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# Introduction of taurine (2-aminoethanesulfonic acid) as a green bio-organic catalyst for the promotion of organic reactions under green conditions<sup>†</sup>

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Taurine (2-aminoethanesulfonic acid), a semi-essential amino acid that exists in the human body and numerous other living creatures, is used as a green bio-organic catalyst for the promotion of the Knoevenagel reaction between aldehydes and malononitrile. In the same way, tetraketones can also be produced through a Knoevenagel reaction, followed by Michael addition. 2-Amino-3-cyano-4H-pyran derivatives are simply prepared via a three-component reaction in the presence of taurine as the catalyst. All these reactions are performed in water, a green solvent. The advantages of using of taurine as the catalyst are it is environmentally friendly, low cost, commercially available, easy to separate from the reaction mixture, and has high reusability. Use of this catalyst results in acceptable reaction times, high yields and high purities of the obtained products without utilizing any organic solvents.

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## Introduction

The progress of science has been more and more towards environmentally compatible, or “green” processes, with particular focus on catalysts and other materials in organic chemistry. One aspect of these topics is the application of an alternative reaction medium that is free from the problems associated with the numerous traditional volatile solvents. Using this type of media may also increase the chance of separation and reuse of the catalyst. From the viewpoint of green chemistry, it is better to perform the reactions under solvent-free conditions, but when solvent is necessary, water is the best choice.<sup>1</sup> In addition to environmental concerns, chemists prefer to use water as a solvent due to its economic benefits and generally easy separation and work-up conditions. Furthermore, all biological reactions are known to occur in aqueous media. Moreover, numerous organic reactions proceed faster and better in water than in organic solvents.<sup>2</sup>

2-Aminoethane sulfonic acid, or taurine (Fig. 1), is an amino acid that is found in high concentration in the tissues of animals. It is one of the constituent members of bile, which can be found in the large intestine and its amount in the average is one-tenth percent of the total weight of the

human body.<sup>3</sup> Taurine appellation refers to its first isolation from ox bile, named *Bos taurus*.<sup>4</sup> The concentration of taurine in mammalian organs is higher in comparison with the other types of the animals; its concentration in insects and arthropods is less than that in mammals, whereas in plants and bacteria, its concentration is negligible.<sup>5</sup> Red algae, although not the brown or green ones, contain high levels of taurine and its *N*-(1-carboxylated) derivatives, but lichens, mushrooms, mosses, and ferns have very low concentrations of this amino acid.<sup>6</sup>

In comparison with other homologue amino acids, taurine is structurally different. It is a  $\beta$ -amino acid with a sulfonic acid group instead of a carboxylic acid group. This difference increases its acidity, *i.e.*, in the range of mineral acids ( $pK_a = 1.5$ ), compared to carboxylic acid homologues, and unlike those homologues that are not dissociated at biological pH, taurine is in a zwitterionic state at this pH level, which leads to distinct biological properties.<sup>7</sup> According to computational investigations, the neutral conformation of taurine exists in the gas phase, whereas its zwitterionic form exists in water media, which is in agreement with the experimental NMR analysis.<sup>8</sup>

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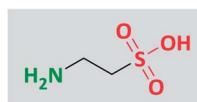


Fig. 1 Taurine.

Taurine has been used for many years as an ingredient in energy drinks and nutrient supplements and has many biological properties such as osmoregulation, immunomodulation and bile salt formation.<sup>9</sup>

Very recently, silica gel supported taurine was used in the oxidation of sulfides to their corresponding disulfides.<sup>10</sup> Moreover, to the best of our knowledge, there are no other reports of the catalytic activity of this  $\beta$ -amino acid for organic transformations.

In recent decades, performance of standard chemical reactions in aqueous media has been often considered, with particular focus on carbon–carbon bond forming transformations.<sup>11</sup> Several reactions that have been investigated in this capacity are Diels–Alder, Claisen and Aldol condensations, and radical additions.<sup>12</sup>

The Knoevenagel reaction, *i.e.*, treatment of an aldehyde with an active methylene group reported by Emil Knoevenagel in 1894, is one of the most important and notable reactions for C=C bond formation.<sup>13</sup> This reaction is suitable for the preparation of alkenes with electron-withdrawing groups and its products can be used as intermediates for many other types of reactions.<sup>14</sup>

