

## COMMUNICATION

[View Article Online](#)  
[View Journal](#) | [View Issue](#)


Cite this: RSC Adv., 2015, 5, 72529

 Received 5th May 2015  
 Accepted 12th August 2015

DOI: 10.1039/c5ra08297h

[www.rsc.org/advances](http://www.rsc.org/advances)

The combination of transfer hydrogenation reaction with the advantages of  $\gamma$ -valerolactone-based ionic liquids could result in an environmentally benign method for the reduction of organic substrates. Ionic liquids containing 4-hydroxyvalerate anion were applied as alternative solvents for the reduction of acetophenone, its substituted forms and different alkenes using transition metal based catalysts. The optimal conditions (e.g. type of catalyst precursor and hydrogen donor) for the transformation were also specified.

## Catalytic transfer hydrogenation in $\gamma$ -valerolactone-based ionic liquids†

 Andrea Strádi,<sup>ab</sup> Márk Molnár,<sup>b</sup> Péter Szakál,<sup>a</sup> Gábor Dibó,<sup>bc</sup> Dániel Gáspár<sup>a</sup> and László T. Mika<sup>\*a</sup>

(RTILs),<sup>8</sup> including industrially important reactions *e.g.* Friedel-Crafts acylation,<sup>9</sup> hydroformylation,<sup>10</sup> and Beckmann-rearrangement.<sup>11</sup>

Hydrogenation reaction as one of the most important and widely studied transition metal catalyzed homogeneous transformation<sup>12</sup> was also performed in ionic liquids.<sup>13,14</sup> The catalytic transfer hydrogenation of organic substrates is a safer and more energy-efficient alternative operation without using hydrogen gas and can generally be performed in common organic solvents.<sup>15</sup> Although, the ionic liquids are widely used as environmentally benign solvents in high-pressure catalysis, there are only a few examples for application in transfer hydrogenation reactions. The combination of the mild conditions of transfer hydrogenation with the advantages of  $\gamma$ -valerolactone (GVL)-based ILs could result in an environment-friendly method for the reduction of organic substrates. In addition, due to the low vapour pressure of GVL based ILs,<sup>14</sup> the risk of environmental emission, therefore the potential hazard to health, could be significantly reduced compared to conventional organic solvents.

First, Berthold *et al.* published the application of ILs as reaction media for transfer hydrogenation of ketones and their  $\alpha,\beta$ -unsaturated forms.<sup>16</sup> Hermecz *et al.* reported the reduction of  $\alpha,\beta$ -unsaturated carbonyl derivatives by using  $[\text{ClRh}(\text{PPh}_3)_3]$  and  $[\text{Rh}(\text{cod})\text{Cl}]_2$  precursors in imidazolium, ammonium, and phosphonium containing ILs, representing outstanding chemoselectivities depending on the nature of the anions, with  $\text{TOF} = 2.5\text{--}20 \text{ h}^{-1}$  at  $90^\circ\text{C}$ .<sup>17</sup> Since, the asymmetric hydrogenation of ketones is a key reaction in the synthesis of biologically active molecules, enantioselective transfer hydrogenation reactions were also performed in ILs resulting in enantiomeric excess of 76–96%.<sup>18</sup> Dyson<sup>19</sup> and Ohta<sup>18</sup> revealed that ILs do not act only as solvents, they have a special role in the stabilization of the catalytically active species. Chakraborti *et al.* proved that ionic liquids could have a special role in the activation of organic substrates *e.g.* ketones and aldehydes *via* hydrogen bond formation between the IL and C=O group.<sup>20,21</sup>

## Introduction

Typically, the commonly used organic solvents are toxic, have high vapour pressure and negative impact on the environment. Their replacement represents a crucial part in the development of greener and environmentally benign chemical technologies,<sup>1</sup> and preference should be given to solvents with no or low toxicity according to the fifth principle of green chemistry.<sup>1b</sup> Recently several alternative solvents *e.g.* water,<sup>2</sup> fluorous solvents,<sup>3</sup> supercritical carbon-dioxide,<sup>4</sup> alcohols,<sup>5</sup> and ionic liquids (ILs) were introduced with significantly different solvent properties.<sup>6</sup> Due to their attractive properties, ILs have attracted considerable attention as environmentally benign reaction media. By the proper variation of their anionic and cationic parts, their physical properties such as vapour pressure, thermal stability, solvation strength can easily be fine-tuned.<sup>7</sup> Accordingly, wide range of chemical transformations has been successfully performed in room temperature ionic liquids

<sup>a</sup>Budapest University of Technology and Economics, Department of Chemical and Environmental Process Engineering, Műegyetem rkp. 3, Budapest H-1111, Hungary. E-mail: laszlo.t.mika@mail.bme.hu; Fax: +36 1 463 3197; Tel: +36 1 463 1263

<sup>b</sup>Eötvös Loránd University, Institute of Chemistry, Pázmány Péter str. 1/A, Budapest H-1117, Hungary

János Selye University, Faculty of Education, Department of Chemistry, 3322 Bratislavská str., Komárno SK-94501, Slovakia

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c5ra08297h



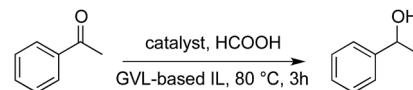
It was recently demonstrated, that selective catalytic hydrogenation can be performed in GVL-based ILs.<sup>14</sup> This generation of ILs can easily be prepared from bio-originated GVL,<sup>14,22</sup> which is a proposed sustainable platform molecule.<sup>23</sup> The reaction rate could be affected by varying the substituents of the valerate anion and the tetraalkylammonium cation.<sup>14</sup> We also reported that various functionalized olefins can be selectively reduced in tetrabutylammonium 4-hydroxyvalerate [TBA][HV] ionic liquid with TOF = 20–41 h<sup>-1</sup>. Accordingly, [TBA][HV] was selected as solvent for the transfer hydrogenation of acetophenone, a typical model substrate.

