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Catalytic asymmetric hydrogenation of (Z)- α -dehydroamido boronate esters: direct route to alkyl-substituted α -amidoboronic esters†

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The direct catalytic asymmetric hydrogenation of (Z)- α -dehydroamino boronate esters was realized. Using this approach, a class of therapeutically relevant alkyl-substituted α -amidoboronic esters was easily synthesized in high yields with generally excellent enantioselectivities (up to 99% yield and 99% ee). The utility of the products has been demonstrated by transformation to their corresponding boronic acid derivatives by a Pd-catalyzed borylation reaction and an efficient synthesis of a potential intermediate of bortezomib. The clean, atom-economic and environment friendly nature of this catalytic asymmetric hydrogenation process would make this approach a new alternative for the production of alkyl-substituted α -amidoboronic esters of great potential in the area of organic synthesis and medicinal chemistry.

Since FDA approval of bortezomib¹ for the treatment of multiple myeloma, chiral α -aminoboronic acids have been recognized as key pharmacophores for the design of proteasome inhibitors.² The incorporation of chiral α -aminoboronic acid motifs at the C-terminal position of a peptide³ to develop potential clinical drug candidates has drawn increasing interest⁴ (Fig. 1). Meanwhile, chiral α -amidoboronic acids and their derivatives are useful synthetic building blocks for the stereospecific construction of chiral amine compounds.⁵ The biological and synthetic value of α -amidoboronic acids has led to considerable efforts for the development of efficient synthetic methods. However, up to now, limited transition-metal-catalyzed asymmetric approaches have been reported. The widely used strategies to synthesize these compounds are stepwise Matteson homologation/N-nucleophilic replacement,⁶ borylation of imines,⁷ and alkene functionalization.⁸ Recently, two other elegant approaches, Ni-catalyzed decarboxylative borylation of α -amino acid derivatives⁹ and enantiospecific borylation of lithiated α -N-Boc species,¹⁰ were reported by the Baran and Negishi groups, respectively. To the best of our knowledge, the majority of the methods relied on either stoichiometric amounts of chiral auxiliaries^{6,7a,b} or substrate-control strategies⁹ and most of these methods enable the construction of aryl-

substituted α -aminoboronic acids. Enantioselective methods to access unfunctionalized alkyl-substituted α -aminoboronic esters are still rarely developed and so far only two examples have been realized by the Miura^{8a} and Scheidt^{7f} groups, respectively. Considering that most therapeutically relevant α -amidoboronic acid fragments contain an alkyl subunit and the fact that the options for the synthesis of alkyl-substituted α -amidoboronic esters in an enantioselective manner are still rare, the development of other distinct approaches would be highly desirable. Herein, we report a new alternative to access these compounds by catalytic asymmetric hydrogenation of (Z)- α -dehydroamidoboronate esters. With this approach, the desired chiral alkyl-substituted α -amidoboronic esters could be obtained in high yields and generally excellent enantioselectivities (up to 99% yield and 99% ee) with simple purification.

Catalytic asymmetric hydrogenation of olefins is an atom-economic, environmentally friendly and clean process for the

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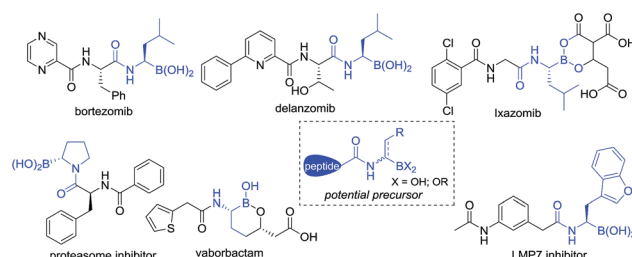
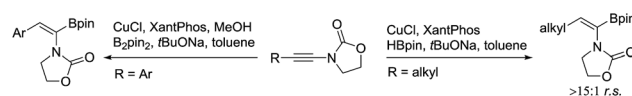


Fig. 1 Selected inhibitors containing chiral alkyl-substituted α -amidoboronic acids.



synthesis of valuable pharmaceuticals, agricultural compounds and feedstock chemicals.¹¹ Recently, hydrogenation of vinyl-boronic compounds has emerged for the preparation of chiral boronic compounds in a regiodefined manner.^{12,13} However, surprisingly α -dehydroamido boronate esters and their derivatives, as elegant precursors to access alkyl-substituted α -amidoboronic compounds, have never been used as substrates in asymmetric hydrogenation and remain a challenging project. To our knowledge, only one efficient hydrogenation approach to (1-halo-1-alkenyl) boronic esters was reported for indirect synthesis of alkyl-substituted α -aminoboronic esters but it was accompanied by inevitable de-halogenated by-products¹⁴ (Scheme 1). Given the catalytic efficiency and atom economy of the hydrogenation method, the development of a new direct hydrogenation approach to construct these important chiral alkyl-substituted α -amidoboronic esters would be very appealing.