Many conditions have been used to promote the Knoevenagel reaction between aldehydes and malononitrile, including grinding, high pressure, microwaves, and ultrasonics.<sup>15</sup> Different types of catalysts have also been utilized; among the most important are cetyltrimethylammonium bromide (CTMAB),<sup>16</sup>  $\text{ReBr}(\text{CO})_5$ ,<sup>17</sup> 1,1,3,3-tetramethylguanidium lactate,<sup>18</sup>  $[\text{C}_4\text{dabco}][\text{BF}_4]$ ,<sup>19</sup> amine-functionalized polyacrylonitrile fiber,<sup>20</sup>  $\text{Fe}_3\text{O}_4$  MNPs–guanidine,<sup>21</sup> sulfonated carbon/silica composites,<sup>22</sup> MP(DNP),<sup>23</sup>  $\text{Na}_2\text{S}/\text{Al}_2\text{O}_3$ ,<sup>24</sup> silica–L-proline,<sup>25</sup>  $\text{ZnO}$ ,<sup>26</sup>  $\text{Ni}_x\text{Mg}_{1-x}\text{Fe}_2\text{O}_4$ ,<sup>27</sup> sodium carbonate,<sup>28</sup> hydroxyapatite supported caesium carbonate,<sup>29</sup> *Tamarindus indica* juice,<sup>30</sup> L-proline–IL,<sup>31</sup> and  $\text{SiO}_2\text{–NH}_4\text{OAc}$ .<sup>32</sup>

Arylmethylene[bis(3-hydroxy-2-cyclohexene-1-ones)] (tetra-ketones) firstly were introduced by Merling through the addition of aldehydes with 1,3-diketones *via* a Knoevenagel reaction followed by a Michael addition.<sup>33</sup> These compounds have been extensively used as intermediates for some other important target compounds such as acridine diones, thianthrenes and xanthene diones.<sup>34</sup> Tetraketones have biological activity as antioxidants, lipoxygenases and tyrosinase inhibitors.<sup>35</sup> Because of the important properties of these products, several methods have been used to achieve them using various catalysts such as  $\text{SnCl}_2/\text{HCl}$ ,<sup>36</sup>  $\text{In}(\text{OTf})_3$ ,<sup>37</sup>  $\text{SmCl}_3$ ,<sup>38</sup>  $\text{Yb}(\text{OTf})_3\text{–SiO}_2$  with aniline,<sup>39</sup>  $\text{Fe}_3\text{O}_4@\text{SiO}_2\text{–SO}_3\text{H}$ ,<sup>40</sup> PVP-stabilized Ni nanoparticles,<sup>41</sup> choline chloride-based deep eutectic,<sup>42</sup> nano Fe/NaY zeolite,<sup>43</sup> EDDA,<sup>44</sup> and Al/MCM-41.<sup>45</sup>

4H-Pyran and its derivatives have attracted much interests because of their important biological activities, such as anticoagulant, spasmolytic, diuretic, anticancer, antianaphylactin,<sup>46</sup> antiallergenic,<sup>47</sup> antiproliferative,<sup>48</sup> antitumor,<sup>49</sup> antibacterial,<sup>50</sup> cytotoxic,<sup>51</sup> mutagenic<sup>52</sup> and sex pheromonal<sup>53</sup> activities. These compounds also are present in the structure of some photoactive materials<sup>54</sup> and natural products<sup>55</sup> and can be used in the synthesis of cosmetics and pigments.<sup>56</sup>

A large number of catalysts were introduced for the synthesis of 2-amino-3-cyano-4H-pyran derivatives under various conditions; notable among them are hexadecyldimethyl benzyl ammonium bromide,<sup>57</sup> 1-butyl-3-methyl imidazolium hydroxide,<sup>58</sup> 2,2,2-trifluoroethanol,<sup>59</sup>  $\text{Ba}(\text{OTf})_2$ ,<sup>60</sup> tetrabutylammonium chloride,<sup>61</sup> triazine functionalized ordered mesoporous organosilica,<sup>62</sup> tungstic acid functionalized mesoporous SBA-15,<sup>63</sup> *p*-dodecylbenzenesulfonic acid,<sup>64</sup> potassium phthalimide,<sup>65</sup> red sea sand,<sup>66</sup> choline hydroxide,<sup>67</sup> IRMOF-3( $\text{Zn}_4\text{O}(\text{H}_2\text{N-TA})_3$ ),<sup>68</sup>  $\beta$ -cyclodextrins–glycerine,<sup>69</sup> tris-hydroxymethylaminomethane,<sup>70</sup> squaramide,<sup>71</sup>  $\text{Ce}(\text{SO}_4)_2\cdot 4\text{H}_2\text{O}$ ,<sup>72</sup> poly(vinylpyrrolidonium)hydrogen phosphate,<sup>73</sup> L-proline,<sup>74</sup>  $\text{I}_2$ ,<sup>75</sup> and glutamic acid.<sup>76</sup>

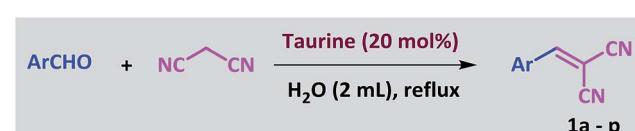
## Experimental

All the chemicals for this study were purchased from Merck, Aldrich, and Fluka Chemical Companies and used without further purification. All the products were separated and characterized by their physical properties in comparison with the reported standards. Both the purity determination of the substrates and reaction monitoring were accomplished by thin layer chromatography (TLC) using SIL G/UV 254 silica gel plates. Melting points were determined using a Buchi B-545 apparatus. FT-IR spectra were obtained by a Perkin-Elmer spectrum BX series spectrophotometer (KBr disks). The  $^1\text{H}$  NMR and  $^{13}\text{C}$  NMR spectra were acquired by a Bruker Avance 400 MHz instrument using deuterated solvents.

