Supporting Information for


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Contents:

<table>
<thead>
<tr>
<th>Section</th>
<th>Pages</th>
</tr>
</thead>
<tbody>
<tr>
<td>General Methods</td>
<td>S1-S2</td>
</tr>
<tr>
<td>Synthesis and characterization of cyclopenta[b]pyrrole derivatives</td>
<td>S2-S8</td>
</tr>
<tr>
<td>X-ray crystal structure of 3c</td>
<td>S9</td>
</tr>
<tr>
<td>NMR spectra of all new compounds</td>
<td>S10-S25</td>
</tr>
</tbody>
</table>

**General Methods.** All reactions were carried out under argon or nitrogen. Dioxane, THF, Toluene was distilled from sodium/benzophenone. Unless noted, all commercial reagents were used without further purification.

\(^1\)H NMR spectra was recorded at 300 or 400 MHz, \(^13\)C NMR spectra was recorded at 75 or 100 MHz, and in CDCl₃ (containing 0.03% TMS) solutions. \(^1\)H NMR spectra was recorded with tetramethylsilane (δ = 0.00 ppm) as internal reference; \(^13\)C NMR spectra was recorded with CDCl₃ (δ = 77.00 ppm) as internal reference. GC-MS analyses were performed on a GC-MS analysis. High-resolution mass spectra were performed on an EI or ESI mass spectrometer.
Typical procedure for the synthesis of 3-benzoyl-1,4,4-triphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one (3a). To a solution of zinc(II) chloride (4.1 mg, 0.03 mmol) in DCE (3 mL) was added furan-2-yldiphenylmethanol (75.1 mg, 0.3 mmol), (Z)-1-phenyl-3-(phenylamino)prop-2-en-1-one (67 mg, 0.3 mmol). The resulting solution was stirred at 80 °C for 2 h. Then the solvent was evaporated under reduced pressure and the residue was purified by Chromatography on silica gel (eluent: petroleum ether / ethyl acetate = 3 : 1) afforded the product in 94% yield as a brown solid. M.p. 211-214 °C. \(^1\)H NMR (400 MHz, CDCl\(_3\), Me\(_4\)Si) \(\delta\) 2.80 (d, \(J = 18.8\) Hz, 1H), 3.05 (dd, \(J = 8.0, 18.8\) Hz, 1H), 5.07-5.11 (m, 1H), 5.29 (d, \(J = 10.0\) Hz, 1H), 6.83-6.89 (m, 4H), 7.02-7.10 (m, 4H), 7.15-7.19 (m, 3H), 7.26-7.44 (m, 8H), 7.84-7.86 (m, 2H); \(^13\)C NMR (100.6 MHz, CDCl\(_3\), Me\(_4\)Si) \(\delta\) 41.66, 52.59, 60.42, 67.22, 116.19, 120.09, 123.12, 126.51, 127.10, 127.47, 127.76, 127.88, 128.55, 128.66, 129.68, 130.33, 130.51, 139.14, 139.48, 140.57, 141.56, 147.93, 189.32, 215.44; HRMS (ESI) calcd for C\(_{32}H\(_{26}\)NO\(_2\) [M+H]\(^+\): 456.1964, found 456.1953.

