


 Cite this: *Chem. Commun.*, 2016, **52**, 9363

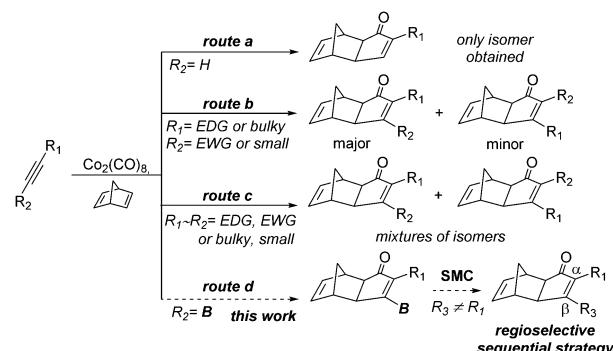
 Received 7th June 2016,
 Accepted 24th June 2016

DOI: 10.1039/c6cc04717c

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The first intermolecular Pauson–Khand reaction, conducted using internal alkynylboronic esters, allows the installation of the boronic ester moiety at the β -position of the cyclopentenone with total regio- and stereoselectivity.

Since its first report by Khand and Pauson in the early seventies,¹ the Pauson–Khand reaction (PKR) has proven to be a very powerful tool to build cyclopentenone cores. The reaction consists of a [2+2+1] cycloaddition involving an alkyne, an alkene and carbon monoxide allowing the assembly of architecturally complex molecules from very simple starting materials.² The intermolecular version has to face challenging issues namely the regioselective addition of alkenes and alkynes. Towards this end, terminal alkynes have been successfully employed yielding the α -substituted adduct in a regioselective manner (Scheme 1, route a). However, internal nonsymmetric alkynes can potentially provide both regioisomers.³ Interestingly, when sterically or electronically biased internal alkynes are used, the largest or the most electrondonating group (EDG) is mainly (but not exclusively) placed at the α -position of the resulting adduct (Scheme 1, route b). However, when non-biased alkynes are used, mixtures of α/β -substituted adducts are observed (Scheme 1, route c).⁴ This fact has been supported by recent computational studies where the lack of electronic or steric differentiation along the triple bond does not allow, in some cases, a clear distinction in the regioselective outcome of the reaction.⁵ Bearing these limitations in mind, we envisioned the possibility of using boron chemistry to solve the lack of selectivity in the PKR reaction for internal alkynes, complementing the current approaches to prepare α,β -diarylcyclopentenones, namely the Nazarov reaction,⁶ Ni- and Pd-catalysed annulations⁷ or Au-catalyzed [4+1] cyclizations.⁸



Scheme 1 Regioselective scenarios of alkynes in intermolecular PKR.

Our proposal is based on a two-step strategy and its usefulness lies in the use of internal alkynylboronic esters, which take advantage of the polarization along the triple bond falling into a specific case of biased internal alkynes (Scheme 1, route d). Internal alkynylboronic esters are versatile synthons that have been successfully used in a vast array of reactions such as standard Suzuki–Miyaura cross-coupling reactions⁹ or cycloaddition reactions,¹⁰ but its use in the PKR is surprisingly unprecedented in the literature.

Herein, we describe, the first PKR using internal alkynylboronic esters, with the exclusive formation of a single *exo* stereoisomer and the β -regioisomer. Subsequently, the *B*-substituted cyclopentenones are subjected to the Suzuki–Miyaura cross-coupling (SMC) allowing the overall sequential installation of α and β substitutions. This strategy allows access to both regioisomers by simply reversing the order of introduction of the substituents, converting it to a very direct and intuitive tool (Scheme 2).

We initiated our study by optimizing the PKR using 4,4,5,5-tetramethyl-2-(phenylethynyl)-1,3,2-dioxaborolane (**1a**) as a model substrate (Table 1). We found that the optimal reaction conditions were the combination of $\text{Co}_2(\text{CO})_8$ (1 equiv.), norbornadiene (NBD, 3 equiv.), and 4-methylmorpholine *N*-oxide (NMO, 6 equiv.) in dichloromethane (0.03 M) at room temperature yielding the corresponding adduct **2a** in 75% isolated yield (Table 1, entry 1). Along with the desired product **2a**, the protodeborylated side-product **3** was

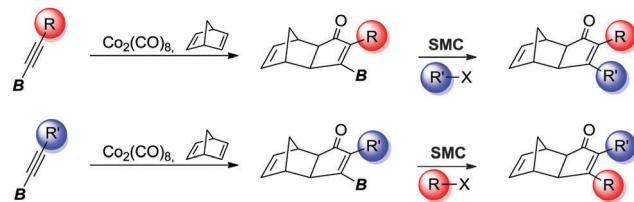
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[†] Electronic supplementary information (ESI) available. CCDC 1483204–1483206. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c6cc04717c





Scheme 2 Selection of the regiosomer by a sequential strategy.

