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## Ruthenium catalyzed transformation of levulinic acid to $\gamma$ -valerolactone in water<sup>†</sup>

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High catalytic activity for selective and efficient transformation of levulinic acid (LA) to  $\gamma$ -valerolactone (GVL) in water was achieved over ( $\eta^6$ -*p*-cymene)Ru(*ii*)-pyridylamine,  $[(\eta^6\text{-C}_{10}\text{H}_{14})\text{RuCl}(\kappa^2\text{-L})]^+$  (*L* = *N*<sub>amine</sub>-substituted pyridylamine ligands) based molecular catalysts. A series of complexes with pyridylamine ligands having different electronic and steric properties were synthesized and characterized. A significant influence of the *N*<sub>amine</sub>-substituents of the pyridylamine ligand on the catalytic activity was observed where the  $[(\eta^6\text{-p-cymene})\text{RuCl}(\kappa^2\text{-pyNH}^n\text{pr})]^+$  catalyst ([Ru]-2) outperformed others with 87% yield and >99% selectivity for GVL at 80 °C in water. Advantageously, the activity of [Ru]-2 was also scaled up to gram scale transformation of LA to GVL. Control experiments, pH dependent NMR and mass studies revealed the involvement of crucial reaction intermediates and catalytic species in the transformation of LA to GVL.

### Sustainability spotlight

With the increase in global population, industrialization, and civilization, there is a rapid growth in energy consumption and environmental pollution. To address these issues, we need to find alternative renewable energy resources and advanced cost-effective technologies. Abundantly available lignocellulosic biomass can serve as an alternative energy resource, as it can be transformed into a variety of bio-platform chemicals including furfural, 5-hydroxymethyl furfural, levulinic acid, and  $\gamma$ -valerolactone (GVL). Herein, we demonstrated low temperature hydrogenation of levulinic acid to GVL using *N*-pyridylamine based Ru catalysts in water. Notably, GVL has wide application in the production of biofuels, fuel additives, and as a green solvent. This work addresses the UN SDG 7 (Affordable and Clean Energy) and SDG 13 (Climate Action).

## Introduction

In the current scenario of depleting fossil fuel resources, abundantly available biomass is being extensively explored as alternative sustainable feedstock for the production of a variety of value-added chemicals and fuel components.<sup>1,2</sup> Among several biomass-derived chemicals, levulinic acid (LA) can be readily produced by the acid catalyzed hydrolysis of sugars and is one of the top ten biomass-derived platform chemicals.<sup>3,4</sup> The valorization of LA can produce a wide variety of value-added chemicals and fuel blenders such as  $\gamma$ -valerolactone (GVL), valeric acid, 2-methyl tetrahydrofuran, and valeric esters.<sup>5</sup> GVL is of interest as it offers tremendous flexibility in downstream applications and upgradation into a variety of value-added chemicals.<sup>6–9</sup> Due to its chemical and physical properties, it can be used in various ways, for example as a gasoline blender, as a natural aromatic substance,<sup>10</sup> as an additive in cosmetics

and in domestic products.<sup>11</sup> Moreover, enantio-pure GVL can be utilized as a chiral building block for the synthesis of various pharmaceutical products.<sup>12</sup>

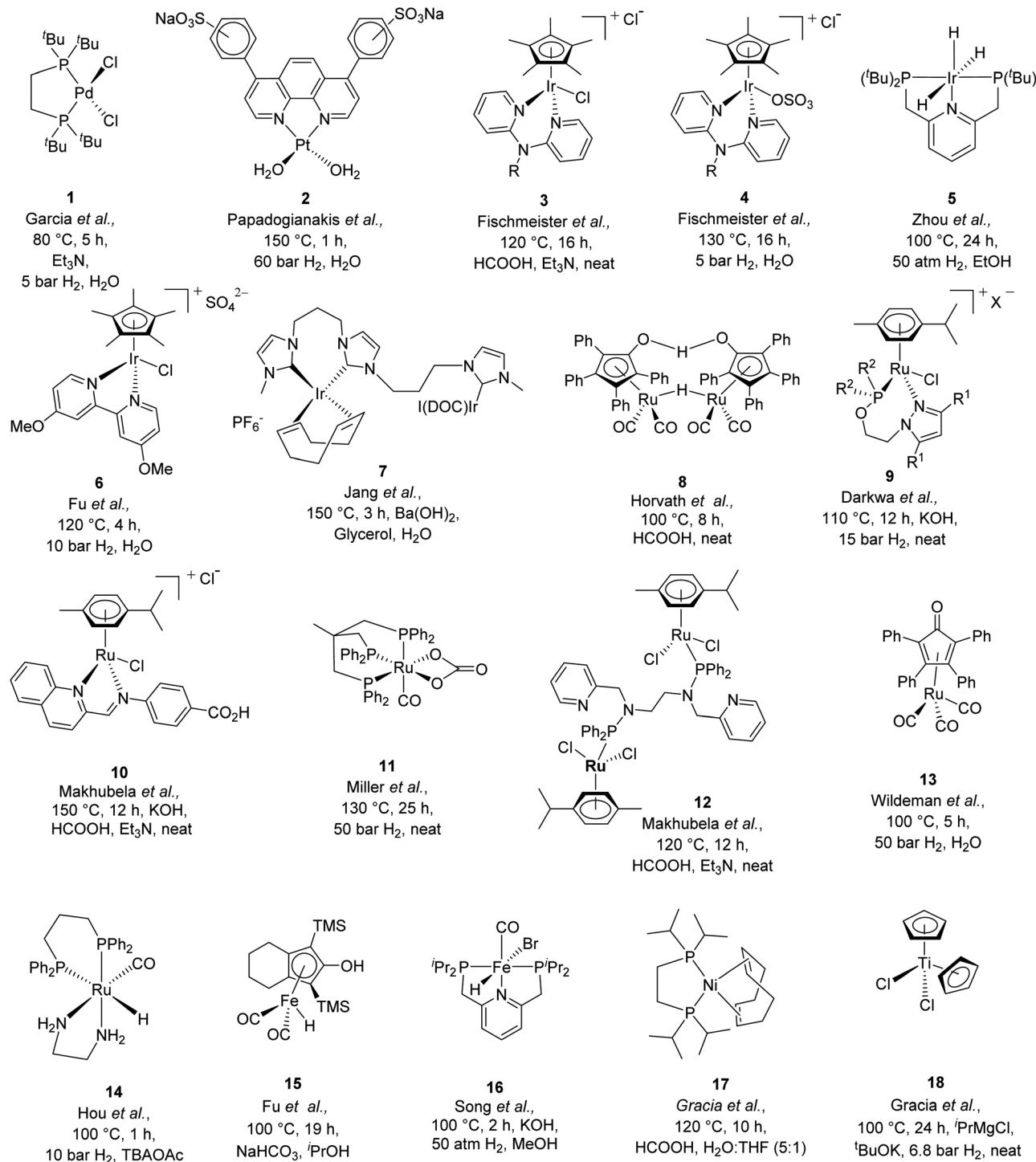
Various homogeneous as well as heterogeneous catalytic systems have been extensively explored for the transformation of LA to GVL. A number of heterogeneous catalysts based on various noble and non-noble metals (such as Pt,<sup>13</sup> Ru,<sup>14</sup> Au,<sup>15</sup> Fe,<sup>16</sup> Pd,<sup>17</sup> Ni,<sup>18</sup> and Cu,<sup>19,20</sup>) were employed and found efficient for the transformation of LA to GVL. However, reaction conditions were relatively harsher (higher temperature, 130–150 °C and higher pressure of H<sub>2</sub> gas, 12–60 bar) while using these catalysts. A number of review articles listing the various heterogeneous catalytic systems for this transformation are published in recent years.<sup>10,21</sup> On the other hand, in the search for efficient homogeneous catalytic systems for the transformation of LA to GVL, several Pd (1 in Scheme 1),<sup>22</sup> Pt (2 in Scheme 1),<sup>23</sup> and Ir (3, 4 in Scheme 1)<sup>24,25</sup> based molecular catalysts were explored. For instance, Zhou *et al.* reported an iridium trihydride complex with the PNP pincer ligand (5 in Scheme 1) for LA to GVL conversion at 100 °C and 50 bar H<sub>2</sub> and achieved a TOF of nearly 1480 h<sup>-1</sup>.<sup>26</sup> Fu *et al.* reported a half sandwich Ir-cp\*-bipyridine based complex (6 in Scheme 1) for the transformation of LA to GVL in water at 10 bar H<sub>2</sub> or with formic acid at 120 °C.<sup>27</sup> Jang *et al.* also reported an

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Scheme 1 Molecular catalysts explored for the transformation of LA to GVL.