We report here the application of GVL-based ILs as reaction media for transfer hydrogenation reactions using various transition metal catalysts.

Initially, to extend the series of GVL-based ionic liquids, novel species were synthesized as follows: tetraethylammonium 4-hydroxyvalerate ([TEA][HV]), tetrapropylammonium 4-hydroxyvalerate ([TPrA][HV]), and tetrapentylammonium 4-hydroxyvalerate ([TEA][HV]). Since the temperature dependence of the vapour pressure of solvents is a key property, it was determined for novel ILs in the temperature range of 0–85 °C. Expectedly, the data obtained was found to be in the same range as measured in case of GVL and the previously published GVL-based ILs<sup>14</sup> as well (Fig. 1).

Furthermore, different catalyst precursors were investigated in the presence of [TBA][HV] and formic acid as hydrogen donor in the transfer hydrogenation of acetophenone at 80 °C (Scheme 1). Expectedly, in the absence of any catalyst no conversion was detected. The activities of the Rh-based catalyst precursors were much higher than that of obtained for the Pd-containing system (Table 1), in accordance with previous results.<sup>17a,24</sup> The activity of the Shvo catalyst, which was found to be very efficient for the conversion of levulinic acid to GVL,<sup>25</sup> was rather low under these conditions (Table 1, entry 7). The highest activity was achieved in the case of using [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>-</sup> similarly to the conventional hydrogenation reactions.<sup>14</sup> The resulted TOF was 20 h<sup>-1</sup> which is significant, compared to literature data reported so far.<sup>17</sup> Consequently, further experiments were performed using [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>-</sup>.

In contrast with the conventional high-pressure hydrogenation system, the reaction rate could not be increased by the



Scheme 1

addition of phosphine ligand. When 10 eq. triphenyl phosphine was added to the reaction mixture, the conversion decreased to 58% under identical conditions. In the presence of the sodium salt of trisulfonated triphenylphosphine (TPPTS), even lower yield (6%) of 1-phenylethanol was obtained. Similar effect was reported for the Pd-catalyzed selective reduction of alkenes.<sup>26</sup>

To optimize the reaction temperature, we repeated the reduction of acetophenone in [TBA][HV] in the temperature range of 60–90 °C. The highest conversion was obtained at 80 °C (Fig. 2).

The catalytic performance of [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>-</sup> was also evaluated utilizing different conventional hydrogen donors (Table 2). In the presence of formic acid and sodium formate >99% conversion was obtained, while by using potassium formate, the hydrogenation was incomplete. It should be noted that no ester formation was detected in the presence of HCOOH. No product formation could be detected after 3 hours by using ammonium formate (which was not even soluble under the applied conditions) and isopropyl alcohol. In accordance with the 7<sup>th</sup> principle of green chemistry, formic acid was selected as hydrogen donor since it can be produced from renewable raw materials;<sup>27</sup> moreover the carbon dioxide produced in the reaction is thermodynamically very stable and can be easily removed from the reaction mixture.

It was shown that the reaction rate of the transfer hydrogenation is determined by the concentration of the H-donor.<sup>28</sup> The negative effect of acidity on reaction rate was clearly demonstrated in case of asymmetric transfer hydrogenations.<sup>29</sup> Accordingly, we investigated this effect in the range of 1.2–6.5 molar excess of HCOOH relative to the substrate (Fig. 3).

The conversion reached a plateau between 5 : 1 and 6 : 1 molar ratio (Fig. 3), similarly to the observed case of asymmetric transfer hydrogenation of ketones, using azeotropic mixture of HCOOH/triethylamine as hydrogen donor.<sup>29g</sup> In extreme cases, such as when the ratio was over 8 : 1, the conversion decreased

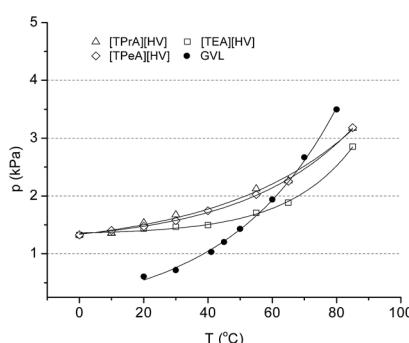


Fig. 1 Vapour pressure of GVL-based ILs compared to data for GVL (ref. 21).

