[4,4'-Bipyridine]-1,1'-dium tricyanomethanide as a nanostructured molten salt and its catalytic application in the synthesis of tetrahydrobenzo[*b*]pyrans, amido and aminoalkyl naphthol derivatives

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Abstract: An efficient process for the synthesis of tetrahydrobenzo[*b*]pyrans derivatives was attained via the reaction of dimedone, various aldehydes and malononitrile using [4,4'-Bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a green mild nano structure molten salt catalyst under solvent-free conditions at room temperature. To further investigation on catalytic properties of {[4,4'-BPyH][C(CN)₃]₂}, it was used in the synthesis of amido and aminoalkyl naphthol derivatives under solvent-free conditions at 75 °C. {[4,4'-BPyH][C(CN)₃]₂} has advantages such as cleaner reaction profile, benign characters, cost effectiveness, reusability of the catalysts and being in agreement with the green chemistry protocols.

Keywords: Bifunctional nano structure molten salt; [4,4'-Bipyridine]-1,1'-dium tricyanomethanide; Tetrahydrobenzo[*b*]pyrans; Amidoalkyl naphthol; Aminoalkyl naphthol; Solvent-free conditions.

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1. Introduction

It is generally acknowledged that there is an increasing significance for more accessible eco-friendly approaches in the chemical and heterocyclic synthesis and pharmaceutical industries. This advanced improvement called 'green chemistry' or 'ecological technology' needs a model alteration taken from customary ideas of procedure efficiency, that focuses chiefly on yield of product, to one that assigns economic value, removing waste and avoiding the usage of dangerous and toxic materials [1]. Room-temperature molten salts (RTMSs), named as room-temperature ionic liquids, (RTILs), which are immiscible or only partially miscible with water, are promising to be replacements of usual organic solvents. Molten salts (MSs) have significant potentials such as electrolyte substances. The molten salts, recently termed ionic liquids, have drawn much consideration in numerous scientific fields [2]. Molten salts or ionic liquids are usually liquid electrolytes combined of ions. More recently, the melting point pattern has been introduced to make identification of molten salts (very corrosive medium, high-melting and highly viscous) and ionic liquids (relatively low viscosity and liquid below 100 °C) [3] possible. Molten salts have noteworthy potentials such as electrolyte substances. Scientists have performed experiments to change solid salts into liquids using appropriate substances and also efforts have been made to polymerize molten salts into solid films. Numerous investigations have been done to improve a superb ion-conductive matrix, among which polymerization of molten salts is a promising process to create highly ion-conductive polymer films [4, 5]. Besides the polarization procedure applied, molecular dynamics simulations of molten salts in which polarization effects are included have exposed that the mobility of liquids increases [6, 7]. Furthermore, other uses of molten salt are in methane reforming and methanol steam reforming [8, 9].

Multicomponent reactions (MCRs) demonstrate a very efficient and economic approach to build up complex structures in a single synthetic process from very simple reactants, most of the time in a one-pot reaction [10]. Such reactions display methodological ease, high selectivity and high atom economy because of the formation of carbon-carbon and carbon-heteroatom bonds in a single step [11]. MCRs are usually very high yielding being a one-pot process and are basically fully different from two-component reactions [12]. One such MCRs is the one-pot three-component tandem Knoevenagel-Michael cyclocondensation synthesis of tetrahydrobenzo[*b*]pyran derivatives that is one of the significant types of organic compounds. Tetrahydrobenzo[*b*]pyran derivatives are significant compounds that have full-fledged remarkable consideration in past years owing to their varied range of biological activities [13] for example diuretic, anticancer, anti-anaphylactic activities and anticoagulant [14]. Correspondingly, tetrahydrobenzo[*b*]pyrans are applied as cognitive enhancers for the treatment of neurodegenerative

illness, for instance Alzheimer's, Huntington's, Parkinson's illnesses and Down's syndrome in addition to the treatment of myoclonus and schizophrenia [15]. Owing to the significance of tetrahydrobenzo[*b*]pyrans, various methodologies have been reported for its synthesis in past years [16].

