Arylation of α-Pivalate Ketones via Ni-Catalyzed sp3 C-O Activation with Arylboroxine

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**General:**

All the reactions were carried out under nitrogen atmosphere using standard Schlenk technique. NiCl$_2$(PCy$_3$)$_2$ was synthesized according to literature method$^1$. Ni(COD)$_2$, boronic acids and NaOtBu were purchased from Alfa Aesar Company. Toluene was freshly distilled over sodium with the use of diphenyl ketone as an indicator under nitrogen. $N,N$-dimethyl formate (DMF) was dried using standard method. Pivalates were prepared by treating the corresponding alcohols with trimethyl amine followed by PivCl in CH$_2$Cl$_2$. $^1$H NMR (300 MHz) and $^{13}$C NMR (50 MHz) were registered on Varian 300 M or 200 M spectrometers with CDCl$_3$ as solvent and tetramethylsilane (TMS) as internal standard. Chemical shifts were reported in units (ppm) by assigning TMS resonance in the $^1$H spectrum as 0.00 ppm and CDCl$_3$ resonance in the $^{13}$C spectrum as 77.0 ppm. All coupling constants ($J$ values) were reported in Hertz (Hz). Column chromatography was performed on silica gel 200-300 mesh. IR, GC, MS, and HRMS were performed by the State-authorized Analytical Center in Peking University.

**Typical procedure:**

**Synthesis of 2-oxo-2-phenylethyl pivalate (1a):**

![Chemical structure](image)

2-hydroxyacetophenone (680 mg, 5 mmol) was placed in a dry 100 ml flask, and DCM (30 mL) was added into the flask. Then Et$_3$N (1.01 g, 10 mmol) and PivCl (720 mg, 6 mmol) were added at room temperature. The resulting mixture was stirred overnight. 40 mL distilled water was added into the mixture and the aqueous phase was extracted with DCM (30 mL). The combined organic layers were washed with brine before being dried over anhydrous Na$_2$SO$_4$. After removal of solvent, the product was obtained in 88% yield by silicon gel column chromatography.

The substrates of 1j-1y were prepared in a similar way.

**The reaction of $\alpha$-pivalate ketones with boronic acids:**

An oven-dried Schlenk tube was charged with $\alpha$-pivalate ketones 1 (0.25 mmol), boronic acid 2 (1.0 mmol) or boroxines (0.33 mmol), Ni(PCy$_3$)$_2$Cl$_2$ (17.3 mg, 0.025 mmol), and NaOtBu (48 mg, 0.5 mmol).
The tube was evacuated and refilled with N₂ and this process was repeated three times. Then freshly distilled toluene (1.0 mL) and DMF (0.5 mL) were injected and the resulting mixture was stirred at 100 °C for 40 minutes. The mixture was cooled to room temperature under N₂, quenched via the addition of saturated aqueous ammonium chloride (4 mL) and water (8 mL). The aqueous phase was extracted with ethyl acetate (10 x 3 mL). The combined organic layers were washed with brine, dried over anhydrous Na₂SO₄, filtered and concentrated. The product was obtained by silicon gel short column chromatography.

The products of 4aa-4af and 4ja-4ya were prepared in the same way.

**The reaction of α-pivalate ketones with boroxines:**

An oven-dried Schlenk tube was charged with α-pivalate ketones 1 (0.25 mmol), boroxines 3 (0.33 mmol), Ni(PCy₃)₂Cl₂ (17.3 mg, 0.025 mmol), and NaOtfBu (48 mg, 0.5 mmol). The tube was evacuated and refilled with N₂ and this process was repeated three times. Then freshly distilled toluene (1.0 mL) and DMF (0.5 mL) were injected and the resulting mixture was stirred at 100 °C for 60 minutes. The mixture was cooled to room temperature under N₂, quenched via the addition of saturated aqueous ammonium chloride (4 mL) and water (8 mL). The aqueous phase was extracted with ethyl acetate (10 x 3 mL). The combined organic layers were washed with brine, dried over anhydrous Na₂SO₄, filtered and concentrated. The product was obtained by silicon gel short column chromatography.