Table 1 Optimization of the conditions for the Knoevenagel reaction [1g]<sup>a</sup>

Entry	Catalyst (mol%)	Solvent	Temp.	Time (min)	Yield <sup>b</sup> (%)
1	—	—	90 °C	90	Trace <sup>c</sup>
2	24	—	90 °C	90	Trace <sup>c</sup>
3	24	$\text{CHCl}_3$	Reflux	90	Trace <sup>c</sup>
4	12	$\text{EtOH}$	Reflux	90	Trace <sup>c</sup>
5	24	$\text{EtOH}$	Reflux	90	Trace <sup>c</sup>
6	24	$\text{CH}_3\text{CN}$	Reflux	90	Trace <sup>c</sup>
7	12	$\text{H}_2\text{O}$	RT	90	Trace <sup>c</sup>
8	24	$\text{H}_2\text{O}$	RT	90	87
9	16	$\text{H}_2\text{O}$	Reflux	13	91
10	20	$\text{H}_2\text{O}$	Reflux	7	98
11	24	$\text{H}_2\text{O}$	Reflux	6	95
12	24	$\text{H}_2\text{O}/\text{EtOH}$ (3 : 1)	Reflux	20	86

<sup>a</sup> Reaction conditions: 4-chlorobenzaldehyde (1.0 mmol), malononitrile (1.1 mmol), solvent (2 mL) and required amount of catalyst. <sup>b</sup> The yields are related to the isolated products. <sup>c</sup> The reaction was not completed.



Scheme 1 The Knoevenagel reaction between aldehydes and malononitrile.

Table 2 The Knoevenagel reaction of aldehydes and malononitrile in the presence of taurine as the catalyst in water medium

Entry	Aldehyde	Product	Symbol	Time (min)	Yield <sup>a</sup> (%)	Mp (°C)	
						(Obs.)	(Lit.)
1			1a	14	86	82–83	82–84 (ref. 16)
2			1b	11	94	92–93	93–94 (ref. 27)
3			1c	9	96	138–140	139–140 (ref. 21)
4			1d	18	90	81–82	80 (ref. 24)
5			1e	15	87	99–101	98–99 (ref. 29)
6			1f	10	93	103–105	103–105 (ref. 27)
7			1g	7	98	161–163	161–162 (ref. 27)
8			1h	9	96	156–158	159–160 (ref. 21)
9			1i	12	92	122–124	122–124 (ref. 28)
10			1j	15	94	161–163	157–160 (ref. 22)
11			1k	20	89	115–117	112–114 (ref. 22)
12			1l	14	91	186–188	185–187 (ref. 27)
13			1m	10	93	136–137	137–138 (ref. 24)
14			1n	8	94	295–297	298–300 (ref. 30)
15			1o	10	86	125–127	124–126 (ref. 22)

Table 2 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
16			1p	6	97	155–157 New product

<sup>a</sup> The yields are related to the isolated products.

### General procedure for the Knoevenagel condensation of aldehydes and malononitrile

In a 25 mL round-bottomed flask, a mixture of aldehyde (1.0 mmol), malononitrile (1.1 mmol) and taurine (0.025 g, 20 mol%) in water (2 mL) was heated at a reflux temperature for the appropriate time. After the conversion, which was monitored by TLC, 10 mL of water was added and stirred for 3 minutes. During this time, the product was precipitated and subsequently separated by filtration. The separated product was washed several times with water. After drying, the pure product was obtained; there was no need for further purification and addition of organic solvent was not necessary. Furthermore, water was evaporated from the filtrate to re-obtain the taurine catalyst.

### General procedure for the synthesis of tetraketones

In a 25 mL round-bottomed flask, a mixture of aldehyde (1.0 mmol), 1,3-cyclodicarbonyl compound (2.0 mmol) and taurine (0.030 g, 24 mol%) in water (2 mL) was heated at reflux for the

Table 3 Optimization of the conditions for the synthesis of tetraketones [2g]<sup>a</sup>

Entry	Catalyst (mol%)	Solvent	Temp.	Time (min)	Yield <sup>b</sup> (%)
1	—	—	90 °C	120	Trace <sup>c</sup>
2	24	—	60–90 °C	120	Mixture <sup>c</sup>
3	24	CHCl <sub>3</sub>	Reflux	120	Trace <sup>c</sup>
4	12	EtOH	Reflux	120	Trace <sup>c</sup>
5	24	EtOH	Reflux	120	30 <sup>c</sup>
6	24	CH <sub>3</sub> CN	Reflux	120	Trace <sup>c</sup>
7	12	H <sub>2</sub> O	RT	120	20 <sup>c</sup>
8	24	H <sub>2</sub> O	RT	120	35 <sup>c</sup>
9	16	H <sub>2</sub> O	Reflux	33	91
10	24	H <sub>2</sub> O	Reflux	20	96
11	28	H <sub>2</sub> O	Reflux	17	92
12	20	H <sub>2</sub> O/EtOH (3 : 1)	Reflux	30	83