Other significant types of organic compounds are the 1-(amino or amidoalkyl)-2-naphthol derivatives, called Betti bases [17]. Recently, attention to the chemistry of the Betti base [18] has increased because of their attractive catalytic [19] and biological [20] properties. These compounds can be transformed into derivatives having hypotensive, antibacterial and bradycardiac activities [21]. There are various processes described for the synthesis of 1-(amino or amidoalkyl)-2-naphthols containing saccharin sulfonic acid (SASA) [22], nano-S (S8-NP) [23], [Msim]Cl or [Dsim]Cl or [Msim]AlCl₄ [24], {[HMIM]C(CN)₃} [25], {[1,4-DHPyrazine][C(NO₂)₃]₂} [26], un-catalyzed in water [27] and sulfanilic acid-functionalized silica-coated magnetite nanoparticles [28].

Following our previous studies linked to the design, synthesis, applications and knowledge-based development of novel solid acids and inorganic acidic salts [29], novel nano structured, green and benign ionic liquids, molten salts and organo-catalysts for organic functional group transformations and as a section of our continuous work to the development of environmentally benign synthetic procedures for MCRs to synthesize various biologically significant heterocyclic compounds [30], herein we wish to report the synthesis of an efficient and reusable bifunctional nano structure molten salt catalyst, namely [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} (Scheme 1) [31] and its application for the highly effective synthesis of tetrahydrobenzo[*b*]pyrans and amido or aminoalkyl naphthol derivatives under solvent-free conditions (Scheme 2).

>Scheme 1<

>Scheme 2<

2. Experimental

The materials were purchased from Merck, Fluka and Sigma-Aldrich and were used without any additional purification. All reactions were monitored by thin layer chromatography (TLC) on gel F254 plates. Spectrometer (¹H NMR 400 MHz and ¹³C NMR 100) in pure deuterated DMSO with tetramethylsilane (TMS) as the internal standard. Characteristics of the synthesized catalyst were determined by IR, ¹H NMR, ¹³C NMR, mass, X-ray diffraction patterns (XRD), scanning electron microscopy (SEM) and transmission electron microscopy (TEM), thermo gravimetric (TG) and derivative thermal gravimetric (DTG) analysis. X-ray diffraction (XRD) patterns of all catalysts were performed on a APD 2000, Ital structure with Cu K_α radiation (λ = 0.1542 nm) operating at 50 kV and 20 mA in a 2 h

range of 10–70° with step size 0.01° and time step 1.0 s to assess the crystallinity of the catalyst. Fourier transform-infrared spectra of the samples were recorded on a Perkin-Elmer FT-IR spectrometer 17259 using KBr disks. Thermo gravimetric analyses using a Perkin-Elmer TGA were performed on catalysts. The weight loss about 152 °C was identified. The SEM analyses were done with a TESCAN/MIRA with a maximum acceleration voltage of the primary electrons between 10 and 15 kV. Transmission electron microscope, TEM measurements were carried out on a Philips CM10 analyzer. Operating at 120 kV.

General procedure for the synthesis of bifunctional nano structure molten salt catalyst: [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂}

The NMS {[4,4'-BPyH][C(CN)₃]₂} as a green catalyst was synthesized according to our previously reported procedure [31]. A pale yellow solid was prepared in high purity and then the physical data of this identified NMS was found to be identical with the reported procedure (Scheme 1).

[4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂}

Pale yellow solid; M.p: >300 °C; Yield: 97% (1.641 g); Spectral data: IR (KBr): $\nu_{\max}$ = 3411, 3070, 3025, 2927, 2169, 2082, 1662, 1486, 1218 cm⁻¹; ¹H NMR (400 MHz, DMSO-*d*₆): δ_{ppm} = 6.92 (d, 4H, *J* = 0.8 Hz, ArH), 7.12 (d, 4H, *J* = 0.8 Hz, ArH), 7.63 (s, 2H, —NH); ¹³C NMR (100 MHz, DMSO-*d*₆): δ_{ppm} 50.1, 109.8, 121.7, 151.0, 167.3; MS: *m/z* = 338 [M]⁺.

