

Supporting Information
for
Cu-Mediated trifluoromethylation of benzyl, allyl and
propargyl methanesulfonates with TMSCF₃

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Experimental details, characterization data of all products and copies of NMR spectra.

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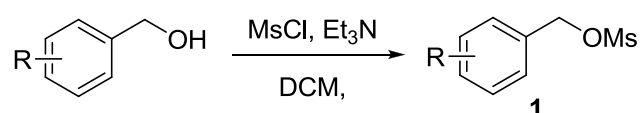
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General information:

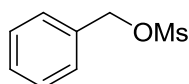
^1H NMR (TMS as the internal standard) and ^{19}F NMR spectra (CFCl_3 as the outside standard and low field is positive) were recorded on a Bruker AM300 or Bruker AM400 spectrometer. ^{13}C NMR was recorded on a Bruker AM400 spectrometer. Chemical shifts (δ) are reported in ppm, and coupling constants (J) are in Hertz (Hz). The following abbreviations were used to explain the multiplicities: s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet, br = broad. Substrates were purchased from commercial sources (Aldrich, Alfa and Chemical Reagent Companies of China) and used as received. Reactions were performed under an atmosphere of nitrogen using glassware that was flame-dried under vacuum. Benzyl alcohols and allylic alcohols were obtained commercially. Propargylic alcohols 1-phenylhex-4-yn-3-ol [1], 1-phenylnon-4-yn-3-ol [2] and 5-phenyl-1-(trimethylsilyl)pent-1-yn-3-ol [3] were prepared according to literature procedures. Compounds **2a** [4], **2b** [5], **2d** [5], **2e** [6], **2f** [7], **2h** [8], **2i-2j** [7] and **2o** [9] have been published previously.

Preparation of substrates:

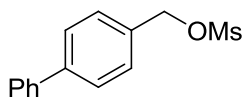
General procedure for the preparation of benzyl, allylic and propargylic methanesulfonates:



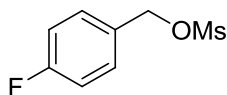
To a stirred solution of benzyl alcohol (5.0 mmol) and triethylamine (10 mmol) in DCM (20 mL) under N_2 atmosphere at $-20\text{ }^\circ\text{C}$ was added methanesulfonyl chloride (7.5 mmol) dropwise. The reaction was stirred for 1 hour (monitored by TLC). The mixture was then allowed to warm to room temperature and washed with water, 2 M HCl and brine, dried over sodium sulfate. The solvent was removed to give the crude product, which was used without purification.



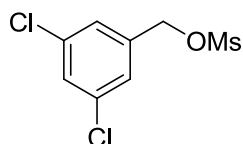
Benzyl methanesulfonate (1a): Colorless oil; yield 91%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.41 (m, 5H), 5.24 (s, 2H), 2.90 (s, 3H).



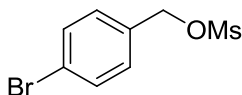
Biphenyl-4-ylmethyl methanesulfonate (1b): White solid; yield 83%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.38-7.65 (m, 9H), 5.29 (s, 2H), 2.95 (s, 3H).



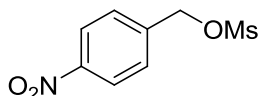
4-Fluorobenzyl methanesulfonate (1c): Colorless oil; yield 91%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.39-7.41 (m, 2H), 7.10-7.13 (m, 2H), 5.13 (s, 2H), 2.93 (s, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -111.5 (m, 1F).



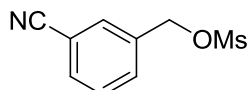
3,5-Dichlorobenzyl methanesulfonate (1d): White solid; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.39 (s, 1H), 7.31 (s, 2H), 5.17 (s, 2H), 3.04 (s, 3H).



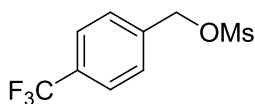
4-Bromobenzyl methanesulfonate (1e): White solid; yield 98%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.55 (d, J = 8.4 Hz, 2H), 7.30 (d, J = 8.4 Hz, 2H), 5.19 (s, 2H), 2.95 (s, 3H).



