OL | Organic Letters

Letter

Cite This: Org. Lett. XXXX, XXX, XXX−XXX

pubs.acs.org/OrgLett

Synthesis and Homologation of an Azetidin-2-yl Boronic Ester with α-Lithioalkyl Triisopropylbenzoates

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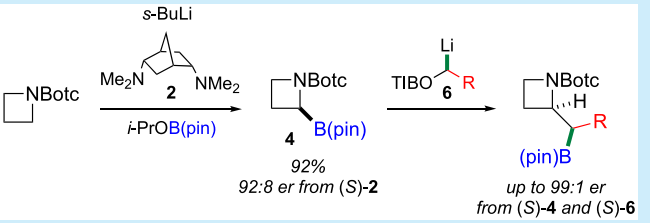
Supporting Information

ABSTRACT: An α-boryl azetidine, obtained by α-lithiation−borylation of *N*-Botc azetidine, undergoes reaction with α-triisopropylbenzoyloxy organolithiums to give homologated boronic esters that can be further oxidized, homologated, arylated, and deprotected to give a range of α-substituted azetidines. Scalemic α-boryl azetidine−α-triisopropylbenzoyloxy organolithium pairings show stereospecific reagent control, providing access to either diastereomeric series of homologated boronic esters with very high er's.



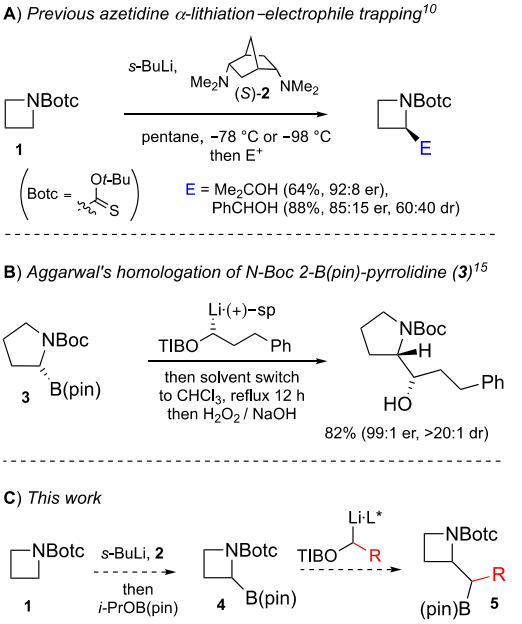
Saturated azacycles are ubiquitous structural motifs in natural products,[1] drug designs,[2] and organocatalysts.[3] Synthetic studies on pyrrolidines[4] and piperidines[5] have been extensive and aziridines[6] have also seen more attention compared with azetidines. However, with azetidine moieties increasingly emerging as constituents of highly bioactive compounds,[7] developing effective syntheses of substituted azetidines has become an important and growing field of chemistry.[8]

Our previous research on azetidine α-lithiation−electrophile trapping gave 2-substituted azetidines through the use of a thiopivaloyl or *tert*-butoxythiocarbonyl (Botc) *N*-protecting/activating group (Scheme 1, A).[9−11] The methodology enables direct diversifying α-functionalization of the azetidine moiety. Good enantioselectivities can be achieved by α-lithiation of *N*-Botc azetidine **1** in the presence of the chiral DIANANE ligand **2**[12] and trapping with certain electrophiles (e.g., Scheme 1, A).[10,13] However, reactions with aldehydes typically suffered from poor diastereoselectivity and enolizable aldehydes, such as hydrocinnamaldehyde, gave no conversion.

Boronic esters are established as highly versatile functional groups, known to undergo many stereospecific transformations.[14] In a recent synthesis of (−)-stemaphylline, Aggarwal, Leonori, and co-workers reported the preparation and some C−C bond-forming reactions of 2-B(pin)-pyrrolidine **3** (e.g., Scheme 1, B).[15] We considered that if access to 2-B(pin)-azetidine **4** was achievable (Scheme 1, C), then this could lead on to providing a broader range of α-functionalized azetidines than previously possible through direct α-lithiation−electrophile trapping. Here we communicate progress on these areas, in both racemic and asymmetric variants.