General procedure for the tandem Knoevenagel-Michael cyclocondensation reaction synthesis of tetrahydrobenzo[b]pyran derivatives

{[4,4'-BPyH][C(CN)₃]₂} (1 mol%; 0.0034 g) as a green mild catalyst was added to a mixture of aromatic aldehydes (1 mmol), malononitrile (0.066 g, 1 mmol) and dimesone (0.140 g, 1 mmol) in a round bottom flask, and the consequent mixture was first stirred magnetically under solvent-free condition at room temperature. After completion of the reaction, as known by TLC *n*-hexane/ethyl acetate (5:2), ethyl acetate (10 mL) was added to the reaction mixture, stirred and refluxed for 5 min, and then was washed with water (10 mL) and decanted to separate catalyst from the reaction mixture (the reaction mixture was soluble in hot ethyl acetate and NMS catalyst was soluble in water). The solvent of organic layer was evaporated and the crude product was purified *via* recrystallization from ethanol/water (10:1). In this investigation, bifunctional NMS catalyst was recycled and reused for four experimental errors with negligible losing of its catalytic activity.

General procedure for the synthesis of amido or aminoalkyl naphthol derivatives

{[4,4'-BPyH][C(CN)₃]₂} (1 mol%; 0.0034 g) as a green mild catalyst was added to a mixture of aromatic aldehydes (1 mmol), β -naphthol (0.144 g, 1 mmol) and amide or amine derivatives (1 mmol) in a round bottom flask, and the subsequent mixture was first stirred magnetically under solvent-free

condition at 75 °C. After completion of the reaction, as identified by TLC *n*-hexane/ethyl acetate (5:3), ethyl acetate (10 mL) was added to a reaction mixture, stirred and refluxed for 5 min, and then was washed with water (10 mL) and decanted to separate catalyst from the reaction mixture (the reaction mixture was soluble in hot ethyl acetate and NMS catalyst was soluble in water). The solvent of organic layer was evaporated and the crude product was purified by recrystallization from ethanol/water (10:1). In this study, bifunctional NMS catalyst were recycled and reused for five experimental errors with a minor loss of its catalytic activity.

Spectral data analysis for compounds

2-Amino-4-(4-fluorophenyl)-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile
(Table 3, entry 8)

White solid; M.p: 188-190 °C; Yield: 96%; IR (KBr): $\nu_{\max}$ = 3358, 3181, 2900, 2190, 1673, 1637, 1507, 1366, 1216 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6): δ_{ppm} = 0.95 (s, 3H—CH₃), 1.04 (s, 3H—CH₃), 2.13 (d, 2H, J = 8.0 Hz, —CH₂), 2.27 (d, 2H, J = 8.0 Hz, —CH₂), 4.20 (s, 1H,—CH aliphatic), 7.11 (s, 2H, —NH₂), 7.16 (d, 2H, J = 6.0 Hz, ArH), 7.20 (d, 2H, J = 6.0 Hz, ArH); ^{13}C NMR (100 MHz, DMSO- d_6): δ_{ppm} = 26.8, 28.3, 31.8, 34.8, 49.9, 58.0, 112.6, 114.9, 115.1, 119.6, 129.0, 129.1, 158.3, 158.4, 159.7, 162.1, 162.5, 195.7.

2-Amino-4-(4-bromophenyl)-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile
(Table 3, entry 14)

White solid; M.p: 205-207 °C; Yield: 92%; IR (KBr): $\nu_{\max}$ = 3319, 3183, 2962, 2169, 1682, 1655, 1367, 1215 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6): δ_{ppm} = 0.95 (s, 3H—CH₃), 1.04 (s, 3H—CH₃), 2.13 (d, 2H, J = 8.0 Hz, —CH₂), 2.27 (d, 2H, J = 8.0 Hz, —CH₂), 3.36 (s, 2H, —NH₂), 4.18 (s, 1H,—CH aliphatic), 7.13 (d, 2H, J = 8.4 Hz, ArH), 7.50 (d, 2H, J = 8.4 Hz, ArH); ^{13}C NMR (100 MHz, DMSO- d_6): δ_{ppm} = 18.5, 26.8, 28.3, 31.8, 35.1, 49.9, 55.9, 57.6, 112.2, 119.5, 119.6, 129.5, 131.2, 144.1, 158.3, 158.4, 162.6, 195.7.

