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Recent advances in the conversion of furfural into bio-chemicals through chemo- and bio-catalysis

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Furfural is a promising renewable platform molecule derived from hemi-cellulose, which can be further converted to fossil fuel alternatives and valuable chemicals due to its highly functionalized molecular structure. This mini-review summarizes the recent progress in the chemo-catalytic and/or bio-catalytic conversion of furfural into high-value-added chemicals, including furfurylamine, C₆ carboxylic acid, *i.e.*, furandicarboxylic acid, furfural alcohol, aromatics, levulinic acid, maleic acid, succinic acid, furoic acid, and cyclopentanone, particularly the advances in the catalytic valorization of furfural into useful chemicals in the last few years. The possible reaction mechanisms for the conversion of furfural into bio-chemicals are summarized and discussed. The future prospective and challenges in the utilization of furfural through chemo- and bio-catalysis are also put forward for the further design and optimization of catalytic processes for the conversion of furfural.

1. Introduction

With the depletion of fuel reserves and increasing demands for fuel, great efforts have been devoted to developing novel sustainable and renewable energies and resources.^{1–5} Biomass is the only renewable organic carbon source in nature, which is environmentally friendly, has abundant reserves and inexpensive, providing unique advantages for the production of fuels and industrially important chemicals.^{5–7} Furfural is a versatile platform molecule, and thus has received considerable attention in recent years. The presence of an aldehyde group in furfural makes it possible to further synthesize higher value chemicals and polymer monomers, including furandicarboxylic acid, furfurylamine, furfural alcohol, cyclopentanone, and levulinic acid and other molecules through oxidation, hydrogenation, and hydrolysis (Fig. 1).⁶ Hence, the high-value utilization of biomass-derived furfural is important for sustainable development in the future.

Furfural is produced from agricultural resources containing xylose or xylan, such as wood wastes and corn cob.⁸ The main producers of furfural are China (~70% total production capacity), The Dominican Republic (Central Romana

Corporation, 32 kton per year), and South Africa (20 kton per year). These three countries account for approximately 90% of the global furfural capacity (280 kton per year).^{9,10} In the past, most studies have been carried out to reveal the reaction mechanism for the conversion carbohydrates into furfural at the molecular level. As shown in Fig. 2, initially, the formation of furfural from hemicellulose-containing biomass includes the acid-catalyzed depolymerization of the hemicellulose component to form xylose. Subsequently, the isomerization of xylose to xylulose occurs followed by a dehydration reaction to yield furfural.¹¹ To date, various reaction systems such as biphasic

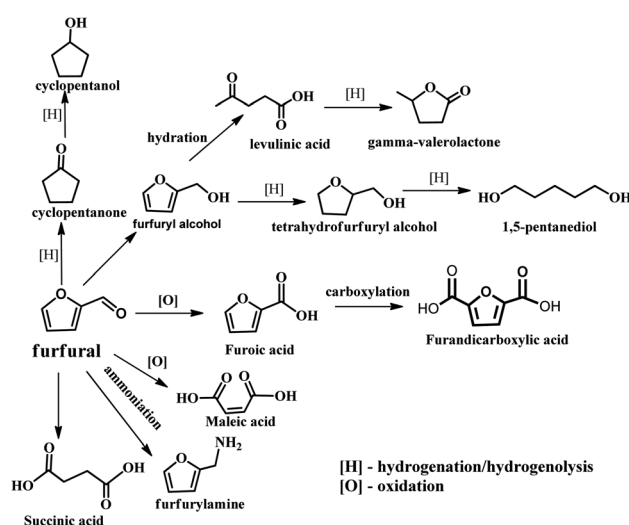


Fig. 1 Furfural as a platform compound for diverse reactions.

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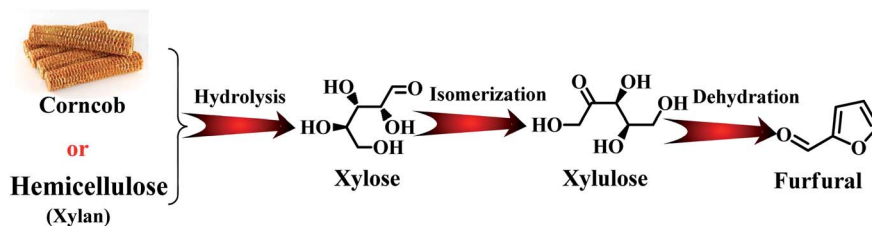


Fig. 2 Conversion pathway of hemicellulose component into furfural.

reaction systems and ionic liquids have been developed to boost the yield of furfural.

Some excellent reviews about the conversion of hemicellulose to furfural and the valorization of furfural into bio-fuels have been published.^{5,6,12–14} A systematic review of the conversion of xylan into furfural is beyond the scope of this manuscript. However, a comprehensive summary of latest advances in the valorization of furfural to value-added chemicals, especially the progress in the last five years, has not been reported to date. This mini-review summarizes the recent advances in the conversion of furfural into valuable chemicals through chemo- and bio-catalysis. This mini-review mainly focuses on the catalytic strategies for the valorization of furfural into useful chemicals including selective oxidation reactions, hydrogenation, hydrogenolysis, reductive amination, and aromatization. The valuable products, include maleic anhydrides, maleic acid, succinic acid, furoic acid, furfurylamine, furandicarboxylic acid, furfural alcohol, aromatics, levulinic acid, and cyclopentanone, especially for advancements in the catalytic conversion of furfural to useful chemicals in the last few years. The possible reaction mechanisms for the conversion of furfural into bio-chemicals are summarized and discussed. In addition, future prospects are also provided to highlight the challenges and opportunities for the valorization of furfural.

2. Catalytic conversion of furfural into value-added chemicals

2.1 Aerobic oxidation of furfural into carboxylic acid

2.1.1 Oxidation of furfural to maleic acid (MA) and fumaric acid (FA). MA and FA are a critical C₄ chemical intermediates in the manufacture of vinyl copolymers, surface coatings, plasticizers, and pharmaceuticals.^{15,20} At present, maleic acid is manufactured industrially through the aerobic oxidation of butane, benzene, and butadiene, whereas FA can be produced *via* the hydrolysis of maleic anhydride or biological transformation with fungi.¹⁶ Alternatively, the production of MA and FA starting from renewable lignocellulose such as furfural and HMF is an emerging topic. Currently, HMF is generated from edible and expensive monosaccharides such as fructose and glucose, thus making it uneconomic to synthesize downstream value-added chemicals on a large scale.^{17,18} In comparison with HMF, furfural is relatively cheaper and is currently manufactured on an industrial scale. Thus, it is highly desirable to consider furfural as a better starting material for the production of MA and FA.

