Montmorillonite K10 Catalysed Condensation of 1-(3-Chlorophenyl)-4-(3-chloropropyl)piperazin and 1,2,4-Triazolo [4,3-a]pyridine-3-(2H)-one: A Proficient and Green Synthesis of Trazodone Hydrochloride and Its Analogues

Abstract
Montmorillonite K10 clay has an extremely significant heterogeneous catalyst for condensation reaction with high yield of the product as compared to other reported reagents.

Background: Novel green, synthetic methodology has been adopted for the synthesis of 2-[3-{4-(3-chlorophenyl)-1-piperazinyl}propyl]-1,2,4-triazolo[4,3-a]pyridine-3-(2H)-one hydrochloride (5) analogues using Montmorillonite K10 (MK10) as an effective-efficient heterogeneous catalyst. The present synthesis offers various significant features with numerous advantages such as outstanding yields, operational simplicity, recyclability and reusability of the catalyst for this synthetic reaction.

Methods and findings: Condensation reaction is the only exceptional tool for the synthesis of numerous biologically significant organic compounds. In the current investigation, condensation of 1-(3-chlorophenyl)-4-(3-chloropropyl)piperazine (3) and 1,2,4-triazolo[4,3-a]pyridine-3-(2H)-one (4) in the presence of MK10 under thermal condition has been carried out with afford the corresponding products i.e., (5) analogues with good yields (88-95%) over conventional reagents. MK10 is such a heterogeneous catalyst which can be completely recovered and reusable for a three times without potential loss in its catalytic activity.

Keywords: Condensation reaction; Montmorillonite K10; Recyclability; Heterogeneous catalyst

Introduction
Multi-component condensation reactions [1] that are performed in heterogeneous catalyst have gained improved interest in synthetic chemistry over the past decade not only for the advantages rendered by avoiding extensive decomposition of reactants, strong reaction conditions and solvents, but also for the development of environmentally benign methodology [2].

Furthermore, when a heterogeneous catalyst [3] is used, the insoluble catalyst can be separated by simple filtration and the catalyst can be reused. Therefore, the improvement of a heterogeneous catalyst in solvent appears enormously desirable.

Commercially available Montmorillonite K10 (MK10) [4] is one such catalyst that can fulfill these requirements. MK10 are environmentally friendly and economically reasonable solid acid catalysts that offers several advantages, such as ease of handling, non-corrosiveness, low cost and regeneration. Green chemistry, believes in replacement of toxic organic solvents [5], which are responsible for excessive waste of chemicals and environmental problems. In response catalysts such as MK10 are presently under dynamic research. These not only avoid the use of toxic acids, high temperature carrying reactions but also prompt important simplifications to the reaction procedures. The main contributing factors are the high atom economy, extensive application in combinatorial chemistry and diversity oriented synthesis [6-10].
The synthetic product (5), known as “Trazodone hydrochloride” [11] is a well-known drug with psychoactive properties synthesized from piperazine and triazolopyridine classes of compounds having antidepressant, anxiolytic and hypnotic properties [12]. The pharmacological potential of Trazodone hydrochloride is due to inhibition of serotonin uptake and also a lower affinity for the serotonin transporter than other drugs which belongs to selective serotonin reuptake inhibitor class [13].

Recently, the Synthesis of Novel Benzofluorenone Derivatives and their HIV-Reverse Transcriptase Inhibitory Activity [14], one-pot synthesis of new N,N’-alkylidene bisamide derivatives under solvent free condition by montmorillonite K10 [15] and the synthesis of Trazodone hydrochloride by sulphamic acid are reported [16].

Thus, in the present study the green synthetic procedure for the synthesis of (5) analogues in the presence of MK10 as an efficient catalyst has been established (Scheme 1). The literature survey reveals that there is reported no prior synthesis of these new compounds using green synthetic methodology.