4,4'-(1,4-Phenylene)bis(2-amino-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile) (Table 3, entry 15)

White solid; M.p: 269-271 °C; Yield: 85%; IR (KBr): $\nu_{\max}$ = 3452, 3331, 3187, 2962, 2198, 1683, 1665, 1364, 1212 cm^{-1} ; ^1H NMR (400 MHz, DMSO- d_6): δ_{ppm} = 0.99 (s, 6H—CH₃), 1.04 (s, 6H—CH₃), 2.19 (d, 4H, J = 8.0 Hz, —CH₂), 2.26 (d, 4H, J = 8.0 Hz, —CH₂), 2.51 (s, 4H, —NH₂), 4.14 (s, 2H,—CH aliphatic), 7.04 (s, 4H, ArH); ^{13}C NMR (100 MHz, DMSO- d_6): δ_{ppm} = 27.2, 28.2, 31.8, 34.9, 49.9, 58.3, 112.7, 112.8, 119.7, 126.9, 142.8, 158.4, 158.5, 162.7, 195.8.

2-Amino-4-(3-chlorophenyl)-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile
(Table 3, entry 17)

White solid; M.p: 230-232 °C; Yield: 94%; IR (KBr): $\nu_{\max}$ = 3321, 3184, 2965, 2191, 1682, 1659, 1372, 1213 cm^{-1} ; ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ_{ppm} = 0.96 (s, 3H—CH₃), 1.04 (s, 3H,—CH₃), 2.15 (d, 2H, J = 8.0 Hz, —CH₂), 2.28 (d, 2H, J = 8.4 Hz, —CH₂), 2.53 (s, 2H, —NH₂), 4.22 (s, 1H,—CH aliphatic), 7.13 (d, 1H, J = 7.6 Hz, ArH), 7.17 (s, 1H, ArH), 7.27 (d, 1H, J = 8.0 Hz, ArH), 7.36 (t, 1H, J = 7.6 Hz, ArH); ^{13}C NMR (100 MHz, $\text{DMSO}-d_6$): δ_{ppm} = 26.8, 28.2, 31.8, 35.3, 49.9, 57.6, 112.0, 119.4, 125.9, 126.6, 127.0, 130.3, 132.9, 147.2, 158.4, 158.5, 162.8, 195.7.

3. Results and discussion

*Application of [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional nano structure molten salt catalyst in the synthesis of tandem Knoevenagel-Michael cyclocondensation synthesis of tetrahydrobenzo[*b*]pyran derivatives.*

Firstly, to optimize the reaction conditions, the condensation reaction of 4-chlorobenzaldehyde, malononitrile and dimedone was chosen as a typical model and various amounts of bifunctional nano structure molten salt catalyst at range of 25–100 °C were tested under solvent-free conditions (Table 1). As revealed in Table 1, the best results were attained when the reaction was achieved in the presence of 1 mol% of bifunctional nano structure molten salt at room temperature (Table 1, entries 5 and 9). No improvement was identified in the yield of reaction via increasing the amount of the nano molten salt catalyst and the temperature. Table 1 clearly displays that in the absence of nano molten salt catalyst, reaction progressed slowly and the product was obtained with low yield (Table 1, entries 1 and 2).

>Table 1<

To compare the effect of the solution with solvent-free conditions, a mixture of 4-chlorobenzaldehyde, malononitrile and dimedone was selected as classic reaction, by 1 mol% of bifunctional nano structure molten salt as a catalyst in numerous solvents for instance H₂O, CH₃CN, C₂H₅OH, CH₂Cl₂, CH₃CO₂Et, *n*-hexane and toluene was studied at room temperature. As it can be detected in Table 2, solvent-free condition was the best one in this reaction. Consequently, solvent-free condition was selected for running condensation reactions.

