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## Introduction

The development of efficient synthetic methodologies to incorporate alkyne motifs has received broad interest, as they are not only important building blocks in natural products, pharmaceuticals and materials<sup>1</sup> but also essential functional groups in cross-coupling, metathesis and cycloaddition reactions.<sup>2</sup> Meanwhile, carboranyl acetylenes have proved to be useful basic units in molecular rods,<sup>3</sup> nonlinear optical materials,<sup>4</sup> supramolecular design,<sup>5</sup> nanovehicles<sup>6</sup> and metal-organic frameworks.<sup>7</sup> As there is a lack of direct and efficient methodologies for the synthesis of B-alkynylated carboranes, the alkyne moieties in the aforementioned materials are generally connected to cage carbon atoms,<sup>3-8</sup> which limits the application scope of the carborane derivatives.

Though cage boron alkynylated carboranes can be prepared by two-step reactions, such as the selective iodination of an *o*carborane, followed by Pd(0)-catalyzed cross-coupling with alkynyl Grignard reagents,<sup>9</sup> the installation of iodo groups to specific positions on the carboranes is necessary (Scheme 1). However, the selective iodination of cage B(4,5,7,11)–H is rather challenging, if not impossible.<sup>8</sup> Thus, we aim to develop new

# Palladium catalyzed regioselective B–C(sp) coupling *via* direct cage B–H activation: synthesis of B(4)-alkynylated *o*-carboranes†

Yangjian Quan, Cen Tang and Zuowei Xie\*

Pd-catalyzed carboxylic acid guided regioselective alkynylation of cage B(4)–H bonds in *o*-carboranes has been achieved for the first time using two different catalytic systems. In the presence of 5 mol% Pd(OAc)<sub>2</sub> and 3 equiv. of AgOAc, the reaction of 1-COOH-2-R<sup>1</sup>-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> with R<sub>3</sub>SiC=CBr in ClCH<sub>2</sub>CH<sub>2</sub>Cl gives 4-(R<sub>3</sub>SiC=C)-2-R<sup>1</sup>-*o*-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> in moderate to high yields. This reaction is compatible with alkynes possessing sterically bulky silyl groups such as <sup>i</sup>Pr<sub>3</sub>Si or <sup>t</sup>BuMe<sub>2</sub>Si. Meanwhile, another catalytic system of Pd(OAc)<sub>2</sub>/AgOAc/K<sub>2</sub>HPO<sub>4</sub> can catalyze the direct B(4)-alkynylation of 1-COOH-2-R<sup>1</sup>-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> with terminal alkynes R<sup>2</sup>C=CH in moderate to high yields. The latter has a broader substrate scope from bulky silyl to aromatic to carboranyl substituents. Desilylation of the resultant products affords carboranyl acetylene 4-(HC=C)-2-R<sup>1</sup>-*o*-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> which can undergo further transformations such as Sonogashira coupling, dimerization and click reactions. It is suggested that the above two catalytic systems may proceed *via* Pd(w)–Pd(w) and Pd(w)–Pd(w) catalytic cycles, respectively. In addition, the silver salt is found to promote the decarboxylation reaction and thereby controls the mono-selectivity.

methodologies for the selective and direct alkynylation of carboranes *via* cage B–H activation.

Directing groups are essential in transition metal catalyzed C-H activation due to their ability to chelate the metal catalyst, position it for selective C-H cleavage, and reduce activation energy by stabilizing the metallacycle intermediates.<sup>10</sup> Nevertheless, strategies using directing groups suffer from limitations when the directing groups are not present in the target molecules. To overcome this problem, the use of traceless directing groups is obviously an ideal method. Recently, the use of -COOH as a weak coordinating yet efficient directing group for transition metal catalyzed phenyl C-H activation has been documented, and has been found to be easily removed by decarboxylation after the reaction.<sup>10h</sup> Subsequently, carboxylic acid directed phenyl C-H olefination,11 arylation,12 alkylation,13 acylation,<sup>14</sup> carboxylation,<sup>15</sup> amination,<sup>16</sup> hydroxylation<sup>17</sup> and halogenation<sup>18</sup> have been successfully developed. However, to the best of our knowledge, the direct alkynylation of C-H bonds guided by -COOH is still elusive, although nitrogen-based directing-group-guided transition-metal catalyzed phenyl C-H alkynylation has been recently documented using alkynyl halides,19 hypervalent iodine-alkyne reagents20 and terminal alkynes<sup>21</sup> as the alkynylating reagents. Meanwhile, oxidative coupling of two C-H bonds for the formation of a C-C bond has received growing interest due to its benefits which include atom-economy, step-economy and less waste.<sup>22</sup> Compared with the achievements of phenyl C-H bond oxidative coupling, the regioselective and direct oxidative coupling of an organic C-H bond with a cage B-H bond in o-carboranes is very rare.23