Ir(triscarbene) catalyst (7 in Scheme 1) for transfer hydrogenation of LA to GVL in water at 150 °C using glycerol as a hydrogen source in the presence of Ba(OH)<sub>2</sub> and observed an appreciably high TON of 500 000.<sup>28</sup> Recently, Tu *et al.* used the self-supporting strategy and reported a solid molecular catalyst based on bis-N-heterocyclic-carbene-iridium complexes for LA to GVL conversion at 100 °C and 1 bar H<sub>2</sub>.<sup>29</sup> Although, Ir based

molecular catalysts are reported to be more active and display a higher TOF than Ru based catalysts, the Ru catalysts have relatively low cost and easy availability and thus used more widely at laboratory and industrial scales. Among them, most of the Ru catalysts explored were associated with the use of phosphine-based ligands (Table S1†). For instance, Horvath *et al.* reported an *in situ* generated catalyst by using Ru(acac)<sub>3</sub>

and the TPPTS (triphenylphosphine trisulfonate) ligand at 140 °C and 70 bar H<sub>2</sub> pressure for the hydrogenation of LA to GVL and achieved complete conversion of LA with a turnover frequency (TOF) of 50 h<sup>-1</sup>.<sup>30</sup> Furthermore, Guo *et al.* used RuCl<sub>3</sub> with PPh<sub>3</sub> for the hydrogenation of LA to GVL using HCOOH as the H<sub>2</sub> source (or 40 bar H<sub>2</sub> with 40 bar CO<sub>2</sub>) in the presence of pyridine at 150 °C and observed a TOF of 155 h<sup>-1</sup>.<sup>31</sup> Beller *et al.*, Mika *et al.*, and Leitner *et al.* also used triphos/biphos ligands with Ru(II) precursors at 140–160 °C and obtained a TOF of 452 h<sup>-1</sup>, 3540 h<sup>-1</sup>, and 144 h<sup>-1</sup> respectively.<sup>32–34</sup> Furthermore, Horvath *et al.* explored a ruthenium based Shvo catalyst (8 in Scheme 1), while using formic acid as a hydrogen source at 100 °C under solventless conditions to achieve a TOF of 177 h<sup>-1</sup>.<sup>35</sup> Darkwa *et al.* reported solvent-free transformation of LA to GVL using pyrazolylphosphite and pyrazolylphosphinite ruthenium(II) complexes (9 in Scheme 1), using both formic acid (TOF 60 h<sup>-1</sup>) and molecular hydrogen (15 bar) as hydrogen sources at 100–120 °C.<sup>11</sup> Makhubela *et al.* replaced the phosphine ligands with *N,N* donor iminopyridine ligands for Ru(II) (10 in Scheme 1) catalyzed LA to GVL transformation and the reaction was performed under solvent free conditions using formic acid as a hydrogen source to achieve a TOF of 81.6 h<sup>-1</sup> at 150 °C.<sup>36</sup> Miller *et al.* reported triphos complexes of ruthenium (11 in Scheme 1) for the transformation of LA to GVL at 130 °C and 50 bar H<sub>2</sub> (TOF 48 h<sup>-1</sup>).<sup>37</sup> Makhubela *et al.* reported a dinuclear Ru complex having multidentate pyridyl-aminophosphinite and pyridyl-phosphoramidite ligands (12 in Scheme 1) for the transfer hydrogenation of LA using formic acid as a hydrogen source at 120 °C under neat conditions and observed complete conversion of LA in 8 h with a TOF of 123 h<sup>-1</sup>.<sup>38</sup> Wildeman *et al.* used a  $\eta^4$ -(2,3,4,5-tetraphenylcyclopentadienone)-Ru(CO)<sub>3</sub> complex (13 in Scheme 1) for LA to GVL transformation under solvent free conditions at 100 °C and 50 bar H<sub>2</sub> pressure and observed a TOF of 144 h<sup>-1</sup>.<sup>39</sup> More recently, Hou *et al.* tested a ruthenium hydride complex (14 in Scheme 1) for solvent assisted hydrogenation of methyl levulinate to GVL at 100 °C and 10 bar H<sub>2</sub> pressure.<sup>40</sup> They used tetrabutylammonium acetate (TBAOAc) as the solvent, which not only acted as a reaction medium but also coordinated to Ru resulting in the formation of Ru-OAc species, which further helped in the activation of H<sub>2</sub>. Some iron complexes (15 and 16 in Scheme 1) were also reported for the transfer hydrogenation<sup>41</sup> and direct hydrogenation<sup>42</sup> of LA and alkyl levulimates. LA to GVL transformation was also explored using *in situ* generated Fe-nanoparticles from the Fe<sub>3</sub>(CO)<sub>12</sub> precursor in the presence of imidozyl, pyridine, triethylamine or KOH at 180 °C yielding GVL in good yield (92%).<sup>43</sup> Similarly, cobalt based catalysts using Co(BF<sub>4</sub>)<sub>2</sub>·6H<sub>2</sub>O and a tetradeятate phosphine ligand P(CH<sub>2</sub>-CH<sub>2</sub>PPh<sub>2</sub>)<sub>3</sub> at 100 °C and 1 bar H<sub>2</sub> in 1,3-dimethyl-2-imidazolidinone<sup>44</sup> and Ni based catalysts using Ni(OAc)<sub>2</sub>·4H<sub>2</sub>O and the triphos ligand at 1 atm H<sub>2</sub> pressure<sup>45</sup> were also utilized for the hydrogenation of LA to GVL. Recently, Gracia *et al.* reported a [(dippe)Ni(COD)] catalyst (17 in Scheme 1) for transfer hydrogenation of LA in a biphasic solvent system (H<sub>2</sub>O: THF (5 : 1)) at 120 °C using formic acid as the hydrogen source.<sup>46</sup> More recently, the same group reported a homogeneous Cu(I) based catalytic system comprising [(PPh<sub>3</sub>)<sub>2</sub>Cu(NO<sub>3</sub>)] and the

dippe ligand for the hydrogenation of LA at 140 °C and 20.6 bar H<sub>2</sub> pressure in hexane and yielded >99% GVL.<sup>47</sup> A titanium based catalyst [Cp<sub>2</sub>TiCl<sub>2</sub>] (18 in Scheme 1) is also reported for the hydrogenation of LA at 100 °C and 6.8 bar H<sub>2</sub>.<sup>48</sup>

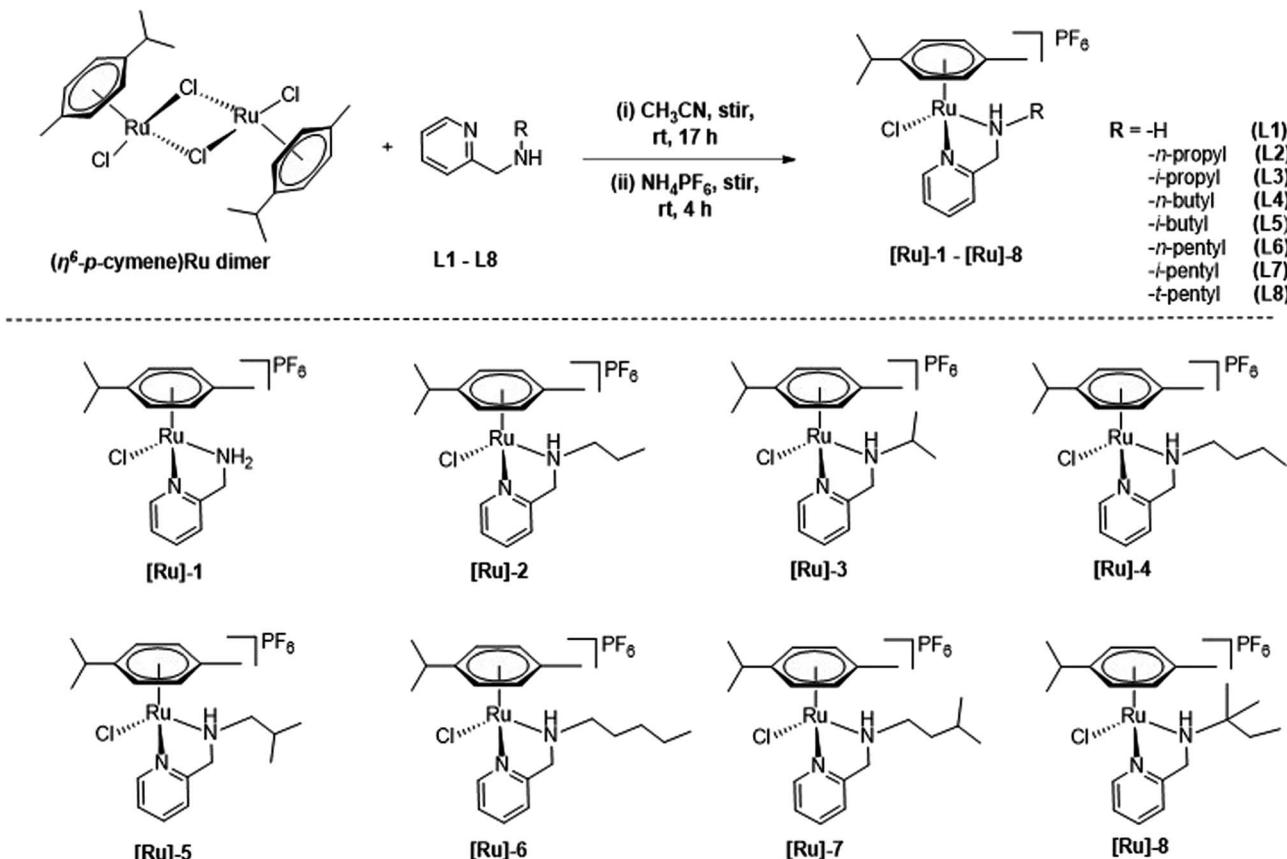
It is thus evident from the rich literature reports that most of the catalytic systems explored for LA to GVL transformation either use external H<sub>2</sub> gas (as high as 100 bar) or organic hydrogen donors such as formic acid, isopropanol, methanol, ethanol, or glycerol (Scheme 1 and Table S1†). Utilizing organic hydrogen donors eliminates the need for pressurized hydrogen gas in pressure reactors. Notably, Guo *et al.* demonstrated that formic acid produced during the transformation of 5-hydroxymethylfurfural (5-HMF) to LA can serve as a hydrogen source for further hydrogenation of LA to GVL.<sup>31</sup> However, a wide range of molecular catalysts explored for the transformation of LA to GVL using HCOOH require high reaction temperatures (as high as 150 °C) (Scheme 1 and Table S1†). Therefore, it is desirable to explore molecular catalysts which can avoid the use of elevated reaction temperatures and higher H<sub>2</sub> pressures for the transformation of LA to GVL. In this regard, herein, we report an efficient transformation of LA to GVL at a lower reaction temperature over water-soluble ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine-based molecular catalysts. A series of water-soluble ( $\eta^6$ -*p*-cymene)Ru(II) complexes containing N<sub>amine</sub> substituted pyridylamine ligands are synthesized and characterized, and the X-ray molecular structure of the complexes [Ru]-4, [Ru]-5, and [Ru]-8 were determined. The synthesized complexes catalyzed the hydrogenation of biomass-derived LA to GVL using HCOOH as a hydrogen source under milder reaction conditions (80 °C) in water. Advantageously, using pyridylamine ligands with substituted N<sub>amine</sub> enabled us to effectively explore and tune the steric bulkiness and basicity of N<sub>amine</sub> of the synthesized Ru complexes and hence the catalytic activity for the transformation of LA to GVL. Mechanistic investigations using mass and NMR studies were performed to identify the crucial catalytic intermediate species involved in the transformation of LA to GVL. Moreover, the optimized methodology for LA to GVL was scaled up to the gram-scale.

