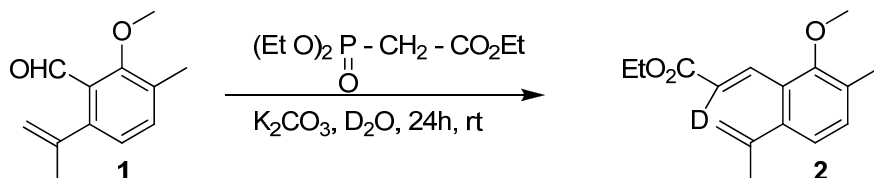
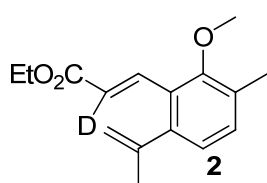


Deuterium Experiment:



Compound 2: Dry potassium carbonate (431 mg, 3.12 mmoles), deuterium oxide (2 ml) and triethylphosphonoacetate (0.265 mg, 1.18 mmoles) are stirred vigorously

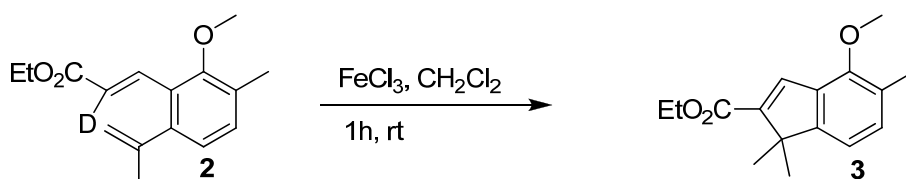


in a dry flask for 20 hours at room temperature under nitrogen atmosphere¹. Aldehyde (150 mg, 0.78 mmoles) is then introduced and stirring is continued for 24 hours. Extraction (diethylether, 10ml x 3) is

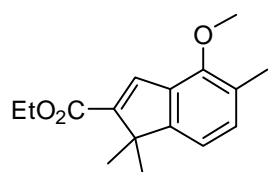
performed after addition of water (5 ml), and the combined organic layers were washed with brine and dried over anhydrous Na₂SO₄. The solvent was removed under reduced pressure. The crude product was purified on silica gel column chromatography using EtOAc-hexane as eluent to furnish the product (185 mg, 91%) as a yellow solid. *R_f* = 0.51 (EtOAc-hexane 3:97); **IR** (neat): $\nu_{\text{max}}/\text{cm}^{-1}$ 1713, 1619, 1593, 1476, 1218; **¹H NMR** (500 MHz, CDCl₃) δ 1.33 (t, *J* = 7.3 Hz, 3H), 2.01 (s, 3H), 2.29 (s, 3H), 3.67 (s, 3H), 4.25 (q, *J* = 7.3 Hz, 2H), 4.90 (br s, 1H), 5.24 (br s, 1H), 6.88 (d, *J* = 7.9 Hz, 1H), 7.12 (d, *J* = 7.9 Hz, 1H), 7.82 (s, 1H); **¹³C NMR** (125 MHz, CDCl₃) δ 14.3, 15.9, 24.6, 59.8, 60.3, 116.7, 122.0(q), 124.3, 125.0, 130.4, 132.0, 139.8, 144.5, 144.8, 157.8, 167.7; **HRMS:** *m/z* calcd for C₁₆H₁₉DNaO₃ [(M+Na)⁺]: 284.1373; Found: 284.1373.

1. Pascale Segueineau, Jean Villieras, *Tetrahedron Letters.*, 1988, **29**, 477

Cyclization reaction on deuterated diene ester



Compound 3: Under inert atmosphere of argon, to a stirred solution of diene ester (50 mg, 0.19



mmol) in CH₂Cl₂ (50ml) and FeCl₃ (62 mg, 0.38 mmol) was added at 0°C and stirred for 1h at RT. The reaction progress was monitored by TLC and after completion the reaction was quenched by sodium bisulphate.

The reaction mixture was extracted with CH₂Cl₂ and the combined organic layers were washed with brine and dried over anhydrous Na₂SO₄. The solvent was removed under reduced pressure. The crude product was purified on silica gel column chromatography using EtOAc-hexane as an eluent to furnish the cyclised product (46 mg, 93%) as a yellow solid. *R_f* = 0.51 (EtOAc-hexane 3:97); **IR** (neat): $\nu_{\text{max}}/\text{cm}^{-1}$ 2979, 2928, 1704, 1568, 1478, 1419, 1244; **¹H NMR** (200 MHz, CDCl₃) δ 1.38 (t, *J* = 7.0 Hz, 3H), 1.44 (s, 6H), 2.29 (s, 3H), 3.94 (s, 3H), 4.3 (q, *J* = 7 Hz, 2H), 7.00 (d, *J* = 7.4 Hz, 1H), 7.17 (d, *J* = 7.4 Hz, 1H), 7.79 (s, 1H); **¹³C NMR** (100 MHz, CDCl₃) δ 14.4, 15.8, 24.2, 49.8, 60.0, 61.5, 116.7, 128.5, 130.5, 131.3, 136.0, 145.0, 153.8, 157.0, 164.4; **HRMS** (C₁₆H₂₁O₃): Calc'd 261.1491 [(M+H)⁺], Found 261.1495.