In the past years, numerous heterogeneous catalysts have been applied to catalyze the oxidation of furfural to MA and FA. This reaction was studied by Yin *et al.* with phosphomolybdic acid as the catalyst in aqueous/organic biphasic reaction systems.¹⁹ In the biphasic reaction system, the aerobic oxidation of furfural took place in the aqueous phase, while the organic phase constituted a reservoir to gradually release the substrate to the aqueous phase through phase equilibrium. A 34.5% yield of MA with a selectivity of 68.6% was obtained with the furfural conversion of 50.4%.¹⁹ In addition, a possible reaction mechanism was also proposed, as illustrated in Fig. 3.¹⁹ Firstly, a furfural radical, compound 1, is formed through the first hydrogen abstraction by either oxygen or phosphomolybdic acid, and subsequently transfers an electron to phosphomolybdic acid to produce the furfural cation (compound 2). Secondly, the furfural cation is attacked by H₂O to generate compound 3, which can undergo 1,4-rearrangement to form compound 4. Eventually, compound 4 is hydrolyzed to produce compound 5, which may be further oxidized to MA.¹⁹ However, the catalysts were prone to deactivation owing to the leaching of their active components. In the case of the Mo–V–O catalyst, a total yield of MA and FA of up to 65% was afforded in acetic acid solvent under 20 bar of O₂, but the leaching of the V species into the solution resulted in a decrease in the yield of MA and FA in the catalyst recycling tests.²¹ Significant leaching of Ti was also observed for titanium silicate-1 (TS-1), which was used to

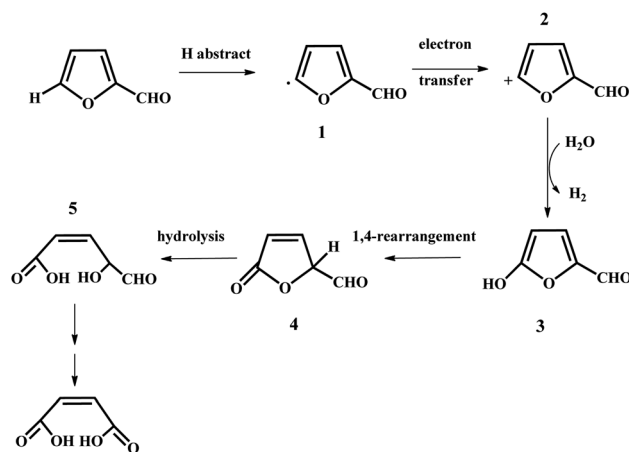


Fig. 3 Proposed mechanism for the oxidation of furfural catalyzed by phosphomolybdic acid. Reprinted from ref. 19. Copyright 2011, the American Chemical Society.

It has been reported that MANh may be generated with 5-HMF as the starting material.⁴³ A high yield of MA of up to 79% was obtained over a cheap V₂O₅ catalyst.⁴³ However, although the yield of MANh from pure 5-HMF is good, the high cost of 5-HMF hinders the production of MANh on an industrial scale.^{18,44} Alternatively, furfural possess a similar structure as 5-HMF and may act as another feedstock for the production of MANh. Compared with HMF with a limited production volume, furfural

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is regarded as a more desirable starting material for the production of MANh.

In the past few years, several catalytic reaction systems including homogeneous and heterogeneous catalysis have been applied to catalyze the oxidation of furfural to MA. Yin *et al.* explored a new homogeneous oxidation strategy in which $\text{H}_5\text{PV}_2\text{Mo}_{10}\text{O}_{40}$ and $\text{Cu}(\text{CF}_3\text{SO}_3)_2$ were used to catalyze the aerobic oxidation of furfural to MANh and furanone.³⁸ Under the optimal conditions, an MANh yield of 54% was obtained with 7.5% of 5-acetoxyl-2(5H)-furanone. Furthermore, a plausible mechanism for the oxidation of furfural was proposed, in which three different reaction routes were put forward. They suggested that the dominant oxidation pathway is initiated by hydrogen abstraction from the 5-position of furfural, and then MANh and 5-acetoxyl-2(5H)-furanone are produced simultaneously. As is well known, homogeneous catalysis has some drawbacks in terms of product separation and environmentally friendliness, especially for catalyst recycling.

In comparison with homogeneous catalysis, heterogeneous catalytic systems have more advantages in the separation of products and catalyst recycling. In 2012, Ojeda *et al.* reported a heterogeneous catalytic system, giving an MANh yield as high as 73% through the gas oxidation of furfural at 593 K over $\text{VO}_x/\text{Al}_2\text{O}_3$.⁴⁵ It was shown that furfural undergoes tandem oxidation to generate furan, 2-furanone, and eventually MANh. It was revealed that Al_2O_3 -supported polyvanadates are more active than monovanadates (VO_4) and V_2O_5 oxides in the production of MANh and furan.⁴⁵ Zhang and co-workers reported that plate-like vanadium phosphorous oxides (VPO) exhibited good activity in the oxidation of furfural to MANh in the gas phase.⁴¹ An MANh yield of up to 97% was achieved at 10 vol% of furfural in the feed with air as the oxidant at 340 °C. The catalyst also showed good stability with no remarkable loss in activity and selectivity toward MANh during 25 h reaction. Based on the product distribution analysis, 2-furoic acid was proposed as the

main side-product and was also a key intermediate for the synthesis of MANh over the VPO catalyst. The possible reaction pathway for the gas oxidation of furfural to MANh was proposed, as depicted in Fig. 4.⁴¹ The authors assumed that the oxidation of furfural to 2-furoic acid is the rate-limiting step for the oxidation of furfural over the VPO catalyst because furan was detected in a very trace amount or not produced.⁴¹

Granados *et al.* studied the deactivation process and mechanism for the gas phase oxidation of furfural to MANh over $\text{V}_2\text{O}_5/\gamma\text{-Al}_2\text{O}_3$ catalysts.⁴⁶ The results showed that the catalyst was readily deactivated because of the formation of maleates and resins on its surface, especially when the fresh catalyst initially contacted the reaction feedstock at a low temperature. Moreover, increasing the temperature from low to high did not lead to the removal of the deposits, but boosted their deposition. Thus, the yield of MANh never exceeded 30% because of the deactivation of the catalyst. In contrast, when the fresh catalyst initially contacted the feedstock at high temperature, the deposition rate of maleates and resins remarkably slowed down. Therefore, the yield of MANh significantly increased and the deactivation rate was reduced substantially.⁴⁶ In addition, VO_x immobilized on Schiff base-modified graphene oxide was utilized to oxidize furfural to MANh with 95.3% yield under the optimized reaction conditions.²⁶ The oxidation reaction results of furfural to MANh are summarized in Table 2.

