Synthesis of some novel 7-(1H-benzimidazol-2-ylazo)-1,3-dimethyl-6,8-disubstituted-1H-pyrimido[4,5-b][1,4]diazepine-2,4-diones as potential anti-anxiety agents

KANTI SHARMA* and RENUKA JAIN

Department of Chemistry, University of Rajasthan, Jaipur 302 004, India

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Abstract: An efficient and mild method for the synthesis of some novel 7-(1H-benzimidazol-2-ylazo)-1,3-dimethyl-6,8-disubstituted-1H-pyrimido[4,5-b][1,4]diazepine-2,4-diones in water has been developed. This method is a good option to obtain the title compounds in quantitative yields in a simple and inexpensive manner. Further, the NH of the title compounds was replaced by various substituents in an ionic liquid, [bmim]PF_6, a recyclable and environmentally benign solvent. The synthesized compounds were characterized by analytical and spectral (IR, ^1H-NMR, ^13C-NMR and FAB mass) data and have been screened for their anti-anxiety activity in mice.

Keywords: 2-aminobenzimidazole, diketones, benzimidazolyl-pyrimido-diazepines, aqueous/ionic liquid mediated synthesis; anti-anxiety activity.

INTRODUCTION

Nitrogen-containing heterocycles represent the central framework of many biologically active compounds. Benzimidazoles show antifungal, anti-inflammatory, analgesic, CNS depressant, antitubercular, anticancer, anti-HIV, etc. activities. Pyrimidines exhibit varied activities, such as anticancer and antibacterial. 5-Fluorouracil is a well-known anticancer agent. Diazepine derivatives are used as tranquilizers, anticonvulsant, anxiolytic, analgesics, sedatives, antidepressives and hypnotic agents. 1,5-Benzodiazepine shows anti-anxiety, analgesic, anticonvulsant and anti-inflammatory activities.

Due to their wide range of pharmacological activity and industrial applications, the development of mild and efficient protocols for the preparation of diazepines continues to be challenging endeavor in synthetic organic chemistry. The common procedure for the synthesis of these compounds is a one-pot condensation between α-phenylenediamines and carbonyl compounds. However, a

*Corresponding author. E-mail: drkanti@gmail.com
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large number of modified methods reported in the literature suffer from several drawbacks, such as the use of a large amount of catalysts, unsatisfactory product yield and critical product isolation procedures. These disadvantages requires the development of an efficient and practically useful process of preparation.

In continuation of work on bioactive heterocycles reported herein is for the first time a rapid efficient, clean and environmentally benign exclusive synthesis of 7-(1H-benzimidazol-2-ylazo)-1,3-dimethyl-6,8-disubstituted-1H-pyrimido[4,5-b][1,4]diazepine-2,4-diones by the reaction of 2-[(1H-benzimidazol-2-yl)hydrazono]-1,3-disubstituted-1,3-dione and 5,6-diamino-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione hydrochloride in aqueous medium in excellent yields (92–97 %). The use of water as a solvent for organic transformations offers green chemistry benefits and is widely used to enable and expedite the synthesis of diverse heterocycles. Compounds were prepared by coupling reaction of a diazonium salt, formed by the diazotization of 2-aminobenzimidazole and 1,3-diketones in presence of sodium acetate and ethanol. Further, the NH of title compounds was substituted by various substituents by reacting with methyl iodide, benzyl chloride, chloroacetyl chloride, formaldehyde and secondary amines, trifluoroacetic anhydride in [bmim]PF₆, an ionic liquid to give the N-substituted derivatives (Scheme 1). The synthesized compounds and were screened for their anti-anxiety activity in mice and they exhibited excellent results.

**EXPERIMENTAL**

Melting points are uncorrected and were taken in open glass capillaries using a Gallenkamp melting point apparatus. The IR spectra were recorded on a Shimadzu FTIR-8400S spectrophotometer in KBr pellets and band positions are recorded in wavenumbers (cm⁻¹). The ¹H- and ¹³C-NMR spectra were recorded on a Bruker DRX-300 instrument at 300.15 and 75 MHz, respectively. CDCl₃ was used as the solvent and TMS as internal reference. The mass spectra were recorded on a JEOL, SX 102 (FAB) mass spectrometer. The mass spectra and elemental analyses were performed at the Central Drug Research Institute, Lucknow, India. All employed chemicals were of analytical reagent grade, purchased from Acros and used without further purification.

**General method for the synthesis of compounds**

2-Aminobenzimidazole (0.02 mol) was dissolved in a mixture of conc. HCl and water (20 mL; 1:1), then cooled to 0 °C and a cold aqueous solution of sodium nitrite (0.02 mol; 1.3 g in 10 mL water) was added slowly maintaining the temperature between 0–2 °C. The cold diazotized solution was added dropwise to a cooled mixture of 1,3-diketone (0.02 mol) and sodium acetate (10 g) in 20 mL of 50 % ethanol. The stirring was continued for 1 h and the crystals that separated were filtered, washed with water, dried and crystallized from ethanol to yield 3a–c.

