

2.5.3 Preparation of 2-methyl-1-[4-(2-methylpiperidin-1-yl)but-2-yn-1-yl]-2,3-dihydro-1H-indole (AZ-4)

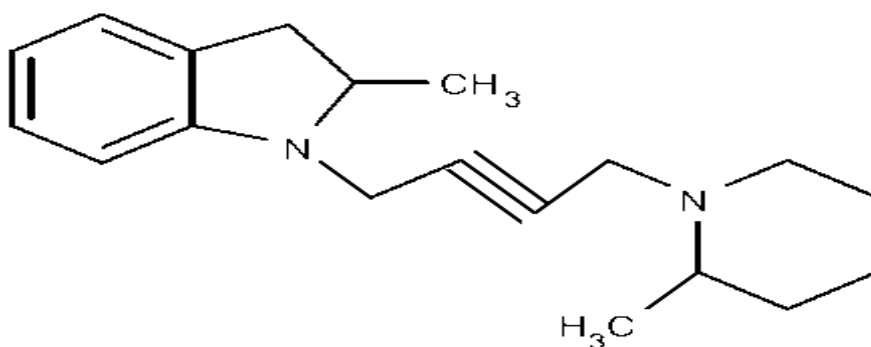


Figure 25: 2-methyl-1-[4-(2-methylpiperidin-1-yl)but-2-yn-1-yl]-2,3-dihydro-1H-indole.

The titled compound was prepared following the general procedure for synthesis of 2-methyl-1-[4-(amino-1-yl)but-2-yn-1-yl]-2,3-dihydro-1H-indole, AZ2-AZ7, yielded 2.7 gm 95.6 %. **IR (NaCl, Cm^{-1}):** 3048, 2963, 2929 (ArH, stretch), 1607, 1481, 1460 (Ar, C=C, stretch), 1374, 1234, 1065 (Ar, C=C, bending), 849, 749 (ArH, bending). **$^1\text{H-NMR}$ (DMSO- d_6):** δ 1.22 (d, 3H, CH-CH₃), 1.26 (d, 3H, $J = 4.4$ Hz, N-C-CH₃), 1.16, 1.33, 1.50, 1.51, 1.62, 1.72, 2.37, 2.95 (m, various protons of cyclicamine), 3.06 (d, 1H, CH₂-CH-N), 3.07, 3.47 (d, 2H, $J = 2.4$ Hz, CH₂-N) due to long range coupling, 3.48, 3.87 (d, 2H, $J = 2.4$ Hz, CH₂-C) due to long range coupling, 3.66 (d, 1H, CH₂-CH-N), 3.82 (m, 1H, $J = 6.15$ Hz, N-CH-CH₃), 6.81-7.28 (m, 4H, ArH). **DSC:** melting point = 98 °C. **$^{13}\text{C-NMR}$ (DMSO- d_6):** δ , 19 (C^{28, 39}), 20 (C³¹), 24 (C¹⁷), 26 (C²¹), 34 (C⁷), 43 (C^{26, 27, 42}), 53 (C³⁰), 59 (C¹⁴), 78 (C²⁴), 80 (C²⁵), 109 (C³), 117 (C⁴), 124 (C^{1, 5}), 127 (C⁶), 151 (C²). **Anal. Calcd: (C₁₉H₂₆N₂):** C (80.8%); H (9.28%); N (9.92%). Found C (81.2%); H (9.22%); N (9.52%).