## 2.5.1 Preparation of 1-[4-(2,6-dimethylpiperidin-1-yl)but-2-yn-1-yl]-2-methyl-2,3-dihydro-1H-indole (AZ-2)

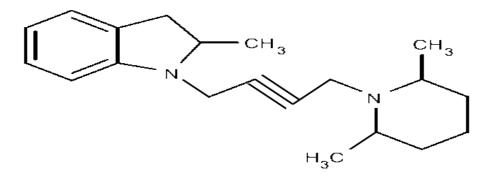


Figure 23: 1-[4-(2,6-dimethylpiperidin-1-yl)but-2-yn-1-yl]-2-methyl-2,3-dihydro-1H-indole.

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 2.72 gm 91.7 %. **IR** (**NaCl**, **Cm**<sup>-1</sup>): 3048, 2964, 2926 (ArH, stretch), 1672, 1607, 1481, 1460 (Ar, C=C, stretch), 1091, 1057 (Ar,C=C, bending), 849, 749 (ArH, bending). <sup>1</sup>**H-NMR** (**DMSO-d**<sub>6</sub>):  $\delta$ , 1.16, 1.22, 1.34, 1.50, 1.66 (m, various protons of cyclicamine), 1.22 (d,3H, CH-C $\underline{\text{H}}_3$ ), 1.23-1.27 (q, 6H, J = 4.4 Hz, N-C-CH<sub>3</sub>) H splitting each one into doublet , 3.05 (m, 2H, N-C $\underline{\text{H}}$ , CH<sub>3</sub>), 3.06 (d, 1H, C $\underline{\text{H}}$ -CH-N), 3.11, 3.51 (t, 2H, J = 2.4 Hz, C-CH<sub>2</sub>-N) due to long range coupling, 3.46, 3.85 (t, 2H, J = 2.4 Hz, CH<sub>2</sub>-C) due to long range coupling, 3.83 (m, 1H, J = 6.15 Hz), 6.81-7.28 (m, 4H, ArH). <sup>13</sup>**C-NMR** (**DMSO-d**<sub>6</sub>):  $\delta$ , 19 (  $C^{28, 39}$ ), 21 ( $C^{17, 31}$ ), 24 ( $C^{21}$ ), 35 ( $C^{7}$ ), 37 ( $C^{26, 27, 39}$ ), 41 ( $C^{29,30}$ ), 53 ( $C^{14}$ ), 77 ( $C^{24}$ ), 80 ( $C^{25}$ ), 109 ( $C^{3}$ ), 118 ( $C^{4}$ ), 124 ( $C^{1, 5}$ ), 127 ( $C^{6}$ ), 151 ( $C^{2}$ ). DSC: melting point = 109  $C^{\circ}$ .