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Copper(i)-catalyzed enantioselective hydroboration of cyclopropenes: facile synthesis of optically active cyclopropylboronates†

Bing Tian, ‡^{a,b} Qiang Liu, ‡^{a,b} Xiaofeng Tong, ^b Ping Tian*^a and Guo-Qiang Lin*^a

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Copper(i)-catalyzed enantioselective hydroboration of 3-aryl substituted cyclopropene-3-carboxylate is described, providing chiral cyclopropylboronates with excellent enantioselectivities (89–95% ee) in moderate to high yields (55–86%). The non-directing effect of the ester group was observed, and the reaction proceeded with solely *trans*-selectivity. The chiral boronates could be conveniently converted into chiral 1,2-diaryl substituted cyclopropane derivatives.

Introduction

The chiral cyclopropane framework represents the smallest carbocycles existing in a wide range of naturally-occurring compounds, chiral drugs, and insecticides, for instance, (+)-Coronatine, Saxagliptin (Onglyza®), EBC-219, Milnacipran, Deltamethrin, and (+)-Tranylcypromine (Fig. 1). These three-membered carbocycles, due to their unique structural and electronic properties, serve as extremely significant versatile building blocks in organic synthesis. Thus, a few interest-

CH₃
(+)-Coronatine

Saxagliptin (Onglyza®)

H₂N

CH₃
(-1)-Coronatine

Saxagliptin (Onglyza®)

EBC-219

H₂N

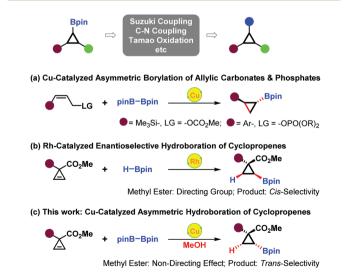
(+)-Tranvlcvpromine

Fig. 1 Cyclopropane-containing natural products, chiral drugs and insecticides.

‡These authors contributed equally to this work.

ing and characteristic transformations have continually emerged.⁹ Owing to their important biological activities and wide applications in organic chemistry, much attention has been paid to their efficient enantioselective syntheses.

Through Suzuki-Miyaura coupling, C-N coupling, Tamao oxidation reaction, *etc.*, cyclopropylboronates could be readily converted into structurally and functionally diverse cyclopropanes.¹⁰ Thus, efficient enantioselective synthesis of optically active cyclopropylboronates has gradually become a spotlight. Recently, Ito and co-workers successfully established copper(1)-catalyzed asymmetric cyclopropanation reactions of allylic phosphates and carbonates with bis(pinacolato)diboron (B₂pin₂), affording optically active *trans*-silyl- and *trans*-aryl-substituted cyclopropylboronates (Scheme 1a).^{11,12} Gevorgyan



Scheme 1 Enantioselective synthesis of optically active cyclopropylboronates.

^aKey Laboratory of Synthetic Chemistry of Natural Substances, Shanghai Institute of Organic Chemistry, Chinese Academy of Sciences, 345 Lingling Road, Shanghai 200032, China. E-mail: tianping@sioc.ac.cn, lingq@sioc.ac.cn; Tel: (+86) 21-54925081

^bShanghai Key Laboratory of Functional Materials Chemistry, East China University of Science and Technology, 130 Meilong Road, Shanghai 200237, China

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and co-workers described rhodium-catalyzed asymmetric hydroboration of 3,3-disubstituted cyclopropenes, directly constructing enantiopure 2,2-disubstituted cyclopropylboronates. The directing effect of the ester group was found to be necessary for achieving cis-selectivity and high enantioselectivity (Scheme 1b). 13,14 Herein, we present our findings in copper(1)catalyzed asymmetric hydroboration of 3,3-disubstituted cyclopropenes. 15 Interestingly, the non-directing effect of the ester group was observed in this case, and the reaction proceeded with solely trans-selectivity (Scheme 1c).