In addition to the gas-phase oxidation of furfural to MA, solid-liquid reaction systems exhibit good performances in the production of MA. For example, a binary Mo–V metal oxide was used to catalyze the aerobic oxidation of furfural to MA, and an MA yield of 65% was afforded on the $\text{Mo}_4\text{VO}_{14}$ catalyst under the optimal reaction conditions.²¹ However, the recyclability test of the $\text{Mo}_4\text{VO}_{14}$ catalyst showed that a certain amount of V and Mo components leaked into the solution, thereby leading to a decrease in the activity and selectivity toward MANh. Unlike the gas-phase oxidation of furfural, the solid-liquid reaction

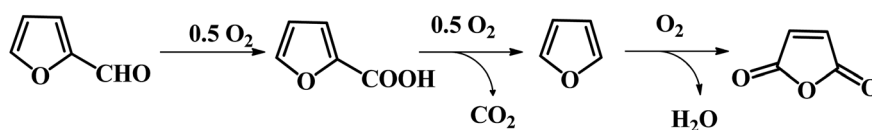


Fig. 4 Possible reaction pathway for the oxidation of furfural to MANh on VPO catalyst. Reprinted from ref. 41. Copyright 2018 Wiley.

Table 2 Various catalysts for the conversion of furfural to MANh

Entry	Catalyst	Reaction type	T (°C)/ t (h)	Oxidant (MPa)	Conversion of furfural (%)	MANh yield (%)	Ref.
1	$\text{H}_5\text{PV}_2\text{Mo}_{10}\text{O}_{40} \cdot x\text{H}_2\text{O}$	Liquid/ CH_3CN	110/3	O_2 (20 bar)	98.7	54.0	47
2	$\text{H}_6\text{PV}_3\text{Mo}_9\text{O}_{40} \cdot x\text{H}_2\text{O}$	Liquid/ CH_3CN	110/3	O_2 (20 bar)	98.7	54.0	47
3	$\text{V}_2\text{O}_5/\text{Al}_2\text{O}_3$	Gas-phase	310	10% O_2/He	100	86	40
4	VPO_{HT}	Gas-phase	340	10% O_2/He	99.2	97.9	41
5	$\text{Mo}_4\text{VO}_{14}$	liquid/ AcOH	120/16	O_2 (20 bar)	100	65	48
6	$\text{VO}_x/\text{Al}_2\text{O}_3$	Gas-phase	320	O_2	100	73	37
7	$\text{VO}_x/\text{Al}_2\text{O}_3$	Gas-phase	300	O_2	—	55	42
8	$\text{VO-NH}_2\text{-GO}$	liquid/ AcOH	90/4	O_2 (20 bar)	99.8	95.3	26



Entry	Catalyst	Reaction type	<i>T</i> (°C)/ <i>t</i> (h)	Oxidant	Conversion of furfural (%)	SA yield (%)	Ref.
1	SO ₃ H-CD ^a		60/1.5	H ₂ O ₂	100	81.2	53
2	Amberlyst-15	Toluene	—	H ₂ O ₂	—	52.3	49
3	Sulfated zirconia	—	75	H ₂ O ₂	100	86.0	52
4	GO ^b	H ₂ O	70/24	H ₂ O ₂	99.8	97.9	50
5	SO ₃ H-carbon	H ₂ O	80/1	H ₂ O ₂	100	93.0	51

Recently, solid acids have been used to catalyze the oxidation of furfural to SA with H_2O_2 as the oxidant, such as Amberlyst-15 cation exchange resin,^{49,53} sulfonated graphene oxide (GO),⁵⁰ $-\text{SO}_3\text{H}$ -functionalized polymer,^{51,53} and sulfated zirconia.⁵⁶ An



Table 4 Summary of the reaction results for the catalytic oxidation of furfural to FCA

Entry	Catalyst	Reaction type	$T (^{\circ}\text{C})/t$ (h)	Oxidant	Conversion of furfural (%)	SA yield (%)	Ref.
1	Au/ZTC ^a		120/6	O ₂	89	89	63
2	N-heterocyclic carbene	DMSO	40/6	O ₂	>99	>99	64
3	AuPd/Mg(OH) ₂	H ₂ O	30	O ₂	99.5	84.5	57
4	ALDHs ^b	Buffer	40	—	>99	>99	62
5	<i>E. coli</i>	Buffer ^c	30	—	100	99	61

^a ZTC – zeolite-templated carbons. ^b ALDHs – aldehyde dehydrogenases; buffer – KH₂PO₄/K₂HPO₄, pH = 8.5. ^c Buffer – KH₂PO₄/K₂HPO₄, pH = 7.

innovative pathway has been reported for the conversion of hemicellulose prehydrolysate to SA using Amberlyst-15 as the catalyst and H₂O₂ as the oxidant in a biphasic system.⁴⁹ In the biphasic system, a key intermediate, furfural, which was produced from the dehydration of xylose, was oxidized to SA with a maximum yield of 52.3%. This work exhibits the potential for the valorization of inexpensive hemicellulose to the value-added SA. Lv and co-workers synthesized a highly active graphene oxide (GO) catalyst decorated with –SO₃H groups for the oxidative conversion of furfural to SA with H₂O₂ as the oxidant.⁵⁰ A maximum SA yield of 97.9% was achieved under the optimized reaction conditions. They revealed that GO with –SO₃H groups exhibited superior activity and SA selectivity for the oxidation of furfural than other common acid catalysts, such as HCl, H₂SO₄, and ZSM-5. Moreover, the GO catalyst has good reusability with no remarkable loss in furfural conversion and SA yield after six consecutive recycles. The decrease in the furfural oxidation activity was attributed to the leaching of the oxygenated-containing groups and –SO₃H groups, as evidenced by the XPS results. However, the active sites of the catalysts are the –SO₃ groups and the oxidizing agent is the expensive H₂O₂.⁵⁰ In addition, Murzin and co-workers reported the oxidation of furfural over sulfated zirconia in the presence of H₂O₂, and many products including MA, SA, furoic acid, and 2(5H)-furanone were generated, but the SA selectivity was low. Thus,

more efforts should be devoted to developing more robust and effective catalytic systems besides the use of –SO₃H group-grafted polymers or their functionalized catalysts with the use of the inexpensive O₂ or air as the oxidant instead of the expensive H₂O₂ for the oxidation of furfural to SA in the future.

2.1.4 Oxidation of furfural to furoic acid (FCA). FCA is a versatile chemical for the production of a variety of pharmaceutical drugs, fragrances, biofuels, *etc.* FCA is currently manufactured commercially through a Cannizzaro reaction with an inorganic base.⁵⁷ However, this industrial process has many drawbacks including low yield of FCA and difficulties in the separation of the product. Thus, several efforts have been undertaken to develop environmentally friendly approaches for the conversion of furfural to FCA in recent years. Chemo- or biocatalysis has been applied to produce FCA *via* the selective oxidation of furfural. It has been shown that the use of heterogeneous catalysts to enhance the FCA yield is very challenging due to the formation of undesirable by-products.