Table 1 Transfer hydrogenation of acetophenone using different catalyst precursors<sup>a</sup>

Entry	Catalyst precursor	Conv. (%)	TON (-)	TOF (h <sup>-1</sup> )
1	ClRh(PPh <sub>3</sub> ) <sub>3</sub>	92	55	18.3
2	[Rh(cod)Cl] <sub>2</sub>	94	56	18.6
3	[Rh(cod) <sub>2</sub> ] <sup>+</sup> [BF <sub>4</sub> ] <sup>-</sup>	>99	60	20.0
4	[Rh(cod) <sub>2</sub> ] <sup>+</sup> [SbF <sub>6</sub> ] <sup>-</sup>	97	58	19.3
5	[Rh(cod) <sub>2</sub> ] <sup>+</sup> [B(Ph(CF <sub>3</sub> ) <sub>2</sub> ) <sub>4</sub> ] <sup>-</sup>	87	52	17.3
6	Pd(OAc) <sub>2</sub>	39	23	7.6
7	Shvo catalyst <sup>b</sup>	35	21	7.0

<sup>a</sup> Reaction conditions: 0.005 mmol catalyst, 0.3 mmol acetophenone,

<sup>b</sup>  $\{2,3,4,5-(C_6H_5)_4(\eta^5-C_4CO)\}_2H\}Ru_2(CO)_4(\mu-H)$ .



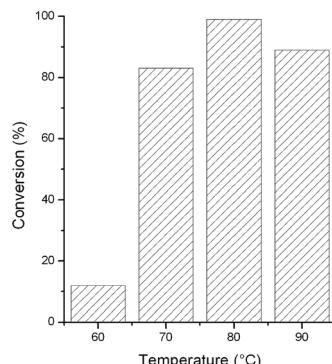


Fig. 2 Effect of reaction temperature on the reduction of acetophenone. Reaction conditions: 0.005 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ , 0.3 mmol acetophenone, 1.6 mmol HCOOH, 0.5 mL [TBA][HV],  $t = 3$  h.

significantly (below 20%) presumably due to the dilution and increased acidity of the system, which could result in the loss of the catalyst's activity. The exact amount of HCOOH remained after the completed reaction was determined by  $^1\text{H-NMR}$  spectroscopy by using GVL as an internal standard in the presence of 0.3 mmol acetophenone, 0.005 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$  in 0.5 mL [TBA][HV]. When the HCOOH to substrate ratio was 1 : 1 and 1.2 : 1, no residual HCOOH was detected after 3 hours, while in case of 3 : 1 and 5 : 1 ratio 40% decrease of the initial amount of H-donor could be obtained. At ratio 6 : 1 more residual amount of formic acid was detected with full conversion of substrate. Further increasing this ratio resulted a slight decrease of the conversion rate. It is important to note that no water formation was detected. Consequently, due to the decomposition of formic acid, an excess of hydrogen donor has to be used to reach full conversion.

It was proven that the GVL-based ILs can be easily fine-tuned by varying the chain length of the alkyl groups on the cation, resulting in significant changes in viscosity and vapour pressure. In order to determine the influence of the structure on the performance of the catalytic system, we screened a series of GVL-based ILs having different tetraalkylammonium cations. Table 3 shows, that negligible conversion could be achieved using [TMA][HV], which has relatively high melting point (75–80 °C), thus it becomes liquid at the reaction temperature and might be more useful at even higher temperatures. The

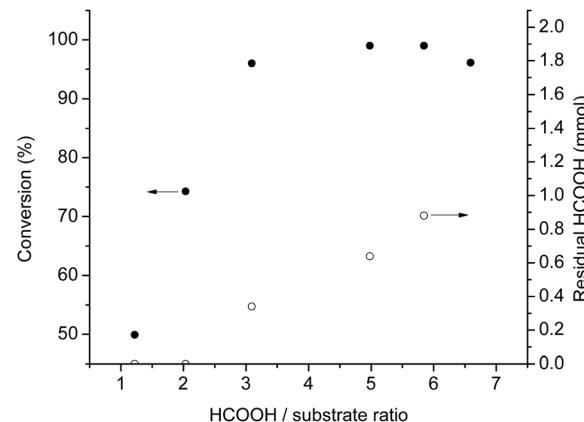


Fig. 3 Formation of 1-phenylethanol at different HCOOH/substrate molar ratio. Reaction conditions: 0.005 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ , 0.3 mmol acetophenone, 0.5 mL [TBA][HV],  $T = 80$  °C,  $t = 3$  h. ●: Conversion, ○: residual HCOOH (mmol).

replacement of methyl to ethyl group resulted in significantly higher conversion (94%, see Table 3, entry 2). Interestingly, further increase of the length of the alkyl chain on the cation had no dramatic effect on the product formation. The reaction rate reached >99% in case of [TBA][HV] (Table 3, entry 4) similarly as in the conventional hydrogenation reactions using GVL-based ILs containing hydroxyvalerate anion.

Chakraborti's studies indicated that carbonyl group(s) could participate in the hydrogen bond interaction between the oxygen atom of carbonyl species and a hydrogen atom of corresponding ionic liquid to form an activated intermediate that readily reacts with the other substrate of the reaction.<sup>20</sup> This observation was also verified for the addition reactions of  $\alpha,\beta$ -unsaturated carbonyl species.<sup>21</sup> To investigate the role of the 4-hydroxyl group of the IL's anion in transfer hydrogenation, the reduction of acetophenone was also performed in tetrabutylammonium 4-methoxyvalerate and 4-ethoxyvalerate, (where the 4-OH group was replaced with 4-OMe and 4-OEt ones). In both cases, full conversion was achieved showing that the hydroxyl group has no role in the substrate activation (Table 3, entries 7 and 8).