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Department of Chemistry, State Key Laboratory of Synthetic Chemistry, The Chinese University of Hong Kong, Shatin N. T., Hong Kong, China. E-mail: zxie@cuhk.edu.hk † Electronic supplementary information (ESI) available. CCDC 1455897 and 1455898. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6sc00901h

previous work



 $\label{eq:scheme1} \begin{array}{l} \mbox{Scheme1} & \mbox{Selected examples of transition metal catalyzed formation} \\ \mbox{of cage B-C(sp) and B-C(sp^2) bonds in o-carboranes.} \end{array}$ 

Very recently, our group has developed a transition metal catalyzed –COOH guided cage B–H alkenylation<sup>23,24</sup> and arylation<sup>25</sup> of *o*-carboranes, in which the carboxyl group is removed in a one-pot fashion. Inspired by these results and other cage B–H activation reactions,<sup>26–29</sup> we have extended our research to investigate direct cage B–H alkynylation by alkynyl halides through a Pd( $\pi$ )–Pd( $\pi$ )–Pd( $\pi$ ) catalytic cycle and by terminal alkynes *via* a Pd( $\pi$ )–Pd( $\pi$ )–Pd( $\pi$ ) catalytic cycle. These new findings are reported in this article (Scheme 1).

### **Results and discussion**

#### Alkynylation using alkynyl halides

The initial reaction of 1-COOH-2-CH<sub>3</sub>-o-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> (**1a**) with <sup>i</sup>Pr<sub>3</sub>SiC=CBr in the presence of 10 mol% Pd(OAc)<sub>2</sub> and 1 equiv. of AgOAc in toluene at 90 °C for 6 h did not give any of the desired product (entry 1, Table 1). Replacement of toluene with 1,2-dichloroethane (DCE) afforded the desired coupling product 4-(<sup>i</sup>Pr<sub>3</sub>SiC=C)-2-CH<sub>3</sub>-o-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> in 40% GC yield (entry 2, Table 1). Increasing the amount of AgOAc to 3 equiv.

GC yield of 3a (entry 4, Table 1). Higher or lower reaction temperatures led to decreased yields of 3a (entries 5 and 6, Table 1). Lowering the catalyst loading to 5 mol% did not change the reaction efficiency (entry 7, Table 1). In view of the yields of 3a, entry 7 in Table 1 was chosen as the optimal reaction conditions.

A variety of carborane monocarboxylic acids (1) were examined under the chosen optimal reaction conditions, and the results are compiled in Table 2. All alkyl, alkenyl and aryl substituents on cage C(2), regardless of electronic properties, afforded the coupling products 3 in high isolated yields (entries 1–10 and 13, Table 2). For the heteroatom containing substrate 1j, the product 3j was isolated in 78% yield (entry 10, Table 2), whereas that bearing a thiophenyl group (1l) afforded the product 3l in 54% yield (entry 12, Table 2) probably due to the interaction of Pd with the S atom. Meanwhile, substrate 1k with a naphthyl substituent on cage C(2) gave 3k in only 40% isolated yield (entry 11, Table 2). For R<sup>1</sup> = H, an inseparable mixture was produced (entry 14, Table 2). When R<sup>1</sup> = Me<sub>3</sub>Si, the desilylation species 3n was isolated in 41% yield after work up (entry 15, Table 2).