**Methods**

**Materials and reagents**

Required chemicals and reagents were procured from Aldrich (Sigma–Aldrich, St. Louis, MO, USA), Lancaster (Alfa Aesar, Johnson Matthey Company, Ward Hill, MA, USA) and used without further purification. Reaction monitoring was done with Thin Layer Chromatography (TLC), performed on silica glass plates 60 GF-254, and visualization was achieved by UV light. Column chromatography was performed by Merck 60–120 mesh silica gel. ¹H spectra were recorded by Bruker UXNMR/XWIN-NMR (300 MHz) instruments. Chemical shifts (δ) were reported in ppm downfield from internal TMS standard. ESI spectra were recorded in Micro mass; Quattro LC uses ESI+ software with capillary voltage 3.98 kV and ESI mode positive ion trap detector. Melting points were determined with an Electro thermal melting point apparatus, and were approximate.

**Novel procedure for synthesis of Intermediate analogues**

Intermediate analogues were synthesized based on practically a similar procedure as reported by Lambat and Deo [16]:

**Step 1:** General procedure for the synthesis of bis-(2-chloroethylamine) hydrochloride (1) [16].

**Step 2:** General procedure for the synthesis of 1-(3-chlorophenyl)piperazine hydrochloride and its derivatives (2) [16].

**Step 3:** General procedure for the synthesis of 1-(3-chlorophenyl)-4-(3-chloropropyl) piperazine and its derivatives (3) [16].

**Step 4:** General procedure for the synthesis of sodium salt of 1, 2, 4-triazolo [4, 3-a] pyridine-3-(2H)-one (4) [16].

**Step 5:** Novel procedure for the synthesis of Trazodone hydrochloride derivative.

In a typical reaction procedure, product (3) (3.6 mmol), (4) (1.15 mol) and Montmorillonite K10 (0.500 g) in acetonitrile (3.00 mL). The resulting mixture was refluxed at 90°C for 5 h. After completion of the reaction as monitored by TLC, the reaction mixture was cooled to room temperature and filtered. The acetonitrile was removed by distillation and toluene (3 mL) was added to the reaction mixture and stirred for 10 min and filtered. Then, the catalyst was isolated by simple filtration, which was recovered later and the remaining supernatant was treated with 20% NaOH followed by 2% brine solution at 50°C. The toluene solution containing desired product which is base, in which HCl (15%) was added and pHS-adjusted between 2-2.5 when salt starts precipitating. The precipitated product was filtered; recrystallization of the crude product from methanol to furnished the pure product. All the synthesized products (SA-5G) are known compounds, which were characterized based on their melting points and spectral (FT-IR, ¹H and ¹³C NMR) data and compared with the reported corresponding data.

**Spectral data of synthesized product**

**Lead compound:** IR (KBr) (νmax): 3000, 2954, 1704, 1650, 1600, 1350.80, 750; ¹H NMR(300 MHz, DMSO-d6) δ: 2.16-2.12 ppm (t, 2H, N-CH₂-N), 2.64-2.60 (t, 2H, CH₂-N), 2.73 (s, 4H, CH₂-N), 3.09 (s, 4H, CH₂-N), 4.12-4.07 (t, 2H, CH₂-N), 6.51-6.46 (m, 1H, -ArH), 7.02-6.93 (m, 2H, -ArH), 7.09-7.08 (d, 2H, -ArH), 7.26-7.17 (m, 1H,-ArH), 7.34-7.31 (d, 1H,-ArH); m/z (ESI): 372 (M⁺).

**SA:** IR (KBr) (νmax): 3000, 2850, 1700, 1650, 1600, 1350, 750; ¹H NMR(300 MHz, DMSO-d6) δ: 2.50-2.49 (m, 2H, -CH₂-); 3.21-3.15 (m, 4H, -CH₂-); 4.12-4.07 (t, 2H, -CH₂-); 6.51-6.46 (m, 1H, -ArH); 7.02-6.93 (m, 2H, -ArH); 7.09-7.08 (d, 2H, -ArH); 7.26-7.17 (m, 1H,-ArH); 7.34-7.31 (d, 1H,-ArH); m/z (ESI): 372 (M⁺).
(m, 8H, CH₂-piperazine), 3.60 (t, 2H, -CH₂), 4.01 (t, 2H, -CH₂), 6.65-6.63 (m, 1H, -ArH), 7.25-7.21 (m, 3H, -ArH), 7.36-7.34 (t, 2H, -ArH), 7.88-7.86 (d, 1H, -ArH); m/z (ESI): 406 (M⁺).

