Energy Advances



REVIEW ARTICLE

View Article Online



Cite this: Energy Adv., 2024, 3.366

a review César Catizane, Ving Jiang * and Joy Sumner

Improving plastic pyrolysis oil quality via an

electrochemical process for polymer recycling:

Electrochemical hydrogenation (ECH) is a novel route for the upgradation of pyrolysis oil from both biomass and plastic feedstocks. Compared with conventional routes, including thermal cracking, ECH can be performed under mild conditions ($<80\,^{\circ}\text{C}$ and 1 atm) and without the requirement of additional H₂ supply. The successful demonstration of this application can be a critical step to enabling a circular plastic economy and low-carbon fuel production. In this review we provide a critical overview of the recent advancements in understanding the variables that influence the ECH process. In addition, we debate how this technology could be optimized and applied to plastic waste pyrolysis oil, assessing concerns such as the selection of cathode material, which needs to be resilient enough to address the complex nature of bio-oil. In addition, we present ideas on how to circumvent the challenge where the commonly used water-based electrolytes are unlikely to be suitable for pyrolysis oil treatment. Finally, we discuss the possible utilization of this product and scalability of this process.

Received 9th August 2023, Accepted 17th December 2023

DOI: 10.1039/d3ya00389b

rsc.li/energy-advances

1 Introduction

With the increasing severity of climate change and global warming, there is a growing recognition for the pressing need

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to mitigate carbon emissions and other greenhouse gases. To address this environmental challenge, several key international targets and agreements have been stablished, including Agenda 2030 and the Paris agreement. The United Nations' Agenda 2030 has outlined a set of sustainable development goals (SDGs). These goals encompass the effective management of waste, the preservation of ocean ecosystems, and the urgent



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César graduated from Pontifical Catholic University, Brazil, in Chemical Engineering and is now a PhD student at Cranfield University. His project is focused on evaluating the extent to which electrochemistry can be used to improve the pyrolysis oil quality and processing in feedstock recycling plants. He has carried out projects focused on wastewater treatment via electrochemistry and production of hydrogen from ethanol, utilizing

steam reforming. César also has a history working with both companies and government agencies, in an effort to help them achieve their carbon neutral goals.



Ying Jiang

Dr Ying Jiang is a Senior Lecturer in Bioenergy within the School of Water, Energy and Environment. Ying has extensive experiences in developing research programmes to improve the technical viability of biofuel production processes and maintain a consistent product quality and higher reaction kinetics. Current he leads the bioenergy research activities at Cranfield University, focusing on the fundamental reactions involved in biochemical and

thermo-chemical conversion routes of biomass/waste to energy. This scientific enquiry is key to address the demand of the bioenergy sector for an improved technical viability of biofuel production processes, consistent product quality and higher reaction kinetics.

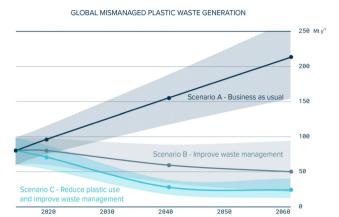


Fig. 1 Mismanaged plastic waste trend for three possible scenarios.³

undertaking of climate change mitigation efforts. 1 Notably, plastic production and utilisation is becoming more prominent in daily life, as it is a material that is utilized in a huge amount of different day-to-day products, increasing plastic pollution, which demands effective and comprehensive solutions. The trend of mismanaged plastic waste is shown in Fig. 1.^{2,3}

Many of these plastic products are designed for single-use, often discarded when they are still in pristine condition.^{2,3} In the EU for instance, in 2020, 29.5 million tonnes of postconsumer waste (PCW) plastics were collected and only 10% was sent to recycling plants, the majority of which via mechanical recycling. 4-6 This presents a significant challenge, as most of the plastic waste is either consigned to landfills or incinerated. There is considerable potential for expanding thermochemical recycling processes, with pyrolysis emerging as the most notable option.5 Furthermore, the degradation and depolymerisation of plastic through weathering process leads to the production of micro and nano-plastics. These particles are ingested by animals, thereby being introduced into the food



Joy Sumner

Dr Joy Sumner is a Reader in Energy Materials and Head of the Centre for Energy Engineering at Cranfield University. Dr Joy Sumner's work is related to the degradation mechanisms of materials used in components, most often in energy systems. Her main focus is the mechanisms of failure for steels and nickel-based alloys in simulated real world environments. To achieve this, Joy is involved in several projects including recently the EPSRC-

funded "Investigating Corrosion in Supercritical Fluids", and the Innovate UK funded "Plastic Recycling Technology and Outputs: Enhancing Material Longevity and Product Quality for Commercialisation".

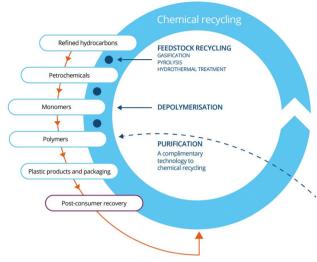


Fig. 2 Closed loop life cycle of plastic. 11

chain, and cause food contamination for both animals and humans.7

Fast pyrolysis is a thermochemical process that occurs in an inert environment, in the range of 450 to 600 $^{\circ}\text{C}$ and can be used for chemical recycling of biomass or plastic.8 This process products are separated into three categories: gas, wax, and oil. The resulting pyrolysis oil is a complex mixture composed of acids, alcohols, aldehydes, esters, ketones, sugars, phenols, guaiacols, syringols, furans, lignin-derived phenols, and extractible terpene with multi-functional groups. 9 This is a product that potentially can be utilized as a biofuel, an alternative to petroleum-based fuels, or as feedstock for the chemical industry, which could help the creation of a circular economy¹⁰ (Fig. 2).

Pyrolysis oil, however, usually has a high viscosity, lowdensity, low pH, high-water content (as high as 15-30 wt%),9 high oxygen-to-carbon (O:C) ratio and low hydrogen-to-carbon (H:C) content. 12 All these factors make the storage, transport, and utilization of the product more difficult, due to corrosion and instability, as well as lower high heating values (HHV).8 For effective and widespread usage, the pyrolysis oil needs to be upgraded.13

2 Pyrolysis oil composition

2.1 Biomass

Biomass is a renewable organic material that comes from plants and animals; its sources include wood, agricultural crops and waste material, municipal solid waste, animal manure and human sewage. Bio-oil, derived from biomass pyrolysis, is a highly intricate mixture whose composition varies with changes in the feedstock, temperature, pressure, residence time and catalysts. Other factors that can lead to different results are pre- and post-treatment of the biomass or the oil.¹⁷ Typical physical-chemical properties of the biomass derived bio-oil are summarized in Table 1.13

Table 1 Bio-oil composition matrix and associated ASTM standards¹²

Property (ASTM standard)	Value (average)	
C	56 wt%	
Н	6 wt%	
0	38 wt%	
N	0.2 wt%	
S	0.02 wt%	
Water content (D95, E203)	25 wt%	
pH acidity (D974, D664, D3339)	2.5	
Specific gravity (density compared to water)	1.2	
High heating value (D240, D4809)	17 MJ kg^{-1}	
Viscosity (D88, D445, D2170)	40–100 mPa	
Solids (char content)	0.10 wt%	
Density (D1298, D4052)	$1.2~\rm kg~L^{-1}$	

Pre-treatment is common practice in the industry in which the biomass goes through several physical-chemical processes before being pyrolyzed, such as torrefaction and trituration. 18,19 Pyrolysis oil from a lignocellulosic (Acacia nilotica) raw biomass (PO-RAW) and torrefied biomass (PO-TB) are composed of furan derivatives, phenol derivatives, acids, alcohols, aldehydes, ketones, sugar derivatives, esters, and other compounds. Table 2 summarizes the most abundant species. 19

Temperature is a critical factor during the pyrolysis process, significantly affecting product yields and altering the composition of the resulting oil. 18,20-23 For example, during the pyrolysis of food waste (FW) and food waste solid digestate (FWSD), increasing the temperature from 500 to 800 °C led to a reduction in oil yield, decreasing from 19.5% to 8.2% for FW and from 15.9% to 8.3% for FWSD. Additionally, the yield of aliphatics in the bio-oil decreased from 18.91% to 16.86% for FW and from 33.78% to 25.64% for FWSD. Notably, as the temperature increased, carbon and hydrogen content in the oil increased, while oxygen content decreased. This desirable outcome for improving the oil stability and fuel quality.²² Rice husk pyrolysis oil composition was also influenced by temperature. Acids concentration varied from 7.07 wt% at 400 °C to 5.69 wt% at 600 °C, although it peaked at 8.46 wt% at 450 °C, ketones presented a similar behaviour, going from 8.65 wt% to 4.14 wt% for the same temperature increase. Aldehydes, phenols and nitrogenated compounds all presented a reduction in the concentration. 23 Using a heavy metal contaminated mangrove as the biomass (temperatures range: 300-700 °C) the main compounds found in all cases were phenol derivatives, carboxylic acids, and alcohols. Analysis by GC-MS showed that independently of the temperature, the most abundant alcohol and phenol derivative were methylpropan-1-ol and catechol, respectively, although concentrations varied. The lower temperature (300 °C) led to a higher concentration of 2-methylpropan-1-ol, catechol, syringol and germanicol. For the higher temperatures (500-700 °C), the most abundant compounds were the same ones. A slight variation of the concentration of 2-methylpropan-1-ol and syringol occurred (0.59 and 0.76% variation, respectively), for catechol and germanicol, the lower temperature resulted in a noticeably higher concentration (2.80 and 2.09% variation, respectively).24