In contrast to R<sup>1</sup> at cage C(2), the scope of R<sup>2</sup> is highly limited in such a coupling reaction. <sup>*t*</sup>BuMe<sub>2</sub>SiC≡CBr worked well to give **3p** in 70% isolated yield (entry 16, Table 2). However, less hindered Me<sub>3</sub>SiC≡CBr was not reactive, probably due to its propensity to coordinate with a Pd center *via* the  $\pi$  bond (entry 17, Table 2). Such a phenomenon was also observed in phenyl C-H alkynylations using R<sub>3</sub>SiC≡CBr as reagents.<sup>30</sup> It was noted that other alkynyl bromides such as PhC≡CBr and <sup>*t*</sup>BuC≡CBr were not compatible with this reaction.

#### Alkynylation using terminal alkynes

As the previous method has a limited substrate scope, we wanted to develop a more atom- and step-economic method for cage B-H alkynylation using terminal alkynes as reagents. We commenced our studies by screening for a suitable base for the oxidative coupling of cage B-H in 1-COOH-2-CH<sub>3</sub>-o-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub> (1a) with  ${}^{i}Pr_{3}SiC \equiv CH$  under the aforementioned optimal reaction conditions. No reaction was observed in the absence of a base (entry 1, Table 3). The addition of 2 equiv. of K<sub>2</sub>HPO<sub>4</sub> afforded the target product 3a in 30% GC yield with <sup>i</sup>Pr<sub>3</sub>SiC=C- $C \equiv C^{i} Pr_{3}Si$  as the side product (entry 2, Table 3). To inhibit the formation of a homocoupling side product, <sup>i</sup>Pr<sub>3</sub>SiC≡CH was added slowly via a syringe pump, leading to a significantly increased yield of 3a to 56% GC yield (entry 3, Table 3). The yield was further improved to 75% if 2 equiv. of the terminal alkyne was used (entry 4, Table 3). Replacement of 1,2-dichloroethane (DCE) with toluene resulted in a slightly higher yield of 3a (entry 5, Table 3). Decreasing the reaction temperature to 80 °C afforded 3a in 86% GC yield (entry 6, Table 3). In view of the yields of 3a, entry 6 in Table 3 was chosen as the optimal reaction conditions.

This reaction has a much broader substrate scope ( $R^2 = silyl$ , phenyl and carboranyl). The results are compiled in Table 4. For  $R^1 = alkyl$  groups, the isolated yields of 3 are comparable to those observed in Table 2. However, if  $R^1 = aryl$  unit such as 1g,

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Entry	Cat (mol%)	Additive (equiv.)	Solvent	Temp (°C)	$\operatorname{Yield}^{b}(\%)$
1	$Pd(OAc)_{2}$ (10)	AgOAc (1)	Toluene	90	N.R.
2	$Pd(OAc)_2$ (10)	AgOAc (1)	DCE	90	40
3	$Pd(OAc)_2$ (10)	AgOAc (2)	DCE	90	67
4	$Pd(OAc)_2$ (10)	AgOAc (3)	DCE	90	90
5	$Pd(OAc)_2$ (10)	AgOAc (3)	DCE	80	70
6	$Pd(OAc)_2$ (10)	AgOAc (3)	DCE	100	75
7	$Pd(OAc)_2$ (5)	AgOAc (3)	DCE	90	89
8	$Pd(OAc)_{2}$ (2.5)	AgOAc (3)	DCE	90	78
9	$Pd(TFA)_{2}$ (5)	AgOAc (3)	DCE	90	82
10	$Pd(OAc)_2$ (5)	$Ag_2CO_3(2)$	DCE	90	75
11	$Pd(OAc)_{2}$ (5)	$Ag_2O(2)$	DCE	90	63
12	$Pd(OAc)_2$ (5)	$\operatorname{AgNO}_{3}(3)$	DCE	90	30

<sup>*a*</sup> Reactions were conducted on a 0.05 mmol scale in 0.5 mL of solvent in a closed flask for 6 h; DCE = 1,2-dichloroethane; TFA = trifluoroacetate. <sup>*b*</sup> GC yields.

Table 2 Synthesis of cage B(4)-alkynylated o-carboranes using alkynyl bromide<sup> $\alpha$ </sup>



<sup>*a*</sup> Reactions were conducted on a 0.2 mmol scale of **1** in a closed flask. <sup>*b*</sup> Me<sub>3</sub>Si was removed after work up. <sup>*c*</sup> N.R. = no reaction.

the isolated yield of **3g** is 30% (entry 4, Table 4), which is significantly lower than that of 70% shown in entry 7, Table 2. On the other hand, compounds **1n** ( $\mathbb{R}^1 = \mathbb{H}$ ) and **1o** ( $\mathbb{R}^1 = \mathbb{M}e_3Si$ ) give **3n** in 35% and 74% yields, respectively (entries 5 and 6, Table 4). These yields are much higher than those found in the previous reaction (entries 14 and 15, Table 2). The reasons for this phenomenon are not clear at this stage.

