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# One-pot synthesis of 3,4-dihydropyrimidin-2(1H)-ones catalyzed by SO<sub>3</sub>H@imineZCMNPs as a novel, efficient and reusable acidic nanocatalyst under solvent-free conditions

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The synthesis of 3,4-dihydropyrimidin-2(1H)-one derivatives was accomplished efficiently *via* a three-component reaction between ethyl acetoacetate, various types of aldehydes, and urea in the presence of 10 mg SO<sub>3</sub>H@imineZCMNPs as a novel, environment friendly, and reusable heterogeneous magnetic nanocatalyst under solvent-free conditions at 90 °C. The desired products were obtained with high quantitative yields. The catalyst was separated by simple isolation from the reaction mixture using a permanent magnet and reused several times without any significant loss of catalytic activity. The synthesized catalyst was fully characterized through various techniques including thermogravimetric analysis, Fourier transform infrared spectroscopy, scanning electron microscopy, transmission electron microscopy, X-ray diffraction, and the Hammett acidity test. This methodology tolerates most substrates and has the salient features of green reaction conditions, lower catalyst loading, high quantitative yields, low cost, the absence of solvents, and easy isolation and reusability of the catalyst.

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## Introduction

Energy and stability are challenging and vital issues from universal economic and social viewpoints. These issues are connected to the concept of green chemistry and make the science of catalysts more creative. The philosophy behind green chemistry deals with encouraging chemical and engineering researchers to design products and processes with minimum usage and production of harmful materials.<sup>1–6</sup> Therefore, green catalysts are considered as an important subset of green chemistry. In recent years, green and environment-friendly syntheses have played a central role in increasing efficiency, preventing the use of volatile solvents and toxic reactants, and reducing reaction times.<sup>7–9</sup> In the dissimilation process, the catalysts are trapped by the surface or pores of a solid bed such as silica or alumina.<sup>10</sup> The connection of the catalyst to the intended surface can occur by simple absorption of the active molecules of the catalyst and/or covalent bonds as the covalent bond of the catalyst and the surface is of utmost importance. The formation of such bonds results in the connection of the catalyst to the surface. This connection will be tight enough, and washing the catalyst from the surface reaches its minimum

in harsh reaction conditions. However, a limited access to the active centers of heterogeneous catalysts leads to the reduction of their activity compared to homogeneous catalysts.<sup>11–13</sup> To resolve this issue, all the centers of the porous solid bed should be activated, as in most cases, only the active centers on the external surface of the porous solid bed are available for reaction. A homogeneous system and a heterogeneous one are similar in terms of higher number of active centers and a facile recovery in the catalyst system. Achieving this dual purpose is possible only by using nanocatalysts. The benefit of nanocatalysts is that it allows for the favorable properties of both catalysts by filling the gap between homogeneous and heterogeneous catalysts.<sup>14–17</sup> Heterogeneous nanocatalysts, which are prepared by specific methods in nanoscale, have many applications in different industries because of the multiplicity of active surfaces and the selectivity, high stability, and extensive use of the heterogeneous catalyst.<sup>18–21</sup> Although nanocatalysts have many benefits compared to typical catalyst systems, the separation and recovery of small nanocatalysts from the reaction mixture does not appear to be an easy task. Moreover, typical methods of filtration do not show good efficacy because of the small size of the nanocatalyst particles.<sup>22,23</sup> To solve this problem, a suitable solution is to employ magnetic nanoparticles. The paramagnetic and insoluble natures of nanoparticles make their recovery possible by an external magnet; consequently, the catalyst could be used in another cycle.<sup>24–32</sup>

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The compounds containing dihydropyrimidinone (DHMP) moiety have wide pharmaceutical and biological activities such as anti-bacterial, anti-oxidant, anti-HIV, and anti-cancer activities.<sup>33,34</sup> In this regard, 3,4-dihydropyrimidin-2(1H)-ones are a common type of these heterocyclic compounds that can be prepared by one-pot condensation of ethyl acetoacetate, urea and an aldehyde under strongly acidic conditions.<sup>35</sup> Previously, different derivatives of 3,4-dihydropyrimidine-2(1H)-one have exhibited calcium channel modulators for the treatment of cardiovascular diseases.<sup>36</sup> Some of the reported catalysts for the preparation of these types of heterocyclic molecules are:  $\text{Ln}(\text{OTf})_3$ ,<sup>37</sup> *p*-dodecylbenzenesulfonic acid,<sup>38</sup> chiral phosphoric acid,<sup>39</sup> sulfated tungstate,<sup>40</sup>  $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ ,<sup>41</sup>  $\text{Mn}(\text{OAc})_3 \cdot 2\text{H}_2\text{O}$ ,<sup>42</sup>  $\text{Ce}(\text{LS})_3$ ,<sup>43</sup> and  $\text{SnCl}_2$ -nano  $\text{SiO}_2$ .<sup>44</sup> Although numerous strategies have been reported for the production of 3,4-dihydropyrimidin-2(1H)-ones, some of these protocols suffer from disadvantages such as the need for excess amounts of the catalyst, the need for microwave or ultrasound irradiation, the use of high temperatures, the use of a toxic solvent, require multiple steps, and long reaction times. Therefore, it is essential to develop an improved procedure for the synthesis of 3,4-dihydropyrimidin-2(1H)-ones under mild reaction conditions.

Herein, we decided to investigate the synthesis of 3,4-dihydropyrimidin-2(1H)-ones with various substituents from the reaction of ethyl acetoacetate, aldehyde and urea under solvent-free conditions using  $\text{SO}_3\text{H@imineZCMNPs}$  as a novel, eco-friendly, reusable and promising nanocatalyst.

## Experimental

### General

All reagents and solvents were obtained from Fluka or Merck chemical companies. The purity determination of the substrates and reaction monitoring were carried out with TLC on silica gel 60 F254 (0.25 mm thick). FTIR spectra were recorded using KBr pellets in the range of 400–4000  $\text{cm}^{-1}$  with a PerkinElmer PXI spectrometer. The powder XRD patterns of samples were carried out in the  $2\theta$  range of 10–80° at room temperature on a Philips X-pert diffractometer (Holland), using Ni-filtered  $\text{Co-K}\alpha$  radiation ( $\lambda = 1.54 \text{ \AA}$ ). Scanning electron microphotography (SEM) was performed on an LEO 1430VP instrument. TEM images were obtained with a Zeiss-EM10C transmission electron microscope with an acceleration voltage of 80 kV (Oberkochen, Germany). Thermogravimetric analyses (TGA) were performed from room temperature to 700 °C at a ramp of 10 °C  $\text{min}^{-1}$  under  $\text{N}_2$  atmosphere using a Linseis STA PT 1000 instrument.