Table 2 Chemical composition of pyrolysis oil identified by GC-MS analysis at the optimum conditions of pyrolysis of raw and TB¹⁹

	Relative co (peak area	
Compound	РО-ТВ	PO-RAW
Furan derivatives		
2-Furancarboxaldehyde	10.46	_
2-Furanmethanol	5.62	6.05
2(3 <i>H</i>)-Furanone, 5-methyl	3.52	5.02
2-Furancarboxaldehyde,-5-methyl	_	6.45
Phenol derivatives		
Phenol, 2-methoxy	5.78	_
Phenol, 4-methoxy	_	8.72
Creosol	3.84	6.05
Phenol, 2,6-dimethoxy	9.46	9.11
3,5-Dimethoxy-4-hydroxytoluene	4.63	4.61
Phenol, 2,6-dimethoxy-4-(2-propenyl)	6.68	0.82
Acids		
n-Hexadecanoic acid	0.60	0.54
Linoelaidic acid	_	1.90
Alcohols 1-Penten-3-ol	0.80	0.65
	0.80	2.15
1,2-Benzenediol,3-methyl	0.82	2.15
1-Octen-3-ol,acetate	0.82	_
Aldehydes		0.0
Benzaldehyde, 4-hydroxy-3-methoxy	_	0.8
Benzaldehyde, 4-hydroxy-3,5-dimethoxy	0.64	0.64
Ketones		
Alpha, beta-crotonolactone	2.16	2.16
3-Methylcyclopentane-1,2-dione	2.86	3.93
6-Methoxycoumaran-7-ol-3-one	3.61	_
Sugar derivatives		
1,4:3,6-Dianhydro-alpha-D-glucopyranose	0.30	1.47
Betap-glucopyranose, 1,6-anhydro	_	6.39
Esters		
2-(Acetyloxy) ethyl acetate	0.53	
Diethyl phthalate	_	0.55
Di- <i>n</i> -octyl phthalate	0.35	0.31
Other compounds		
1 <i>H</i> -Pyrazole, 3,5-dimethyl	_	9.34
Benzene, 1,2,3-trimethoxy-5-methyl	3.18	1.91

Phenol derivatives, lignin derivatives, benzene derivatives, alcohols and carboxylic acids are the main components of biooil, and account for the high oxygen concentration and instability of the oil due to low pH, when compared to conventional oils.19,24-27 The viscosity is also highly dependent on the feedstock, as reported by Luo et al.25 when comparing lignocellulosic biomass fast pyrolysis bio-oils. The results of the characterization show a total phenolic, lignin derivative and benzene derivative compound content of 32.26%, 32.36%, 15.78% respectively. All main compounds reported are oxygenated, most of which are aromatics. The results are summarized in Table 3.

Due to its highly complex nature, it is common to separate the bio-oil into hydrophilic, also referred to as water soluble bio-oil (WSBO) and hydrophobic shares. This process can increase the number of quantifiable compounds in the

Table 3 Relative content of main compounds in the organic composition of bio-oil produced from P. indicus²⁵

Compound	Relative content (%)
Furfural	9.06
Acetoxyacetone, 1-hydroxyl	1.21
Furfural, 5-methyl	1.82
Phenol	2.55
2-Cyclopentane-1-one, 3-methyl	1.58
Benzaldehyde, 2-hydroxyl	2.70
Phenol, 2-methyl	5.04
Phenol, 4-methyl	0.51
Phenol, 2-methoxyl	0.27
Phenol, 2,4-dimethyl	9.62
Phenol, 4-ethyl	2.18
Phenol, 2-methoxy-5-methyl	4.15
Phenol, 2-methoxy-4-methyl	0.55
Benzene, 1,2,4-trimethoxyl	3.80
Phenol, 2,6-dimethyl-4-(1-propenyl)	4.25
1,2-Benzenedicarboxylic acid, diisooctyl ester	1.80
2-Furanone	5.70
Levoglucosan	6.75
Phenol, 2,6-dimethoxy-4-propenyl	3.14
Furanone, 5-methyl	0.49
Acetophenone, 1-(4-hydroxy-3-methoxy)	2.94
Vanillin	6.35
Benzaldehyde, 3,5-dimethyl-4-hydroxyl	4.54
Cinnamic aldehyde, 3,5-demethoxy-4-hydroxyl	2.19

analysis, although it remains difficult to identify all components.^{26,27} Naturally, the water-soluble fraction presents a higher concentration of water, alcohols, and carboxylic acids, due to their polar nature. Sipila et al. reported that WSBO from straw, pine and ensyn (hardwood) oils show a high water content (19.9, 11.1 and 23.2% respectively), low pH (3.7, 2.6 and 2.8 respectively), and high carboxylic acid concentration (10.80, 7.30 and 6.20% respectively). Acetic and formic acids were the most significant components and the main reason for the low pH of the oils.²⁷ When two commercial wood pyrolysis oils (BTG and Amaron) were separated in water-soluble (WS) and hydrophobic (oil), the authors reported only 34 directly quantifiable compounds, and the main components are shown in Table 4.26

2.2 Plastics

Characterization of plastic pyrolysis oil shows that the composition is highly dependent on the feedstock and other pyrolysis process factors.¹⁷ Dyer et al. studied co-pyrolysis of biomass with several types of plastics and reported on the composition of different plastic pyrolysis oils; the results are summarized in Table 5. These results show that PET is one of the main sources of oxygen in plastic waste, both being highly produced plastics

Table 4 Compounds identified in the pyrolysis oils and WS fractions by GC-MS (wt%)²⁶

Compound	BTG oil	Amaron oil	BTG WS	Amaron WS
Glycolaldehyde	5.60	1.0	16.10	2.90
Acetic acid	3.90	5.50	0.40	0.70
Acetol	5.60	5.50	1.30	1.20
Levoglucosan	3.50	3.00	0.20	0.20
Syringaldehyde	0.10	0.10	8.10	6.80

Table 5 Characteristics of the pyrolysis oil for different plastic wastes

Proximate analysis (wt%)	HDPE	LDPE	PET	PP	PS
Moisture ^a	0.0	0.0	0.0	0.0	0.0
Ash^b	4.1	0.3	42.7	0.8	1.6
Volatile ^c	98.1	100.0	94.6	100.0	100.0
Fixed carbon ^c	1.9	0.0	5.4	0.0	0.0
Elemental analysis (%)					
C^c	82.5	83.3	69.0	81.0	87.8
\mathbf{H}^{c}	13.1	12.6	5.0	11.4	9.5
O^c	1.8	0.7	32.7	0.6	0.0
N^c	0.1	0.1	0.0	0.1	0.0

^a As received. ^b Dry. ^c Dry ash-free (HDPE: high-density polyethylene; LDPE: low-density polyethylene; PET: polyethylene terephthalate; PP: polypropylene; PS: polystyrene).²⁸

(10 wt% of total resin production)⁴ and having high oxygen composition (32.7%). Although polyolefins (like PE and PP) present a low oxygen concentration, its major role in the plastic production matrix (57 wt% of all resin produced) make it the largest contributor of oxygenated compounds in plastic oil.²⁸

Plastic waste pyrolysis oil usually presents a highly aromatic compound share, when compared to bio-oil, in which can be found monoaromatics, diaromatics, naphthenoaromatics and triaromatics (51.3, 7.5, 5.9 and 2.3 wt% respectively). Some of the main components of the aromatic share of the oil are benzene, toluene, ethylbenzene, xylene, and styrene. The detailed composition of the oil can be found in Table 6.²⁹

The feedstock variation is the main factor in plastic-oil composition In a process treating PET as the feedstock, the resulting pyrolysis oil contains a high concentration of benzene derivatives (10.6 wt%), ketones (5.90 wt%) and acids/esters (62.0 wt%). The most abundant compound in PET pyrolysis oil is benzoic acid and its derivatives, constituting 49.9 wt% of the oil.³⁰ Cit et al. found similar results when comparing polyolefins (PP and LDPE) and PET, where PET pyrolysis oil main components were benzoic acid derivatives (benzoic acid = 23.7, 4-methyl-benzoic acid = 1.6, 4-ethylbenzoic acid = 1.3 and 4-acetylbenzoic acid = 10.3 wt%).³¹ LDPE pyrolysis oil has a high share of carbon doble bonds, as 39% of its composition are aromatic compounds and other unsaturated carbon molecules.³² A mix of LDPE and HDPE led to naphthalene derivates as the main components (approximately 24 wt%) in the plastic-oil, followed by p-xylene, toluene, naphthalene, benzene, 1-ethyl-4-methyl-benzene, m-xylene, 1,2,4-trimethylbenzene, indane, indene and 1-methyl-3H-indene (approximately 19, 14, 9, 6, 3.5, 3, 2, 1.5, 1.5 and 1 wt% respectively).³³ For PS, monoaromatics are the main components, followed by

Table 6 Detailed composition of the plastic pyrolysis oil. Edited with permission from ref. 29

Component	Composition (wt%)
Paraffins	5.0
Iso-paraffins	8.2
Olefins	12.3
Aromatics	67.1