More importantly, this catalytic system is compatible with phenyl acetylene, producing the corresponding product **3r** in 52% isolated yield (entry 8, Table 4). The coupling efficiency was largely enhanced from  $R^2 = Ph$ , 2-MeC<sub>6</sub>H<sub>4</sub>, 2,6-(Me)<sub>2</sub>C<sub>6</sub>H<sub>3</sub>, 2-<sup>i</sup>PrC<sub>6</sub>H<sub>4</sub> to 1-<sup>*n*</sup>C<sub>6</sub>H<sub>13</sub>-*o*-C<sub>2</sub>B<sub>10</sub>H<sub>10</sub>, affording the corresponding products, **3s**, **3t**, **3u** and **3v**, in 65%, 73%, 80% and 82% isolated yields, respectively (entries 9–12, Table 4). It should be noted that *o*-carboranyl is a strong electron-withdrawing unit.<sup>4c</sup> In view of the isolated yields of **3w** and **3x** (entries 13 and 14, Table 4), the electronic effects on the reactions are not obvious as -CH<sub>3</sub> and -CF<sub>3</sub> have significantly different electronic properties. The above data (entries 8–14, Table 4) indicate strongly that bulkier substituents favor the formation of coupling products.

#### Transformation of 3a

To demonstrate the applications of the resultant compounds **3** as building blocks, further transformation of **3a** was carried out. The <sup>i</sup>Pr<sub>3</sub>Si group in **3a** was readily removed by treatment with TBAF (TBAF = tetra-*n*-butylammonium fluoride) to afford quantitatively the terminal alkyne **4a** (Scheme 2). Like other terminal alkynes, compound **4a** can undergo various transformations to give different kinds of carborane-incorporated functional molecules. Sonogashira coupling of **4a** with iodobenzene or 2-bromothiophene generated **3r** or **5a** in 92% and



Entry	Cat (mol%)	Additive (equiv.)	Solvent	Temp (°C)	$\operatorname{Yield}^{b}(\%)$
1 <sup><i>c</i></sup>	$Pd(OAc)_2$ (5)	AgOAc (3)	DCE	90	N.R.
2	$Pd(OAc)_2$ (5)	AgOAc (3)	DCE	90	30
3	$Pd(OAc)_2$ (5)	AgOAc (3)	DCE	90	$56^d$
4	$Pd(OAc)_2$ (5)	AgOAc (3)	DCE	90	$75^{d,e}$
5	$Pd(OAc)_2(5)$	AgOAc (3)	Toluene	90	$78^{d,e}$
6	$Pd(OAc)_2$ (5)	AgOAc (3)	Toluene	80	$86^{d,e}$
7	$Pd(OAc)_2(5)$	AgOAc (3)	Toluene	70	Trace
8	$Pd(OAc)_2$ (3)	AgOAc (3)	DCE	90	18
9	$Pd(TFA)_2(5)$	AgOAc (3)	DCE	90	26
10	$Pd_2(dba)_3(5)$	AgOAc (3)	DCE	90	21
11	$Pd(OAc)_2(5)$	$Ag_2CO_3(2)$	DCE	90	15
12	$Pd(OAc)_2(5)$	$Ag_2O(2)$	DCE	90	12
13	$Pd(OAc)_2(5)$	$AgNO_3(3)$	DCE	90	Trace

<sup>*a*</sup> Reactions were conducted on a 0.05 mmol scale of **1a** in 0.5 mL of solvent in the presence of 2 equiv. of  $K_2HPO_4$  in a closed flask for 10 h; DCE = 1,2-dichloroethane; TFA = trifluoroacetate; dba = dibenzylideneacetone. <sup>*b*</sup> GC yields. <sup>*c*</sup> Without  $K_2HPO_4$ . <sup>*d*</sup> Terminal alkyne was added dropwise by a syringe pump over a period of 10 h. <sup>*e*</sup> Two equiv. of terminal alkyne was added.