## Results and discussion

### Synthesis and characterization of ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine complexes ([Ru]-1-[Ru]-8)

Cationic ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine complexes [Ru]-1-[Ru]-8 having *N*-pyridylamine ligands L1-L8 were synthesized by treating  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  with the respective N<sub>amine</sub> substituted pyridylamine ligand in acetonitrile at room temperature (Scheme 2), following our previously reported process.<sup>49</sup> The synthesized ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine complexes vary in terms of the substituents at N<sub>amine</sub> ([Ru]-1: N<sub>amine</sub>H<sub>2</sub>; [Ru]-2: N<sub>amine</sub>-*n*-propyl; [Ru]-3: N<sub>amine</sub>-*i*-propyl; [Ru]-4: N<sub>amine</sub>-*n*-butyl; [Ru]-5: N<sub>amine</sub>-*i*-butyl; [Ru]-6: N<sub>amine</sub>-*n*-pentyl; [Ru]-7: N<sub>amine</sub>-*i*-pentyl; [Ru]-8: N<sub>amine</sub>-*t*-pentyl). Varying the substituents at the N<sub>amine</sub> group may play a crucial role in tuning the electronic and steric properties and hence further tune the catalytic performance of the synthesized complexes [Ru]-1-[Ru]-8 for the transformation of LA to GVL.





Scheme 2 Synthesis of  $(\eta^6\text{-}p\text{-cymene})\text{Ru(II)}$  complexes  $[\text{Ru}]\text{-1}\text{-}[\text{Ru}]\text{-8}$ , containing pyridylamine ligands L1–L8.

The spectro-analytical analysis of the synthesized complexes corroborated well with the proposed structures. The  $^1\text{H}$  NMR of the  $[\text{Ru}]\text{-1}\text{-}[\text{Ru}]\text{-8}$  displayed a similar trend of downfield chemical shift (8.93–9.12 ppm) as compared to that of the free pyridylamine ligands (8.42–8.48 ppm) for the ortho  $-\text{CH}$  protons.<sup>49</sup> The significant downfield shifts for the  $-\text{CH}_2$  protons indicated the ligand coordination to the ruthenium metal center. Upon introducing an electron rich alkyl substituent at the aminic nitrogen ( $\text{N}_{\text{amine}}$ ), a significant upfield shift of resonance is observed for the ortho  $-\text{CH}$  protons of the pyridine ring from 9.12 ppm in  $[\text{Ru}]\text{-1}$  to 8.93–9.03 ppm in  $[\text{Ru}]\text{-2}\text{-}[\text{Ru}]\text{-8}$ . The coordination of  $(\eta^6\text{-}p\text{-cymene})$  to the Ru(II) center is also confirmed by the presence of peaks for aromatic protons in the range of 5.65–5.92 ppm, the methyl proton in the field of 1.88–1.98 ppm, and the methine and methyl protons of the isopropyl group, respectively in the ranges of 2.59–3.65 and 1.00–1.12 ppm.<sup>49</sup> The counter anion  $\text{PF}_6^-$  resonated at  $-144$  ppm in the  $^{31}\text{P}$  NMR spectrum for  $[\text{Ru}]\text{-1}\text{-}[\text{Ru}]\text{-8}$ . The characteristic isotopic pattern of ruthenium was observed with a significantly intense base peak at  $[\text{M}]^+$  for all the synthesized complexes  $[\text{Ru}]\text{-1}\text{-}[\text{Ru}]\text{-8}$ .

Furthermore, to confirm the molecular identity of  $(\eta^6\text{-}p\text{-cymene})\text{Ru(II)}$ -pyridylamine complexes, we tried to obtain single crystals using various methods, including the slow evaporation of methanol-dichloromethane (1 : 1) solution of the respective complexes, but we were able to grow the crystals only for the

complexes  $[\text{Ru}]\text{-4}$ ,  $[\text{Ru}]\text{-5}$ , and  $[\text{Ru}]\text{-8}$ . The single crystal X-ray diffraction studies inferred that the complexes  $[\text{Ru}]\text{-4}$ ,  $[\text{Ru}]\text{-5}$  and  $[\text{Ru}]\text{-8}$ , respectively crystallize in the triclinic ( $P\bar{1}$  space group), monoclinic ( $P2_1/c$  space group) and orthorhombic ( $Pbca$  space group) crystal systems. The solid-state structure clearly demonstrates the pseudo-octahedral geometry for all the complexes which is characteristic of half-sandwich arene-ruthenium complexes. In typical piano-stool geometry around the ruthenium metal center, the three legs are occupied by the bis-chelating *N*-pyridylamine ligand coordinated to the ruthenium metal center in a  $\kappa^2$ -bidentate fashion through pyridine ( $\text{N}_{\text{py}}$ ) and amine nitrogen ( $\text{N}_{\text{amine}}$ ), and the chloro ligand ( $-\text{Cl}$ ) while the arene ring is placed at the apex of the structure in a  $\eta^6$  manner. The displacement of the  $(\eta^6\text{-}p\text{-cymene})$  ring from the ruthenium metal center for the synthesized complexes lies in the range of 1.67–1.69 Å. The  $\text{Ru-N}_{\text{py}}$  and  $\text{Ru-N}_{\text{amine}}$  bond distances are also in the range of 2.07–2.19 Å and are comparable to those for analogous *N*-pyridylamine arene–ruthenium complexes.<sup>49,50</sup>  $\text{Ru-Cl}$  bond distances are observed in the range of 2.39–2.40 Å. The  $\text{N}_{\text{py}}\text{-Ru-N}_{\text{amine}}$  bond angles lie in the range of 76.1–78.1°, while  $\text{N}_{\text{py}}\text{-Ru-Cl}$  and  $\text{N}_{\text{amine}}\text{-Ru-Cl}$  bond angles are observed between 83.8–85.6° and 82.5–84.6°, respectively. The bond angle between  $\text{N}_{\text{py}}\text{/N}_{\text{amine}}$  and ruthenium metal with the centroid of the  $\eta^6$ -arene ring ( $C_{\text{ct}}$ ) is in the range of 129.2–137.3° for complexes  $[\text{Ru}]\text{-4}$ ,  $[\text{Ru}]\text{-5}$  and  $[\text{Ru}]\text{-8}$ , which further supports the piano-stool geometry of the  $(\eta^6\text{-}p\text{-cymene})\text{Ru(II)}$ -

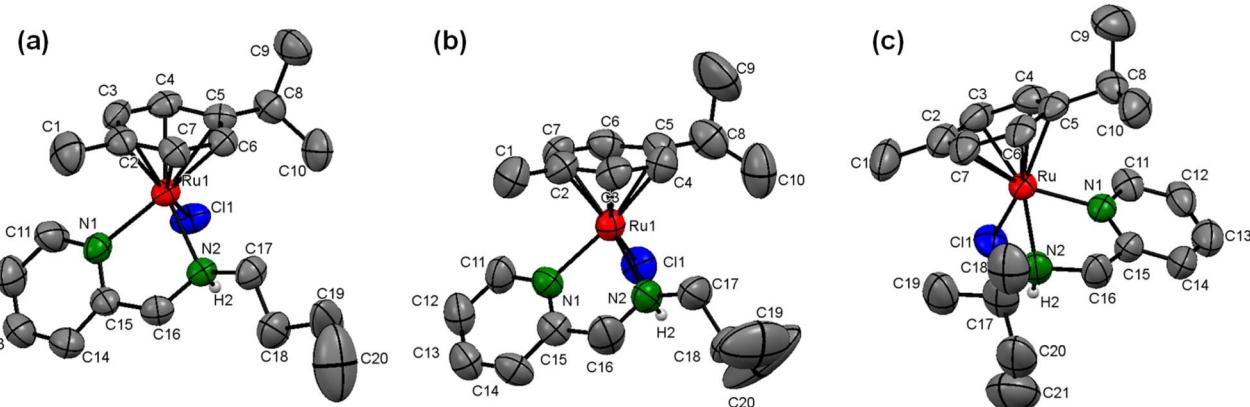


Fig. 1 X-ray crystal structures of complexes (a) [Ru]-4, (b) [Ru]-5, and (c) [Ru]-8 with 30% ellipsoid probability. Counteranions ( $\text{PF}_6^-$ ) and H atoms (except for those on N) are omitted for the sake of clarity.

pyridylamine complexes (Fig. 1). The crystal refinement data and some other important bond lengths and bond angles are summarized in Tables S2–S4<sup>†</sup> for the complexes [Ru]-4, [Ru]-5 and [Ru]-8.

### Catalytic transformation of LA to GVL

At an outset, the catalytic activity of the synthesized ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine complexes ([Ru]-1–[Ru]-8) for the transformation of biomass derived LA to GVL was investigated using formic acid as a hydrogen source along with triethylamine as a base at 80 °C in water (Table 1).

Results inferred that ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylmethyl amine ([Ru]-1) exhibited only 14% conversion of LA to GVL with 80% selectivity for GVL and 20% for 4-hydroxypentanoic acid (HPA) (Table 1, entry 1). A significantly enhanced conversion (>99%) of LA selectively to GVL (yield 87%) with a TOF of 20 h<sup>-1</sup>

was achieved over the [Ru]-2 containing *N*<sub>amine</sub>-*n*-propyl substituted pyridyl methyl amine ligand in 2 h (Table 1, entry 2). In contrast to [Ru]-2, the [Ru]-3 having *N*<sub>amine</sub>-*i*-propyl substituted pyridyl methyl amine displayed a very poor conversion (TOF 2.4 h<sup>-1</sup>) (Table 1, entry 3). Notably, enhancement in the catalytic activity (75–95% conversion of LA, 77–97% selectivity for GVL and a TOF of 15–19.4 h<sup>-1</sup>) was observed while performing the catalytic reaction over ( $\eta^6$ -*p*-cymene)Ru(II) complexes having other *N*-substituted pyridyl methyl amine ligands ([Ru]-4: *N*<sub>amine</sub>-*n*-butyl; [Ru]-5: *N*<sub>amine</sub>-*i*-butyl; [Ru]-6: *N*<sub>amine</sub>-*n*-pentyl; [Ru]-7: *N*<sub>amine</sub>-*i*-pentyl; [Ru]-8: *N*<sub>amine</sub>-*t*-pentyl) (Table 1, entries 4–8). In contrast to the high activity observed for ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine, the precursor  $[(\eta^6$ -*p*-cymene)RuCl<sub>2</sub>]<sub>2</sub> ([Ru]-PC) exhibited almost negligible activity (Table 1, entry 9). The results show that the basicity of *N*<sub>amine</sub> as well as steric crowding played an important role in tuning the catalytic activity of these ( $\eta^6$ -*p*-cymene)Ru(II)-pyridylamine complexes.