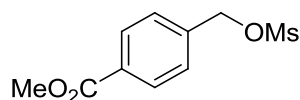
4-Nitrobenzyl methanesulfonate (1f): White solid; yield 97%. ^1H NMR (300 MHz, CDCl_3) δ ppm 8.29 (d, J = 8.4 Hz, 2H), 7.74 (m, J = 8.4 Hz, 2H), 5.46 (s, 2H), 3.34 (s, 3H).



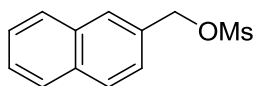
3-Cyanobenzyl methanesulfonate (1g): White solid; yield 95%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.52-7.72 (m, 4H), 5.26 (s, 2H), 3.05 (s, 3H).



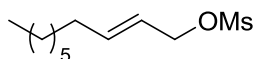
4-(Trifluoromethyl)benzyl methanesulfonate (1h): Colorless oil; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.68 (d, $J = 7.2$ Hz, 2H), 7.54 (m, $J = 7.2$ Hz, 2H), 5.29 (s, 2H), 3.01 (s, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -62.8 (s, 3F).



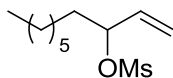
Methyl 4-((methanesulfonyloxy)methyl)benzoate (1i): White solid; yield 98%. ^1H NMR (300 MHz, CDCl_3) δ ppm 8.08 (d, $J = 8.4$ Hz, 2H), 7.49 (d, $J = 8.4$ Hz, 2H), 5.29 (s, 2H), 2.94 (s, 3H), 2.98 (s, 3H).



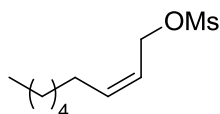
Naphthalen-2-ylmethyl methanesulfonate (1j): White solid; yield 93%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.85-7.91 (m, 4H), 7.50-7.55 (m, 3H), 5.41 (s, 2H), 2.91 (s, 3H).



(E)-dec-2-enyl methanesulfonate (1k): Colorless oil; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 5.88-5.97 (m, 1H), 5.56-5.66 (m, 1H), 4.69 (d, $J = 6.6$ Hz, 2H), 3.01 (s, 3H), 2.06-2.12 (m, 2H), 1.27-1.41 (m, 10H), 0.88 (t, $J = 6.9$ Hz, 3H).

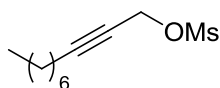


Dec-1-en-3-yl methanesulfonate (1l): Colorless oil; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 5.80-5.92 (m, 1H), 5.32-5.44 (m, 2H), 4.97-5.04 (m, 1H), 2.98 (s, 1H), 1.28-1.82 (m, 12H), 0.88 (t, $J = 6.6$ Hz, 3H).

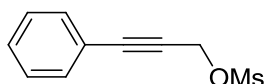


(Z)-non-2-enyl methanesulfonate (1m): Colorless oil; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 5.76-5.85 (m, 1H), 5.55-5.63 (m, 1H), 4.80 (d, $J = 6.9$ Hz, 2H), 3.01 (s, 3H), 2.10-2.17 (m,

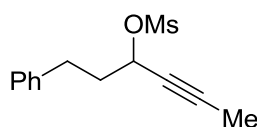
2H), 1.28-1.39 (m, 8H), 0.88 (t, $J = 6.6$ Hz, 3H).



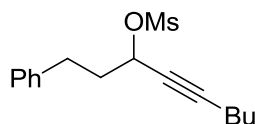
Dec-2-ynyl methanesulfonate (1n): Colorless oil; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 4.86 (m, 2H), 3.11 (s, 3H), 2.25 (t, $J = 6.9$ Hz, 2H), 1.28-1.58 (m, 10H), 0.89 (t, $J = 6.9$ Hz, 3H).



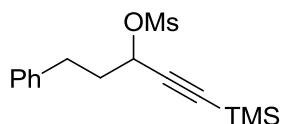
3-Phenylprop-2-ynyl methanesulfonate (1o): Yellow oil; yield 97%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.35-7.48 (m, 5H), 5.09 (s, 2H), 3.17 (s, 3H).