Table 7 Main components of plastic-oil³⁵

Compound	Composition (wt%)
Xylene	8.80
Benzene	5.30
Chlorinated phenols	4.20
Naphthalene derivatives	4.10
Cresols	3.90
Naphthalene	3.70
Cyclopentene	2.90
Toluene	2.70
Benzene derivatives	1.00
Cyclobutane	0.90

polyaromatics and aliphatics (77.62, 12.86 and 0.01 wt% respectively). As expected, the main component is styrene (73.60 wt%), other noticeable compounds being 2,4-diphenyl-1-butene (7.80 wt%), 2,4,6-triphenyl-1-hexene (2.21 wt%), α-methylstyrene (1.61 wt%), and toluene (1.58 wt%).³⁴ When conducting a study on the performance of plastic pyrolysis oil as fuel, Kalagaris et al. reported on the composition of the oil. The feedstock was a mixture of styrene butadiene, polyester, clay, ethylene-vinyl acetate, rosin, polyethylene, and polypropylene (47, 26, 12, 7, 6, 1 and 1 wt% respectively). The main components of the oil are presented in Table 7.35

The addition of different feedstocks may also change the composition of the oil; that is, the oil composition of a PS/PP mixture may not be just the sum of PS and PP oils. Many of the components found in PP pyrolysis oil, such as methylnaphthalene (8.4 wt%), xylene (7.8 wt%) and phenanthrene (7.6 wt%), completely disappeared when the feedstocks were mixed. Fig. 3 shows the oil composition for several different feedstock compositions. The ratio of each plastic in each mixture was 50/50%, and in the last mixture (PS, PP, PE and PET) the ratio was 40/20/ 20/20%, respectively.36

Both temperature and pressure are vital factors in the pyrolysis process. Pyrolysis of LDPE under different conditions $(425 \, ^{\circ}\text{C}/1.60 \, \text{MPa}, 450 \, ^{\circ}\text{C}/2.45 \, \text{MPa} \text{ and } 500 \, ^{\circ}\text{C}/4.31 \, \text{MPa}) \, \text{led to}$ variation in both the classes of compounds and their concentration. The carbon chain length also changed, significantly reducing at higher temperatures/pressures. Table 8a shows the variation of the macro composition and Table 8b the specific compounds.³⁷

The same variation can be observed in the pyrolysis of PS (process conditions: 400 °C/1.14 MPa, 425 °C/1.26 MPa, 450 °C/ 1.47 MPa, and 500 °C/1.60 MPa). Results are summarized in Table 9.

Long chain alkanes and alkenes are common products in plastic-oil, and the carbon chain lengths can reach as high as 55 carbons (C₅₅) when the feedstock is a mixture of PP, LDPE, HDPE and styrene. 14,15 The carbon number range of alkanes (Table 10a) and alkenes (Table 10b) are as follows:

Plastic waste (polyethylene film and mixed polyolefin) pyrolysis oils blended with fossil naphtha led to an even higher carbon number range. For polyethylene film (PE-film), was C₅-C₅₅ and for mixed polyolefin (MPO) was C₅-C₅₂. The composition groups of the resulting pyrolysis oils consisted of

paraffins, iso-paraffins, α-olefins, iso-olefins, diolefins, naphthenes, monoaromatics and naphthenoaromatics. 15 Both oils also present a substantial amount of several halogen and metal contaminants, among which are found Cl, Br, Al, Ca, Cr, Cu, Fe, Li, Na, Ni, Sb, Si, Sr and Zn. Both Cl and Br can be explained from the initial composition of the feedstock, as polyvinyl chloride (PVC) is a common plastic and bromine is an element that is used in fire retardants. Metal contamination can be traced to plastic enhancers, as metals are commonly used as additives. 15 The detailed composition of the plastic-oil and the distribution of paraffins, olefins, naphthenes and aromatics over the light, medium and heavy product fractions of all post-consumer plastic waste pyrolysis oils can be found in Table 11.16

2.3 Pyrolysis oil comparison

Although both biomass pyrolysis oil (bio-oil) and plastic pyrolysis oil (plastic-oil) share common components such as paraffins, olefins, aromatics, and oxygenated compounds, they exhibit significant differences in their composition. Notably, bio-oil water content is much higher than plastic oil.8,12,14-16 On the other hand, plastic-oil presents a higher aromatic share, longer carbon chains and various metal contaminants, added during its production as stabilizers and colouring, for example. The presence of long chain alkanes and alkanes in plastic-oil can lead to a heavier and more viscous product and is a factor that will affect the upgradation process, making it slower. 15

For utilization as biofuel, HHV is a vital factor. Bio-oil has a lacklustre HHV when compared to transportation fuels (bio-oil HHV: 13-19 MJ kg⁻¹ compared to gasoline: 44-46 MJ kg⁻¹ or coal: 25-35 MJ kg⁻¹). Plastic-oil, on the other hand, presents a high HHV (41-47 MJ kg⁻¹).8

The main components from both bio and plastic oil are summarized in Fig. 4.

3 Upgrading routes

Bio and plastic oils usually present subpar physical-chemical properties when compared to traditional fuels, such as acidity, O:C and H:C ratios and, for bio-oils, low HHV. This means that for its utilization as an alternative fuel, an upgradation process is necessary. The same is true when it comes to recycling, where these products need to be upgraded to be utilized as feedstock for the petrochemical or chemical industries. Several processes for bio-oil upgradation have been the focus of many studies throughout the years. 9,10,13,38-50 The most common method is deoxygenation, which is commonly accomplished through catalytic cracking and hydroprocessing routes such as hydrotreatment and hydrodeoxygenation.³⁸

Upgrading routes can be separated into chemical (hydrotreatment, catalytic cracking, hydrocracking, steam reforming, hydrodeoxygenation and hydrogenation) and physical (distillation, supercritical fluid extraction, liquid-liquid extraction, and emulsification) processing. There are several advantages and disadvantages of each upgrading route; some of the

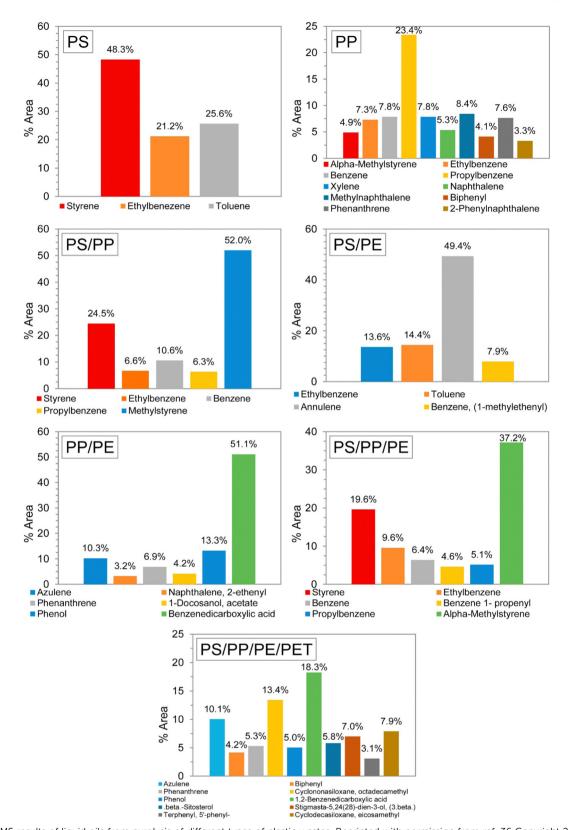


Fig. 3 GC-MS results of liquid oils from pyrolysis of different types of plastic wastes. Reprinted with permission from ref. 36 Copyright 2017 Elsevier.

disadvantages of the more traditional routes are the operational conditions, high costs and coke production. Catalytic cracking is accomplished at medium temperature and atmospheric pressure, but it leads to catalyst deactivation and the formation of carbon double bonds, which would require further improvement.⁵¹ Decarbonylation and decarboxylation leads to the loss of

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Table 8 (a) Composition of the oil products from the pyrolysis of LDPE (% w/w oil).37 (b) Main components of the oil products from the pyrolysis of LDPE (% w/w oil)³⁷

Composition	425 °C/1.60 MP	a 450 °C/2.45 Mpa	500 °C/4.31 Mpa
(a)			
Naphthenes	2.69	5.56	1.50
Alkanes	46.20	31.7	17.8
Alkenes	12.40	13.1	3.58
Aromatics	12.00	22.9	68.0
Unknowns	19.10	17.9	_
(b)			
Heptane	2.26	3.16	0.46
Toluene	0.35	3.05	24.30
Octane	2.45	3.78	0.19
Ethylbenzene	0.25	0.89	3.06
p/m-Xylene	0.62	1.79	3.90
Allylbenzene	0.21	1.02	4.84
Methylnaphthalene	0.90	0.36	4.22
C ₂₁ -Č ₃₀ alkanes	9.98	1.85	1.20

Table 9 Main components in the pyrolysis oil products from PS (%w/w)³⁷

Compound	400 °C/ 1.60 MPa	425 °C/ 2.45 MPa	450 °C/ 4.31 MPa	500 °C/ 4.31 MPa
Benzene	0.38	0.73	1.27	1.63
Toluene	21.70	23.10	27.00	28.40
Ethylbenzene	32.60	39.30	39.00	36.60
Styrene	1.09	0.80	0.39	0.61
Cumene	10.20	9.10	9.36	9.60
Propylbenzene	0.60	0.77	0.90	1.29
Methylstyrene	1.37	0.39	0.33	0.55
Diphenylmethane	0.91	0.83	1.00	1.61
1,2-Diphenylethane	1.44	1.16	0.60	0.37
Phenylnaphthalene	0.98	1.02	1.01	1.25
1,2-Diphenylbenzene	1.27	1.26	1.87	1.89
Triphenylbenzene	6.30	5.13	3.83	3.18