Table 2 Transfer hydrogenation of acetophenone using different hydrogen donors<sup>a</sup>

Entry	Hydrogen donor	Conversion (%)	TOF (h <sup>-1</sup> )
1	HCOOH	>99	20.0
2	HCOONa	>99	20.0
3	HCOOK	89	17.8
4	HCOONH <sub>4</sub>	0	0
5	iPrOH	0	0

<sup>a</sup> Reaction conditions: 0.005 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ , 0.3 mmol acetophenone, 1.6 mmol H-donor, 0.5 mL [TBA][HV],  $T = 80$  °C,  $t = 3$  h.

Table 3 Transfer hydrogenation of acetophenone using different GVL-based ionic liquids<sup>a</sup>

Entry	Ionic liquid	Conv. (%)	TOF (h <sup>-1</sup> )
1	$[(\text{CH}_3)_4\text{N}]^+[\text{HV}]^-$ , [TMA][HV]	6	1.2
2	$[(\text{C}_2\text{H}_5)_4\text{N}]^+[\text{HV}]^-$ , [TEA][HV]	93	18.7
3	$[(\text{C}_3\text{H}_7)_4\text{N}]^+[\text{HV}]^-$ , [TPtA][HV]	96	19.2
4	$[(\text{C}_4\text{H}_9)_4\text{N}]^+[\text{HV}]^-$ , [TBA][HV]	>99	20.0
5	$[(\text{C}_5\text{H}_{11})_4\text{N}]^+[\text{HV}]^-$ , [TPtA][HV]	96	19.3
6	$[(\text{C}_6\text{H}_{13})_4\text{N}]^+[\text{HV}]^-$ , [TBA][HV]	94	18.8
7	$[(\text{C}_4\text{H}_9)_4\text{N}]^+[\text{MeOV}]^-$ , [TBA][MeOV]	>99	20.0
8	$[(\text{C}_4\text{H}_9)_4\text{N}]^+[\text{EtOV}]^-$ , [TBA][EtOV]	>99	20.0

<sup>a</sup> Reaction conditions: 0.005 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ , 0.3 mmol acetophenone, 1.6 mmol HCOOH, 0.5 mL ionic liquid,  $T = 80$  °C,  $t = 3$  h.



Table 4 Reuse of catalyst phase for reduction of acetophenone<sup>a</sup>

Cycle	Time (h)	Conversion
1	3	>99
2	3	>99
3	3	>99
4	4	>99

<sup>a</sup> Reaction conditions: 0.01 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ , 0.3 mmol acetophenone, 1.6 mmol HCOOH in 1 mL [TBA][HV],  $T = 80^\circ\text{C}$ .

It was established that the C2-hydrogen of the imidazolium cation could activate the C=O group *via* hydrogen bond formation, as well.<sup>20a,b</sup> In order to further demonstrate the possible effect of the IL's structure on the efficiency of the system, butylmethyl imidazolium-based ionic liquids  $[\text{BMIM}]^+[\text{X}]^-$  ( $\text{X} = \text{Cl}^-$ ,  $\text{PF}_6^-$ ,  $\text{OH}^-$ , octylsulphonate) were investigated for comparison. Unexpectedly, no conversion could be detected under the previously optimized conditions. Presumably the proposed interaction could not be established under the applied acidic conditions. Similar observation was reported for the selective reduction of chalcone performed in dialkyl imidazolium type ILs.<sup>17a</sup> It should be noted that the reduction of the carbonyl group could not be achieved even in tetraalkylphosphonium-based ILs. Analogously, no product formation was detected applying conventional organic solvents such as toluene, methanol, ethanol, propanol, and dimethyl sulfoxide.

To check the activity and stability of the catalytic system, a second portion of acetophenone (0.3 mmol) and HCOOH (1.6 mmol) was added to the mixture of the completed reaction after 3 hours. It was observed that the catalyst remained active and the substrate was completely converted in the consecutive runs, as well. However, a slight decrease in the activity could be detected in the fourth cycle, in which 4 hours was needed to reach the full conversion. Since [TBA][HV] was miscible with the product, it was extracted from the catalyst phase by pentane ( $2 \times 0.5$  mL) extraction. Subsequently, HCOOH (1.6 mmol) and the substrate (0.3 mmol) were added and the next cycle was started. This procedure was repeated for three times. Although, full conversion was achieved in cycles 1–3, a slight decrease in the activity was detected in the 4<sup>th</sup> run (Table 4). Similar remarks were described for Rh-catalyzed transfer hydrogenation of chalcone, where toluene was used for the extraction.<sup>17</sup> The isolated yield of 1-phenylethanol was 58%.

As already stated, the electronic properties of the substituents on the carbonyl group had significant effect on the reactivity. The 4-substituted acetophenone derivatives having electron donating substituent such as methyl or methoxy group, were found to be less reactive compared to acetophenone (Table 5, entries 1–3). Consequently, the results indicated that the introduction of electron withdrawing groups led to higher conversion rate (Table 5, entries 4–6), as expected. The 2- and 3-substituted derivatives also showed lower reactivity (Table 5, entries 7 and 8). Identical observations were reported for both conventional and asymmetric transfer hydrogenations.<sup>30</sup> It is noteworthy that the reduction of 4-nitroacetophenone led to the

formation of 1-(4-nitrophenyl)ethanol. Concerning non-aromatic carbonyl compounds, cyclic ketones were reduced in [TBA][HV] under identical conditions and, as it was predicted, full conversion was achieved (Table 6, entries 1 and 2). Regarding the levulinic acid and its methyl ester, it was revealed that both species can be selectively reduced to gamma-valerolactone (Table 6, entries 3 and 4).