### Catalyst synthesis

**Preparation of the magnetic  $\text{Fe}_3\text{O}_4$  nanoparticles (MNPs).** The magnetite nanoparticles were synthesized according to the method reported in ref. 45. Briefly, 7.0 g of  $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$  and 2.4 g of  $\text{FeCl}_2 \cdot 4\text{H}_2\text{O}$  were dissolved in 40 mL of distilled water in a three-necked round-bottom flask (250 mL) under stirring at 90 °C and a continuous flow of Ar atmosphere. Thereafter, under intense mechanical stirring, 10 mL of ammonia (25%)

was added to the reaction mixture. The resulting nanoparticles were magnetically separated and subsequently washed three times with deionized water and then dried under vacuum at 60 °C for 12 h.

**Preparation of zirconia coated magnetic nanoparticles (ZCMNPs).** The  $\text{Fe}_3\text{O}_4@\text{ZrO}_2$  core-shell nanoparticles were prepared according to the method proposed by Wu with slight modifications.<sup>46</sup> Briefly, 1 g of  $\text{ZrOCl}_2 \cdot 8\text{H}_2\text{O}$  was dissolved in 20 mL of a 5 : 3 (v/v) mixture of ethanol and water and heated at 70 °C for 2 h. Then, under continuous mechanical stirring, 5.0 g of  $\text{Fe}_3\text{O}_4$  nanoparticles was added into the zirconia sol. The suspension was homogenized under ultrasonication for 1 h. After 12 h, the coated particles were finally separated from the liquid by magnetic decantation and dried under vacuum at 300 °C for 2 h.

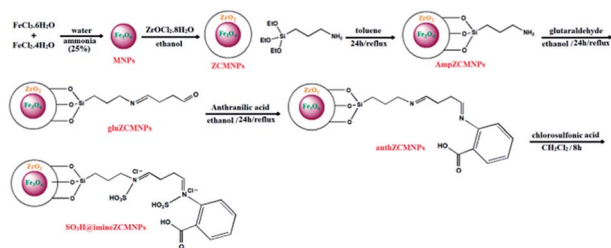
**Functionalization of  $\text{Fe}_3\text{O}_4@\text{ZrO}_2$  with 3-aminopropyltriethoxysilane (AmpZCMNPs).** Core-shell  $\text{Fe}_3\text{O}_4@\text{ZrO}_2$  nanoparticles (1 g) were dispersed in dry toluene (20 mL) for 20 min using ultrasound. Then, 3-aminopropyltriethoxysilane (Amp, 2 mL) was added to the resulting mixture. The mixture was refluxed for 24 h under argon atmosphere. After 24 h, the functionalized magnetite nanoparticles (AmpZCMNPs) were separated from the reaction mixture using a permanent magnet, washed several times with ethanol and distilled water, and dried in a vacuum oven.

**Immobilization of anthranilic acid on the surface of zirconia-coated magnetite nanoparticles (anthZCMNPs).** For the preparation of anthZCMNPs, AmpZCMNPs (2 g) was dispersed in ethanol (100 mL) for 30 min using ultrasonication. About 4 mmol of excess glutaraldehyde was added to the mixture and refluxed for 24 h. After this period of time, the resultant solid (named as gluZCMNPs) was separated by an external magnet and washed with ethanol several times; the unreacted glutaraldehyde residue was removed, and the solid was then dried in a vacuum oven. Afterward, the prepared gluZCMNPs (2 g) were suspended in 100 mL of ethanol. Anthranilic acid (4 mmol) was added to this solution under dry nitrogen atmosphere, and the mixture was refluxed for 24 h. After 24 h, the resultant solid, named as anthZCMNPs, was filtered, and 500 mL ethanol was added to it in order to remove the unreacted reagents. The resultant material was then dried in a vacuum oven.

**Immobilization of  $\text{SO}_3\text{H}$  on the surface of magnetite nanoparticles ( $\text{SO}_3\text{H@imineZCMNPs}$ ).** For the preparation of  $\text{SO}_3\text{H@imineZCMNPs}$ , 2 g of anthZCMNPs was suspended in 50 mL dry dichloromethane; 10 mmol chlorosulfonic acid was added dropwise to the vessel, and the reaction mixture was stirred for 8 h. After separation with an external magnet, the product was washed with dry dichloromethane to remove the unreacted chlorosulfonic acid. The resultant  $\text{SO}_3\text{H@imineZCMNPs}$  material was then dried under vacuum at 80 °C. All stages of the  $\text{SO}_3\text{H@imineZCMNPs}$  synthesis are shown in Scheme 1.

**General procedure for the synthesis of 3,4-dihydropyrimidin-2(1H)-one.** A mixture of ethyl acetoacetate (1 mmol), aldehyde (1 mmol), urea (1.2 mmol), and  $\text{SO}_3\text{H@imineZCMNPs}$  (10 mg) was stirred under solvent-free conditions at 90 °C for the required period of time (Table 3). After completion of the



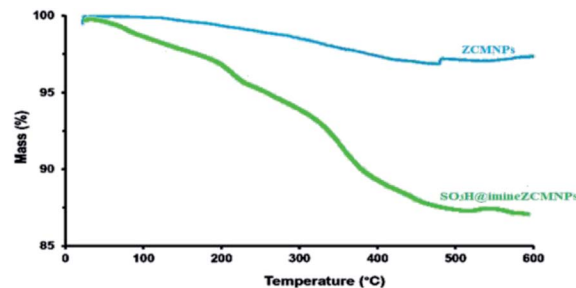
Scheme 1 All stages of the  $\text{SO}_3\text{H@imineZCMNPs}$  synthesis.

reaction (monitored by TLC), it was observed that most of the nanoparticles were adsorbed onto the magnetic stirring bar when the magnetic stirring was turned off. After cooling to room temperature, the catalyst was separated using a permanent magnet and reused for the next experiment. The pure product was obtained *via* recrystallization from ethanol.