1-Phenylhex-4-yn-3-yl methanesulfonate (1p): Colorless oil; yield 83%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.19-7.32 (m, 5H), 5.12-5.17 (m, 1H), 3.11 (s, 3H), 2.78-2.84 (m, 2H), 2.12-2.22 (m, 2H), 1.91 (d, $J = 1.8$ Hz, 3H).



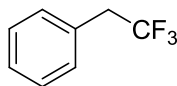
1-Phenylnon-4-yn-3-yl methanesulfonate (1q): Colorless oil; yield 93%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.19-7.32 (m, 5H), 5.16 (t, $J = 6.6$ Hz, 1H), 3.11 (s, 1H), 2.81-2.82 (m, 2H), 2.18-2.29 (m, 4H), 1.42-1.52 (m, 4H), 0.92 (t, $J = 6.9$ Hz, 3H).



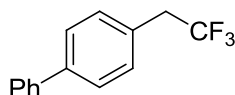
5-Phenyl-1-(trimethylsilyl)pent-1-yn-3-yl methanesulfonate (1r): Colorless oil; yield 100%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.19-7.32 (m, 5H), 5.13 (t, $J = 6.6$ Hz, 1H), 3.14 (s, 3H), 2.78-2.85 (m, 2H), 2.14-2.25 (m, 2H), 0.20 (s, 9H).

General procedure for the Cu-mediated trifluoromethylation of benzyl methanesulfonates:

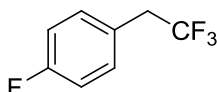
Into a Schlenk tube equipped with a magnetic stir bar were added CuI (2.2 mmol) and KF (4.0 mmol) under Ar atmosphere. DMF (5.0 mL) and Me₃SiCF₃ (2.0 equiv) were added. After stirring for 20 minutes, the mixture was heated to 60 °C and then the benzyl methanesulfonate (2.0 mmol) was added under N₂ atmosphere. The reaction mixture was kept at 60 °C for 4 hours and then allowed to cool down to room temperature. The resulting mixture was diluted with diethyl ether, washed with water and brine, dried over sodium sulfate, and concentrated. The crude products were purified by column chromatography on silica gel to give the products.



(2,2,2-Trifluoroethyl)benzene (2a): Colorless liquid; yield 76% after distillation on 10 mmol scale. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.27-7.32 (m, 5H), 3.31 (q, *J* = 10.8 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -66.0 (t, *J* = 11.6 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 130.2, 128.7, 128.1, 125.9 (q, *J* = 275.5 Hz), 40.2 (q, *J* = 29.6 Hz). HRMS (EI) Calculated for C₈H₇F₃; 160.0500; Found: 160.0502.

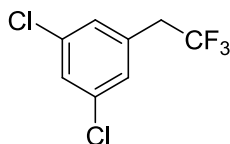


4-(2,2,2-Trifluoroethyl)biphenyl (2b): White solid; yield 79%; mp 81-82 °C. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.36-7.60 (m, 9H), 3.41 (q, *J* = 10.5 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -65.8 (t, *J* = 11.3 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 141.1, 140.5, 130.6, 129.2, 128.9, 127.6, 127.4, 127.1, 125.9 (q, *J* = 275.6 Hz), 39.9 (q, *J* = 29.6 Hz). Anal. Calcd. for C₁₄H₁₁F₃: C, 71.18; H, 4.69. Found: C, 71.17; H, 4.69.

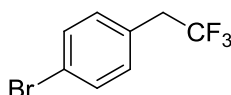


1-Fluoro-4-(2,2,2-trifluoroethyl)benzene (2c): Colorless liquid; yield 75% after distillation on 10 mmol scale. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.22-7.27 (m, 2H), 7.00-7.05 (m, 2H), 3.32 (q, *J* = 10.8 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -66.4 (t, *J* = 11.3 Hz, 3F), -114.2 (s, 1F). ¹³C

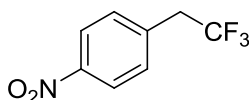
NMR (100 MHz, CDCl₃) δ ppm 162.9 (d, J = 245.9 Hz), 132.0 (d, J = 8.4 Hz), 126.1, 125.8 (q, J = 275.5 Hz), 115.8 (d, J = 22.0 Hz), 39.6 (q, J = 29.6 Hz). HRMS (EI) Calculated for C₈H₆F₄: 178.0406; Found: 178.0403.



