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Zhang, Y., Nielsen, J., Liu, Z. (2021). Yeast based biorefineries for oleochemical production. Current Opinion in Biotechnology, 67: 26-34. http://dx.doi.org/10.1016/j.copbio.2020.11.009

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# Biotechnology

### **Yeast based biorefineries for oleochemical production** Yiming Zhang<sup>1</sup>, Jens Nielsen<sup>1,2,3</sup> and Zihe Liu<sup>1</sup>



Biosynthesis of oleochemicals enables sustainable production of natural and unnatural alternatives from renewable feedstocks. Yeast cell factories have been extensively studied and engineered to produce a variety of oleochemicals, focusing on both central carbon metabolism and lipid metabolism. Here, we review recent progress towards oleochemical synthesis in yeast based biorefineries, as well as utilization of alternative renewable feedstocks, such as xylose and L-arabinose. We also review recent studies of C1 compound utilization or coutilization and discuss how these studies can lead to third generation yeast based biorefineries for oleochemical production.

#### Addresses

<sup>1</sup> Beijing Advanced Innovation Center for Soft Matter Science and Engineering, College of Life Science and Technology, Beijing University of Chemical Technology, Beijing, 100029, China

<sup>2</sup> Department of Biology and Biological Engineering, Chalmers University of Technology, Gothenburg, Sweden

<sup>3</sup> BioInnovation Institute, Ole Maaløes Vej 3, DK2200 Copenhagen N, Denmark

Corresponding authors:

Nielsen, Jens (nielsenj@chalmers.se), Liu, Zihe (zihe@mail.buct.edu.cn)

#### Current Opinion in Biotechnology 2021, 67:26–34

This review comes from a themed issue on Energy biotechnology

Edited by Huimin Zhao and Yasuo Yoshikuni

#### https://doi.org/10.1016/j.copbio.2020.11.009

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#### Introduction

Fatty acids and their derivatives, also termed oleochemicals, are used as drop-in fuels, lubricants, additives in foods, polymers, cosmetics, and pharmaceuticals, ranging from bulk chemicals to fine chemicals. Traditional production of oleochemicals relies on feedstocks of vegetable oils and animal fats, but microbial synthesis offers sustainable production from a wide range of renewable sources [1]. Besides, it allows for production of nonnatural oleochemicals having novel chemical features. Oleochemical synthesis has been well studied and intensively engineered in many microorganisms, including bacterial, yeast and microalgae [2]. This review will only focus on yeast based production of oleochemicals, whereas we refer to other recent reviews on oleochemical production from other microbial cell factories [3,4]. Oleaginous veast hosts, such as Yarrowia lipolytica, Rhodosporidium toruloides and Cryptococcus curvatus, are often chosen as industrial workhouses for oleochemical production, because of their efficient lipid synthesis and accumulation [5–9]. Non-oleaginous yeasts, such as *Saccharomyces cerevi*siae, is also preferred because of its well-studied cell metabolism and strain engineering tools. So far, a variety of oleochemicals have been synthesized in yeast, including free fatty acids [10], fatty alcohols, triacylglycerols (TAGs) [11], alkanes, alkenes [12], fatty acid methyl-esters (FAMEs), fatty acid ethyl-esters (FAEEs) [13], oleoylethanolamide [14] and wax esters [15,16]. Engineering efforts on deregulating lipid metabolism, redirecting carbon flux towards lipid synthesis and rebalancing cytosolic redox factors have been performed to increase the titre, rate and yield of oleochemicals. Meanwhile, discovering enzymes that are capable of synthesizing unnatural or natural oleochemicals with controlled chain length has gained more attention [17,18<sup>••</sup>,19].