2.1.4.1 Selective oxidation of furfural to FCA through chemical conversion. Various heterogeneous catalysts such as Ag₂O/CuO,⁵⁸ AuPd/Mg(OH)₂,⁵⁷ and Au/ZTC⁵⁹ catalysts, have been utilized to catalyze the oxidation of furfural into FCA. A summary of the solid catalysts for the selective oxidation of furfural to FCA is provided in Table 4. Giordano *et al.* reported the excellent performances of Au/ZTC, a carbon support

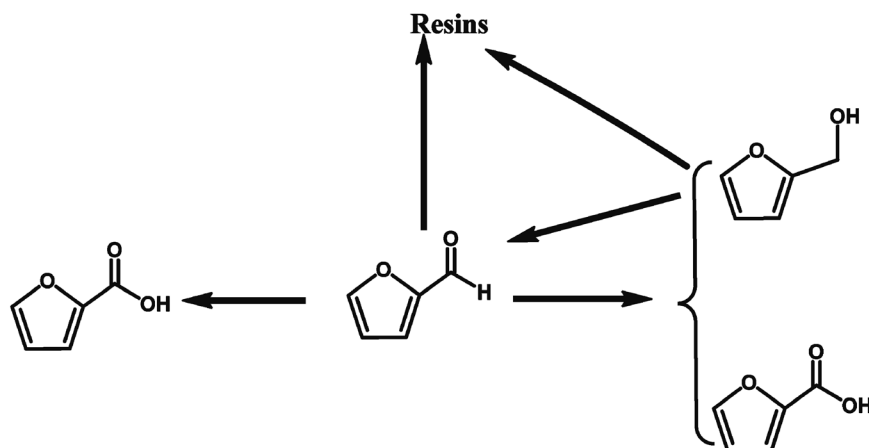


Fig. 6 Proposed reaction routes for the selective oxidation of furfural to FCA over AuPd/Mg(OH)₂ in the presence of NaOH. Reprinted from ref. 57. Copyright 2017, The Royal Society of Chemistry.



synthesized using a carbon replica of BETA zeolite, in the aerobic oxidation of furfural to FCA in the absence of base.⁵⁹ An 89% furfural conversion with full selectivity toward FCA was achieved in the absence of base, which provides a green approach for the production of FCA from furfural. Hutchings *et al.* reported that a 1% AuPd/Mg(OH)₂ catalyst catalyzed the selective oxidation of furfural to FCA with a yield of 84.5% in the presence of NaOH.⁵⁷ They revealed that the reaction proceeds *via* many different pathways, and the addition of NaOH promotes the Cannizzaro reaction between the furfural compounds, leading to stoichiometric quantities of furfural alcohol and FCA, as illustrated in Fig. 6. The catalyst enhances both the direct conversion of aldehyde to the acid and the tandem oxidation of furfural alcohol to furfural and then to FCA.⁵⁷ However, this reaction requires the addition of NaOH as the base catalyst to promote the selective oxidation of furfural to FCA, making it difficult to separate the products downstream and increasing the production costs. Compared to the current industrial process through the Cannizzaro reaction catalyzed by an inorganic base, the use of solid catalysts for the selective oxidation of furfural to FCA has more advantages such as easy separation of the products and higher FCA yields.

2.1.4.2 Selective oxidation of furfural to FCA through biocatalysis. Besides the chemical synthesis of FCA, Peng *et al.* reported an efficient one-pot chemo-enzymatic approach for the conversion of rice straw to form FCA through a tandem catalyst with Sn-sepiolite and whole-cell harboring horse liver alcohol dehydrogenase (HLADH) under mild reaction conditions.⁶⁰ The Sn-sepiolite could effectively convert rice straw to furfural in 42.2% yield at 170 °C for 20 min. Subsequently, the resulting furfural could be completely oxidized to FCA with *E. coli* TS whole-cells within 96 h at room temperature. In addition, the inorganic catalyst and immobilized TS whole-cell biocatalyst were recycled and reused. Li and co-workers also reported a biocatalytic route for the selective oxidation of furfural to FCA using recombinant *E. coli* cells harboring ASPDH from *C. testosteroni* SC1588.⁶¹ An FCA yield as high as 99% was achieved

under the optimized reaction conditions. Moreover, the recombinant strain showed good tolerance to FF up to 100 mM.⁶¹ Mutti *et al.* presented a study on the oxidation of furfural to FCA using three recombinant aldehyde dehydrogenases (ALDHs).⁶² A very high FCA yield of >99% could be obtained in phosphate buffer at pH 8.5 and 40 °C.⁶² However, for the biocatalytic oxidation of furfural toward FCA, the long-term stability should be further improved through reaction engineering strategies. In addition, the enhanced substrate tolerance is highly favorable for industrial application in future.

2.2 Catalytic transformation of furfural to C6 carboxylic acid

FDCA-based polyesters represent one of the important types of furan-based polymers. At present, the production of FDCA is mainly through the catalytic oxidation of HMF.⁶³ However, as mentioned above, HMF has high production costs, which impedes the large-scale commercial production of downstream products such as FDCA and DFF. Furfural, a commercial bulk biomass-based chemical, is a more desirable starting material in the production of FDCA than HMF. Efforts have been undertaken to explore some novel pathways for the generation of FDCA from furfural to address the bottleneck of HMF production.

Fu and co-workers reported the multistep catalytic conversion of furfural to poly(butylene 2,5-furandicarboxylate) (PBF) *via* the disproportionation of furoate to furan and 2,5-furandicarboxylate with total carbon utilization, as shown in Fig. 7.⁶⁶ The reaction pathway includes four steps as follows: firstly, the oxidation of furfural to furoate is catalyzed. Secondly, the resulting furoate undergoes disproportionation to give rise to furan and 2,5-furandicarboxylate. Subsequently, furan is subjected to hydrogenation, and then hydrolysis to form 1,4-butanediol (1,4-BDO). Finally, polymerization of 1,4-BDO with 2,5-FDCA is performed to form PBF. The critical step of this pathway is the selective disproportionation of furoate into furan and 2,5-furandicarboxylate, which provides a key linkage

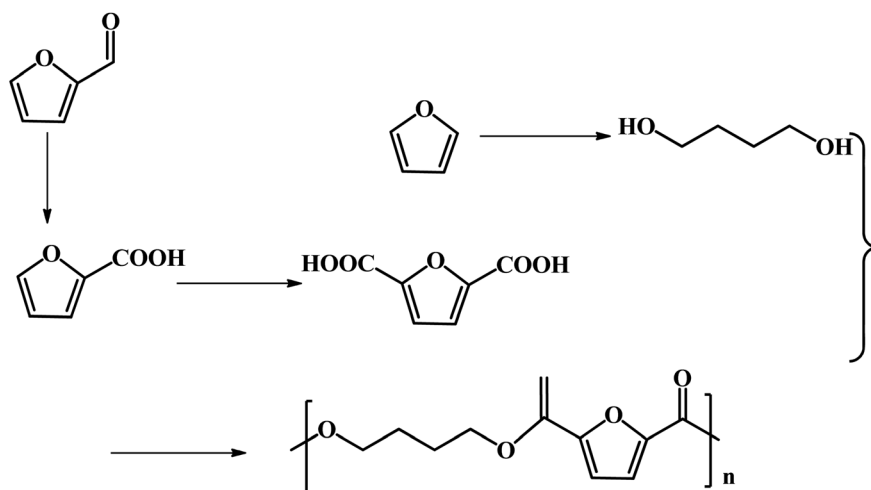


Fig. 7 Multistep conversion of furfural to the 2,5-furandicarboxylic acid-based polyester PBF. Reprinted from ref. 66. Copyright 2013 Wiley.