## Results and discussion

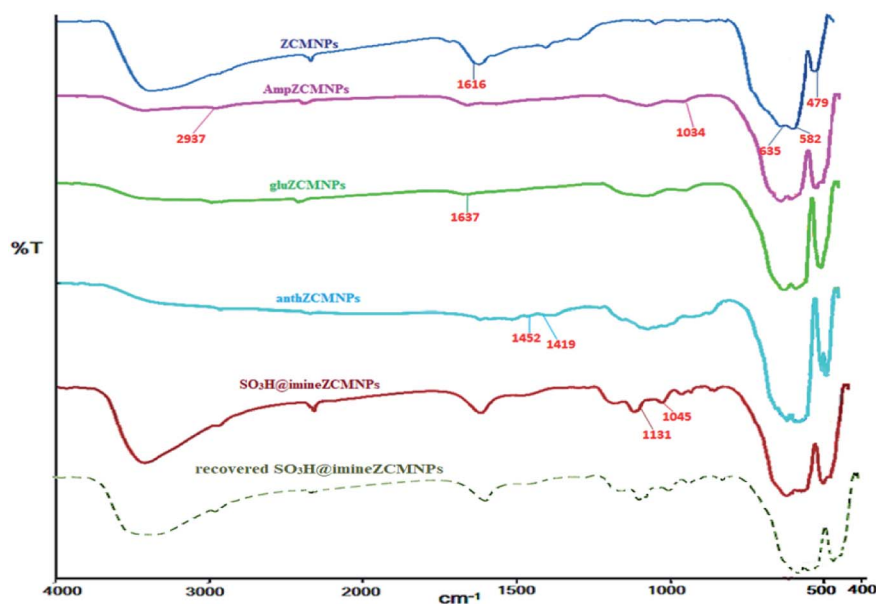
### Catalyst characterization

**IR analysis of  $\text{SO}_3\text{H@imineZCMNPs}$ .** The infrared spectra of the ZCMNPs, AmpZCMNPs, gluZCMNPs, anthZCMNPs, and  $\text{SO}_3\text{H@imineZCMNPs}$  samples in the region of  $400\text{--}4000\text{ cm}^{-1}$  are presented in Fig. 1. In the ZCMNPs spectrum, the absorption peaks at  $479\text{ cm}^{-1}$  and  $582\text{ cm}^{-1}$  can be indexed to the Fe–O bonds in the crystalline lattice of  $\text{Fe}_3\text{O}_4$ . The peak observed at  $635\text{ cm}^{-1}$  is ascribed to the Zr–O groups. The peak observed at  $1616\text{ cm}^{-1}$  is related to the H–OH bending vibrations of the physically adsorbed water, while the peaks appearing in the  $3100\text{--}3500\text{ cm}^{-1}$  region can be attributed to the stretching vibrations of the OH bonds in water molecules. In the case of the AmpZCMNPs, the bands at  $1034\text{ cm}^{-1}$  can be attributed to

Fig. 2 TGA curves of ZCMNPs and  $\text{SO}_3\text{H@imineZCMNPs}$ .

Si–O stretching vibration mode. The absorption bands appearing at  $2937\text{ cm}^{-1}$  (aliphatic C–H group) demonstrate the successful modification of the magnetite nanoparticles. In the gluZCMNPs spectrum, the stretching vibration of the imine C=N group appears at  $1637\text{ cm}^{-1}$ , which confirms the successful modification of the surface of the AmpZCMNPs. In the case of the anthZCMNPs, the absorption peaks at  $1419$  and  $1452\text{ cm}^{-1}$  can be assigned to the stretching vibration of the C=C bond. The presence of the  $\text{SO}_3\text{H}$  groups was confirmed by the stretching vibrations appearing at  $1045$  and  $1131\text{ cm}^{-1}$  in the FTIR spectrum of the compound named  $\text{SO}_3\text{H@imineZCMNPs}$ . Also, the increase in the intensities of the peaks at  $2700\text{--}3600\text{ cm}^{-1}$  suggests that there were more OH functional groups under the magnetic nanoparticle surface after sulfonation. It should be noted that the spectrum of recovered  $\text{SO}_3\text{H@imineZCMNPs}$  after the first recovery and reuse did not show any change.

**Thermal studies of  $\text{SO}_3\text{H@imineZCMNPs}$ .** TGA analyses were carried out to characterize the ZCMNPs and  $\text{SO}_3\text{H@imineZCMNPs}$  samples toward each other. The results of these analyses are presented in Fig. 2. In the TGA curve of

Fig. 1 FT-IR spectra of  $\text{SO}_3\text{H@imineZCMNPs}$ .

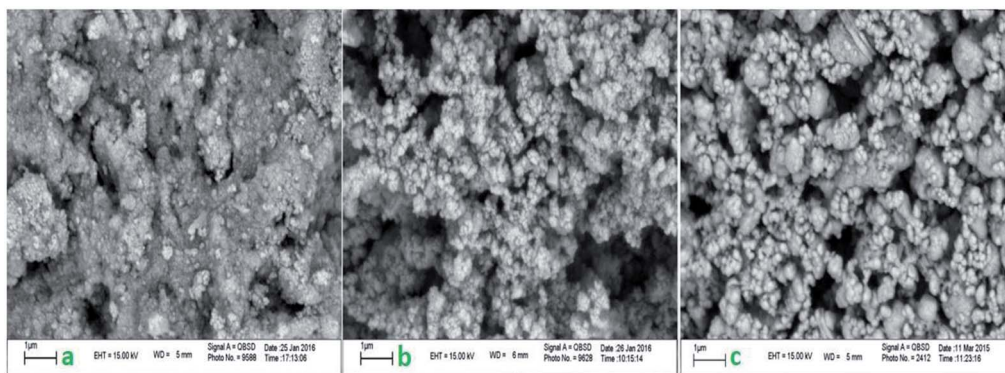


Fig. 3 SEM images of ZCMNPs (a), anthZCMNPs (b), and  $\text{SO}_3\text{H@imineZCMNPs}$  (c).

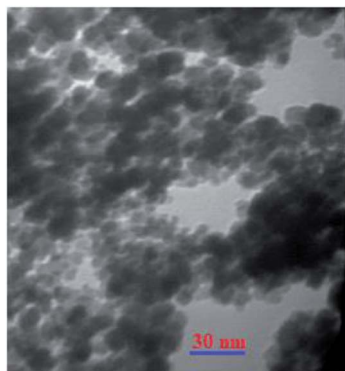


Fig. 4 TEM images of  $\text{SO}_3\text{H@imineZCMNPs}$ .