Results and discussion

At the outset, a set of representative chiral phosphorus ligands were investigated for the Cu-catalyzed asymmetric hydroboration of the cyclopropene substrate 1f, and the screening results are summarized in Table 1. The chiral bisphosphine ligand, (R,S_p) -Josiphos (L1), has been successfully employed in the

Table 1 Initial evaluation of various ligands and solvents^a

CI CO₂Me + B₂Pin₂ CO₂Me + B₂Pin₂ (2 equiv)
$$\frac{L^*(12 \text{ mol}\%)}{\text{MeOH}(2 \text{ equiv})}$$
 Solvent, rt $\frac{L^*(12 \text{ mol}\%)}{\text{MeOH}(2 \text{ equiv})}$ Solvent, rt $\frac{L^*(12 \text{ mol}\%)}{\text{MeOH}(2$

Entry	L*	Solvent	Time (h)	$Yield^{b}$ (%)	ee ^c (%)
1	L1	Toluene	6	62	40
2	L2	Toluene	8	16	22
3	L3	Toluene	8	30	51
4	L4	Toluene	6	75	94
5	L5	Toluene	8	58	93
6	L6	Toluene	10	85	-89
7	L7	Toluene	12	60	93
8	L8	Toluene	16	40	79
9^d	L4	Toluene	24	46	90
10	L4	THF	6	44	84
11	L4	DCM	6	32	94
12^e	L4	Toluene	6	80	95
13^f	L4	Toluene	6	78	94

^aThe reaction was carried out with 1f (0.15 mmol), B₂Pin₂ (2, 0.3 mmol), CuCl (10 mol%), chiral ligand (L*, 12 mol%) and NaOtBu (11 mol%) in anhydrous toluene (1.0 mL) at room temperature under a N2 atmosphere, unless otherwise noted. bYield of the isolated product. c Determined by HPLC analysis using a chiral stationary phase. d At 0 o C. e L4 (15 mol%) was used. f L4 (20 mol %) was used. $B_2Pin_2 = bis(pinacolato)diboron$.

Cu-catalyzed asymmetric conjugate hydroboration reaction of α,β-unsaturated compounds.16 However, only 62% yield and 40% ee were observed in our hydroboration (Table 1, entry 1). Phosphoramidite ((R)-MonoPhos, L2)¹⁷ and (R)-MOP (L3) ligands were subsequently subjected to this reaction, but no promising outcomes were obtained (Table 1, entries 2 and 3). To our delight, the ligand (R)-BINAP (L4) could dramatically improve the yield and ee of hydroboration product 3f to 75% and 94%, respectively (Table 1, entry 4). Several electronically different bisphosphine ligands (L5-L8) were applied in this reaction, but no better results were achieved (Table 1, entries 5-8).

Next, the reaction temperature and the solvent were investigated to further improve the enantioselectivity. Unfortunately, they led to different levels of erosion in yields and ee values (Table 1, entries 9–11). Increasing the ligand loading to 15 mol% resulted in a slight improvement of both yield and ee values (Table 1, entry 12). However, further increasing the ligand loading failed to give better results (Table 1, entry 13).

With the optimal reaction conditions identified, various aryl-substituted cyclopropenes were investigated, and the results are summarized in Table 2. All 4-substituted phenyl substrates, regardless of the electron-donating or electron-

Table 2 Substrate scope of various aryl-substituted cyclopropenes^a

^a The reaction was carried out with 1 (0.15 mmol), B₂Pin₂ (2, 0.3 mmol), CuCl (10 mol%), (R)-BINAP (L4, 15 mol%) and NaOtBu (11 mol%) in anhydrous toluene (1.0 mL) at room temperature under a N_2 atmosphere. Beaction time. Yield of the isolated product. ^d Determined by HPLC analysis using a chiral stationary phase.

withdrawing properties of the substituent at the phenyl ring, afforded the hydroboration products in moderate to high yields (55-86%) and with high to excellent enantioselectivities (89-95% ee, Table 2, entries 1-7). Interestingly, p-, m-, and o-bromophenyl substituted cyclopropene substrates (1g, 1h, and 1i) gave almost the same yields with high to excellent levels of enantioselectivities (Table 2, entries 7-9). As for 2-naphthyl and disubstituted phenyl substrates, the hydroboration reaction also proceeded smoothly with high yields and excellent enantioselectivities (Table 2, entries 10-12). In general, cyclopropene substrates bearing electron-withdrawing phenyl substituents provided better yields (Table 2, entries 1-3 vs. 4-6, 12).

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Given the highly enantioselective nature of this hydroboration reaction, the methyl substituted substrates 1n and 10 were tested under the standard conditions. Unfortunately, no desired products were observed, indicating that the α-substituent played an important role in the cyclopropene reactivity (Scheme 2, eqn (1) and (2)). As for the diester substrate 1p, the hydroboration reaction readily occurred with excellent enantioselectivity, albeit in a lower yield. This was partially attributed to the decomposition of the starting material (Scheme 2, eqn (3)).

