

Synthesis of 6-chloro-2-oxo-1,2-dihydroquinoline-4-carbohydrazide

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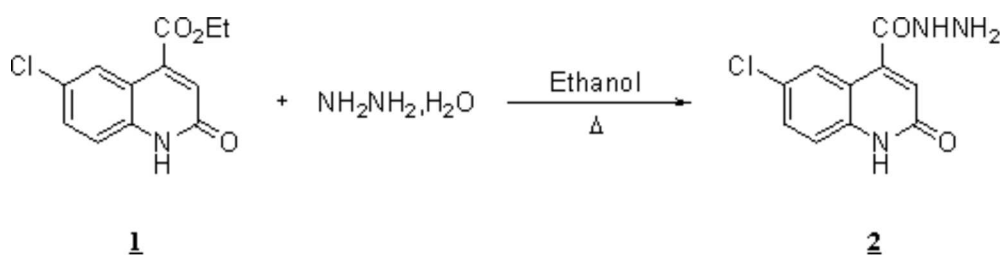
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In previous works we have shown that hydrazides are highly useful starting materials and intermediates in the synthesis of several heterocyclic compounds¹⁻³ of potential biological activities.

The aim of this work is to describe the preparation of a novel compound entitled 6-chloro-2-oxo-1,2-dihydroquinoline-4-carbohydrazide.



To a solution of ethyl 2-oxo-1,2-dihydroquinoline-4-carboxylate **1** (1g, 3.9 mmol) in ethanol, was added hydrazine hydrate 80 % (0.22 mL, 4.6 mmol). The mixture was refluxed for 24h and ice-water was added. The precipitate was filtered and recrystallised from ethanol to afford 0.73 g (70 % yields) of product **2**.

Melting point: > 250 °C.

¹H-NMR (300 MHz, DMSO): δ= 4.67 (NH₂); 6.54 (s, 1H, =CH); 7.35-7.78 (m, 3H, H_{Ar}); 9.93 (NH).

¹³C-NMR (300 MHz, DMSO): δ= 121.8 (=CH); 118.0, 125.4, 131.2 (CH_{Ar}); 114.8, 126.3, 139.3, 139.9 (Cq); 161.3 (C=O); 165.0 (CON₂H₃).

MS (EI, m/z): 237.

Elemental analysis: Calculated for C₁₀H₈ClN₃O₂: C, 50.54 %; H, 3.39 %; N, 17.68 %; Found: C, 50.60 %; H, 3.42 %; N, 17.74 %;

Reference

- 1- Essassi E.M., Fifani J., Bull. Soc. Chem. Belg. 1987, 96, 63.
- 2- El Otmani B., El Mahi M., Essassi E.M., C. R. Chimie, 2002, 5, 517.
- 3- El Otmani B., El Hakmaoui A., Fifani J., Essassi E.M., Gueiffier A., C. R. Acad. Sci. Paris t-2, série IIc, 2001, 4, 285.

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