ZCMNPs, the weight loss observed at temperatures below  $320^\circ\text{C}$  is related to water desorption from the surface of the zirconia layer while weight loss appearing above  $600^\circ\text{C}$  can be attributed to the release of the water structure. The TGA of  $\text{SO}_3\text{H@imineZCMNPs}$  illustrates an approximately 9% weight loss

in the temperature range from  $200$  to  $600^\circ\text{C}$  apart from two water loss events, resulting from the decomposition of the organic groups grafted to the zirconia surface and the degradation of the sulfonic acid parts.

**SEM analysis of  $\text{SO}_3\text{H@imineZCMNPs}$ .** The surface morphology, particle shape and size distribution of the  $\text{SO}_3\text{H@imineZCMNPs}$  were examined by SEM (scanning electron microscopy) as shown in Fig. 3. The SEM image of the  $\text{SO}_3\text{H@imineZCMNPs}$  clearly demonstrates variations in the surfaces of these nanoparticles. The SEM images illustrate that these nanoparticles are of nearly spherical morphology with a mean diameter of about  $45\text{ nm}$ .

**TEM analysis of  $\text{SO}_3\text{H@imineZCMNPs}$ .** The transmission electron micrograph of the prepared  $\text{SO}_3\text{H@imineZCMNPs}$  determined that they appear to have an almost spherical shape with a narrow size distribution (the average particle size is  $35\text{ nm}$ , Fig. 4), thereby retaining a nanocrystalline appearance. Therefore, the major active sites of the  $\text{SO}_3\text{H@imineZCMNPs}$  may allow for high activity in organic transformations, even with a low catalyst loading.

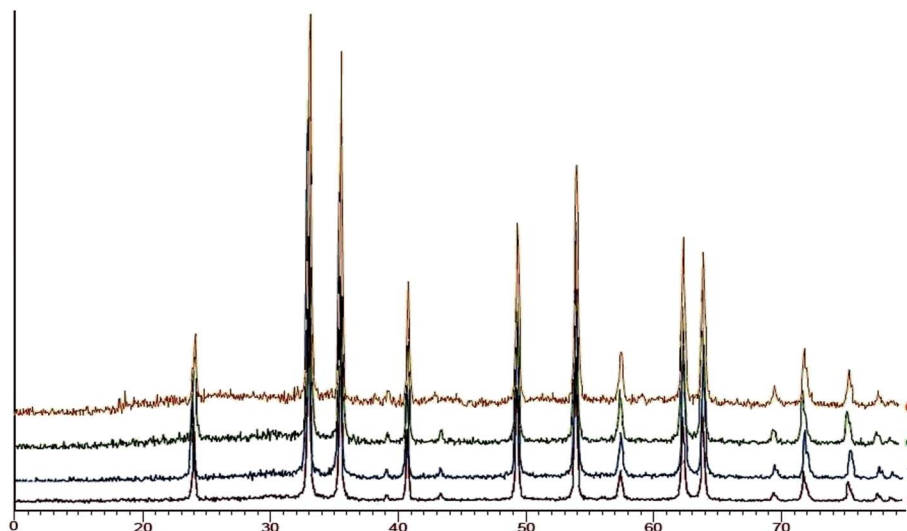


Fig. 5 XRD patterns of ZCMNPs (a), anthZCMNPs (b),  $\text{SO}_3\text{H@imineZCMNPs}$  (c) and recovered  $\text{SO}_3\text{H@imineZCMNPs}$  (d).



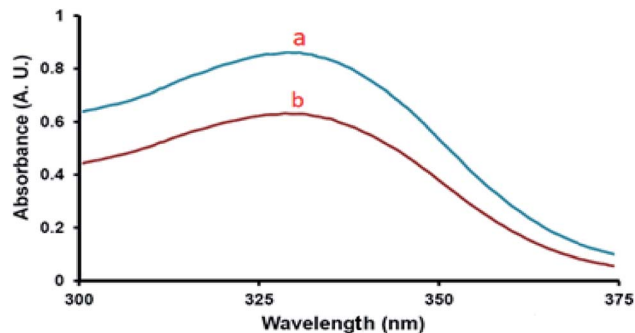


Fig. 6 Absorption spectra of (a) 4-nitroaniline (indicator) and (b)  $\text{SO}_3\text{H@imineZCMNPs}$  (catalyst) in  $\text{CCl}_4$ .

Table 1 Calculation of the Hammett acidity function ( $H_0$ ) of  $\text{SO}_3\text{H@imineZCMNPs}^a$

Entry	Catalyst	$[\text{I}]_s$ (%)	$[\text{IH}^+]_s$ (%)	$H_0$
1	—	100.0	0.0	—
2	$\text{SO}_3\text{H@imineZCMNPs}$	75.0	25.0	1.47

<sup>a</sup> Conditions for UV-visible spectrum measurement: solvent,  $\text{CCl}_4$ ; indicator, *p*-nitroaniline ( $\text{p}K(\text{I})_{\text{aq}} = 0.99$ ),  $1.44 \times 10^{-4} \text{ mol L}^{-1}$ ; catalyst (10 mg)  $25^\circ\text{C}$ .

**XRD analysis of  $\text{SO}_3\text{H@imineZCMNPs}$ .** The X-ray diffraction patterns (XRD) for ZCMNPs, anthZCMNPs, and  $\text{SO}_3\text{H@imineZCMNPs}$  are given in Fig. 5. As can be seen, the XRD patterns of anthZCMNPs and  $\text{SO}_3\text{H@imineZCMNPs}$  are similar to those of ZCMNPs, suggesting that the modification does not have any considerable effect on the phase of ZCMNPs. Also, the XRD patterns of recovered  $\text{SO}_3\text{H@imineZCMNPs}$  after the first recovery and reuse do not show any change.

The acidity strength of our catalyst can be expressed by the Hammett acidity function ( $H_0$ ). It can be obtained from the following equation:

$$H_0 = \text{p}K(\text{I})_{\text{aq}} + \log([\text{I}]_s/[\text{IH}^+]_s)$$

where 'I' is the indicator base and  $[\text{I}]_s$  and  $[\text{IH}^+]_s$  are the unprotonated and protonated indicator molar concentrations, respectively. The  $\text{p}K(\text{I})_{\text{aq}}$  value of 4-nitroaniline as a basic indicator to abstract proton is 0.99. Also, the value of  $[\text{I}]_s/[\text{IH}^+]_s$  according to the Lambert–Beer's law could be determined using the UV-visible spectrum. In our experiment, an unprotonated

form of the 4-nitroaniline in  $\text{CCl}_4$  showed an absorbance maximum at 330 nm.