Further investigations were focused on the applicability of  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ /HCOOH catalyst system for the hydrogenation of different substrates with olefinic carbon–carbon double bond, under the optimized conditions. It was shown that alk-1-enes could be quantitatively converted to the corresponding alkanes with  $\text{TOF} = 20 \text{ h}^{-1}$  (Table 7, entries 1–3), as expected. For cycloalkenes full conversions were observed (Table 7, entries 4 and 5), similarly to the catalytic hydrogenation of olefins in these ILs using molecular hydrogen. When crotyl bromide and cinnamic acid were subjected to reduction, >99% conversion was achieved for both cases, however, in case of crotyl bromide the reaction was already completed in 1.5 hour (Table 7, entries 6 and 7). For the sterically hindered *trans*-stilbene slightly lower conversion was observed (Table 7, entry 8).

The selective reduction of  $\alpha,\beta$ -unsaturated carbonyl compounds were investigated by Hermecz *et al.* in different imidazolium-based ILs using  $\text{Pd}(\text{OAc})_2$ ,  $[\text{Rh}(\text{cod})\text{Cl}]_2$  and  $\text{ClRh}(\text{PPh}_3)_3$  as catalyst precursors in the presence of  $\text{HCOONH}_4$  or HCOOH as H-donor.<sup>17a</sup> When 0.6 mmol of 1-cyclohex-2-enone in the presence of 3.2 mmol HCOOH and 0.01 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$  in 1 mL [TBA][HV] was reduced at  $80^\circ\text{C}$ , the complete conversion to cyclohexanone was detected after 15 min; however, after 3 hours both the carbon–carbon double bonds and the carbonyl groups were completely reduced to cyclohexanol. The same results were obtained in case of using chalcone as substrate indicating that the hydrogenation of  $\text{C}=\text{C}$  bond is much faster than  $\text{C}=\text{O}$  bond. However, by applying longer reaction time both functional groups were reduced. The reduction of cinnamic aldehyde (0.3 mmol) led to the formation of 3-phenylpropanal with conversion of 66% under identical conditions.

Table 5 Transfer hydrogenation of substituted acetophenones<sup>a</sup>

Entry	R	Time (h)	Conv. (%)	TOF ( $\text{h}^{-1}$ )
1	H	3	>99	20.0
2	4-CH <sub>3</sub>	3	17	3.4
3	4-OCH <sub>3</sub>	3	27	5.4
4	4-NO <sub>2</sub>	1.5	>99	40.0
5	4-F	2.5	>99	15.0
6	4-Br	2.5	>99	20.0
7	3-Br	3	94	18.8
8	2-Br	3	60	12.0

<sup>a</sup> Reaction conditions: 0.005 mmol  $[\text{Rh}(\text{cod})_2]^+[\text{BF}_4]^-$ , 0.3 mmol substrate, 1.6 mmol HCOOH, 0.5 mL [TBA][HV],  $T = 80^\circ\text{C}$ .



Table 6 Transfer hydrogenation of non-aromatic carbonyl species<sup>a</sup>

Entry	R	Time (h)	Conv. (%)	TOF (h <sup>-1</sup> )
1		3	>99	20.0
2		3	>99	20.0
3 <sup>b</sup>		3	96	19.2
4 <sup>b</sup>		3	>99	20.0

<sup>a</sup> Reaction conditions: 0.005 mmol [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>-</sup>, 0.3 mmol substrate, 1.6 mmol HCOOH, 0.5 mL [TBA][HV], T = 80 °C. <sup>b</sup> Product:  $\gamma$ -valerolactone.

Table 7 Transfer hydrogenation of several alkenes<sup>a</sup>

Entry	Substrate	Time (h)	Conv. (%)	TOF (h <sup>-1</sup> )
1		3	>99	20.0
2		3	>99	20.0
3		3	>99	20.0
4		3	>99	20.0
5		3	>99	20.0
6		1.5	>99	40.0
7		3	>99	20.0
8		3	72	14.4

<sup>a</sup> Reaction conditions: 0.005 mmol [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>-</sup>, 0.3 mmol substrate, 1.6 mmol HCOOH, 0.5 mL [TBA][HV], T = 80 °C.

The reproducibility of the catalytic reactions were confirmed by repeating the reduction of acetophenone, hex-1-ene, oct-1-ene and dec-1-ene (0.3 mmol) in the presence of HCOOH (1.6 mmol) and [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>-</sup> (0.005 mmol) in [TBA][HV]. Complete conversions were achieved after 3 h at 80 °C in all cases.

## Experimental

Tetramethylammonium 4-hydroxyvalerate [TMA][HV], tetrabutylammonium 4-hydroxyvalerate [TBA][HV], tetrahexylammonium 4-hydroxyvalerate [THA][HV], tetrabutylammonium 4-methoxyvalerate [TBA][MeOV] and tetrabutylammonium 4-ethoxyvalerate [TBA][4EtOV] were prepared by published methods.<sup>14,20</sup> 1-Butyl-3-methylimidazolium

hexafluorophosphate, 1-butyl-3-methylimidazolium tetrafluoroborate, 1-butyl-3-methylimidazolium octyl sulfate, chalcone, 1-cyclohex-2-enone, cinnamic acid were purchased from Sigma-Aldrich Kft. and used as received. 1-Butyl-3-methylimidazolium chloride, 1-butyl-3-methylimidazolium hydroxide and 1-ethyl-3-methylimidazolium hydroxide were prepared by published methods.<sup>31</sup> The water content of the ILs was determined by Karl-Fischer titration.

