

SHORT  
COMMUNICATIONS

## A Simple Synthesis of 2-(2-Chlorophenyl)benzimidazole from *o*-Phenylenediamine and 2-Chlorobenzaldehyde\*

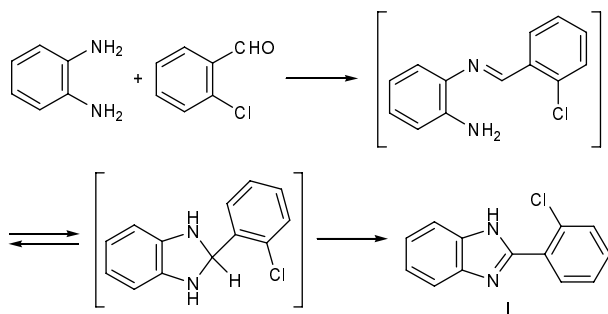
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Benzimidazoles are well known as herbicides, fungicides, and both medicines and veterinary agents. One of the most convenient methods for the preparation of benzimidazoles is condensation of *o*-phenylenediamine with carboxylic acids [1]. Another way of synthesis of these compounds is reaction of *o*-phenylenediamine with benzaldehydes, followed by cyclization of intermediate Schiff base in the presence of various oxidants, such as nitrobenzene [1, 2], 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) [3], benzofuroxan [4], and MnO<sub>2</sub> [2]. The latter procedure requires the presence of an oxidant and correspondingly severe reaction conditions (high temperature, strongly acidic medium [1], or microwave irradiation [5]).



We have found that 2-(2-chlorophenyl)-1H-benzimidazole (I) can be synthesized in a moderate yield in the absence of an oxidant under mild conditions by direct condensation of *o*-phenylenediamine with 2-chlorobenzaldehyde. The product was reliably identified by the <sup>1</sup>H NMR and mass spectra. The molecular structure of I, determined by X-ray analysis, was identical to that reported previously [6].

\* The text was submitted by the authors in English.

**2-(2-Chlorophenyl)-1H-benzimidazole (I).** 2-Chlorobenzaldehyde, 2.25 ml (20 mmol), was added to a solution of 2.16 g (20 mmol) of *o*-phenylenediamine in 50 ml of acetonitrile. The mixture was heated for 1.5 h under reflux and cooled to room temperature, and the precipitate was filtered off and washed with acetonitrile (2 × 1 ml). Yield 1.30 g (28%), mp 228°C [6]. IR spectrum,  $\nu$ , cm<sup>-1</sup>: 3400 (NH), 1624 (C=N). <sup>1</sup>H NMR spectrum,  $\delta$ , ppm: 5.43 s (1H), 7.1–8.0 m (8H). Mass spectrum,  $m/z$ : 228 [ $M$ ]<sup>+</sup>, 193 [ $M - Cl$ ]<sup>+</sup>.

The IR spectrum was recorded in KBr on a Nicolet 20SXR instrument. The <sup>1</sup>H NMR spectrum was obtained on a Varian Unity spectrometer (300 MHz) in CDCl<sub>3</sub>. The mass spectrum (fast atom bombardment) was recorded on a JEOL SX-102 mass spectrometer.

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