Divergent Pd-catalyzed cross-coupling of allenyloxazolidinones to give chiral 1,3-dienes and vinyloxazolidinones

Ronald Brown
*University of Wollongong, rb976@uowmail.edu.au*

Farzad Zamani
*University of Wollongong, fz774@uowmail.edu.au*

Michael G. Gardiner
*Australian National University*

Haibo Yu
*University of Wollongong, hyu@uow.edu.au*

Stephen G. Pyne
*University of Wollongong, spyne@uow.edu.au*

See next page for additional authors

Follow this and additional works at: [https://ro.uow.edu.au/smhpapers1](https://ro.uow.edu.au/smhpapers1)

**Publication Details Citation**
Brown, R., Zamani, F., Gardiner, M. G., Yu, H., Pyne, S. G., & Hyland, C. J. (2019). Divergent Pd-catalyzed cross-coupling of allenyloxazolidinones to give chiral 1,3-dienes and vinyloxazolidinones. Faculty of Science, Medicine and Health - Papers: Part B. Retrieved from [https://ro.uow.edu.au/smhpapers1/974](https://ro.uow.edu.au/smhpapers1/974)

Research Online is the open access institutional repository for the University of Wollongong. For further information contact the UOW Library: research-pubs@uow.edu.au
Divergent Pd-catalyzed cross-coupling of allenyloxazolidinones to give chiral 1,3-dienes and vinyloxazolidinones

Abstract
The divergent reactivity of 5-allenyloxazolidinones has been explored. This novel building block undergoes Pd(0)-catalyzed cross-coupling with boronic acids to form a wide range of chiral 1,3-dienes and pharmaceutically useful vinyloxazolidinones, the chemoselectivity being tightly controlled by a simple switch in additive.

Publication Details
Brown, R. W., Zamani, F., Gardiner, M. G., Yu, H., Pyne, S. G. & Hyland, C. J.T. (2019). Divergent Pd-catalyzed cross-coupling of allenyloxazolidinones to give chiral 1,3-dienes and vinyloxazolidinones. Chemical Science, 10 (39), 9051-9056.

Authors
Ronald Brown, Farzad Zamani, Michael G. Gardiner, Haibo Yu, Stephen G. Pyne, and Christopher J. T Hyland

This journal article is available at Research Online: https://ro.uow.edu.au/smhpapers1/974
Divergent Pd-catalyzed cross-coupling of allenyloxazolidinones to give chiral 1,3-dienes and vinyloxazolidinones†

Ronald W. Brown,a Farzad Zamani,a Michael G. Gardiner,b Haibo Yu,a Stephen G. Pynea and Christopher J. T. Hyland*a

The divergent reactivity of 5-allenyloxazolidinones has been explored. This novel building block undergoes Pd(0)-catalyzed cross-coupling with boronic acids to form a wide range of chiral 1,3-dienes and pharmaceutically useful vinyloxazolidinones, the chemoselectivity being tightly controlled by a simple switch in additive.

The synthetic utility of strained vinylic building blocks such as vinylaziridines and vinyloxazolidinones (Scheme 1A) in constructing high-value N-containing scaffolds is well-established. Upon activation by low-valent transition metals such as palladium(0), 2-vinylaziridines and 5-vinylloxazolidinones are known to undergo direct cross-coupling and cycloaddition reactions,1–5 which often proceed via a Pd–π-allyl intermediate. Systems with cumulated double bonds, most notably the allene unit, are valuable synthetic tools as the reactive 1,2-diene structure opens up reaction pathways otherwise inaccessible to simple vinyl systems. The high degree of unsaturation facilitates tandem, multistep reactivity, and can be retained in the products of cycloaddition and carbometalation processes.6 It is therefore surprising that the alleny analogues of vinylaziridines and vinyloxazolidinones have not been investigated to date.