3-benzoyl-1-(4-chlorophenyl)-4,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one (3b). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate = 2 : 1) afforded the product in 78% yield as a brown solid. M.p. 113-116 °C. \(^1\)H NMR (400 MHz, CDCl\(_3\), Me\(_4\)Si) \(\delta\) 2.73 (d, \(J = 18.4\) Hz, 1H), 3.05 (dd, \(J = 8.0, 18.8\) Hz, 1H), 5.03-5.08 (m, 1H), 5.30 (d, \(J = 10.0\) Hz, 1H), 6.75-6.78 (m, 2H), 6.86-6.88 (m, 2H), 7.04-7.10 (m, 4H), 7.18-7.20 (m, 2H), 7.24-7.44 (m, 8H), 7.80 (d, \(J = 7.6\) Hz, 2H); \(^13\)C NMR (100.6 MHz, CDCl\(_3\), Me\(_4\)Si) \(\delta\) 41.58, 52.75, 60.39, 67.23, 117.21, 120.66, 126.62, 127.18, 127.61, 127.85, 127.96, 128.13, 128.62, 128.65, 129.72, 130.38, 130.73, 137.95, 139.27, 140.43, 141.41, 146.99, 189.28, 214.91; HRMS (ESI) calcd for C\(_{32}H\(_{25}\)ClNO\(_2\) [M+H]\(^+\): 490.1574, found 490.1572.
3-benzoyl-4,4-diphenyl-1-(3,4,5-trimethoxyphenyl)-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3c). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 93% yield as a yellow solid. M.p. 217-220 °C. ¹H NMR (400 MHz, CDCl₃, Me₄Si) δ 2.85 (d, J = 18.8 Hz, 1H), 3.06 (dd, J = 8.0, 18.6 Hz, 1H), 3.80 (s, 3H), 3.83 (s, 6H), 5.09 (dd, J = 8.0, 8.4 Hz, 1H), 5.28 (d, J = 9.6 Hz, 1H), 6.05 (s, 2H), 6.87-6.90 (m, 2H), 7.05-7.11 (m, 4H), 7.18-7.20 (m, 2H), 7.26-7.44 (m, 6H), 7.83 (d, J = 7.6 Hz, 2H); ¹³C NMR (100.6 MHz, CDCl₃, Me₄Si) δ 41.75, 52.59, 56.29, 60.91, 61.01, 67.15, 95.08, 119.84, 126.51, 127.03, 127.46, 127.71, 127.87, 128.52, 128.65, 130.34, 130.49, 134.75, 135.60, 139.57, 140.55, 141.53, 148.21, 154.02, 189.20, 215.23; HRMS (ESI) calcd for C₃₅H₃₂NO₅ [M+H]+: 546.2281, found 546.2280.

3-benzoyl-1-(2,6-dimethylphenyl)-4,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3d). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate = 5 : 1) afforded the product in 85% yield as a white solid. M.p. 247-250 °C. ¹H NMR (400 MHz, CDCl₃, Me₄Si) δ 2.21 (s, 3H), 2.42 (d, J = 16.4 Hz, 1H), 2.44 (s, 3H), 2.77 (dd, J = 7.6, 18.0 Hz, 1H), 4.99 (dd, J = 8.0, 10.0 Hz, 1H), 5.42 (d, J = 10.0 Hz, 1H), 6.77 (s, 1H), 7.01-7.06 (m, 4H), 7.10-7.23 (m, 8H), 7.27-7.34 (m, 2H), 7.42 (t, J = 7.6 Hz, 2H), 7.79 (d, J = 7.6 Hz, 2H); ¹³C NMR (100.6 MHz, CDCl₃, Me₄Si) δ 18.32, 19.45, 41.27, 52.79, 64.06, 66.98, 117.20, 126.38, 126.86, 127.43, 127.64, 127.67, 128.36, 128.49, 128.67, 129.04, 129.44, 130.06, 130.89, 136.54, 136.77, 136.95, 139.88, 140.38, 141.43, 154.52, 188.84, 215.81; HRMS (ESI) calcd for C₃₄H₃₀NO₂ [M+H]+: 484.2277, found 484.2258.