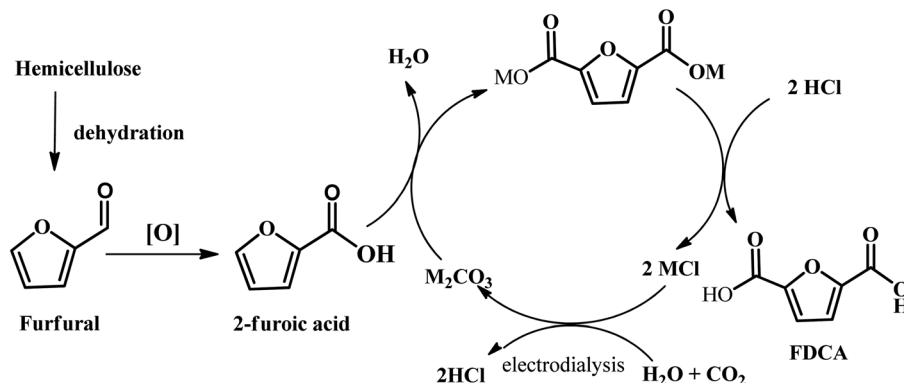


Fig. 8 Schematic illustration of the overall diagram for the production of FDCA from biomass through furfural using CO_3^{2-} -promoted C–H carboxylation. Recovery of the alkali cations is achieved by protonation of the FDCA salt and bipolar membrane electrodialysis of the resulting alkali chloride salt. Reprinted from ref. 67. Copyright 2016, The Royal Society of Chemistry.

between commercialized furfural and the furan-based polymer monomer FDCA.⁶⁶

Another emerging pathway has been reported by Kanan *et al.*, where a C–H carboxylation reaction was firstly carried out to generate 2-furonic acid and CO_2 in high yield followed by protonation to convert 2-furoic acid to FDCA (see Fig. 8).⁶⁷ This reaction was performed by heating a salt mixture consisting of caesium furoate (the deprotonated form of 2-furoic acid) and Cs_2CO_3 under a CO_2 atmosphere. A maximum FDCA yield of 89% was achieved on a 1–10 mmol scale after 5 h of reaction, with only 5% degradation product. After carboxylation, protonation with strong acid (*e.g.*, HCl) in water precipitated the FDCA product and an aqueous salt solution (*e.g.*, MCl). The stoichiometric base and acid consumed in this synthesis could be regenerated *via* bipolar membrane electrodialysis of the salt solution. This CO_3^{2-} -promoted carboxylation pathway to FDCA has a number of great potential advantages over conventional hexose-based routes. Firstly, the oxidation of furfural into 2-furoic acid is easier than oxidizing HMF to FDCA. Next, no solvents or transition metal catalysts are required in this strategy, which is an environmentally friendly route to synthesize FDCA. Considering that 2-furoic acid can be readily generated from lignocellulose, CO_3^{2-} -promoted C–H

carboxylation opens a new avenue to convert inedible biomass and CO_2 into a valuable chemical.

Recently, Liu and co-workers reported an integrated process for the full conversion of lignocellulosic materials and CO_2 to form FDCA using a single-atom Co on N-doped carbon catalyst (Co SAS/N@C).⁶⁸ During the carboxylation reaction of furfural, Cs_2CO_3 was employed as a base additive, which boosted the carboxylation reaction of the 5-position of furoic acid. An FDCA yield of 71.7% with a selectivity of 82.9% was obtained from the oxidation-derived product of furfural, which is comparable to the previous work (76% FDCA yield with 82.6% selectivity). Moreover, the Co SAS catalyst was found to afford an FA yield as high as 96.9% in the aerobic oxidation of furfural to FA, outperforming the state-of-the-art non-noble metal catalysts.⁶⁸

In addition to the disproportionation and carboxylation routes of furoate, Yin *et al.* explored a new approach to synthesize FDCA from 2-furoic acid through consecutive bromination, esterification, carbonylation, and hydrolysis with 65% total yield in four steps and above 80% isolated yield in each step (Fig. 9).⁶⁹ The first step, *i.e.*, bromination, was performed by reacting furoic acid with bromine to obtain 5-bromofuroic acid with a yield of 86%. In the next step, ethyl 5-bromofuran-2-carboxylate (3) was feasibly generated by refluxing 5-bromofuroic acid (2) in ethanol with the use of concentrated

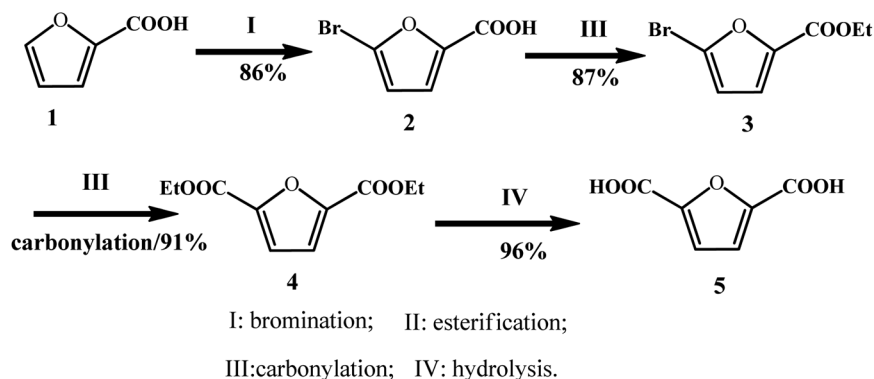


Fig. 9 Synthetic strategies for FDCA from furoic acid. Reprinted from ref. 69. Copyright 2017, the American Chemical Society.

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alanine as the amine donor.⁸⁰ In this approach, an acidified Sn/ZrO₂ catalyst was applied to convert pretreated corncob into furfural. Subsequently, *E. coli* CVPRSF Duet whole cells were added to hydrolysate for the bio-catalytic conversion of furfural without separation of the Sn/ZrO₂ catalyst, which afforded 84.0% yield of furfurylamine with 100% selectivity. Therefore, a novel tandem chemoenzymatic synthesis pathway from lignocellulose has been exploited by the combination of heterogeneous catalysis and bio-catalysis for the first time. In comparison with the use of heterogeneous catalysts, the bio-catalytic amination reaction does not require the use of H₂ and the reaction conditions are very mild. However, in the first step of the process, the conversion of corncob into furfural requires the addition of homogeneous acids, and thus involves a neutralizing step using alkali prior to enzymatic catalysis.

