

Synthesis of 3-[(Z)-2-piperazin-1-yl-ethylimino]-1,3-dihydro indol-2-one as a novel Schiff base

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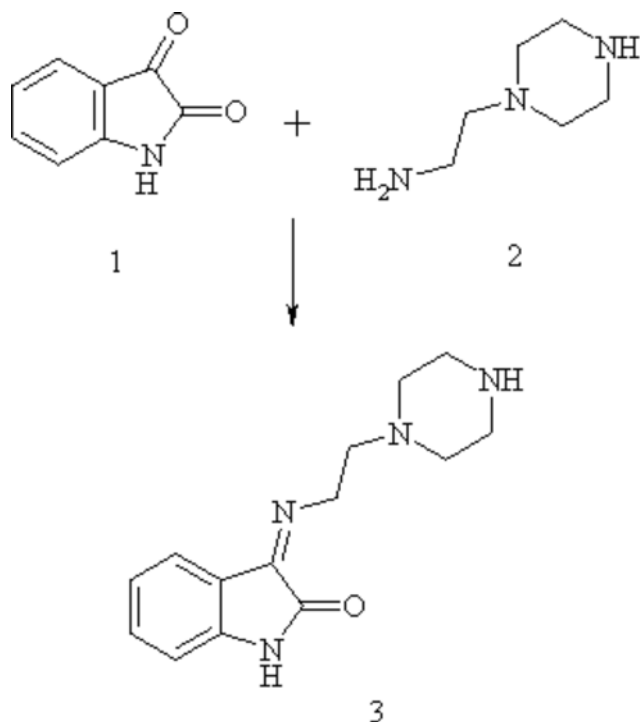
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Isatin (1H-indole-2,3-dione) was first obtained by Erdman and Laurent in 1841 as a product from the oxidation of indigo by nitric and chromic acids [1]. The synthetic versatility of isatin has led to the extensive use of this compound in organic synthesis. In nature, isatin is found in plants of the genus *Isatis* [2], in *Calanthe discolor* LINDL [3]. Isatin is the biologically active chemical produced by an *Altermones* sp. Strain inhabiting the surface of embryos of the cardiean shrimp *Palaemon macrodectylus*, which protect them from the pathogenic fungus *Lagenidium callinectes* [4]. Schiff bases of isatin were reported to possess interesting biological activities such as anti-HIV [5-7], anticonvulsant [8], antibacterial [9-11], antiprotozoal [12, 13], antifungal [14-16], antiviral [17-19], anthelmintic [20,21], antidepressant [22] anti-inflammatory and antitumour [23] activities. The Schiff bases of isatin have also been used as a ligand for complexation with various metal ions [24]. In view of the potent biological activities, we decided to synthesize a new isatin Schiff base by the condensation of isatin with 1-(2-aminoethyl)piperazine (AEP). Its biological activities and chelating behaviour with various metal ions is under study.

Ethanol solutions of isatin (1) (0.01mol, 1.47g) in 50 ml and 1-(2-aminoethyl) piperazine (2) (0.01mol, 1.8ml, 1.79g) in 50 ml were mixed and refluxed for about 2 hours. The reaction mixture was evaporated to a small volume and left to cool. The resulting Schiff base (3) ligand precipitated on cooling and then was filtered off, washed with ethanol and recrystallised from ethanol. The purity of the Schiff base ligand was monitored on TLC using eluants 1:1 ethyl acetate and petroleum ether and separated by column chromatography (Yield =90%).



Melting point: 220 °C

Elemental analysis: Calculated for $C_{14}H_{18}N_4O$ C, 60.86; H, 7.24; N, 20.29. Found: C, 60.28; H, 7.20; N, 19.97.

IR (KBr, cm^{-1}): 3239.9 (N-H), 1714.6 (C=O), 1618.6 (C=N)

MS (ESI, m/z): 259 [M^+]

1H -NMR (400MHz, DMSO- d_6): 10.3(1H, s), 7.6-6.8 (4H, m), 2.8-2.1(13H, m).

^{13}C -NMR (400MHz, DMSO- d_6): 38.3; 45.6; 54.3; 61.8; 110.6; 121.9; 122.6; 123.3; 131.6; 134.3; 141.6; 162.0.

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