The relative configuration of hydroboration products 3 was determined using NOE interactions; for example, the NOE interactions between the aryl group and the boronate group in 3g and 3j clearly revealed that both of them were on the same side of the cyclopropane plane (Fig. 2). Thus trans-cyclopropylboronates were achieved in this Cu(1)-catalyzed asymmetric hydroboration of cyclopropenes.

To determine the absolute configurations of the hydroboration product 3a in Table 2,18 we converted cyclopropylboronate

B₂Pin₂ (1)(2 equiv) BPin Standard Condition B₂Pin₂ (2 equiv) BPin Standard Condition B₂Pin₂ (2 equiv) 'BPin 3p (39%, 95% ee)

Scheme 2 Cu-catalyzed asymmetric hydroboration of cyclopropenes 1n. 1o and 1p.

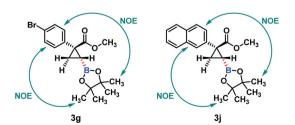


Fig. 2 The NOE interactions between the aryl group and the boronate group in 3g and 3j.

3a, through Suzuki-Miyaura coupling with iodobenzene (4), into a known compound (1R,2S)-5 in almost quantitative yield with no loss of the enantiomeric excess. 19 Thus, the absolute configuration of cyclopropylboronate 3a was unambiguously assigned as 1R,2R. The absolute configurations of other hydroboration products in Table 2 were assigned on the basis of their chemical correlation with (1R,2R)-3a (Scheme 3).

To probe the 'hydrogen' source of this hydroboration reaction, [D4]-methanol experiment was investigated. cis-Deuterated product 3a (50%) was observed, suggesting that the proton partially came from methanol and this hydroboration reaction was a syn-addition process (Scheme 4).

Piecing together the above details and preceding results,²⁰ a plausible reaction mechanism is proposed in Fig. 3. Initiation of the reaction through the transmetallation of a (pinacolato)boron group (BPin) from boron to copper species A generated the borylated copper B, which subsequently underwent syn-addition from the aryl group side²¹ to the double bond of the cyclopropene substrate 1 to afford the borylated cyclopropyl-copper intermediate D. The intermediate D

Scheme 3 Determining the absolute configuration of cyclopropylboronate 3a

Scheme 4 [D4]-Methanol experiment.

Proposed mechanism.

was readily protonated by trace water or methanol to regenerate A and liberate the trans-product 3. Due to the bigger steric hindrance of the methyl ester group (C vs. E), the weak coordination between copper and carboxyl groups could not overcome this energy barrier. Therefore, the cis-product was not observed.

Conclusions

In summary, copper-catalyzed asymmetric hydroboration reaction of 3-aryl, 3-methylester substituted cyclopropenes has been successfully established. This reaction proceeded smoothly at room temperature, affording optically active transcyclopropylboronates with excellent enantioselectivities (89-95% ee) in moderate to high yields (55-86%). The nondirecting effect of the methylester group was observed and this method was actually complimentary to the earlier reported cisborylated cyclopropane products through rhodium catalysis. The chiral boronates could be readily transformed to chiral 1,2-diaryl substituted cyclopropanes through Suzuki-Miyaura coupling reaction. Further studies on the applications of cyclopropylboronates are in progress in our laboratories.

Experimental section

General information

All solvents were dried before use by following the standard procedures. Unless otherwise indicated, all starting materials purchased from commercial suppliers were used without further purification. The 1H and 13C NMR spectra were recorded on a Bruker AV 400 MHz in the indicated solvents. Chemical shifts are reported in δ (ppm) referenced to the internal standard TMS for 1 H NMR and to CDCl₃ (δ = 77.10 ppm) for ¹³C NMR. Coupling constants (*J*) are quoted in Hz. Optical rotations were measured on a JASCO P-1030 polarimeter. IR spectra were recorded on a Nicolet iN 10 MX. ESI mass spectra were recorded on an Agilent 1200/G6100A. HRMS of boron-containing compounds is based on ¹⁰B. For the preparation of substrates 4a, see the ESI.†

General procedure for Cu-catalyzed hydroboration of 3,3-disubstituted cyclopropenes

A dried Schlenk flask was charged with CuCl (1.5 mg, 0.015 mmol, 10 mol%), (R)-(+)-BINAP (14 mg, 0.0225 mmol, 15 mol%), B₂pin₂ (2, 76.2 mg, 0.3 mmol, 2.0 equiv.), NaOtBu (1.6 mg, 0.0165 mmol, 11 mol%) and anhydrous toluene (1.0 mL) under a nitrogen atmosphere. After the mixture was stirred at room temperature for 40 min, a solution of cyclopropene 1 (0.15 mmol) in anhydrous toluene (0.5 mL) was added, followed by anhydrous MeOH (12.2 µL, 0.30 mmol, 2.0 equiv.). The resulting mixture was stirred at room temperature for the time indicated in Table 2, then filtered through Celite®, and concentrated in vacuo. The residue was purified by silica gel (300-400 mesh) column chromatography using hexane-ethyl acetate (15:1) as an eluent to afford the desired product 3.