<sup>a</sup> Reaction conditions: 4-chlorobenzaldehyde (1.0 mmol), dimedone (2.0 mmol), solvent (2 mL) and required amount of the catalyst. <sup>b</sup> The yields are related to the isolated products. <sup>c</sup> The reaction was not completed.

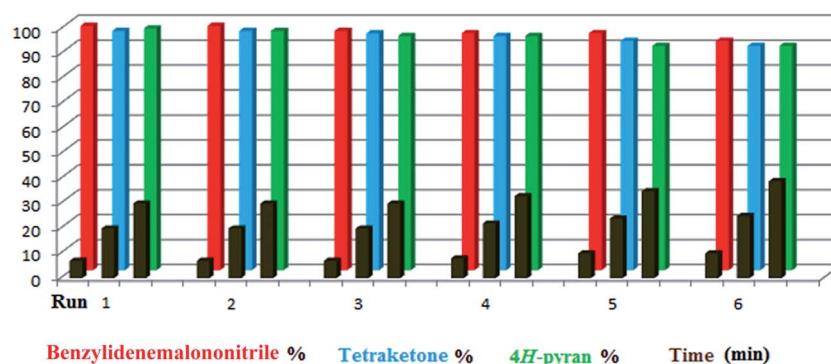
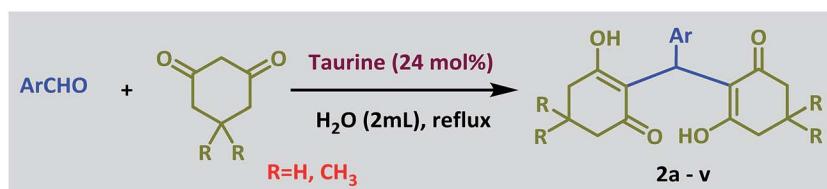


Fig. 2 Reusability of taurine in the model reactions.



Scheme 2 Synthesis of tetraketones.

Table 4 Synthesis of tetraketones in the presence of taurine catalyst in water

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
1			2a	10	95	181–183 185 (ref. 42)
2			2b	15	93	194–196 199–200 (ref. 43)
3			2c	10	92	242–243 244–246 (ref. 38)
4			2d	25	87	181–183 184–186 (ref. 43)
5			2e	20	91	209–211 208–209 (ref. 40)
6			2f	25	94	195–197 197–199 (ref. 40)
7			2g	20	97	139–141 138–141 (ref. 42)

Table 4 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
8			2h	15	96	159–161 154–156 (ref. 43)
9			2i	15	90	183–185 184–186 (ref. 43)
10			2j	45	91	185–187 188–190 (ref. 43)
11			2k	15	90	142–144 146–148 (ref. 38)
12			2l	20	95	182–183 180 (ref. 40)
13			2m	25	85	133–135 132 (ref. 41)
14			2n	25	85	220–221 225 (ref. 42)

Table 4 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
15			2o	15	92	151–153 146 (ref. 41)
16			2p	30	92	215–218 213–215 (ref. 38)
17			2q	45	93	174–176 New product
18			2r	15	89	219–220 214–216 (ref. 43)
19			2s	20	93	203–205 199–200 (ref. 42)
20			2t	30	96	207–209 209–211 (ref. 36)
21			2u	25	91	199–200 201–203 (ref. 43)

Table 4 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
22			2v	30	90	189–191 190–191 (ref. 39)

<sup>a</sup> The yields are related to the isolated products.

appropriate time. After completion of the reaction, which was monitored by TLC, 10 mL of water was added and stirred for 3 minutes. During this time, the product was precipitated and subsequently separated by filtration. The separated product was washed several times with water. After drying, the pure product was obtained; there was no need for further purification and addition of organic solvent was not necessary. Furthermore, water was evaporated from the filtrate to re-obtain the taurine catalyst.