**Water mediated synthesis of compounds**

A solution of 5,6-diamino-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione hydrochloride (0.01 mol) in water (200 mL) was neutralized with ammonia to pH 7 then 3a–c (0.01 mol),
was added and the mixture heated under reflux for 30 min. The crystalline product, which started to separate out just after cooling the reaction mixture, was washed with water and found to be pure by TLC with no need for further purification. All compounds were synthesized similarly to yield 4a–c.
### Scheme 1. Synthesis of compounds 3a–c, 4a–c and 5a–i.

**Ionic liquid mediated synthesis of compounds 5a–i**

A mixture of 4a–c (0.01 mol), methyl iodide/benzoyl chloride/chloroacetyl chloride/formaldehyde and secondary amine (1:1) or trifluoroacetic anhydride (0.01 mol) and ionic liquid [bmim]PF$_6$ (5 mL) were taken in a round bottom flask and heated at 60–70 °C under N$_2$ protection for 1 h. The progress of the reaction was monitored by TLC. After completion of reaction, the mixture was extracted with diethyl ether (6×10 mL). The organic extract was washed with 5% Na$_2$CO$_3$ (40 mL) and water (40 mL), dried with anhydrous sodium sulfate and evaporated under vacuum. The residual product was purified by recrystallization from AcOEt/petroleum ether (60–80 °C) or by column chromatography (silica gel, 60–120 mesh, eluent petroleum ether/AcOEt = 4:1) to give 5a–i (the ionic liquid layer was washed with water (3×5 mL) and kept for 2 h. at 80–85 °C under reduced pressure, the ionic liquid was recycled).

**Anti-anxiety activity in mice**

Synthesized compounds 4a–c and 5a–i were screened for their anti-anxiety activity in mice on a plus maze apparatus devised by Crawley and Godwin$^{20}$ modified by Kilfoil$^{21}$ using sodium pentobarbitone as the standard.

The apparatus consisted of a Plexiglas base (102 cm×53 cm×53 cm) divided into two chambers by a black Plexiglas partition. The box was placed within a layer of a soundproof box, which was equipped with a one-way observation window. The partition dividing the sub chamber had a 33 cm×13 cm opening, through which animal could easily pass. The dark chamber (36 cm×53 cm×53 cm) was made up of dark Plexiglas except for the side ferry observation window. This side was clear and covered with black plastic. The testing was performed between 12.00 noon to 6.00 p.m. in an isolated darkened laboratory.

Mice weighing 25 g were chosen and were sorted into five animals in a group. They were allowed free access to food and water *ad libitum*. Animals were given 60 min to acclimatize to the environment prior to the administration of the drug. Drugs or test samples in DMF were given at a dose of 20 mg kg$^{-1}$ body weight, after which each animal was individually placed in the centre of the light area of the apparatus and observed for 10 min. The total amount of time spent in the dark area was measured.

**RESULTS AND DISCUSSION**

2-Aminobenzimidazole (1) was reacted with sodium nitrite and HCl at 0–5 °C. The so-formed diazonium salt was further reacted with 1,3-diketones (2) and sodium acetate in ethanol to afford 2-[(1H-benzimidazole-2-yl)hydrazono]-1,3-disubstituted 1,3-diones 3a–c. The formation of compound 3 was confirmed by
their IR spectra, in which band appeared at 3200, 3030, 1680 and 1620 cm\(^{-1}\) due to –NH benzo, NH hydrazono, C=O and C=N, respectively. The \(^1\)H-NMR spectra showed peaks at \(\delta\) 10.62 and 9.52 ppm due to the NH hydrazono and NH of benzimidazolyl moieties, respectively. The \(^{13}\)C-NMR spectra showed peaks at \(\delta\) 178.2, 180.1 and 162.4 ppm due to two C=O and the C=N, respectively. Further, the mass spectrum shows M\(^+\) at \(m/z\) 244 (3a).

The advantage of using water as a solvent is its cost, safety (it is non-inflammable and is devoid of any carcinogenic effects) and simple operation. It has the highest specific heat value of all substances as well as unique enthalpic and entropic properties, which have led us to its use. Water has abnormally low volatility because its molecules are associated with each other by means of hydrogen bonding.

In view of this, compounds 3a–c were reacted with 5,6-diamino-1,3-dimethyl-2,4(1H,3H)-pyrimidinedione hydrochloride in aqueous medium to afford 7-(1H-benzimidazol-2-ylazo)-1,3-dimethyl-6,8-disubstituted-1H-pyrimido[4,5-b]-[1,4]diazepine-2,4-diones 4a–c.