Table 1 Optimization table for catalytic transformation of biomass derived LA to GVL in water<sup>a</sup>

Entry	Catalyst	Ru catalyst (2.5 mol%)			Conv. <sup>b</sup> (%)	Sel. <sup>b</sup> (%) GVL/HPA	TOF (h <sup>-1</sup> )
		HCOOH (6 equiv.)	Et <sub>3</sub> N (1 equiv.), T (°C),	t (h)			
1	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sub>2</sub> )] <sup>+</sup> [Ru]-1	80	2	14	80/20		2.8
2	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>i</sup> pr)] <sup>+</sup> [Ru]-2	80	2	>99	>99(87%) <sup>c</sup> /-		20
3	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>i</sup> pr)] <sup>+</sup> [Ru]-3	80	2	12	96/4		2.4
4	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>n</sup> bu)] <sup>+</sup> [Ru]-4	80	2	97	93/7		19.4
5	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>i</sup> bu)] <sup>+</sup> [Ru]-5	80	2	75	77/23		15
6	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>n</sup> pen)] <sup>+</sup> [Ru]-6	80	2	95	97/3		19
7	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>i</sup> pen)] <sup>+</sup> [Ru]-7	80	2	93	92/8		18.6
8	$[(\eta^6$ - <i>p</i> -cymene)RuCl( $\kappa^2$ -pyNH <sup>t</sup> pen)] <sup>+</sup> [Ru]-8	80	2	95	95/5		19
9	$[(\eta^6$ - <i>p</i> -cymene)RuCl <sub>2</sub> ] <sub>2</sub> [Ru]-PC	80	2	4	n.d.		0.8

<sup>a</sup> Reaction conditions: Ru catalyst (2.5 mol%), LA (1 mmol), HCOOH (6 mmol), Et<sub>3</sub>N (1 mmol), and water (5 mL). <sup>b</sup> Conversion and selectivity determined by <sup>1</sup>H NMR. <sup>c</sup> Isolated yield, TOF – turnover frequency, and n.d. – not determined.



The strong interaction of the  $\text{NH}_2$  group with the solvent molecule ( $\text{H}_2\text{O}$ ) in **[Ru]-1** permanently retarded the availability of the  $-\text{NH}$  group for the reaction, leading to lower activity. Notably, the basicity of  $\text{N}_{\text{amine}}$  increases with increase in the alkyl chain length or branching in the ligands.<sup>40,49–52</sup> However, with increase in the carbon chain length or branching the steric hindrance at  $\text{N}_{\text{amine}}$  may also increase. For instance, in the case of **[Ru]-3** having the  $\text{N}_{\text{amine}}$ -*i*-propyl substituted pyridylmethyl amine ligand, the methyl groups are placed closer to  $\text{N}_{\text{amine}}$ , which increased the steric crowding resulting in the lower activity of **[Ru]-3** as compared to **[Ru]-2**. Furthermore, the increase in the *N*-substituted alkyl chain length or branching causes increase in the steric crowding around the  $-\text{NH}$  group, which is expected to play a crucial role in the observed trend in the catalytic activity of **[Ru]-4**–**[Ru]-8** for LA to GVL transformation (Fig. S1†).<sup>40,49,50</sup> From single crystal X-ray crystallography, the  $\text{Ru-N}_{\text{amine}}$  bond length in **[Ru]-4** ( $\text{Py-NH}^{\text{H}}\text{bu}$ ) is shorter than that of **[Ru]-5** ( $\text{Py-NH}^{\text{H}}\text{bu}$ ) and **[Ru]-8** ( $\text{Py-NH}^{\text{H}}\text{pen}$ ). Moreover, the  $\text{N}_{\text{amine}}-\beta\text{C}$  and  $\text{N}_{\text{amine}}-\gamma\text{C}$  bond lengths in **[Ru]-4**, **[Ru]-5**, and **[Ru]-8** suggested more crowding around  $\text{N}_{\text{amine}}$  for **[Ru]-5** ( $\text{N}_{\text{amine}}-\beta\text{C}$  2.540 Å and  $\text{N}_{\text{amine}}-\gamma\text{C}$  3.181 Å) and **[Ru]-8** ( $\text{N}_{\text{amine}}-\beta\text{C}$  2.521 Å and  $\text{N}_{\text{amine}}-\gamma\text{C}$  3.048 Å) as compared to **[Ru]-4** ( $\text{N}_{\text{amine}}-\beta\text{C}$  2.544 Å and  $\text{N}_{\text{amine}}-\gamma\text{C}$  3.895 Å). Also, it was reported in the literature that when more electron rich substituents are attached to the metal center, transfer hydrogenation using formic acid is facilitated.<sup>53</sup> Hence, in accordance with these reports and our findings, it is clear that the increased basicity due to different steric and electronic properties of different ligands attached to  $\text{N}_{\text{amine}}$  catalyzed the dehydrogenation of formic acid which in turn increased the rate of hydrogenation of LA to GVL.

Further optimization of the reaction conditions to get the best suited reaction conditions for LA to GVL was performed with **[Ru]-2**. Though the reaction performed with 2.5 mol% catalyst loading yields the best result with a TOF value of  $20\text{ h}^{-1}$ , reactions with lower catalyst loadings of 1 and 0.25 mol% were also performed and we achieved 98 and 92% conversion of LA to GVL in a longer duration (Table S5†). The results showed that upon performing the reaction at lower temperatures (below 80 °C), lower conversion was observed and while increasing the reaction temperature to 90 and 100 °C, the conversion of LA increased (Table S6†). Over **[Ru]-2**, the estimated activation energy was found to be 46.89 kJ mol<sup>−1</sup> (Fig. S2†). The presence of a base is crucial for the activation and deprotonation of  $\text{HCOOH}$  to form formate that can coordinate to the metal centre and forms the Ru-formato species, which is an important intermediate to initiate the transfer hydrogenation reactions. Unlike other inorganic bases, transfer hydrogenation is more facile with  $\text{Et}_3\text{N}$ , presumably due to its ability to facilitate the reversible proton abstraction/release from  $\text{HCOOH}$ .<sup>54</sup> Therefore, change in the base from  $\text{Et}_3\text{N}$  to inorganic bases such as  $\text{K}_3\text{PO}_4$ ,  $\text{NaOH}$ ,  $\text{NaHCO}_3$  and  $\text{HCOONa}$  could not improve the conversion of LA to GVL significantly, while in the absence of the base ( $\text{Et}_3\text{N}$ ) reaction could not occur (Table S7†). The reaction performed for various time intervals showed that the reaction proceeds *via* the formation of 4-hydroxypentanoic acid (4-HPA) as an intermediate with continuous consumption of LA (Table S8† and Fig. 2).

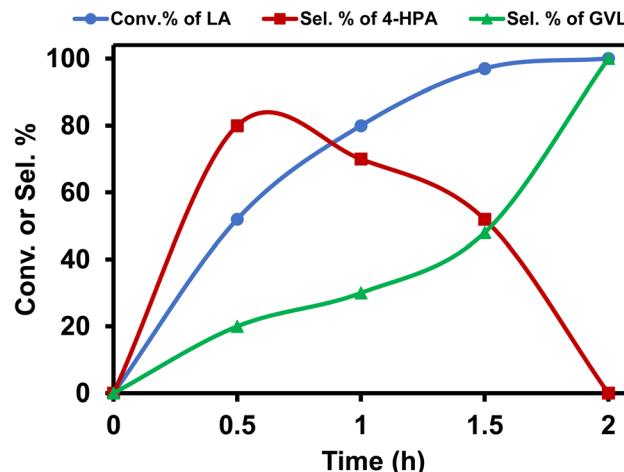


Fig. 2 Time dependent reaction profile for catalytic transformation of LA to GVL over the **[Ru]-2** catalyst. Reaction conditions: **[Ru]-2** (2.5 mol%), LA (1 mmol),  $\text{HCOOH}$  (6 mmol),  $\text{Et}_3\text{N}$  (1 mmol), water (5 mL), 80 °C, 0–2 h.

Furthermore, pH dependent  $^1\text{H}$  NMR studies showed that lower pH is favorable for the high selectivity of GVL. It was observed that at pH 6.1, ( $\text{HCOOH} : \text{Et}_3\text{N} = 6 : 1$ ), 4-HPA which is the hydrogenated form of LA, was observed. With the further decrease in the pH to 4.7, 3.6, and 2.7, the intensity of peaks corresponding to 4-HPA decreased, while those corresponding to GVL became more prominent (Fig. 3 and Table S9†). We observed complete conversion of LA with complete selectivity for GVL in 2 h in the pH range of 2.5–2.7. The identification of  $^1\text{H}$  NMR peaks corresponding to 4-HPA, further reinforced that the transformation of LA to GVL proceeded through the intermediate 4-HPA.