The products of 4ag-4an were prepared in the same way.
**Table S1.** Investigation of the Suzuki-Miyaura Reaction under Various Conditions. \(^a\)

![Chemical Structure](image)

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<th>time</th>
<th>solvent</th>
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<td>100</td>
<td>40 (min)</td>
<td>PhMe/DMF (2 : 1)</td>
<td>71(^c)</td>
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</tbody>
</table>

\(^a\) All the reactions were carried out on the scale of 0.25 mmol of 1a and 1.0 mmol of 2a or 0.33 mmol of 3a under N₂. \(^b\) GC yield with the use of n-dodecane as an internal standard. \(^c\) Isolated yield.
Analytical and spectral data of compounds 4aa-4an and 4ja-4ya

1,2-diphenylethanone (4aa)²
The product was obtained (39.7 mg, 81%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; ¹H-NMR (CDCl₃, 300 MHz): δ 8.08-8.09 (m, 2H), 7.28-7.64 (m, 8H), 4.35 (s, 2H); ¹³C NMR (CDCl₃, 50 MHz): δ 197.5, 136.4, 134.6, 133.0, 129.3, 128.5, 128.3, 126.7, 45.3; MS (EI) m/z: 196 (M⁺).

1-phenyl-2-p-tolylethanone (4ab)³
The product was obtained (39.4 mg, 75%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; ¹H-NMR (CDCl₃, 300 MHz): δ 7.99-8.02 (m, 2H), 7.52-7.54 (m, 1H), 7.42-7.47 (m, 2H), 7.14-7.24 (m, 4H), 4.24 (s, 2H), 2.31 (s, 3H); ¹³C NMR (CDCl₃, 50 MHz): δ 197.9, 136.4, 133.0, 131.5, 129.4, 129.3, 128.6, 128.54, 128.53, 45.1, 21.0; MS (EI) m/z: 210 (M⁺).

1-phenyl-2-m-tolylethanone (4ac)⁴
The product was obtained (35.1 mg, 67%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; ¹H-NMR (CDCl₃, 300 MHz): δ 8.06-8.09 (m, 2H), 7.51-7.61 (m, 3H), 7.27-7.32 (m, 2H), 7.11-7.14 (m, 2H), 4.30 (s, 2H), 2.38 (s, 3H); ¹³C NMR (CDCl₃, 50 MHz): δ 198.1, 138.3, 133.1, 130.1, 128.62, 128.6, 128.5, 127.6, 126.5, 45.5, 21.4; MS (EI) m/z: 210 (M⁺).

2-(4-fluorophenyl)-1-phenylethanone (4ad)⁵
The product was obtained (40.7 mg, 76%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; ¹H-NMR (CDCl₃, 300 MHz): δ 8.00-8.03 (m, 2H), 7.56-7.59 (m, 1H), 7.46-7.51 (m, 2H), 7.21-7.27 (m, 2H), 7.00-7.06 (m, 2H), 4.28 (s, 2H); ¹³C NMR (CDCl₃, 50 MHz): δ 197.6, 164.4, 159.5, 136.5, 133.3, 131.1, 130.9, 130.2, 128.7, 128.5, 115.7, 115.3, 44.5; MS (EI) m/z: 214 (M⁺).
The product was obtained (36.4 mg, 68%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 7.92-8.19 (m, 2H), 7.55-7.60 (m, 1H), 7.40-7.52 (m, 2H), 7.25-7.32 (m, 1H), 6.92-7.05 (m, 3H), 4.26 (s, 2H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.0, 165.7, 136.4, 133.3, 130.1, 130.0, 128.7, 128.5, 125.2, 116.7, 116.3, 114.1, 113.6, 45.0; MS (EI) m/z: 214 (M$^+$).

The product was obtained (44.7 mg, 71%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 8.01-8.04 (m, 2H), 7.46-7.56 (m, 3H), 7.33-7.36 (m, 2H), 7.19-7.25 (m, 2H), 4.26 (s, 2H), 1.29 (s, 8H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.9, 149.5, 136.9, 133.0, 129.1, 128.6, 128.5, 125.6, 44.9, 34.4, 31.3; MS (EI) m/z: 252 (M$^+$).

The product was obtained (45.5 mg, 67%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 8.03-8.06 (m, 2H), 7.25-7.59 (m, 12H), 4.33 (s, 2H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.7, 140.7, 139.9, 136.7, 133.2, 129.9, 128.7, 128.6, 128.5, 127.4, 127.2, 127.0, 45.1 ; MS (EI) m/z: 272 (M$^+$).