2.4 Hydrogenation of furfural to furfuryl alcohol (FOL)

Furfuryl alcohol is the most important hydrogenated product derived from furfural and accounts for nearly 62% of the annual downstream products of furfural.¹¹ Currently, its industrial production in the liquid or gas phase is mainly through the selective hydrogenation of furfural.⁸¹ Although the cost-effective gas reaction process is more suitable for industrial applications, this process only gives a low yield of FOL, which cannot be properly addressed.¹¹ Therefore, the production of FOL in the liquid phase has become the trend of current research.

For liquid phase hydrogenation, a variety of catalysts can achieve an FOL yield greater than 90%. The commonly used catalysts are Cu–Cr catalysts.⁸² Although this type of catalyst can give high FOL yields, the catalyst tends to deactivation because of the strong adsorption of byproducts on its active metal sites, and the Cr element is toxic. Furthermore, the hydrogenation process of furfural with Cu–Cr catalysts is environmentally unfriendly. Thus, in recent years, environmentally benign catalysts for the hydrogenation of furfural have been extensively explored using green reaction systems, and some excellent works are summarized in Table 5.

Li *et al.* used a Pd/H-UiO-6 catalyst to catalyze the hydrogenation of furfural to FOL and a high yield was achieved under mild reaction conditions.⁸⁴ The Pd-Cu/MgO and Ru/UiO-66 catalysts afforded excellent FOL yields of 99% and 94%, respectively.^{87,88} Halilu *et al.* synthesized low-cost, recyclable Fe(NiFe)O₄-SiO₂ for the hydrogenation of furfural, which resulted in 93% FOL yield in heptane solvent.⁸⁹ In addition, some durable, efficient catalysts such as Cu-Fe oxides,⁹⁵ Ni-Fe alloys,^{96,97} and PtSn alloy⁹⁴ have also been employed in the hydrogenation of furfural to produce FOL in recent years. It is

Table 5 Various catalysts for the conversion of furfural into FOL

Entry	Catalyst	Solvent	T (°C)/ t (h)	H ₂ (MPa)	Furfural con. (%)	FOL yield (%)	TOF (h ⁻¹)	Ref.
1	Cu-SiO ₂ -I	2-Propanol	110	10	66.3	—	0.009	83
2	3% Pt/SiC-C	H ₂ O	25/3	1.0	80.2	26.5	—	
3	Pd/H-UiO-6	H ₂ O	60/3	0.5	>99	>99	66.7	84
4	Pt/(Fe,Co)-BTC	2-Propanol	70/2	1.0	98.0	97.0	1044	85
5	PtNi nanoframes	2-Propanol	100/1	1	100	99	2.56	86
6	Pd-Cu/MgO	H ₂ O	110/1.3	0.6	100	99	—	87
7	Ru/UiO-66	H ₂ O	20/4	0.5	95	94	—	88
8	Fe(NiFe)O ₄ -SiO ₂	Heptane	90/4	2	94	93	—	89
9	Ru-PES ^a	H ₂ O	70/7	0.7	26.1	>99	48 000	90
10	Cu/C	2-Propanol	170/3	—	75.0	72.5	14.4	91
11	Pt-Sn/SiO ₂	2-Propanol	100/8	—	100	96	—	92
12	Ni-SAs/NS ^b	2-Propanol	130/3	—	95.6	92.5	623	93
13	PtSn@HMSNs	2-Propanol	100/5	1.0	100	94.5	—	94

^a PES refers to polyethersulfone. ^b SAs refer to single atom catalysts.

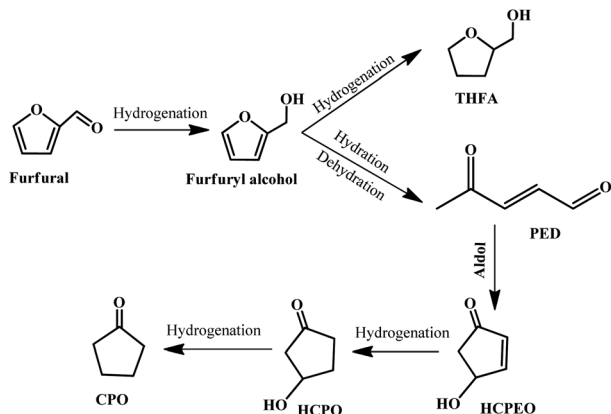


Fig. 12 Reaction pathway for the hydrogenative ring-rearrangement of furfural into cyclopentanone. Reprinted from ref. 115. Copyright 2019, Elsevier.

highly desirable to design and develop cheap, robust, efficient catalysts for the hydrogenation of furfural, but noble metal catalysts should not be the first choice because they are expensive and increasingly exhausted.

Besides the use of H_2 as a hydrogen source, catalytic transfer hydrogenation (CTH) is a potential alternative protocol for the hydrogenation of furfural to FOL.^{93,98–101} Zr-based catalysts,^{102–105} CuNi@C,¹⁰¹ single-atom Ni catalyst,⁹³ metal–organic framework (MOF) materials,^{101,103,106,107} noble metal catalysts such as Pd, Pt, and Ru,^{108–111} and transition metal catalysts such as Fe oxides¹¹² have been used to catalyze the reaction. MOF-808 has been shown to be the most active catalyst in the CTH of FUR to FOL among the Zr-MOFs due to the facile modification of the metal node mediating the acid–base properties by the introduction of coordinatively unsaturated sites, hydroxyl and methoxy groups in the framework of the Zr-MOF, which favor the adsorption of furfural and 2-propanol. For the in-depth understanding of the CTH mechanism, a computational study based on DFT was conducted, and two feasible geometries of Zr nodes were proposed, *i.e.*, singly and doubly modified Zr nodes of MOF-808.¹⁰³ It was disclosed that the hydride transfer from alkoxy carbon to carbonyl carbon is the rate-limiting step based on DFT studies.¹⁰³ Considering that the hydrogenation of furfural

is a relatively easy reaction, some cheap and robust supported catalysts are more desirable for the reaction compared to expensive noble metal catalysts.

2.5 Furfural to cyclopentanone (CPO)

Cyclopentanone (CPO) is a versatile compound derived from furfural, which is widely used in the synthesis of fuels, pharmaceuticals, rubber chemicals, fungicides, and flavor and fragrance chemicals.^{113–116} It is well-accepted that the most important reaction step in the conversion of furfural to cyclopentanone is the hydrogenative ring-rearrangement of furfural, but the reaction mechanism is still under debate, and various reaction routes and reaction intermediates have been proposed, which may be dependent on the catalyst type, reaction conditions, *etc.*^{116–121} In general, furfural is firstly hydrogenated to produce furfuryl alcohol, followed by a ring-opening reaction to give 2-pentene-1,4-dione (2-PED) accompanied by the formation of a tetrahydrofurfuryl alcohol (THFA) byproduct. Subsequently, 2-PED undergoes an aldol reaction to give 4-hydroxy-2-cyclopentenone (HCPEO), and then further hydrogenated to form 3-hydroxycyclopentanone (HCPO). Eventually, HCPO is converted to CPO by hydrogenation.¹¹⁵ The possible reaction pathway is displayed in Fig. 12.