1-phenyl-2-(4,4,5,5-tetramethyl-1,3,2-di-(1R,2R)-Methyl oxaborolan-2-yl)cyclopropanecarboxylate (3a). Colorless oil. 29.9 mg, 66% yield. $[\alpha]_D^{28}$ –206.8 (c 1.0, CHCl₃) for 94% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.35–7.21 (m, 5H), 3.60 (s, 3H), 1.70 (dd, J = 10.4 Hz, 3.2 Hz, 1H), 1.62 (dd, J = 8.0 Hz, 3.2 Hz, 1H), 1.28 (dd, I = 10.0 Hz, 8.0 Hz, 1H), 1.05 (s, 6H), 0.82 (s, 6H); 13 C NMR (75 MHz, CDCl₃) δ (ppm) 174.96, 137.76, 131.17 (2C), 127.87 (2C), 127.15, 83.37 (2C), 52.56, 33.83, 24.89 (2C), 24.44 (2C), 18.78. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: $[M + Na]^{\oplus}$ 325.1; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{23}^{10}BO_4Na^{\oplus}$ 324.1618, found 324.1614; IR (KBr) ν (cm⁻¹) 3451, 3086, 3047, 3027, 2979, 2954, 1961, 1726, 1602, 1429, 1372, 1264, 1166, 1142, 1062, 971, 858, 733, 698, 637, 503; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; *n*-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 6.4 min (R,R-isomer), 8.2 min (S,S-isomer).

(1R,2R)-Methyl 2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2vl)-1-p-tolylcyclopropanecarboxylate (3b). Colorless 26.1 mg, 55% yield. $[\alpha]_D^{26}$ –158.5 (c 1.0, CHCl₃) for 94% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.30–7.21 (m, 2H), 7.07 (d, J =7.6 Hz, 2H), 3.60 (s, 3H), 2.31 (s, 3H), 1.68 (dd, J = 10.0 Hz, 3.2 Hz, 1H), 1.58 (dd, J = 8.4 Hz, 3.2 Hz, 1H), 1.32-1.20 (m, 1H), 1.06 (s, 6H), 0.84 (s, 6H); 13 C NMR (100 MHz, CDCl₃) δ (ppm) 175.16, 136.74, 134.74, 131.01 (2C), 128.56 (2C), 83.37 (2C), 52.58, 33.43, 24.89 (2C), 24.45 (2C), 21.21, 18.86. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + Na][⊕] 339.2; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{18}H_{25}^{-10}BO_4Na^{\oplus}$ 338.1774, found 338.1758; IR (KBr) ν (cm⁻¹) 2978, 2951, 2924, 1723, 1515, 1436, 1410, 1371, 1329, 1285, 1263, 1215, 1165, 1142, 963, 858, 821, 751, 583, 504; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane-ipropanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 6.7 min (*R*,*R*-isomer), 7.8 min (*S*,*S*-isomer).

(1R,2R)-Methyl 1-(4-methoxyphenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3c). Colorless oil. 28.9 mg, 58% yield. $[\alpha]_D^{26}$ –190.1 (c 1.0, CHCl₃) for 93% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.25 (d, J = 8.8 Hz, 2H), 6.81 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 3.60 (s, 3H), 1.69 (dd, J =10.0 Hz, 3.2 Hz, 1H), 1.57 (dd, *J* = 8.0 Hz, 3.2 Hz, 1H), 1.24 (dd, $J = 10.0 \text{ Hz}, 8.0 \text{ Hz}, 1\text{H}, 1.07 (s, 6\text{H}), 0.86 (s, 6\text{H}); ^{13}\text{C NMR}$ (100 MHz, CDCl₃) δ (ppm) 175.08, 158.65, 132.08 (2C), 129.95, 113.20 (2C), 83.26 (2C), 55.26, 52.42, 32.93, 24.87 (2C), 24.36 (2C), 18.62. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: $[M + H]^{\oplus}$ 333.3; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{18}H_{25}^{10}BO_5Na^{\oplus}$ 354.1724, found 354.1729; IR (KBr) ν (cm⁻¹) 3542, 2979, 2952, 2837, 1723, 1614, 1582, 1517, 1440, 1409, 1331, 1264, 1247, 1165, 1143, 1034, 858, 834, 689, 548; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; *n*-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 9.6 min (R,R-isomer), 12.4 min (S,S-isomer).