3-benzoyl-1-(2,6-diisopropylphenyl)-4,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3e). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 92% yield as a white solid. M.p. 257-260 °C. ¹H NMR (400 MHz, CDCl₃, Me₄Si) δ 1.05 (d, J = 6.8 Hz, 3H), 1.23-1.26 (m, 6H), 1.47 (d, J = 6.8 Hz, 3H), 2.52 (d, J = 18.0 Hz, 1H), 2.81 (dd, J = 7.6, 17.7 Hz, 1H), 2.91-2.98 (m, 1H), 3.30-3.37 (m, 1H), 4.85 (dd, J = 7.6,
10.2 Hz, 1H), 5.45 (d, J = 10.4 Hz, 1H), 6.74 (s, 1H), 7.03-7.05 (m, 3H), 7.11-7.36 (m, 11H), 7.43 (t, J = 7.2 Hz, 2H), 7.83 (d, J = 8.0 Hz, 2H); 13C NMR (100.6 MHz, CDCl3, Me4Si) δ 22.40, 24.11, 24.64, 25.57, 28.22, 28.63, 41.01, 53.14, 66.55, 67.03, 117.12, 124.22, 125.00, 126.37, 126.92, 127.36, 127.53, 127.61, 128.50, 128.65, 129.41, 130.01, 130.78, 133.84, 139.90, 140.45, 141.48, 147.75, 147.90, 155.56, 189.09, 215.81; HRMS (ESI) calcd for C38H38NO2 [M+H]+: 540.2903, found 540.2901.

3-benzoyl-1-benzyl-4,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3f). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 83% yield as a brown oil. 1H NMR (400 MHz, CDCl3, Me4Si) δ 2.67-2.75 (m, 2H), 4.18 (d, J = 14.8 Hz, 1H), 4.27-4.31 (m, 1H), 4.36 (d, J = 15.2 Hz, 1H), 5.11 (d, J = 10.0 Hz, 1H), 6.81 (s, 1H), 6.87-6.89 (m, 2H), 7.02-7.08 (m, 5H), 7.13 (d, J = 6.8 Hz, 2H), 7.20-7.39 (m, 9H), 7.83 (d, J = 7.6 Hz, 2H); 13C NMR (100.6 MHz, CDCl3, Me4Si) δ 40.96, 52.08, 53.21, 62.40, 66.51, 117.06, 126.28, 126.88, 127.16, 127.58, 127.64, 127.82, 128.17, 128.40, 128.64, 128.90, 130.01, 130.23, 134.78, 139.98, 141.06, 141.39, 155.15, 188.65, 216.64; HRMS (ESI) calcd for C33H28NO2 [M+H]+: 470.2120, found 470.2120.

3-benzoyl-1-hexyl-4,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3g). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate = 5:1) afforded the product in 65% yield as a brown oil. 1H NMR (400 MHz, CDCl3, Me4Si) δ 0.88 (t, J = 6.4 Hz, 3H), 1.27-1.28 (m, 6H), 1.50-1.58 (m, 2H), 2.76 (d, J = 18.4 Hz, 1H), 2.89 (dd, J = 7.2, 18.4 Hz, 1H), 3.10 (t, J = 7.2 Hz, 2H), 4.43 (dd, J = 6.8, 9.6 Hz, 1H), 5.13 (d, J = 10.0 Hz, 1H), 6.69 (s, 1H), 6.84-6.86 (m, 2H), 7.00-7.09 (m, 5H), 7.20-7.25 (m, 2H), 7.29-7.33 (m, 2H), 7.41 (t, J = 7.6 Hz, 2H), 7.89 (d, J = 7.6 Hz, 2H); 13C NMR (100.6 MHz, CDCl3, Me4Si) δ 13.84, 22.37, 26.27, 27.94, 31.17, 41.29, 47.51, 53.24, 63.21, 66.48, 116.59, 126.28, 126.87, 127.20, 127.53, 127.65, 128.48, 128.76, 129.80, 130.22, 140.35, 141.23, 141.45, 155.10, 188.45, 216.89; HRMS (ESI) calcd for C32H34NO2 [M+H]+: 464.2590, found 464.2574.
3-benzoyl-1-(tert-butyl)-4,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3h). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / dichloromethane = 3 : 1 : 1) afforded the product in 90% yield as a yellow oil. $^1$H NMR (400 MHz, CDCl$_3$, Me$_4$Si) δ 1.25 (s, 9H), 2.78 (dd, $J = 1.2$, 18.4 Hz, 1H), 3.02 (dd, $J = 8.4$, 18.8 Hz, 1H), 4.60-4.65 (m, 1H), 5.25 (d, $J = 10.8$ Hz, 1H), 6.86-6.90 (m, 3H), 6.99-7.06 (m, 3H), 7.19-7.42 (m, 8H), 7.79 (d, $J = 7.6$ Hz, 2H); $^{13}$C NMR (100.6 MHz, CDCl$_3$, Me$_4$Si) δ 29.26, 45.89, 54.92, 55.39, 59.47, 66.42, 116.57, 126.23, 126.77, 127.29, 127.65, 127.91, 128.44, 128.62, 130.11, 130.26, 139.88, 140.84, 141.98, 152.55, 188.20, 217.06; HRMS (ESI) calcd for C$_{30}$H$_{30}$NO$_2$ [M+H]$^+$: 436.2277, found 436.2259.