Table 1 Intermolecular PKR of **1a**^a

Entry	Change from optimized conditions	2a ^b (%)	3 (%)
1	None	89 (75) ^c	9
2 ^d	Using thermal conditions	38	12
3 ^d	Using thermal conditions ^e with 1 equiv. PPh ₃	20	13
4	With acetone instead of DCM	23	37
5	0 °C instead of r.t.	70	15
6	With slow addition of NMO ^f	75	20
7	Using complex 4 ^g	82 (75) ^c	6

^a **1a** (0.6 mmol), Co₂(CO)₈ (1 equiv.), NBD (3 equiv.), and NMO (6 equiv.) in DCM (0.03 M) at r.t. for 16 h. ^b NMR yield using 1,1,2,2-tetrachloroethane as an internal standard. ^c Isolated yield. ^d Thermal conditions: **1a**, Co₂(CO)₈ (1 equiv.), NBD (3 equiv.) in toluene (0.1 M) at 70 °C. ^e At 100 °C. ^f Via syringe pump. ^g **4** (0.2 mmol), NBD (3 equiv.), NMO (6 equiv.) in DCM (0.03 M) at r.t.

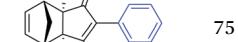
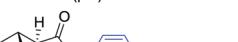
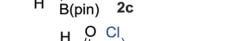
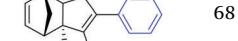
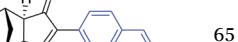
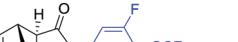
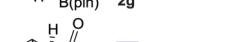
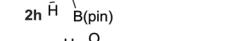
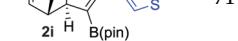
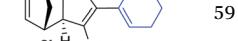
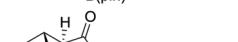
obtained in 9% yield. However, when thermal conditions were used in toluene, the yield of adduct **2a** dropped to 38% together with 12% yield of side-product **3** (Table 1, entry 2), probably due to decomposition of intermediate species during the reaction. Attempts to stabilize the cobalt complex with PPh₃ had a detrimental impact on the yield lowering it to 20% of **2a** (entry 3). Furthermore, changing from dichloromethane to acetone resulted in the major formation of the side-product **3** (entry 4). Lower temperatures or slow addition of NMO did not provide any benefit to the reaction outcome (entries 5 and 6). The side-product **3** is presumably formed during the complexation step of alkyne **2a** with the cobalt carbonyl complex. In order to shed light on the origin of **3**, we synthesized the hexacarbonylic complex **4**¹¹ to test it in the PKR. Adduct **2a** was obtained in comparable yields to the one of the optimized conditions (compare entry 1 with 7). It is plausible to think that **1a** forms two hexacarbonylic complexes, namely complex **4** and the corresponding standard hexacarbonylic complex bearing the terminal (deborylated)phenylacetylene, in which both undergo PKR forming **2a** and **3**, respectively. Experiments based on the decomposition of **2a** by treatment with silica-gel or activated charcoal did not provide any traces of **3**.

We next turned our attention to study the preparative scope of this reaction (Table 2). Towards this end, we succeeded to convert a wide range of terminal alkynes into their corresponding alkynylboronic pinacol esters (**1a–1o**), using the methodology developed by Brown and Srebnik.¹² The fifteen internal alkynes reacted with norbornadiene, in the presence of Co₂(CO)₈ and NMO, to give a unique single *exo*-stereoisomer and the β -regioisomer of the α , β -substituted cyclopentenones (Table 2).