We began by applying our azetidine α-lithiation chemistry in a racemic sense with *N*-Botc azetidine **1** and TMEDA in THF[10] using *i*-PrOB(pin) as the electrophile; this gave boronic ester **4** in moderate yield (53%, Scheme 2). A factor contributing to the modest yield was likely product loss during purification by column chromatography. Despite testing a variety of different deactivated silica gels, chromatographed boronic ester **4** proved impossible to isolate without partial decomposition.[16] ^{1}H NMR analysis of the crude material indicated pinacol as a major impurity, which could not be easily removed via aqueous wash. It was reasoned that running the reaction in a more hydrophobic solvent than THF, such as pentane, could enable easier removal of impurities such as pinacol during aqueous workup. Conditions for lithiation/

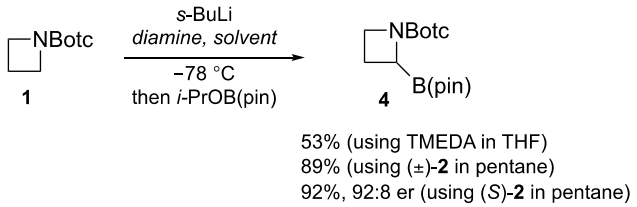
Scheme 1. α-Substitution of Azetidines and Pyrrolidines through Organolithium Chemistry



Received: October 31, 2019

DOI: 10.1021/acs.orglett.9b03901
Org. Lett. XXXX, XXX, XXX−XXX

Scheme 2. Racemic and Asymmetric α-Lithiation–Borylation of *N*-Botc Azetidine 1

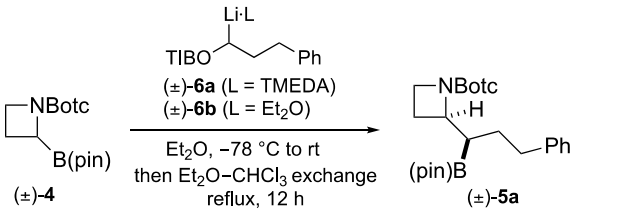


electrophile trapping in pentane using (*S*)-**2** as a ligand (Scheme 1, A)[10] were applied with (±)-**2**, to give 2-B(pin)-azetidine **4** in excellent yield (89%) and sufficiently high purity to negate the need for chromatographic purification. Following acid/base extraction, the diamine ligand could also be efficiently recovered (88%) and recycled.

Applying the above conditions using (*S*)-**2** gave the enantioenriched boronic ester (+)-**4** also in excellent yield (92%) and high enantioenrichment (92:8 er). The absolute configuration of boronic ester (+)-**4** was determined to be *S* by derivatization to an alcohol enantiomeric to that derived from (*S*)-azetidine-2-carboxylic acid.[16] The sense of asymmetric induction is opposite to that previously seen with MeI,[10] but the same as found for acetone and benzaldehyde.[13] These findings parallel observations by O'Brien and co-workers, who noted that the sense of asymmetric induction following lithiation of *N*-thiopivaloyl azetidine in the presence of (−)-sparteine is dependent on the nature of the electrophile.[17]

An attractive functionalization of boronic ester **4** would be homologation via ate complex formation with an α-triisopropylbenzoyloxy (TIBO) organolithium and subsequent 1,2-metalate rearrangement (cf. Scheme 1, B and C).[18] This should in principle allow assembly of two contiguous stereocenters with, ultimately, control arising from choice of reactant configurations. Homologations of α-to-nitrogen boronic esters have often been found to be difficult,[15,19] although such transformations can benefit from a solvent exchange to chloroform following ate complex formation to facilitate the migration step.[15,20] Following such a protocol, 2-B(pin)-azetidine (±)-**4** was reacted with α-lithiobenzoate (±)-**6a** (1.3 equiv) to give homologated boronic ester **5a** in moderate yield (45%, Scheme 3) as a readily separable mixture

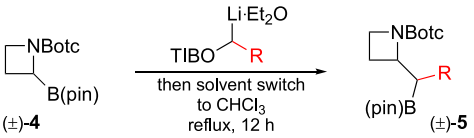
Scheme 3. Homologation of 2-B(pin)-azetidine (±)-4

