

**(±)-1-(4-Hydroxy-3-methoxyphenyl)-3-butanol****Guy L. Plourde**

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The discussion and purpose for the synthesis of this compound has been reported elsewhere [1]. To a cold (0°C) solution of 1-(4-hydroxy-3-methoxyphenyl)-3-butanone (735 mg, 3.8 mmol) in EtOH (35 mL) was added sodium borohydride (145 mg, 3.8 mmol, 1 eq). The solution was stirred at 0°C for 30 min., then at room temperature for 2 h. 10% HCl (15 mL) was added and the solution was stirred at room temperature for 2 h. The solution was concentrated in vacuo and the aqueous residue was extracted with ethyl acetate (3 x 20 mL). The organic fractions were combined, dried (MgSO<sub>4</sub>) and the solvent was evaporated in vacuo. Chromatography on silica gel (40% EtOAc/hexanes) afforded a colorless oil (600 mg, 81%).

IR (neat) cm<sup>-1</sup>: 3311 (OH)<sup>1</sup>H-NMR (CDCl<sub>3</sub>) δ: 1.24 (d, 3H, J=6.2 Hz, H-4), 1.76 (m, 2H, H-2), 2.68 (m, 2H, H-1), 3.82 (m, 1H, H-3), 3.89 (s, 3H, OCH<sub>3</sub>), 5.53 (s, 1H, exchangeable with D<sub>2</sub>O, OH), 6.67 (d, 1H, J=7.4 Hz, ArH-6), 6.72 (s, 1H, ArH-2), 6.84 (d, 1H, J=7.4 Hz, ArH-5).<sup>13</sup>C-NMR (CDCl<sub>3</sub>) δ: 23.7 (C-4), 32.1 (C-2), 41.3 (C-1), 56.1 (OCH<sub>3</sub>), 67.8 (C-3), 111.1 (ArC-2), 114.4 (ArC-5), 121.1 (ArC-6), 134.2 (ArC-1), 143.9 (ArC-4), 146.6 (ArC-3).MS m/e (rel %): 196 [M<sup>+</sup>] (97), 178 (12), 163 (27), 138 (86), 137 (100), 134 (42), 123 (22), 106 (16), 91 (15).Anal. calc. for C<sub>11</sub>H<sub>16</sub>O<sub>3</sub>: C 67.31, H 8.74; found: C 67.03, H 8.55.**Acknowledgment**

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**Reference**1. Plourde, G.L. *Tetrahedron Letters* **2002**, *43*, 3597-3599.

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