Yeast based biorefineries can rely on using different renewable feedstocks, for example, glucose from food crops (1st-generation feedstock), xvlose and L-arabinose from plant biomass and energy crops (2nd-generation feedstock), and possibly C1 compounds like CO<sub>2</sub>, methane or methanol (3rd-generation feedstock) [20]. Glucose is the preferred carbon source for yeast, but use of 1stgenetation feedstock for oleochemical production threatens food supply. Utilization of xylose, the second most abundant monosaccharide in lignocellulose after glucose, has been studied for years to avoid food competition [21]. Utilization of L-arabinose from 2nd generation feedstock has also been studied for oleochemical production [1,22<sup>•</sup>]. Recently, utilization of 3rd-generation feedstock has started to be considered even though it is a challenging alternative for microbial synthesis and oleochemical production [23]. This review highlights recent advances in yeast based biorefineries towards production of fatty acids and their derivatives and end with prospects of utilization C1 compounds for microbial synthesis of oleochemicals (Figure 1).

#### Engineering on synthesis of oleochemicals

Feedstocks can be assimilated into the central carbon metabolism, including Embden–Meyerhof–Parnas (EMP) pathway, pentose phosphate (PP) pathway and tricarboxylic acid (TCA) cycle, to provide the building block acetyl-CoA and redox factor NADPH for fatty acid synthesis. Acetyl-CoA is converted to malonyl-CoA by acetyl-CoA carboxylase (ACC), which is directly used for fatty acid synthesis and elongation catalysed by fatty acid synthase in a cyclic manner. Engineering for



Sustainable feedstocks are utilized in yeast-based biorefineries for production of various oleochemicals. Yeast cell factories are exploited and optimized for oleochemical production via the Design-Build-Test cycle pipeline.

oleochemical production has been comprehensively reviewed in [5,24,25], and we therefore here focus on recent key points of particular interests for oleaginous and non-oleaginous yeasts, as summarized in Table 1.

#### Acetyl-CoA pool and NADPH supply

Generation of cytosolic acetyl-CoA varies between oleaginous and non-oleaginous yeasts. In non-oleaginous yeasts, cytosolic acetyl-CoA is synthesized via the pyruvate dehydrogenase bypass, which is a branch of the fermentative pathway with strong competition with ethanol pathway. In oleaginous yeast, cytosolic acetyl-CoA is shuttled from the mitochondria acetyl-CoA pool via a citrate-malate antiport cycle, and under nitrogen starvation TCA activity decreases and excess citrate is shuttled into the cytosol, which is converted to acetyl-CoA via ATP citrate lyase. Therefore, in oleaginous yeasts under nitrogen starvation high flux through the acetyl-CoA precursor allows for high lipid accumulation and advances oleochemical production, compared with non-oleaginous yeasts. When similar pathways were engineered into S. cerevisiae, that is, construction of acetyl-CoA shuttling cycle and deletion of fermentative pathway, cell metabolism was reported to convert from alcoholic fermentation to lipogenesis [26<sup>••</sup>]. Moreover, increasing the acetyl-CoA pools of oleaginous yeasts further enhanced lipid production under conditions with both high and low C/N ratios, confirming that efficient supply of cytosolic acetyl-CoA is required for high-level lipid synthesis [27–29,30<sup>•</sup>].

Cytosolic NADPH is either predominantly generated from the oxidative PP pathway in most yeasts, or primarily through cytosolic malic enzymes (MEs) in some oleaginous yeasts [25]. Increasing NADPH availability has successfully promoted fatty acid synthesis in both *S. cerevisiae* and *Y. lipolytica* via increasing PP pathway flux, decreasing glycolysis flux, expressing NADPH<sup>+</sup>-dependent glyceraldehyde 3-phosphate dehydrogenase or malic enzyme, or engineering oxidative defense pathways [26<sup>••</sup>,31<sup>••</sup>,32].

#### Fatty acid synthetase

Yeast fatty acid synthetases (FASs), Type I FASs, are large proteins with multiple modules of acetyl transferase (AT), malonyl-palmitoyl transferase (MPT), keto synthase (KS), ketoacyl reductase (KS), dehydratase (DH), enoyl reductase (ER), enoyl reductase (ER) and acyl carrier protein (ACP) [33]. In yeasts, FASs catalyze the synthesis of a range of acyl-ACPs (mainly C16 and C18) from acetyl-CoA and malonyl-CoA, and it is suggested that the chain length of