Various monometallic catalysts such as supported Cu,¹²¹⁻¹²³ Pt,¹²⁴ Ru,^{125,126} and bimetallic catalysts, *e.g.*, supported Co-Ni,¹²⁷ Cu-Co,¹²⁸ and Pd-Co,¹²⁹ have been shown to have high catalytic activity for the transformation of furfural to CPO, and their catalytic performances are listed in Table 6. Among the catalysts used in this reaction, Cu-based catalysts are the most thoroughly studied because of their low cost and high catalytic activity. Wang and co-workers demonstrated that a CuNi_{0.5}@C bimetallic catalyst fabricated *via* the pyrolysis of Cu-based metal-organic frameworks provided a high CPO yield of 96.9% with the almost complete conversion of furfural.¹¹⁸ 2-Cyclopentenone is prospected as the reaction intermediate involved in the reaction,¹¹⁸ which is consistent with the assumption by some researchers that 4-hydroxy-2-cyclopentenone is the reaction intermediate.^{115,130}

In addition to Cu-based catalysts, other noble metal catalysts also possess appreciable activity for the generation of CPO from furfural.^{114,116,120,124–126,134–136} Anatase TiO₂-supported Au

Table 6 The conversion of furfural into CPO over various catalysts

Entry	Catalyst	Solvent	<i>T</i> (°C)/ <i>t</i> (h)	H ₂ pressure (MPa)	Conversion of furfural (%)	CPO yield (%)	Ref.
1	CuZnAl-LDH	H ₂ O	140/2	2	100	86.5	131
2	Cu/ZrO ₂	H ₂ O	150/4	1.5	100	91.3	121
3	Cu/Fe ₃ O ₄	H ₂ O	170/4	3	100	91	123
4	CuNi@C	H ₂ O	130/5	5	99.3	96.9	132
5	Cu–Co@C-500	H ₂ O	150	0.5		90.2	128
6	PtCo/C	Toluene/H ₂ O	180	1	100	75	133
7	Pt/N-doped carbon	H ₂ O	150/4	3	>99	>76	124
8	Pd–Co@UIO-66	H ₂ O	120/2	3	>99	96	116
9	Pd/FeZn-DMC	H ₂ O	150	4	100	96.6	114
10	Co–Ni/TiO ₂	H ₂ O	150	4	100	53.3	127
11	Ru/CNTs	H ₂ O	160/3	1	99	91	126

nanoparticles were found to effectively inhibit the side reaction and achieve nearly 100% CPO yield in the conversion of furfural to CPO.¹³⁴ Deng *et al.* showed that a double-metal cyanide (DMC)-supported bi-functional Pd catalyst had high efficiency for the hydrogenative ring-rearrangement of furfural to cyclopentanone.¹¹⁴ It has been revealed that Lewis acid sites play an important role, *e.g.*, Pd/FeZn-DMC with moderate Lewis acidity gave a cyclopentanone yield of 96.6%, whereas Pd/FeNi-DMC and Pd/FeCo-DMC with weak Lewis acidity afforded a furfuryl alcohol yield of 91.2%.

2.6 Hydrogenolysis of furfural to pentanediol

Pentanediol, which is considered to be a promising bio-fuel, monomer of polyesters, and fine chemical intermediate, can be generated *via* the one-pot reaction of furfural or the hydrogenolysis of THFA.^{137–139} However, compared to THFA hydrogenolysis, the direct transformation of furfural to pentanediol is more challenging because it is very difficult to achieve high selectivity toward pentanediol and it tends to yield many byproducts.^{138,140}

Much effort has been focused on exploring some novel catalysts for the hydrogenolysis of furfural to pentanediol. For instance, a Pd–Ir–ReO_x/SiO₂ bi-functional catalyst was used for the one-pot hydrogenolysis of furfural to 1,5-pentanediol with THFA as the intermediate and gave a maximum 1,5-pentanediol yield of 71.4%.¹⁴¹ Hydrotalcite-supported Pt nanoparticles achieved a 1,5-pentanediol yield of up to 73% from furfural without any additives.¹⁴² Besides 1,5-pentanediol and 1,2-pentanediol, 1,4-pentanediol (1,4-PeD) could also be produced by the direct hydrogenolysis of furfural catalyzed by hydroxyapatite-supported Pt–Mo bimetallic catalysts.¹⁴³

2.7 Aromatization of furfural to aromatics

Aromatics are an important type of chemicals, especially for the production of polymers, solvents, pharmaceutical, fragrances, *etc.* Aromatics can be produced from fossil resources, either *via* the catalytic reforming of naphtha or steam cracking of hydrocarbons.^{5,6,144} This is an emerging and alternative way to produce aromatics using biomass as starting materials. Bio-based aromatics can be typically produced *via* the catalytic fast pyrolysis of lignocellulose, depolymerization or cracking of lignin, and conversion of carbohydrates *via* the furanic platform.^{144–146} In many cases, these processes mainly generate benzene, toluene, and xylene (BTX) through sequential Diels–Alder (DA)/aromatization reactions with ethylene over acid catalysts. Bio-based BTX can be further converted without any

modification of current downstream processes and applications.

To date, very few studies have been reported on the production of functionalized and industrially relevant aromatics from biomass, such as benzonitrile, benzaldehyde, and benzylamine derivatives. Among the biomass-derived aromatics, *meta*-xylylenediamine (MXD) is an important chemical, which is widely used in the production of polymers including polyamide and polyurethane. To date, MXD is produced at 100 kT per year from fossil feedstocks. Recently, the renewable MXD was synthesized from biomass-derived furfural and acrylonitrile.¹⁴⁷ However, the D–A reaction of furfural with acrylonitrile did not occur because the electron-withdrawing formyl group in furfural is thermodynamically unfavorable with a significantly positive free energy according to DFT results. Therefore, a novel strategy was employed *via* the acetalization derivatization of furfural with ethylene glycol to give 2-(furan-2-yl)-1,3-dioxolane, which afforded a thermodynamically favorable reaction when it reacted with acrylonitrile (as shown in Fig. 13).¹⁴⁷ Subsequently, aromatization of D–A with acrylonitrile gave 86% yield of aromatic cycloadducts catalyzed by CH₃ONa in DMSO. After aromatization, the *meta*-aromatic isomer underwent deprotection in the presence of HCl to produce ethylene glycol and *meta*-cyanobenzaldehyde in quantitative yield. Finally, reductive amination of the aldehyde with ammonia using RANEY® Co yielded 70% MXD. Moreover, all the steps (protection, DA/aromatization, deprotection and hydrogenation) are highly selective and chemical intermediates can be directly used in the next step without purification. Therefore, this novel strategy provides a sustainable, alternative way to produce MXD with high yield from biomass-derived furfural and acrylonitrile.