We have recently developed a novel class of enantioenriched 5-allenyloxazolidinones (Scheme 1B) derived from l-α-amino aldehydes,7 serving as an entirely new building block with complementary reactivity to the aforementioned vinylic systems. We have recently reported that these systems can be converted in situ to 1,3-(E)-enyne and preclude the dual gold-catalyzed cycloaromatization of unconjugated-(E)-enedynes to enantioenriched isoindolines.8 However, direct utilization of the allene unit of this building block in transition metal-catalyzed reactions has the potential to unlock new reactivity. Like its vinyl counterpart, the reactivity of the allene moiety can be harnessed through palladium catalysis,9 notably Pd(0)-catalyzed cross-coupling reactions with boronic acids and alkenylcarbinol derivatives to give substituted 1,3-dienes,10 a process first reported by Miyaura in 1994.10 The mild conditions associated with such routes to these invaluable precursors to complex molecular architectures have led to more recent examples in the literature,10c however all examples to date have provided achiral products. We envisioned that the N-nosyl derivatives (1) of our newly-developed 5-allenyloxazolidinones could undergo decarboxylative cross-coupling under Suzuki-Miyaura conditions to generate 2-substituted 1,3-dienes 2

Scheme 1 (A) Known reactivity of 2-vinylaziridines and 5-vinyl-oxazolidinones. (B) Previously unknown 5-allenyloxazolidinones developed in the Hyland and Pyne groups. (C) Novel, divergent reactivity of 5-allenyloxazolidinones reported herein.

† Electronic supplementary information (ESI) available. CCDC 1919153 and 1919154. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9sc03215k

A. Known:

2-vinylaziridines

5-vinyl-
oxazolidinones

cross-coupling
dipolarophiles
cycloaddition

B. Unknown:

5-alleny-
loxazolidinones

Pd(0)–Pd

C. This work:

2

3

Pd(0)-K2CO3

Pd(0)-AcOH

1

MRX-I antibiotic

cytchalasin

actin inhibitors

1,3-diene intermediate

isoindolines

Scheme 1 (A) Known reactivity of 2-vinylaziridines and 5-vinyl-oxazolidinones. (B) Previously unknown 5-allenyloxazolidinones developed in the Hyland and Pyne groups. (C) Novel, divergent reactivity of 5-allenyloxazolidinones reported herein.

This journal is © The Royal Society of Chemistry 2019

Chem. Sci., 2019, 10, 9051–9056 | 9051
bearing both a stereogenic centre and pendant a nitrogen atom: a system that is perfectly tailored for Type I intramolecular Diels–Alder (IMDA) reactions to give densely functionalized, stereodefined isoindoline scaffolds (Scheme 1C). A critical limitation of the powerful IMDA reaction is the lack of synthetic routes to complex and stereodefined 1,3-diene substrates, highlighting the importance of the work reported herein. Interestingly, while optimizing the construction of these 1,3-dienes, exchanging the base for AcOH led to the exclusive formation of the substituted 5-vinylxazolidinone scaffold 3, bearing structural resemblance to oxazolidinone antibiotics such as linezolid, ranizolid and the recently developed MRX-L. It is now apparent that the novel class of enantioenriched allenylxazolidinones developed in our group exhibits complementary reactivity to 2-vinylaziridines and 2-vinylxazolidinones in the pursuit of stereodefined N-heterocyclic scaffolds. Moreover, this system is capable of an elegant divergence in reactivity, exclusively giving rise to either N-tethered, stereodefined 2-substituted 1,3-dienes or medicinally relevant substituted 5-vinylxazolidinones by virtue of a single switch in additive, under mild conditions, in good to excellent yields and with high stereoselectivity and enantioretention.