#### Figure 1

|--|

Feedstocks	Products	Host	Goals and genetic modifications	Titer, yield, productivity	Reference
Glucose	<b>F</b>		<b>Constructing ACL route</b> AnACL↑, MmusACL↑, RtME↑, MDH3'↑, CTP1↑		10.0 <b>0</b>
	Fatty acids	Sc	<b>Enhancing fatty acid synthesis</b> Δ <i>pox1</i> , Δ <i>faa1</i> , Δ <i>faa4</i> , FAS↑ <b>Optimizing central carbon flux</b> PGI1↓, IDH2↓	FB 33.4 g/L, 0.1 g/g	[26**]
Glucose	Fatty alcohols	ΥI	<b>Enabling oleochemical synthesis</b> MmarCAR↑, BsSfp↑, EcAHR↑, EcFadD↑	SF, 205.4 mg/L B, 2.15 g/L	[30 <b>°</b> ]
Glucose	Fatty alcohols	Sc	Increasing fatty acid flux: AnACL <sup>↑</sup> , MmusACL <sup>↑</sup> , RtME <sup>↑</sup> . MDH3' <sup>↑</sup> , CTP1 <sup>↑</sup> , Δpox1, FAS <sup>*</sup> <sup>↑</sup> Enabling oleochemical synthesis Δhfd1, Δtoo1, MmarCAR <sup>*</sup> <sup>↑</sup>	SF, 252 mg/L	[19]
Glucose	Alkanes/ alkenes	Sc	Targeting peroxisomes for oleochemical production Per- MmarCAR1, Per-SeADO1, Per-SeFd/FNR1	SF: 3.55 mg/L	[59]
Glucose	Alkanes/ alkenes	ΥI	Targeting ER for oleochemical production ER-AbFAR↑, ER- PmADO↑	SF: 23.3 mg/L	[30 <b>°</b> ]
Glucose	Lipids	ΥI	Increasing fatty acid flux, ACC↑, DGA↑ Increasing NADPH supply, CaGapC↑, McMCE2↑	FB, 98.9 g/L, 0.269 g/ g, 1.3 g/L/h	[31**]
Xylose	Fatty alcohols	Sc	Increasing fatty acid flux ACC↑, FAS↑, Δrpd3 Enabling oleochemical synthesis TaFAR↑ Enhancing xylose utilization: CsXR↑, CtXDH↑, PpXK↑	B, 1.2 g/L, 0,1 g/g, 0.02 g/L/h	[40]
Xylose	Lipids	ΥI	Increasing fatty acid flux Δpox1-6, Δtgl4, GDP1↑, DGA2↑, Enhancing xylose utilization SsXR↑, SsXDH↑, YIXK↑	FB, 20.1 g/L, 0.08 g/g, 0.19 g/L/h	[44]
Xylose	Lipids	ΥI	Enhancing xylose utilization SsXR↑, SsXDH↑, SsXK↑ Starvation-enabled adaptive evolution	B, 15.0 g/L, 0.08 g/g, 0.19 g/L/h	[43]
Agave bagasse hydrolysate	Lipids	ΥI	Increasing fatty acid flux Δpox1-6, Δtg/4, GDP1↑, DGA2↑, Enhancing xylose utilization YIXR, YIXDH, YIXK, AnXPKA, AnACK	FB, 16.5 g/L, 0.34 g/g, 0.19 g/L/h	[45]