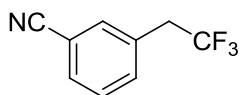
1,3-Dichloro-5-(2,2,2-trifluoroethyl)benzene (2d): White solid; yield 72%; mp 46-47 °C. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.36 (s, 1H), 7.20 (s, 2H), 3.33 (q, J = 10.5 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -65.7 (t, J = 10.0 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 135.1, 133.3, 128.8, 128.7, 125.3 (q, J = 276.1 Hz), 39.7 (q, J = 30.0 Hz). Anal. Calcd. for C₈H₅Cl₂F₃: C, 49.95; H, 2.20. Found: C, 41.89; H, 2.21.



1-Bromo-4-(2,2,2-trifluoroethyl)benzene (2e): White solid; yield 78%; mp 55-57 °C. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.48 (d, J = 6.6 Hz, 2H), 7.15 (d, J = 7.8 Hz, 2H), 3.30 (q, J = 10.5 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -66.0 (t, J = 10.2 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 132.0, 131.9, 129.2, 125.6 (q, J = 274.8 Hz), 39.8 (q, J = 29.6 Hz). HRMS (EI) Calculated for C₈H₆BrF₃: 237.9605; Found: 237.9607.

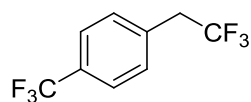


1-Nitro-4-(2,2,2-trifluoroethyl)benzene (2f): White solid; yield 40%; mp 65-66 °C. ¹H NMR (300 MHz, CDCl₃) δ ppm 8.24 (d, J = 8.4 Hz, 2H), 7.50 (d, J = 8.4 Hz, 2H), 3.50 (q, J = 10.4 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -65.5 (t, J = 10.9 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 148.1, 137.4, 131.3, 125.2 (q, J = 276.6 Hz), 124.0, 40.1 (q, J = 30.6 Hz). HRMS (EI) Calculated for C₈H₆F₃NO₂: 205.0351; Found: 205.0347.

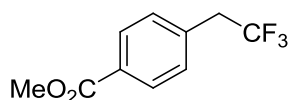


3-(2,2,2-Trifluoroethyl)benzonitrile (2g): White solid; yield 81%; mp 65-67 °C. ¹H NMR (300

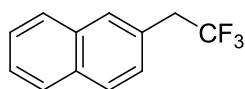
MHz, CDCl₃) δ ppm 7.47-7.66 (m, 4H), 3.43 (q, J = 10.8 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -65.9 (t, J = 10.2 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 134.6, 133.6, 131.8, 125.2 (q, J = 275.5 Hz), 118.2, 113.0, 39.7 (q, J = 30.4 Hz). HRMS (EI) Calculated for C₉H₆F₃N: 185.0452; Found: 185.0452.



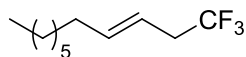
1-(2,2,2-Trifluoroethyl)-4-(trifluoromethyl)benzene (2h): Colorless liquid; yield 80%. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.63 (d, J = 8.1 Hz, 2H), 7.43 (d, J = 7.8 Hz, 2H), 3.44 (q, J = 10.5 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -62.7 (s, 3F), -65.7 (t, J = 11.3 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 134.3, 130.7, 129.0, 125.8 (q, J = 4.2 Hz), 125.6 (q, J = 275.3 Hz), 124.2 (q, J = 270.8 Hz), 40.2 (q, J = 30.1 Hz). HRMS (EI) Calculated for C₉H₆F₆: 228.0374; Found: 228.0375.