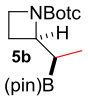
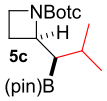
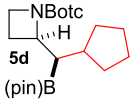
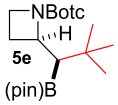
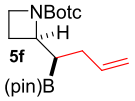
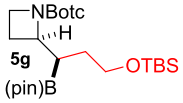
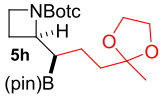


of diastereomers (74:26 dr, major diastereomer shown[16]). Using boronic ester **4** in slight excess (1.5 equiv) significantly improved the yield of **5a** (71%, 83:17 dr, Scheme 3). While these latter conditions would be appropriate when the benzoate is the more precious material, studies to establish an efficient method for homologation with boronic ester **4** as the limiting reagent were also undertaken. Aggarwal and co-workers previously employed stannanes in a diamine-free tin–lithium exchange approach to α-lithiated benzoates and carbamates.[21] This latter method has the benefit of reduced steric hindrance around α-TIBO organolithiums such as **6b**, as the only coordinating species on their generation is solvent (Et_2O). Pleasingly, following this strategy with 1.3 equiv of tributylstannyl-derived organolithium (±)-**6b** gave the homologated boronic ester **5a** in good yield (69%, 72:28 dr, Scheme 3).

Having established conditions for homologation of azetidine boronic ester **4** where it could be used as the limiting reagent, a study of substrate scope with respect to the stannyl-derived α-TIBO organolithium was carried out to examine reaction tolerance toward steric hindrance and functional groups. Reaction using a smaller α-TIBO ethyl stannane proceeded smoothly; a shorter reflux time of 3 h was sufficient to give

Table 1. Homologation Scope of 2-B(pin)-azetidine 4 with Stannane-Derived α-TIBO Organolithiums



Entry[a]	Homologated boronate **5**[b]	Yield	dr
1[c]	 **5b**	68%	56:44
2	 **5c**	65%	86:14
3	 **5d**	55%	72:28
4	 **5e**	0%[d]	n/a
5	 **5f**	50%	71:29
6[e]	 **5g**	65%	58:42
7[f]	 **5h**	56%	67:33

[a]Unless otherwise noted, (i) reactions used 2-B(pin)-azetidine **4** (0.42 mmol), stannane (0.55 mmol), 12 h reflux in $CHCl_3$ and (ii) amount of individual diastereomers isolated after chromatography used to give dr's and combined yields. [b]Major diastereomer shown (**5c,d,f–h** assigned by analogy to **5a,b**).[16] [c]3 h reflux. [d]Tin–B(pin) exchange observed (71%).[16] [e]Using 2-B(pin)-azetidine **4** (0.21 mmol) and stannane (0.28 mmol). [f]Yield for inseparable diastereomeric mixture and dr from ^{1}H NMR analysis.

DOI: 10.1021/acs.orglett.9b03901
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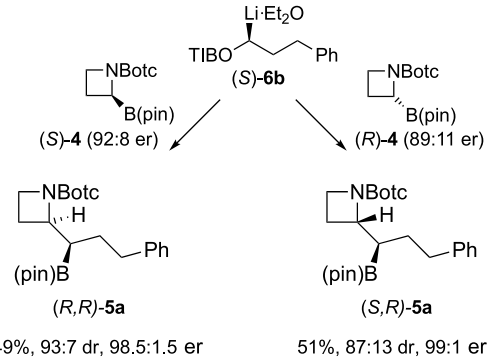
homologated boronic ester **5b** in 68% yield (Table 1, entry 1). Applying this shorter reflux time with α-TIBO organolithium **6b** also gave **5a** in 65% yield and unchanged dr. More sterically demanding β-branched stannanes remained effective, giving homologated isobutyl and methylcyclopentyl boronic esters **5c** and **5d** in good yields (Table 1, entries 2 and 3). However, further increase of the steric bulk at the β-position, using an α-TIBO neopentyl stannane, failed to give any of the desired homologated product **5e** (entry 4). In this last case, isolation of the B(pin) neopentyl benzoate (71%)[16] from overall tributyltin−B(pin) exchange suggests generation of the intermediate ate complex proceeded, but its collapse occurred without 1,2-metalate rearrangement. Homologations performed using stannanes bearing alkenyl, silyloxy and ketal functionality all proved viable (entries 5−7).