**Table 4** Synthesis of cage B(4)-alkynylated *o*-carboranes using terminal alkynes<sup>*a*</sup>

	H COOH R <sup>1</sup> 1	5% Pd(OAc) <sub>2</sub> 9 R <sup>2</sup> — H ( <b>2</b> ) 3 eq AgOAc 2 eq K <sub>2</sub> HPO <sub>4</sub> fol./80 °C/10 h	$R^2$ H $R^1$ 3
Entry	R <sup>1</sup>	$R^{2}(2)$	Isolated yield (%)
1	Me (1a)	<sup>i</sup> Pr <sub>3</sub> Si	79 ( <b>3a</b> )
2	<sup>i</sup> Pr (1c)	<sup>i</sup> Pr <sub>3</sub> Si	86 ( <b>3c</b> )
3	Bn ( <b>1d</b> )	<sup>i</sup> Pr <sub>3</sub> Si	70 ( <b>3d</b> )
4	$3,5-(CH_3)_2C_6H_3$ (1g)	<sup>i</sup> Pr <sub>3</sub> Si	30 ( <b>3g</b> )
5	Н ( <b>1n</b> )	<sup>i</sup> Pr <sub>3</sub> Si	35 ( <b>3n</b> )
6	$Me_3Si(10)$	<sup>i</sup> Pr <sub>3</sub> Si	$74^{b}$ (3n)
7	Me (1a)	<sup>t</sup> BuMe <sub>2</sub> Si	72 ( <b>3p</b> )
8	Me (1a)	Ph	$52^{c}$ (3r)
9	Me (1a)	$2-CH_3C_6H_4$	$65^{c}$ (3s)
10	Me (1a)	$2,6-(CH_3)_2C_6H_3$	$73^{c}$ (3t)
11	Me (1a)	2- <sup>i</sup> PrC <sub>6</sub> H <sub>4</sub>	$80^{c}$ (3u)
12	Me (1a)	Сбн <sub>13</sub>	$82^{c}$ (3v)
13	Me (1a)	4-CH <sub>3</sub> C <sub>6</sub> H <sub>4</sub>	$48^{c}$ ( <b>3w</b> )
14	Me (1a)	$4\text{-}\mathrm{CF}_3\mathrm{C}_6\mathrm{H}_4$	$44^{c}$ (3x)

<sup>a</sup> Reactions were conducted on a 0.2 mmol scale of 1 in a closed flask.
 <sup>b</sup> Me<sub>3</sub>Si was removed after work up. <sup>c</sup> 3 equiv. of terminal alkyne was used.

90% isolated yields, respectively. Glaser–Hay homocoupling of **4a** gave 1,4-dicarboranyldiacetylene (**6a**) in 84% isolated yield. A click reaction of **4a** with phenyl azide afforded carborane-functionalized 1,2,3-triazole (**7a**) in 95% isolated yield.

All new compounds 3 and 4a–7a were fully characterized by <sup>1</sup>H, <sup>13</sup>C, and <sup>11</sup>B NMR spectroscopy as well as high-resolution mass spectrometry (HRMS).<sup>31</sup> Molecular structures of 4a and 6a were further confirmed by single-crystal X-ray analyses and are shown in Fig. 1. Experimental details are included in the ESI.†

#### **Reaction mechanism**

To gain some insight into the reaction mechanism, the following control experiments were carried out. No reaction was observed if **1a** was treated with 1 equiv. of  ${}^{i}Pr_{3}SiC \equiv CBr$  in the presence of 20 mol% Pd(dba)<sub>2</sub> (dba = dibenzylideneacetone) in DCE at 90 °C for 6 h in the absence of AgOAc. On the other hand, under the same reaction conditions, replacement of Pd(dba)<sub>2</sub> with Pd(OAc)<sub>2</sub> gave the alkynylation product **3a** in 30% GC yield (Scheme 3a). Similarly, in the presence of 20 mol% Pd(OAc)<sub>2</sub>, the reaction of **1a** with 2 equiv. of  ${}^{i}Pr_{3}SiC \equiv CH$  afforded **3a** in 16% GC yield without AgOAc as the oxidant. While, no **3a** was observed when 20 mol% Pd(dba)<sub>2</sub> was used instead of Pd(OAc)<sub>2</sub> (Scheme 3b). These results suggest that both cross-coupling reactions are initiated by Pd(n) not Pd(0).