>Table 2<

After optimization of the reaction conditions, to consider the efficacy and the scope of the presented procedure, various of tetrahydrobenzo[*b*]pyran derivatives were prepared by the one-pot three-component tandem Knoevenagel-Michael cyclocondensation reaction between aromatic aldehydes, malononitrile and dimedone in the presence of a catalytic amount of [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a novel bifunctional nano structure molten salt catalyst at

room temperature under solvent-free reaction conditions. The results are presented in Table 3. The effect of substituents on the aromatic ring displayed was estimated strong in terms of yields under same reaction conditions. All aromatic aldehydes including electron-releasing substituents and electron-withdrawing substituents on their aromatic ring provided the desired products in high to excellent yields in short reaction times. The reaction times of aromatic aldehydes possessing electron withdrawing groups were rather faster than electron donating groups.

>Table 3<

A plausible mechanism for the creation of the tetrahydrobenzo[*b*]pyran (**4**) is well-defined in Scheme 3 [16a]. Firstly, {[4,4'-BPyH][C(CN)₃]₂} as a nano molten salt (NMS) catalyst activates the carbonyl group of the aromatic aldehyde to provide intermediate (**8**) and malononitrile (**2**) tautomerized to (**9**). The tandem Knoevenagel condensation of intermediate (**8**) and (**9**) was happened to form the arylidene malononitrile (**11**). Subsequently, dimedone (**3**) tautomerized to (**10**) tolerates nucleophilic attack to (**11**) and affords the Michael adduct (**12**). The Michael adduct (**12**) tautomerizes using NMS catalyst to make intermediate (**13**) which cyclizes to provide compound (**14**) and which then tautomerized to afford the completely aromatized tetrahydrobenzo[*b*]pyran (**4**).

>Scheme 3<

Furthermore, reusability of the {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS catalyst was approved *via* condensation of 4-chlorobenzaldehyde, malononitrile and dimedone as a model. At the end of the reaction, ethyl acetate was added to the reaction mixture and heated to extract product from remained substances. This solution was washed with water to separate bifunctional NMS catalyst from other materials (the product is soluble in hot ethyl acetate and bifunctional NMS catalyst is soluble in water). The aqueous layer was decanted, separated and used for additional reaction after eliminating water. It is identified that the catalytic activity of the catalyst was restored within the limits of the experimental errors for four continuous runs (Table 4). The deactivation of the bifunctional NMS catalyst is low. The reaction was scaled up to 10 mmol of 4-chlorobenzaldehyde, malononitrile and dimedone by 10 mol% of {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS catalyst at room temperature. The yield of the reaction was 94% after 7 min and 78% after 12 min the fourth run.

>Table 4<

Application of [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional nano structure molten salt catalyst in the synthesis of amido or aminoalkyl naphthol derivatives.

After the synthesis of {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional nano structure molten salt catalyst, to optimize the reaction conditions, we approved the effect of the NMS catalyst in the synthesis of amino or amidoalkyl naphthols (Scheme 2). For this purpose, as a typical reaction, the condensation between acetamide, 4-chlorobenzaldehyde and β -naphthol was studied using different amounts of the catalyst at range of 25–125 °C under solvent-free conditions (Table 5). As Table 5 displays, the best results were attained when the reaction was achieved in the presence of 1 mol% of NMS catalyst at 75 °C (Table 5, entry 7). Increasing the reaction temperature and catalyst loading did not improve the rate of the reaction (Table 5, entries 9-13). In the absence of catalyst a low yield of the products was attained after 120 min (Table 5, entries 1 and 2).

>Table 5<

To compare the result of the solution with that of solvent-free conditions, a mixture of acetamide, 4-chlorobenzaldehyde and β -naphthol as a model reaction, using 1 mol% of NMS catalyst in numerous solvents such as H₂O, C₂H₅OH, CH₃CN, CH₃CO₂Et, CH₂Cl₂, *n*-Hexane and toluene was investigated under reflux conditions. Also, reaction was performed under solvent-free condition at 75 °C. As it can be seen in Table 6, solvent-free condition was the best one in this reaction.