Table 5 Optimization of the conditions for the synthesis of 2-amino-3-cyano-4H-pyran derivatives [3i]<sup>a</sup>

Entry	Catalyst (mol%)	Solvent	Temp.	Time (min)	Yield <sup>b</sup> (%)
1	—	—	90 °C	120	Trace <sup>c</sup>
2	24	—	60–90 °C	120	Trace <sup>c</sup>
3	24	CHCl <sub>3</sub>	Reflux	120	Trace <sup>c</sup>
4	12	EtOH	Reflux	120	Trace <sup>c</sup>
5	24	EtOH	Reflux	120	Trace <sup>c</sup>
6	24	CH <sub>3</sub> CN	Reflux	120	Trace <sup>c</sup>
7	12	H <sub>2</sub> O	RT	115	90
8	24	H <sub>2</sub> O	RT	90	91
9	16	H <sub>2</sub> O	Reflux	65	90
10	28	H <sub>2</sub> O	Reflux	30	97
11	32	H <sub>2</sub> O	Reflux	26	94
12	28	H <sub>2</sub> O/EtOH (3 : 1)	Reflux	45	88

<sup>a</sup> Reaction conditions: 4-chlorobenzaldehyde (1.0 mmol), malononitrile (1.1 mmol) and dimedone (1.0 mmol), solvent (2 mL) and required amount of the catalyst. <sup>b</sup> The yields are related to the isolated products. <sup>c</sup> The reaction was not completed.

### General procedure for the synthesis of 2-amino-4H-chromenes

In a 25 mL round-bottomed flask, a mixture of aldehyde (1.0 mmol), 1,3-cyclodiacarbonyl compound (1.0 mmol), malononitrile, (1.1 mmol) and taurine (0.030 g, 28 mol%) in water (2 mL) was heated at reflux for the appropriate time. After the reaction was completed, which was monitored by TLC, 10 mL of water was added and stirred for 3 minutes. During this time, the product was precipitated and subsequently separated by filtration. The separated product was washed several times with water. After drying, the pure product was obtained; there was no need for further purification and addition of organic solvent was not necessary. Furthermore, water was evaporated from the filtrate to re-obtain the taurine catalyst.

### Spectroscopic data of the new compounds

**2-(4-(Methylthio)benzylidene)malononitrile (1p).** IR (KBr, cm<sup>-1</sup>): 3040, 2217, 1648, 1564, 1094; <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  = 2.58 (s, 3H), 7.48 (d, *J* = 8.4 Hz, 2H), 7.89 (d, *J* = 8.4 Hz, 2H), 8.42 (s, 1H) ppm; <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  = 14.30, 79.10, 114.14, 115.10, 125.87, 127.75, 131.41, 148.92, 160.86 ppm.

**2,2'-(3-(2-Nitrophenyl)prop-2-ene-1,1-diy)bis(3-hydroxy-5,5-dimethylcyclohex-2-en-1-one) (2q).** IR (KBr, cm<sup>-1</sup>): 3445, 3070, 2960, 2868, 1600, 1590, 1519, 1380. <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  = 1.10 (s, 6H), 1.17 (s, 6H), 2.30–244 (m, 8H), 4.98 (m, 1H), 6.39 (dd, *J* = 16.0, 4.0 Hz, 1H), 6.89 (dd, *J* = 16.0, 2.4 Hz, 1H), 7.35 (dt, *J* = 8.0, 1.6 Hz, 1H), 7.56 (dt, *J* = 8.0, 1.2 Hz, 1H), 7.94 (dd, *J* =



Scheme 3 Synthesis of 2-amino-3-cyano-4H-pyran derivatives.

Table 6 Synthesis of 2-amino-3-cyano-4*H*-pyran derivatives in the presence of taurine as the catalyst in water

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
1			3a	65	82	231–233 232–234 (ref. 73)
2			3b	25	92	209–211 211–213 (ref. 73)
3			3c	15	94	222–225 228 (ref. 69)
4			3d	70	81	204–206 196–199 (ref. 73)
5			3e	45	98	211–213 211–213 (ref. 72)
6			3f	30	96	224–226 228–230 (ref. 72)
7			3g	25	93	287–289 289–291 (ref. 72)
8			3h	35	92	190–192 188–190 (ref. 72)

Table 6 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
9			3i	30	97	208–210 208–210 (ref. 73)
10			3j	30	95	203–205 201–203 (ref. 73)
11			3k	25	97	210–212 214–215 (ref. 73)
12			3l	30	96	180–182 181–183 (ref. 73)
13			3m	65	91	193–195 196–198 (ref. 70)
14			3n	45	87	217–219 217–219 (ref. 70)
15			3o	50	94	220–222 222–224 (ref. 70)

Table 6 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.) (Lit.)
16			3p	35	93	266–269 270–275 (ref. 77)
17			3q	30	90	223–226 223–225 (ref. 73)
18			3r	60	89	211–212 208–209 (ref. 58)
19			3s	50	93	212–214 212–213 (ref. 79)
20			3t	20	91	209–210 210–212 (ref. 57)
21			3u	50	94	202–204 202–204 (ref. 57)
22			3v	40	93	221–223 224–226 (ref. 57)

Table 6 (Contd.)

Entry	Aldehyde	Product	Symbol	Time (min)	Mp (°C)	
					Yield <sup>a</sup> (%)	(Obs.)
23			3w	35	98	237–240 234–235 (ref. 63)
24			3x	45	85	202–204 206–208 (ref. 63)
25			3y	50	92	217–219 New product

<sup>a</sup> The yields are related to the isolated products.