3-(4-chlorobenzoyl)-1,4,4-triphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3i). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / DCM = 2 : 1 : 1) afforded the product in 82% yield as a red solid. M.p. 100-103 °C. $^1$H NMR (400 MHz, CDCl$_3$, Me$_4$Si) δ 2.80 (d, $J = 18.8$ Hz, 1H), 3.06 (dd, $J = 7.6$, 19.0 Hz, 1H), 5.11 (dd, $J = 8.0$, 9.2 Hz, 1H), 5.28 (d, $J = 10.0$ Hz, 1H), 6.85-6.89 (m, 4H), 7.04-7.13 (m, 7H), 7.24-7.35 (m, 5H), 7.42 (t, $J = 7.6$ Hz, 2H), 7.82 (d, $J = 8.0$ Hz, 2H); $^{13}$C NMR (100.6 MHz, CDCl$_3$, Me$_4$Si) δ 41.62, 52.53, 60.46, 67.72, 116.25, 119.90, 123.34, 126.57, 127.12, 127.55, 128.17, 128.60, 129.18, 129.75, 130.34, 136.60, 137.79, 139.01, 140.44, 141.58, 147.76, 187.87, 215.19; HRMS (ESI) calcd for C$_{32}$H$_{25}$ClNO$_2$ [M+H]$^+$: 490.1574, found 490.1574.

1,4,4-triphenyl-3-(3,4,5-trimethoxybenzoyl)-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3j). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 87% yield as a brown solid. M.p. 194-196 °C. $^1$H NMR (400 MHz, CDCl$_3$, Me$_4$Si) δ 2.86 (d, $J = 18.8$ Hz, 1H), 3.07 (dd, $J = 7.2$, 19.0 Hz, 1H), 3.81 (s, 6H), 3.85 (s, 3H), 5.10 (dd, $J = 8.0$, 9.6 Hz, 1H), 5.29 (d, $J = 10.0$ Hz, 1H), 6.40 (s, 2H), 6.83 (d, $J = 8.4$ Hz, 2H), 6.88-6.90 (m, 2H), 7.04-7.13 (m, 4H), 7.20 (s, 1H), 7.33 (t, $J = 7.6$ Hz, 3H), 7.42 (t, $J = 7.6$ Hz,
2H), 7.88 (d, J = 8.0 Hz, 2H); $^{13}$C NMR (100.6 MHz, CDCl$_3$, Me$_4$Si) δ 41.59, 52.24, 56.02, 60.63, 60.69, 66.98, 104.83, 116.18, 120.04, 123.20, 126.42, 126.99, 127.36, 128.43, 128.59, 129.66, 130.21, 134.88, 139.03, 139.88, 140.46, 141.89, 148.03, 152.46, 188.52, 215.60; HRMS (ESI) calcd for C$_{35}$H$_{32}$NO$_5$ [M+H]$^+$: 546.2281, found 546.2264.