The inspiration for our approach to the hydrogenation of α -dehydroamido boronates came from the molecular structures of relevant biologically active inhibitors containing alkyl-substituted α -amidoboronic acid fragments. Due to the limited stability of free α -aminoboronic acids, an electron-withdrawing carboxylic N-substituent is often required.¹⁵ Thus, we envisaged that *N*-carboxyl protected α -dehydroamido boronate esters could serve as a potential precursor for the synthesis of alkyl-substituted α -amidoboronic esters through Rh-catalyzed asymmetric hydrogenation of the C=C bond¹⁶ (Fig. 1), a strategically distinct approach to the construction of unfunctionalized alkyl-substituted α -amidoboronic esters. However, challenges still remain, including: (1) how to synthesize α -dehydroamido boronates; (2) the facile transmetalation process of the starting materials leading to deboronated by-products in the hydrogenation process;¹⁷ (3) the unknown stability of α -amidoboronic compounds in the presence of a transition-metal catalyst and hydrogen molecules. As part of our continuous efforts to develop efficient hydrogenation



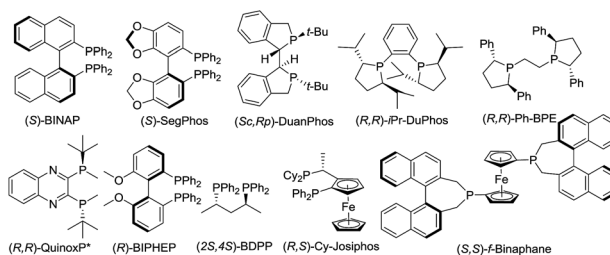
Scheme 2 Synthetic route to (*Z*)- α -dehydroamino boronates.

approaches to construct valuable motifs,¹⁸ here we present the results of the investigation to address the aforementioned challenges.

The desired aryl-substituted (*Z*)- α -dehydroamido boronates could be obtained by Cu-catalyzed regioselective hydroborylation of ynamide according to a previous report.¹⁹ However, different α/β -regioselectivity was observed for the preparation of alkyl-substituted (*Z*)- α -dehydroamido boronate esters and a new synthetic route was developed (Scheme 2, see the ESI† for details). Of note, (*Z*)- α -dehydroamido boronate esters should be purified with deactivated silica gel,^{7c} or else protodeborylation would occur readily with flash chromatography.

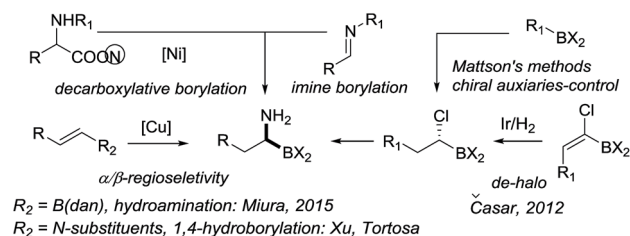
Table 1 Condition optimization for catalytic asymmetric hydrogenation of 1^a

Entry	Sub	Ligand	Conv. ^b (%)	ee ^c (%)
1 ^d	1a	(<i>Rc,Sp</i>)-DuanPhos	89	n.d.
2 ^e	1b	(<i>Rc,Sp</i>)-DuanPhos	>99	98
3	1c	(<i>Rc,Sp</i>)-DuanPhos	n.r.	n.d.
4	1b	(<i>R,R</i>)-QuinoxP*	>99	97
5	1b	(<i>S</i>)-SegPhos	>99	17
6	1b	(<i>S</i>)-BINAP	>99	10
7	1b	(<i>R,R</i>)- ⁱ Pr-DuPhos	>99	3
8	1b	(<i>R,S</i>)-Cy-JosiPhos	>99	14
9	1b	(<i>R</i>)-BIPHEP	>99	−30
10	1b	(<i>R,R</i>)-Ph-BPE	>99	−86
11	1b	(<i>S,S</i>)-f-Binaphane	>99	61
12	1b	(<i>2S,4S</i>)-BDPP	>99	59
13 ^{e,f,g}	1b	(<i>Rc,Sp</i>)-DuanPhos	>99(99)	99