The formation of compounds 4 was confirmed by their IR spectra, which showed the disappearance of the peaks due to the hydrazono group and the ketonic group of the 1,3-diketone part with the appearance of a peak due to the NH of diazepine at 3340 cm\(^{-1}\) and at 1720 and 1700 cm\(^{-1}\) due to the C=O of the pyrimidine ring. The \(^1\)H-NMR spectra showed peaks at \(\delta\) 9.51 and 8.46 ppm due to the NH of benzimidazole and diazepine moiety, respectively. The NCH\(_3\) methyl groups present at the positions 4 and 6 appeared at \(\delta\) 3.20 and 3.62 ppm, respectively. The \(^{13}\)C-NMR spectra showed peaks at \(\delta\) 33.5 and 34.8 ppm due to the two NCH\(_3\) groups of the pyrimidine ring, while two C=O groups appeared at \(\delta\) 190.2 and 195.1 ppm. Further, the mass spectrum shows M\(^+\) at \(m/z\) 378 (4a).

In recent years ionic liquids\(^{22}\) (ILs) have emerged, which are organic salts the ions of which do not pack well and remain liquid at room temperature. They have unique properties, such as, for example, a wide liquid range, good solvency, tunable polarity, high thermal stability, negligible vapor pressure and recyclability. Ionic liquids could be recycled and reused as opposed to traditional solvent catalyst systems.

In view of the above, compounds 4a–c were reacted with: i) methyl iodide, ii) benzyl chloride, iii) chloroacetyl chloride, iv) formaldehyde and a secondary amine and v) trifluoroacetic anhydride in the ionic liquid \([\text{bmim}]\text{PF}_6\) to give N-benzimidazolyl substituted derivatives 5a–i.

Although compounds 4a–c have two NH groups, one in the benzimidazolyl moiety and another in the diazepine ring, it was observed that only NH of benzimidazolyl moiety was replaced by various substituents. The NH of the diazepine moiety could not be replaced due to hydrogen bonding with the C=O of the pyrimidine ring.
The formation of compounds 5a–i were confirmed by the disappearance of peak due to the NH of the benzimidazolyl moiety and the appearance of a peak due to N-substituent in their IR, 1H-NMR and 13C-NMR spectra. The mass spectrum also shows M\(^+\) at m/z 392 (5a).

**Anti-anxiety activity in mice**

Synthesized compounds 4a–c and 5a–i were screened for their anti-anxiety activity in mice on a plus maze apparatus devised by Crawley and Godwin\(^\text{20}\) modified by Kilfoil.\(^\text{21}\) The less time a mouse spent in the dark space, the greater was the anti-anxiety activity of the drug. The results of the anti-anxiety activity of the compounds are given in Table I. Compounds 4c, 5b, 5e, 5h and 5i showed better activity than the standard, sodium pentobarbitone, and the other compounds were moderately active. The good activity of these compounds is due to presence of trifluoromethyl, chloro and trifluoroacetyl groups, which enhance their activity.

**TABLE I. Results of anti-anxiety activity in mice of compounds 4a–c and 5a–i**

<table>
<thead>
<tr>
<th>Compound</th>
<th>R(^1)</th>
<th>R(^2)</th>
<th>R(^3)</th>
<th>Time spent in dark space, s</th>
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<tr>
<td>4a</td>
<td>CH(_3)</td>
<td>CH(_3)</td>
<td>–</td>
<td>300±9.2</td>
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<tr>
<td>4b</td>
<td>CH(_3)</td>
<td>C(_6)H(_5)</td>
<td>–</td>
<td>298±8.4</td>
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<tr>
<td>4c</td>
<td>CF(_3)</td>
<td>C(_6)H(_5)</td>
<td>–</td>
<td>291±8.2</td>
</tr>
<tr>
<td>5a</td>
<td>CH(_3)</td>
<td>CH(_3)</td>
<td>CH(_3)</td>
<td>304±10.5</td>
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<tr>
<td>5b</td>
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<td>CH(_3)</td>
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<tr>
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<tr>
<td>5d</td>
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<tr>
<td>5e</td>
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<td>Pentobarbitone</td>
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<td>295±11.50</td>
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</table>

**CONCLUSIONS**

Novel 7-(1\(H\)-benzimidazol-2-ylazo)-1,3-dimethyl-6,8-disubstituted-1\(H\)-pyrimido[4,5-b][1,4]diazepine-2,4-diones 4a–c were synthesized in aqueous medium by the reaction of 2-[(1\(H\)-benzimidazol-2-yl)hydrazono]-1,3-disubstituted-1,3-diones 3a–c and 5,6-diamino-1,3-dimethyluracil hydrochloride. Further, the title compounds were reacted with various reagents/reactants in [bmim]PF\(_6\) to determine whether all NH groups of the title compounds could be substituted. It was observed that only monosubstitution was possible yielding 5a–i as the diazepine NH is involved in hydrogen bonding with the C=O of the pyrimidine ring. Compounds 4a–c and 5a–i were screened for their anti-anxiety activity in
mice and the data revealed that the compounds showed good to moderate activity. Some of the compounds were more active than the standard and could act as potential anti-anxiety agents.

SUPPLEMENTARY MATERIAL

Analytical and spectral data for the compounds 3a–c, 4a–c and 5a–i are available electronically from http://www.shd.org.rs/JSCS/, or from the corresponding author on request.

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REFERENCES