3-(1-naphthoyl)-1,4,4-triphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3k). Chromatography on silica gel (elucent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 91% yield as a yellow solid. M.p. 207-209 °C. $^1$H NMR (400 MHz, CDCl$_3$, Me$_4$Si) δ 2.85 (d, J = 18.8 Hz, 1H), 3.05 (dd, J = 7.6, 18.6 Hz, 1H), 5.13 (dd, J = 8.4, 9.0 Hz, 1H), 5.36 (d, J = 10.0 Hz, 1H), 6.77-6.79 (m, 3H), 6.94-6.96 (m, 2H), 7.01-7.04 (m, 2H), 7.18-7.48 (m, 11H), 7.67 (d, J = 8.4 Hz, 1H), 7.75-7.80 (m, 2H), 7.97 (d, J = 7.6 Hz, 2H); $^{13}$C NMR (100.6 MHz, CDCl$_3$, Me$_4$Si) δ 41.51, 52.81, 61.50, 67.21, 116.72, 122.39, 123.57, 124.13, 125.57, 125.71, 126.06, 126.54, 126.67, 127.30, 127.51, 127.87, 128.66, 128.85, 129.69, 130.41, 130.67, 133.49, 137.67, 139.01, 141.00, 141.86, 149.92, 190.27, 215.79; HRMS (ESI) calcd for C$_{36}$H$_{28}$NO$_2$ [M+H]$^+$: 506.2120, found 506.2106.

3-(cyclohexanecarbonyl)-1,4,4-triphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one(3l). Chromatography on silica gel (elucent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 99% yield as a light yellow solid. M.p. 235-238 °C. $^1$H NMR (400 MHz, CDCl$_3$, Me$_4$Si) δ 0.79 (bs, 2H), 1.01-1.07 (m, 3H), 1.18-1.29 (m, 1H), 1.44-1.69 (m, 4H), 2.28 (bs, 1H), 2.78 (d, J = 18.4 Hz, 1H), 2.91 (dd, J = 6.8, 18.6 Hz, 1H), 4.86-4.96 (m, 2H), 6.64 (bs, 2H), 6.86 (d, J = 7.6 Hz, 2H), 6.98-6.70 (m, 4H), 7.22-7.33 (m, 6H), 7.80 (d, J = 7.19 Hz, 2H); $^{13}$C NMR (100.6 MHz, CDCl$_3$, Me$_4$Si) δ 25.40, 25.46, 25.53, 28.80, 29.48, 41.33, 45.75, 52.61, 60.75, 66.82, 116.82, 120.23, 123.37, 126.47, 127.01, 127.52, 128.73, 129.02, 129.97, 130.77, 139.86, 141.17, 141.72, 145.70, 198.57, 216.85; HRMS (ESI) calcd for C$_{32}$H$_{32}$NO$_2$ [M+H]$^+$: 462.2433, found 462.2421.
3-benzoyl-1,2,4,4-tetraphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one (3m). Chromatography on silica gel (eluuent: petroleum ether / ethyl acetate / dichloromethane = 3 : 1 : 1) afforded the product in 53% yield as a dark blue solid. M.p. 256-259 ºC. 1H NMR (400 MHz, CDCl₃, Me₄Si) δ 2.87 (d, J = 18.4 Hz, 1H), 2.95 (dd, J = 6.0, 18.4 Hz, 1H), 5.14 (dd, J = 6.0, 8.8 Hz, 1H), 5.32 (d, J = 8.8 Hz, 1H), 6.35 (bs, 2H), 6.58 (d, J = 8.0 Hz, 2H), 6.66-6.72 (m, 6H), 6.82-6.86 (m, 2H), 7.03-7.06 (m, 3H), 7.12-7.24 (m, 5H), 7.32-7.33 (m, 1H), 7.42 (t, J = 7.2 Hz, 2H), 7.99 (d, J = 8.0 Hz, 2H); 13C NMR (100.6 MHz, CDCl₃, Me₄Si) δ 41.84, 53.43, 64.92, 66.71, 117.89, 125.51, 125.58, 126.44, 126.54, 127.00, 127.04, 127.11, 127.30, 128.28, 128.38, 128.55, 128.93, 129.01, 129.87, 129.96, 130.58, 139.01, 140.61, 141.28, 142.52, 161.29, 192.28, 216.78; HRMS (ESI) calcd for C₃₈H₃₀NO₂ [M+H]+: 532.2277, found 532.2262.