The product was obtained (28.9 mg, 55%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 8.02-8.05 (m, 2H), 7.46-7.59 (m, 3H), 7.23-7.15 (m, 4H), 4.32 (s, 2H), 2.28 s, 3H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.5, 136.8, 133.4, 133.1, 130.3, 130.2, 128.6, 128.3, 127.2, 126.1, 43.4, 19.7; MS (EI) m/z: 210 (M$^+$).
2-(4-methoxyphenyl)-1-phenylethanone (4ai)\(^5\)
The product was obtained (34.5 mg, 61%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines; 
\(^1H\)-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 7.97-8.00 (m, 2H), 7.43-7.56 (m, 3H), 7.15-7.18 (m, 2H), 6.83-6.87 (m, 2H), 4.21 (s, 2H), 3.76 (s, 3H); \(^13C\) NMR (CDCl\(_3\), 50 MHz): \(\delta\) 197.9, 158.6, 133.0, 130.4, 128.6, 128.54, 128.52, 126.5, 114.1, 55.2, 44.6; MS (EI) m/z: 226 (M\(^+\)).

2-(3-methoxyphenyl)-1-phenylethanone (4aj)\(^5\)
The product was obtained (36.1 mg, 64%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines; 
\(^1H\)-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 7.98-8.02 (m, 2H), 7.44-7.54 (m, 3H), 7.19-7.29 (m, 1H), 6.80-6.86 (m, 3H), 4.24 (s, 2H), 3.76 (s, 3H); \(^13C\) NMR (CDCl\(_3\), 50 MHz): \(\delta\) 197.2, 159.6, 136.4, 135.8, 132.8, 129.3, 128.4, 128.3, 128.2, 121.5, 114.8, 112.1, 54.8, 45.2; MS (EI) m/z: 226 (M\(^+\)).

2-(4-fluoro-3-methylphenyl)-1-phenylethanone (4ak)
The product was obtained (38.8 mg, 68%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines; 
\(^1H\)-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 7.99-8.02 (m, 2H), 7.44-7.57 (m, 3H), 6.95-7.08 (m, 3H), 4.22 (s, 2H), 2.25 (s, 3H); \(^13C\) NMR (CDCl\(_3\), 50 MHz): \(\delta\) 197.6, 162.0, 158.9, 136.5, 133.2, 132.5, 128.7, 128.6, 128.4, 128.2, 128.1, 115.2, 114.9, 44.5, 14.5; MS (EI) m/z: 228 (M\(^+\)); HRMS (ESI): Calcd. (M+H\(^+\)) 229.10232, Found:229.10196. IR (cm\(^{-1}\)): ν 2989, 2911, 2900, 1788, 1393, 1077, 1066, 740.

2-(2-fluorophenyl)-1-phenylethanone (4al)\(^5\)
The product was obtained (32.6 mg, 61%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines; 
\(^1H\)-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 8.02-8.07 (m, 2H), 7.45-7.58 (m, 3H), 7.22-7.28 (m, 2H), 7.05-7.13 (m, 2H), 4.33 (s, 2H); \(^13C\) NMR (CDCl\(_3\), 50 MHz): \(\delta\) 196.3, 136.2, 133.7, 131.7, 131.5, 128.9, 128.8, 128.7, 128.4, 124.2, 124.1, 115.6, 115.1, 38.6; MS (EI) m/z: 214 (M\(^+\)). HRMS (ESI): Calcd. (M+H\(^+\)), Found; IR (cm\(^{-1}\)): ν 2989, 2911, 2900, 1788, 1393, 1077, 1066, 740.
2-(3,5-dimethylphenyl)-1-phenylethanone (4am)\(^{10}\)

The product was obtained (41.4 mg, 74%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines;
\(^1\)H-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 8.02-8.05 (m, 2H), 7.46-7.59 (m, 3H), 6.81-6.92 (m, 3H), 4.20 (s, 2H), 2.29 (s, 6H); \(^13\)C NMR (CDCl\(_3\), 50 MHz): \(\delta\) 198.0, 138.1, 136.6, 134.3, 133.1, 128.6, 125.7, 124.3, 45.3, 21.2; MS (EI) m/z: 224 (M\(^+\)).

2-(naphthalen-1-yl)-1-phenylethanone (4an)\(^{6}\)

The product was obtained (40.6 mg, 66%) starting from 0.25 mmol of 1a and 0.33 mmol of boroxines;
\(^1\)H-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 8.07-8.10 (m, 2H), 7.79-7.89 (m, 3H), 7.57-7.59 (m, 1H), 7.25-7.51 (m, 6H), 4.74 (s, 2H); \(^13\)C NMR (CDCl\(_3\), 50 MHz): \(\delta\) 197.6, 136.6, 133.8, 133.2, 132.2, 131.3, 128.7, 128.6, 128.4, 128.0, 126.2, 125., 123.8, 43.0; MS (EI) m/z: 246 (M\(^+\)).