8.0, 1.2 Hz, 1H), 11.27 (br, 1H), 12.12 (s, 1H) ppm; <sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>):  $\delta$  = 20.86, 29.73, 31.42, 31.44, 46.20, 46.83, 115.96, 124.53, 125.37, 127.72, 128.80, 133.12, 133.16, 134.64, 147.34, 189.43, 190.05 ppm.

**2-Amino-4-(4-(methylthio)phenyl)-5-oxo-5,6,7,8-tetrahydro-4H-chromene-3-carbonitrile (3y).** IR (KBr, cm<sup>-1</sup>): 3318, 3171, 2961, 2913, 2194, 1682, 1647, 1364; <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  = 1.92–2.00 (m, 2H), 2.24–2.34 (m, 2H), 2.35 (s, 1H), 2.61–2.63 (m, 2H), 4.16 (s, 1H), 7.08 (s, 2H), 7.02 (d, *J* = 8.4 Hz, 2H), 7.10 (d, *J* = 8.4 Hz, 2H) ppm; <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  = 15.29, 20.29, 26.95, 35.46, 36.81, 58.50, 114.17, 120.23, 126.53, 128.31, 136.53, 142.09, 158.88, 158.92, 164.87, 196.35 ppm.

## Results and discussion

In recent years, introduction of sulfonic acid based catalysts for the promotion of organic transformations has been an important part of our ongoing research program.<sup>77,78</sup> In continuation of these studies, we were interested in investigating the applicability of taurine, a natural, green and commercially available amino acid containing a sulfonic acid group, in the acceleration of organic reactions. The Knoevenagel reaction was selected as the model reaction. In the optimization study, the reaction was initially carried out in the absence of solvent and catalyst and

only a little product was obtained. This result was achieved again even when the catalyst was used under solvent-free conditions. Then, the reaction was tested in chloroform, acetonitrile and ethanol with no significant change in the progress of the reaction. In continue the reaction was studied in water as the solvent. The progress of the reaction at room temperature in water was slow, and became even slower as it proceeded; however, the rate was greatly increased at reflux, so the decision was made to pursue these conditions instead. At the end of this study, the optimal amount of catalyst was determined by systematically altering the amount of taurine (Table 1). The Knoevenagel reaction was then performed with various aromatic aldehydes and malononitrile using the optimized amount of the catalyst (20 mol%, 24 mg) in water and under reflux conditions (Scheme 1). As shown in Table 2, various types of aldehydes containing electron-donating and electron-withdrawing groups were successfully converted to the corresponding nitrile products. No distinct substitution effect was observed for this reaction. The isolated products were extremely pure without the need for any costly purification steps, which can be expensive in terms of time, materials and overall yield.

Since taurine is soluble in water, it was easily separated from the products by simple filtration. The filtered solution could be