(1R,2R)-Methyl 2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2yl)-1-(4-(trifluoromethyl)phenyl)cyclopropanecarboxylate (3d). White semisolid. 47.7 mg, 86% yield. $[\alpha]_{D}^{25}$ -128.2 (c 1.0,

CHCl₃) for 92% ee; 1 H NMR (400 MHz, CDCl₃) δ (ppm) 7.54 (d, J = 8.0 Hz, 2H, 7.45 (d, J = 8.0 Hz, 2H), 3.62 (s, 3H), 1.76 (dd, J = 10.0 Hz, 3.2 Hz, 1H), 1.64 (dd, J = 8.4 Hz, 3.2 Hz, 1H), 1.31 (dd, J = 10.0 Hz, 8.4 Hz, 1H), 1.04 (s, 6H), 0.81 (s, 6H);¹³C NMR (100 MHz, CDCl₃) δ (ppm) 174.13, 142.00, 131.58 (2C), 129.45, 124.79 (q, $J_{CF} = 3.5$ Hz, 2C), 83.54 (2C), 52.68, 33.69, 24.77 (2C), 24.38 (2C), 18.82. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) EI-MS, 370 (M. $^{\oplus}$), 355 (M $^{\oplus}$ – CH₃), 312 (M[⊕] - CO₂CH₃). HRMS (FTMS-EI) calcd for C₁₈H₂₂¹⁰BF₃O₄ $(M.^{\oplus})$ 369.1600, found 369.1597; IR (KBr) ν (cm⁻¹) 3430, 2980, 1726, 1607, 1514, 1437, 1372, 1332, 1287, 1263, 1223, 1165, 1143, 1102, 971, 858, 837, 689, 579, 543; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; *n*-hexane-i-propanol = 99/1; flow rate = 1.0 mL min⁻¹; retention time: 5.4 min (R,R-isomer), 7.4 min (S,S-isomer).

(1R,2R)-Methyl 1-(4-fluorophenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3e). White semisolid. 39.8 mg, 83% yield. $[\alpha]_D^{26}$ -163.0 (c 1.0, CHCl₃) for 95% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.32–7.28 (m, 2H), 7.00-6.92 (m, 2H), 3.60 (s, 3H), 1.72 (dd, J = 10.0 Hz, 3.2 Hz, 1H), 1.58 (dd, J = 8.0 Hz, 3.2 Hz, 1H), 1.26 (dd, J = 10.0 Hz, 8.0 Hz, 1H), 1.07 (s, 6H), 0.86 (s, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm) 174.61, 161.92 (d, $J_{\rm CF}$ = 244.4 Hz), 160.70, 133.61 (d, $J_{\rm CF}$ = 3.0 Hz), 132.66 (d, $J_{\rm CF}$ = 8.4 Hz, 2C), 114.56 (d, $J_{\rm CF}$ = 21.3 Hz, 2C), 83.36 (2C), 52.46, 33.01, 24.81 (2C), 24.34 (2C), 18.86. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: $[M + H]^{\oplus}$ 321.0; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{22}^{10}BFO_4Na^{\oplus}$ 342.1524, found 342.1519; IR (KBr) ν (cm⁻¹) 2983, 1960, 1720, 1618, 1430, 1392, 1382, 1327, 1296, 1268, 1165, 1141, 1115, 1064, 1018, 877, 837, 765, 608; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; *n*-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 4.5 min (R,R-isomer), 5.9 min (S,S-isomer).