Decarboxylation of carboranyl carboxylic acids (**1b** and **3b**-COOH) was also examined (Scheme 3c). Compound **1b** was stable after heating at 90 °C for 12 h in DCE, whereas **3b**-COOH underwent complete decarboxylation within one hour under the same reaction conditions. Notably, it only took ten minutes to



convert **3b**-COOH to **3b** in the presence of 1 equiv. of AgOAc. These results clearly indicate that the introduction of an alkynyl group at the cage B(4) site can induce the decarboxylation, and the addition of a silver salt can accelerate such decarboxylation, which is crucial for controlling the mono-selectivity.

On the basis of the aforementioned experimental data, two plausible reaction mechanisms are proposed in Scheme 4. For the  $Pd(\pi)-Pd(\pi)-Pd(\pi)$  catalytic cycle: an exchange reaction of 1 with  $Pd(OAc)_2$ , followed by regioselective electrophilic attack at



Fig. 1 Molecular structures of 4a (top) and 6a (bottom) (only the terminal alkyne H atom is shown for clarity).



Scheme 3 Control experiments.



Scheme 4 Proposed reaction mechanism.

the more electron-rich cage B(4) site yields the intermediate A as the charge distribution on the cage follows the trend B(9,12) >B(8,10) > B(4,5,7,11) > B(3,6).<sup>32</sup> Oxidative addition of R<sup>2</sup>C=CBr affords a Pd(IV) intermediate B.25,33 Reductive elimination produces the intermediate C, which undergoes a salt metathesis reaction, protonation and decarboxylation to give the final product 3 and regenerates the catalyst Pd(OAc)<sub>2</sub>. Meanwhile, another catalytic system involves a  $Pd(\pi)-Pd(0)-Pd(\pi)$  cycle. An acid-base reaction between K2HPO4 and carboranyl carboxylic acid 1 gives the potassium salt 1'.34 Coordination of the oxygen atom of  $\mathbf{1}'$  to the Pd(II) center, followed by subsequent regioselective electrophilic attack at the more electron-rich cage B(4)site generates the intermediate D. Ligand exchange by acetylide gives a carboranyl-palladium acetylide intermediate E.<sup>21b,35</sup> Reductive elimination affords the cage B(4)-alkynylated intermediate F and Pd(0). Decarboxylation of F results in the formation of the final product 3, meanwhile Pd(0) is oxidized by AgOAc to regenerate  $Pd(OAc)_2$ . It is noted that AgOAc acts as a bromide captor in the  $Pd(\pi)-Pd(\pi)-Pd(\pi)$  catalytic cycle, but as an oxidant to regenerate  $Pd(\pi)$  from Pd(0) in the  $Pd(\pi)-Pd(0)$ - $Pd(\pi)$  catalytic cycle. However, in both cross-coupling reactions, AgOAc plays a crucial role in promoting decarboxylation and thereby controlling the mono-selectivity.

# Conclusion

We have developed two catalytic systems for regioselective and efficient alkynylation of cage B(4)–H bonds in *o*-carboranes using alkynyl bromides or terminal alkynes as alkynylating agents, where –COOH acts as a traceless directing group. A series of new cage B(4)-alkynylated *o*-carborane derivatives has been prepared for the first time, which could find many applications in the synthesis of carborane-based materials.<sup>3–7</sup> This opens up a new window for the functionalization of carboranes by direct oxidative coupling of the cage B–H and organic C–H bonds. This work also offers a useful reference for selective C–H alkynylation using carboxylic acid as a traceless directing group in other aromatic systems.

On the basis of control experiments and literature work, two catalytic cycles are proposed for the above two reactions: a  $Pd(\pi)-Pd(\pi)-Pd(\pi)$  cycle for using alkynyl bromides as coupling agents and a  $Pd(\pi)-Pd(0)-Pd(\pi)$  cycle for employing terminal alkynes as coupling partners. The latter has a broader substrate scope than the former. This work also gives some hints for the development of new catalytic systems for the functionalization of carboranes.

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