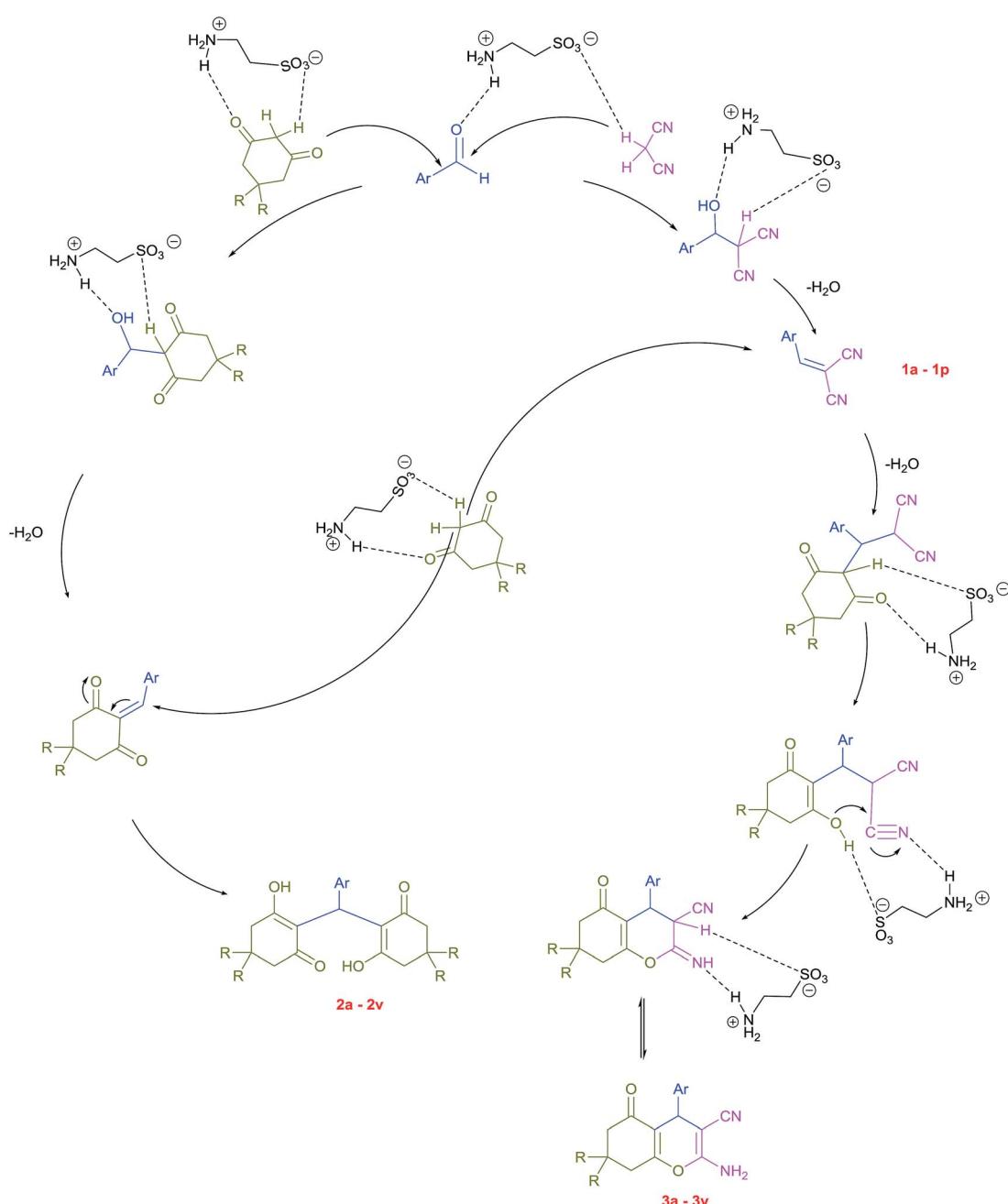
reused in the same reaction without significant loss of catalytic activity. This procedure was repeated six times for the model reaction; each time, the cycle was completed with no significant change in reaction time or yield. During the recycling studies, the product was separated and its melting point was verified after every run to ensure that the purity remained excellent (Fig. 2). It should be mentioned that since the purity of most of the derivatives was too high, we preferred not to use any organic solvent at all; thus, the melting points in this work are related to the non-recrystallized products.

Next, the efficiency of taurine for promoting the synthesis of tetraketone derivatives was investigated *via* the Knoevenagel

reaction, followed by Michael addition of an aldehyde with two equivalents of dimedone or 1,3-cyclohexanedione (Scheme 2).

It should be mentioned that taurine is only suitable to catalyze formation of open chain xanthenes in water and any effort to achieve closed chain xanthenes as single products using this catalyst was not successful.

Only a small amount of product was formed in the absence of a catalyst and solvent at the first step of the optimization of the conditions. When the catalyst was used under solvent-free conditions, a mixture of the products was achieved at different temperatures. As shown in Table 3, the reaction was tested in different solvents and conditions to determine the



Scheme 4 The plausible mechanisms of the studied reactions in the presence of taurine.

optimum conditions (entry 10). Furthermore, various aldehydes were used to prepare diverse tetraketone structures. The aldehydes selected for this purpose produced generally high yields of the desired products with acceptable reaction times and do not require further purification (Table 4). Just like the previous reaction, reusability of the catalyst was studied by selecting a typical reaction and using the catalyst-containing filtrate solution in a new reaction mixture. The process showed good reusability over 6 cycles (Fig. 2).

Ultimately, a one-pot, three-component reaction system was designed containing an aldehyde, either dimedone or 1,3-cyclohexanedione, and malononitrile, to test the ability of taurine to catalyze the formation of 2-amino-3-cyano-4*H*-pyran derivatives.

Optimization of the conditions was performed using a typical reaction of 4-chlorobenzaldehyde, dimedone and malononitrile under a variety of conditions and in the presence of different amounts of taurine as the catalyst. As shown in Table 5, the best results were obtained in refluxing water using 35 mg (28 mol%) catalyst (entry 10).

After optimization of the reaction conditions (Scheme 3), different types of aldehydes containing electron-donating and electron-withdrawing groups were subjected to the same reaction. The results showed that all types of aldehydes performed

well to give the corresponding products in good to excellent yields.

1,3-Cyclohexanedione was also used in some reactions in place of dimedone and the desired products were obtained (Table 6, entries 20–26). The purity of the isolated products was again such that there was no need for further purification or even recrystallization of the products.

As we have mentioned previously, the reusability of the catalyst was investigated in the model reactions for all three transformations studied (products were **1g**, **2g** and **3i**). In each case, six consecutive runs showed excellent reusability (Fig. 2).