3-pentanoyl-1,2,4,4-tetraphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one (3n). Chromatography on silica gel (eluuent: petroleum ether / ethyl acetate = 5 : 1) afforded the product in 95% yield as a light yellow solid. M.p. 205-208 ºC. 1H NMR (400 MHz, CDCl₃, Me₄Si) δ 0.47 (t, J = 7.6 Hz, 3H), 0.65-0.73 (m, 2H), 0.90-1.00 (m, 2H), 1.18-1.26 (m, 1H), 1.38-1.44 (m, 1H), 2.65 (d, J = 18.4 Hz, 1H), 2.77 (dd, J = 5.6, 18.4 Hz, 1H), 5.02-5.07 (m, 2H), 6.60 (d, J = 7.6 Hz, 2H), 6.66 (d, J = 7.6 Hz, 2H), 6.86-7.32 (m, 14H), 7.84 (d, J = 7.6 Hz, 2H); 13C NMR (100.6 MHz, CDCl₃, Me₄Si) δ 13.16, 21.77, 26.55, 38.53, 41.42, 51.88, 63.75, 66.42, 119.74, 124.88, 125.31, 126.45, 126.99, 127.15, 128.21, 128.42, 128.97, 129.09, 129.43, 130.77, 131.90, 138.88, 141.63, 142.66, 159.23, 196.29, 217.27; HRMS (ESI) calcd for C₃₆H₃₄NO₂ [M+H]+: 512.2590, found 512.2574.

3-benzoyl-4,4-bis(4-chlorophenyl)-1-phenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one (3o). Chromatography on silica gel (eluuent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product in 85% yield as a brown solid. M.p. 85-87 ºC. 1H NMR (400 MHz, CDCl₃, Me₄Si) δ 2.86 (d, J = 18.8 Hz, 1H), 3.04 (dd, J = 7.6, 19.2 Hz, 1H), 5.07-5.11 (m, 1H), 5.17 (d, J = 10.4 Hz, 1H).
3-benzoyl-4-(4-chlorophenyl)-1,4-diphenyl-3a,4,6,6a-tetrahydrocyclopenta[b]pyrrol-5(1H)-one (3p). Chromatography on silica gel (eluent: petroleum ether / ethyl acetate / dichloromethane = 2 : 1 : 1) afforded the product as a mixture of two diastereomers (d.r. = 1 : 1.2) in 91% yield as a brown solid. M.p. 91-93 °C. 1H NMR (400 MHz, CDCl3, Me4Si) δ 2.75 (d, J = 18.8 Hz), 2.87 (d, J = 19.2 Hz), 2.99-3.06 (m), 5.05-5.10 (m), 5.17 (d, J = 10.0 Hz), 5.27 (d, J = 10.0 Hz), 6.80-6.84 (m), 7.01-7.13 (m), 7.18-7.39 (m), 7.79 (d, J = 7.6 Hz), 7.83 (d, J = 9.2 Hz); 13C NMR (100.6 MHz, CDCl3, Me4Si) δ 41.42, 41.71, 52.35, 52.44, 60.28, 60.68, 66.62, 116.19, 116.39, 119.57, 119.96, 123.26, 123.34, 126.65, 127.08, 127.19, 127.53, 127.62, 127.68, 127.88, 128.00, 128.41, 128.44, 128.69, 129.68, 129.69, 129.94, 130.29, 130.50, 130.64, 131.75, 132.39, 133.36, 138.96, 138.97, 139.21, 139.34, 139.46, 140.00, 140.18, 141.67, 148.01, 148.48, 189.26, 189.45, 214.76, 215.63; HRMS (ESI) calcd for C32H25ClNO2 [M+H]+: 490.1574, found 490.1563.
X-ray crystal structure of 3c
Electronic Supplementary Material (ESI) for Chemical Communications
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