2.8 Furfural to other useful chemicals

Tetrahydrofurfuryl alcohol (THFA) is a liquid compound with a high boiling point, which is transparent and completely miscible with water. It is a green solvent for agricultural applications, electronic cleaners and printing inks.¹⁴⁸ THFA can be produced directly from furfural. The typical approach for the production of THFA from furfural is a two-step hydrogenation reaction through the furfuryl alcohol intermediate over Cu–Cr and noble metal catalysts separately.¹⁴⁹ However, this cumbersome process increases the production cost of THFA. Recently, the synthesis of THFA from furfural *via* a “one-pot method” addressed this problem. For example, in the study conducted by Rode *et al.*, 95% THFA yield was achieved in the presence of an

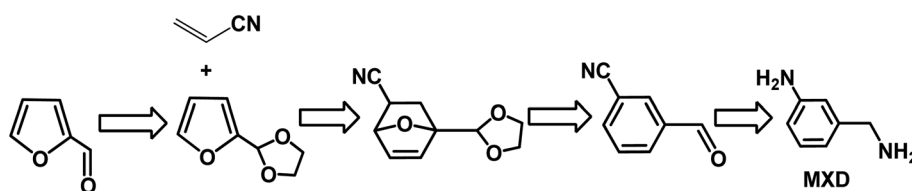


Fig. 13 Pathways to *meta*-xylylenediamine (MXD) from furfural and acrylonitrile. Reprinted from ref. 147. Copyright 2018, Wiley.



Fe catalyst by direct hydrogenation of furfural using a “one-pot process”.¹⁵⁰

Levulinic acid can also be produced from furfural, where the conversion of furfural to LA firstly involves the selective hydrogenation of furfural to form FA, followed by hydrolysis to yield levulinic acid.¹¹ Maldonado *et al.* used different experimental methods to study the hydrolysis of FA to LA with Amberlyst-15 in the aqueous phase.¹⁵¹ Different reaction pathways were systematically investigated in this process, in which 4,5,5-trihydropentan-2-one was considered to be a key intermediate.¹⁵¹ Sulfuric acid was employed to hydrolyze hemicellulose to LA in a bi-phasic reactor consisting of alkylphenol solvents and water.¹⁵² The bi-phasic reaction system effectively suppressed the undesirable polymerization reactions and obtained a yield of 72%. Recently, an LA yield greater than 70% was achieved using H-ZSM-5 as a catalyst in an H₂O/tetrahydrofuran (THF) biphasic reaction system.¹⁵³

γ-Butyrolactone (GBL), an important intermediate, is widely used as a fine chemical, medicine, *etc.* At present, GBL is produced in the petrochemical industry such as the hydrogenation of maleic anhydride. Wang *et al.* reported a two-step approach for the synthesis of GBL from furfural.¹⁵⁴ Firstly, furfural oxidation was conducted to form 2-furanone with HCOOH as the catalyst and H₂O₂ as the oxidant. Subsequently, the purified 2-furanone was further converted to GBL *via* hydrogenation reaction. The oxidation products include maleic acid and succinic acid besides 2-furanone, and formic acid was found to give a high yield of 2-furanone of up to 61.5%. It was revealed that the TOF values of supported group VIII metals for the hydrogenation of 2-furanone exhibit a volcano-type correlation with the d-band center relative to the Fermi level, and Pd/SiO₂ shows the best performance. This work provides a novel approach for transforming biomass-derived furfural to a useful chemical. It is of great importance to develop heterogeneous catalysts to afford a higher yield of 2-furanone in the first step, and thus decrease its downstream separation cost to realize the large-scale production of GBL in the future.

3. Conclusions and outlook

This review presented a comprehensive overview on the latest advancements in the catalytic valorization of furfural into value-added chemicals through chemo- and bio-catalysis approaches, focusing on the development of catalysts and the reaction mechanisms of furfural conversion. The wide utilization of biomass-derived furfural opens a new avenue to reduce the dependence on oil and fossil resources and provides a sustainable development for mankind. Despite the great progress achieved in the efficient utilization of furfural to produce useful chemicals including furandicarboxylic acid, aromatics, and cyclopentanone, and the fact that some bottlenecks have been addressed and some emerging strategies have been exploited, there are still critical challenges that hinder the utilization of furfural to produce value-added chemicals on a large scale.

(1) The bio-catalytic conversion of furfural provides a green protocol to valuable chemicals. Compared to chemo-catalysis, bio-catalysis has some advantages including high product selectivity and environmentally friendly nature, but it has some limitations, such as low productivity and low feedstock concentration, leading to extensive energy consumption for product separation and purification in the downstream processes. Thus, it is urgent to develop some novel processes by integrating bio-catalysis and chemo-catalysis to make full use of their advantages and overcome their drawbacks in the transformation of biomass into useful chemicals.

(2) Significant challenges remain in designing and developing industrially feasible reaction pathways through heterogeneous catalysts and/or bio-catalysts for the conversion of furfural into useful chemicals including diacids, diols, aromatics, and cyclopentanone. To date, most research on the utilization of furfural is still in its infancy, such as the production of furandicarboxylic acid, aromatics, succinic acid, and cyclopentanone. Thus, great efforts need to be further undertaken to meet the commercial-scale application requirements. Recently, artificial intelligence (AI)-driven synthesis has been shown to be a very strong tool in organic synthesis, materials design, *etc.* This novel technique will remarkably facilitate the screening of excellent catalysts and the development of efficient and economic pathways for the valorization of furfural to value-added chemicals.

(3) In general, the catalytic conversion of furfural to bio-chemicals requires cascade reactions, especially for hydrogenation and hydrodeoxygenation, which are usually conducted under harsh reaction conditions at high temperatures and pressures. These harsh reaction conditions result in the degradation of furfural to produce other by-products or humins, which will increase the production cost of the desired products. In addition, the hydrothermal reaction conditions easily lead to leaching of the active components and even deactivation of solid catalysts. Thus, it is of great significance to develop efficient and robust heterogeneous catalysts for the conversion of furfural. Recently, the photo-, electro-, and photoelectron-catalytic conversion of biomass into bio-fuels and bio-chemicals have attracted increasing attentions due to their mild reaction conditions, high selectivity, and environmentally friendly nature. The development of photo-electrochemical processes can minimize the energy input, while affording high conversion efficiency and selectivity. It can be expected that some innovative reaction pathways and novel valuable bio-chemicals will be yielded through these new techniques.

(4) It is necessary to have an in-depth understanding of the reaction mechanism for further improving the design and catalytic properties of solid catalysts. Besides the identification of the key reaction intermediates, the surface/interface reaction kinetics also need thorough studies. However, it is a significant challenge to *in situ* dynamically identify and follow the evolution of reaction intermediates, especially under the harsh hydrothermal reaction conditions. Thus, *in situ* and even *operando* characterization techniques and theoretical calculation may act as useful tools for this goal, which can propel our



understanding of the structure–activity relationship of catalysts and promote the development of high-efficient catalysts for the valorization of furfural.

Conflicts of interest

There are no conflicts to declare.

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