The required 5-allenyxazolidinone building blocks 1a–d were prepared via our previously developed zinc-catalyzed allenylation of α-α-amino aldehydes, then an N-nosyl group introduced to promote decarboxylative ring-opening. In order to develop and optimize the 1,3-diene synthesis, the valine-derived allenylxazolidinone 1a was subjected to Suzuki–Miyaura conditions using p-methoxyphenylboronic acid as the cross-coupling partner and Pd2dba3–CHCl3 as the Pd(0) source (Table 1). The desired 1,3-diene 2a was obtained in good yield and stereoselectivity (E : Z) in the absence of an additional ligand (Table 1, entry 1), however the introduction of a phosphine ligand significantly increased the stereoselectivity (entries 2–6). A bulky phosphine (entry 4) or phosphite (entry 6) ligand improved yield and stereoselectivity, prompting the use of the bulky phosphine ligand L (tris(2,4-di-tert-butylphenyl)phosphate). While the previous ligands left the product requiring multiple purifications, ligand L cleanly afforded the 1,3-diene after column chromatography while maintaining yield and stereoselectivity (entry 7). This, as well as being a relatively cheap, non-toxic and air-stable solid, left phosphite ligand L the optimal choice. While investigating the effect of other bases typically used in Suzuki–Miyaura type cross-couplings, it was found that KOAc promoted the formation of the directly cross-coupled 5-vinylxazolidinone 3a (entry 8). Given the existing literature on Pd(0)/AcOH-catalyzed cross-coupling of boronic acids to allenes, it was suspected that this reaction was catalyzed by trace AcOH. This was confirmed by the addition of 1.5 equivalents of AcOH (entry 9) and unlocked a divergent path to the 5-vinylxazolidinone in excellent yield and stereoselectivity with low catalyst loading (entry 10), though the 1,3-diene required a slightly higher catalyst loading to maintain stereoselectivity and chemoselectivity (entries 11 and 12). The E stereochemistry of the 1,3-diene 2a could be confirmed from its 1H NMR spectrum, while X-ray crystallography provided unambiguous proof for the E stereochemistry of the 5-vinylxazolidinone 3a. It was confirmed via chiral HPLC that both reactions proceed with complete enantioretention (>99% ee).

Once the optimal conditions for the divergent synthesis of the 1,3-dienes or 5-vinylxazolidinones had been established, a crystal structure was obtained for 1a (CCDC 1919153) and 3a (CCDC 1919154).

Table 1 Selected optimization of the divergent synthesis

| Entry | Ligand | Mol% [L Cat.−1] | Additive | Yield 2a [%] (E : Z) | Yield 3a [%] (E : Z) |
|-------|--------|----------------|----------|-------------------|-------------------|
| 1     | —      | 0/10           | K2CO3    | 71 (90 : 10)      | 0 (—)            |
| 2     | PPh3   | 20/10          | K2CO3    | 64 (93 : 7)       | 0 (—)            |
| 3     | TTMP   | 20/10          | K2CO3    | 54 (91 : 9)       | 0 (—)            |
| 4     | (t-Bu)3P–BF4 | 20/10 | K2CO3 | 62 (<97 : 3) | 0 (—) |
| 5     | dppe   | 20/10          | K2CO3    | 34 (95 : 5)       | 0 (—)            |
| 6     | (PhO)2P | 20/10         | K2CO3    | 74 (<95 : 5)      | 0 (—)            |
| 7     | L      | 20/10          | KOAc     | 0 (—)             | 91 (95 : 5)      |
| 8     | L      | 10/5           | KOAc     | 0 (—)             | 85 (<95 : 5)     |
| 9     | L      | 10/5           | AcOH     | 0 (—)             | 93 (<95 : 5)     |
| 10    | L      | 5/2.5          | AcOH     | 0 (—)             | 93 (<95 : 5)     |
| 11    | L      | 10/5           | K2CO3    | 74 (<95 : 5)      | 0 (—)            |
| 12    | L      | 5/2.5          | K2CO3    | 75 (95 : 5)       | Trace (ND)       |