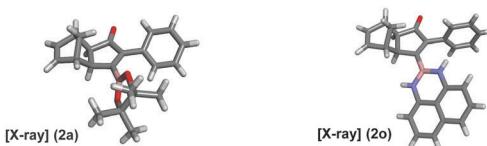
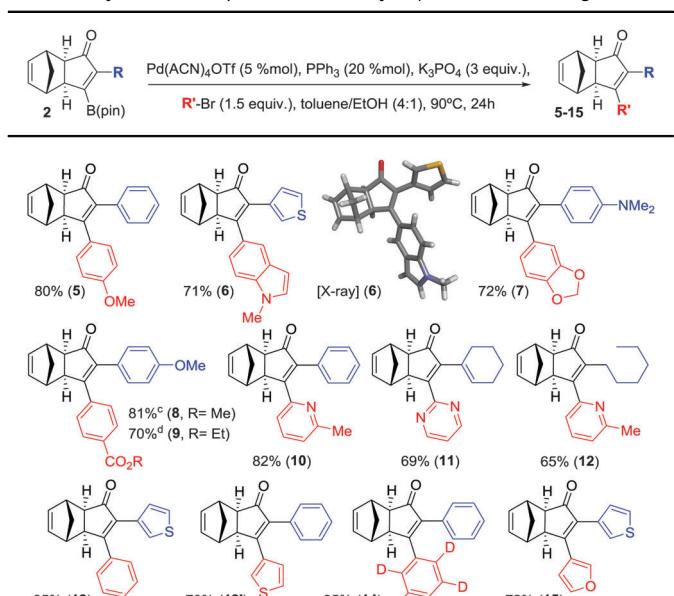
As shown in Table 2, the substitution pattern includes aromatic groups with a vast array of substituents with different electronic properties (entries 1–8), heteroaryl group (entry 9), olefinic (entry 10) or aliphatic groups (entries 11–13). However, when hindered alkynes such as *tert*-butylalkynylboronic pinacol ester or trimethylsilyl-alkynylboronic pinacol ester were used, the PKR failed to yield the corresponding adduct. Alkynes bearing Bbzpin (bzpin: benzo-pinacol) or Bdan (dan: 1,8-diaminonaphthalene) moieties were also tolerated (entries 14 and 15). In all cases, the resulting adducts were obtained with total stereo- and regioselectivity since none of the other possible isomers were detected by analytical means.¹³ Suitable crystals of **2a** and **2o** for X-ray diffraction analysis were successfully obtained for the first time from pentane and the selectivity of the pinacolboryl moiety at the β position of cyclopentenones could be unequivocally confirmed (Fig. 1). It confirms that the total regioselectivity observed is outweighed by electronic effects over the steric ones since adducts **2a** and **2o**, both bearing different steric requirements, exclusively placed the boryl moiety at the β position. Overall, we believe that the results presented in Table 2 not only demonstrate a high reactivity and functional group compatibility, but also the potential of alkynylboronic esters in the stereo- and regioselective PKR, comparable to the [2+2+1] cycloaddition from biased nonsymmetric internal alkynes,¹⁴ and the two-step protocol to form β -substituted cyclopentenones using the trifluoromethyl group as a removable steering group.¹⁵

In order to demonstrate the usefulness of the β -boryl α -substituted cyclopentenones, we next turned our attention to their application towards the SMC, in line with our tunable synthetic strategy (Scheme 2) as a reliable and straightforward methodology.¹⁶ The reaction conditions were optimized using adduct **2a** as a model substrate and 4-methoxyphenyl bromide as a reagent to furnish the desired α , β -disubstituted cyclopentenone **5** in 80% isolated yield (Table 3), in the presence of Pd(ACN)₄OTf (5 mol%), PPh₃ (20 mol%), and K₃PO₄ (3 equiv.) in a mixture of toluene/ethanol (4 : 1) at 90 °C during 24 hours. Our protocol was optimal for the assembly of several compounds regardless of their electronic properties or substitution patterns, providing the selective α , β -disubstituted cyclopentenones (**5–15**) with moderate to high isolated yields ranging 65–85% (Table 3). More interestingly, our methodology enabled the regioselective construction of challenging substrates that cannot be accessed through a standard PKR, for instances: (1) compounds **6**, **7** and **10–12** were synthesized by installing the most hindered substituent at the β position, (2) compounds **5** and **7–9** bearing electronically or sterically non-differentiable substituents in both positions were successfully obtained, (3) isomers that are usually not accessible when carrying out the standard PKR such in the case of compound **12**, holding an alkyl group at the α position. It should be highlighted that a number of heterocycles could be easily made. Crystals of **6** were successfully analysed by X-ray diffraction confirming the regioselectivity of the overall strategy claimed here.¹⁷ However, the strength of this methodology lies in the possibility of preparing specific regioisomers bearing negligible distinction in the substitution between α and β positions which would yield no selectivity through a standard PKR. Thus, compounds **14** and **15**, containing minimal differentiation between both positions, such as phenyl

Table 2 Synthesis of alkynylboronic esters and their application in the PKR^{a,b}

Entry	Alkenes 1	Yield 1 (%)	PK adduct 2	Yield 2 (%)
1	(pin)B— 	72		75
2	(pin)B— 	64		53
3	(pin)B— 	61		60
4	(pin)B— 	67		66
5	(pin)B— 	67		68
6	(pin)B— 	72		65
7	(pin)B— 	75		55
8	(pin)B— 	81		65
9	(pin)B— 	69		71
10	(pin)B— 	78		59
11	(pin)B— 	69		64
12	(pin)B— 	64		72
13	(pin)B— 	89		68
14 ^{c,d}	(bzpin)B— 	64		61
15 ^{c,e}	(dan)B— 	53		81

^a As for Table 1, entry 1. ^b Isolated yields. ^c See the ESI for detailed synthetic procedure. ^d bzpin: benzopinacol. ^e dan: 1,8-diaminonaphthalene.