>Table 6<

With the purpose of approving this main process, we decided to synthesize several amido or aminoalkyl naphthol derivatives from three-component condensation reaction of numerous aromatic aldehyde, β -naphthol and several amine or amide under solvent-free conditions at 75 °C in the presence of a catalytic amount of {[4,4'-BPyH][C(CN)₃]₂} as a NMS catalyst. The results are presented in Table 7. The effect of substituents on the aromatic ring exhibitions was estimated strong in terms of yields under these reaction conditions. All aromatic aldehydes bearing electron-withdrawing and electron-releasing substituents on their aromatic ring provided the related products in high to excellent yields in short reaction time. The reaction time of aromatic aldehydes having electron-withdrawing groups was rather faster than that of electron donating groups.

>Table 7<

In Scheme 4, we have proposed a probable mechanism for the synthesis of amido or aminoalkyl naphthol derivatives using {[4,4'-BPyH][C(CN)₃]₂} as a NMS catalyst. The reaction of aromatic aldehydes with β -naphthol in the presence of NMS catalyst is plausible to produce ortho-quinone methides (*o*-QMs). The prepared *o*-QMs, reacted with amide or amine derivatives to provide amido or aminoalkyl naphthols. A practical explanation for this result can be identified by studying the

nucleophilic addition of amide or amine to **o-QMs** intermediate and **eventually** the desired product was attained after aromatization [25, 26].

>Scheme 4<

Furthermore, **recyclability** and reusability of the {[4,4'-BPyH][C(CN)₃]₂} as a NMS catalyst was approved from the condensation between acetamide, 4-chlorobenzaldehyde and β -naphthol. After the completion of the reaction, ethyl acetate was added to the reaction mixture and heated to extract the product. This solution was washed with water to separate catalyst from other materials (the product is soluble in hot ethyl acetate and NMS catalyst is soluble in water). The aqueous layer was decanted, separated and used for **alternative** reaction after removing of water. It has been **proved** that the catalytic activity of the catalyst was changed within the limits of the experimental errors for five continuous runs. The deactivation of the NMS catalyst is low. The reaction was scaled up to 10 mmol of acetamide, 4-chlorobenzaldehyde and β -naphthol using 10 mol% of NMS catalyst at 75 °C. The yield of the reaction was 93% after 20 min and 91% after the fifth run. The results are summarized in Fig. 1.

>Fig. 1<

4. Conclusion

In summary, an environmentally friendly, green, mild and efficient bifunctional nano structure molten salt catalyst namely [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} was used and investigated in the synthesis of tandem Knoevenagel-Michael cyclocondensation synthesis of tetrahydrobenzo[b]pyran derivatives by three-component condensation reaction of various aldehydes, malononitrile and dimedone under solvent-free conditions at room temperature. **In another** study catalytic properties of {[4,4'-BPyH][C(CN)₃]₂}, it was applied in the synthesis of amido and aminoalkyl naphthol derivatives under solvent-free conditions at 75 °C. Other works offered that the NMS basicity and acidity displays a significant role to play in the dual-catalyzed reactions. Different noteworthy advantages of this study are reasonably ease of product isolation, cleaner reaction profile, short reaction time, high yield, recyclability of the NMS catalyst and accordance with the green chemistry disciplines. More study for systematic and knowledge-based development of nano molten salt is going on in our research group.

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Appendix A. Supplementary data

Supplementary data to this article can be found online at <http://dx.doi.org.10.1016/j.molliq.2016.xx.xxx>.

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Scheme, Table and Figure section:

Scheme 1. The synthesis of [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional nano structure molten salt catalyst.

Scheme 2. The synthesis of tetrahydrobenzo[*b*]pyrans and amido or aminoalkyl naphthol derivatives in the presence of [4,4'-bipyridine]-1,1'-diium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} (1 mol%) as a green bifunctional nano structure molten salt catalyst.

Table 1. Result of amount of the catalyst and temperature on the condensation reaction among 4-chlorobenzaldehyde, malononitrile and dimedone under solvent-free conditions.^a

Entry	Catalyst amount (mol%)	Reaction temperature (°C)	Reaction time (min)	Yield ^b (%)
1	—	r.t.	120	33
2	—	100	120	33
3	0.5	r.t.	60	81
4	0.5	100	60	81
5	1	r.t.	7	94
6	1	50	7	94
7	1	75	7	94
8	1	100	7	94
9	2	r.t.	7	94
10	2	100	7	94
11	5	r.t.	10	89
12	10	r.t.	10	89

Reaction conditions: ^a4-Chlorobenzaldehyde (1 mmol), malononitrile (1 mmol), dimedone (1 mmol); ^bIsolate yield.