Plausible mechanisms for the abovementioned reactions as catalyzed by taurine are shown in Scheme 4. Taurine acts as a bifunctional donor–acceptor reagent in which the aldehyde carbonyl site is activated by taurine and then attacked by the negatively activated methylene group in malononitrile. Elimination of water in the Knoevenagel reaction results in arylidene malononitriles that can be separated by filtration (compounds **1a**–**1p**). Moreover, it is believed that the taurine-activated aldehyde can be attacked by a taurine-enolized  $\beta$ -dicarbonyl. Then, after losing water and forming an arylidene dicarbonyl in the Knoevenagel reaction, this species can react with another enolized  $\beta$ -dicarbonyl, leading to a tetraketone (products **2a**–**2v**). For the three component system, the Knoevenagel product

Table 7 Comparison between existing literature reports and some of the taurine-catalyzed reactions in the current work

Product	Catalyst	Amount	Conditions	Time (min)	Yield (%)
	SiO <sub>2</sub> –L-proline <sup>25</sup> L-Proline–IL <sup>31</sup> MNPs–guanidine <sup>21</sup> SiO <sub>2</sub> –NH <sub>4</sub> OAc <sup>32</sup> HAP–Cs <sub>2</sub> CO <sub>3</sub> (ref. 29) CTMAB <sup>16</sup> ZnO <sup>26</sup> Na <sub>2</sub> S/Al <sub>2</sub> O <sub>3</sub> (ref. 24) Taurine <sup>a</sup> SmCl <sub>3</sub> (ref. 38) EDDA <sup>44</sup> Fe <sub>3</sub> O <sub>4</sub> @SiO <sub>2</sub> –SO <sub>3</sub> H <sup>40</sup> Al/MCM-41 (ref. 45) ChCl : urea <sup>42</sup> Fe/NaY <sup>43</sup> Taurine <sup>a</sup>	10 mol% (0.100 g) 30 mol% 0.39 mol% (0.005 g) 0.200 g 0.200 g 0.50 mmol 0.500 g 20 mol% on 0.500 g <b>20 mol% (0.025 g)</b> 20 mol% 30 mol% 0.010 g 0.100 g 1 mL 0.025 g <b>24 mol% (0.030 g)</b>	CH <sub>3</sub> CN/80 °C 80 °C H <sub>2</sub> O–PEG/RT CH <sub>2</sub> Cl <sub>2</sub> /reflux H <sub>2</sub> O/80 °C H <sub>2</sub> O/RT H <sub>2</sub> O/RT CH <sub>2</sub> Cl <sub>2</sub> /reflux <b>H<sub>2</sub>O/reflux</b> 120 °C THF/reflux H <sub>2</sub> O/RT EtOH/reflux 80 °C EtOH/reflux <b>H<sub>2</sub>O/reflux</b>	540 1440 150 450 180 90 90 30 7 20 240 80 120 70 20	95 96 96 90 86 94 86 90 <b>98</b> 95 97 83 88 86 98 <b>97</b>
	$\beta$ -Cyclodextrin <sup>69</sup> IRMOF–Zn complex <sup>68</sup> [Ch][OH] <sup>67</sup> Red sea sand <sup>66</sup> Glutamic acid <sup>74</sup> DBSA <sup>64</sup> TFE <sup>62</sup> I <sub>2</sub> (ref. 75) L-Proline <sup>76</sup> HDMBAB <sup>57</sup> TBAC <sup>61</sup> Taurine <sup>a</sup>	0.227 g 4 mol% 10 mol% 0.500 g 20 mol% 20 mol% 2 mL 10 mol% 10 mol% 2.4 mol% 10 mol% <b>28 mol% (0.035 g)</b>	Aq glycerin/40 °C 60 °C H <sub>2</sub> O/80 °C EtOH/reflux EtOH/reflux H <sub>2</sub> O/reflux Reflux DMSO/120 °C EtOH/reflux H <sub>2</sub> O/80–90 °C H <sub>2</sub> O/reflux <b>H<sub>2</sub>O/reflux</b>	30 300 60 280 40 240–420 300 210 120 450 120 30	90 90 86 85 91 69 95 88 72 90 98 <b>97</b>

<sup>a</sup> Current work.

is again formed, followed by Michael addition with a  $\beta$ -dicarbonyl. Lastly, enolization occurs, followed by amine-enamine tautomerization in the presence of taurine to produce the 4*H*-pyran derivative (products 3a–3y).

The results of this study were compared with some existing literature reports in order to better illustrate the utility of taurine in accelerating the reactions under study (Table 7). In each case, the taurine-catalyzed reaction had an advantage wherein it utilized a green catalyst and a nontoxic solvent, featured easy separation of product and catalyst, offered catalyst reusability, and resulted in lower reaction times and higher yields.

## Conclusions

The general advantages of taurine as a catalyst in the studied reactions are as follows: it is non-toxic for humans and other living organisms, it offers ease of separation of products and catalyst, it does not require use of organic solvents, it features excellent catalyst recyclability and reusability, it is highly stable and easy to store, and finally, it is commercially available at a low price. All of these points are in full compliance with the requirements of green chemistry.

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