The listed examples report the highest titers for corresponding products within the reviewed publications. Symbols and prefixes: ↑, overexpression; ↓, downregulation;', truncated version; \*, mutated version. General abbreviations: SF, shake-flask; FB, fed-batch bioreactor; B, batch bioreactor. Species abbreviations: Ab, *Acinetobacter baylyi*; An, *Aspergillus nidulans*; Bs, *Bacillus subtilis*; Ca, *Clostridium acetobutylicum*; Cs, *Candida shehatae*; Ct, *Candida tropicalis*; Ec, *E. coli*; Mmar, *Mycobacterium marinum*; Mmus, *Mus musculus*; Mc, *Mucor circinelloides*; Pm, *Prochlorococcus marinus*; Pp, *Pichia pastoris*; Rt, *Rhodosporidium toruloides*; Sc, *Saccharomyces cerevisiae*; Se, *Synechococcus elongatus*; Ss, *Scheffersomyces stipites*; Ta, *Tyto alba*; YI, *Yarrowia lipolytica*. Gene/Enzyme abbreviations: ACC, acetyl-CoA carboxylase; ACK, acetate kinase; ADO, fatty-aldehyde deformylating oxygenase; AHR, aldehyde reductase; CAR, carboxylic acid reductase; DGA, diacylglycerol acyltransferase; FAR, fatty acyl-CoA-ACP or fatty acyl-ACP reductase; FAA, fatty acyl-CoA synthetase; FAS: fatty-acid synthase; Fd/FNR, ferredoxin and ferredoxin/NADP + reductase; GPD, glycerol-3-phosphate dehydrogenase; ME, malic enzyme; POX, fatty-acyl coenzyme A oxidase; Sfp, phosphopantetheinyl transferase; XR, xylose reductase; XDH, NADH-dependent xylitol dehydrogenase; XK, xylulokinase; XPKA, phosphoketolase.

oleochemicals are mainly controlled via FASs. Based on biochemical mechanism and structural analysis of FAS complexes, rational protein engineering on FASs from Y. *lipolytica, S. cerevisiae, R. toruloides, Mycobacterium vaccae* and *Aplanochytrium kerguelense* enabled production of short/ medium chain fatty acids (S/MCFAs, C6-C12) in S. cerevisiae [18<sup>••</sup>,34,35], with several mutations introduced into KS, AT and MPT or heterologous thioesterase (TE) domain introduced to FAS. Engineered fungal and bacterial FAS with mutated KS and TE domains were co-expressed in a strain with enhanced MCFA tolerance and optimized carbon flux redirection, resulting in production of  $2.87 \pm 0.06$  g L<sup>-1</sup> MCFAs in fed-batch cultivations [18<sup>••</sup>].

Moreover, heterologous FASs from *R. toruloides* and *M. vaccae* have been overexpressed in *S. cerevisiae* to produce very long chain fatty acids and fatty alcohols [36], or enhance fatty acid synthesis  $[26^{\bullet\bullet}]$ .

#### Lipid metabolism

Fatty acyl-ACPs synthesized from FAS are esterified to form three major lipid classes, including triacylglycerols

(TAGs), sterol esters (SEs) and phospholipids (PLs), or released as free fatty acids (FFAs), or degraded to acetyl-CoA via β-oxidation, as shown in Figure 2. Lipid metabolism is tightly regulated in yeast via several regulatory nodes [5,37<sup>•</sup>]. Strategies with blocking β-oxidation and downregulating fatty acid activation and storage lipid formation in S. cerevisiae resulted in accumulation of FFAs and PLs [37<sup>•</sup>], which ensured high flux into synthesis of downstream oleochemicals, including fatty alcohols, alkanes, alkenes and oleoylethanolamide (OEA) [14,21,26<sup>••</sup>]. In Y. lipolytica, blocking B-oxidation was also performed to enhance lipid synthesis, and in combination with expressing heterologous enzymes for fatty acid synthesis, fatty acyl-ACPs were redirected to synthesis FFAs instead of TAGs, as successfully demonstrated in production of omega-3-eicosapentaenoic acid (EPA) [25]. The interconversion between FFAs and different lipids pools offers varied opportunities for synthesis of various oleochemicals.

#### **Oleochemical tolerance and cell fitness**

High tolerance towards oleochemicals is required for efficient producing strains. Transporter engineering and





Yeast based biorefineries for oleochemical production utilizing 1st generation feedstock glucose. Carbon flux is redirected to fatty acid synthesis by engineering central carbon metabolism and lipid metabolism with upregulated (as shown in bold arrows) and downregulated (as shown in dashed grey arrows) reactions, and a variety of oleochemicals are synthesized by expression of heterologous pathways (as shown in blue arrows).

adaptive evolution are usually adopted for improving strain tolerance towards toxic biochemicals. For example, expression of a fatty acid transporter FATP1 from human in *S. cerevisiae* mediated fatty acid uptake and facilitated fatty alcohol export, leading to enhanced fatty alcohol production and improved cell growth [38]. Moreover, protein engineering on a membrane transporter Tpo1 involved in tolerance against C10 fatty acid enhanced MCFA production by 1.3 fold, while adaptative evolution enhanced MCFA production by 1.7 fold [18<sup>••</sup>]. Meanwhile, adaptive evolution was adopted in balancing oleochemical production of free fatty acids [26<sup>••</sup>].

