

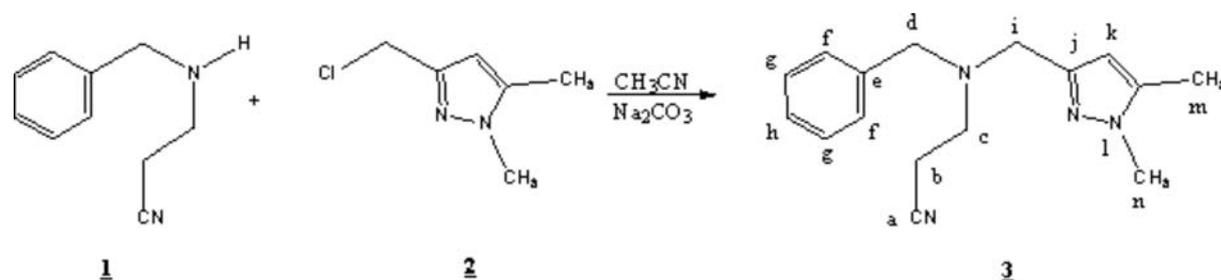
3-[Benzyl-(1,5-dimethyl-1H-pyrazol-3-ylmethyl)-amino]-propionitrileLeila Herrag,^{1,2*} Rachid Touzani,^{1,3} Abdelkrim Ramdani¹ and Belkheir Hammouti²¹ Laboratoire de Chimie Organique Physique, Département de Chimie, Faculté des Sciences, Université Mohammed Premier, B.P. 524, 60000 Oujda, Maroc.² Laboratoire de Chimie des Eaux et Corrosion, Département de Chimie, Faculté des Sciences, B.P. 717, Oujda, Maroc.³ Université Mohammed Premier, Faculté Pluridisciplinaire de Nador B.P. 300, 62700 Selouane, Nador, MarocE-mail: herragleila@yahoo.com

*Author to whom correspondence should be addressed

Received: 19 July 2006 / Accepted: 31 July 2006 / Published: 1 September 2006

Keywords: pyrazole; propionitrile and tridentate nitrogen ligand.

The products of aza-type *Michael* addition, *i.e.*, β -amino carbonyl compounds and their derivatives, are often used as peptide analogs or precursors of optically active amino acids, amino alcohols, diamines, and lactams [1]. Moreover, β -amino carbonyl functionalities are ubiquitous motifs in natural products such as alkaloids and polyketides [2]. Herein, we report the synthesis of new product using aza-type *Michael* reactions under mild conditions.



To a mixture of 3-chloromethyl-1,5-dimethylpyrazole **1** [3] (0.51 g, 3.53 mmol) and sodium carbonate (1.5 g, 14.12 mmol) in 32 mL of acetonitrile was added slowly 3-(benzylamino)propionitrile **2** [4] (0.56 g, 3.53 mmol). The mixture was stirred at room temperature for four days. The solid material (Na₂CO₃, NaCl) was filtered and the filtrate was concentrated under reduced pressure to give the title compound **3** as yellow oil (85%).

¹H-NMR (300 MHz, CDCl₃): δ = 7.24-7.40 (CH_{arom}, 5H, m); 6.02 (CH_{pyr}, 1H, s); 3.68 (N-CH₃, 3H, s); 3.66 (N-CH₂-C, 2H, s); 3.62 (C₆H₅-CH₂, 2H, s); 2.76-2.81 (CH₂-CH₂-CN, 2H, t, J = 7.66 Hz); 2.42-2.47 (N-CH₂, t, J = 7.66 Hz); 2.24 (CH₃, 3H, s).

¹³C-NMR (CDCl₃, 75 MHz): δ = 147.91(j); 139.85(l); 138.93(e); 129.29 and 129.19 (f); 128.79(g); 127.65(h); 119.37(a); 105.64(k); 58.67(d); 51.22(i); 48.82(c); 36.28(n); 16.63(b); 11.64(m).

EI-MS (70 eV, m/z): 228; 173 (2.4); 119; 95(4.7); 91 (37.3).

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