a A crystal structure was obtained for 1a (CCDC 1919153) and 3a (CCDC 1919154). b Ratio (E : Z) determined after purification. All other ratios determined from the 1H NMR spectrum of the crude reaction mixture. c Required further purification. Yield after recrystallization = 32% (entry 4), 20% (entry 6). d Average yield.
the substrate scope with respect to the boronic acid and the substituent at the stereogenic centre was investigated for both reactions. A range of arylboronic acids bearing electron-donating and withdrawing substituents were well-tolerated in the synthesis of the 1,3-dienes (Table 2). The sterically hindered nature of the ortho-substituted boronic acids likely impedes transmetallation\textsuperscript{\textregistered} (2c and 2f) and is well known to drive protodeboronation, the resulting boric acid likely catalyzing the unexpected formation of the corresponding vinylazolidinone in the case of 2c. In contrast, the highly electron-poor m- and p-nitrophenylboronic acids gave the highest yields (2d and 2e). An ester functionality, however, was not as well-tolerated (2i), though electron-poor boronic acids are known to undergo homocoupling\textsuperscript{\textregistered} and protodeboronation.\textsuperscript{\textregistered} Aryl substituents with halogen handles were able to be introduced (2g and 2h), though competitive oxidative addition to the C–Br bond may explain the lower yield of desired product in the case of 2g. Pleasingly, the scope was extended to heteroarylboronic acids (2k and 2l) and 1-phenylvinylboronic acid (2m), albeit in lower yield and stereoselectivity for the latter. Interestingly, n-butylboronic acid was successfully employed in the reaction (2n), which is significant considering the reduced reactivity of alkylboronic acids under Suzuki–Miyaura conditions (though protocols have been developed to accommodate these).\textsuperscript{\textregistered} With respect to the substituent at the stereoeric centre, the corresponding alanine, leucine and phenylalanine-derived 1,3-dienes were synthesized in moderate yield and excellent stereoselectivity, using m-nitrophenylboronic acid as the coupling partner (2o–q).\textsuperscript{\textregistered}

A range of arylboronic acids were also well-tolerated in the synthesis of the 5-vinylazolidinones (Table 3), with some notable differences in reactivity in comparison to the 1,3-diene synthesis. For example, while the electron-rich boronic acids (3a–e, including the sterically hindered ortho-substituted example 3e) furnished the corresponding products in good to excellent yields, the electron-poor examples were mostly unreactive (3d–f) apart from m-nitrophenylboronic acid (3e, where the resonance effect of the nitro group was less effective at the boron-substituted carbon), though a higher catalyst loading and resubjection to the reaction conditions was required for this reaction to proceed to completion.\textsuperscript{\textregistered} Aryl substituents with functionalizable handles such as halogens (3g and 3h) and an ester (3i) were introduced in high yields, as were heteroaryl groups (3k and 3l), with the benzo[b]thiophene analogue 3l furnished in almost quantitative yield. It was found that 1-phenylvinylboronic acid did not cleanly afford the corresponding product as in the 1,3-diene synthesis, however (E)-stylylboronic acid was well tolerated with excellent stereoselectivity (3m). In contrast to the 1,3-diene synthesis, n-butylboronic acid was unreactive (3n). With respect to the substituent at the stereoeric centre, the alanine, leucine and

| Table 2 | Substrate scope for the 1,3-diene synthesis\textsuperscript{\textregistered} |
| --- | --- |
| **R’-B(OH)\textsubscript{2} (1.5 equiv), L/Pd\textsubscript{2}bpy, CH\textsubscript{2}Cl\textsubscript{2} (10/1 mol %), toluene, rt. 24 h |  |
| 1 | 2 |
| 2a 4-Ome: 74% (>95:5) | 2a 4-Ome: 74% (>95:5) |
| 2b 3-Ome: 82% (>93:7) | 2b 3-Ome: 82% (>93:7) |
| 2c 2-Ome: 32% (>99:1)\textsuperscript{a} | 2c 2-Ome: 32% (>99:1)\textsuperscript{a} |
| 2d 4-NO\textsubscript{2} 79% (90:10) | 2d 4-NO\textsubscript{2} 79% (90:10) |
| 2e 3-NO\textsubscript{2} 80% (>99:1) | 2e 3-NO\textsubscript{2} 80% (>99:1) |
| 2f 2-NO\textsubscript{2} 47% (90:10) | 2f 2-NO\textsubscript{2} 47% (90:10) |
| 2g X = Br: 27% (>95:5) | 2g X = Br: 27% (>95:5) |
| 2h X = Cl: 66% (>95:5) | 2h X = Cl: 66% (>95:5) |
| 2i X = CO\textsubscript{2}Me: 34% (>95:5) | 2i X = CO\textsubscript{2}Me: 34% (>95:5) |
| 2j 61% (>95:5) | 2j 61% (>95:5) |
| 2k 58% (>95:5) | 2k 58% (>95:5) |
| 2l 76% (>99:1) | 2l 76% (>99:1) |
| 2m 43% (85:15) | 2m 43% (85:15) |
| 2n 30% (>95:5)\textsuperscript{f} | 2n 30% (>95:5)\textsuperscript{f} |
| 2o R = Me: 47% (>95:5) | 2o R = Me: 47% (>95:5) |
| 2p R = i-Pr: 58% (>95:5) | 2p R = i-Pr: 58% (>95:5) |
| 2q R = Br: 46% (>95:5) | 2q R = Br: 46% (>95:5) |