1-(2,3-dihydrobenzo[b][1,4]dioxin-6-yl)-2-phenylethanone (4ja)\(^{11}\)

The product was obtained (48.9 mg, 77%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid;
\(^1\)H-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 7.54-7.57 (m, 2H), 7.23-7.34 (m, 5H), 6.88-6.91 (m, 1H), 4.22-4.31 (m, 4H), 4.19 (s, 2H); \(^13\)C NMR (CDCl\(_3\), 50 MHz): \(\delta\) 196.1, 148.1, 143.4, 134.9, 130.5, 129.4, 129.3, 128.6, 126.8, 126.7, 122.79, 122.78, 118.1, 117.3, 117.2, 64.7, 64.1, 45.2; MS (EI) m/z: 254 (M\(^+\)).

1-(benzo[d][1,3]dioxol-5-yl)-2-phenylethanone (4ka)\(^{12}\)

The product was obtained (43.2 mg, 72%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid;
\(^1\)H-NMR (CDCl\(_3\), 300 MHz): \(\delta\) 7.61-7.65 (m, 1H), 7.47-7.48 (m, 1H), 7.24-7.35 (m, 5H), 6.82-6.85 (m, 1H), 6.04 (t, \(J = 6\) Hz, 2H), 4.21 (s, 2H); \(^13\)C NMR (CDCl\(_3\), 50 MHz): \(\delta\) 195.7, 151.8, 148.2, 131.5, 129.3, 128.6, 126.8, 125.0, 108.3, 107.9, 101.8, 45.3; MS (EI) m/z: 240 (M\(^+\)).
The product was obtained (40.1 mg, 71%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 7.51-7.60 (m, 3H), 7.07-7.37 (m, 6H), 4.25 (s, 2H), 3.82 (s, 3H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.3, 134.5, 129.6, 129.4, 128.6, 126.9, 121.3, 119.6, 112.9, 55.4, 45.6; MS (EI) m/z: 226 (M$^+$).

The product was obtained (36.7 mg, 65%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 7.99-8.02 (m, 2H), 7.26-7.33 (m, 5H), 6.91-6.94 (m, 2H), 4.24 (s, 2H), 3.86 (s, 3H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 196.2, 163.5, 135.0, 130.9, 129.6, 129.4, 129.3, 128.6, 128.5, 126.7, 113.8, 113.7, 55.4, 45.2; MS (EI) m/z: 226 (M$^+$).

The product was obtained (37.8 mg, 72%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 7.89-7.92 (m, 2H), 7.23-7.29 (m, 7H), 4.25 (s, 2H), 2.39 (s, 3H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.1, 143.9, 134.7, 134.3, 129.4, 129.2, 128.7, 128.6, 126.8, 45.4, 21.6; MS (EI) m/z: 210 (M$^+$).

The product was obtained (36.2 mg, 69%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
$^1$H-NMR (CDCl$_3$, 300 MHz): $\delta$ 7.79-7.82 (m, 2H), 7.24-7.36 (m, 7H), 4.27 (s, 2H), 2.40 (s, 3H); $^{13}$C NMR (CDCl$_3$, 50 MHz): $\delta$ 197.9, 156.7, 138.4, 136.7, 134.7, 133.9, 129.4, 129.1, 128.6, 128.4, 126.8, 125.8, 45.5, 21.3; MS (EI) m/z: 210 (M$^+$).
2-phenyl-1-o-tolyethanone (4pa)\textsuperscript{6}

The product was obtained (35.1 mg, 67\%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; \textsuperscript{1}H-NMR (CDCl\textsubscript{3}, 300 MHz): \(\delta\) 7.69-7.72 (m, 1H), 7.20-7.36 (m, 8H), 4.20 (s, 2H), 2.40 (s, 3H); \textsuperscript{13}C NMR (CDCl\textsubscript{3}, 50 MHz): \(\delta\) 201.6, 138.4, 134.5, 131.9, 131.3, 129.5, 128.6, 128.5, 126.8, 125.6, 48.6, 21.2; MS (EI) m/z: 210 (M\textsuperscript{+}).

\[
\begin{align*}
\text{Ph} & \quad \text{O} & \quad \text{C}
\end{align*}
\]