To further investigate a plausible mechanistic pathway for the transformation of LA to GVL over the **[Ru]-2** catalyst, we performed extensive mass studies to identify several intermediate species involved during the course of reaction (Fig. 4). In a control experiment, the **[Ru]-2** catalyst was stirred for 5–10 min in water where a prominent peak at  $m/z = 385.1$  corresponding to the species  $[(\eta^6\text{-}p\text{-cymene})\text{Ru}(\kappa^2\text{-PyNpr})]^+$  (**[Ru]-2A**) having a vacant coordination site on Ru was observed. We further noticed that the formation of species **[Ru]-2A** was independent of the presence of  $\text{HCOOH}$  and  $\text{Et}_3\text{N}$ . In the subsequent step, the addition of 6 mmol of  $\text{HCOOH}$ , 1 mmol of  $\text{Et}_3\text{N}$  and 1 mmol of LA followed by stirring at 60 °C for 30 minutes resulted in the appearance of new peaks at  $m/z = 431.1$  and  $m/z = 501.2$  corresponding to the  $[(\eta^6\text{-}p\text{-cymene})\text{Ru}(\kappa^2\text{-PyNHpr})(\text{HCOO})]^+$  (**[Ru]-2B**) and  $[(\eta^6\text{-}p\text{-cymene})\text{Ru}(\kappa^2\text{-PyNHpr})(\text{levulinate})]^+$  (**[Ru]-2C**) coordinated species.

Based on extensive mass studies, we anticipated that the generation of active catalytic species **[Ru]-2A** from **[Ru]-2** is the first step in the mechanistic cycle for the transformation of LA to GVL. Furthermore, the addition of formic acid,  $\text{Et}_3\text{N}$ , and LA resulted in the formation of the Ru-formato (**[Ru]-2B**) and Ru-levulinate (**[Ru]-2C**) species. The **[Ru]-2B** species then undergoes decarboxylation to form the Ru-hydrido species (**[Ru]-2D**), which is responsible for the transfer hydrogenation of LA to 4-HPA (identified as an intermediate during the catalytic reaction, Fig. 2) and the product GVL was finally formed by



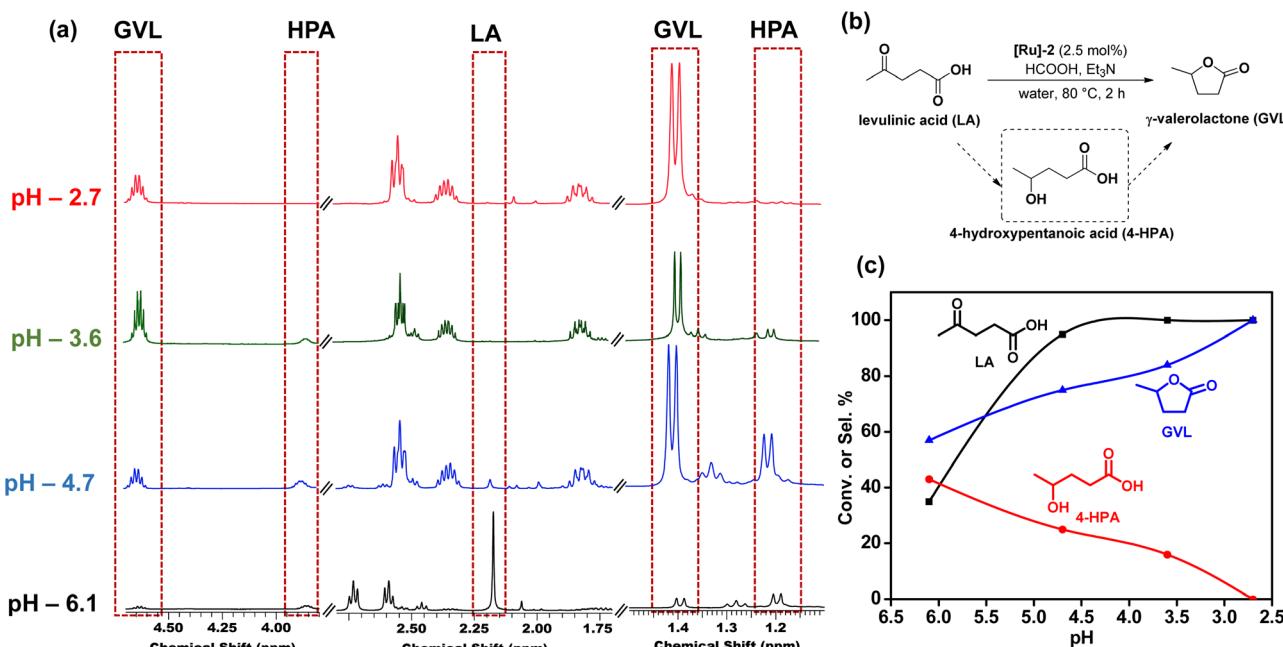


Fig. 3 (a)  $^1\text{H}$  NMR for pH dependent transformation in  $\text{CDCl}_3$ , (b) schematic representation, and (c) pH dependent transformation of biomass derived LA to GVL through the formation of intermediate 4-HPA over the  $[\text{Ru}]\text{-2}$  catalyst. Reaction conditions:  $[\text{Ru}]\text{-2}$  (2.5 mol%), LA (1 mmol), HCOOH (1–6 mmol),  $\text{Et}_3\text{N}$  (1 mmol), water (5 mL),  $80^\circ\text{C}$ , 2 h.

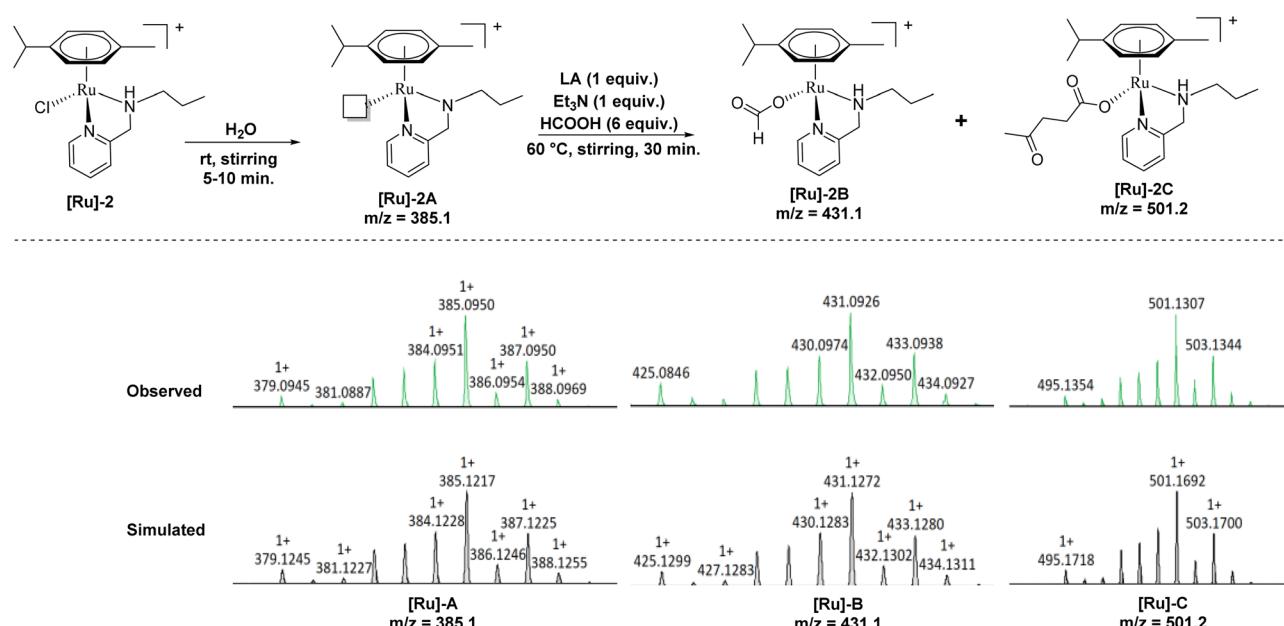


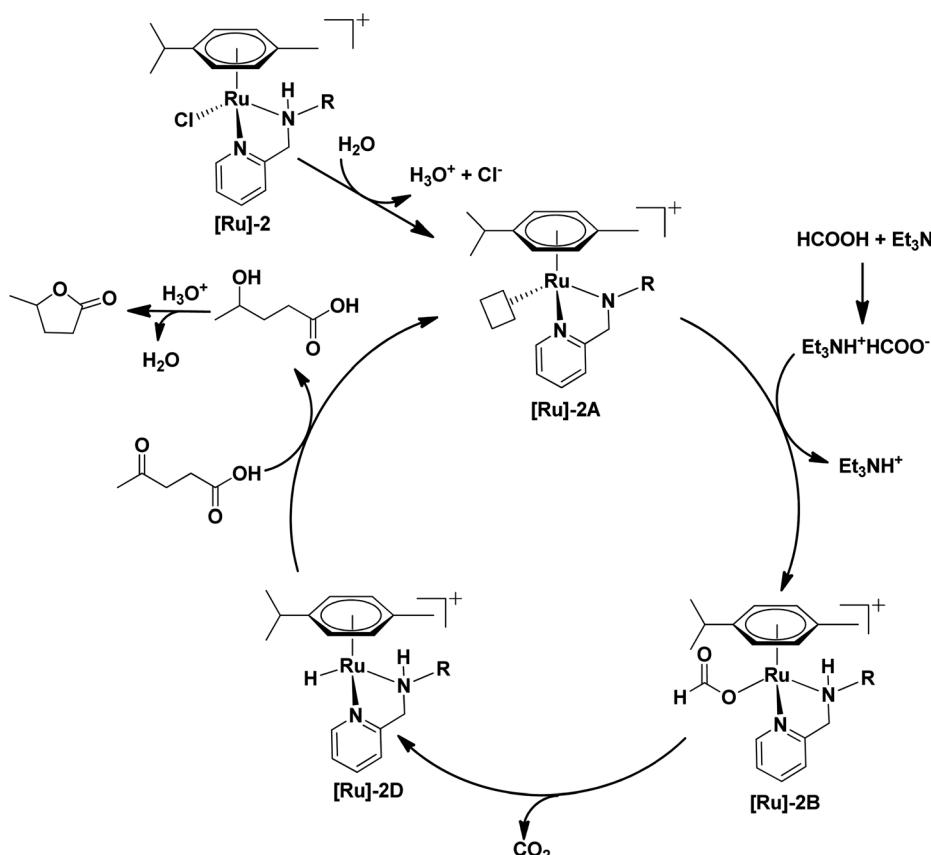
Fig. 4 Ru-formato and Ru-levulinate species observed during mass investigation of the catalytic reaction aliquots.

dehydration and cyclization of 4-HPA (Scheme 3). Upon further investigation, we noticed that the formation of the  $[\text{Ru}]\text{-2C}$  species depends on the ratio of HCOOH :  $\text{Et}_3\text{N}$ . Notably, mass studies inferred that high intensity mass peaks of the  $[\text{Ru}]\text{-2C}$  species appeared for a HCOOH :  $\text{Et}_3\text{N}$  ratio of 3 : 1, while the intensity of this species greatly reduced with a 6 : 1 ratio of HCOOH :  $\text{Et}_3\text{N}$  (Fig. S3 $\dagger$ ). Therefore, presumably the  $[\text{Ru}]\text{-2C}$  species is not directly involved in the LA hydrogenation to GVL.