### Preparation of tetraethylammonium 4-hydroxyvalerate [TEA][HV]

The mixture of 14.73 g (100 mmol) of tetraethylammonium 4-hydroxide (in 25% aqueous solution) and 10.51 g (105 mmol) of  $\gamma$ -valerolactone was stirred at room temperature for 1 hour. After the reaction was completed, the residual water was removed *in vacuo* at 80 °C. The product was isolated as a colourless, highly viscous liquid. Yield 22.11 g (87.6%). <sup>1</sup>H-NMR (D<sub>2</sub>O)  $\delta$ , ppm: 0.98 (d, 3H, J = 6.5 Hz), 1.07 (t, 12H, J = 7.1 Hz), 1.52 (q, 2H, J = 7.2 Hz), 2.04 (m, 4H), 3.07 (q, 8H, J = 7.1 Hz), 3.62 (m, 1H, J = 6.1 Hz) <sup>13</sup>C-NMR (D<sub>2</sub>O)  $\delta$ , ppm: 10.51, 25.7, 37.95, 38.72, 55.80 (t, J = 3.3 Hz), 71.5, 186.8 (ESI Fig. S1 and S2†). H<sub>2</sub>O (wt%): 2.1%.

### Preparation of tetrapropylammonium 4-hydroxyvalerate [TPrA][HV]

The mixture of 10.2 g (50 mmol) of tetrapropylammonium 4-hydroxide (in 20% aqueous solution) and 5.26 g (53 mmol) of  $\gamma$ -valerolactone was stirred at room temperature for 1 hour. After the reaction was completed, the residual water was removed *in vacuo* at 80 °C. The product was isolated as a colourless, highly viscous liquid. Yield 15.25 g (98.6%). <sup>1</sup>H-NMR (D<sub>2</sub>O)  $\delta$ , ppm: 0.81 (t, 12H, J = 7.1 Hz), 1.04 (d, 3H, J = 6.3 Hz), 1.57 (m, 8H), 2.09 (m, 2H), 3.02 (m, 8H), 3.67 (m, 1H, J = 6.3 Hz) <sup>13</sup>C-NMR (D<sub>2</sub>O)  $\delta$ , ppm: 9.79, 14.76, 21.65, 33.92, 34.69, 59.8 (t, J = 2.6 Hz), 67.57, 183.05 (ESI Fig. S3 and S4†). H<sub>2</sub>O (wt%): 4.2%.

### Preparation of tetrapentylammonium 4-hydroxyvalerate [TPeA][HV]

The mixture of 15.8 g (50 mmol) of tetrapentylammonium 4-hydroxide (in 20% aqueous solution) and 5.50 g (55 mmol) of  $\gamma$ -valerolactone was stirred at room temperature for 1 hour. After the reaction was completed, the residual water was removed *in vacuo* at 80 °C. The product was isolated as a colourless, highly viscous liquid. Yield 19.83 g (93.1%). <sup>1</sup>H-NMR (D<sub>2</sub>O)  $\delta$ , ppm: 0.80 (t, 12H, J = 6.5 Hz), 1.06 (d, 3H, J = 6.3 Hz), 1.24 (m, 16H), 1.55 (m, 10H), 2.29 (t, 2H, J = 7.1 Hz), 3.08 (m, 8H, J = 7.3 Hz), 3.7 (m, 1H) <sup>13</sup>C-NMR (D<sub>2</sub>O)  $\delta$ , ppm: 13.17, 20.81, 21.58, 21.80, 27.79, 34.08, 34.82, 58.26 (t, J = 3.60 Hz), 67.71, 183.03 ESI (Fig. S5 and S6†). H<sub>2</sub>O (wt%): 2.6%.

The vapour pressure of the ionic liquids were measured by using a Rosemount manometer attached to a 25 mL Hastelloy-C high-pressure reactor (Parr Inst., IL, USA) equipped with magnetic stirrer and a thermocouple. The latter was connected to the computer through an AD interface to register the actual temperature. Initially, the reactor was charged with 1 mL of the corresponding ionic liquid, followed by the attachment of the



manometer, and de-aerated at room temperature. The reactor was then placed into ice–water mixture. The exact value of the vapour pressure was recorded when it was stabilized at the desired temperature.

The transfer hydrogenation reactions were performed in Hach tubes (100 × 13 mm) closed with screw-cap. In a typical experiment, 0.005 mmol of the correspondent catalyst was dissolved in 0.5 mL ionic liquid, then 1.5 mmol H-donor and 0.3 mmol substrate was added to the solution. The reaction mixture was stirred at 80 °C for 3 h. The conversion and yield were determined by GC analysis and <sup>1</sup>H-NMR spectroscopy.

In order to determine the amount of residual HCOOH in the reaction, equimolar amount of HCOOH and GVL (as internal standard) was added to the mixture of 0.5 mL [TBA][HV] and 0.3 mmol acetophenone. A sample was taken before heating up the mixture and another one after the reaction was completed. The samples were dissolved in d<sub>6</sub>-acetone, then analyzed by <sup>1</sup>H-NMR.