\textsuperscript{\textregistered} E : Z ratio shown in parentheses. \textsuperscript{a} Corresponding 5-vinylazolidinone 3e obtained as major product (83%, >95 : 5, E : Z). \textsuperscript{\textregistered} Inseparable mixture (86 : 14) of product and dimerized starting material 4 (Scheme 2B), respectively. Yield of 2f calculated via 1H NMR spectrum of the mixture. \textsuperscript{\textregistered} Required resubjection to the reaction conditions to consume starting material and enable purification. |
phenylalanine-derived products were able to be synthesized in high yield and stereoselectivity (3o–q).

The proposed reaction mechanisms for both syntheses are shown in Scheme 2A. The origin of divergence in the reactivity of the 5-allenylazolidinone arises from the formation of either intermediate I or V. During 1,3-diene formation, Pd(0) likely undergoes decarboxylative oxidative addition to the internal allene carbon, furnishing the Pd(II) intermediate I as the E stereoisomer, stabilized by a carbonate anion. This intermediate is thought to bind η⁴ rather than η₃, which is supported both in the literature and by the appearance of dimerized starting material 4 (Scheme 2B) during the synthesis of 2f, the dendraline core of 4 likely forming via the same η¹ coordination mode. In contrast, the mechanism for the formation of the 5-vinyloxazolidinone begins with oxidative addition of Pd(0) to AcOH to form complex IV, which undergoes a concerted hydropalladation to form the Pd(II) intermediate V. This mechanism is supported by mass spectrometry studies conducted by Guo and Ma, and previous work by Ma supports that the E stereochemistry arises from facially-selective coordination of Pd(II) opposite to the substituent at the stereogenic centre, and that the regioselectivity is controlled by delivery of the hydride to the relatively less sterically hindered terminal allene carbon (Scheme 2C). The mechanism of transmetalation differs between the two syntheses and this may explain the differing relative reactivity of the electron-poor arylboronic acids. Under basic conditions, the formation of the activated boronate II likely precedes transmetalation. It is well known that electron-poor arylboronic acids facilitate this pathway due to their increased Lewis acidity, and this is reflected in the high yields of the m- and p-nitrophenyl 1,3-dienes. The 5-vinyloxazolidinone synthesis, however, occurs in the absence of an added base and therefore a direct transmetalation likely occurs instead, in which the electron-poor nature of the nitrophenylboronic acids renders them unreactive. Both syntheses conclude with a reductive elimination, delivering the organic moiety to the 1,3-diene III or 5-vinyloxazolidinone VI and regenerating the Pd(0) catalyst. Critically, a control experiment showed that treatment of the 5-vinyloxazolidinone with the Pd(0)-catalyst system and K₂CO₃ did not result in formation of the 1,3-diene via elimination, indicating the 5-vinyloxazolidinone is not a precursor to the 1,3-diene.

As previously mentioned, the 5-vinyloxazolidinones may serve as structural analogues to existing oxazolidinone antibiotics, bearing the correct stereochemical scaffold and an unsaturated alkene moiety suitable for further functionalization. The 1,3-dienes, however, can act as powerful substrates in the Type I IMDA reaction to give more complex molecular scaffolds. In order to demonstrate this, a dienophile was tethered to the p-methoxyphenyl system 2a via N-alkylation with ethyl 2-(bromomethyl)acrylate (Scheme 3). It was found that the alkylation and subsequent IMDA reaction occurred in a one-pot