### Catalyst recyclability and gram-Scale reaction for the transformation of LA to GVL

Notably, the  $[\text{Ru}]\text{-2}$  catalyst also displayed good recyclability for five consecutive catalytic runs, where complete conversion was achieved for the initial three cycles with some decline in the conversion for the fourth and fifth catalytic runs to 82 and 74%, respectively (Table S10 and Fig. S4 $\dagger$ ). We performed the ICP-AES





**Scheme 3** Plausible mechanism for the transformation of biomass derived LA to GVL.

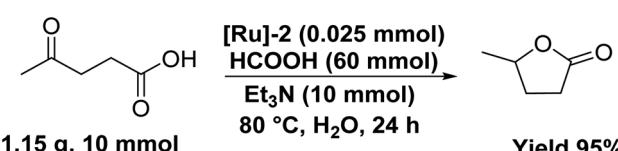
analysis of the organic and aqueous phases of the reaction mixture. The ICP-AES analysis of the organic portion inferred a loss of 10–12 ppm Ru in each catalytic run, which may account for the observed loss in catalytic activity during recyclability experiments. Furthermore, the aqueous layer inferred a prominent mass peak at *m/z* 385 (corresponding to [Ru]-2A) (Fig. S5†), suggesting the high stability of the catalyst under catalytic reaction conditions. Advantageously, the studied catalytic system can be scaled up to the gram-scale transformation of LA to GVL over the [Ru]-2 (0.025 mmol) catalyst to yield 95% GVL (TON 380) with complete conversion of LA in 24 h at 80 °C (Scheme 4).

#### Catalytic one-pot two step transformation of furfural (FAL) to GVL

We also explored the catalytic one-pot transformation of furfural (FAL) to GVL over the [Ru]-2 catalyst (Fig. 5a). For this,

to an aqueous suspension of [Ru]-2 (2.5 mol% in 5 mL water) in a graduated 10 mL test tube, 1 mmol FAL was added and heated at 80 °C in an oil bath for 12 h in the presence of 12 equivalents of formic acid. FAL is first hydrogenated to furfuryl alcohol (FA) and then acid catalyzed ring opening of FA resulted in the formation of LA, which is further confirmed by the <sup>1</sup>H NMR analysis of the reaction aliquot (Fig. 5b). HCOOH served as both a hydrogenating agent and an acid source during the progress of reaction. To the same reaction vessel, 4 equiv. of HCOOH and 1 equiv. of Et<sub>3</sub>N were further added and the reaction mixture was allowed to stir at 80 °C for 4 h. <sup>1</sup>H NMR analysis confirmed the formation of GVL as the product. Thus, we achieved complete conversion of FAL with 100% selectivity for GVL (yield 47%). The observed lower yield of GVL can be attributed to the poor stability of FAL in aqueous medium. These results demonstrated that the [Ru]-2 catalyst can be used for the production of GVL from the biomass platform chemical FAL with the aid of HCOOH in one pot without any need for the isolation of LA.

Therefore, our findings inferred that the [Ru]-2 catalyst demonstrated high catalytic activity for the complete and selective transformation of LA to GVL in water under much milder reaction conditions (80 °C) using HCOOH as a H<sub>2</sub> source, as compared to previously reported Ru based catalytic systems (Table S1†). For instance, the literature revealed the use of Ru-phosphine-based catalysts that are active only at high H<sub>2</sub>



**Scheme 4** Gram scale catalytic transformation of LA to GVL over the [Ru]-2 catalyst.



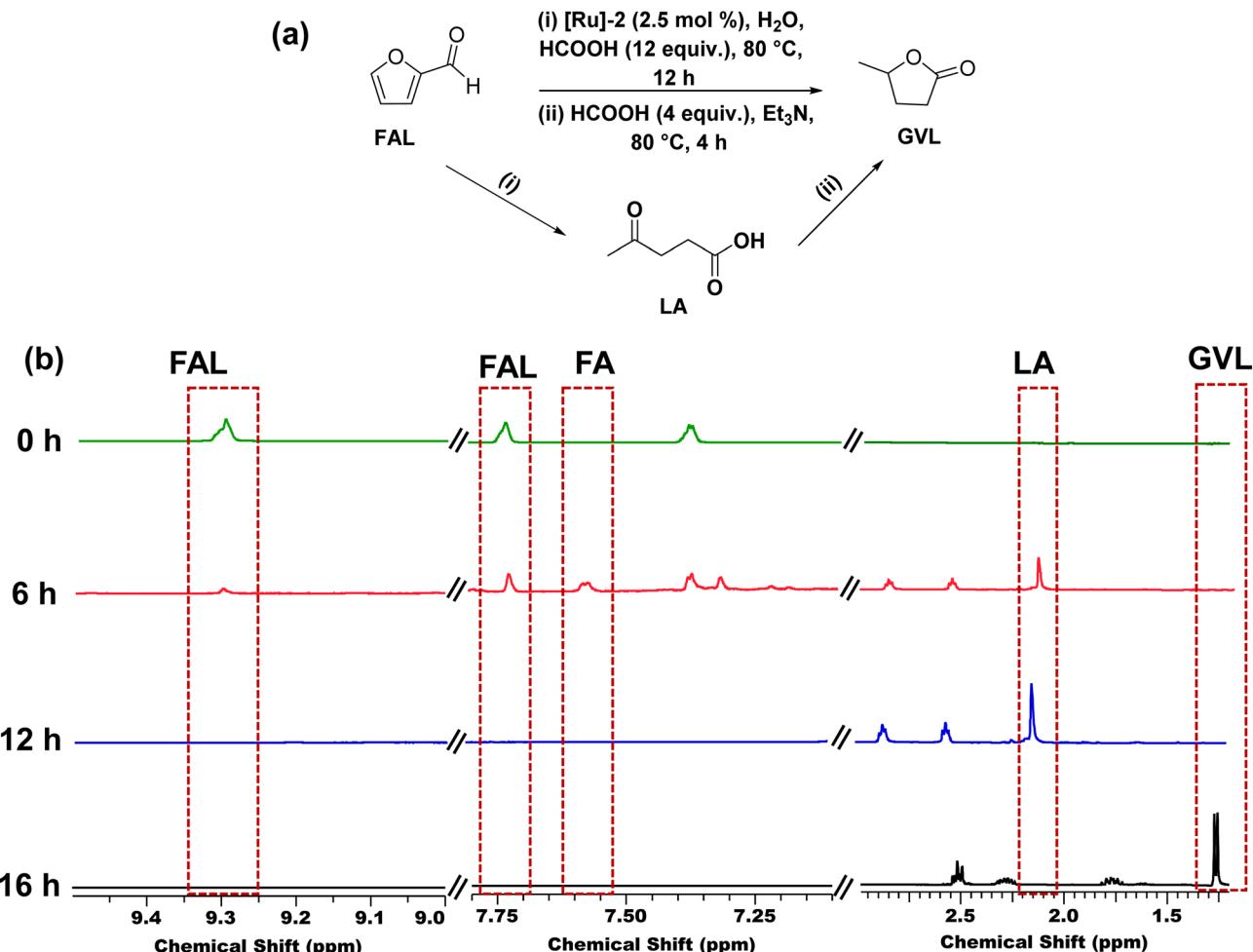


Fig. 5 (a) Schematic representation and (b) time-dependent <sup>1</sup>H NMR (in D<sub>2</sub>O) of one pot-two step catalytic transformation of FAL to GVL over the [Ru]-2 catalyst. Reaction conditions: FAL (1 mmol), [Ru]-2 (2.5 mol%), HCOOH (12 mmol), H<sub>2</sub>O (5 mL), 80 °C, and 12 h, and after 12 h Et<sub>3</sub>N (1 mmol) and HCOOH (4 mmol) were added, and the reaction continued for another 4 h.

pressure (as high as 100 bar)<sup>37,39</sup> and high reaction temperature (as high as 160 °C) to achieve the hydrogenation of LA to GVL (Table S1†). Advantageously, we could achieve one-pot transformation of furfural to GVL over the [Ru]-2 catalyst, which clearly demonstrated the advancement of our catalytic system for future applications.

## Conclusions

We demonstrated herein a series of (*η*<sup>6</sup>-*p*-cymene)Ru(*ii*)-pyridylamine based complexes for the efficient catalytic transformation of biomass-derived LA to GVL in water using formic acid at 80 °C. The water soluble (*η*<sup>6</sup>-*p*-cymene)Ru(*ii*)-pyridylamine complexes with varying substituents at N<sub>amine</sub> were synthesized and their molecular structure was established using various characterization techniques including NMR, mass spectrometry and single crystal X-ray diffraction. The substituents at N<sub>amine</sub> of the pyridyl methyl amine ligands exerted a significant effect on the catalytic activity by tuning the steric crowding and the basicity at N<sub>amine</sub>. Among the studied complexes,  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-pyNH}^n\text{pr})]^+$  ([Ru]-2)

outperformed with complete conversion of LA selectively to GVL (87% yield) at 80 °C in 2 h. Detailed mass investigations and control experiments provided a detailed mechanistic insight by revealing the possible involvement of [Ru]-2A and [Ru]-2B (Ru-formato) species in the process of LA hydrogenation, where the ratio of HCOOH : Et<sub>3</sub>N was found to be crucial to generate the Ru-formato ([Ru]-2B) and subsequently the Ru-hydrido ([Ru]-2D) species to facilitate the transformation of LA to GVL in water. Furthermore, time dependent studies revealed that 4-hydroxypentanoic acid (4-HPA) is formed as an intermediate during the catalytic transformation of LA to GVL. [Ru]-2 also exhibited high recyclability up to 4 cycles for LA to GVL transformation. Moreover, the developed methodology was also scaled up to gram-scale transformation of LA to GVL, highlighting the practical application of the system.