Table 10 (a) Carbon number range of alkane products of produced pyrolysis oil in wt%. Adapted with permission from ref. 14. (b) Carbon number range of alkene products of produced pyrolysis oil in wt%. Adapted with permission from ref. 14

Feedstock	C_{11} – C_{20}	C_{21} – C_{30}	C_{31} – C_{40}	C_{41} – C_{50}
(a)				
Polypropylene	2	63	0	0
LDPE	6	54	12	13
HDPE	1	67	18	0
Styrene	0	4	27	8
(b)				
Polypropylene	9.9	8.9	0	0
LDPE	3.3	4.2	0	0.8
HDPE	0.5	6.2	1	1
Styrene	0	0	0	10

valuable carbon. Hydrocracking, hydrodeoxygenation, and hydrogenation require an outside source of H2, which is costly and difficult to store (see Fig. 5). 43,44

Electrochemical hydrogenation

One recently emerging route for pyrolysis oil upgradation is electrochemical processing. Electrochemical hydrogenation

Table 11 (a) Detailed composition of the plastic pyrolysis oil in wt%. Edited with permission from ref. 15. (b) Detailed composition of the plastic waste pyrolysis oil in wt%¹⁶

Composition	MPO pyrolysis oil	PE-film pyrolysis oi
Paraffins	32.2	41.9
Iso-paraffins	3.7	3.5
α-Olefins	23.5	33.0
Diolefins	5.3	4.5
Naphthenes	12.4	5.5
Monoaromatics	2.7	1.0
Naphthenoaromatics	0.2	0.1

(6)			
Composition	Light fraction ^a	Medium fraction b	Heavy fraction ^c
Paraffins	28	29	44
Olefins	32	40	29
Naphthenes	18	14	17
Aromatics	19	12	11
Others	4	5	0

^a Carbon number up to C₂₀ and boiling point up to 250 °C. ^b Carbon number up to C₃₀ and boiling point up to 450 °C. ^c Carbon number >C₃₀ and FBP >450 $^{\circ}$ C.¹⁶

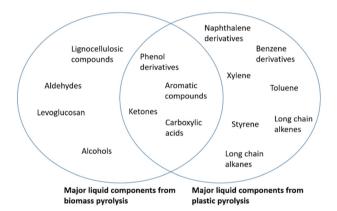


Fig. 4 Intersection in the composition between bio-oil and plastic-oil.

(ECH) does not require H₂ from an outside source, which is produced during the process, reducing costs, and mitigating the risks associated with H₂ storage. ^{18,39,51} Most setups for ECH consist of two chambers separated by a cathode exchange membrane (CEM). The most prevalent cell design is H-type cells (Fig. 6), although there are some studies utilizing flow type cells. 40,52,53

During ECH, water is oxidised at the anode (eqn (1)), the resulting H+, that can flow through a cation exchange membrane into the cathode cell. The H⁺ is then chemisorbed by the catalytic cathode (eqn (2)), where M is the metal surface and H_(ads)M is the chemisorbed hydrogen.^{54,55}

$$H_2O \leftrightarrow H_{(aq)}^+ + 2e^- + \frac{1}{2}O_2$$
 (1)

$$H_{(aq)}^{+} + e^{-} + M \leftrightarrow H_{(ads)}M$$
 (2)

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Cracking:
$$R_1 \xrightarrow{CH_2} CH_2 \xrightarrow{CH_2} CH_2 \xrightarrow{R_2} \longrightarrow R_1 \xrightarrow{CH} CH_2 + \xrightarrow{H_2C} CH_2$$

Decarbonylation: $R_1 - C \xrightarrow{H_2CH_2} \longrightarrow R_1 - H + CO$

Decarboxylation: $R_1 - C \xrightarrow{H_2CH_2} \longrightarrow R_1 - H + CO_2$

Hydrocracking: $R_1 \xrightarrow{CH_2} CH_2 \xrightarrow{R_2} + H_2 \xrightarrow{H_2CH_2} R_1 - CH_3 + H_3C - R_2$

Hydrodeoxygenation: $R - OH + H_2 \longrightarrow R - H + H_2O$

Hydrogenation: $R_1 \xrightarrow{H_2CH_2} \xrightarrow{H_2$

Fig. 5 Examples of reactions associated with catalytic bio-oil upgradation. Reprinted with permission from ref. 44 Copyright 2011 Elsevier Inc.

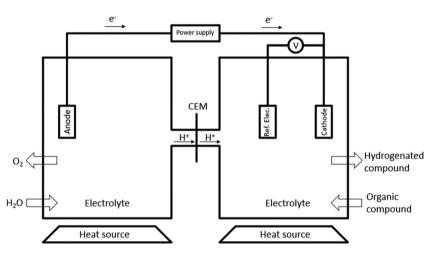


Fig. 6 A typical H-type cell, where the two chambers are separated by a membrane

In the cathode chamber, the organic unsaturated compound is chemisorbed onto the cathode (eqn (3)), where the hydrogenation process occurs (eqn (4)), followed by the desorption of the stabilized product (eqn (5)). Eqn (6) is the net reaction. 54,55

$$(Y = Z)_{aq} + S \leftrightarrow (Y = Z)_{(ads)}S \tag{3}$$

$$2H_{(ads)}M + (Y = Z)_{(ads)}S \leftrightarrow (YH-ZH)_{(ads)}S + 2 M$$
(4)

$$(YH-ZH)_{(ads)}S \leftrightarrow (YH-ZH)_{(aq)} + S$$
 (5)

Unsaturated compound + nH^+ + $ne^{-1} \leftrightarrow$ stabilized product

where $(Y = Z)_{aq}$ is the unsaturated compound, S is the absorption site of the cathode, $(Y = Z)_{(ads)}S$ is the adsorbed organic compound, (YH-ZH)(ads) is the hydrogenated product before desorption, (YH-ZH)(aq) is the final product and n denotes any arbitrary numerical value. 54,55 The hydrogen evolution reaction (HER) is a common side reaction, that can lead to the loss of hydrogen, rather than reaction with bio-oil, thus reducing the faradaic efficiency (FE), smaller conversion rates, and a poorer overall process. This reaction can occur via either the Tafel step (eqn (7)) or the Heyrovsky step (eqn (8)). 54,55

$$2(H_{(ads)}M) \leftrightarrow 2H_2 + 2M \tag{7}$$

$$H_{(ads)}M + H_{(aq)}^{+} + e^{-} \leftrightarrow H_{2} + M$$
 (8)

ECH can be performed under mild conditions (<80 °C and 1 atm) (Table 12). Lower temperatures can also avoid catalyst deactivation by coke formation and membrane fouling, potentially reducing the costs associated with catalyst purchase and recycle.⁵⁶ However, due to high viscosity and low conductivity, it requires the utilization of membranes, catalysts, and electrolytes to facilitate the reaction. 10,13,38,39 That leads to a vast spectrum of different arrangements that can be organized to optimize the process.

3.1.1 Key ECH performance parameters. To understand and quantify the ECH process, the most important parameters **Energy Advances** Review

Table 12 Methods for bio-oil upgradation comparison

Method	T (°C)	P (bar)	H ₂ need
Cracking	330-700	1.0	No
Hydrocracking	> 400	68-96	Yes
Hydrodeoxygenation	300-600	80-300	Yes
Electrochemical hydrogenation	< 80	≈1	No

are faradaic efficiency (eqn (9) and (10)), conversion (eqn (11)), selectivity (eqn (12)) and current density (eqn (13)). Due to difficulties in measuring the surface area of the cathode, and therefore the current density, some authors report this parameter as current (usually in mA), these will be highlighted in the following sections.

FE (%) =
$$\frac{(I \times T - 2 \times n \times F)}{(I \times T)}$$
 (9)

where I = current in A; T = time in s; n = moles of hydrogen; F = Faraday's constant.