(1R,2R)-Methyl 1-(4-chlorophenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3f). White semisolid. 40.3 mg, 80% yield. $[\alpha]_D^{26}$ -175.7 (c 1.0, CHCl₃) for 95% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.29–7.23 (m, 4H), 3.60 (s, 3H), 1.72 (dd, J = 10.0 Hz, 3.2 Hz, 1H), 1.57 (dd, J = 10.0 Hz, 1H), 1.57 8.0 Hz, 3.2 Hz, 1H), 1.27 (dd, J = 10.0 Hz, 8.0 Hz, 1H), 1.07 (s, 6H), 0.86 (s, 6H); 13 C NMR (100 MHz, CDCl₃) δ (ppm) 174.36, 136.35, 132.86, 132.47 (2C), 127.88 (2C), 83.42 (2C), 52.50, 33.16, 24.79 (2C), 24.34 (2C), 18.78. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + Na][⊕] 359.0; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{22}^{10}B^{35}ClO_4Na^{\oplus}$ 358.1228, found 358.1241; IR (KBr) ν (cm⁻¹) 2978, 2955, 1918, 1723, 1490, 1446, 1372, 1337, 1279, 1259, 1192, 1146, 1098, 1067, 1010, 967, 864, 752, 664, 542; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane-ipropanol = 98/2; flow rate = 1.0 mL min^{-1} ; retention time: 5.2 min (R,R-isomer), 6.5 min (S,S-isomer).

(1*R*,2*R*)-Methyl 1-(4-bromophenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3g). White semisolid. 42.9 mg, 75% yield. $[\alpha]_D^{26}$ -79.8 (*c* 1.0, CHCl₃) for

89% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.40 (d, J = 8.4 Hz, 2H), 7.21 (d, J = 8.4 Hz, 2H), 3.60 (s, 3H), 1.71 (dd, J = 10.0 Hz, 3.2 Hz, 1H), 1.57 (dd, J = 8.0 Hz, 3.2 Hz, 1H), 1.27 (dd, J = 10.0 Hz, 8.0 Hz, 1H), 1.07 (s, 6H), 0.86 (s, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm) 174.27, 136.87, 132.84 (2C), 130.84 (2C), 120.98, 83.42 (2C), 52.51, 33.25, 24.79 (2C), 24.34 (2C), 18.74. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + Na][⊕] 403.1; HRMS (FTMS-ESI): [M + Na][⊕] calcd for $C_{17}H_{22}^{10}B^{79}BrO_4Na^{⊕}$ 402.0723, found 402.0719; IR (KBr) ν (cm⁻¹) 3062, 3045, 2987, 2945, 2848, 1724, 1486, 1424, 1325, 1265, 1193,1141, 1012, 856, 826, 768, 757, 539, 510; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane–i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 7.3 min (R,R-isomer), 9.6 min (S,S-isomer).

1-(3-bromophenyl)-2-(4,4,5,5-tetramethyl-(1R,2R)-Methyl 1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3h). White semisolid. 43.9 mg, 77% yield. $[\alpha]_D^{26}$ –152.5 (c 1.0, CHCl₃) for 93% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.48 (s, 1H), 7.36 (d, J = 7.6 Hz, 1H), 7.28-7.25 (m, 1H), 7.14 (t, J = 7.6 Hz, 1H),3.61 (s, 3H), 1.72 (dd, J = 10.0 Hz, 3.6 Hz, 1H), 1.56 (dd, J =8.0 Hz, 3.6 Hz, 1H), 1.26 (dd, J = 10.0 Hz, 8.0 Hz, 1H), 1.09 (s, 6H), 0.88 (s, 6H); 13 C NMR (100 MHz, CDCl₃) δ (ppm) 174.20, 140.19, 134.34, 130.17, 129.70, 129.33, 121.71, 83.46 (2C), 52.56, 33.48, 24.87 (2C), 24.42 (2C), 18.85. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + Na][⊕] 403.1; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{22}^{10}B^{79}BrO_4Na^{\oplus}$ 402.0733, found 402.0723; IR (KBr) ν (cm⁻¹) 3419, 3048, 2983, 2951, 1723, 1597, 1566, 1479, 1404, 1260, 1281, 1260, 1166, 1138, 998, 977, 854, 716, 695, 686, 574, 564; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 5.4 min (R,R-isomer), 6.9 min (S,S-isomer).

(1R,2R)-Methyl 1-(2-bromophenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate semisolid. 42.2 mg, 74% yield. $[\alpha]_{D}^{26}$ -219.3 (c 1.0, CHCl₃) for 94% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.51 (d, J = 8.0 Hz, 1H), 7.31-7.24 (m, 2H), 7.13-7.11 (m, 1H), 3.62 (s, 3H), 1.79-1.44 (m, 2H), 1.31-1.26 (m, 1H), 1.07 (s, 6H), 0.88 (s, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm) 173.78, 132.28, 128.42, 126.81, 83.18 (2C), 52.59, 35.46, 24.76 (2C), 24.25 (2C), 21.37. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: $[M + Na]^{\oplus}$ 403.0; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{22}^{10}B^{79}BrO_4Na^{\oplus}$ 402.0723, found 402.0719; IR (KBr) ν (cm⁻¹) 3061, 3018, 2978, 2954, 2931, 1721, 1592, 1567, 1431, 1411, 1332, 1285, 1169, 1143, 993, 860, 759, 666, 561; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; *n*-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 7.9 min (R,R-isomer), 10.9 min (S,S-isomer).