As can be seen from Fig. 6, the absorbance of the unprotonated form of the indicator in  $\text{SO}_3\text{H@imineZCMNPs}$  was weak compared with the indicator sample in  $\text{CCl}_4$ , suggesting that the indicator was partially in the  $[\text{IH}^+]$  form. The obtained results of the acidity strength of  $\text{SO}_3\text{H@imineZCMNPs}$  are listed in Table 1.

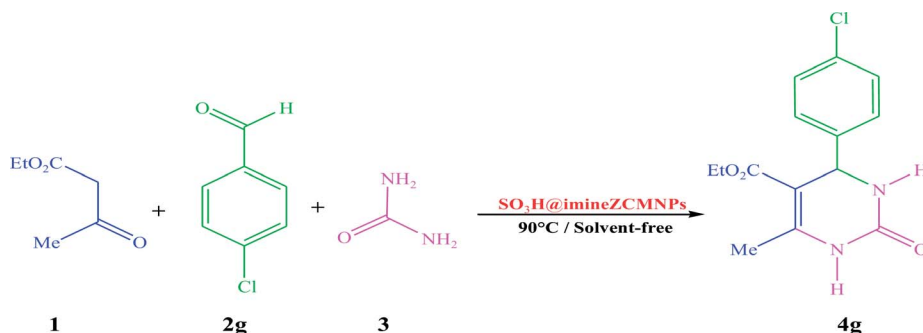
In continuation of our efforts on the synthesis of novel catalysts in performing organic transformations,<sup>47–49</sup> we decided to investigate the synthesis of 3,4-dihydropyrimidin-2(1*H*)-one derivatives from the reaction of ethyl acetoacetate, various aldehyde, and urea under solvent-free conditions using  $\text{SO}_3\text{H@imineZCMNPs}$  as a new, environment friendly, and reusable heterogeneous magnetic nanocatalyst (Scheme 2).

Solvents are widely applied in a myriad of applications as reactants, reaction mediums or carriers in chemical transformations.<sup>2,50</sup> In order to screen the reaction conditions for the synthesis of 3,4-dihydro-5-etoxy carbonyl-4-(4-chlorophenyl)-6-methylpyrimidine-2(1*H*)-one (**4g**), the reaction of ethyl acetoacetate (**1**), 4-chlorobenzaldehyde (**2g**) and urea (**3**) (molar ratio 1.0 : 1.0 : 1.2) toward the corresponding product was considered as a model reaction. Next, different variables affecting the reaction yield, including the amount of the catalyst, solvent, and temperature were investigated. The results are summarized in Table 2. To obtain the optimal reaction media, different solvents such as  $\text{H}_2\text{O}$ ,  $\text{CH}_2\text{Cl}_2$ ,  $\text{CHCl}_3$ , DMSO, DMF, EtOH, toluene, and solvent-free conditions in the presence of a catalytic amount (10 mg) of  $\text{SO}_3\text{H@imineZCMNPs}$  were used. Surprisingly, a short reaction time and the highest yield of the product (98%) was achieved when the same reaction was carried out while utilizing the same amount of catalysts at  $90^\circ\text{C}$  under solvent-free conditions (entry 8). The initial optimization experiments revealed that low yields of the desired products were obtained in the presence of DMSO and DMF as coordinating solvents even after 1 h under reflux conditions (entries 3 and 4). Moderate to good yields were obtained when the same reaction was performed while utilizing  $\text{H}_2\text{O}$  and EtOH, respectively, instead of coordinating solvents (entries 1 and 2). It is also remarkable that with EtOH, a lower reaction time was required compared to  $\text{H}_2\text{O}$ . These solvents play a negative role by retarding the multi-component pathway. This might be due to the solvent adsorption on the acidic catalyst surface or the solvent–reactant interactions. In contrast, using  $\text{CH}_2\text{Cl}_2$  and  $\text{CHCl}_3$  gave the desired product in lower yields (entries 5 and 6). A closer examination indicated that when toluene was replaced



Scheme 2 One-pot three-component reaction of ethyl acetoacetate (**1**), arylaldehyde (**2**) and urea (**3**) catalyzed by  $\text{SO}_3\text{H@imineZCMNPs}$  under solvent-free conditions.



**Table 2** Optimization of the three-component reaction of ethyl acetoacetate (**1**), 4-chlorobenzaldehyde (**2g**) and urea (**3**) under various conditions<sup>a</sup>

Entry	Solvent	Amount of catalyst (mg)	Temperature ( $^\circ\text{C}$ )	Reaction time (min)	Yield <sup>b</sup> (%)
1	H <sub>2</sub> O	10	Reflux/100 $^\circ\text{C}$	30	52
2	EtOH	10	Reflux/78 $^\circ\text{C}$	20	86
3	DMSO	10	Reflux/189 $^\circ\text{C}$	60	41
4	DMF	10	Reflux/153 $^\circ\text{C}$	60	37
5	CH <sub>2</sub> Cl <sub>2</sub>	10	Reflux/40 $^\circ\text{C}$	60	31
6	CHCl <sub>3</sub>	10	Reflux/61 $^\circ\text{C}$	60	25
7	Toluene	10	Reflux/110 $^\circ\text{C}$	30	74
8	—	10	90 $^\circ\text{C}$	8	98
9	—	5	90 $^\circ\text{C}$	8	81
10	—	15	90 $^\circ\text{C}$	8	92
11	—	10	25 $^\circ\text{C}$	8	21
12	—	10	80 $^\circ\text{C}$	8	92
13	—	10	100 $^\circ\text{C}$	8	90

<sup>a</sup> Reaction conditions: a mixture of ethyl acetoacetate (**1**), 4-chlorobenzaldehyde (**2g**), urea (**3**) (molar ratio 1 : 1 : 1.2) and  $\text{SO}_3\text{H@imineZCMNPs}$  (10 mg). <sup>b</sup> Isolated yield.

with the above-mentioned chlorinated solvents, the yield was markedly improved (entry 7). In the next phase of the survey, the effect of different catalyst amounts on the completion of the

reaction was evaluated (entries 8–10). As noted above, the best result was obtained for 10 mg loading (entry 8). Decreasing the amount of catalyst required for the reaction produced a lower