This equation can also be simplified:

FE (%) =
$$\frac{\text{Electrons used to generate products}}{\text{Total electrons passed}} \times 100$$
 (10)

Conversion (%) =
$$\left(\frac{\text{Moles reactant consumed}}{\text{Initial moles reactant}}\right) \times 100$$
 (11)

Selectivity (%) =
$$\left(\frac{\text{Moles desired product}}{\text{Total moles product}}\right) \times 100$$
 (12)

Current density (mA cm⁻²) =
$$\frac{\text{Current}}{\text{Surface area}}$$
 (13)

3.2 ECH reactor design considerations

3.2.1 Electrodes. Electrodes usually consists of conductive materials and catalyst, which play a vital part in the ECH process. Different electrodes can increase bio-oil conversion and interfere with product selectivity.⁵¹ Noble metals,^{39,51} base metals, 41,57,58 and graphite, 59 besides catalyst-coated carbon cathodes (e.g. Ru/ACC) 39,41,56 have been reported for ECH biooil and model compounds upgrading. Pt, Rh, Ru, Au and Pd have been employed as cathode materials during ECH of aldehyde and phenolic model compounds. 51,60,61

The choice of cathode can greatly interfere with the process, as early transition metals (e.g. Ti, Mn, Nb) and platinum group metals (Ru, Rh, Ir, and Pt) have higher affinity toward hydrogen formation, leading to more H+ in solution, which increases collision probability and, in turn, leads to higher conversion rates.⁵¹ Cathode surface coverage (or adsorption rate) can lead to the same problem of low collision probability, and is dependent on cathode/compound pairing, for example, the adsorption of phenol on Pt is stronger compared to guaiacol. 42,62

Cathode selection can also influence the selectivity of the process and the desirable product yield. For the ECH of glucose, late transition metals (e.g. Fe, Cu, Pd, Ag and Au) led to higher sorbitol formation and post-transition metals (In, Sn,

Sb, Pb, and Bi), Zn and Cd (d metals) favoured d 2-deoxysorbitol formation. Ni had the lowest and Pb the highest sorbitol formation potential.63

Platinum group metals (Pt, Pb, Ru, Rh and Ir) are commonly used in hydrogenation processes, due their stability and high affinity towards hydrogen production and have been employed under several conditions and for different compound classes. 63 For the ECH of indigo, a highly unsaturated compound that contains both oxygen and nitrogen (reaction conditions: 1 M NaOH, 50 °C, 20 mA cm⁻²) using RANEY[®] nickel electrodes doped with different noble metal catalysts (Pt, Pd and Pd/C), Pt coating led to both optimal conversion and FE (10.2 and 8.16% respectively), followed by Pd (8.6, 6.88%), Pd/C (4.7, 3.76%) and RANEY[®] nickel without coating (4.4, 3.5%).⁶⁴ Nobel metals are also capable of hydrogenating phenolic compounds at high conversion rates and FE. ECH of phenol on several cathodes (Pt/C, Rh/C, and Pd/C) demonstrated that Rh/C exhibited the highest hydrogenation rate, the order was Rh/C > Pt/C > Pd/C.65 Du et al. reported on the ECH of phenol on several different electrocatalysts: Pt, Ru, Pt₃Ru, Pt₃Sn, Ru₃Sn, and Pt₃RuSn supported on carbon cloth (reaction conditions: 0.2 M H₂SO₄ containing 10 mM phenol at −100 mA, 50 °C for 100 min). Pt₃Ru/CC presented the highest conversion rate, at 96.3%, followed by Pt₃RuSn/CC and Pt/CC (91.5 and 90.2% respectively). 66 The concentration of the metal on the surface of the carbon can also lead to different results, although graphite alone was unable to react (reaction conditions: phenol, electrolyte 0.1 mol L^{-1} H₂SO₄ (30 mL), 30 mA, Pt sheet as the anode). 1% Pt/G and 1.5% Pt/G presented the highest conversion rates, both 95%. The highest yield of cyclohexane and the highest FE were 30.4% and 20.3%, respectively, both occurred with the same cathode (1.5% Pt/G). The results are summarized in Table 13.55 As shown in previous sections, lignin pyrolysis oil can contain phenolic compounds (phenol, p-cresol, 4-ethylphenol, and 4-propylphenol, as well as guaiacol) which can be hydrogenated into their corresponding alkylcyclohexanols at high conversion rates using Ru/ACC as the catalytic cathode. The same cathode is effective for transforming model lignin monomers into cyclohexanol.⁵⁶

Metals outside the platinum group can also be utilized as the catalytic cathode, although it still requires more investigation. The ECH of guaiacol and related lignin models utilizing a

Table 13 ECH of phenol on various cathodes⁵⁵

Cathode	Conversion (%)	F.E.	1	2	3	4
Pt	0	0	0	0	0	0
Ni/G	0	0	0	0	0	0
Rh/G	90	0	0	65.7	24.3	0
Pd/G	18	0	0	12.6	5.4	0
0.5% Pt/G	92	11.1	16.6	57	12	5.5
1% Pt/G	95	15.2	22.8	52.3	16.2	3.8
1.5% Pt/G	95	20.3	30.4	58.9	5.7	0
2% Pt/G	90	9.1	13.5	60.3	9	7.2
Graphite	0	0	0	0	0	0

Where [1] is cyclohexane, [2] cyclohexanol, [3] cyclohexanone and [4]

nickel cathode and a Co/phosphate anode presented high selectivity towards cyclohexanol formation (>99%), with only traces of 2-methoxycyclohexanol. Other experiments were done utilizing cobalt and copper as catalyst cathodes, and only Co formed trace amounts of cyclohexanol and phenol, Cu completely failed in producing these products. The ECH of aldehyde compounds shows that aromatic aldehydes have a higher conversion rate than aliphatic aldehydes. Au, Ag, Cu, and C catalysts exhibit the highest conversion to alcohol products.

3.2.2 Electrolyte. Electrolytes can affect the ECH of bio-oil in three major ways. First, the adsorption of anions from the electrolyte into the electrode can make the process difficult. Second, the pH of the solution has a direct relation to product selectivity. Lastly, alcohol has been reported to reduce conversion rate and faradaic efficiency. The addition of different alcohols (methanol, ethanol, or isopropanol) to the electrolyte led to lower conversion rates and faradaic efficiency as the alcohol concentration increased. Carbon chain length was a factor, as the impact was greater for longer chains.

Electrolytes are needed during ECH to increase the conductivity of the system, although they can interfere with the process. Acids, alkalis, and salts can achieve these results, and the selection of electrolytes should take into consideration both the organic compound and the cathode. Another challenge during this selection is that the electrolyte that presents the highest conversion rate and the one that leads to the highest FE might not be the same one. For the ECH of guaiacol (reaction conditions: 3-CE-NH₃ cathode, 80 °C and 100 mA), the conversion rates, FE and product selectivity varied greatly. The highest conversion rates were NaOH (62 \pm 2.4%), NaCl (64 \pm 8.8%), and HCl (75 \pm 3.9%). For FE, the order was NaCl $(20 \pm 2.2\%)$ < NaOH $(28 \pm 1.0\%)$ < HCl $(30 \pm 4.3\%)$. Similar results were reported for phenol and syringol.³⁹ The pH of the system can interfere with the results, as demonstrated by the ECH of guaiacol where HClO4 as the electrolyte led to the highest guaiacol conversion rates, HCl and H2SO4 led to moderate conversion rates (PtNiB/CMK-3 cathode, 0.2 M HClO₄, 0.2 M HCl and 0.2 M H₂SO₄ as electrolytes). In addition, HClO₄ achieved optimal selectivity for the desirable product (KA oil). However, increasing the concentration of sulfuric acid from 0.2 M to 0.5 M reduced the ECH activity. 42

Different catholyte/anolyte pairings can also lead to vastly different results. Acid/acid pair (catholyte/anolyte respectively) lead to higher guaiacol conversion. (Conditions: $T=50\,^{\circ}\mathrm{C}$, $I=300\,\mathrm{mA}$ ($j=-109\,\mathrm{mA}$ cm $^{-2}$), catalyst = 5 wt% Pt/C (0.133 g), reaction time = 2 h, concentration of all electrolytes = 0.2 M) followed by salt/acid and acid/alkali. Alkali/acid completely failed in generating products. ECH of phenol led to similar results. 62

3.2.3 Temperature. Temperature is a factor that can greatly influence the results of ECH. FE, conversion rates and product selectivity have been reported to be affected by temperature. ^{39,51,69} Lower temperatures favours electrocatalytic hydrogenation, whilst high system temperature is favourable for hydrogen desorption, which leads to the low ECH performance.

This condition increases the water oxidation rate on the anode, which contributes to hydrogen evolution reaction (HER), a competing reaction, due to $\rm H^+$ generation and transfer to the cathode. For the ECH of indigo, increasing the temperature from 30 to 60–80 °C led to an FE increase, although it substantially dropped down at higher values. Optimal reported temperature was 70 °C. Gauiacol (HCl as catholyte, T=2 h, 100 mA) behaves in a similar way, where raising the temperature from 25 °C to 50 °C increased FE from 8% to 17% but further increase to 80 °C dropped it back to 10%. Increasing temperature from 5 to 40 °C during phenol to cyclohexanol ECH upgrading significantly increased FE, upon further heating to 60 °C, FE increase was continuous, but minimal.

3.2.4 Current and potential. Current density can have a direct impact on product selection, FE, and conversion rates, as it can lead to higher H_{ads} desorption from the cathode and thus become an obstacle for the main reaction. 51,54,55 Although higher current densities could decrease the overall process quality, insufficient current can do the same as insufficient current can lead to a lower concentration of Hads on the cathode surface. This reduces collision probability and, consequently, reduces FE.51 Current is directly linked to potential values, and therefore can also interfere in the reaction pathways and product selectivity. The major side reaction that can occur during the ECH process is the Hydrogen Evolution Reaction (HER), which leads to lower FE and product yield. The HER usually occurs by Tafel or Heyrovsky reactions (eqn (7) and (8)). 51,57 The surface coverage of Hads is determined by the kinetics of eqn (2) and side reactions, eqn (7) and (8), which can be increased or decreased depending on potential. Therefore, potential is an essential factor of the ECH process. 54,57 For the ECH upgrading of phenol, an increase on current from the range of 10-30 to 30-70 mA slightly lowered product yields and FE. Further increase on the current to over 70 mA highly decreased the overall process quality.55 Furfural behaves in a similar way (desirable product: furfuryl alcohol and 2methylfuran) where the increase of current density from 300 to 1200 mA dm⁻² led to a decrease in FE.⁵⁴ Current density increase during the ECH of lactic acid to lactaldehyde and propylene glycol (0.01 M H₂SO₄ for 9 h at 90 °C) led to a lower product yield⁷¹ (Tables 14 and 15).