Scheme 2 (A) Proposed reaction mechanism for the divergent syntheses. (B) Proposed reaction mechanism for the formation of dimer 4. (C) Origin of E : Z stereoselectivity during hydropalladation.
fashion to furnish the isoindoline derivative 5a. The stereochemistry of the major diastereoisomer of this isoindoline system was determined via 1D NOE experiments and supported by computational data. The major diastereoisomer was able to be isolated via column chromatography after reducing the ethyl ester and introducing an O-tosyl group to the resulting alcohol to give derivative 5b. The 1D NOE experiments conducted on 5b revealed strong correlations between the bridgehead proton and the CH$_2$-OTs protons, indicating the formation of a cis-fused system. The strong correlation between the bridgehead proton and the isopropyl methyl protons indicated that both substituents were positioned on the same face, as did the lack of correlation between the existing stereogenic proton and the CH$_2$-OTs protons. The interpretation of the 1D NOE data was further supported by the interproton distances estimated by density functional theory calculations (see ESI†). The stereochemical outcome can be rationalized in terms of the proposed transition state TS1, the exo configuration of which is not uncommon in IMDA processes. This isoindoline core, including the stereochemical outcome, is structurally similar to that of the cytochalasin natural products (Scheme 1C), which are known to exhibit activity in actin polymerization and other cellular processes, giving heavy precedent to the medicinal relevance of these systems.

Conclusions

In summary, the Pd(0)-catalyzed cross-coupling of boronic acids to enantioenriched allenloxazolidinolones has been explored, giving rise to chiral 1,3-dienes and vinyl-oxazolidinones. Not only is this a divergent process that can be tightly controlled by a single switch in additive, but the allenloxazolidinolone itself is an entirely new building block that can now complement known related systems such as vinylaziridines and vinylloxazolidinones. A synthetic route to highly complex and stereodefined vinylloxazolidinolones has been established, which can now serve as potential analogues to existing oxazolidinone antibiotics. Furthermore, we have developed a route to stereodefined, N-tethered 1,3-dienes under mild conditions, the complex nature of which has left such systems underexplored in the literature, as they are inherently difficult to synthesize. We have demonstrated that these substrates can now be employed in the powerful intramolecular Diels–Alder reaction to form highly complex and medicinally relevant isoindoline scaffolds.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This research was supported in part by the Australian Government through the Australian Research Council’s Discovery Projects funding scheme (project DP180101332). The University of Wollongong is also acknowledged for generous partial funding of this research. The NCI National Facility is acknowledged for providing the computing time.