The NMR spectra were recorded on a Brucker Avance 250 spectrometer.

In the recyclability experiments 0.3 mmol of acetophenone was reduced by using 0.01 mmol of [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>−</sup> in 1 mL [TBA][HV] at 80 °C. After completing the reaction the product was extracted with 2 × 0.5 mL pentane. To initiate the following cycle, acetophenone (0.3 mmol) and HCOOH (1.6 mmol) were added to the catalyst phase followed by heating up to 80 °C. An intensive gas evolution indicated the start of the reaction. The procedure was repeated 3 times and the isolated products from the consecutive cycles were analyzed by GC.

GC analyses were performed on an HP 5890N instrument with a HP-5 capillary column (15 m × 0.25 μm) using H<sub>2</sub> as a carrier gas. For the analysis, 5 μL of reaction mixture was dissolved in 1 mL of methylene chloride followed by the addition of 5 μL toluene as internal standard.

## Conclusions

γ-Valerolactone-based ionic liquids (ILs) containing hydroxylvalerate anion were successfully applied as alternative solvents for homogenous catalytic transfer hydrogenation of acetophenone and its substituted forms, functionalized ketones and alkenes. The highest conversions were achieved by using [Rh(cod)<sub>2</sub>]<sup>+</sup>[BF<sub>4</sub>]<sup>−</sup> as catalyst precursor and formic acid as hydrogen donor at 80 °C, with the molar ratio of HCOOH/substrate between 5 : 1 and 6 : 1. A series of GVL-based ILs with tetraalkylammonium cations were tested showing negligible influence of the cation structure on the catalytic activity. The potential recyclability of the catalytic system was demonstrated in four consecutive cycles for the reduction of acetophenone.

## Acknowledgements

This work was funded by the Budapest University of Technology and Economics under project number KMR\_12-1-2012-0066 and the Bolyai János Research Scholarship of the Hungarian Academy of Sciences.

## Notes and references

- (a) F. M. Kerton, *Alternative Solvents for Green Chemistry*, RSC, Cambridge, 2009; (b) P. T. Anastas and J. C. Warner, *Green Chemistry: Theory and Practice*, Oxford University Press, Oxford, UK, 1998.
- (a) *Water in Organic Synthesis*, ed. S. Kobayashi, Georg Thieme Verlag KG, Stuttgart, 2012; (b) F. Joó, *Aqueous Organometallic Catalysis*, Kluwer Academic Press, Dordrecht, The Netherlands, 2001.
- (a) *Handbook of Fluorous Chemistry*, ed. J. A. Gladysz, D. P. Curran and I. T. Horváth, Wiley-VCH, Weinheim, 2005; (b) I. T. Horváth, *Acc. Chem. Res.*, 1998, **31**, 641.
- Handbook of Green Chemistry, in *Supercritical Solvents*, ed. W. Leitner and P. G. Jessop, Wiley-VCH, Weinheim, vol. 4, 2010.
- W. Keim, *Chem. Ing. Tech.*, 1984, **56**, 850.
- D. J. Adams, P. J. Dyson and S. J. Taverner, *Chemistry in Alternative Reaction Media*, John Wiley & Sons Ltd, Chichester, 2004.
- (a) H. Olivier-Bourbigou, L. Magna and D. Morvan, *Appl. Catal., A*, 2010, **373**, 1; (b) Handbook of Green Chemistry, *Green Solvents, Ionic Liquids*, ed. P. T. Anastas, P. Wasserscheid and A. Stark, vol. 6, Wiley-VCH, Weinheim, 2005.
- (a) T. Welton, *Chem. Rev.*, 2009, **99**, 2071; (b) V. A. Parvulescu and C. Hardacre, *Chem. Rev.*, 2007, **107**, 2615; (c) J. P. Hallet and T. Welton, *Chem. Rev.*, 2011, **111**, 3508; (d) Q. Zhang, S. Zhang and Y. Deng, *Green Chem.*, 2011, **13**, 2619.
- (a) S. Csihony, H. Mehdi and I. T. Horváth, *Green Chem.*, 2001, **3**, 307; (b) C. J. Adams, M. J. Earle, G. Roberts and K. R. Seddon, *J. Chem. Soc., Chem. Commun.*, 1998, 2097.
- C. P. Mehnert, R. A. Coock, N. C. Dispenziere and E. J. Mozeleski, *Polyhedron*, 2004, **23**, 2679.
- V. Fábos, D. Lantos, A. Bodor, A. M. Bálint, L. T. Mika, O. E. Sielcken, A. Cuiper and I. T. Horváth, *ChemSusChem*, 2008, **1**, 189.
- (a) P. A. Chaloner, M. A. Esteruelas, F. Joó and L. Oro, *Homogeneous Hydrogenation*, Kluwer Academic Press, Dordrecht, 1994; (b) L. Oro, Hydrogenation – Homogeneous, in *Encyclopedia of Catalysis*, ed. I. T. Horváth, John Wiley & Sons, Hoboken, New Jersey, 2003.
- (a) Y. Chauvin, L. Mussmann and H. Olivier, *Angew. Chem., Int. Ed. Engl.*, 1995, **34**, 2698; (b) P. A. Z. Suarez, J. E. L. Dullius, S. Einloft, R. F. de Souza and J. Dupont, *Polyhedron*, 1996, **15**, 1217; (c) P. J. Dyson, G. Laurenczy, C. A. Ohlin, J. Vallance and T. Welton, *Chem. Commun.*, 2003, 2418.
- A. Strádi, M. Molnár, M. Óvári, G. Dibó, F. U. Richter and L. T. Mika, *Green Chem.*, 2013, **15**, 1857.
- (a) G. Brieger and T. J. Nestrick, *Chem. Rev.*, 1974, **74**, 567; (b) R. A. W. Johnstone and A. H. Wilby, *Chem. Rev.*, 1985, **85**, 129.
- H. Berthold, T. Schotten and H. Hönig, *Synthesis*, 2002, **11**, 1607.
- (a) Z. Baán, Z. Finta, G. Keglevich and I. Hermecz, *Green Chem.*, 2009, **11**, 1937; (b) Z. Baán, Z. Finta, G. Keglevich and I. Hermecz, *Tetrahedron Lett.*, 2003, **46**, 6203.