Table 2. The effect of numerous solvents on the reaction between 4-chlorobenzaldehyde, malononitrile and dimedone catalyzed by {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS (1 mol%) at room temperature.^a

Entry	Solvent	Reaction time (min)	Yield ^b (%)
1	Solvent-free	7	94
2	H ₂ O	7	93
3	CH ₃ CN	7	91
4	C ₂ H ₅ OH	7	93
5	CH ₂ Cl ₂	15	89
6	CH ₃ CO ₂ Et	20	83
7	<i>n</i> -Hexane	30	77
5	Toluene	30	77

Reaction conditions: ^a4-Chlorobenzaldehyde (1 mmol), malononitrile (1 mmol), dimedone (1 mmol); ^bIsolate yield.

Table 3. The tandem Knoevenagel-Michael cyclocondensation reaction synthesis of tetrahydrobenzo[*b*]pyran derivatives using [4,4'-bipyridine]-1,1'-diium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS catalyst.^a

Entry	Aldehyde	Time (min)	Yield ^b (%)	M.p (°C) [Lit.] ^{Ref.} (Color)
1		7	94	210-212 [208-210] ^{16a} (White solid)
2		10	90	227-229 [229-230] ^{16a} (White solid)
3		8	91	180-182 [175-177] ^{16p} (White solid)
4		7	95	226-228 [225-228] ¹⁶ⁿ (White solid)
5		18	87	219-221 [214-216] ^{16a} (White solid)

6		11	92	211-213 [202-204] ^{16a} (White solid)
7		15	88	212-214 [218-219] ^{16a} (White solid)
8		5	96	188-190 [178-180] ^{16a} (White solid)
9		10	90	195-197 [208-212] ^{16a} (White solid)
10		10	91	120-122 [183-186] ^{16a} (White solid)
11		15	89	179-181 [158-160] ^{16p} (White solid)
12		10	90	215-217 [216-217] ^{16a} (White solid)
13		9	91	157-159 [150-152] ^{16a} (White solid)
14		8	92	205-207 [207-208] ^{16a} (White solid)
15		17	85	269-271 (White solid)
16		14	90	206-208 [212-213] ^{16a} (Orange solid)

17		9	94	230-232 [235-237] ^{16p} (White solid)
18		10	92	235-237 [215-217] ^{16p} (White solid)

Reaction conditions: ^aAromatic aldehyde (1 mmol), malononitrile (1 mmol), dimedone (1 mmol), {[4,4'-BPyH][C(CN)₃]₂} as a NMS catalyst (1 mol%), solvent-free, r.t.; ^bIsolate yield.

Scheme 3. The proposed mechanism for the tandem Knoevenagel-Michael cyclocondensation synthesis of tetrahydrobenzo[*b*]pyran derivatives using {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS catalyst.

Table 4. Reusability of {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS catalyst on the synthesis of 2-amino-4-(4-chlorophenyl)-7,7-dimethyl-5-oxo-5,6,7,8-tetrahydro-4*H*-chromene-3-carbonitrile.^a

Experimental errors	Time (min)	Isolated yields ^b
1	7	94
2	8	93
3	11	85
4	12	78

Reaction conditions: ^a4-Chlorobenzaldehyde (10 mmol), malononitrile (10 mmol), dimedone (10 mmol), {[4,4'-BPYH][C(CN)₃]₂} as a NMS catalyst (10 mol%), Solvent-free, r.t.; ^bIsolate yield.