(1*R*,2*R*)-Methyl 1-(naphthalen-2-yl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3j). Colorless oil. 31.2 mg, 59% yield. [α]²⁶ -194.7 (c 1.0, CHCl₃) for 92% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.80–7.73 (m, 4H), 7.50–7.42 (m, 3H), 3.60 (s, 3H), 1.79 (dd, J = 10.0 Hz, 3.2 Hz,

1H), 1.74 (dd, J = 8.0 Hz, 3.2 Hz, 1H), 1.35 (dd, J = 10.0 Hz, 8.0 Hz, 1H), 1.00 (s, 6H), 0.67 (s, 6H); 13 C NMR (100 MHz, CDCl₃) δ (ppm) 174.86, 135.28, 133.05, 132.61, 129.58, 129.43, 127.76, 127.50, 127.25, 125.80, 125.65, 83.28 (2C), 52.47, 33.92, 24.73 (2C), 24.27 (2C), 18.94. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + Na] $^{\oplus}$ 375.0; HRMS (FTMS-ESI): [M + Na] $^{\oplus}$ calcd for C₂₁H₂₅ 10 BO₄Na $^{\oplus}$ 374.1774, found 374.1771; IR (KBr) ν (cm $^{-1}$) 2999, 2982, 1724, 1618, 1438, 1410, 1265, 1168, 1147, 1128, 1113, 1070, 1016, 974, 858, 759, 660, 608, 531; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane–i-propanol = 98/2; flow rate = 1.0 mL min $^{-1}$; retention time: 8.7 min (R,R-isomer), 12.3 min (S,S-isomer).

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(1R,2R)-Methyl 1-(3,4-dichlorophenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3k). White semisolid. 43.3 mg, 78% yield. $[\alpha]_{D}^{26}$ -98.4 (c 1.0, CHCl₃) for 92% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.43 (d, J = 1.2 Hz, 1H), 7.35 (d, J = 8.4 Hz, 1H), 7.17 (dd, J = 8.4 Hz, 1.2 Hz, 1H), 3.62 (s, 3H), 1.74 (dd, J = 10.0 Hz, 3.2 Hz, 1H), 1.54 (dd, J =8.0 Hz, 3.2 Hz, 1H), 1.30-1.24 (m, 1H), 1.09 (s, 6H), 0.90 (s, 6H); 13 C NMR (100 MHz, CDCl₃) δ (ppm) 173.95, 138.32, 133.38, 131.70, 131.19, 130.57, 129.77, 83.66 (2C), 52.71, 33.16, 24.93 (2C), 24.49 (2C), 19.08. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: $[M + Na]^{\oplus}$ 393.0; HRMS (FTMS-ESI): $[M + Na]^{\oplus} \ calcd \ for \ C_{17} H_{21}{}^{10} B^{35} Cl_2 O_4 Na^{\oplus} \ 392.0838, \ found$ 392.0830; IR (KBr) ν (cm⁻¹) 2979, 2952, 1727, 1558, 1474, 1435, 1411, 1380, 1372, 1333, 1262, 1224, 1193, 1167, 1140, 1104, 1071, 1031, 971, 945, 857, 833, 758, 737, 666, 597; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 5.2 min (R,R-isomer), 6.5 min (S,S-isomer).

(1R,2R)-Methyl 1-(3,5-difluorophenyl)-2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)cyclopropanecarboxylate (3m). White semisolid. 42.1 mg, 83% yield. $[\alpha]_{D}^{26}$ -96.1 (c 1.0, CHCl₃) for 93% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 6.89–6.86 (m, 2H), 6.69 (t, J = 9.2 Hz, 1H), 3.62 (s, 3H), 1.73 (dd, J = 10.0 Hz, 3.6 Hz, 1H), 1.55 (dd, J = 8.4 Hz, 3.6 Hz, 1H), 1.27 (dd, J =10.0 Hz, 8.4 Hz, 1H), 1.10 (s, 6H), 0.92 (s, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm) 173.78, 162.46 (dd, J_{CF} = 246, 12.9 Hz, 2C), 141.81 (t, J_{CF} = 9.1 Hz), 114.30 (dd, J_{CF} = 18.2 Hz, 6.1 Hz, 2C), 102.68 (t, J_{CF} = 25.0 Hz), 83.62 (2C), 52.71, 33.69, 24.93 (2C), 24.46 (2C), 19.02. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + Na][⊕] 361.3; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{21}BF_2O_4Na^{\oplus}$ 360.1429, found 360.1434; IR (KBr) ν (cm⁻¹) 3438, 3085, 1981, 1728, 1624, 1599, 1435, 1409, 1372, 1334, 1269, 1216, 1142, 1100, 1077, 990, 966, 857, 759, 736, 685, 532, 511; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane-i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 4.4 min ($R_{\star}R_{\star}$ isomer), 5.6 min (S,S-isomer).