1-(biphenyl-4-yl)-2-phenylethanone (4qa)\textsuperscript{13}

The product was obtained (49.6 mg, 73\%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; \textsuperscript{1}H-NMR (CDCl\textsubscript{3}, 300 MHz): \(\delta\) 8.06-8.09 (m, 2H), 7.60-7.69 (m, 4H), 7.25-7.46 (m, 8H), 4.31 (s, 2H); \textsuperscript{13}C NMR (CDCl\textsubscript{3}, 50 MHz): \(\delta\) 197.2, 145.8, 139.8, 134.6, 129.4, 129.2, 128.9, 128.6, 127.2, 126.8, 45.5; MS (EI) m/z: 272 (M\textsuperscript{+}).

\[
\begin{align*}
\text{Ph} & \quad \text{O} & \quad \text{C}
\end{align*}
\]

1-(naphthalen-2-yl)-2-phenylethanone (4ra)\textsuperscript{14}

The product was obtained (41.8 mg, 68\%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; \textsuperscript{1}H-NMR (CDCl\textsubscript{3}, 300 MHz): \(\delta\) 8.55-8.56 (m, 1H), 8.05-8.09 (m, 1H), 7.86-7.98 (m, 3H), 7.55-7.61 (m, 2H), 7.33-7.35 (m, 5H), 4.43 (s, 2H); \textsuperscript{13}C NMR (CDCl\textsubscript{3}, 50 MHz): \(\delta\) 197.6, 135.7, 134.7, 132.5, 130.4, 129.6, 129.5, 128.7, 128.5, 127.7, 126.9, 126.8, 124.2, 45.5; MS (EI) m/z: 246 (M\textsuperscript{+}).

\[
\begin{align*}
\text{Ph} & \quad \text{O} & \quad \text{C}
\end{align*}
\]

3,3-dimethyl-1-phenylbutan-2-one (4sa)\textsuperscript{15}

The product was obtained (41.8 mg, 68\%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; \textsuperscript{1}H-NMR (CDCl\textsubscript{3}, 300 MHz): \(\delta\) 8.55-8.56 (m, 1H), 8.05-8.09 (m, 1H), 7.86-7.98 (m, 3H), 7.55-7.61 (m, 2H), 7.33-7.35 (m, 5H), 4.43 (s, 2H); \textsuperscript{13}C NMR (CDCl\textsubscript{3}, 50 MHz): \(\delta\) 197.6, 135.7, 134.7, 132.5, 130.4, 129.6, 129.5, 128.7, 128.5, 127.7, 126.9, 126.8, 124.2, 45.5; MS (EI) m/z: 246 (M\textsuperscript{+}).
The product was obtained (43.8 mg, 82%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
\[^1\text{H}]\text{NMR (CDCl}_3, 300 \text{ MHz): } \delta 7.80-8.06 (m, 2H), 7.24-7.35 (m, 5H), 7.08-7.15 (m, 2H), 4.25 (s, 2H); 
\[^{13}\text{C}\text{ NMR (CDCl}_3, 50 \text{ MHz): } \delta 196.1, 168.3, 163.3, 134.0, 131.3, 131.1, 129.3, 128.7, 126.9, 115.9, 
115.5, 45.5; \text{MS (EI) m/z: } 214 (M^+).} 

The product was obtained (39.6 mg, 57%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
\[^1\text{H}]\text{NMR (CDCl}_3, 300 \text{ MHz): } \delta 7.91-7.98 (m, 2H), 7.11-7.39 (m, 7H), 4.23 (s, 2H), 2.51-2.55 (m, 1H), 
1.74-1.92 (m, 6H), 1.36-1.46 (m, 4H); \[^{13}\text{C}\text{ NMR (CDCl}_3, 50 \text{ MHz): } \delta 197.2, 153.8, 134.9, 134.5, 129.4, 
128.8, 128.6, 128.5, 127.1, 127.0, 126.7, 45.4, 44.6, 34.0, 26.7, 26.0; \text{MS (EI) m/z: } 278 (M^+).} 

The product was obtained (64.9 mg, 86%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
\[^1\text{H}]\text{NMR (CDCl}_3, 300 \text{ MHz): } \delta 7.59-7.61 (m, 2H), 7.14-7.45 (m, 7H), 5.11 (s, 2H), 4.25 (s, 2H); \[^{13}\text{C}\text{ NMR (CDCl}_3, 50 \text{ MHz): } \delta 197.2, 158.8, 137.7, 136.2, 134.2, 129.3, 129.1, 128.4, 128.3, 127.8, 127.2, 126.5, 
121.2, 120.1, 113.7, 69.9, 45.3; \text{MS (EI) m/z: } 302 (M^+).} 