The observation of diastereoselectivity in the above boronate homologations suggests possible matched/mismatched effects depending on reactants configurations.[22] This could potentially render the asymmetric approach less effective in enabling access to the enantiomers of both diastereomers. To examine this aspect, separate reactions were carried out (Scheme 4) of *S* and *R* boronic ester **4** with α-

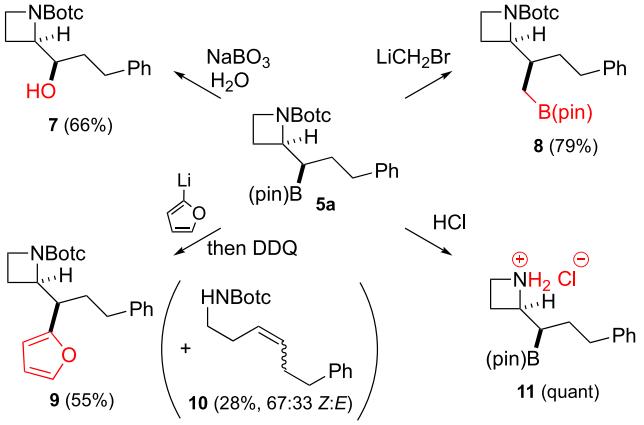
Scheme 4. Asymmetric Homologation of *S* and *R* Boronic Ester 4



lithiobenzoate (*S*)-**6b** [generated by stereoretentive Sn−Li exchange[23] from the corresponding *S* stannane (95:5 er)].[18] Pleasingly, both reactions proceeded to give homologated boronic esters (*R*,*R*)- and (*S*,*R*)-**5a**, respectively, in high diastereoselectivities and excellent enantioselectivities (up to 99:1 er), albeit with slightly lower yields compared with the racemic homologation. Similar selectivities (up to 99:1 er) and yields were also achieved for **5b**.[16]

To demonstrate synthetic versatility of the homologated boronic esters, further transformations of boronic ester **5a** were performed [Scheme 5, shown for (*R**,*R**)-**5a**; separately examined (*R**,*S**)-**5a** behaved similarly].[16] Oxidation of boronic ester **5a** to alcohol **7** was achieved using sodium perborate in good yield (66%). Matteson homologation, using dibromomethane, successfully gave the one-carbon homologated boronic ester **8** in high yield (79%). Arylation was achieved using DDQ as the activating electrophile[24] to give the furanylated product **9** in moderate yield (55%). This reaction also resulted in the formation of homoallylic amine **10** (28%). The latter likely arises from a competitive 1,2-elimination/ring-opening pathway, indicating that these C−B derivatizations, on a system with a β-electron withdrawing functional group incorporated within a strained ring, are not always straightforward. Finally, Botc deprotection of **5a** in ethereal HCl gave the

Scheme 5. Transformations of Homologated Boronic Ester (*R,*R**)-5a**



azetidinium chloride salt **11** (quant), which further demonstrates the utility/lability of this recently introduced protecting/activating group.[10] Whiting and co-workers have shown homoboroprolines to be effective organocatalysts for asymmetric aldol reactions,[19] and the current formation of a similar azetidinium boronic ester could lead to applications in this area.

In conclusion, we have developed a synthetic route to enantioenriched 2-B(pin)-azetidine **4**, and the latter has been converted to homologated azetidine boronic esters, by boronate formation with α-lithiobenzoates and subsequent 1,2-metalate rearrangement. This process can be performed with high diastereo- and enantioselectivity. The homologated boronic esters can be further transformed into α-substituted azetidines which cannot be accessed through direct α-lithiation−electrophile trapping chemistry.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.orglett.9b03901.

Full experimental procedures and characterization data (PDF)

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Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

We thank the EPSRC for studentship support (to P.K.D.).

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DOI: 10.1021/acs.orglett.9b03901
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