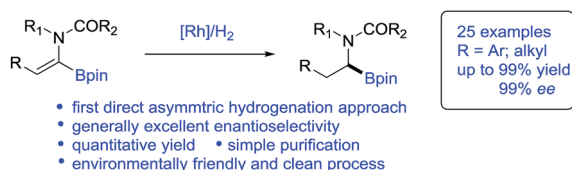


^a Unless otherwise mentioned, the reactions were performed with **1** (0.1 mmol), Rh(NBD)₂BF₄ (10 mol%), and a ligand (11 mol%) in 1.0 mL THF at 50 °C for 15 h. ^b Determined by crude ¹H NMR. ^c Determined with chiral HPLC. ^d The reaction was performed in ⁱPrOH. ^e Rh(NBD)₂BF₄ (1.0 mol%) and ligand (1.05 mol%) were used. ^f Isolated yield in parentheses. ^g 1,2-DCE was used as the solvent. Pin = 2,3-dimethyl-2,3-butanediol; dan = 1,8-diaminonaphthalene.

a) Previous approaches for alkyl-substituted α -aminoboronic esters



b) Our approaches: hydrogenation of alkene



Scheme 1 Approaches towards the synthesis of chiral alkyl-substituted α -aminoboronic esters.



In order to check the feasibility of our hypothesis, three substrates were prepared with $\text{Rh}(\text{NBD})_2\text{BF}_4$ and examined and our group prepared (*Rc,Sp*)-DuanPhos under 50 atm hydrogen pressure (Table 1). Gratifyingly, substrate **1b** reacted smoothly to provide the desired product **2b** in high yield and enantioselectivity (>99% conv., 98% ee, entry 2) whilst the reaction with substrate **1a** yielded a mixture of deborylation products and **1c** did not work at all (entries 1 and 3). Of note, we did not observe deborylation products with **1b** under the current reaction conditions and we did not select (*Z*)- α -dehydroamido boronic

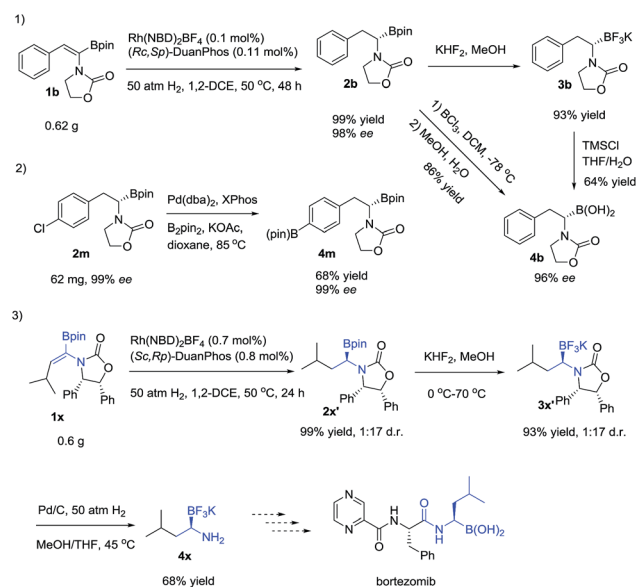
Table 2 Substrate scope.^{a,b,c}

1	2
2b 98% yield ^b 99% ee ^c	2d 96% yield 99% ee
2e 98% yield 95% ee	2f 96% yield 96% ee
2g 99% yield 99% ee	2h 95% yield 96% ee
2i 98% yield 99% ee	
2j 97% yield 99% ee	2k 96% yield 98% ee
2l 92% yield 90% ee	2m 99% yield 99% ee
2n 98% yield 99% ee	2o 94% yield 99% ee
2p 98% yield 99% ee	2q 92% yield 97% ee
2r 95% yield 99% ee	2s 99% yield 18:1 d.r. ^d
2t 99% yield 2:1 d.r. ^d	2u 97% yield 55% ee
2v (<i>Rc, Sp</i>)-DuanPhos: 99% yield, 26:1 d.r. ^d (<i>Sc, Rp</i>)-DuanPhos: 99% yield, 1:7 d.r. ^d	2w (<i>Rc, Sp</i>)-DuanPhos: 99% yield, 7:1 d.r. ^d (<i>Sc, Rp</i>)-DuanPhos: 99% yield, 1:27 d.r. ^d
2x (<i>Rc, Sp</i>)-DuanPhos: 99% yield, 44:1 d.r. ^d (<i>Sc, Rp</i>)-DuanPhos: 99% yield, 1:17 d.r. ^d	2y (<i>Rc, Sp</i>)-DuanPhos: 99% yield, 40:1 d.r. ^d (<i>Sc, Rp</i>)-DuanPhos: 99% yield, 1:11 d.r. ^d
2z 92% yield 57% ee	1r n.r. 1s n.r. 1t Boc n.r.