# Alternative feedstocks for oleochemical production

Compared with 1st-generation biorefineries, efficient feedstock uptake and utilization is usually a major issue for 2nd-generation biorefineries, in particular as most yeasts do not naturally use pentoses and other components from biomass. There has therefore been much work on engineering yeasts for utilization of pentoses and more recently C1 feedstocks like methanol and carbon dioxide.

#### **Xylose utilization**

Xylose can be converted to xylulose-5-phosphate (Xu5P) via two different assimilation pathways. One is composed of NAD(P)H-dependent xylose reductase (XR), NADH-dependent xylitol dehydrogenase (XDH) and xylulokinase (XK), and the other is relying on xylose isomerase (XI) and XK. Xu5P is then channelled via the PP pathway or phosphoketolase (PK) pathway into central carbon metabolism for cell growth and biochemical synthesis (Figure 3).

Recently, engineering efforts have been performed in non-xylose metabolizing yeasts to enable oleochemical production [1,39]. For example, in a fatty alcohol producing strain of *S. cerevisiae*, expression of XR from *Candida shehatae*, XDH from *Candida tropicalis* and XDH from *Pichia pastoris* resulted in higher yields on xylose compared to glucose during batch and fed-batch cultivations [40].

In Y. lipolytica, the dormant pentose metabolism resulted in poor and unstable xylose utilization [41,42]. Expression of XR, XDH and XK from Scheffersomyces stipitis together with starvation adaptation enabled efficient xylose utilization and lipid production of 15 g/L [43]. In another study, expression of XR, XDH from S. stipitis and native XK in an engineered lipid producing strain resulted in lipid production of 20.1 g/L [44]. By overexpressing heterologous and native xylose utilizing enzymes, Y. lipolytica was reported to produce lipids with a high productivity (0.185 g/L/h) and a high yield (0.344 g lipids/g sugars) in xylose-rich agave bagasse hydrolysate [45]. A recent study demonstrated that Y. lipolytica could combine the xylose utilization phenotype with the metabolite overproduction phenotype via a mating approach, and allowed 1.42 g/L  $\alpha$ -linolenic acid production on xylose [46].

Oleaginous yeasts that can naturally grow on xylose appear to be promising candidate hosts for 2nd-generation biorefineries of oleochemical production, including R. toruloides [47] and C. curvatus [48]. However, limited knowledge of xylose metabolism and engineering tools have restricted their application in oleochemical production. A recent study on comparative analysis of R. toruloides revealed that cells grown on xylose achieved almost 50% lower growth rate and sugar consumption rate, lower final biomass yield whereas similar final cellular lipid content [49<sup>•</sup>]. Proteome analysis then identified a number of putative sugar transporters for xylose and glucose and suggested that xylose import might be the limiting step during xylose conversion into lipids. NADPH regeneration relied primarily on the PP pathway, and may also involve malic enzyme, alcohol dehydrogenases and aldehyde dehydrogenases. The PK pathway with higher efficiency and carbon conservation, however, seemed to have limited role in xylose conversion into lipids, possibly due to the inefficient xylose





Utilization pathways of pentoses and C1 compounds. Abbreviations used: XR, xylose reductase; XDH, xylitol dehydrogenase; XK, Xylulokinase; XI, xylose isomerase; AR, Arabinose reductase; LAD, L-arabinitol 4-dehydrogenase, LXR, L-xylulose reductase; AI, L-arabinose isomerase, RK, L-ribulokinase; RPE, L-ribulose-5-P-4-epimerase; DAS, dihydroxyacetone synthase; AOX, alcohol oxidase; HPS, 3-hexulose-6-phosphate synthase; PHI, 6-phospho-3-hexuloisomerase; Xu5P, xylulose-5-phosphate; Ru5P, ribulose-5-phosphate; GAP, glyceraldehde-3-phosphate; DHA, dihydroxyacetone.

uptake. These findings are valuable for developing lipid production processes on xylose-containing substrates and further optimization of xylose utilization.