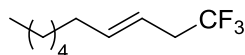
Methyl 4-(2,2,2-trifluoroethyl)benzoate (2i): White solid; yield 68%; mp 54-55 °C. ¹H NMR (300 MHz, CDCl₃) δ ppm 8.03 (d, J = 8.7 Hz, 2H), 7.38 (d, J = 8.1 Hz, 2H), 3.92 (s, 2H), 3.43 (q, J = 10.5 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -65.6 (t, J = 10.7 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 166.7, 135.2, 130.3, 130.2, 130.0, 125.6 (q, J = 275.7 Hz), 52.2, 40.2 (q, J = 29.3 Hz).



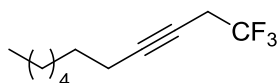
2-(2,2,2-Trifluoroethyl)naphthalene (2j): White solid; yield 78%; mp 54-55 °C. ¹H NMR (300 MHz, CDCl₃) δ ppm 7.39-7.85 (m, 7H), 3.53 (q, J = 11.1 Hz, 2H). ¹⁹F NMR (282 MHz, CDCl₃) δ ppm -65.6 (t, J = 11.3 Hz, 3F). ¹³C NMR (100 MHz, CDCl₃) δ ppm 133.4, 133.0, 129.6, 128.5, 127.9, 127.8, 127.7, 126.5, 126.4, 126.1 (q, J = 275.6 Hz), 40.4 (q, J = 29.8 Hz). Anal. Calcd. for C₁₂H₉F₃: C, 68.57; H, 4.32. Found: C, 68.57; H, 4.36.



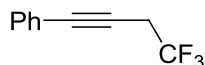
(E)-1,1,1-Trifluoroundec-3-ene (2k): Colorless liquid; yield 78% from **1k**, 80% from **1l**. ^1H NMR (300 MHz, CDCl_3) δ ppm 5.65-5.73 (m, 1H), 5.33-5.40 (m, 1H), 2.70-2.79 (m, 2H), 2.02-2.07 (m, 2H), 1.28-1.39 (m, 10H), 0.88 (t, $J = 6.8$ Hz, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -67.0 (t, $J = 9.4$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 138.7, 126.3 (q, $J = 274.9$ Hz), 117.7 (q, $J = 4.3$ Hz), 37.6 (q, $J = 29.0$ Hz), 32.7, 32.1, 29.4, 29.3, 29.2, 22.9, 14.2. HRMS (EI) Calculated for $\text{C}_{11}\text{H}_{19}\text{F}_3$: 208.1439; Found: 208.1438.



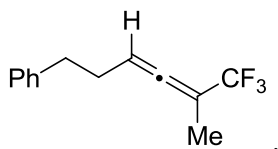
(E)-1,1,1-Trifluorodec-3-ene (2m): Colorless liquid; yield 87%. ^1H NMR (300 MHz, CDCl_3) δ ppm 5.66-5.73 (m, 1H), 5.33-5.40 (m, 1H), 2.71-2.81 (m, 2H), 2.02-2.07 (m, 2H), 1.28-1.39 (m, 8H), 0.88 (t, $J = 4.8$ Hz, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -66.8 (t, $J = 8.2$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 138.7, 126.3 (q, $J = 275.0$ Hz), 117.7 (q, $J = 3.6$ Hz), 37.6 (q, $J = 28.6$ Hz), 32.7, 31.9, 29.1, 28.9, 22.8, 14.2. IR (neat) 2960, 2930, 1367, 1253, 1138 cm^{-1} . HRMS (EI) Calculated for $\text{C}_{10}\text{H}_{17}\text{F}_3$: 194.1282; Found: 194.1279.



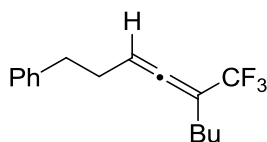
1,1,1-Trifluoroundec-3-yne (2n): Colorless liquid; yield 64%. ^1H NMR (300 MHz, CDCl_3) δ ppm 2.98 (q, $J = 7.5$ Hz, 2H), 2.16 (t, $J = 5.4$ Hz, 2H), 1.28-1.54 (m, 10H), 0.88 (t, $J = 5.1$ Hz, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -67.3 (t, $J = 9.8$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 124.7 (q, $J = 274.2$ Hz), 85.2, 68.3 (q, $J = 4.8$ Hz), 31.9, 28.94, 28.89, 28.6, 26.3 (q, $J = 33.7$ Hz), 22.8, 18.7, 14.2. IR (neat) 2933, 2859, 2245, 1366, 1258, 1113 cm^{-1} . Anal. Calcd. for $\text{C}_{11}\text{H}_{17}\text{F}_3$: C, 64.06; H, 8.31. Found: C, 64.09; H, 8.32.