Fig. 1 X-ray structures of compounds **2a** and **2o**.Table 3 Synthesis of α,β -disubstituted cyclopentenones through SMC^{a,b}

^a 2 (0.14 mmol), Pd(ACN)₄OTf (5 mol%), PPh₃ (20 mol%), K₃PO₄ (3 equiv.), R'-Br (1.5 equiv.) in toluene/EtOH (4:1) at 90 °C during 24 h. ^b Isolated yield.

^c With MeOH instead of EtOH. ^d From methyl 4-bromobenzoate.

versus *d*₅-phenyl or 3-thionyl versus 3-furyl, could be successfully prepared.

In order to better illustrate the potential of this strategy, we selected a few examples from the literature, which suffer from regioselectivity issues when synthesized through a standard PKR (Fig. 2). Direct comparison with representative closely-related compounds showed an substantial improvement in regioselectivity. Compounds **5** and **8** were obtained as a single regioisomer while a standard methodology afforded mixtures that are usually difficult to handle. Finally, cyclopentenones **13** and **13'** were independently obtained by simple reversal in the order of addition of the reagents, thus illustrating the ability to access both regioisomers.

We eventually focus our strategy to the synthesis of a known scaffold of cyclooxygenase-2 (COX-2)¹⁸ (Scheme 3). COX-2 inhibitors are a type of non-steroidal anti-inflammatory drug that directly targets COX-2 enzymes, which are responsible for inflammation and pain. We tackled its synthesis using our sequential strategy starting from the alkynylboronic pinacol esters **1h**. PKR afforded exclusively the regioisomer **2h** (Table 2, entry 8), which was submitted to the SMC reaction with 4-bromophenyl methyl sulfone, yielding **16** in 88% isolated yield as a key scaffold.

In summary, we have described an intermolecular Pauson–Khand reaction conducted using internal alkynylboronic esters installing the boronic ester moiety at the β -position of the cyclo-

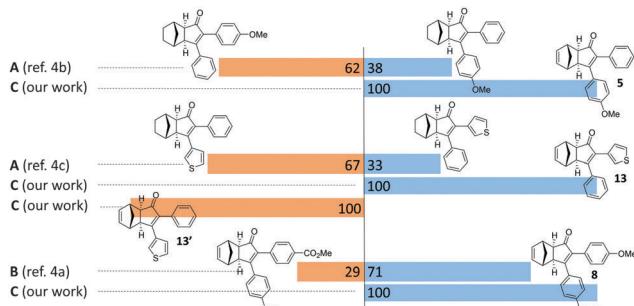
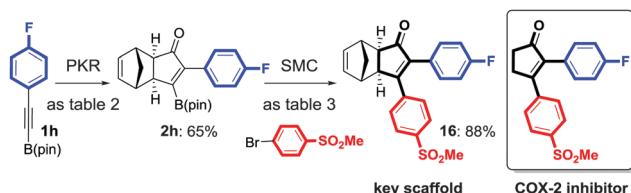


Fig. 2 Regioselective outcome comparison of selected examples towards the application of our strategy. Reaction conditions: Method A: (a) $\text{Co}_2(\text{CO})_8$, DCE; (b) norbornene, microwave, 90 °C. Method B: (a) $\text{Co}_2(\text{CO})_8$, hexane; (b) norbornadiene, 60 °C. Method C: (a) as for Table 2; (b) as for Table 3.



Scheme 3 Synthesis of a key scaffold **16**.

pentenone with total regio- and stereoselectivity. More interestingly, the resulting adduct can be subsequently derivatised by Suzuki–Miyaura cross-coupling reaction allowing any combination of α,β -disubstituted cyclopentenones, solving a well-known and long-standing drawback about regioselective Pauson–Khand reactions. Overall, our approach also complements existing PKR protocols providing access to regioisomers that are not formed as the major isomers in the standard PKR. Currently, the methodology is being exploited using other challenging substrates.

We thank the Spanish Ministerio de Economía y Competitividad (CTQ2013-43395-P) for financial support. T. L. thanks ACCIÓ for the TecnoiSpring funding (TECSPR13-1-0040).

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