The product was obtained (45.4 mg, 71%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; 
\[^1\text{H}]\text{NMR (CDCl}_3, 300 \text{ MHz): } \delta 7.65-7.68 (m, 2H), 7.24-7.39 (m, 7H), 5.20 (s, 2H), 4.26 (s, 2H), 3.48 (s, 
3H); \[^{13}\text{C}\text{ NMR (CDCl}_3, 50 \text{ MHz): } \delta 197.2, 157.5, 137.9, 134.4, 129.6, 129.4, 128.6, 126.8, 122.2, 121.2, 
116.0, 94.4, 56.1, 45.6; \text{MS (EI) m/z: } 256 (M^+). \text{ HRMS (ESI): Calcd. } 257.11722, \text{ (M+H}^+)\text{, Found: } 
257.11708; \text{IR (cm}^{-1}): \nu 2917, 2849, 1761, 1554, 1393, 1231, 1027, 852, 727.
The product was obtained (32.0 mg, 48%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; ¹H-NMR (CDCl₃, 300 MHz): δ 8.03-8.13 (m, 4H), 7.24-7.33 (m, 5H), 4.36 (q, J = 7.2, 2H), 4.31 (s, 2H), 1.41 (t, J = 7.2 Hz, 3H); ¹³C NMR (CDCl₃, 50 MHz): δ 197.2, 165.8, 139.6, 134.2, 1339, 129.8, 129.4, 128.7, 128.5, 127.0, 61.4, 45.8, 14.3; MS (EI) m/z: 268 (M⁺).

The product was obtained (41.4 mg, 56%) starting from 0.25 mmol of 1a and 1 mmol of boronic acid; ¹H-NMR (CDCl₃, 300 MHz): δ 8.08-8.11 (m, 2H), 7.30-7.38 (m, 5H), 7.19-7.22 (m, 2H), 4.32 (s, 2H), 1.41 (s, 9H); ¹³C NMR (CDCl₃, 50 MHz): δ 196.7, 176.1, 154.6, 134.2, 133.6, 129.9, 128.4, 126.6, 121.4, 45.2, 38.8, 26.7; MS (EI) m/z: 296 (M⁺).
1,2-diphenylethanone (4aa)
1-phenyl-2-m-tolylethanone (4ac)
2-(4-fluorophenyl)-1-phenylethanone (4ad)
2-(3-fluorophenyl)-1-phenylethanone (4ae)
2-(4-tert-butyphenyl)-1-phenylethanone (4af)
2-(biphenyl-4-yl)-1-phenylethanone (4ag)
1-phenyl-2-o-tolylethanone (4ah)
2-(4-methoxyphenyl)-1-phenylethanone (4ai)
2-(3-methoxyphenyl)-1-phenylethanone (4aj)
2-(4-fluoro-3-methylphenyl)-1-phenylethanone (4ak)
2-(2-fluorophenyl)-1-phenylethanone (4al)
2-(3,5-dimethylphenyl)-1-phenylethanone (4am)
2-(naphthalen-1-yl)-1-phenylethanone (4an)
1-(2,3-dihydrobenzo[b][1,4]dioxin-6-yl)-2-phenylethanone (4a)
1-(benzo[d][1,3]dioxol-5-yl)-2-phenylethanone (4ka)
1-(3-methoxyphenyl)-2-phenylethanone (4la)
1-(4-methoxyphenyl)-2-phenylethanone (4ma)
2-phenyl-1-p-tolylethanone (4na)
2-phenyl-1-m-tolylethanone (4oa)
2-phenyl-1-\text{-}o\text{-}tolylethanone (4pa)
1-(biphenyl-4-yl)-2-phenylethanone (4ca)
1-(naphthalen-2-yl)-2-phenylethanone (4ra)
3,3-dimethyl-1-phenylbutan-2-one (4sa)
1-(4-fluorophenyl)-2-phenylethanone (4ta)
1-(4-cyclohexylphenyl)-2-phenylethanone (4ua)
1-(3-(benzyloxy)phenyl)-2-phenylethanone (4va)
1-(3-(methoxymethoxy)phenyl)-2-phenylethanone (4wa)
ethyl 4-(2-phenylacetyl)benzoate (4xa)
4-(2-phenylacetyl)phenyl pivalate (4ya)
References