Other alternative routes for xylose catabolism, such as the XI pathway, Dahms pathway, Weimberg pathway, and synthetic pathways, may also be useful for future optimization in oleochemical production [21,50]. Besides, engineering strategies to facilitate the conversion of xylose to ethanol in *S. cerevisiae*, like improving xylose uptake, balancing redox factors, as well as evolutionary engineering and transcriptional factor engineering, could also contribute to engineer yeast for oleochemical production from xylose [51,52].

#### L-arabinose utilization

L-arabinose is another abundant pentose in lignocellulose feedstock, and it can be assimilated into the central carbon metabolism via two different routes in prokaryotes and eukaryotes (Figure 3). The prokaryotic route involves arabinose isomerase (AI), L-ribulokinase (RK) and Lribulose-5-P4-epimerase (RPE), and the eukaryotic involves arabinose reductase (AR), L-arabitol dehydrogenase (LDH), L-xylulose reductase (LXR), XDH and XK [53].

Expression of heterologous utilization pathways with subsequent evolution engineering in *S. cerevisiae* enabled efficient assimilation of L-arabinose, as well as efficient co-fermentation of xylose and L-arabinose [54]. A recent study found that the underlying mechanism of fast co-

fermenting capacity of L-arabinose and xylose was a high number of copies of the L-arabinose utilization pathways, which will benefit oleochemical production [55]. In *Y. lipolytica*, the dormant L-arabinose assimilation pathway was identified through transcriptomic and metabolic analyses, and it can be activated by overexpression of pentose transporters, XDH and LAD, shedding light on oleochemical production on 2nd generation feedstock [22<sup>\*</sup>].

#### Methanol utilization

 $\rm CO_2$  and methanol are gaining increasing interests as 3rdgeneration feedstocks for bioproduction, because of their abundance in nature and cheap price, as well as the urgent need to reduce the threat of the global warming and human reliance on fossil fuels [20,56,57]. However, biochemical production with these C1 compounds is still challenging due to low efficiencies of utilization pathways and high demands of energy and reducing power. Currently, C1 compound utilization pathways are still under evaluation with and without utilization pathways of other feedstocks.

*Pichia pastoris* can grow on methanol as the sole carbon and energy source. During methanol cultivation, peroxisomes amplificate and dominate the cell volume of *P. pastoris* [58], which make it a promising methanol utilizing host for oleochemical production, as enhanced alkane titres were achieved when the biosynthetic pathway was targeted to the peroxisomes of *S. cerevisiae* [59]. In a recent study, *P. pastoris* can also be converted into an autotroph strain with  $CO_2$  as the carbon source and methanol as the energy source, by engineering the methanol assimilation pathway, xylose monophosphate (XuMp) cycle (Figure 3) or called as dihydroxyacetone (DHA) cycle, to a CO<sub>2</sub> fixation pathway [60<sup>••</sup>]. With the efficient CRISPR-Cas9 mediated genome editing toolkit developed in *Hansenula polymorpha* [61], this thermotolerant methylotrophic yeast could also be a promising chassis for oleochemical production utilizing methanol.

To direct utilize methanol in *S. cerevisiae*, the XuMP cycle from *P. pastoris* was reconstructed in *S. cerevisiae*, and the resulting strain could consume 1.04 g/L methanol, with slow growth and 0.26 g/L pyruvate produced [62]. Meanwhile, expression of the ribulose monophosphate (RuMP) cycle, another methanol assimilation pathway identified in methylotrophic prokaryotes (Figure 3), failed to result in methanol utilization for cell growth in both *S. cerevisiae* and *Y. lipolytica* [62,63]. Clearly, the poor cell growth and slow methanol utilization imposes requirement for more engineering work. This may require improved subcellular expression to reduce the toxicity of the intermediate formaldehyde or establish efficient regeneration of xylose-5-phosphate.