4 ECH systems for model compound and bio-oil upgradation

4.1 Model compounds

Zhou *et al.* proposed a new catalyst for electrocatalytic hydrogenation (ECH) of guaiacol, a boron-doped PtNiB/ordered mesoporous carbon (CMK-3) cathode on a Nafion 117 membrane and IrO_2/C as the anode. The goal was the production of KA oil (a mixture of cyclohexanol and cyclohexanone) which is highly utilized as industrial feedstock. ⁴² The working temperature was 60 °C. The anolyte was 10 mL 0.2 M $HClO_4$ and catholyte was 10 mL 0.2 M $HClO_4$ with 10 mM guaiacol.

Table 14 Previous studies comparison table

Cathode	Feedstock	Conversion	Electrolyte	Temperature (°C)	J (mA cm ⁻²)	F.E.	Duration (h)	Ref.
Ru/ACC	Guaiacol	64 ± 8.8	NaCl			20 ± 2.2		
		75 ± 3.9	HCl	80	100^{a}	30 ± 4.3	2	39
	Phenol	89	HCl			29		
	Syringol	58	HCl			29		
PtNiB/CMK-3	Guaiacol	>99%	$HClO_4$	60	20^a	86.2	2	42
RANEY [®] -Ni	Guaiacol	>99%	BK_3O_3	75	8	26	6	58
Ru/ACC	Phenol	100				20	4	
	4-Methylphenol	100	HCl	80	100^a	9		56
	4-Ethylphenol	98				9	6	
	4-Proprylphenol	100				6		
RANEY [®] -Ni		4.4				3.5		64
RANEY®-Ni with Pt	Indigo	10.2	NaOH	50	20	8.16	_	
RANEY®-Ni with Pd	Ü	8.6				6.88		
RANEY®-Ni with Pd/C		4.7				3.76		
Iron	Furfural	97 ± 1.4	NH_4Cl	_	6	29 ± 3.5	5	54
Pt/CC		90.2	•			34.6		
Ru/CC		73.4				28.2		
Pt ₃ Ru/CC	Phenol	96.3	H_2SO_4	50	100^{a}	37.6	1.67	66
Pt ₃ Sn/CC		73.3	2 4			29.7		
Ru ₃ Sn/CC		62.3				26.7		
Pt ₃ RuSn/CC		91.5				39.5		
^a Reported as mA.								

Table 15 Different setups and studies comparison table

Feedstock	Cathode	Catholyte	Anode	Anolyte	T (°C)	Duration (h)	J (mA cm ⁻²)	Ref.
Guaiacol	Boron-doped PtNiB/ordered mesoporous carbon (CMK-3)	0.2 M HClO ₄	IrO ₂ /C	0.2 M HClO ₄	60	5	40^a	42
Phenol derivatives	Ru/ACC	0.2 M HCl	Pt wire	0.2 M phosphate	80	_	100^{a}	56
WSBO	Ru/ACC	0.2 M NaCl	Pt wire	1 M H ₂ SO ₄	27	6.5	480^{a}	72
WSBO	Ru nanoparticles on mesoporous carbon-Ru@OMC	WSBO	Fe(III)/Fe(II) + carbon felt	Wastewater or lignin	_	3	_	41
Resin wood (ponderosa pine) bio-oil	Metal-free (carbon)	1 M Na ₂ SO ₄	Ti sheets/ Pt coating	Purified water	35(±2)	_	_	40
^a Reported as mA.								

The flow rate was 50 mL min⁻¹. Constant current of 20 mA for 2 h. The setup is shown in Fig. 7.42

Faradaic efficiency (FE) of the process was 86.2%, in comparison to the 6.3% that was found when the reaction was carried out in a PtNi/CMK-3 cathode, that is, without boron doping. The conversion rate was respectively <99.0% and 11.7%. The study also reported on the ECH of other phenolic compounds using PtNiB/CMK-3. All achieved high (<95%) conversion rates and high (<80%) FE. The temperature increase effect on conversion rate and FE on range of 20-60 °C was positive (conversion rate: from 58% to 98.9% and FE: from 45% to 86.2%), but further heating to 80 $^{\circ}$ C reduced these parameters to 78% and 57% respectively (Fig. 8).

Garedew et al. studied the ECH of phenolic compounds. Several model compounds were selected, including catechol, anisole, phenol, guaiacol, 3-methoxyphenol, 4-methoxyphenol, syringol, 1,3-dimethoxybenzene, o-cresol, m-cresol, p-cresol, creosol, vanillin, syringaldehyde, eugenol, 4-ethylphenol, 4-propylphenol, 4-propylguaiacol and 4-ethylguaiacol.⁵⁶

The cell setup was a H cell, separated by a Nafion 117 membrane. The cathode was Ru/ACC and the anode a Pt wire. Catholyte was a 0.2 M HCl solution (30 mL) and the analyte a 0.2 M phosphate buffer (30 mL). Temperature was maintained at 80 °C and the experiment was run at 100 mA. The cell setup is shown on Fig. 9.56

The authors reported that increasing guaiacol concentration from 5 to 20 mM increased the FE and product selectivity. Further increases from 20 to 60 mM had led to a steep decline in the conversion rate, although this was mitigated by increasing the duration of the process (Fig. 10).

Phenol, p-cresol, 4-ethylphenol, and 4-propylphenol, as well as guaiacol, were transformed into their corresponding alkylcyclohexanols at high conversion rates (>98%). Carbon chain length was a determinant factor in reaction speed, as increased lengths resulted in slower conversions.⁵⁶

4.2 Water soluble bio-oil (WSBO)

Li et al. proposed ruthenium (Ru) supported on activated carbon cloth (ACC) as the catalytic cathode as the electrocatalyst.

0, H HO PtNiB/CMK-3

Fig. 7 Schematic of ECH of quaiacol into KA oil integrated with O2 production, equipped with PtNiB/CMK-3 and IrO2/C as the cathode and anode, respectively. Reprinted with permission from ref. 42 Copyright 2019 Wiley.

The bio-oil was mixed with water, in water to bio-oil ratio of 4:1, with bio-oil final concentration of 15 wt%. The mixture was then centrifuged, and the top layer was separated as the water-soluble bio-oil (WSBO).⁷² The same group has proved that this cathode was capable of hydrogenating guaiacol, syringol and phenol, as showed in the previous sessions.³⁹

Electrochemical hydrogenation of WSBO was carried out in a two-chamber glass H-cell, separated with a DuPont® Nafion117 membrane (see Fig. 11). The reaction temperature was 27 °C and the pressure was 1 atm. Analyte consisted of 1 M sulfuric acid and Pt wire was used as the anode. A mixture of WSBO with 0.2 M NaCl served as the catholyte. Current was 480 mA.⁷²

After the 6.5 h treatment, most of the organic compounds were depleted or fully converted by hydrogenation, the proposed mechanisms are shown in Fig. 12. Removal of >50% of acetic acid is an expressive result, as it is a major contributor to bio-oil corrosiveness. No coke formation was observed. The same experiments were done using only ACC as the cathode and showed that Ru is an essential part of the proposed ECH process.³⁹

Although a noble metal catalyst (e.g., Pt and Ir) as the anode is normally used to ensure effective water oxidation, Zhang et al. found that a Fe(III)/Fe(II) redox pair can be utilized as a substitute when combined with carbon felt as the anode. This change can make the process less expensive and can further its usage. The authors presented a Ru-based ECH cathode catalyst by uniformly embedding Ru nanoparticles into the surface of the ordered mesoporous carbon (OMC) support (noted as Ru@OMC). The catholyte solution was WSBO, duration was 3 h.41 The cell setup is shown in Fig. 13.

The EHC treatment led to an increase in the H: C ratio and a decrease in the O:C ratio. Another result was the content reduction in several organic groups, such as acids, esters, carbonyls, phenols, sugars, and furans. The alcohol content increase was substantial, and the process elevated the selectivity of polyhydric alcohols. The results infer that this technique provides de ability to saturate unsaturated components in WSBO.41

4.3 Whole oil

Lister et al. employed both an anion exchange membrane (AEM) and a cation exchange membrane (CEM), in what is described as a dual exchange membrane (DEM) to upgrade a whole pyrolysis bio-oil mixed with an aqueous phase containing acetic acid and formic acid. The concentrate vessel was filled with 10 mL solution of 1 M Na₂SO₄. The oil was filtered to remove any char particles, a common product from pyrolysis. Fig. 14 shows a diagram of the electrolysis system used in this work.40

The cell temperature was held at 35 °C (± 2 °C), the fluids flow rate was 2 mL min⁻¹. A metal-free carbon catalytic cathode was chosen to prevent hydrogen evolution reaction (HER). The process achieved an increase in the pH from 2.6 to >4and a significant reduction in total acid number (TAN). There were no signs of coke formation, improving carbon efficiency and catalyst lifetime. Although membrane degradation was noticed.40