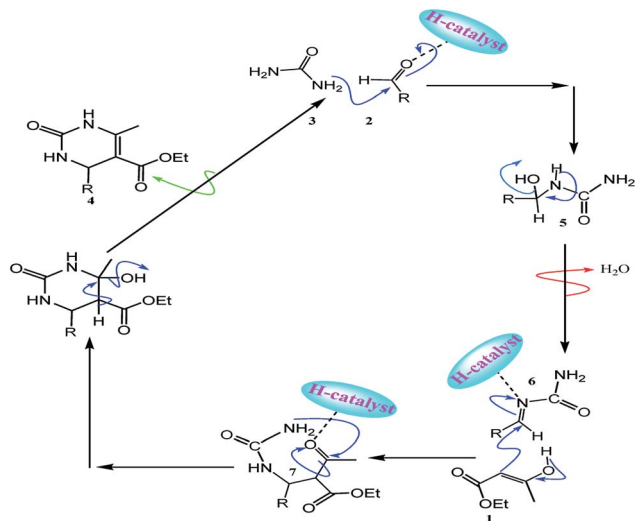
**Table 3** The  $\text{SO}_3\text{H@imineZCMNPs}$  catalyzed three-component Biginelli coupling<sup>a</sup>

Entry	Product	R	Yield <sup>b</sup> (%)	Time (min)	Mp <sup>c</sup> ( $^\circ\text{C}$ ) found	Mp ( $^\circ\text{C}$ ) (Lit.)
1	<b>4a</b>	C <sub>6</sub> H <sub>5</sub> –	93	8	202–204	202–203 (57)
2	<b>4b</b>	2-(NO <sub>2</sub> )–C <sub>6</sub> H <sub>4</sub> –	95	8	218–220	220–222 (42)
3	<b>4c</b>	3-(NO <sub>2</sub> )–C <sub>6</sub> H <sub>4</sub> –	95	8	226–228	226–228 (57)
4	<b>4d</b>	4-(NO <sub>2</sub> )–C <sub>6</sub> H <sub>4</sub> –	98	8	211–213	208–211 (63)
5	<b>4e</b>	2-(Cl)–C <sub>6</sub> H <sub>4</sub> –	97	8	217–219	215–218 (44)
6	<b>4f</b>	3-(Cl)–C <sub>6</sub> H <sub>4</sub> –	96	8	192–195	192–193 (56)
7	<b>4g</b>	4-(Cl)–C <sub>6</sub> H <sub>4</sub> –	98	8	211–214	212–214 (51)
8	<b>4h</b>	2,4-(Cl) <sub>2</sub> –C <sub>6</sub> H <sub>3</sub> –	98	8	252–255	251–252 (60)
9	<b>4i</b>	3,4-(Cl) <sub>2</sub> –C <sub>6</sub> H <sub>3</sub> –	97	8	222–225	222–223 (62)
10	<b>4j</b>	4-(F)–C <sub>6</sub> H <sub>4</sub> –	95	8	178–180	175–177 (35)
11	<b>4k</b>	4-(CF <sub>3</sub> )–C <sub>6</sub> H <sub>4</sub> –	98	8	177–179	173–175 (58)
12	<b>4l</b>	3-(Br)–C <sub>6</sub> H <sub>4</sub> –	94	8	186–189	185–186 (61)
13	<b>4m</b>	2-(OH)–C <sub>6</sub> H <sub>4</sub> –	94	8	198–201	199–201 (59)
14	<b>4n</b>	4-(OH)–C <sub>6</sub> H <sub>4</sub> –	95	8	229–231	231–233 (57)
15	<b>4o</b>	2-(OCH <sub>3</sub> )–C <sub>6</sub> H <sub>4</sub> –	93	8	258–260	259–260 (53)
16	<b>4p</b>	3-(OCH <sub>3</sub> )–C <sub>6</sub> H <sub>4</sub> –	91	8	208–210	207–208 (52)
17	<b>4q</b>	4-(OCH <sub>3</sub> )–C <sub>6</sub> H <sub>4</sub> –	95	8	203–205	203–204 (57)

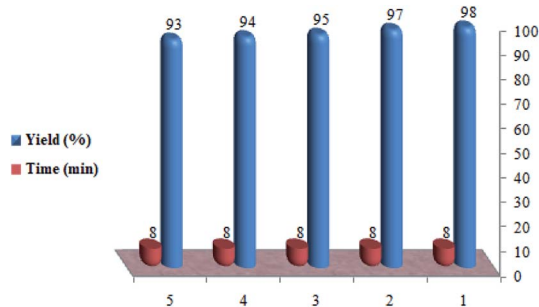
<sup>a</sup> Reaction conditions: ethyl acetoacetate (**1**), aldehyde (**2**), urea (**3**) (molar ratio 1 : 1 : 1.2) and  $\text{SO}_3\text{H@imineZCMNPs}$  (10 mg), 90  $^\circ\text{C}$ . <sup>b</sup> Isolated yield.

<sup>c</sup> Melting points are uncorrected.





**Scheme 3** A plausible mechanism for the one-pot three-component reaction of ethyl acetoacetate (**1**), aldehyde (**2**), urea (**3**) catalyzed by the  $\text{SO}_3\text{H@imineZCMNPs}$  under solvent-free conditions.



**Fig. 7** Reusability of the catalyst.

yield of the product, while higher catalyst loading did not considerably affect the duration of the reaction or the product yield (entries 9 and 10). To explore the effect of temperature with the same amount of catalyst (10 mg) under solvent-free conditions, different temperatures (25, 80, 90, and 100 °C) were used to compare the reaction efficiency (entries 8 and 11–13). It was observed that room temperature was not suitable for the formation of the desired product **4g** under solvent-free conditions (entry 11). It is noteworthy that the use of heat

with the same conditions leads to a faster reaction and a higher yield (entries 8, 12 and 13). Increasing the temperature beyond this point (90 °C) led to no substantial improvement in the yield (entry 13).

Using the optimized reaction conditions (Table 2, entry 8) in order to generalize the effectiveness and acceptability of the strategy, a variety of aryl aldehydes containing electron-withdrawing as well as electron-donating groups in the *ortho*-, *meta*-, and *para*- positions of the benzene ring (**2a–q**) were selected for the reaction with ethyl acetoacetate (**1**) and urea (**3**), furnishing the desired products with high to excellent isolated yields (Table 3, entries 1–17). It is observed that with electron-withdrawing substituents, a relatively higher reaction yield of the products was achieved compared to electron-donating substituents.