Notes and references

1 For a review on the general reactivity of vinylaziridines, see: H. Ohno, Chem. Rev., 2014, 114, 3703.
2 For examples of Pd-catalyzed cross-coupling reactions of vinylaziridines, see: (a) J. Kjellgren, J. Aydin, O. A. Wallner, I. V. Saltanova and K. J. Szabó, Chem.–Europ. J., 2005, 11, 5260; (b) J. Yin, T. Mekelburg and C. J. T. Hyland, Org. Biomol. Chem., 2014, 12, 9113.
3 For reviews on cycloaddition reactions of vinylaziridines, see: (a) A. L. Cardoso and T. M. V. D. Pinho e Melo, Eur. J. Org. Chem., 2012, 6479; (b) E. A. Ilardi and J. T. Njardarson, J. Org. Chem., 2013, 78, 9533; (c) J.-J. Feng and J. Zhang, ACS Catal., 2016, 6, 6651; (d) B. D. W. Allen, C. P. Lakeland and J. P. A. Harrity, Chem.–Europ. J., 2017, 23, 13830.
4 For a recent example from our group of a Pd-catalyzed [3 + 2] cycloaddition of N-tosyl-2-vinylaziridine to 3-nitroindoles to give a medicinally relevant pyrroloindoline scaffold, see: D. J. Rivinoja, Y. S. Gee, M. G. Gardiner, J. H. Ryan and C. J. T. H. Hyland, ACS Catal., 2017, 7, 1053.
5 For examples of Pd-catalyzed cycloaddition reactions of 5-vinloxazolidinolones, see: (a) J. G. Knight, P. A. Stoker, K. Tchabanenko, S. J. Harwood and K. W. M. Lawrie, Tetrahedron, 2008, 64, 3744; (b) K. Ohmatsu, N. Imagawa and T. Ooi, Nat. Chem., 2014, 6, 47; (c) K. Ohmatsu, S. Kawai, N. Imagawa and T. Ooi, ACS Catal., 2014, 4, 4304.
6 For reviews on allene reactivity, see: (a) S. Ma, Chem. Rev., 2005, 105, 2829; (b) S. Yu and S. Ma, Angew. Chem., Int. Ed., 2012, 51, 3074. For a recent example of a palladium-catalyzed reaction of allenes leading to a wide range of synthetic applications, see: (c) B. M. Trost, D. Zell, C. Hohn, G. Mata and A. Maruniak, Angew. Chem., Int. Ed., 2018, 57, 12916; Angew. Chem., 2018, 130, 13098.
7 F. Zamani, S. G. Pyne and C. J. T. Hyland, J. Org. Chem., 2017, 82, 6819.
8 F. Zamani, R. Babaahmadi, B. F. Yates, M. G. Gardiner, A. Ariafard, S. G. Pyne and C. J. T. Hyland, Angew. Chem., Int. Ed., 2019, 58, 2114; Angew. Chem., 2019, 131, 2136.
9 For reviews on Pd-catalyzed reactions of allenes, see: (a) R. Zimmer, C. U. Dinesh, E. Nandanand and F. A. Khan, Chem. Rev., 2000, 100, 3067; (b) T. Lechel, F. Pfrengle, H.-U. Reissig and R. Zimmer, ChemCatChem, 2013, 5, 2100.
10 For selected examples of Pd(0)-catalyzed reactions of allenylcarbinol derivatives with boronic acids to form 1,3-dienes, see: (a) T. Moriya, T. Furuuchi, N. Miyaura and A. Suzuki, Tetrahedron, 1994, 50, 7961; (b) M. Yoshida, T. Gotou and M. Ihara, Chem. Commun., 2004, 1124; (c) T. Liu, J. Dong, S.-J. Cao, L.-C. Guo and L. Wu, RSC Adv., 2014, 4, 61722; (d) Y.-Z. Chen, L. Zhang, A.-M. Lu, F. Yang and L. Wu, J. Org. Chem., 2015, 80, 673; (e) D. J. Lippincott, R. T. H. Linstadt, M. R. Maser, F. Gallou and B. H. Lipshutz, Org. Lett., 2018, 20, 4719.
11 For reviews on the intramolecular Diels–Alder reaction, see: (a) G. Brieger and J. N. Bennett, Chem. Rev., 1980, 80, 63; (b)
A. G. Fallis, Can. J. Chem., 1984, 62, 183; (c) D. Craig, Chem. Soc. Rev., 1987, 16, 187; (d) K.-I. Takao, R. Munakata and K.-I. Tadano, Chem. Rev., 2005, 105, 4779; (e) M. Juhl and D. Tanner, Chem. Soc. Rev., 2009, 38, 2983; (f) P. T. Parvatkar, H. K. Kadam and S. G. Tilve, Tetrahedron, 2014, 70, 2857.

12 A number of other synthetic routes to complex, chiral 1,3-diienes have previously been reported. For selected examples using rhodium catalysis, see: (a) P.-S. Wang, P. Liu, Y.-J. Zhai, H.-C. Lin, Z.-Y. Han and L.-Z. Gong, J. Am. Chem. Soc., 2015, 137, 12732; (b) S. E. Korkis, D. J. Burns and H. W. Lam, J. Am. Chem. Soc., 2016, 138, 12252; (c) J. D. Dooley and H. W. Lam, Chem.–Eur. J., 2018, 24, 4050; (d) B. Bai, J. Xia, F. Song, X. Li, B. Liu, L. Liu, G. Zheng, X. Yang, J. Sun and X. Li, Chem. Sci., 2019, 10, 3987. For an example using palladium catalysis, see: (e) G. T. Crisp and M. G. Gebauer, J. Org. Chem., 1996, 61, 8425. For an additional example, see ref. 6c. For an example using a chiral vinylstannane, see: (f) A. D. Brosius, L. E. Overman and L. Schwink, J. Am. Chem. Soc., 1999, 121, 700. For an example using a TiCl4-promoted electrophilic substitution of axially chiral allenes, see: (g) M. Ogasawara, K. Ueyama, T. Nagano, Y. Mizuhata and T. Hayashi, Org. Lett., 2003, 5, 217. For an example using ring-closing metathesis of enantioenriched enynes, see: (h) M. J. Garcia-Munoz, A. Sirvent, F. Foubelo and M. Yus, An. Acad. Bras. Cienc., 2018, 90, 1059. For selected transition metal-free examples, see: (i) M. J. Lilly and M. S. Sherburn, Chem. Commun., 1997, 967; (j) R. Fernández de la Pradilla, I. Colomer and A. Viso, Org. Lett., 2012, 14, 3068 (Overman rearrangement).