5 Outlook of previous studies

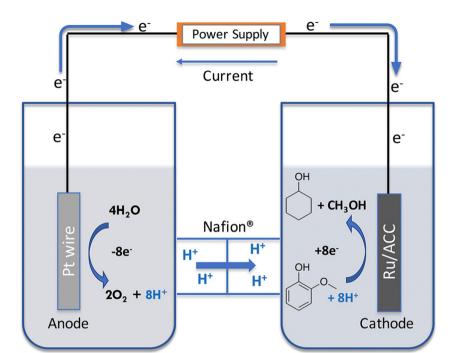
The electrochemical approach to upgrading bio-oils is a technique that can be performed at mild temperatures (<80 °C) and pressure (1 atm). This is an essential condition for a

Entry	Structural formula	t (min)	Conv. (%)	FE (%)	Product (sel. %)
1	OH OH	50	97.5	90.8	OH OH 10.1
2	OCH3	60	95.8	92.6	OH OH 39.3
3	OH OCH ₃	60	95.1	88.1	OH 0 34.4
4	OH OCH ₃	90	98.7	84.3	OH 0 12.1
5	OCH ₃	90	97.5	86.2	83.5 16.1 O
6	H ₃ CO OH OCH ₃	90	98.6	80.2	OH OH

Fig. 8 ECH of other phenolic compounds using PtNiB/CMK-3. Reprinted with permission from ref. 42 Copyright 2019 Wiley.

cheaper product, as traditional methods are energy intensive. Also, in thermal stabilization and deoxygenation, oxygen is released as CO_2 , CO, and H_2O . These products represent a loss of expensive carbon and H₂. 40 The research gap that exists is on what are the ideal catalysts, membranes, cell configuration and how to increase the oil's low conductivity. Understanding the mechanism of the reaction and the behaviour of the compounds is a key part of the process, so model compound studies are necessary. Another way of simplifying the process so it can

be more easily studied is utilizing WSBO as a feedstock; this allows the study to circumvent the challenges brought by the complexity of a mixture that contains both polar and non-polar compounds. Most studies have focused on either model compounds or WSBO, and therefore ECH of whole pyrolysis oil still needs further investigation. Zhenglong et al. demonstrated that the separation of WSBO can lead to a higher organic compound conversion. Ruthenium was used as the basis for the catalyst cathode, this metal was also utilized by Zhang et al., Zhou et al.



Electrocatalytic hydrogenation in a two-chambered H cell. Reprinted with permission from ref. 56. Copyright 2019 American Chemical Society.

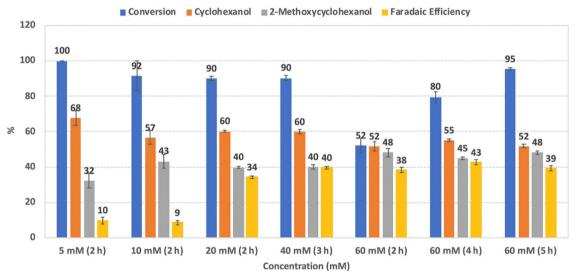


Fig. 10 Conversion, product selectivity, and faradaic efficiency of quaiacol reduction using different substrate concentrations (5, 10, 20, 40, and 60 mM) at 100 mA and 80 °C. Error bars are standard error. Reprinted with permission from ref. 56. Copyright 2019 American Chemical Society.

and Gardew et al. The authors also proposed reactions for some compounds (see Fig. 12), this shows that hydroxyl groups, such as ketones, carboxylic acids and alcohols are resistant to hydrogenation, whereas double bonds were almost depleted. Gardew et al. also reported that, for phenolic compounds, the double bonds inside the aromatic ring were hydrogenated, when the hydroxyl group was not (e.g., phenol was converted into cyclohexanol and cyclohexanone rather than benzene). These data were extrapolated for the main components in plastic pyrolysis oil (see Fig. 15). It is worth noting that,

although cyclohexane is a possible product of phenol hydrogenation, the process of C-O cleavage, which would involve the hydrogenation of cyclohexanol into cyclohexene (C₆H₁₂O → C_6H_{10}), followed by cyclohexene into cyclohexane ($C_6H_{10} \rightarrow$ C₆H₁₂) would require much higher temperatures and overpotentials.62

Carbon loss during the ECH process was reported by Zhenglong et al., as only 80% of the carbon was retained in the cathode chamber. 6.0% was found in the anode chamber, due to migration through the membrane, 2.6% was adsorbed by the

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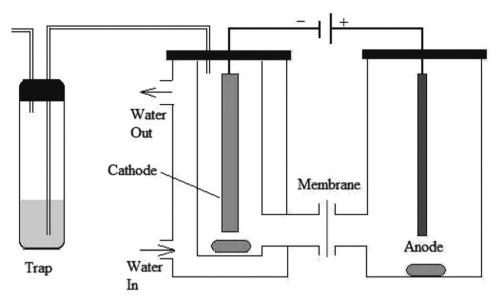


Fig. 11 Electrochemical hydrogenation cell setup for ECH of water-soluble bio-oil.⁷²

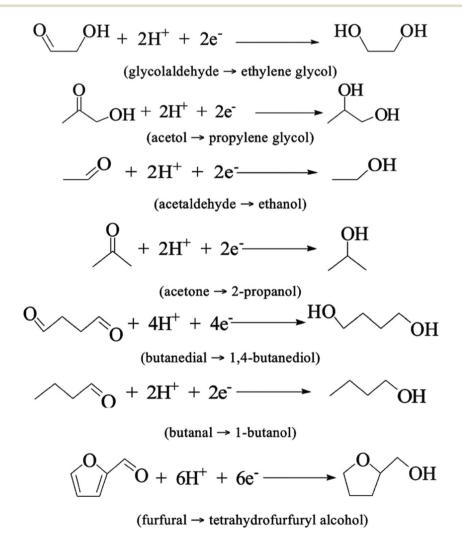


Fig. 12 Reductions via ECH of organic compounds in water-soluble bio-oil. 72

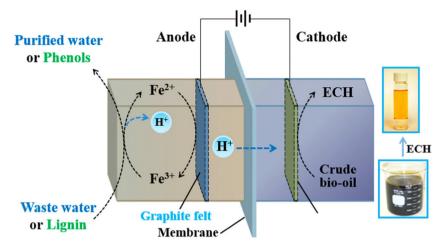


Fig. 13 Schematic illustration of low-energy ECH of bio-oil: experimental setup. Reprinted with permission from ref. 41. Copyright 2018 American Chemical Society.

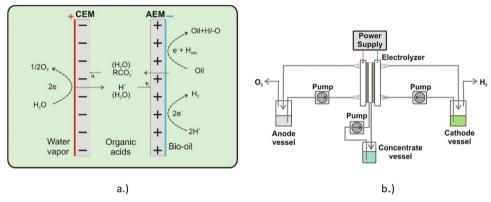


Fig. 14 Spatial chemical representation (a) and diagram of the separation—hydrogenation process (b). Reprinted with permission from ref. 40. Copyright 2018 American Chemical Society.

Main components from plastic pyrolysis oil and the possible products from ECH.

cathode and 0.2% was trapped in the downstream water trapping system. The system's total carbon loss was 11.2%. The

authors suggest that the rest could be inside the membrane, as it turned black after the process.⁷² This exemplifies how different membranes could lead to a better result, as the carbon loss could possibly be avoided. Also, the process cost is increased by constantly changing the system's components. Zhang $et\ al.$ showed that a cheaper anode, the Fe(III)/Fe(II) redox pair, can be utilized as a substitute for the commonly used noble metal (e.g. Pt) as a way to make the overall process cheaper and more competitive.

The work of Zhou *et al.* demonstrates how different catalyst cathodes can interfere with the final product, as the difference in FE and conversion rates were quite significant. The authors also proposed that the guaiacol upgrading was favoured by a chemisorption selectiveness to this compound. This indicates that different feedstocks need different catalytic cathodes to be more effective. Gardew *et al.* exemplified how substrate concentration can affect the FE and selectivity of the product, although it is not a direct relationship. Coke formation was not a major problem, although Zhenglong *et al.* reported finding decanted materials on the cathode chamber.

To avoid the problems related to the commonly used H-type cell, Lister *et al.* proposed a carbon catalyst cathode and DEM as a cheap and effective way to achieve upgrading. Results were promising, and the product had a higher economic value, as it could be used as commodity chemicals, but the authors found that further upgrading would be necessary for utilization as biofuel.

6 Plastic recycling

For the creation of a circular plastic economy, the plastic-oil needs to be upgraded into a valuable product, examples include biofuels and feedstock for industry (refined hydrocarbons, petrochemicals, or monomers). To achieve both the calorific values and chemical composition required, heteroatoms (O, S, N, *etc.*) should be removed. Some other elements that are present in the oil were listed in previous sections.

One area where a process of generating feedstock could be helpful includes plastic recycling, where the waste product can be returned to oil and then upgraded. Taking into consideration the highly variable nature of real plastic waste, Kusenberg *et al.* analysed municipal plastic waste and concluded that, to achieve industrial standards, the nitrogen concentration must be reduced by 87.7%, and oxygen by 98.0% (from 825 to 100 and 5200 to 100 ppmw respectively).⁷³

Therefore, the focus of the upgrading process is the oxygenated compounds, that is, reducing the O:C ratio of the oil. Oxygenated compounds can lead to lower stability and heating values, as well as a more corrosive product, due to carboxylic acids. Most of the oxygen content comes from PET but to accomplish an effective plastic waste recycling process, PET cannot be excluded. Benzoic acid and its derivatives are the most common carboxylic acids in plastic-oil, and therefore should be the focus of the ECH process.