A reasonable pathway for the synthesis of 3,4-dihydropyrimidin-2(1*H*)-one derivatives is presented in Scheme 3. Initially, aldehyde **2** and urea **3** as the reactant components react with each other by nucleophilic addition. After forming the acylimine intermediate **5**, it creates an open-chain ureide **6** after a dehydration step. The intermediate **6** may further undergo nucleophilic addition with ethyl acetoacetate **1** to give the intermediate **7**. The final product **4** was then formed by the intramolecular cyclization and dehydration of the intermediate **7**.

In the next phase of the study, the feasibility of the recovery and reuse cycle of the catalyst was checked by performing the reaction of ethyl acetoacetate, 4-chlorobenzaldehyde, and urea in the presence of  $\text{SO}_3\text{H@imineZCMNPs}$  under solvent-free conditions. After reaction completion, as monitored by TLC, the catalyst was separated from the reaction mixture using a permanent magnet, and the mixture was washed with ethanol ( $2 \times 15$  mL). As shown in Fig. 7,  $\text{SO}_3\text{H@imineZCMNPs}$  can be recycled five times with a negligible change in terms of the reaction time and with a yield range of 98 to 93%.

Furthermore, to investigate and compare the presented method for the synthesis of 3,4-dihydro-5-etoxy carbonyl-4-(4-phenyl)-6-methylpyrimidine-2(1*H*)-one (**4a**), a comparison was made with some of the other earlier reported homogeneous and heterogeneous catalysts. The obtained results are shown in Table 4. The collected data demonstrate our reported protocol in terms of using solvent-free conditions, product yield, and reaction time.

**Table 4** Comparison of the results of the synthesis of compound **4a** using different catalysts

Entry	Catalyst and conditions	Amount of catalyst	Reaction time	Yield (%)	Ref.
1	$\text{BF}_3 \cdot \text{OEt}_2/\text{CuCl}/\text{THF}/\text{reflux}$	10 mol%	18 h	94	52
2	$\text{PPA-SiO}_2/\text{CH}_3\text{CN}/\text{reflux}$	0.3 g	30 min	85	55
3	Zeolite/toluene/reflux	15 mol%	12 h	80	65
4	$\text{CD-SO}_3\text{H}/\text{solvent-free}/100^\circ\text{C}$	0.04 g	2 h	83	54
5	$\text{FeCl}_3 \cdot 6\text{H}_2\text{O}/\text{solvent-free}/\text{reflux}$	0.16 g	4 h	94	56
6	Silica sulfuric acid/EtOH/heat	0.23 g	6 h	91	53
7	$\text{SnCl}_2\text{-nano SiO}_2/\text{EtOH}/\text{reflux}$	0.45 mol%	40 min	92	64
8	Sulfated tungstate/solvent-free/80 °C	10 mol%	1 h	92	40
9	$\text{SO}_3\text{H@imineZCMNPs}/\text{solvent-free}/90^\circ\text{C}$	10 mg	8 min	93	This work



