, Organometallics, 2010, 29, 1222; (g) N. Williams, W. Huang, K. L. Miller and B. P. L. Diaconescu, Inorg. Chem., 2010, 49, 11493.
- 13 (a) N. R. Kelly, S. Goetz, S. R. Batten and P. E. Kruger, *CrystEngComm*, 2008, **10**, 68; (b) A. P. Shaw, M. K. Ghosh, K. W. Törnroos, D. S. Wragg, M. Tilset, O. Sawng,

R. H. Heyn and S. Jakobsen, *Organometallics*, 2012, **31**, 7093; (*c*) T. Kawashima, T. Takao and H. Suzuki, *J. Am. Chem. Soc.*, 2007, **129**, 11006.

- 14 J. Eppinger, K. R. Nikolaides, M. Zhang-Presse, F. A. Riederer, G. W. Rabe and A. L. Rheingold, *Organometallics*, 2008, 27, 736.
- 15 (a) D. Roitershtein, A. Domingos, L. C. J. Pereira, J. R. Ascenso and N. Marques, *Inorg. Chem.*, 2003, 42, 7666;
 (b) A. V. Karpov, A. S. Shavyrin, A. V. Cherkasov, G. K. Fukin and A. A. Trifonov, *Organometallics*, 2012, 31, 5349;
 (c) Y. Zhang, J. Zhang, J. Hong, F. Zhang, L. Weng and X. Zhou, *Organometallics*, 2014, 33, 7052;
 (d) H. Nagae, Y. Shibata, H. Tsurugi and K. Mashima, *J. Am. Chem. Soc.*, 2015, 137, 640.
- 16 (a) H. Bock, J.-M. Lehn, J. Pauls, S. Holl and V. Krenzel, Angew. Chem., Int. Ed., 1999, 38, 952; (b) E. Gore-Randall,

M. Irwin, M. S. Denning and J. M. Goicoechea, *Inorg. Chem.*, 2009, **48**, 8304.

- 17 Molecular structures of **4b** and **4c** can be found in the ESI.†
- 18 (a) B.-J. Deelman, W. M. Stevels, J. H. Teuben, M. T. Lakin and A. L. Spek, Organometallics, 1994, 13, 3881; (b) R. Duchateau, E. A. C. Brussee, A. Meetsma and J. H. Teuben, Organometallics, 1997, 16, 5506; (c) B. R. Elvidge, S. Arndt, P. M. Zeimentz, T. P. Spaniol and J. Okuda, Inorg. Chem., 2005, 44, 6777; (d) S. Arndt, B. R. Elvidge, P. M. Zeimentz, T. P. Spaniol and J. Okuda, Organometallics, 2006, 25, 793.
- 19 B. N. Williams, D. Benitez, K. L. Miller, E. Tkatchouk, W. A. Goddard III and P. L. Diaconescu, J. Am. Chem. Soc., 2011, 133, 4680.