Table 5. Result of amount of the catalyst and temperature on the condensation reaction between acetamide, 4-chlorobenzaldehyde and β -naphthol under solvent-free conditions.^a

Entry	Catalyst amount (mol%)	Reaction temperature (°C)	Reaction time (min)	Yield ^b (%)
1	—	r.t.	120	5
2	—	100	120	7
3	0.5	r.t.	120	23
4	0.5	100	60	25
5	1	r.t.	60	51
6	1	50	60	78
7	1	75	20	93
8	1	100	20	93
9	1	125	20	93
10	2	r.t.	60	51
11	2	100	20	93
12	5	r.t.	60	51
13	5	100	20	93

Reaction conditions: ^a4-Chlorobenzaldehyde (1 mmol), β -naphthol (1 mmol), acetamide (1 mmol); ^bIsolate yield.

Table 6. Reaction of acetamide, 4-chlorobenzaldehyde and β -naphthol catalyzed using {[4,4'-BPYH][C(CN)₃]₂} as a bifunctional NMS (1 mol%) in various solvents under reflux conditions. Reaction was done under solvent-free condition at 75 °C .^a

Entry	Solvent	Reaction time (min)	Yield ^b (%)
1	Solvent-free	20	93
2	H ₂ O	60	18
3	CH ₃ CN	45	78
4	C ₂ H ₅ OH	60	35
5	CH ₂ Cl ₂	60	71
6	CH ₃ CO ₂ Et	60	29

7	<i>n</i> -Hexane	60	15
5	Toluene	60	21

Reaction conditions: ^a4-Chlorobenzaldehyde (1 mmol), β -naphthol (1 mmol), acetamide (1 mmol); ^bIsolate yield.

Table 7. The synthesis of amino or amidolalkyl naphthol derivatives in the presence of [4,4'-bipyridine]-1,1'-dium tricyanomethanide {[4,4'-BPyH][C(CN)₃]₂} as a bifunctional NMS catalyst.^a

Entry	Aldehyde	R'	Time (min)	Yield ^b (%)	M.p (°C) [Lit.] ^{Ref.} (Color)
1		COMe	15	95	243-245 [248-250] ²⁵ (Yellow solid)
2		COPh	20	93	248-250 [253-255] ²⁵ (White solid)
3		CONH ₂	25	92	195-197 [200-202] ²⁵ (Yellow solid)
4		2-Pyridyl	15	95	181-183 [189-191] ²⁶ (Cream solid)
5		4-Pyridyl	10	96	281-283 [290-292] ²⁶ (Brown solid)
6		COMe	20	93	254-256 [258-260] ²⁵ (White solid)
7		COPh	25	90	201-203 [205-207] ²⁵ (White solid)

8		CONH ₂	30	88	171-173 [161-163] ²⁵ (White solid)
9		2-Pyridyl	20	93	201-203 [209-221] ²⁶ (Cream solid)
10		4-Pyridyl	15	95	217-219 [218-220] ²⁶ (Cream solid)
11		COMe	25	91	274-276 [280-282] ²⁵ (White solid)
12		COPh	30	88	264-266 [271-273] ²⁵ (Cream solid)
13		CONH ₂	35	87	203-205 [206-208] ²⁵ (Cream solid)
14		2-Pyridyl	25	91	163-165 [172-174] ²⁶ (Cream solid)
15		4-Pyridyl	20	93	253-255 [263-265] ²⁶ (Cream solid)
16		CONH ₂	35	85	181-183 [177-179] ²⁵ (White solid)

Reaction conditions: ^aAromatic aldehyde (1 mmol), β -naphthol (1 mmol), amine or amide derivatives (1 mmol), {[4,4'-BPyH][C(CN)₃]₂} as a NMS catalyst (1 mol%), solvent-free, 75 °C; ^bIsolate yield.

Scheme 4. Suggested mechanism for the synthesis of amido or aminoalkyl naphthol derivatives using {[4,4'-BPyH][C(CN)₃]₂} as a NMS catalyst.

Fig. 1. Reusability of the NMS catalyst in 20 minutes in the synthesis of corresponded amidoalkyl naphthol.

[4,4'-Bipyridine]-1,1'-diium tricyanomethanide as a nanostructured molten salt and its catalytic application in the synthesis of tetrahydrobenzo[*b*]pyrans, amido and aminoalkyl naphthol derivatives

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Graphical Abstract