18 I. Kawasaki, K. Tsunoda, T. Tsuji, T. Yamaguchi, H. Shibuta, N. Uchida, M. Yamashita and S. Ohta, *Chem. Commun.*, 2005, 2134.

19 T. J. Geldbach and P. Dyson, *J. Am. Chem. Soc.*, 2004, **126**, 8114.

20 (a) A. K. Chakraborti and S. R. Roy, *J. Am. Chem. Soc.*, 2009, **131**, 6902; (b) A. K. Chakraborti, S. R. Roy, D. Kumar and P. Chopra, *Green Chem.*, 2008, **10**, 1111; (c) A. Sarkar, S. R. Roy, N. Parikh and A. K. Chakraborti, *J. Org. Chem.*, 2011, **76**, 7132; (d) S. R. Roy, P. S. Jadhavar, K. Seth, K. K. Sharma and A. K. Chakraborti, *Synthesis*, 2011, **14**, 2261.

21 (a) A. Sarkar, S. A. Roy and A. K. Chakraborti, *Chem. Commun.*, 2011, **47**, 4538; (b) S. A. Roy and A. K. Chakraborti, *Org. Lett.*, 2010, **12**, 3866.

22 D. Fegyverneki, L. Orha, G. Láng and I. T. Horváth, *Tetrahedron*, 2010, **66**, 1078.

23 I. T. Horváth, H. Mehdi, V. Fábos, L. Boda and L. T. Mika, *Green Chem.*, 2008, **10**, 238.

24 C. Comyns, N. Karodia, S. Zeler and J. A. Andersen, *Catal. Lett.*, 2000, **67**, 113.

25 V. Fábos, L. T. Mika and I. T. Horváth, *Organometallics*, 2014, **33**, 181.

26 J. M. Brunel, *Tetrahedron*, 2007, **63**, 3899.

27 R. Xing, W. Qi and G. W. Huber, *Energy Environ. Sci.*, 2011, **4**, 2193.

28 A. Bényei and F. Joó, *J. Mol. Catal.*, 1990, **58**, 151.

29 (a) C. Wang, X. F. Wu and J. L. Xiao, *Chem.-Asian J.*, 2008, **3**, 1750; (b) L. Jiang, T. F. Wu, Y. C. Chen, J. Zhu and J. G. Deng, *Org. Biomol. Chem.*, 2006, **4**, 3319; (c) F. Wang, H. Liu, L. F. Cun, J. Zhu, J. G. Deng and Y. Z. Jiang, *J. Org. Chem.*, 2005, **70**, 9424; (d) Y. Xing, J. S. Chen, Z. R. Dong, Y. Y. Li and J. X. Gao, *Tetrahedron Lett.*, 2006, **47**, 4501; (e) Z. Q. Zhou and L. H. Wu, *Catal. Commun.*, 2008, **9**, 2539; (f) X. F. Wu, X. G. Li, F. King and J. L. Xiao, *Angew. Chem., Int. Ed.*, 2005, **44**, 3407; (g) X. F. Wu, J. K. Liu, D. Di Tommaso, J. A. Iggo, C. R. A. Catlow, J. Baesa and J. L. Xiao, *Chem.-Eur. J.*, 2008, **14**, 7699.

30 (a) M. Aydemir, N. Meric, C. Kayan, F. Ok and A. Baysal, *Inorg. Chim. Acta*, 2013, **398**, 1; (b) R. Guo, X. Chen, C. Elpet, D. Song and R. H. Morris, *Org. Lett.*, 2005, **7**, 1757; (c) J. W. Faller and A. R. Lavoie, *Organometallics*, 2001, **20**, 5245; (d) S. Hashiguchi, A. Fujii, J. Takehara, T. Ikariya and R. Noyori, *J. Am. Chem. Soc.*, 1995, **117**, 7562.

31 (a) P. J. Dyson, M. C. Grossel, N. Srinivasan, T. Vine, T. Welton, D. J. Williams, A. J. P. White and T. Zigras, *J. Chem. Soc., Dalton Trans.*, 1997, 3465; (b) P. A. Z. Suarez, S. Einloft, J. E. L. Dullius, R. F. de Souza and J. Dupont, *J. Chim. Phys.*, 1998, **95**, 1626; (c) A. R. Hajipoura and F. Rafieeb, *J. Iran. Chem. Soc.*, 2009, **6**, 647.