## Experimental section

### Synthesis of complexes [Ru]-1–[Ru]-8

Complexes [Ru]-1–[Ru]-4 and [Ru]-7 are known in the literature and synthesized by our previously reported procedures.<sup>49</sup>



Complexes **[Ru]-5**, **[Ru]-6** and **[Ru]-8** are not known and synthesized using our previously reported procedures.<sup>49</sup> Typically, for the synthesis of **[Ru]-1**–**[Ru]-8** complexes,  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  was suspended in acetonitrile (25 mL), stirred for 30 min at room temperature, and *N*-pyridylamine ligands (**L1**–**L8**) were added. After stirring the reaction mixture for 17 h at room temperature,  $\text{NH}_4\text{PF}_6$  (3 equiv.) was added, and the solution was stirred for another 4 h at room temperature. Then, the solvent was removed under vacuum and the obtained residue was dissolved in a minimum amount of dichloromethane, filtered to remove unreacted  $\text{NH}_4\text{PF}_6$  salt, and an excess of diethyl ether was poured into the above filtered solution to precipitate out the desired complexes as yellow solids. Precipitates were washed with diethyl ether (3 × 10 mL) and dried in air. Single crystals of **[Ru]-4**, **[Ru]-5** and **[Ru]-8** were obtained by diffusion of diethyl ether into methanolic solution of respective complexes. The CCDC deposition numbers of the complexes **[Ru]-4**, **[Ru]-5**, and **[Ru]-8** are 1971292, 1971295 and 1971294, respectively.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-}\text{pyNH}_2)]\text{PF}_6$ (**[Ru]-1**)

Complex **[Ru]-1** was synthesized by stirring 2-(aminomethyl)-pyridine (**L1**) (0.083 g, 0.8 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL) at room temperature.  $\text{NH}_4\text{PF}_6$  (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Yellow-brown solid, yield 0.165 g (63%). <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 9.11 (d, 1H, *J* = 4.00 Hz), 7.96 (t, 1H, *J* = 8.00 Hz), 7.53 (t, 1H, *J* = 8.00 Hz), 7.51 (d, 1H, *J* = 4.00 Hz), 5.91 (d, 1H, *J* = 8.00 Hz), 5.85 (d, 1H, *J* = 8.00 Hz), 5.72 (d, 1H, *J* = 8.00 Hz), 5.66 (d, 1H, *J* = 8 Hz), 4.45–4.39 (m, 1H), 4.21–4.07 (m, 1H), 2.74–2.66 (m, 1H), 1.95 (s, 3H), 1.11 (d, 6H, *J* = 4 Hz). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 161.88, 155.19, 139.78, 125.51, 121.79, 103.69, 98.66, 85.47, 83.71, 82.92, 82.35, 52.68, 30.87, 22.97, 22.07, 18.28. <sup>31</sup>P NMR (161.97 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) –144.19 (sep,  $\text{PF}_6^-$ ). MS (ESI): *m/z* calculated: 379.0150 [M]<sup>+</sup>, found: [M]<sup>+</sup> 379.0776.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-}\text{pyNH}^i\text{pr})]\text{PF}_6$ (**[Ru]-2**)

Complex **[Ru]-2** was synthesized by stirring *N*-(pyridine-2-ylmethyl)-propan-1-amine (**L2**) (0.083 g, 0.55 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL).  $\text{NH}_4\text{PF}_6$  (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Pale yellow solid, yield 0.255 g (90%). <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 9.03 (d, 1H, *J* = 4.00 Hz), 8.02–7.98 (t, 1H, *J* = 8.00 Hz), 7.62–7.60 (d, 1H, *J* = 8 Hz), 7.58–7.54 (t, 1H, *J* = 8 Hz), 5.88–5.85 (t, 2H, *J*<sub>1</sub> = 4 Hz, *J*<sub>2</sub> = 8 Hz), 5.83–5.82 (d, 1H, *J* = 4 Hz), 5.75–5.73 (d, 1H, *J* = 8 Hz), 4.35–4.18 (m, 2H), 3.43–3.40 (m, 1H), 3.29–3.27 (m, 1H), 2.62–2.56 (m, 1H), 1.90 (s, 3H), 1.74–1.64 (m, 2H), 1.09–1.04 (m, 6H), 0.98–0.94 (t, 3H, *J* = 8 Hz). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 159.45, 154.88, 139.22, 125.13, 121.62, 105.14, 96.20, 85.54, 83.85, 82.45, 81.60, 61.12, 59.00, 30.53, 22.15, 21.40, 21.13, 17.40, 11.25. <sup>31</sup>P NMR (161.97 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) –144.42 (sep,  $\text{PF}_6^-$ ). MS (ESI): *m/z* calculated: 421.0981 [M]<sup>+</sup>, found: 421.0976 [M]<sup>+</sup>.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-}\text{pyNH}^i\text{pr})]\text{PF}_6$ (**[Ru]-3**)

Complex **[Ru]-3** was synthesized by stirring *N*-(pyridine-2-ylmethyl)-propan-2-amine (**L3**) (0.083 g, 0.55 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$

(0.153 g, 0.25 mmol) in acetonitrile (25 mL).  $\text{NH}_4\text{PF}_6$  (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Pale yellow solid, yield 0.179 g (63%). <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 9.00 (d, 1H, *J* = 4.00 Hz), 8.02–7.98 (t, 1H, *J* = 8.00 Hz), 7.65 (d, 1H, *J* = 8.00 Hz), 7.58–7.55 (t, 1H, *J*<sub>1</sub> = 8.00 Hz, *J*<sub>2</sub> = 4.00 Hz), 5.91 (d, 1H, *J* = 4.00 Hz), 5.82 (s, 2H), 5.71 (d, 1H, *J* = 4.00 Hz), 4.32–4.20 (m, 2H), 3.85–3.82 (m, 1H), 2.11–2.07 (m, 1H), 1.98 (s, 3H), 1.38 (d, 3H, *J* = 4.00 Hz), 1.31 (d, 3H, *J* = 4.00 Hz), 1.09 (d, 3H, *J* = 8.00 Hz), 1.00 (d, 3H, *J* = 8.00 Hz). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 159.57, 154.86, 139.17, 125.09, 121.79, 105.28, 96.58, 86.19, 84.92, 82.41, 80.38, 58.60, 56.18, 30.45, 23.42, 21.75, 21.64, 20.26, 17.38. <sup>31</sup>P NMR (161.97 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) –144.19 (sep,  $\text{PF}_6^-$ ). MS (ESI): *m/z* calculated: 421.0981 [M]<sup>+</sup>, found: 421.0977 [M]<sup>+</sup>.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-}\text{pyNH}^i\text{bu})]\text{PF}_6$ (**[Ru]-4**)

Complex **[Ru]-4** was synthesized by stirring *N*-(pyridin-2-ylmethyl)-butan-1-amine (**L4**) (0.090 g, 0.55 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL).  $\text{NH}_4\text{PF}_6$  (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Pale yellow solid, yield 0.263 g (91%). <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 9.03–9.01 (d, 1H, *J* = 8 Hz), 8.02–7.98 (t, 1H, *J* = 8 Hz), 7.62–7.60 (d, 1H, *J* = 8 Hz), 7.58–7.55 (t, 1H, *J*<sub>1</sub> = 8 Hz, *J*<sub>2</sub> = 4 Hz), 5.88–5.86 (d, 2H, *J* = 8 Hz), 5.83–5.81 (d, 1H, *J* = 8 Hz), 5.75–5.73 (d, 1H, *J* = 8 Hz), 4.35–4.18 (m, 2H), 3.46–3.44 (m, 2H), 2.60–2.55 (m, 1H), 1.89 (s, 3H), 1.69–1.63 (m, 2H), 1.39 (m, 2H), 1.09–1.05 (t, 6H, *J* = 8 Hz), 0.95–0.92 (t, 3H, *J*<sub>1</sub> = 8 Hz, *J*<sub>2</sub> = 4 Hz). <sup>13</sup>C NMR (100 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) 159.43, 154.87, 139.21, 125.13, 121.62, 105.25, 95.96, 85.71, 83.84, 82.41, 81.57, 61.22, 57.00, 30.55, 30.12, 22.14, 21.09, 19.62, 17.35, 13.79. <sup>31</sup>P NMR (161.97 MHz, DMSO-*d*<sub>6</sub>): δ (ppm) –144.20 (sep,  $\text{PF}_6^-$ ). MS (ESI): *m/z* calculated: 435.1138 [M]<sup>+</sup>, found: 435.1112 [M]<sup>+</sup>.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-}\text{pyNH}^i\text{bu})]\text{PF}_6$ (**[Ru]-5**)