5B: IR (KBr) (νmax): 3050, 2980, 2947, 1704, 1643, 1635, 1530, 750; ¹H NMR(300 MHz, DMSO-d6) δ: 2.26-2.19 (m, 2H, CH₂), 3.23-3.01 (m, 6H, CH₂-piperazine), 3.40-3.50 (d, 2H, CH₂-piperazine), 3.90-3.86 (d, 2H, CH₂), 4.01-3.97 (t, 2H, CH₂), 6.67-6.60 (m, 1H, ArH), 7.01-6.97 (m, 1H, -ArH), 7.21-7.20 (m, 3H, -ArH), 7.40 (d, 1H, -ArH), 7.88-7.859 (d, 1H, -ArH); m/z (ESI): 463 (M⁺).
little or no effect of substituents as observed in compound 5F (Table 2) as far as the yield of product is concerned. It is because two electron donating methoxy groups were present on R₁ and R₄ of product (3) which makes the aromatic ring more electron rich. During the condensation reaction the chloro group present on the side chain of (3) was removed via SN₂ displacement process and hence inversion products were expected in the reaction. The possible role of MK10 is depicted in the proposed mechanism (Scheme 2).

The recyclability potential of the MK10 catalyst was examined for the model reaction product of lead compound. The recycling procedure involved the separation of the catalyst from the reaction mixture simply by usual filtration. The recovered catalyst was purified by washing with ethyl acetate followed by drying
Table 2 Synthesis of 2-[3-{4-(3-chlorophenyl)-1-piperazinyl} propyl]-1,2,4-triazolo[4,3-a]pyridine-3-(2H)-one hydrochloride (5) analogues. *Yields refer to those of pure isolated products characterized by IR, 1H NMR and Mass spectra; **Lambat and Deo [16].

| Entry | Product | R_1 | R_2 | R_3 | R_4 | R_5 | n | Yield (%)* | Melting Point (°C) |
|-------|---------|-----|-----|-----|-----|-----|---|------------|-------------------|
| 1     | Lead (5) | -H  | -Cl | -H  | -H  | -H  | 3 | 94         | 222 [223]         |
| 2     | 5A      | -Cl | -Cl | -H  | -H  | -H  | 3 | 92         | 253 [>250]        |
| 3     | 5B      | -H  | -Cl | -Cl | -H  | -H  | 3 | 91         | 251 [>250]        |
| 4     | 5C      | -H  | -Br | -H  | -H  | -H  | 3 | 92         | 211 [211-213]     |
| 5     | 5D      | -H  | -H  | -F  | -H  | -H  | 3 | 90         | 243 [240-242]     |
| 6     | 5E      | -Cl | -H  | -Cl | -Cl | -H  | 3 | 89         | 252 [>250]        |
| 7     | 5F      | 0   | -H  | -H  | 0   | -H  | 3 | 95         | 232 [230-232]     |
| 8     | 5G      | -C_2H_5 | -H | -H  | -H  | -H  | 3 | 93         | 207 [204-206]     |
| 9     | 5H      | -H  | -Cl | -H  | -H  | -H  | 2 | 91         | 219 [218-220]     |
| 10    | 5I      | -H  | -Cl | -H  | -H  | -H  | 4 | 90         | 227 [227-230]     |
| 11    | 5J      | Naphthyl | -H | -H  | -H  | -H  | 3 | 92         | 226 [225-227]     |

Table 3 Recycling of MK 10 for the synthesis of model product (lead compound). *Isolated yields.

| Sr. no. | Cycle | Yield (%)* |
|---------|-------|------------|
| 1       | Fresh | 94         |
| 2       | 1st   | 91         |
| 3       | 2nd   | 90         |
| 4       | 3rd   | 87         |

in an oven. The results summarized in (Table 3) reveals that the catalyst has to be used for three successive times without loss of its activity. The reliability of the recovered catalyst was detected and proved to be as active as the new catalyst (Figure 5).

**Conclusion**

We have synthesized a novel extremely significant pharmacologically active scaffold by using MK10 catalyst, which
proficiently activates the two component condensation for the production of (S) derivatives. The main advantages of the present synthetic methodologies are efficiency, versatility, good yield, short reaction times, cleaner reaction profile, convenient work-up, easy catalytic recyclability, and reusability with no loss of catalytic activity, which makes this protocol valuable and attractive in the improving of benign chemical processes and products.

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