#### CO<sub>2</sub> utilization

The enzymes ribulose-1,5-bisphosphate carboxylase (RuBisCO) and phosphoribulokinase (PRK) from the Calvin cycle have been expressed in *S. cerevisiae*, and increased the ethanol yields on both glucose and xylose. Besides, the decreased accumulation of by-products glycerol and xylitol suggested that  $CO_2$  could be used as an external electron acceptor to balance cytosolic redox factors [64,65].

The synthetic reductive glycine pathway has also been demonstrated functional in *S. cerevisiae* by overexpressing endogenous enzymes to synthesis glycine from formate and CO<sub>2</sub>. The pathway with high activity, high affinity and tolerance of formate suggested *S. cerevisiae* might be especially suitable for formate utilization [66]. Recent studies found that CO<sub>2</sub> can be efficiently converted to formate with electrochemical and photochemical methods [67], and these findings may enable yeast based 3rd biorefineries for oleochemical production.

#### **Conclusion and perspectives**

Much progress has been achieved in both oleaginous yeasts and non-oleaginous yeasts utilizing glucose, including production improvements and oleochemical portfolio expansions. However, current titres, rates and yields of most bulk oleochemicals cannot meet the requirements of commercial production. A promising strain for commercial production is the engineered *Y*. *lipolytica*, which can produce FAMEs with a high titre, yield and rate of 98.9 g/L, 1.3 g/L/h and 0.27 g/g glucose, respectively [31<sup>••</sup>]. With omega-3 production in *Y. lipolytica* commercialized by DuPont [68], biosynthesis of

high-value oleochemicals have attracted more attention, and more enzymes and pathways for novel oleochemicals have been evaluated, including polyunsaturated fatty acids [10,69], flavour lactones [16], jojoba-like wax esters [15], and oleoylethanolamide [14].

Efficient xylose utilization would be a step further for commercial production of oleochemicals. An engineered Y. lipolytica has demonstrated its potential capacity for using xylose-rich agave bagasse hydrolysates as feedstock resulting in a high lipid yield of 0.344 g/g sugars, titre of 16.5 g/L and rate of 1.85 g/L/h [45]. Non-conventional yeasts, like R. toruloides and C. curvatus, could also be promising chassis for 2nd-generation oleochemical production. For example, C. curvatus could accumulate lipids up to 69.5% of dry cell weight when growing on aromatic substrates, representing one of the promising yeast cell factories for oleochemical production from depolymerized lignin [70]. Although limited genetic engineering tools restrict their usages, multi-omics analysis uncovers their metabolic capabilities of lipid synthesis and possible regulation mechanisms on different feedstocks, which can guide future strain performance on oleochemical production [71,72].

So far it has been demonstrated that S. cerevisiae can expand the range of oleochemicals it can produce, and it therefore represents a ready-to-use chassis for development of novel oleochemical synthesis. Y. lipolytica also seems to be an attractive chassis for both 1st generation and 2nd generation refineries for oleochemical production due to its higher flux from cytosolic acetyl-CoA towards fatty acid synthesis. The capacity may be further enhanced by introducing the Calvin cycle to utilize CO<sub>2</sub> as an electron acceptor, as previously used to enhance ethanol production and yield with high carbon-conversion and energy-conversion in S. cerevisiae [64,65]. Moreover, the PP pathway is highly involved with both the utilization pathways of xylose and C1 compounds, and the previous study found that overexpression of non-oxidative PP enzymes to ensure sufficient pool of ribulose-5-phosphate was required for implementation of the Calvin cycle [65]. Therefore, engineering of the PP pathway to balance the carbon and energy flux is desired for oleochemical production on 2nd-generation and 3rd generation feedstocks.

#### **Conflict of interest statement**

Nothing declared.

#### Acknowledgements

This work was supported by National Natural Science Foundation of China [grant no. 21908004, 2019; 21808008, 2018]; the Fundamental Research Funds for the Central Universities [grant no. buctrc201801, 2018], the Novo Nordisk Foundation [NNF10CC1016517], and the Knut and Alice Wallenberg Foundation.

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