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Anhydrous ZnCl₂ Catalyzed Synthesis of 2–Aryl Substituted Benzimidazole Derivatives

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Abstract

Anhydrous $ZnCl_2$ was used to be a catalyst for the synthesis of 2-arylsubstituted benzimidazoles efficiently. In this method is very simple convenient procedure, easy purification and shorter reaction time. So we prepared the many benzimidazole derivatives from this method. All the synthesized compounds were characterized melting point, IR, ¹H NMR, ¹³C NMR spectral data conformed the structure of the newly synthesized compounds.

Keywords: Anhydrous ZnCl,; Chromatography; o-phenylenediamine benzimidazole; Spectroscopy.

1. INTRODUCTION

Benzimidazole is one of the important heterocyclic groups manifesting various biological properties such as antibacterical, antifungal and antihelmintic activities. Compound containing benzimidazole ring which are formed via the fusion of imidazole and benzene ring, have been used extensively for pharmaceutical purposes since 1960 (Buu-Hoi et al. 1963). Benzimidazole derivatives exhibit significant activity against several viruses such as HIV (Jones et al. 1990) herpes (HSV-1) (Grimmet, 1996), RNA (Preston, 1980), influenza (Kanazawa et al. 2006) and human cytomegalovirus (HCMV). The widespread interest in benzimidazoles - containing structure has promoted. In recent times we have shown KHSO, and p-TSOH can be used as promoters and catalysts for the synthesis of benzimidazole (Gracias et al. 2005; Tamm et al. 1978; Siddiqui, et al. 2005). So we tried to synthesis benzimidazole using inorganic catalyst. In this paper

*V. Mohanraj E-mail: chemistrymohan@gmail.com anhydrous $ZnCl_2$ was used for the synthesis of 2-arylsubstituted benzimidazole by the condensation of aryl aldehyde with *o*-phenylenediamine.

2. EXPERIMENTAL SECTION

2.1 General Procedure

All melting points were determined on a kofler micro melting point apparatus and were uncorrected. IR spectra were recorded on a Perkin Elmer spectrophotometer using KBr pellet method. ¹H NMR and ¹³C NMR spectra were measured on a Bruker spectrophotometer using TMS as internal standard. Typical procedure for synthesis of benzimidazole.

3. RESULT & DISCUSSION

Anhydrous zinc chloride hydrolyzes with moisture to form hydrochloric acid. It also forms complex ions with water, ammonia and some organic solvents. Alkaline materials precipitate zinc hydroxide from zinc chloride solutions. Zinc laurate, linoleate or resinate can be formed from zinc chloride solutions and solutions of the corresponding sodium salt. Zinc chloride is a Lewis acid and therefore electrophilic in character. Its catalytic activity is milder than that of aluminium chloride in, for example, Friedel-Crafts type reactions. Zinc chloride is particularly effective in catalyzing reactions that eliminate molecules of water, ammonia or mercaptans. As a part of our ongoing investigation as developing versatile and efficient method for synthesis of heterocyclic compounds. We discussed here synthetic method of benzimidazoles form aldehydes and O – phenylenediamine in presence of anhydrous ZnCl₂.





4. REACTION OF O-PHENYLENEDIAMINE WITH ANISALDEHYDE IN PRESENCE OF ANHYDROUS



4.1 1H NMR - Spectral Data of compound

Proton	Signal (ppm)
H – 2 ['] ,6 [']	7.6(dd,2H)
H – 3 ['] ,5 [']	7.22(m,1H)
H – 4 [°]	6.82(m,1H)
H – 3 [°] ,5 [°]	7.32(m,1H)
H – 4	6.62(m,1H)
H – 5	7(m,1H)
H – 6	7.45(m,2H)
H – 7	7.42 (m,2H)
N – H	5.10(6rs, 1H)
OCH ₃	3.82(m,3H)





Carbon	Signal (ppm)
C – 2	155.5
C – 4	119.924
C – 5	127.174
C – 6	130.97
C – 7	110.71
C – 8	133.82
C – 9	139.28
C – 1	130.97
C – 2 [°]	131.74
C – 3 [°]	123.93
C – 4 [°]	127.17
C – 5'	126.8
C – 6	131.92
OCH3	55.32

¹³C NMR Spectral Data of the Compound

All the carbon signals are in between $\delta 100 - 155$ suggesting only aromatic and unsaturated carbon atom are present. Out of 16 carbon signals four signals $\delta 155.5, \delta 130.97, \delta 139.28$ and $\delta 133.82$ are weak intensity showing the presence four quaternary carbon atoms.



4. IR spectral data of compound

In IR spectral data of compound was exhibits an absorption of 3217.04 N- H starching, 1605.43 for C=C 1302 for C – N group.

5. CONCLUSION

We have developed a simple one pot synthesis of 2-aryl substituted benzimidazoles by the condensation of o-phenylenediamine with aryl aldehydes catalyzed by anhydrous ZnCl₂. The simplicity of the reaction condition pot with shorter reaction time and with column chromatography to get the pure product. Simple and convenient procedure, easy purification shorter reaction time, and very good yield are the advantageous features of these methods and we have finished the spectral studies of IR ¹H NMR, ¹³C NMR of spectral data confirmed the structure of the newly synthesized compounds.

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