13 C.-R. Li, Q.-Q. Zhai, X.-K. Wang, X.-X. Hu, G.-Q. Li, W.-X. Zhang, J. Pang, X. Lu, H. Yuan, M. F. Gordeev, L.-T. Chen, X.-Y. Yang and X.-F. You, Antimicrob. Agents Chemother., 2014, 58, 2418.

14 For selected examples of Pd/AcOH-catalyzed addition of boronic acids to allenes, see: (a) C. Ho Oh, T. W. Ahn and R. Reddy, Chem. Commun., 2003, 2622; (b) S. Ma, N. Jiao and L. Ye, Chem.–Eur. J., 2003, 9, 6049; (c) S. Ma, H. Guo and F. Yu, J. Org. Chem., 2006, 71, 6634; (d) H. Guo and S. Ma, Synthesis, 2007, 17, 2731; (e) M. Yoshida, K. Matsuda, Y. Shoji, T. Gotou, M. Ihara and K. Shishido, Org. Lett., 2008, 10, 5183. For applications of this reactivity in the synthesis of natural products, see: (f) M. Yoshida, Y. Shoji and K. Shishido, Org. Lett., 2009, 6, 1441; (g) M. Yoshida, Y. Shoji and K. Shishido, Tetrahedron, 2010, 66, 5053; (h) M. Yoshida, T. Kasai, T. Mizuguchi and K. Namba, Synlett, 2014, 25, 1160.

15 The internal alkene proton adjacent to the aryl group exhibited a coupling constant of 15.6 Hz for the E isomer and 11.8 Hz for the Z isomer.

16 T. E. Barder, S. D. Walker, J. R. Martinelli and S. L. Buchwald, J. Am. Chem. Soc., 2003, 127, 4685.

17 M. S. Wong and X. L. Zhang, Tetrahedron Lett., 2001, 42, 4087.

18 H. G. Kuivila, J. F. Reuwer Jr and J. A. Mangravite, J. Am. Chem. Soc., 1964, 86, 2666.

19 For selected examples of Suzuki–Miyaura cross-coupling procedures accommodating alkylboronic acids and their derivatives, see: (a) J. H. Kirchhoff, M. R. Netherton, I. D. Hills and G. C. Fu, J. Am. Chem. Soc., 2002, 124, 13662; (b) I. Kondolf, H. Doucet and M. Santelli, Tetrahedron, 2004, 60, 3813; (c) L. Li, S. Zhao, A. Joshi-Pang, M. Diane and M. R. Biscoe, J. Am. Chem. Soc., 2014, 136, 14027. For a nickel-catalysed example, see: (d) Z. Lu, A. Wilsily and G. C. Fu, J. Am. Chem. Soc., 2011, 133, 8154.

20 It is suspected that these products were more prone to polymerization than the corresponding valine derivatives, as reflected in the isolated yields, despite the use of pH-neutral conditions during purification.

21 It should be noted that other examples required resubjection to the reaction conditions despite high conversion, as even traces of starting material were unable to be separated from the products via column chromatography. See ESi for details.

22 R. Qian, H. Guo, Y. Liao, Y. Guo and S. Ma, Angew. Chem., Int. Ed., 2005, 44, 4771; Angew. Chem., 117, 2005, 4849.

23 For a review covering mechanisms and factors affecting transmetalation, see: A. J. J. Lennox and G. C. Lloyd-Jones, Angew. Chem., Int. Ed., 2013, 52, 7362.

24 For an example of an intramolecular Diels–Alder reaction proceeding through an exo transition state, see: C. I. Turner, R. M. Williamson, M. N. Paddon-Row and M. S. Sherburn, J. Org. Chem., 2001, 66, 3963.

25 K. Scherlach, D. Boettger, N. Remme and C. Hertweck, Nat. Prod. Rep., 2010, 27, 869.