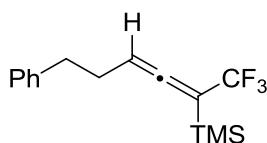
(4,4,4-Trifluorobut-1-ynyl)benzene (2o): Yellow liquid; yield 50%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.46-7.31 (m, 5H), 3.27 (q, $J = 9.6$ Hz, 2H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -66.0 (t, $J = 11.6$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 131.0, 128.9, 128.5, 124.4 (q, $J = 275.1$ Hz), 122.3, 84.5, 77.5 (q, $J = 4.1$ Hz), 26.9 (q, $J = 35.1$ Hz). HRMS (EI) Calculated for $\text{C}_{10}\text{H}_7\text{F}_3$: 184.0500; Found: 184.0498.



(6,6,6-Trifluoro-5-methylhexa-3,4-dienyl)benzene (2p): Colorless liquid; yield 89%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.17-7.31 (m, 5H), 5.56 (m, 1H), 2.74 (t, $J = 7.5$ Hz, 2H), 2.37-2.44 (m, 2H), 1.73 (d, $J = 2.7$ Hz, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -65.8 (d, $J = 4.2$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 202.7, 141.1, 128.6, 128.5, 126.2, 124.1 (q, $J = 271.9$ Hz), 96.5, 94.3 (q, $J = 34.3$ Hz), 34.9, 29.7, 13.0. IR (neat) 3065, 3029, 2932, 1981, 1455, 1307, 1118 cm^{-1} . Anal. Calcd. for $\text{C}_{13}\text{H}_{13}\text{F}_3$: C, 69.02; H, 5.79. Found: C, 69.00; H, 5.72.



(5-(Trifluoromethyl)nona-3,4-dienyl)benzene (2q): Colorless liquid; yield 93%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.17-7.31 (m, 5H), 5.60-5.65 (m, 1H), 2.74 (t, $J = 7.8$ Hz, 2H), 2.37-2.46 (m, 2H), 2.05 (m, 2H), 1.31-1.34 (m, 4H), 0.89 (t, $J = 7.2$ Hz, 3H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -64.3 (d, $J = 3.1$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 202.4, 141.1, 128.6, 128.5, 126.2, 124.2 (q, $J = 273.2$ Hz), 99.5 (q, $J = 34.3$ Hz), 97.9, 35.1, 29.9, 29.5, 26.0, 22.2, 13.9. IR (neat) 3066, 2959, 2864, 1976, 1455, 1298, 1118 cm^{-1} . HRMS (EI) Calculated for $\text{C}_{16}\text{H}_{19}\text{F}_3$: 268.1439; Found: 268.1440.



Trimethyl(1,1,1-trifluoro-6-phenylhexa-2,3-dien-2-yl)silane (2r): Colorless liquid; yield 70%. ^1H NMR (300 MHz, CDCl_3) δ ppm 7.18-7.32 (m, 5H), 5.34-5.38 (m, 1H), 2.73 (t, $J = 7.8$ Hz, 2H), 2.36-2.43 (m, 2H), 0.16 (s, 9H). ^{19}F NMR (282 MHz, CDCl_3) δ ppm -55.8 (d, $J = 4.2$ Hz, 3F). ^{13}C NMR (100 MHz, CDCl_3) δ ppm 208.7, 141.2, 128.63, 128.58, 126.3, 125.2 (q, $J = 271.6$ Hz), 94.4 (q, $J = 34.7$ Hz), 90.9, 35.5, 29.2, -1.2. IR (neat) 3066, 2960, 1956, 1269, 1113 cm^{-1} . Anal. Calcd. for $\text{C}_{15}\text{H}_{19}\text{F}_3\text{Si}$: C, 63.35; H, 6.73. Found: C, 62.98; H, 6.82.

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