6.1 Selecting a catalytic cathode

Catalysts can be very selective over organic functional groups, and this presents a challenge for the ECH of whole pyrolysis oil as it is a complex mixture (more than 300 compounds), therefore the cathode needs to be more versatile than the ones chosen for model compounds and be capable of hydrogenating various functional groups. Recent reports shows that catalysts like alloy and carbon-metal hybrids are possible solutions.⁷⁴

Although carboxylic acids are resistant to hydrogenation, ruthenium shows high activity for this process. ⁷⁵ As shown in the previous section, Ru was also the chosen metal as the cathode for Zhenglong *et al.* (Ru/ACC), Zhang *et al.* (Ru nanoparticles on mesoporous carbon) and Gardew *et al.* (Ru/ACC), which proves its effectiveness in hydrogenating WSBO and phenolic compounds, setting it as a prime candidate for the hydrogenation of whole plastic-oil. The possible mechanism for the hydrogenation of carboxylic acids is shown in Fig. 16.

The hydrogenation of toluene into methylcyclohexane was carried out in a proton exchange membrane (PEM) system by Fukazawa et al. and found that PtRu/C was a better catalyst than individual metals such as Pt and Ru and presented high FE (>90%).⁵³ In a later study, the same group reported on the ECH of benzoic acids into cyclohexane acids (PtRu/C catalyst) and presented high FE (99%) at a current density of 1.5 mA cm $^{-2}$. The authors also reported that in less mild conditions, the carboxyl group could also be hydrogenated (see Fig. 12). As mentioned in previous sections, Du et al. also reported on the activity of different cathodes and found that Pt₃Ru/ACC led to a higher conversion rate than Pt and Ru separately. The authors suggested that this could be the result of a synergistic effect between the two metals, combining platinum's excellent hydrogenation potency and the predominant direct hydrogenolysis ability of ruthenium.66 In addition, Salakhum et al. suggested that combining both metals can facilitate the cleavage of C-O and O-H bonds, as it reduces their heterolytic activation energies.⁷⁶ This indicates that PtRu/ ACC is a prime candidate for the ECH of carboxylic acids, and therefore, a possible choice of cathode for the hydrogenation of plastic-oil.⁵²

6.2 Conductivity challenge

The pyrolysis oil has low conductivity values, which is a challenge for an electrochemical process. To circumvent this, an electrolyte is used, as a way of increasing the conductivity. However, the commonly used water-based electrolytes (acid/water; alkali/water and salt/water) are unlikely to be suited for pyrolysis oil treatment, due to the large number of water-insoluble compounds. The removal of both water and the electrolyte (*e.g.*, HCl) after the treatment is also necessary, as it lowers the quality of the product. This would increase the costs and the complexity of the process and could render it unviable.⁷⁴

Possible solutions consist in either increasing oil conductivity *via* additives (*e.g.*, tetrabutylammonium hexafluorophosphate (NBu₄PF₆); LiCl; ammonium carbonate) or using a solvent that is capable of dissolving both polar and nonpolar compounds (*e.g.*, ionic liquids, DES (Deep Eutectic Solvents); biomass-derived organic solvents). Studies are necessary to understand the changes in the physicochemical properties of

Fig. 16 Mechanism for the hydrogenation of carboxylic acids. Edited with permission from ref. 75. Copyright 2020 Wiley

the oil and its behavior during ECH that an additive could bring. LiCl could increase the corrosivity of the product as Li⁺ and Cl⁻ are prone to corrosion. Ammonium carbonate could lead to membrane clogging via carbonate deposition, as well as increase in the nitrogen composition of the oil, and therefore, its corrosivity. NBu₄PF₆ is a chemically inert powder that is commercially available and could provide the much-needed conductivity for the system.

6.2.1 Ionic liquid and DES. Ionic liquids and DESs are quite similar in their uses and could serve the same purpose and although DESs, especially natural deep eutectic solvents (NADES), are considered greener and cheaper, it is a very recent class of mixtures and still needs time to develop. Some of its uses are on metallurgy and electrodeposition, due to its high electrical conductivity and solubility, NADES have also been utilized as precursor for biomass processing into more valuable chemicals.77-79

6.2.2 Biomass-derived organic solvents. Biomass-derived organic solvents are emerging and very promising alternatives as green solvents, that are produced from renewable sources and are non-hazardous to the environment. From this class of solvents, one of the most prominent is 2-methyltetrahydrofuran (2-MeTHF), which has levulinic acid and furfural, two of biooil's main components, as its precursors.80 That sets both NADES and biomass-derived organic solvents as candidates for alternative solvents for ECH of pyrolysis oils⁷⁴ (Fig. 17).

6.3 Plastic oil utilization

6.3.1 Biofuel. Although not an ideal solution, giving plastic waste a second life usage would be immensely beneficial for the environment as the utilization of the upgraded plastic oil as biofuel could lower the world's consumption of petroleum. For a business-as-usual approach, global plastic waste generation will increase from 260 million tons in 2018 to 460 million tons in 2030. For the same time frame, if the amount of plastic waste collected for recycling achieves 50%, chemical recycling could be responsible for 13% of all plastic production (considering a "high adoption" scenario and oil barrel price around \$75 dollars).5 This means that 59.8 million tons of plastics would go through pyrolysis, producing approximately 39.0 million tons of plastic oil (65.28 wt% average yield from several plastic pyrolysis reactors as reported by Siu Hua Chang in 2023).8

Fig. 17 2-MeTHF production via renewable sources. Edited with permission from ref. 80. Copyright 2012 ChemSusChem.

Biofuel energy production

Total biofuel production is measured in terawatt-hours (TWh) per year. Biofuel production includes both bioethanol and biodiesel.

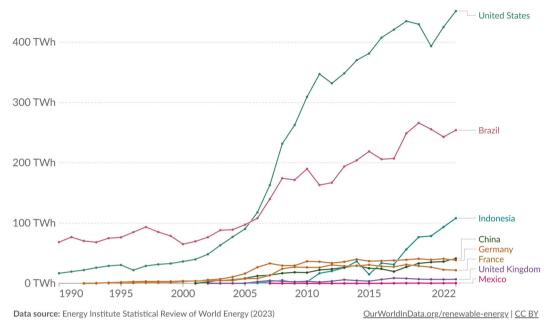


Fig. 18 Total biofuel production in terawatt-hours (TW h) per year. Biofuel production includes both bioethanol and biodiesel. Chart constructed by Our world in data.83

As the ECH of plastic oil still needs to be tested, the following calculations will be done using the heating value of the pretreatment plastic oil (41-47 MJ kg⁻¹, averaged to 44 MJ kg⁻¹).⁸¹ This means that approximately 476.67 TW h (terawatt-hours) per year of biofuel (Fig. 18) could be produced via ECH, which is more than the combined production of the whole of Asia, Europe, and Africa in 2021 (aprox. 380 TW h) and higher than the world's largest producer, the United States (399 TW h).

6.3.2 Plastic production. As stated in the previous sections, for the creation of a circular economy, plastic waste needs to be recycled into feedstock for plastic production. In 2019, 368 million tonnes of plastics were produced worldwide82 16% of the waste was collected for recycling and, after accounting for losses, 12% went through mechanical recycling. Landfills were the destination for 40%, and incineration for 25%; 19% of all plastic waste produced was mismanaged.⁵ This means that, from all the plastic waste, 65% of it, 239.2 million tonnes, was collected but not recycled. If this share went through chemical recycling, 156.15 million tonnes of plastic oil would be produced (65.28 wt% average yield).8 As the ECH of plastic oil still needs to be tested, three scenarios of conversion rates will be considered: low, fair, and high conversion (50, 75 and 90% respectively). For low conversion rates, 78.08 million tonnes of plastic production feedstock would be produced. Fair and high conversion rates would lead to 117.11 and 140.53 million tonnes respectively. This means that, at the current technology state, up to 38.18% of all resin could come from a sustainable source.

7 Conclusions

Fast pyrolysis is a thermochemical process that can be used to produce oil from both biomass and plastic. This material is a complex mixture containing acids, alcohols, aldehydes, esters, phenols, guaiacols, and a great number of other organic compounds. As it has high water content, low C: O ratios, high viscosity, and low density it needs to be upgraded for widespread usage. The electrochemical process for bio-oil upgradation is still new and needs further study to be fully optimized. But previous studies show that it can be done under mild conditions of temperature and pressure. ECH can be done utilizing low-cost materials and can achieve high FE and conversion rates. ECH shows an exciting path for the upgradation process of pyrolysis oils from chemical recycling of both biomass and plastics, a low cost, mild hydrogenation process that can help to achieve the necessary circular and greener economy. The main takeaways from this review are

- Noble metals are reactive species for the ECH of bio-oils. Rh, Pt and Ru loaded carbon cloth have all been reported to hydrogenate alcohols, carboxylic acids, olefins, and aromatic compounds, and presented high efficiency and FE.
- Current, potential, pH, and temperature are all essential factors in the ECH process. The fine tuning of these variables is necessary and directly linked to the quality of the final product.
- Conductivity of the oil presents a major challenge for ECH, alternative electrolytes and additives might be required to mke the process viable.

• The hydrogenation of plastic-oils is a possibility that still needs testing, although, as this technology matures, it may become one alternative for both the production of green plastics and biofuels.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors wish to thank UK Engineering and Physical Sciences Research Council (EPSRC) for supporting the work published in the paper through an EPSRC Doctoral Training Partnership Funding.

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