Complex **[Ru]-5** was synthesized by stirring 2-methyl-*N*-(pyridine-2-ylmethyl)-propan-1-amine (**L5**) (0.090 g, 0.55 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL).  $\text{NH}_4\text{PF}_6$  (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Pale yellow solid, yield 0.168 g (58%). <sup>1</sup>H NMR (400 MHz, methanol-*d*<sub>4</sub>): δ (ppm) 8.95–8.93 (d, 1H, *J* = 8 Hz), 7.95–7.91 (t, 1H, *J* = 8 Hz), 7.51–7.45 (m, 2H), 5.79–5.75 (t, 2H, *J* = 8 Hz), 5.68–5.66 (d, 1H, *J* = 8 Hz), 5.59–5.58 (d, 1H, *J* = 4 Hz), 4.40–4.35 (dd, 1H, *J* = 4 Hz), 4.25–4.18 (t, 1H, *J*<sub>1</sub> = 12 Hz, *J*<sub>2</sub> = 16 Hz), 3.38–3.28 (m, 2H), 2.66–2.56 (m, 1H), 2.18–2.11 (m, 1H), 1.97 (s, 3H), 1.13–1.08 (m, 6H), 0.99–0.97 (m, 6H). <sup>13</sup>C NMR (100 MHz, methanol-*d*<sub>4</sub>): δ (ppm) 160.45, 159.19, 140.63, 126.78, 123.08, 107.73, 98.44, 87.19, 85.30, 84.02, 83.12, 66.92, 62.37, 32.49, 27.94, 22.85, 21.62, 20.96, 20.15, 18.18. <sup>31</sup>P NMR (161.97 MHz, methanol-*d*<sub>4</sub>): δ (ppm) –144.58 (sep,  $\text{PF}_6^-$ ). MS (ESI): *m/z* calculated: 435.1138 [M]<sup>+</sup>, found: 435.1141 [M]<sup>+</sup>.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-}\text{pyNH}^i\text{pen})]\text{PF}_6$ (**[Ru]-6**)

Complex **[Ru]-6** was synthesized by stirring *N*-(pyridine-2-ylmethyl)-pentan-1-amine (**L6**) (0.098 g, 0.55 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL).  $\text{NH}_4\text{PF}_6$  (0.244 g, 1.5 mmol) was added and stirred for another

4 h. Pale yellow solid, yield 0.241 g (81%).  $^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) 9.03–9.01 (d, 1H,  $J$  = 8 Hz), 8.02–7.98 (t, 1H,  $J$  = 8 Hz), 7.62–7.60 (d, 1H,  $J$  = 8 Hz), 7.58–7.54 (t, 1H,  $J$  = 8 Hz), 5.88–5.86 (d, 2H,  $J$  = 8 Hz), 5.83–5.81 (d, 1H,  $J$  = 8 Hz), 5.74–5.73 (d, 1H,  $J$  = 4 Hz), 4.35–4.18 (m, 2H), 3.45–3.41 (m, 2H), 2.62–2.57 (m, 1H), 1.89 (s, 3H), 1.69–1.64 (m, 2H), 1.35–1.30 (m, 4H), 1.09–1.05 (t, 6H,  $J$  = 8 Hz), 0.90 (t, 3H).  $^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) 159.45, 154.87, 139.20, 125.12, 121.62, 105.21, 95.96, 85.71, 83.84, 82.43, 81.57, 61.19, 57.24, 30.55, 28.53, 27.75, 22.14, 21.96, 21.10, 17.35, 13.89.  $^{31}\text{P}$  NMR (161.97 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) –144.18 (sep, PF<sub>6</sub>). MS (ESI):  $m/z$  calculated: 449.1294 [M]<sup>+</sup>, found: 449.1266 [M]<sup>+</sup>.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-pyNH}^t\text{pen})]\text{PF}_6$ ([Ru]-7)

Complex [Ru]-7 was synthesized by stirring 3-methyl-*N*-(pyridine-2-ylmethyl)-butan-1-amine (**L7**) (0.098 g, 0.75 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL). NH<sub>4</sub>PF<sub>6</sub> (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Pale yellow solid, yield 0.187 g (63%).  $^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) 9.02–9.01 (d, 1H,  $J$  = 4 Hz), 8.00–7.98 (t, 1H,  $J$  = 8 Hz), 7.62–7.50 (m, 2H), 5.88 (m, 2H), 5.83–5.81 (d, 1H,  $J$  = 8 Hz), 5.74–5.73 (d, 1H,  $J$  = 4 Hz), 4.36–4.20 (m, 2H), 3.48–3.47 (m, 2H), 2.65–2.58 (m, 1H), 1.88 (s, 3H), 1.70–1.55 (m, 3H), 1.10–1.08 (t, 6H,  $J$  = 8 Hz), 0.94–0.92 (t, 6H,  $J$  = 8 Hz).  $^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) 159.40, 154.84, 139.20, 125.13, 121.63, 105.34, 95.78, 85.86, 83.81, 82.37, 81.61, 61.31, 55.67, 36.78, 30.58, 25.60, 22.57, 22.36, 22.17, 21.02, 17.30.  $^{31}\text{P}$  NMR (161.97 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) –144.08 (sep, PF<sub>6</sub>). MS (ESI):  $m/z$  calculated: 449.1294 [M]<sup>+</sup>, found: 449.1294 [M]<sup>+</sup>.

#### Synthesis of $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}(\kappa^2\text{-pyNH}^t\text{pen})]\text{PF}_6$ ([Ru]-8)

Complex [Ru]-8 was synthesized by stirring 2-methyl-*N*-(pyridine-2-ylmethyl)-butan-2-amine (**L8**) (0.098 g, 0.75 mmol) and  $[(\eta^6\text{-}p\text{-cymene})\text{RuCl}_2]_2$  (0.153 g, 0.25 mmol) in acetonitrile (25 mL). NH<sub>4</sub>PF<sub>6</sub> (0.244 g, 1.5 mmol) was added and stirred for another 4 h. Pale yellow solid, yield 0.258 g (86%).  $^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) 9.03–9.02 (d, 1H,  $J$  = 4 Hz), 8.77–8.66 (d, 1H,  $J_1$  = 40 Hz,  $J_2$  = 4 Hz), 8.02–8.00 (t, 1H, 8 Hz), 7.65–7.55 (m, 1H), 6.01–5.90 (m, 4H), 4.35–4.29 (m, 2H), 3.65–3.57 (m, 1H), 1.92 (s, 3H), 1.72–1.65 (m, 2H), 1.36–1.30 (d, 6H,  $J$  = 24 Hz), 1.06–1.00 (dd, 6H,  $J$  = 4 Hz), 0.90–0.86 (t, 3H,  $J$  = 8 Hz).  $^{13}\text{C}$  NMR (100 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) 159.50, 155.02, 139.23, 125.12, 122.08, 105.40, 96.41, 85.69, 84.55, 82.52, 79.79, 60.70, 55.40, 32.53, 30.53, 24.75, 24.57, 21.88, 17.18, 8.66.  $^{31}\text{P}$  NMR (161.97 MHz, DMSO- $d_6$ ):  $\delta$  (ppm) –144.20 (sep, PF<sub>6</sub>). MS (ESI):  $m/z$  calculated: 449.1294 [M]<sup>+</sup>, found: 449.1232 [M]<sup>+</sup>.

#### General procedure for the catalytic transformation of LA to GVL in water

All the reactions were performed in a 10 mL graduated reaction test tube using water as solvent. Typically for the catalytic reaction, 1.0 mmol of LA, 6 mmol of HCOOH, and 1 mmol of Et<sub>3</sub>N were added to an aqueous suspension of the catalyst (2.5 mol% in 5 mL water). The reaction test tube was fitted to a condenser, and the reaction mixture was stirred at 80 °C in an oil bath for a specified time. All the catalytic reactions were

performed using formic acid as the source of hydrogen without using any external hydrogen gas pressure. After reaction completion, the reaction mixture was extracted with diethyl ether (12 × 10 mL) and then acidified with 1 M HCl solution till pH 1. Then it was again extracted with ethyl acetate (6 × 10 mL). The combined organic portions were dried under reduced pressure. Conversion and selectivity were determined by  $^1\text{H}$  NMR.

#### Control experiments for mechanistic study

The [Ru]-2 (0.025 mmol) catalyst was dissolved in 2.5 mL of water and stirred for 5–10 minutes, and the analysis of the reaction aliquot by mass spectrometry revealed the formation of [Ru]-A after dissociation of one Cl. Furthermore, the addition of LA (1 mmol), Et<sub>3</sub>N (1 mmol) and HCOOH (6 mmol) followed by stirring for 30 min at 60 °C and the subsequent analysis of the reaction aliquot by mass spectrometry inferred the presence of the Ru-formato ([Ru]-B) and Ru-levulinate ([Ru]-C) species. Reaction aliquots were taken out at definite time intervals and analyzed by mass spectrometry to detect the active catalytic species involved in the transformation of LA to GVL. The pH dependent studies were carried out by varying the ratios of HCOOH : Et<sub>3</sub>N to obtain a specific pH and the reaction aliquots were analyzed by  $^1\text{H}$  NMR to know the presence of intermediates involved in the transformation of LA to GVL every 30 minutes.

#### General procedure for the catalytic one-pot two-step transformation of FAL to GVL in water

To an aqueous suspension of the catalyst (2.5 mol% in 5 mL water) in a 10 mL graduated reaction test tube fitted with a condenser, 1.0 mmol of FAL and 12 mmol of HCOOH were added and the reaction mixture was stirred at 80 °C for a specified time in an oil bath. Reaction progress was monitored by TLC and after complete conversion of FAL, 4 mmol of formic acid and 1 mmol of triethylamine were added to the same reaction vessel and the reaction mixture was continued to stir at 80 °C for another 4 h until complete conversion of LA to GVL was achieved.

#### Gram-scale and recyclability experiments

For the catalytic gram-scale reaction, to an aqueous suspension of [Ru]-2 catalyst (2.5 mol% in 5 mL water), 10 mmol (1.15 g) of LA, 60 mmol of HCOOH, and 10 mmol of Et<sub>3</sub>N were added, and the reaction mixture was stirred for 24 h at 80 °C in an oil bath to achieve complete conversion of LA. For the recyclability experiment, to an aqueous suspension of the catalyst (2.5 mol% in 5 mL water), 1.0 mmol of LA, 6 mmol of HCOOH and 1 mmol of Et<sub>3</sub>N were added and the reaction mixture was stirred at 80 °C in an oil bath for a specified time. After each run, the reaction mixture was extracted with diethyl ether and ethyl acetate, while the catalyst remains in the aqueous phase. Before performing the consecutive run, the required amount of HCOOH and Et<sub>3</sub>N was added to adjust pH to 2.7 and then the hydrogenation reaction of LA was carried out.



## Conflicts of interest

There are no conflicts to declare.

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