(*R*)-Dimethyl 2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-cyclopropane-1,1-dicarboxylate (3p). Colorless oil. 16.6 mg, 39% yield. $[\alpha]_{\rm D}^{24}$ –59.8 (*c* 1.06, CHCl₃) for 95% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 3.73 (s, 3H), 3.71 (s, 3 H), 1.54–1.52

(m, 2H), 1.23 (s, 6H), 1.22 (s, 6H), 1.12–1.08 (m, 1 H); 13 C NMR (100 MHz, CDCl₃) δ (ppm) 170.88, 169.23, 83.93 (2C), 52.77, 52.54, 33.71, 24.84 (2C), 24.81 (2C), 18.98. (The carbon directly attached to the boron atom was not detected, likely due to quadrupole relaxation.) ESI-MS: [M + H]^{\oplus} 285.2; HRMS (FTMS-ESI): [M + H]^{\oplus} calcd for C₁₃H₂₂¹⁰BO₆^{\oplus} 285.1504, found 285.15; IR (KBr) ν (cm⁻¹) 2980, 2954, 1735, 1436, 1414, 1381, 1373, 1338, 1290, 1271, 1234, 1208, 1167, 1142, 1079, 971, 879, 858, 835, 772, 758, 669; HPLC: Phenomenex Lux 5u Cellulose-2 (PC-2) Column; detected at 220 nm; n-hexane–i-propanol = 98/2; flow rate = 1.0 mL min⁻¹; retention time: 11.8 min (S-isomer), 12.9 min (R-isomer).

(1R,2S)-Methyl 1,2-diphenylcyclopropanecarboxylate (5). A mixture of 3a (30.2 mg, 0.1 mmol), iodobenzene (4, 30.6 mg, 0.15 mmol), Pd(PPh₃)₄ (11.6 mg, 0.01 mmol), and Cs₂CO₃ (97.5 mg, 0.3 mmol) in toluene (1 mL) was stirred at 80 °C under a N₂ atmosphere overnight. After cooling to room temperature, the reaction mixture was filtered and washed with ethyl acetate. The filtrate was concentrated under reduced pressure and purified by flash column chromatography using hexane-ethyl acetate (8:1) as an eluent to afford the desired product 5 (25 mg, 99% yield). $\left[\alpha\right]_{D}^{26}$ -38.9 (c 1.0, CHCl₃) for 94% ee; ¹H NMR (400 MHz, CDCl₃) δ (ppm) 7.05–6.94 (m, 8H), 6.70-6.68 (m, 2H), 3.58 (s, 3H), 3.06-3.01 (m, 1H), 2.06 (dd, J = 9.2 Hz, 4.8 Hz, 1H), 1.82-1.78 (m, 1H); ¹³C NMR (100 MHz, $CDCl_3$) δ (ppm) 174.42, 136.42, 134.80, 131.99, 128.40, 128.10, 127.76, 127.09, 126.37, 52.68, 37.45, 33.19, 20.54; ESI-MS: $[M + Na]^{\oplus}$ 275.1; HRMS (FTMS-ESI): $[M + Na]^{\oplus}$ calcd for $C_{17}H_{16}O_2Na^{\oplus}$ 275.1043, found 275.1037; IR (KBr) ν (cm⁻¹) 3648, 3412, 3061, 3086, 3029, 2953, 1966, 1897, 1720, 1602, 1496, 1456, 1447, 1428, 1376, 1342, 1255, 1205, 1189, 1104, 1050, 989, 865, 788, 760, 742, 702, 650, 545; HPLC: OJ-H Column; detected at 214 nm; n-hexane-i-propanol = 95/5; flow rate = 0.7 mL min^{-1} ; retention time: 11.5 min (R,S-isomer), 16.3 min (*S*,*R*-isomer).

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