## Conclusion

In conclusion, we described a simple, efficient, and green methodology for the one-pot preparation of a wide range of biologically and pharmacologically important 3,4-dihydropyrimidin-2(1H)-one derivatives in the presence of an eco-friendly and reusable nanocatalyst (SO<sub>3</sub>H@imineZCMNPs). The most important advantages of this procedure are green reaction conditions, low catalyst loading, high quantitative yields, low cost, the absence of solvents, and facile isolation and reusability of the catalyst.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 V. Poshettiwar and R. S. Varma, *Chem. Soc. Rev.*, 2008, **37**, 1546.
- 2 P. T. Anastas and J. C. Warner, *Green Chemistry Theory and Practice*, 1998.
- 3 A. S. Matlack, *Introduction to Green Chemistry*, 2001.
- 4 J. H. Clark and D. J. Macquarrie, *Handbook of Green Chemistry and Technology*, 2002.
- 5 M. Lancaster, *Green Chemistry: An Introductory Text*, RSC Editions, 2002.
- 6 M. Poliakov, J. M. Fitzpatrick, T. R. Farren and P. T. Anastas, *Science*, 2002, **297**, 807.
- 7 M. Doble and A. K. Kruthiventi, *Green Chemistry and Engineering*, Elsevier Science & Technology Books, 2007.
- 8 J. A. Loch and R. H. Crabtree, *Pure Appl. Chem.*, 2001, **73**, 119.
- 9 R. A. Sheldon, I. Arends and U. Hanefeld, *Green Chemistry and Catalysis*, 2007.
- 10 R. T. Baker and W. Tumas, *Science*, 1999, **284**, 1477.
- 11 F. Hoffmann, M. Cornelious, J. Moreli and M. Froba, *Angew. Chem.*, 2006, 3216.
- 12 M. Jaroniec, T. W. Kim, M. Kruk and R. Ryoo, *Chem. Mater.*, 2003, **15**, 2815.
- 13 H. Yang and D. Zhao, *J. Mater. Chem.*, 2005, **15**, 1217.
- 14 H. U. Blaser, A. Indolese and A. Schnyder, *Curr. Sci.*, 2000, **78**, 1336.
- 15 Y. Chauvin and J. C. Vedrine, *Actual. Chim.*, 1996, **58**, 7.
- 16 J. P. Collman, L. S. Hegedus, M. P. Cooke, J. R. Norton, G. Doleeti and D. N. Marquardt, *J. Am. Chem. Soc.*, 1972, **94**, 1789.
- 17 W. Dumont, J. C. Poulin, T. P. Daud and H. B. Kagan, *J. Am. Chem. Soc.*, 1973, **95**, 8295.
- 18 L. Yin and J. Liebscher, *Chem. Rev.*, 2007, **107**, 133.
- 19 H. U. Blaser, *Tetrahedron: Asymmetry*, 1991, **9**, 843.
- 20 P. McMorn and G. J. Hutchings, *Chem. Soc. Rev.*, 2004, **33**, 108.
- 21 M. Heitbaum, F. Glorius and I. Escher, *Angew. Chem.*, 2006, **45**, 4732.
- 22 D. Astruc, *Nanoparticles and Catalysis*, Wiley-VCH, Weinheim, 2008.
- 23 G. A. Somorjia, H. Frei and J. Y. Park, *J. Am. Chem. Soc.*, 2009, **131**, 16589.
- 24 S. Shylesh, L. Wang and W. R. Thiel, *Adv. Synth. Catal.*, 2010, **352**, 425.
- 25 A. J. Amali and R. K. Rana, *Green Chem.*, 2011, **11**, 1781.
- 26 J. Hu, Y. Wang, M. Han, Y. Zhou, X. Jiang and P. Sun, *Catal. Sci. Technol.*, 2011, **2**, 2332.
- 27 Q. Zhang, H. Su, J. Luo and Y. Wei, *Catal. Sci. Technol.*, 2013, **3**, 235.
- 28 A. H. Lu, E. L. Salabas and F. Schüth, *Angew. Chem., Int. Ed.*, 2007, **46**, 1222.
- 29 S. Shylesh, V. Schünemann and W. R. Thiel, *Angew. Chem., Int. Ed.*, 2010, **49**, 3428.
- 30 Y. Zhu, L. P. Stubbs, F. Ho, R. Liu, C. P. Ship, J. A. Maguire and N. S. Hosmane, *ChemCatChem*, 2010, **2**, 365.
- 31 V. Polshettiwar, R. Luque, A. Fihri, H. Zhu, M. Bouhrara and J. M. Basset, *Chem. Rev.*, 2011, **111**, 3036.
- 32 R. B. N. Baig and R. S. Varma, *Chem. Commun.*, 2013, **49**, 752.
- 33 S. S. Kim, B. S. Choi, J. H. Lee, K. K. Lee, T. H. Lee, Y. H. Kim and S. Hyunik, *Synlett*, 2009, 599.
- 34 C. O. Kappe, *Eur. J. Med. Chem.*, 2000, **35**, 1043.
- 35 P. G. Biginelli, *Gazz. Chim. Ital.*, 1893, **23**, 360.
- 36 K. S. Atwal, B. N. S. Wanson, S. E. Unger, D. M. Floyd, S. Mereland, A. Hedberg and B. C. J. O'Reilly, *J. Med. Chem.*, 1991, **34**, 806.
- 37 Y. Huang, F. Yang and C. Zhu, *J. Am. Chem. Soc.*, 2005, **127**, 16386.
- 38 M. A. Bigdeli, G. Gholami and E. Sheikhsosseini, *Chin. Chem. Lett.*, 2011, **22**, 903.
- 39 X. H. Chen, X. Y. Xu, H. Liu, L. F. Cun and L. Z. Gong, *J. Am. Chem. Soc.*, 2006, **128**, 14802.
- 40 S. D. Salim and K. G. Akamanchi, *Catal. Commun.*, 2011, **12**, 1153.
- 41 S. Rostamnia and A. Morsali, *RSC Adv.*, 2014, **4**, 10514.
- 42 K. A. Kumar, M. Kasthuraiah, C. S. Reddy and C. D. Reddy, *Tetrahedron Lett.*, 2001, **42**, 7873.
- 43 Y. Qiu, H. Sun, Z. Ma and W. Xia, *J. Mol. Catal. A: Chem.*, 2014, **392**, 76.
- 44 C. O. Kappe, D. Kumar and R. S. Varma, *Synthesis*, 1999, **10**, 1799.
- 45 M. Yamaura, R. L. Camilo, L. C. Sampaio, M. A. Macedo, M. Nakamura and H. E. Toma, *J. Magn. Magn. Mater.*, 2004, **279**, 210.
- 46 Y. W. Wu, J. Zhang, J. F. Liu, L. Chen, Z. L. Deng, M. X. Han, X. S. Wei, A. M. Yu and H. L. Zhang, *Appl. Surf. Sci.*, 2012, **258**, 6772.
- 47 E. Abbaspour-Gilandeh, M. Aghaei-Hashjin, A. Yahyazadeh and H. Salemi, *RSC Adv.*, 2016, **6**, 55444.
- 48 A. Yahyazadeh, E. Abbaspour-Gilandeh and M. Aghaei-Hashjin, *Catal. Lett.*, 2018, **148**, 1254.
- 49 M. Sheykhan, A. Yahyazadeh and Z. Rahemizadeh, *RSC Adv.*, 2016, **6**, 34553.
- 50 J. Zhang and N. Yan, *ChemCatChem*, 2017, **9**, 2790.



- 51 U. B. More, *Asian J. Chem.*, 2012, **24**, 1906.
- 52 E. H. Hu, D. R. Sidler and U. H. Dolling, *J. Org. Chem.*, 1998, **63**, 3454.
- 53 P. Salehi, M. Dabiri, M. A. Zolfigol and M. A. B. Fard, *Tetrahedron Lett.*, 2004, **44**, 2889.
- 54 S. Asghari, M. Tajbakhsh, B. Jafarzadeh-Kenari and S. Khaksar, *Chin. Chem. Lett.*, 2011, **22**, 127.
- 55 M. Zeinali-Dastmalbaf, A. Davoodnia, M. M. Heravi, N. Tavakoli-Hoseini, A. Khojastehnezhad and H. A. Zamani, *Bull. Korean Chem. Soc.*, 2011, **32**, 656.
- 56 J. Lu and H. Ma, *Synlett*, 2000, 63.
- 57 R. Ghosh, S. Maiti and A. Chakraborty, *J. Mol. Catal. A: Chem.*, 2004, **217**, 47.
- 58 Y. Ma, C. Qian, L. Wang and M. Yang, *J. Org. Chem.*, 2000, **65**, 3864.
- 59 D. Russowsky, F. A. Lopes, V. S. S. Silva, K. F. S. Canto, M. G. Montes and M. N. Godoi, *J. Braz. Chem. Soc.*, 2004, 165.
- 60 A. D. Patil, N. V. Kumar, W. C. Kokke, M. F. Bean, A. J. Freyer, C. D. Brossi, S. Mai, A. Truneh, D. J. Faulkner, B. Carte, A. L. Breen, R. P. Hertzberg, R. K. Johnson, J. W. Westly and B. C. M. Potts, *J. Org. Chem.*, 1995, **60**, 1182.
- 61 N. S. Nandurkar, M. J. Bhanushali, M. D. Bhor and B. M. Bhanage, *J. Mol. Catal. A: Chem.*, 2007, **271**, 14.
- 62 J. S. Yadav, B. V. S. Reddy, R. Srinivas, C. V. enugopal and T. Ramalingam, *Synthesis*, 2001, 1341.
- 63 K. K. Pasunooti, H. Chai, C. N. Jensen, B. K. Gorityala, S. Wang and X. W. Liu, *Tetrahedron Lett.*, 2011, **52**, 80.
- 64 J. Safaei-Ghomi, R. Teymuri and A. Ziarati, *Monatsh. Chem.*, 2013, **144**, 1865.
- 65 V. Radha Rani, N. Srinivas, M. Radha Kishan, S. J. Kulkarni and K. V. Raghavan, *Green Chem.*, 2001, **3**, 305.