^a Unless otherwise mentioned, the reactions were performed with **1** (0.1 mmol), $\text{Rh}(\text{NBD})_2\text{BF}_4$ (1.0 mol%), and a ligand (1.05 mol%) in 1.0 mL 1,2-DCE at 50 °C under 50 atm H_2 for 15 h. ^b Isolated yield. ^c Determined with chiral HPLC. ^d Determined by crude ^1H NMR.

acid **1a** as the model substrate because of its poor solubility in most solvents. Then, a variety of chiral diphosphine ligands were investigated along with $\text{Rh}(\text{NBD})_2\text{BF}_4$ and the results are shown in Table 1. In most cases, the reaction proceeded smoothly to furnish the desired products and the best results were obtained when (*Rc,Sp*)-DuanPhos was used as the ligand (entries 2 and 4–12). Poor results were obtained with axially bidentate phosphine ligands (entries 5, 6 and 9). (*R,R*)-QuinoxP* and (*R,R*)-Ph-BPE also gave good conversion with a slightly decreased ee whilst (*R,R*)-ⁱPr-DuPhos exhibited poor results (entries 4, 7 and 10). Subsequent solvent screening revealed that the desired products could be obtained in most of the solvents and 1,2-DCE was the best solvent. (Entry 13, see the ESI†).

With the optimized reaction conditions in hand, a series of (*Z*)- α -dehydroamido boronate esters were tested and the results are summarized in Table 2. All the substrates reacted smoothly to give the corresponding alkyl-substituted α -amidoboronates in high yields with good to excellent enantioselectivities (**2b**, **2d–2r**, and **2u**, 99% yield, 57–99% ee). Alkyl-substituted (*Z*)- α -dehydroamido boronate esters were well tolerated in the current reaction, providing the corresponding α -amidoboronates in high yields and excellent enantioselectivities (**2d–2i**, 99% yield, 96–99% ee). Aryl-substituted (*Z*)- α -dehydroamido boronate esters with electron-donating (**2j–l**, **2n** and **2p–r**) and withdrawing (**2m** and **2o**) substituents could also give the desired products in excellent yield with excellent enantioselectivities (90–99% ee). The *ortho*-methyl-substituted substrate **1r** reacted smoothly to give the desired product with excellent enantioselectivity, but the 2,6-dimethyl-substituted substrate **1z** could not react at all. Functional groups such as ether, halo and benzyl were well tolerated in the current reaction (**2k**, **2l**, **2m** and **2o–q**). Replacement of the N-substituents with acyclic carbamate was also tolerated but with a decreased ee (**2u** and **2z**). Substrates containing a chiral oxazolidin-2-one unit bearing bulky Ph-substituents around the nitrogen and oxygen were also



Scheme 3 Scale-up synthesis and synthetic utility.



competent, yielding the desired products with good to excellent diastereoselectivities (**2s**, **2t**, and **2v–y**). Of note, the substrate **1s** bearing an N-Ms substituent and the cyclic substrate **1t** did not work in the current reaction. The absolute configuration of generated α -amidoboronates was assigned as (*S*) by X-ray crystallographic analysis of **2i** (Scheme 3).²⁰

To demonstrate the utility of the products, a scale-up reaction (0.62 g) was successfully performed with 0.1 mol% catalytic loading, giving **2b** in 99% yield and 98% ee, and **2b** could be easily transformed to a more stable α -amidoborate **3b** with KHF₂,^{6d,21} followed by hydrolysis with TMSCl to yield α -amido boronic acid **4b** in 46% yield,²² which could also be obtained from **2b** by treating it with BCl₃ in 84% yield, without loss of the optical purity.^{8b} **2m** could easily be transformed to **4m** in 68% yield by a Pd-catalyzed borylation reaction. Meanwhile, after hydrogenation of **1x** to **2x'** and transformation of **2x'** to its trifluoroborate derivative **3x'**, removal of the benzyl group of **3x'** with Pd/C under hydrogenation conditions²³ yielded the primary α -aminoborate **4x** in 62% yield in three steps, which could serve as a potential precursor¹⁵ to synthesize bortezomib.

Conclusions

In summary, we have presented a strategically distinct alternative for the direct synthesis of chiral alkyl-substituted α -amidoboronates by Rh-catalyzed asymmetric hydrogenation. A series of (*Z*)- α -dehydroamido boronate esters could be hydrogenated to the corresponding alkyl-substituted α -amidoboronic esters in excellent yields with generally excellent enantioselectivities. The boronic acid derivatives and the potential precursor of bortezomib could be facilely obtained with this approach. Additional asymmetric transformations of the alkyl-substituted α -amidoboronates are underway in our laboratory.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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